

A Danish Diabetes Register

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Bendix Carstensen Steno Diabetes Center, Gentofte, Denmark
& Department of Biostatistics, University of Copenhagen
<bcar0029@regionh.dk> <b@bxc.dk>
<http://BendixCarstensen.com>

Contents

1	Background	1
1.1	NDR criteria	1
1.2	RUKS	1
1.3	A new register	2
1.3.1	Type of diabetes	3
1.3.2	GDM	3
1.3.3	PCOS	3
1.3.4	Summary	4
1.3.5	Differences to RUKS	4
2	Data acquisition	5
2.1	Diabetes patients	5
2.2	Population data	5
2.3	Follow-up	5
2.3.1	Analysis dataset	5
2.4	Prevalences	6
2.5	Datasets of follow-up and prevalence	6
2.5.1	Rate data	7
2.5.2	Prevalence data	8
2.5.3	Crude tables for teaching	9
3	Prevalence and occurrence rates	10
3.1	Prerequisites	10
3.2	Diabetes prevalences	11
3.2.1	Trends in prevalence	13
3.2.2	Prevalence age-period interaction	16
3.3	DM incidence	19
3.3.1	Age-Period-Cohort modelling	19
3.3.2	Time-trends in rates	20
3.3.3	Summary of the APC modelling	21
3.3.4	Prediction by age for different cohorts	21
3.3.5	Prediction by age across periods	23
3.3.6	Saving the fitted models	24
3.4	Mortality	28
3.4.1	Mortality in non-diabetics	28
3.4.2	Mortality among DM patients	30
3.5	Summary of APC models for incidence and mortality	35

3.5.1	Time-trends in mortality rates	35
4	Prediction of rates	38
4.1	Analysis date	38
4.1.1	Datasets for rate modeling	41
4.2	Models for incidence and mortality rates	41
4.3	Incidence rates	43
4.3.1	Incidence rate predictions	44
4.3.2	A damping extrapolation	44
4.4	Mortality rates	47
4.4.1	Diabetes patients	47
4.4.2	Persons without diabetes	49
4.5	Average trends	49
4.6	Trends in estimated rates	50
4.7	Extrapolation of rates	51
5	Components of prevalence	54
5.1	Transition probabilities	55
5.2	Prediction of the observed prevalences	57
5.2.1	Checking the prediction	59
5.3	How much is attributable to what?	63
5.4	Number of diabetes patients in Denmark	68
5.5	Timetrend in the components	72
6	Predicting prevalence of diabetes	76
6.1	Predicted rates	76
6.2	Transition probabilities	77
6.3	Prediction of the observed prevalences	79
6.4	The actual numbers of diabetes patients in Denmark	81
6.5	Population forecast from DST	82
6.6	Timetrend in prevalent number of DM patients	85
	References	94
7	SAS programs	95
7.1	Rationale and overview	95
7.2	Program execution	96
7.3	Program documentation	96
7.3.1	optslibs.sas	97
7.3.2	xgdm.sas	97
7.4	00-base	98
7.4.1	00-base.lst	98
7.5	01-npr	98
7.5.1	01-npr.lst	105
7.6	02-dvdd	114
7.6.1	02-dvdd.lst	118
7.7	03-ndr	123
7.7.1	03-ndr.lst	125
7.8	04-rmps	126

7.8.1	04-rmps.lst	131
7.9	05-diab	138
7.9.1	05-diab.lst	141
7.10	06-define	143
7.10.1	06-define.lst	147
7.11	06a-comp	161
7.11.1	06a-comp.lst	162
7.12	Tabulation of follow-up and prevalence	165
7.13	08-mkFU	165
7.13.1	08-mkFU.lst	169
7.14	09-mkPr	171
7.14.1	09-mkPr.lst	173

Chapter 1

Background

The maintenance of the National Diabetes Register (NDR) has been discontinued by the Health Data Authority (Sundhedsdatastyrelsen). It is being replaced by the Register of Selected Chronic Diseases (RUKS) which however does not encompass precisely the same persons.

1.1 NDR criteria

The “old” NDR, established 2006, covering the period from 1995 (in terms of incidence) was based on the following criteria [2]:

lpr: recording of diabetes as diagnosis in the NPR

fodt: use of the service “foot-therapy for diabetes patients” in the National Health Services Register (NHSR).

b15i1: the date of the 5th blood glucose measurement within a period of one year in the NHSR.

b12i5: two measurements of blood glucose per year in 5 consecutive years. The date is defined as the 2nd blood glucose measurement within the 5th period of one year.

oad: date of 2nd purchase of OAD as recorded in the Register of Medicines Products Statistics (RMPS).

ins: date of 2nd purchase of insulin as recorded in the RMPS.

The inclusion date was the earliest of the dates where any of these 6 criteria were met.

It has been pointed out that the two blood-glucose (purely *procedural*) criteria included many persons that were unlikely to be diabetic patients, notably women only being *tested* for gestational diabetes (GDM) [3].

1.2 RUKS

The alleged replacement of the NDR is the Register of Selected Chronic Diseases (Register over Udvalgte Kroniske Sygdomme, RUKS). Among the 8 diseases selected for the register, are T1 diabetes and T2 diabetes.

From the register description, the RUKS seems to be a register of repeated prevalences, in the sense that the register has a separate version as of each 1 January (?).

- Type 2 DM:
 - Persons recorded with ICD10 code E11 in NPR, as the latest diagnosis.
Persons are not included on the basis of a single NPR contact with code E11, at least one more contact (E10 or E11) or purchase of OAD or insulin is required.
 - Persons who have purchased OADs (A10B from the RMPS), and at least two purchases of either A10A (insulins) or A10B (other antidiabetic drugs).
 - Women who have a diagnosis of PCOS or have only purchased metformin (and no other OADs or insulin) and have purchased either clomifen (G03GB02) or estrogen (G03HB) are *excluded*.
 - Persons who have had no diabetes recordings in NPR or RMPS during the last 10 years are *excluded*.
 - The term “latest” for the NPR criterion and the exclusion referring to “last 10 years” seems to indicate that the register is defined relative to a particular creation date for the register, although this is not explicitly stated.
- Type 1 diabetes:
 - Persons recorded with ICD10 code E10 in NPR, as the latest diagnosis.
Persons are not included on the basis of NPR contacts with code E11, at least one purchase of insulins is required
 - Persons who have purchased insulins (A10A from the RMPS), and at least two purchases of either A10A (insulins) or A10B (OADs).
 - Persons already classified as T2 above are *excluded*.
 - Women with a diagnosis of GDM (ICD10 024.4) and only have purchased anti diabetic medication in a window from 280 days before the first till 280 days after the last recording of GDM are excluded.

1.3 A new register

The following is an attempt to reconstruct the NDR, using (almost) the same criteria as in the original NDR, with an additional effort to define persons as either T1 or T2.

The basic content of the register is one record per person with sex, type of diabetes and dates of birth, DM and death. Additionally, the register will have the dates for meeting each of the defining criteria (name of the date in the register):

doOAD Date of first recorded purchase of OAD (A10B).

doIns Date of first recorded purchase of insulin (A10A).

doNPR Date of the first recorded contact date with a diagnosis of diabetes in the NPR.

doDVD Recorded date of diagnosis in the Danish adult diabetes register (DADD). If none recorded the date of reporting is used.

doDiab Recorded date of eye-screening in the Danish eye-screening database for diabetes patients.

1.3.1 Type of diabetes

The classification of patients as T1 or T2 based on register date only is not accurate, and the approach chosen here is therefore to identify persons with T1D with reasonably high specificity, and classify the rest as T2D. Thus T2D will be equivalent to “cannot be classified as T1D with reasonable certainty”, and hence the classification should be used cautiously; the persons recorded as T1D are very likely to be T1D, but there is some under-reporting. Consequently, some T1D patients that are erroneously classified as T2D, but the precise size of this problem is unknown.

The practical implementation of the type classification is:

- use the DADD recordings of diabetes type (T1,T2,Other,Unkn) and classify persons as T1 resp. T2 if more than half of the recordings are T1 resp. T2. The rest are left unclassified.
- use the NPR to classify persons as T1 resp. T2 if more than half of the recordings are T1 (E10) resp. T2 (E11). The other codes (E12-E13) are ignored, and thus some persons are left unclassified.
- The classification from DADD is always used. If a person is not classified or does not appear in DADD, the classification from NPR is used.
- Finally, persons that have purchased OAD before age 15 or insulin before age 30 are classified as T1 (except if classified as T2 in DADD); all other are classified as T2.

1.3.2 GDM

If a person is recorded with a diagnosis of GDM in the NPR, the person cannot enter the register on any criterion during the next 365 days. To account for registration delays the window starts 30 days prior to the date of GDM.

GDM diagnoses that are less than 200 days from the previous one are disregarded; so from the earliest GDM for a given person, no GDM diagnosis in the next 200 days is counted, from the next after this, another window of 200 days is used etc.

1.3.3 PCOS

If a person is recorded with PCOS in the NPR, this person cannot be included on the basis of metformin purchase in the period from the earliest PCOS diagnosis until the person's 45th birthday.

Moreover, if a person's only drug dispensations are metformin between age 20 and 45, the person is considered a possible PCOS case and these dispensations are not counted. Dispensations after age 45 for women with a PCOS diagnosis are considered as diabetes medication and lead to inclusion in the register at the first date of purchase after the 45th birthday.

1.3.4 Summary

The register we attempt to build is a register where persons are captured by a set of criteria and then kept in the register with this inclusion date. In principle we might use the health registers to define an exit date as well (for example 10 years), but we have chosen to let prospective users define this type of dates themselves.

1.3.5 Differences to RUKS

The proposed approach differs from RUKS in the following points:

- Persons may be included in our register even if only one purchase of OAD/Insulin is recorded.
- Only NPR diagnosis of PCOS is used, and persons deemed to suffer from PCOS can actually be included at a time after age 40 based in metformin purchase alone.
It is not entirely clear whether persons who meet the criteria for PCOS before age 40 and only meet other criteria after age 40 are included in RUKS with a date of inclusion equal to the first purchase of metformin.
- RUKS maintains a window no inclusion of 280 days *before* date of GDM till 280 days after. Our approach only use a window of 30 days before (to account for registration delays) and of 365 days after the date of GDM.
- The RUKS approach to definition T1/T2 is based on recordings in the NPR and classify persons as T1/T2 according to the most frequently occurring of E10 and E11, whereas our approach only classify persons if more than half of the recordings are E10 resp E11, also taking other codes into account. It seems that RUKS do not include the code E12–E14 as diabetes.
- It appears as if RUKS is a prevalence register, where persons are included with reference to a given date. RUKS seems not to be available as a research register — which is quite logical for a register that have different persons recorded for different reference dates.

Chapter 2

Data acquisition

2.1 Diabetes patients

We have reconstructed the Danish Diabetes Register based on information from the National Patient Register, the Prescription Register (officially: Register of Medicines Products Statistics) and the Danish Adult Diabetes Database (DADD) and the eye examination databas (Diabase). The register contains id, date of birth, date of DM and date of death (among other things).

In the register we have defined T1 diabetes as those persons we are reasonably certain to be T1 patients namely those under 15 at first purchase of any anti-diabetic drugs, those under 30 at first purchase of insulin, and those recorded as T1 patients in the DADD or in NPR (the latter excluding those recorded as T2 in DADD).

2.2 Population data

We have had access to records for the entire population containing date of birth, death, emigration and immigration. From this we constructed a dataset with one record per period spent in Denmark; a new record is started whenever a person enters the study population (by birth, immigration or crossing 1995-01-01 alive). The follow-up represented by this record is terminated by emigration death or the end of the study period, currently 2015-12-31.

2.3 Follow-up

We used the Diabetes Register to subdivide the population records of follow-up by state of follow up in “noDM”, “T1” and “T2”. Also the register information was used to count the number of DM-events; that is diagnoses of T1 resp. T2 and deaths.

2.3.1 Analysis dataset

The constructed dataset or follow-up records was further subdivided by current age and calendar time in 1-year intervals, and the resulting dataset was tabulated by sex, state, age and calendar time of follow-up and date of birth; the latter three in 1-year categories. Each entry in the tabulation contains the person-years at risk, the number of deaths and the

number of diagnoses of T1 and T2, respectively. Obviously, the number of diagnoses of T1 and T2 in the states “T1” and “T2” is 0.

This dataset allows us to model incidence rates of T1 and T2 diabetes as well as mortality rates separately for the three states. In this report we will pool the two types of diabetes and thus base predictions etc. on incidence rates of any type of DM and mortality rates among persons.

2.4 Prevalences

Further when we make projections we shall also need the prevalence of DM at different dates (1 January each year 1996–2016). These numbers were also constructed from the follow-up dataset, by simply extracting those that were alive at the dates and classify these a being either “noDM”, “T1” or “T2”.

The construction of the follow-up and prevalence data from the register is documented in the SAS-programs 08-mkFU (p. 165 ff.) and 09-mkPr (p. 171 ff.).

2.5 Datasets of follow-up and prevalence

First load the relevant package:

```
> options( width=95 )
> library( Epi )
> library( splines )
> library( haven )
> print( sessionInfo(), l=F )
R version 3.3.2 (2016-10-31)
Platform: x86_64-w64-mingw32/x64 (64-bit)
Running under: Windows Server 2012 R2 x64 (build 9600)

attached base packages:
[1] splines  stats      graphics  grDevices  utils      datasets  methods   base

other attached packages:
[1] haven_1.0.0 Epi_2.7

loaded via a namespace (and not attached):
 [1] cmprsk_2.2-7      MASS_7.3-45      assertthat_0.1   Matrix_1.2-7.1   plyr_1.8.4
 [6] parallel_3.3.2   tools_3.3.2      survival_2.40-1  tibble_1.2       etm_0.6-2
[11] Rcpp_0.12.8      grid_3.3.2       numDeriv_2016.8-1 lattice_0.20-34
```

We first load the follow-up dataset and then the prevalence dataset:

```
> FU <- read_sas( "../data/futab.sas7bdat" )
> str( FU )
Classes 'tbl_df', 'tbl' and 'data.frame':      25723 obs. of  10 variables:
 $ sex  : num  1 1 1 1 1 1 1 1 1 1 ...
 $ state: chr  "T1" "T1" "T1" "T1" ...
 $ A    : atomic 0 0 0 0 0 0 0 0 0 0 ...
 ..- attr(*, "label")= chr "Age at FU"
 $ P    : atomic 1995 1996 1997 1997 1998 ...
 ..- attr(*, "label")= chr "Date of FU"
 $ C    : num  1995 1995 1996 1997 1997 ...
 $ D    : num  0 0 0 0 0 0 0 0 0 0 ...
```

```

$ T1 : num 0 0 0 0 0 0 0 0 0 0 ...
$ T2 : num 0 0 0 0 0 0 0 0 0 0 ...
$ Y : num 0.001563 0.000476 0.00017 0.000461 0.000517 ...
$ U : num 0 1 1 0 1 0 1 0 1 0 ...
- attr(*, "label")= chr "FUTAB"
> prv <- read_sas( "../data/prv.sas7bdat" )
> str( prv )
Classes 'tbl_df', 'tbl' and 'data.frame': 13545 obs. of 5 variables:
 $ pdat : num 1995 1995 1995 1995 1995 ...
 $ state: chr "T1" "T1" "T1" "T1" ...
 $ sex : num 1 1 1 1 1 1 1 1 1 1 ...
 $ age : num 1 2 3 4 5 6 7 8 9 10 ...
 $ n : num 3 8 7 17 22 28 40 29 33 42 ...
- attr(*, "label")= chr "PRV"

```

2.5.1 Rate data

We want a dataframe with number of DM cases (X) and deaths among DM patients (D.DM) and non-DM persons (D.nD) and person-years among DM patients (Y.DM) and non-DM persons (Y.nD), classified by sex, age and date.

However, the data from the futab dataset is clearly not usable for analysis of rates till 1996 (see the output tabel from 08-mkFU (page 165), so we restrict the follow-up data to the period after 1996-01-01.

```

> DM <- subset( FU, P>1995 & state %in% c("T1","T2") )[,c("sex","A","P","U","D","Y")]
> DM <- aggregate( DM[,c("D","Y")], by=DM[,c("sex","A","P","U")], FUN=sum )
> names( DM )[5:6] <- c("D.DM","Y.DM")
> head( DM )
  sex A    P U D.DM      Y.DM
1   2 0 1996 0    0 0.001073922
2   1 1 1996 0    0 0.002034223
3   2 1 1996 0    0 0.003585216
4   1 2 1996 0    0 0.006972621
5   2 2 1996 0    0 0.004438741
6   1 3 1996 0    0 0.004505818

> nD <- subset( FU, P>1995 & state %in% c("Well") )[,c("sex","A","P","U","T1","T2","D","Y")]
> nD$X <- nD$T1 + nD$T2
> nD <- aggregate( nD[,c("X","D","Y")], by=nD[,c("sex","A","P","U")], FUN=sum )
> names( nD )[6:7] <- c("D.nD","Y.nD")
> head( nD )
  sex A    P U X D.nD      Y.nD
1   1 0 1996 0 0    0 17.56979
2   2 0 1996 0 5    0 16.38617
3   1 1 1996 0 4   15 18.34906
4   2 1 1996 0 6   10 17.48155
5   1 2 1996 0 7    2 18.06545
6   2 2 1996 0 1    2 17.26346

> TT <- merge( nD, DM, all=TRUE )
> TT[is.na(TT)] <- 0
> TT <- transform( TT, A = A + (1+U)/3,
+                 P = P + (2-U)/3,
+                 sex = factor( sex, labels=c("M","F") ) )
> head( TT )

```

```

sex      A      P U X D.nD      Y.nD D.DM      Y.DM
1  M 0.3333333 1996.667 0 0      0 17.56979      0 0.0000000000
2  M 0.6666667 1996.333 1 1      28 17.88386      0 0.0004757016
3  M 0.3333333 1997.667 0 1      1 17.72566      0 0.0004613279
4  M 0.6666667 1997.333 1 1      18 17.32057      0 0.0001704312
5  M 0.3333333 1998.667 0 2      2 17.11636      0 0.0009267625
6  M 0.6666667 1998.333 1 2      18 17.09326      0 0.0007015743

```

```
> summary( TT )
```

```

sex      A      P      U      X      D.nD
M:4000  Min.   : 0.3333  Min.   :1996  Min.   :0.0  Min.   : 0.0  Min.   : 0
F:4000  1st Qu.:25.1667  1st Qu.:2001  1st Qu.:0.0  1st Qu.: 6.0  1st Qu.: 7
        Median :50.0000  Median :2006  Median :0.5  Median :21.0  Median :51
        Mean   :50.0000  Mean   :2006  Mean   :0.5  Mean   :41.2  Mean   :119
        3rd Qu.:74.8333  3rd Qu.:2011  3rd Qu.:1.0  3rd Qu.:68.0  3rd Qu.:205
        Max.   :99.6667  Max.   :2016  Max.   :1.0  Max.   :280.0  Max.   :590

      Y.nD      D.DM      Y.DM
Min.   : 0.02235  Min.   : 0.00  Min.   :0.00000
1st Qu.: 8.50906  1st Qu.: 0.00  1st Qu.:0.06703
Median :16.05813  Median : 5.00  Median :0.24374
Mean   :13.22582  Mean   :18.92  Mean   :0.41975
3rd Qu.:17.90357  3rd Qu.:33.00  3rd Qu.:0.63896
Max.   :23.34236  Max.   :116.00  Max.   :2.70543

```

The numbers may look a bit odd but it is mainly the mortality among 0-year old that appear in the upper triangle, which indeed *is* a bit strange. Finally we save the data for analysis:

```
> save( TT, prv, file="../data/TT.Rda" )
```

2.5.2 Prevalence data

We need a version of the prevalence dataset in 1-year classes for analysis of prevalence:

```

> P1tab <- aggregate( data.frame( N = prv$n ),
+                    data.frame( typ = Relevel( prv$state, list( DM=1:2, Pop=3 ) ),
+                               sex = factor(prv$sex,labels=c("M","F")),
+                               A = floor(prv$age),
+                               P = prv$pd ),
+                    FUN=sum )

```

```
WARNING: prv$state has been converted to a factor with levels:
T1 T2 Well
```

```
> str( P1tab )
```

```

'data.frame':      9345 obs. of  5 variables:
 $ typ: Factor w/ 2 levels "DM","Pop": 2 1 2 1 2 1 2 1 2 1 ...
 $ sex: Factor w/ 2 levels "M","F": 1 2 2 1 1 2 2 1 1 2 ...
 $ A  : num  0 0 0 1 1 1 1 2 2 2 ...
 $ P  : num  1995 1995 1995 1995 1995 ...
 $ N  : num  35821 1 34257 3 34787 ...

```

```

> PD <- subset( P1tab, typ=="DM" )[, -1]
> PN <- subset( P1tab, typ=="Pop" )[, -1]
> names( PD )[grep("N",names(PD))] <- "X"
> Ptab <- merge( PD, PN, all=TRUE )
> Ptab[is.na(Ptab)] <- 0
> Ptab$N <- Ptab$X + Ptab$N
> head( Ptab )

```

```

sex A    P X    N
1  M 0 1995 0 35821
2  M 0 1996 2 36258
3  M 0 1997 0 34920
4  M 0 1998 1 34936
5  M 0 1999 2 34151
6  M 0 2000 2 34035
> save( Ptab, file="../data/Ptab.Rda" )

```

2.5.3 Crude tables for teaching

```

> FUt看 <- aggregate( TT[,5:9], data.frame( sex=TT$sex, A=floor(TT$A/5)*5, P=floor(TT$P) ),
> str( FUt看 )
'data.frame':
      800 obs. of  8 variables:
 $ sex : Factor w/ 2 levels "M","F": 1 2 1 2 1 2 1 2 1 2 ...
 $ A   : num  0 0 5 5 10 10 15 15 20 20 ...
 $ P   : num  1996 1996 1996 1996 1996 1996 ...
 $ X   : num  27 27 29 33 43 47 44 42 61 65 ...
 $ D.nD: num  72 61 27 18 30 23 95 40 166 51 ...
 $ Y.nD: num  177 168 159 151 141 ...
 $ D.DM: num  0 0 0 0 0 0 0 0 1 0 ...
 $ Y.DM: num  0.051 0.045 0.179 0.132 0.267 ...
> head( FUt看 )
  sex  A    P  X D.nD    Y.nD D.DM    Y.DM
1  M  0 1996 27   72 177.1678    0 0.05100890
2  F  0 1996 27   61 167.8859    0 0.04497604
3  M  5 1996 29   27 158.9996    0 0.17851403
4  F  5 1996 33   18 150.6390    0 0.13218891
5  M 10 1996 43   30 140.8958    0 0.26694867
6  F 10 1996 47   23 134.9699    0 0.25601985
> Ptab <- aggregate( data.frame( N = prv$n ),
+                   data.frame( typ = Relevel( prv$state, list( DM=1:2, Pop=3 ) ),
+                               sex = factor(prv$sex,labels=c("M","F")),
+                               A = floor(prv$age/5)*5,
+                               P = prv$pdat ),
+                   FUN=sum )
WARNING: prv$state has been converted to a factor with levels:
T1 T2 Well
> str( Ptab )
'data.frame':
      1925 obs. of  5 variables:
 $ typ: Factor w/ 2 levels "DM","Pop": 1 2 1 2 1 2 1 2 1 2 ...
 $ sex: Factor w/ 2 levels "M","F": 1 1 2 2 1 1 2 2 1 1 ...
 $ A   : num  0 0 0 0 5 5 5 5 10 10 ...
 $ P   : num  1995 1995 1995 1995 1995 ...
 $ N   : num  41 172220 37 163686 163 ...
> head( Ptab )
  typ sex A    P    N
1  DM  M 0 1995   41
2 Pop  M 0 1995 172220
3  DM  F 0 1995   37
4 Pop  F 0 1995 163686
5  DM  M 5 1995   163
6 Pop  M 5 1995 150446
> save( FUt看, Ptab, file="DM5x1.Rda" )

```

Chapter 3

Prevalence and occurrence rates

3.1 Prerequisites

We will make models for the rates as observed in small intervals of age and calendar time, so we start by specifying the interval length, and then the points at which we want to predict. The transition rates are labeled by the midpoints of the Lexis squares (of width `int`) where we predict them (`a.pt` and `p.pt`), and the prevalences by the midpoints of the age-classes (`a.pt` and the time-points `t.pt` — note that we shall make predictions of rates all the way to 2030 — this will mainly be with the purpose of predicting the total number of DM persons in the population:

```
> int <- 1/12
> a.pt <- seq(0,100,int)[-1] - int/2
> t.pt <- seq(1996,2040,int)
> p.pt <- t.pt[-1] - int/2
```

We shall model all the rates by age-period-cohort models separately for men and women, both on a multiplicative and an additive scale. As a sensitivity analysis we will also model the rates only by an age-period model.

We will use natural splines to model the effects of age, period and cohort, and for all analyses we will use the same **number** of knots for these three effects, but of course place them differently based on the location of information, *i.e.* the events:

```
> nk.a <- 8
> nk.p <- 6
> nk.c <- 8
```

For the practical location of the spline knots we also define a small function which from the number of knots derives reasonable quantiles:

```
> qn <- function( nk, bd=2 ) seq( from = 1/(bd*nk),
+                               to = 1-1/(bd*nk),
+                               length = nk )
> qn( 10, 1 )
[1] 0.1000000 0.1888889 0.2777778 0.3666667 0.4555556 0.5444444 0.6333333 0.7222222
[9] 0.8111111 0.9000000
> qn( 10, 2 )
[1] 0.05 0.15 0.25 0.35 0.45 0.55 0.65 0.75 0.85 0.95
> qn( 10, 4 )
```

```
[1] 0.0250000 0.1305556 0.2361111 0.3416667 0.4472222 0.5527778 0.6583333 0.7638889
[9] 0.8694444 0.9750000
```

So this function generates `nk`, equidistant points in the interval (0,1) where the outer points are $1/(2 \times nk)$ from the end. The second parameter is for modifying the multiplier 2, so that the outer intervals are $1/(nk \times bd)$:

```
> save( qn, int, a.pt, t.pt, p.pt, nk.a, nk.p, nk.c , file="../data/inits.Rda" )
```

3.2 Diabetes prevalences

Finally, in order to get the machinery working, we need the observed prevalences and population size at the starting point, that is at 1.1.1996. These are available from the same tabulation of the diabetes register as before — note that the `N` in the `Ptab` is the number of persons *without* diabetes, and that the remaining code relies on `N` being the **total** number of persons:

```
> load( file="../data/Ptab.Rda" )
> load( file="../data/inits.Rda" )
> pr <- subset( Ptab, P>1995 )
> summary( pr )
```

sex	A	P	X	N
M:2276	Min. : 0.00	Min. :1996	Min. : 0.0	Min. : 1
F:2304	1st Qu.: 27.00	1st Qu.:2001	1st Qu.: 85.0	1st Qu.:12503
	Median : 54.00	Median :2006	Median : 388.0	Median :31840
	Mean : 54.06	Mean :2006	Mean : 774.3	Mean :25061
	3rd Qu.: 81.00	3rd Qu.:2011	3rd Qu.:1198.5	3rd Qu.:36366
	Max. :117.00	Max. :2016	Max. :5348.0	Max. :46309

```
> head( pr )
```

sex	A	P	X	N	
2	M	0	1996	2	36258
3	M	0	1997	0	34920
4	M	0	1998	1	34936
5	M	0	1999	2	34151
6	M	0	2000	2	34035
7	M	0	2001	1	34432

These are empirical prevalences (`X`—no. of cases of DM, `N`—population size) for each of the 21 dates 1.1.1996 – 1.1.2016 in 1-year intervals, but to get the machinery running we will need the number of diabetes cases in age intervals of length `int`.

So we model the prevalences as of 1 January each of the years 1996—2016, as a smooth function of age, and use the predicted prevalences to produce the prevalence of diabetes in each of the smaller age-classes that we use for the simulation. We use a log-link binomial model with a smooth spline with 12 knots:

```
> ( kp.a <- c( 15, with( pr, quantile( rep(A,X), qn(12) ) ) ) ) )
```

	4.166667%	12.5%	20.833333%	29.166667%	37.5%	45.833333%	54.166667%	62.5%
15	30	43	50	55	59	62	65	68
70.833333%	79.166667%	87.5%	95.833333%					
71	75	79	86					

```
> pr.fit <- NArray( list( sex = c("M","F"),
+                        A = a.pt,
+                        P = sort(unique(pr$P)) ) )
> str( pr.fit )
logi [1:2, 1:1200, 1:21] NA NA NA NA NA NA ...
- attr(*, "dimnames")=List of 3
..$ sex: chr [1:2] "M" "F"
..$ A : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.2916666666666667"
..$ P : chr [1:21] "1996" "1997" "1998" "1999" ...
> prod( dim(pr.fit) )
[1] 50400
```

Note that the array of the fitted prevalences has 1200 ages — we used the single-year *observations* of prevalence to predict the prevalences at a set of narrowly spaced ages.

So once we have set up the array to hold the smoothed empirical prevalences we can fill them into `pr.fit`:

```
> for( sx in dimnames(pr.fit)[["sex"]] )
+ for( dt in dimnames(pr.fit)[["P"]] )
+ {
+ pr.mod <- glm( cbind(X,N-X) ~ Ns( A, kn=kp.a ),
+              family = binomial(link="log"),
+              data = subset( pr, sex==sx & P==as.numeric(dt) ) )
+ pr.fit[sx,,dt] <- predict( pr.mod,
+                          newdata = data.frame( A = a.pt ),
+                          type = "response" )
+ }
> tt <- pr.fit[,c(1:2,NA,floor(dim(pr.fit)[2]*3/4)+1:5),]
> dimnames( tt )[[2]] <- round( as.numeric( dimnames( tt )[[2]] ), 2 )
> str( tt )
num [1:2, 1:8, 1:21] 0.000626 0.000579 0.00063 0.000583 NA ...
- attr(*, "dimnames")=List of 3
..$ sex: chr [1:2] "M" "F"
..$ A : chr [1:8] "0.04" "0.12" NA "75.04" ...
..$ P : chr [1:21] "1996" "1997" "1998" "1999" ...
> round( ftable( tt[,,-(5:10)], row.vars=2:1 ) * 100, 1 )
      P 1996 1997 1998 1999 2006 2007 2008 2009 2010 2011 2012 2013 2014 2015 2016
A sex
0.04 M    0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1
     F    0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1
0.12 M    0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1
     F    0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1  0.1
NA  M     NA   NA   NA   NA   NA   NA   NA   NA   NA   NA   NA   NA   NA   NA
     F     NA   NA   NA   NA   NA   NA   NA   NA   NA   NA   NA   NA   NA   NA   NA
75.04 M   5.7  6.1  6.5  6.8 10.6 11.3 12.0 12.9 13.7 14.6 15.5 16.0 16.2 16.4 16.6
      F   4.9  5.2  5.4  5.6  8.2  8.7  9.1  9.6 10.0 10.4 11.0 11.5 11.6 11.7 11.8
75.12 M   5.8  6.1  6.5  6.8 10.6 11.3 12.0 12.9 13.7 14.6 15.5 16.0 16.2 16.4 16.6
      F   4.9  5.2  5.4  5.6  8.2  8.7  9.1  9.7 10.0 10.4 11.1 11.5 11.7 11.7 11.9
75.21 M   5.8  6.1  6.5  6.9 10.6 11.3 12.0 12.9 13.7 14.6 15.5 16.0 16.2 16.4 16.6
      F   4.9  5.2  5.4  5.6  8.2  8.7  9.1  9.7 10.1 10.5 11.1 11.5 11.7 11.7 11.9
75.29 M   5.8  6.1  6.5  6.9 10.6 11.3 12.0 12.9 13.7 14.6 15.5 16.1 16.3 16.4 16.7
      F   5.0  5.2  5.5  5.6  8.2  8.7  9.1  9.7 10.1 10.5 11.1 11.6 11.7 11.8 11.9
75.38 M   5.8  6.1  6.5  6.9 10.6 11.3 12.0 12.9 13.7 14.6 15.5 16.1 16.3 16.4 16.7
      F   5.0  5.2  5.5  5.6  8.2  8.7  9.2  9.7 10.1 10.5 11.1 11.6 11.7 11.8 12.0
```

We can plot how the age-specific prevalences have evolved over time:


```

> plp <- function(grid=FALSE){
+ par( mfrow=c(1,2), mar=c(1,0,1,0), mgp=c(3,1,0)/1.6, las=1,
+      oma=c(2,3,0,1), bty="n" )
+ matplot( a.pt, pr.fit["M",,]*100,
+          ylim=c(0,20), xlim=c(20,90), yaxs="i", xaxt="n", yaxt="n", xlab="", ylab="",
+          type="l", lty=1, col="blue", lwd=c(1,2) )
+ if( grid ) abline(h=0:22,v=2:9*10,col=gray(0.9))
+ matlines( a.pt, pr.fit["M",,]*100,
+          type="l", lty=1, col="blue", lwd=c(1,2) )
+ text( 39, 19.5, "Men", adj=1, col="blue", cex=1.2 )
+ a89 <- grep( "89.", dimnames(pr.fit)[[2]] )[1]
+ a80 <- grep( "80.", dimnames(pr.fit)[[2]] )[1]
+ text( 89, pr.fit["M",a89,"1996"]* 99, "1996", col="blue", adj=c(1,1) )
+ text( 80, pr.fit["M",a80,"2016"]*101, "2016", col="blue", adj=c(0,0) )
+ axis( side=1 )
+ axis( side=2 )
+ axis( side=2, at=0:20, labels=NA )
+ matplot( a.pt, pr.fit["F",,]*100,
+          ylim=c(0,20), xlim=c(20,90), yaxs="i", xaxt="n", yaxt="n", xlab="", ylab="",
+          type="l", lty=1, col="red", lwd=c(1,2) )
+ if( grid ) abline(h=0:22,v=2:9*10,col=gray(0.9))
+ matlines( a.pt, pr.fit["F",,]*100,
+          type="l", lty=1, col="red", lwd=c(1,2) )
+ text( 39, 19.5, "Women", adj=1, col="red", cex=1.2 )
+ text( 89, pr.fit["F",a89,"1996"]* 99, "1996", col="red", adj=c(1,1) )
+ text( 80, pr.fit["F",a80,"2016"]*101, "2016", col="red", adj=c(1,0) )
+ axis( side=1 )
+ mtext( "Age", side=1, line=1, outer=T )
+ mtext( "DM prevalence (%)", side=2, line=2, outer=T, las=0 )
+ }
> plp()

> plp(grid=TRUE)

```

For the calculations we shall only use the estimated prevalences as of 1.1.1996 as starting point for the simulation:

```
> save( pr, pr.fit, file="../data/prFit.Rda" )
```

3.2.1 Trends in prevalence

A crude way of summarizing the prevalences is to assume that relative change is constant from year to year. So we set up a model that does this separately for men and women, and store the predicted values for comparison with those from the model with no assumption about the time evolution:

```

> p.pr <- as.numeric( dimnames( pr.fit )[[ "P" ] ] )
> pr.lfit <- pr.fit
> pr.chg <- NArray( list( dimnames(pr.fit)[[ "sex" ] ],
+                        c("% chg/y", "lo", "hi" ) ) )
> for( sx in dimnames(pr.fit)[[ "sex" ] ] )
+ {
+ lmod <- glm( cbind(X,N-X) ~ Ns( A, kn=kp.a ) + P,
+            family = binomial(link="log"),
+            data = subset( pr, sex==sx ) )

```

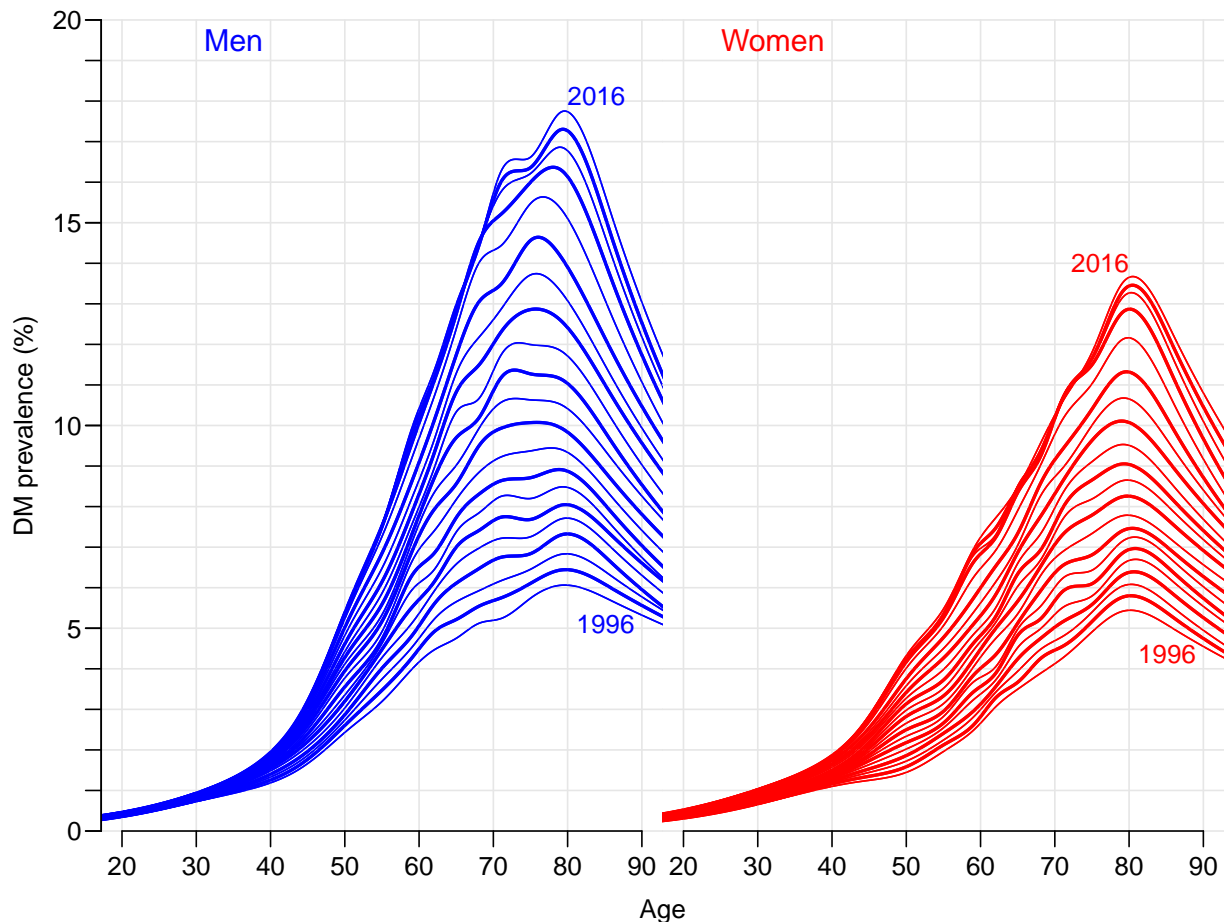


Figure 3.1: Smoothed age-specific prevalences for the years 1996–2016. Blue is men, red is women.

```
+ pr.chg[sx,] <- ( ci.exp( lmod, subset="P" ) - 1 ) * 100
+ pr.lfit[sx,,] <- predict( lmod,
+                          newdata = data.frame( A=rep(a.pt,      length(p.pr)),
+                                                P=rep(p.pr,each=length(a.pt)) ),
+                          type = "response" )
+ }
```

This model is of course a simplification of the model above, which has a totally unspecified age-date interaction, so we can have a peep at how the predicted prevalences looks:

```
> par( mfrow=c(1,2), mar=c(1,0,1,0), mgp=c(3,1,0)/1.6, las=1,
+       oma=c(2,3,0,1), bty="n" )
> lblu <- rgb( 3,3,4,max=4 )
> lred <- rgb( 4,3,3,max=4 )
> matplot( a.pt, pr.fit["M",,]*100,
+          ylim=c(0,22), xlim=c(20,90), yaxs="i", xaxt="n", yaxt="n", xlab="", ylab="",
+          type="n", lty=1, col="blue", lwd=c(1,2) )
> abline( h=0:25, v=seq(0,100,5), col=gray(0.9) )
> matlines( a.pt, pr.fit["M",,]*100, type="l", lty=1, col=lblu , lwd=c(2,3) )
> matlines( a.pt, pr.lfit["M",,]*100, type="l", lty=1, col="blue", lwd=c(2,3) )
> text( 25, 22, "Men", adj=c(0,1), col="blue", cex=1.2 )
> a89 <- grep( "89.", dimnames(pr.fit)[[2]] ) [1]
> a80 <- grep( "80.", dimnames(pr.fit)[[2]] ) [1]
```

```

> text( 89, pr.fit["M",a89,"1996"]* 99, "1996", col="blue", adj=c(1,1) )
> text( 80, pr.fit["M",a80,"2016"]*101, "2016", col="blue", adj=c(0,0) )
> axis( side=1 )
> axis( side=2 )
> matplot( a.pt, pr.fit["F",,]*100,
+         ylim=c(0,22), xlim=c(20,90), yaxs="i", xaxt="n", yaxt="n", xlab="", ylab="",
+         type="n", lty=1, col="red", lwd=c(1,2) )
> abline( h=0:25, v=seq(0,100,5), col=gray(0.9) )
> matlines( a.pt, pr.fit["F",,]*100, type="l", lty=1, col=lred , lwd=c(2,3) )
> matlines( a.pt, pr.lfit["F",,]*100, type="l", lty=1, col="red", lwd=c(2,3) )
> text( 25, 22, "Women", adj=c(0,1), col="red", cex=1.2 )
> text( 89, pr.fit["F",a89,"1996"]* 99, "1996", col="red", adj=c(1,1) )
> text( 80, pr.fit["F",a80,"2016"]*101, "2016", col="red", adj=c(1,0) )
> axis( side=1 )
> mtext( "Age", side=1, line=1, outer=T )
> mtext( "DM prevalence (%)", side=2, line=2, outer=T, las=0 )

```

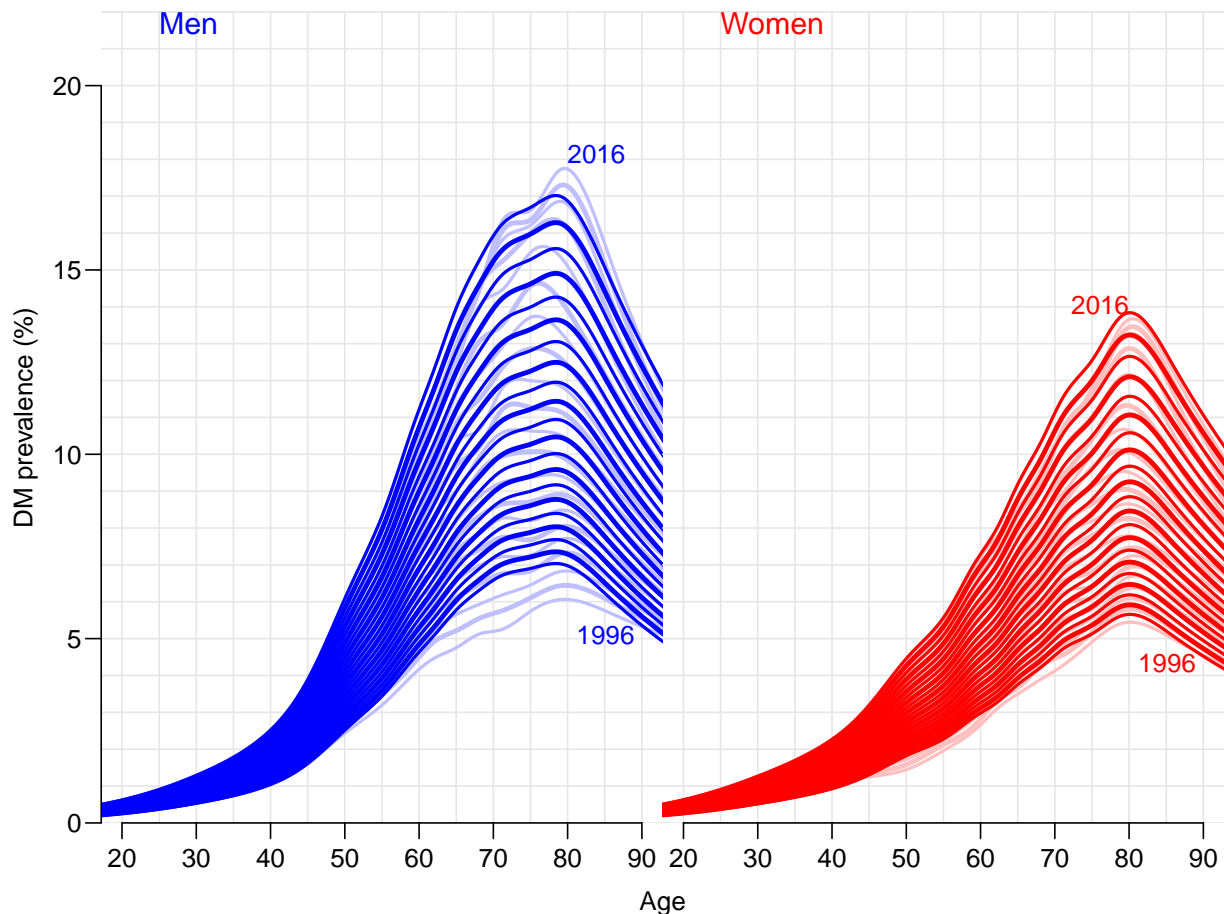


Figure 3.2: *Smoothed age-specific prevalences for the 17-year period 1995–2015 using a model with constant annual relative change in prevalences (full color) compared to the smoothing of the single years (pale color). Blue is men, red is women.*

From figure 3.2 we see that for men the summary using a constant relative change in prevalence is not a very good summary of the change in prevalences; it does not capture the change in the age of peak prevalence of men from 85 in 1995 to 75 in 2015. So the

overall estimate of some 6% in relative annual increase of prevalences over the 20-year period 1995–2015, is *not* providing an adequate summary:

```
> round( pr.chg, 2 )
  % chg/y  lo  hi
M    4.51 4.49 4.54
F    4.57 4.55 4.60
```

3.2.2 Prevalence age-period interaction

Hence the relevant description of average changes per year would be using a model for the prevalences where we allowed the relative change to vary smoothly by age. This is done by including an interaction between a spline term in age and period, and the subsequently fishing out the relative change using a spline basis with a bit fewer knots to fish out the period multiplier.

It goes as follows, where we also as before extract the predicted values for comparison with the prevalence curves fitted separately for each year:

```
> ( kx.a <- c( 10, with( pr, quantile( rep(A,X), qn(5) ) ) ) )
  10% 30% 50% 70% 90%
  10  41  55  64  71  81
> CA <- Ns( 1:99, kn=kx.a, intercept=TRUE )
> A.chg <- NArray( list( A=1:99, c("Est","lo","hi"), sex=c("M","F") ) )
> for( sx in dimnames(pr.fit)[["sex"]] )
+   {
+   limod <- glm( cbind(X,N-X) ~ Ns( A, kn=kp.a ) +
+                 I(P-2000):Ns( A, kn=kx.a, intercept=TRUE ),
+                 family = binomial(link="log"),
+                 data = subset( pr, sex==sx ) )
+   A.chg[, ,sx] <- ci.exp( limod, subset="P", ctr.mat=CA )
+   pr.lfit[sx, ,] <- predict( limod,
+                             newdata = data.frame( A=rep(a.pt, length(p.pr)),
+                                                    P=rep(p.pr, each=length(a.pt)) ),
+                             type = "response" )
+   }
> par( mar=c(3,3,1,1), mgp=c(3,1,0)/1.6, bty="n" )
> matplot( 1:99, (cbind( A.chg[, , "M"], A.chg[, , "F"] )-1)*100,
+          col=rep(c("blue","red"), each=3), lwd=c(3,1,1), lty=1, type="l",
+          ylim=c(0,8), yaxs="i",
+          ylab="Annual change in DM prevalence (%)", xlab="Age" )
> abline( h=pr.chg[,1], col=c("blue","red") )
```

We can also as with the naïve linear change model show how the fitted values under this interaction model looks relative to the separate analyses by year (or full interaction model). The code is exactly as before, because we put the fitted values into the same structure as before:

```
> par( mfrow=c(1,2), mar=c(1,0,1,0), mgp=c(3,1,0)/1.6, las=1,
+       oma=c(2,3,0,1), bty="n" )
> lblu <- rgb( 3,3,4,max=4 )
> lred <- rgb( 4,3,3,max=4 )
> matplot( a.pt, pr.fit["M", ,]*100,
+          ylim=c(0,22), xlim=c(20,90), yaxs="i", xaxt="n", yaxt="n", xlab="", ylab="",
+          type="n", lty=1, col="blue", lwd=c(1,2) )
```

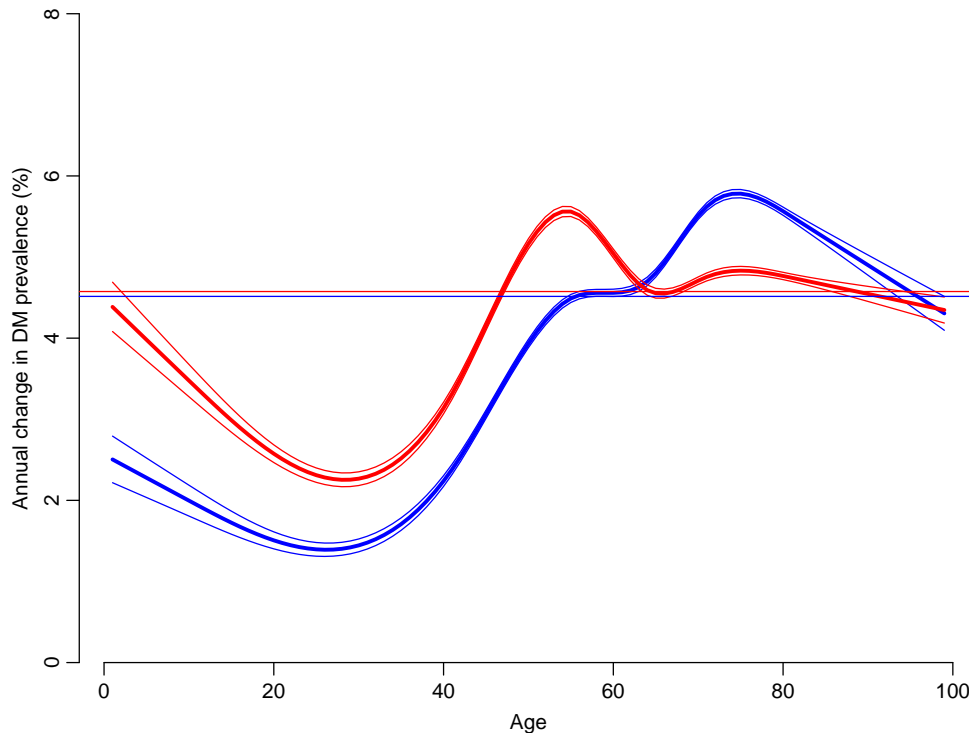


Figure 3.3: The estimated change in prevalence in different ages, separately for men (blue) and women (red). The horizontal lines indicate the estimate from the naïve model with constant change for all ages.

```

> abline( h=0:25, v=seq(0,100,5), col=gray(0.9) )
> matlines( a.pt, pr.fit["M",,]*100, type="l", lty=1, col=lblu , lwd=c(2,3) )
> matlines( a.pt, pr.lfit["M",,]*100, type="l", lty=1, col="blue", lwd=c(2,3) )
> text( 25, 21.5, "Men", adj=0, col="blue", cex=1.2 )
> a89 <- grep( "89.", dimnames(pr.fit)[[2]] ) [1]
> a80 <- grep( "80.", dimnames(pr.fit)[[2]] ) [1]
> text( 89, pr.fit["M",a89,"1996"]* 99, "1996", col="blue", adj=c(1,1) )
> text( 80, pr.fit["M",a80,"2016"]*101, "2016", col="blue", adj=c(0,0) )
> axis( side=1 )
> axis( side=2 )
> matplot( a.pt, pr.fit["F",,]*100,
+         ylim=c(0,22), xlim=c(20,90), yaxs="i", xaxt="n", yaxt="n", xlab="", ylab="",
+         type="n", lty=1, col="red", lwd=c(1,2) )
> abline( h=0:25, v=seq(0,100,5), col=gray(0.9) )
> matlines( a.pt, pr.fit["F",,]*100, type="l", lty=1, col=lred , lwd=c(2,3) )
> matlines( a.pt, pr.lfit["F",,]*100, type="l", lty=1, col="red", lwd=c(2,3) )
> text( 25, 21.5, "Women", adj=0, col="red", cex=1.2 )
> text( 89, pr.fit["F",a89,"1996"]* 99, "1996", col="red", adj=c(1,1) )
> text( 80, pr.fit["F",a80,"2016"]*101, "2016", col="red", adj=c(1,0) )
> axis( side=1 )
> mtext( "Age", side=1, line=1, outer=T )
> mtext( "DM prevalence (%)", side=2, line=2, outer=T, las=0 )

```

From figure 3.4 is seen that the model captures the actual pattern much better than the simple model with an annual change common across ages.

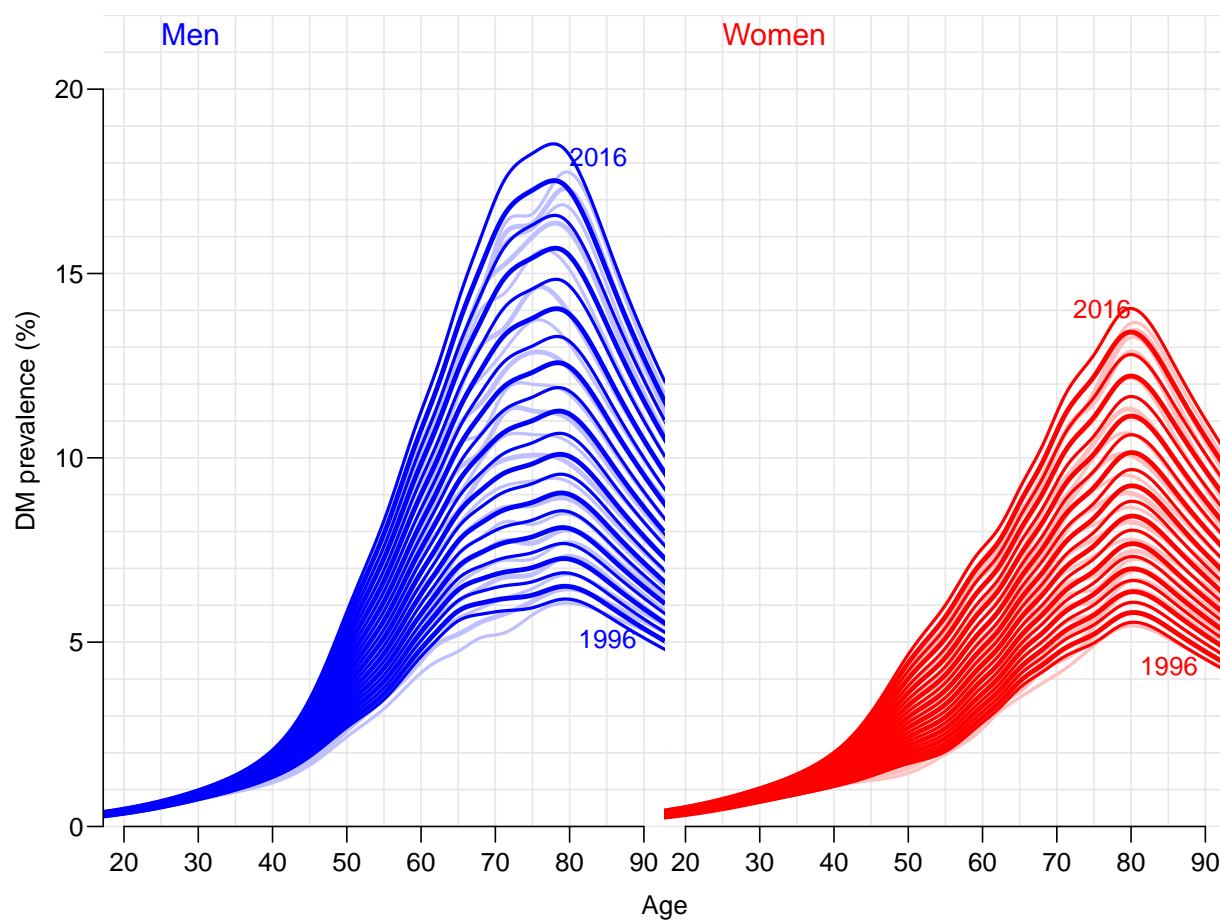


Figure 3.4: *Smoothed age-specific prevalences for the 20-year period 1996–2016 using a model with age-specific constant annual relative change in prevalences (full color) compared to the smoothing of the single years (pale color). Blue is men, red is women.*

3.3 DM incidence

In this chapter we use the constructed follow-up dataset for the entire population:

```
> options( width=95 )
> library( Epi )
> library( splines )
> load( file="../data/TT.Rda" )
```

3.3.1 Age-Period-Cohort modelling

We will use `X` and `Y.nd` as response variables in the analysis of diabetes incidence rates; we fit separate APC-models for total diabetes incidence men and women and plot the estimates together:

```
> DD <- TT[,c("sex", "A", "P", "X", "Y.nd")]
> names(DD)[4:5] <- c("D", "Y")
> ( A.kn <- quantile( rep(DD$A      ,DD$D), probs=(1:8-0.5)/8 ) )
 6.25%  18.75%  31.25%  43.75%  56.25%  68.75%  81.25%  93.75%
35.33333 47.66667 54.33333 59.66667 64.33333 68.66667 74.33333 82.66667
> ( P.kn <- quantile( rep(DD$P      ,DD$D), probs=(1:6-0.5)/6 ) )
8.333333%    25% 41.66667% 58.33333%    75% 91.66667%
1998.333 2002.333 2005.667 2009.333 2011.667 2014.333
> ( C.kn <- quantile( rep(DD$P-DD$A,DD$D), probs=(1:8-0.5)/8 ) )
 6.25%  18.75%  31.25%  43.75%  56.25%  68.75%  81.25%  93.75%
1922.333 1931.667 1938.333 1943.333 1947.667 1953.333 1960.333 1972.667
> acpM <- apc.fit( subset(DD,sex=="M"), ref.c=1950, parm="ACP",
+                 npar=list(A=A.kn,P=P.kn,C=C.kn) )
[1] "ML of APC-model Poisson with log(Y) offset : ( ACP ):\n"
```

Analysis of deviance for Age-Period-Cohort model

	Resid. Df	Resid. Dev	Df	Deviance	Pr(>Chi)
Age	3992	16129			
Age-drift	3991	13747	1	2382.46	< 2.2e-16 ***
Age-Cohort	3985	13683	6	63.27	9.741e-12 ***
Age-Period-Cohort	3981	11179	4	2504.16	< 2.2e-16 ***
Age-Period	3987	11259	-6	-79.66	4.207e-15 ***
Age-drift	3991	13747	-4	-2487.77	< 2.2e-16 ***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```
> acpF <- apc.fit( subset(DD,sex=="F"), ref.c=1950, parm="ACP",
+                 npar=list(A=A.kn,P=P.kn,C=C.kn) )
[1] "ML of APC-model Poisson with log(Y) offset : ( ACP ):\n"
```

Analysis of deviance for Age-Period-Cohort model

	Resid. Df	Resid. Dev	Df	Deviance	Pr(>Chi)
Age	3992	16229			
Age-drift	3991	13964	1	2265.17	< 2.2e-16 ***
Age-Cohort	3985	13678	6	285.53	< 2.2e-16 ***
Age-Period-Cohort	3981	11291	4	2386.63	< 2.2e-16 ***
Age-Period	3987	11743	-6	-451.46	< 2.2e-16 ***
Age-drift	3991	13964	-4	-2220.70	< 2.2e-16 ***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1


```
> apcM <- apc.fit( subset(DD,sex=="M"), ref.p=2000, parm="APC",
+                 npar=list(A=A.kn,P=P.kn,C=C.kn) )
```

```
[1] "ML of APC-model Poisson with log(Y) offset : ( APC ):\n"
```

```
Analysis of deviance for Age-Period-Cohort model
```

	Resid. Df	Resid. Dev	Df	Deviance	Pr(>Chi)
Age	3992	16129			
Age-drift	3991	13747	1	2382.46	< 2.2e-16 ***
Age-Cohort	3985	13683	6	63.27	9.741e-12 ***
Age-Period-Cohort	3981	11179	4	2504.16	< 2.2e-16 ***
Age-Period	3987	11259	-6	-79.66	4.207e-15 ***
Age-drift	3991	13747	-4	-2487.77	< 2.2e-16 ***

```
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
> apcF <- apc.fit( subset(DD,sex=="F"), ref.p=2000, parm="APC",
+                 npar=list(A=A.kn,P=P.kn,C=C.kn) )
```

```
[1] "ML of APC-model Poisson with log(Y) offset : ( APC ):\n"
```

```
Analysis of deviance for Age-Period-Cohort model
```

	Resid. Df	Resid. Dev	Df	Deviance	Pr(>Chi)
Age	3992	16229			
Age-drift	3991	13964	1	2265.17	< 2.2e-16 ***
Age-Cohort	3985	13678	6	285.53	< 2.2e-16 ***
Age-Period-Cohort	3981	11291	4	2386.63	< 2.2e-16 ***
Age-Period	3987	11743	-6	-451.46	< 2.2e-16 ***
Age-drift	3991	13964	-4	-2220.70	< 2.2e-16 ***

```
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

If we stick to the period-major parametrization as in figure 3.5, we are essentially referring to cross-sectional rates, and they seem to have a peak around age 80. However since there is an increasing trend the peak incidence for a given generation is more likely at 85 years as shown in figure ??, using the cohort major parametrization, the longitudinal approach.

3.3.2 Time-trends in rates

The overall time trend in the rates are in the `Drift` component of the `apc` object, here we give the average annual increase in incidence rates among men and women:

```
> pctchg <- (cbind( apcM$Drift, apcF$Drift )-1)*100
> colnames( pctchg ) <- c("Men", "lo", "up", "Women", "lo", "up")
> round( pctchg, 2 )
```

	Men	lo	up	Women	lo	up
APC (D-weights)	2.15	2.06	2.24	2.64	2.54	2.74
A-d	2.00	1.92	2.09	2.20	2.10	2.29

Thus we see that the average annual trend in rates negligible for men and only slightly more pronounced for women.

3.3.3 Summary of the APC modelling

The deviance analysis of the model did not surprisingly show that both cohort and period have non-linear effects, however this formal significance is largely due to the large data base.

```
> par( mfrow=c(2,1), mar=c(0,4,0,4), oma=c(3,0,1,0), mgp=c(3,1,0)/1.6, las=1 )
> apc.frame( a.lab=seq(20,80,20), a.tic=c(5,seq(10,90,10)),
+           cp.lab=seq(1900,2020,20), cp.tic=seq(1900,2020,10),
+           r.lab=outer(c(1,2,5),10^(-1:1)), tic.fac=2,
+           r.tic=c(outer(c(5:9/10,1:5),10^(-1:1)),60), rr.ref=1,
+           gap=15, r.txt="DM incidence per 1000 PY", side=c(2,4) )
> lines( acpM, col="blue", ci=TRUE )
> lines( acpF, col="red" , ci=TRUE )
> apc.frame( a.lab=seq(20,80,20), a.tic=c(5,seq(10,90,10)),
+           cp.lab=seq(1900,2020,20), cp.tic=seq(1900,2020,10),
+           r.lab=outer(c(1,2,5),10^(-1:1)), tic.fac=2,
+           r.tic=c(outer(c(5:9/10,1:5),10^(-1:1)),60), rr.ref=1,
+           gap=15, r.txt="DM incidence per 1000 PY", side=c(1,2,4) )
> lines( apcM, col="blue", ci=TRUE )
> lines( apcF, col="red" , ci=TRUE )
```

In figure 3.5 is shown the same model in two different parametrizations, one with longitudinal and one with cross-sectional age-specific rates. Another way of visualizing the model is to show the estimated age-specific incidence rates for different birth cohorts.

3.3.4 Prediction by age for different cohorts

To that end we use the model-objects returned by the `apc.fit` function to produce predicted rates. So we set up a prediction frame with ages for 15 different cohorts:

```
> prf <- data.frame( A = rep( c(NA,0:98), 8 ),
+                   C = rep( seq(1910,1980,10), each=100 ),
+                   Y = 1 )[-1,]
> prf <- transform( prf, P = C + A )
```

The we can make a fit of the models of relevance and make predictions based on this new frame. ¹

```
> Mapc <- glm( D ~ Ns( A, kn=apcM$Knots$Age ) +
+             Ns( P-A, kn=apcM$Knots$Coh ) +
+             Ns( P , kn=apcM$Knots$Per ),
+             offset = log( Y ),
+             family = poisson,
+             data = subset( DD, sex=="M" ) )
> Map <- glm( D ~ Ns( A, kn=apcM$Knots$Age ) +
+            Ns( P, kn=apcM$Knots$Per ),
+            offset = log( Y ),
+            family = poisson,
+            data = subset( DD, sex=="M" ) )
> Mac <- glm( D ~ Ns( A, kn=apcM$Knots$Age ) +
+            Ns( P-A, kn=apcM$Knots$Coh ),
+            offset = log( Y ),
+            family = poisson,
```

¹Note that we cannot use the returned model from the `apc` object since this is defined in terms specific matrices and *not* in terms of A, P and C:

```

+           data = subset( DD, sex=="M" )
> Fapc <- glm( D ~ Ns( A, kn=apcF$Knots$Age ) +
+           Ns( P-A, kn=apcF$Knots$Coh ) +
+           Ns( P , kn=apcF$Knots$Per ),
+           offset = log( Y ),
+           family = poisson,
+           data = subset( DD, sex=="F" ) )
> Fap <- glm( D ~ Ns( A, kn=apcF$Knots$Age ) +
+           Ns( P, kn=apcF$Knots$Per ),
+           offset = log( Y ),
+           family = poisson,
+           data = subset( DD, sex=="F" ) )
> Fac <- glm( D ~ Ns( A, kn=apcF$Knots$Age ) +
+           Ns( P-A, kn=apcF$Knots$Coh ),
+           offset = log( Y ),
+           family = poisson,
+           data = subset( DD, sex=="F" ) )
> summary( fitted( apcM$Model ) - fitted( Mapc ) )
      Min.      1st Qu.      Median      Mean      3rd Qu.      Max.
-5.116e-13 -1.299e-14  0.000e+00  5.030e-14  8.171e-14  1.023e-12
> summary( fitted( apcF$Model ) - fitted( Fapc ) )
      Min.      1st Qu.      Median      Mean      3rd Qu.      Max.
-4.121e-13  0.000e+00  4.086e-14  4.674e-14  8.171e-14  6.537e-13

```

From the last summary we see that the models are the same as those fitted by `apc.fit`, an moreover we can use this latter to make predictions, regardless of the overparametrization (we will get a warning, though). Recall that the `Y` was scaled to be person-millenia, so we get fitted values as rates per 1000 (namely the expected numbers based on the model for a data point where `Y` is equal to 1, as specified in `prf`):

```

> prr <- subset( prf, (P<2016 & P>1995) | is.na(P) )
> Mfit.apc <- predict( Mapc, newdata=prr )
> Mfit.ap <- predict( Map , newdata=prr )
> Mfit.ac <- predict( Mac , newdata=prr )
> Ffit.apc <- predict( Fapc, newdata=prr )
> Ffit.ap <- predict( Fap , newdata=prr )
> Ffit.ac <- predict( Fac , newdata=prr )

```

For comparison we overlay empirical rates, which we compute for the cohorts 1910 (born 1905–15), ..., 1980 (born 1975–85) calculated in C-sets (\diamond); the `gc` and `gp` are the midpoints of the cohort and period in the C-sets:

```

> DD.x <- transform( DD,
+                 gc = floor(((P-A)-1905)/10)*10+1910,
+                 gp = floor(P)+0.5 )
> ee <- data.frame( xtabs( cbind(D,Y) ~ sex + gp + gc,
+                 data = subset( DD.x, gc>1905 & gc<1985 ) ) )
> ee <- reshape( ee, timevar = "Var4",
+                 idvar = c("sex","gp","gc"),
+                 dir = "wide" )
> names( ee )[4:5] <- c("D","Y")
> ee <- transform( ee, gp = as.numeric(as.character(gp)),
+                 gc = as.numeric(as.character(gc)) )
> str( ee )
'data.frame':      320 obs. of  5 variables:
 $ sex: Factor w/ 2 levels "M","F": 1 2 1 2 1 2 1 2 1 2 ...

```

```

$ gp : num 1996 1996 1998 1998 1998 ...
$ gc : num 1910 1910 1910 1910 1910 1910 1910 1910 1910 1910 ...
$ D  : num 363 645 322 477 266 495 242 382 175 326 ...
$ Y  : num 45.8 95.1 39.2 84.8 33.2 ...

```

We then overlay the empirical over the fitted rates from the three different models, the age-period, the age-cohort and the apc-model:

```

> par( mfrow=c(2,1), mar=c(0,0,0,0), oma=c(3,3,1,1), mgp=c(3,1,0)/1.6 )
> matplot( prr$A, exp(Mfit.apc), type="l", lty=1,
+         log="y", ylim=c(0.2,25), lwd=3, xaxt="n", xlab="", ylab="" )
> matlines( prr$A, exp(Mfit.ap), type="l", lty="11", lwd=2 )
> matlines( prr$A, exp(Mfit.ac), type="l", lty=1, lwd=2 )
> with( subset(ee,sex=="M"),
+       points( gp-gc, D/Y, pch=16, col=rainbow(8)[factor(gc)], cex=0.8 ) )
> text( 20, 14, "Men", col="blue" )
> matplot( prr$A, exp(Ffit.apc), type="l", lty=1,
+         log="y", ylim=c(0.2,25), lwd=3, xlab="", ylab="" )
> matlines( prr$A, exp(Ffit.ap), type="l", lty="11", lwd=2 )
> matlines( prr$A, exp(Ffit.ac), type="l", lty=1, lwd=2 )
> with( subset(ee,sex=="F"),
+       points( gp-gc, D/Y, pch=16, col=rainbow(8)[factor(gc)], cex=0.8 ) )
> text( 20, 14, "Women", col="red" )
> mtext( "DM incidence rate per 1000 PY", side=2, outer=TRUE, line=2, las=0 )
> mtext( "Age (years)", side=1, outer=TRUE, line=2 )

```

3.3.5 Prediction by age across periods

Alternatively we could show the rates in different ages as a function of calendar time; we shall do this for ages 20, 30, ..., 90, note that we use the natural spline property of linearity to boldly predict rates beyond 2016-01-01.

```

> prf <- data.frame( A = rep( 2:8*10, each=31 ),
+                   P = rep( 1995:2025, 7 ),
+                   Y = 1 ) [outer(c(NA,1:31),0:6*31,"+")[-1],]

```

Then we can show the predicted rates:

```

> plp <-
+ function( mod, clr, pr, ya="l" )
+ {
+   matplot( prf$P, ci.pred( mod, prf ),
+           type="l", lty=c("solid","22","66","11","33")[c(1,1,1,1,3,3,4,5,5)],
+           lwd=c(3,1,1), col=clr,
+           log="y", xaxt="n", xlab="", ylab="", ylim=c(0.2,20), yaxt=ya )
+   abline( v = 2016, col="gray" )
+ }
> par( mfcol=c(3,2), mar=c(0,0,0,0), oma=c(3,3,1,1), mgp=c(3,1,0)/1.6 )
> plp( Mapc, "blue", prf ) ; text( 1995, 20, "APC", adj=c(0,1) )
> plp( Map, "blue", prf ) ; text( 1995, 20, "AP", adj=c(0,1) )
> plp( Mac, "blue", prf ) ; text( 1995, 20, "AC", adj=c(0,1) )
> axis( side=1 )
> plp( Fapc, "red", prf, "n" )
> plp( Fap, "red", prf, "n" )
> plp( Fac, "red", prf, "n" )
> axis( side=1 )
> mtext( "DM incidence rate per 1000 PY", side=2, outer=TRUE, line=2, las=0 )
> mtext( "Date of FU", side=1, outer=TRUE, line=2 )

```

3.3.6 Saving the fitted models

We then save these fitted APC-models with different parametrizations:

```
> save( DD, file="../data/incdata.Rda" )  
> save( acpM, apcM, acpF, apcF,  
+       Mapc, Mac, Map, Fapc, Fac, Fap, file="../data/inc.Rda" )
```

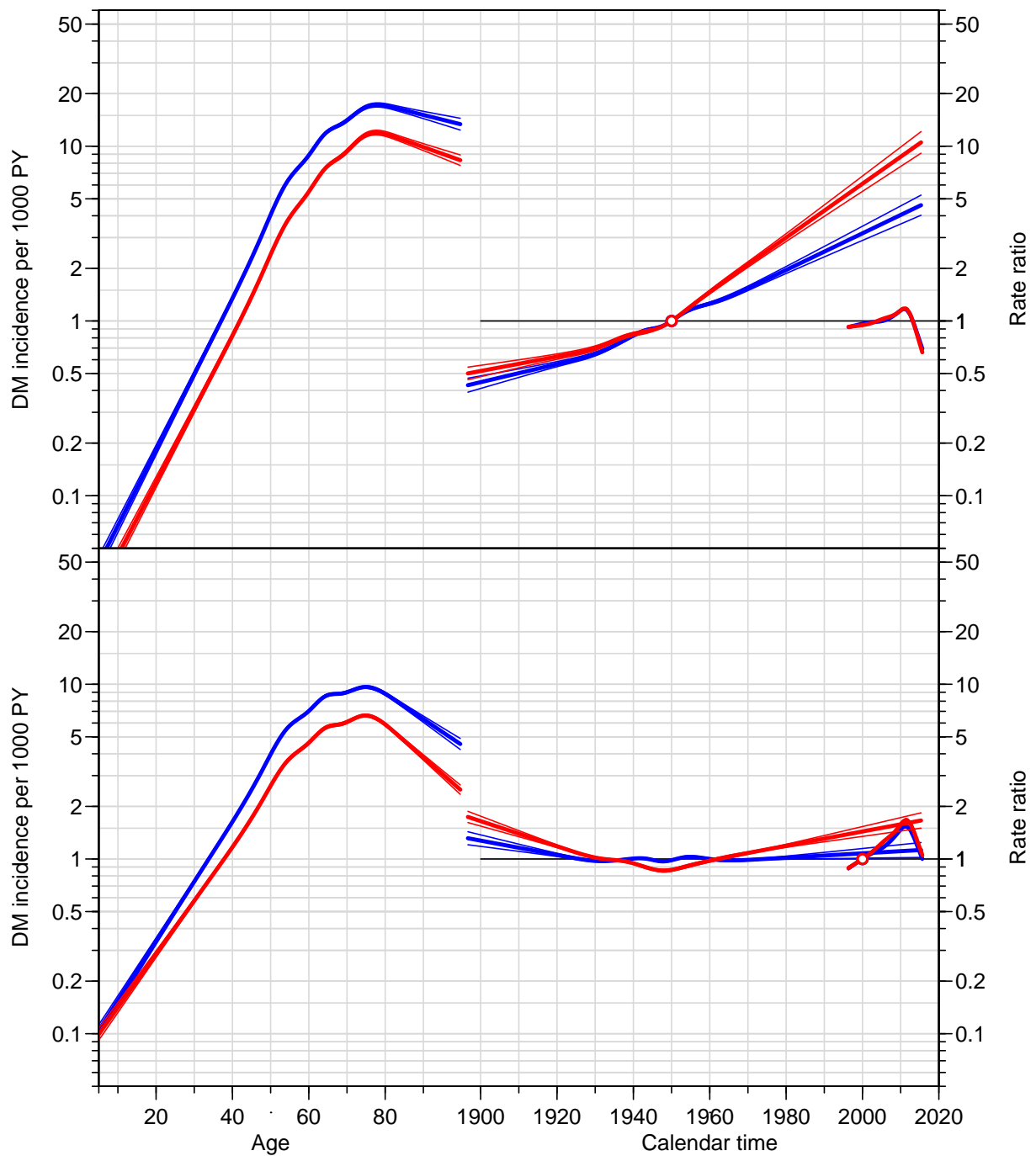


Figure 3.5: Age-Period-Cohort models for DM incidence among men (blue) and women (red), using the same scaling in the two plots. The top panel is the parametrization with horizontal period effect and cohort reference 1950, bottom panel is the parametrization with horizontal cohort effect and period reference 2000.

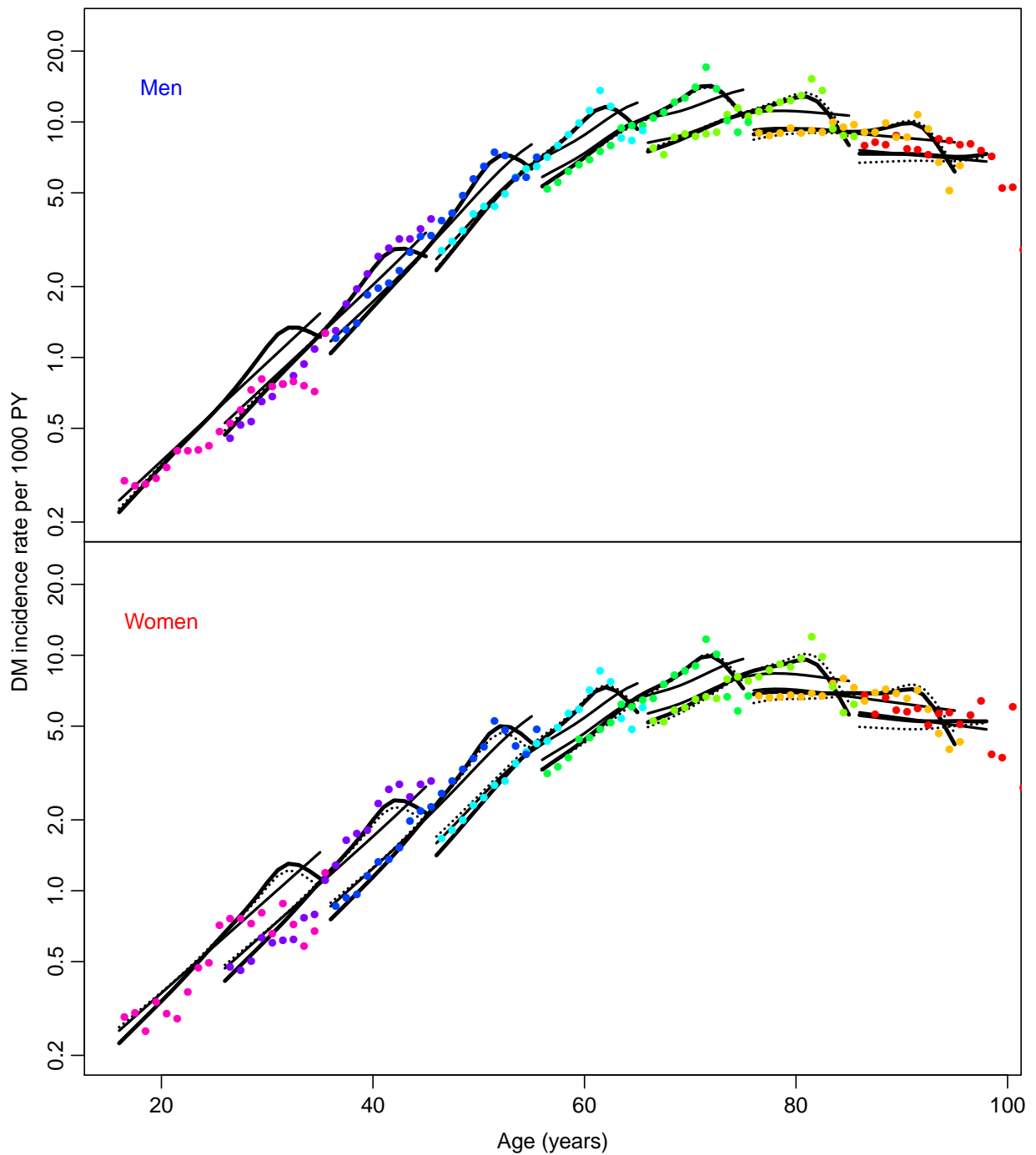


Figure 3.6: *Fitted age-specific incidence rates for the cohorts 1910, . . . , 1980: Full thick line: APC-model, broken line: AP model and full thin line: AC-model. Empirical age-specific rates from C-sets for 1-year period and 10-year cohorts are given as colored dots, colored separately for each cohort.*

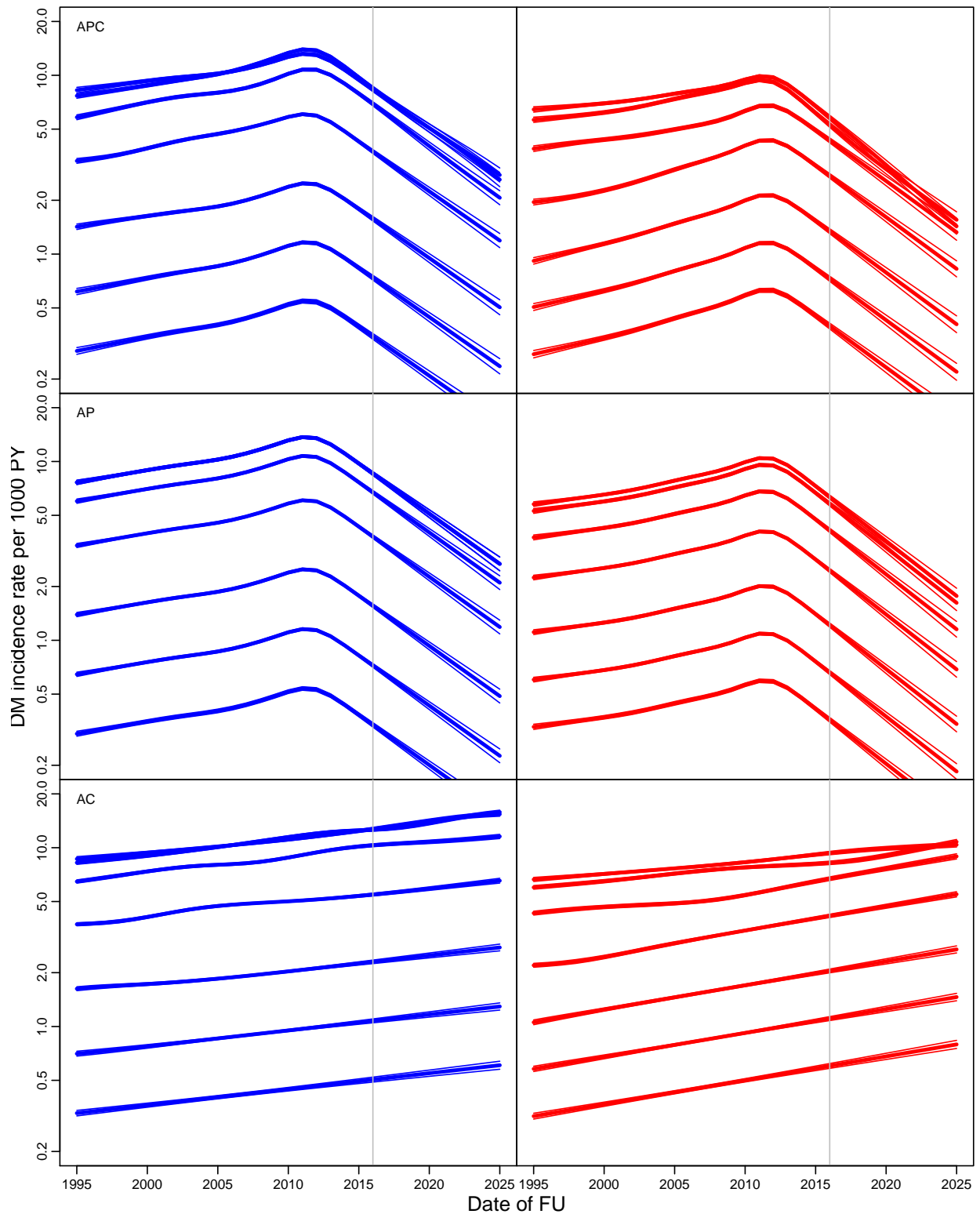


Figure 3.7: Fitted age-specific incidence rates for ages 20, 30, ..., 80. Blue: man, red: women, model from top: APC, AP, AC.

3.4 Mortality

3.4.1 Mortality in non-diabetics

We are going to use `Y.nD` and `Y.nD` as response variables in the analysis of mortality rates; we fit separate models for men and women and plot the estimates together:

```
> nD <- TT[,c("sex", "A", "P", "D.nD", "Y.nD")]
> names( nD )[4:5] <- c("D", "Y")
> ( A.nD.kn <- c( 10, 20,
+               quantile( rep(nD$A      ,nD$D), probs=(1:8-0.5)/8 ) ) )
10.00000 20.00000 50.66667 64.33333 71.66667 77.33333 81.33333 85.33333 88.66667 93.66667
> ( P.nD.kn <- quantile( rep(nD$P      ,nD$D), probs=(1:6-0.5)/6 ) )
8.333333%    25% 41.66667% 58.33333%    75% 91.66667%
1997.333 2000.667 2003.667 2006.667 2010.333 2014.333
> ( C.nD.kn <- quantile( rep(nD$P-nD$A,nD$D), probs=(1:8-0.5)/8 ) )
 6.25% 18.75% 31.25% 43.75% 56.25% 68.75% 81.25% 93.75%
1909.667 1915.667 1920.333 1924.333 1928.667 1934.333 1942.667 1955.667
> nDacpM <- apc.fit( subset(nD,sex=="M"),
+                   ref.c=1950,
+                   parm="ACP",
+                   npar=list(A=A.nD.kn,P=P.nD.kn,C=C.nD.kn) )
[1] "ML of APC-model Poisson with log(Y) offset : ( ACP ):\n"
```

Analysis of deviance for Age-Period-Cohort model

	Resid. Df	Resid. Dev	Df	Deviance	Pr(>Chi)
Age	3990	18794.7			
Age-drift	3989	7763.3	1	11031.4	< 2.2e-16 ***
Age-Cohort	3983	6581.4	6	1181.9	< 2.2e-16 ***
Age-Period-Cohort	3979	6568.2	4	13.3	0.01011 *
Age-Period	3985	7709.3	-6	-1141.2	< 2.2e-16 ***
Age-drift	3989	7763.3	-4	-54.0	5.313e-11 ***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```
> nDacpF <- apc.fit( subset(nD,sex=="F"),
+                   ref.c=1950,
+                   parm="ACP",
+                   npar=list(A=A.nD.kn,P=P.nD.kn,C=C.nD.kn) )
[1] "ML of APC-model Poisson with log(Y) offset : ( ACP ):\n"
```

Analysis of deviance for Age-Period-Cohort model

	Resid. Df	Resid. Dev	Df	Deviance	Pr(>Chi)
Age	3990	13636.1			
Age-drift	3989	7014.9	1	6621.3	< 2.2e-16 ***
Age-Cohort	3983	5469.6	6	1545.3	< 2.2e-16 ***
Age-Period-Cohort	3979	5444.9	4	24.7	5.784e-05 ***
Age-Period	3985	6905.4	-6	-1460.5	< 2.2e-16 ***
Age-drift	3989	7014.9	-4	-109.5	< 2.2e-16 ***

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```
> par( mar=c(3,4,1,4), mgp=c(3,1,0)/1.6, las=1 )
> plot( nDacpF, lty=1, ci=TRUE, col="red", r.txt="Rate per 1000 PY" )
```



```

cp.offset  RR.fac
  1790      10
> lines( nDacpM, lty=1, ci=TRUE, col="blue" )

```

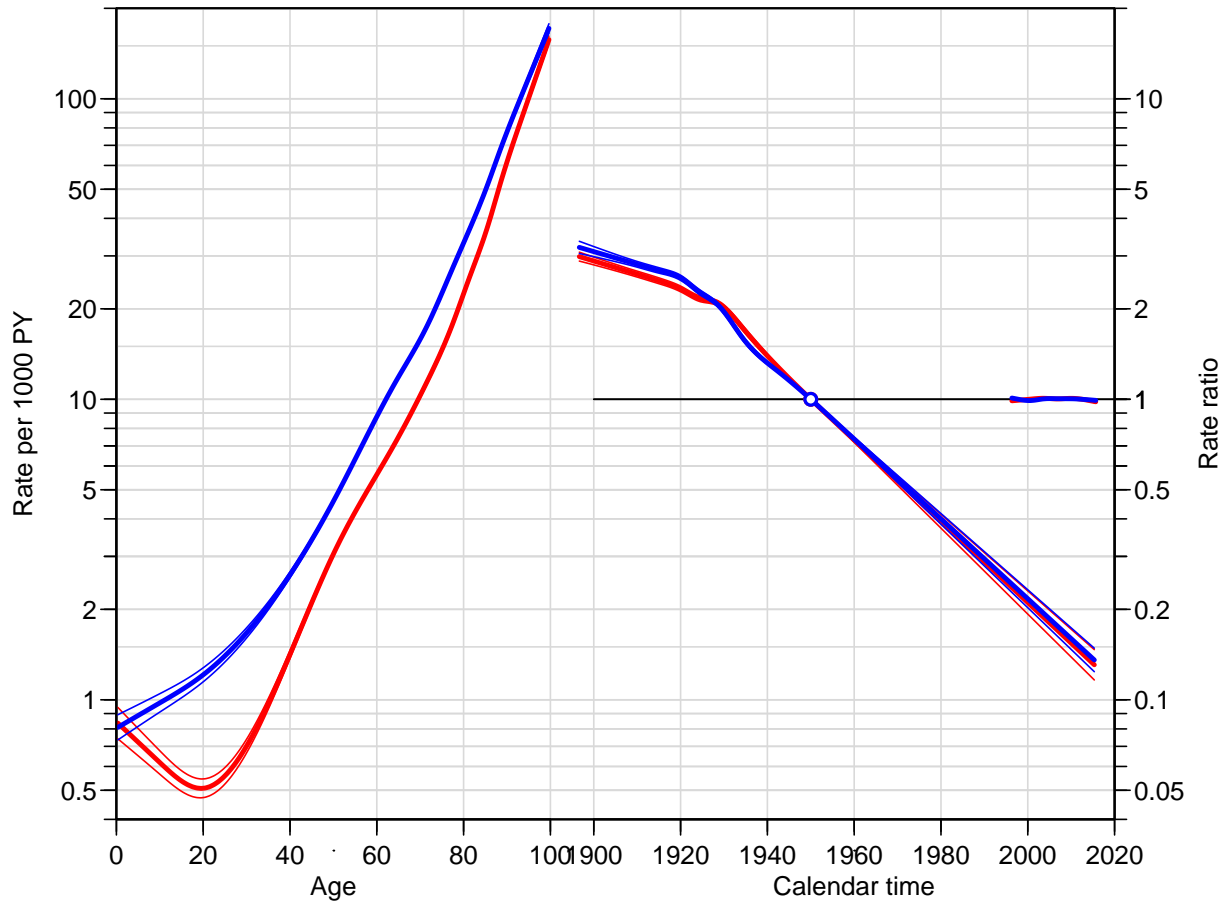


Figure 3.8: Estimates from an APC-model for mortality among non-diabetic individuals in Denmark 1995–2011 (original definition of DM), cohort effects constrained to be 1 at 1950, period slope to be 0. Blue: Men; red: Women.

We also fit using the period-major parametrization:

```

> nDapcM <- apc.fit( subset(nD,sex=="M"),
+                   ref.p=2000,
+                   parm="APC",
+                   npar=list(A=A.nD.kn,P=P.nD.kn,C=C.nD.kn) )
[1] "ML of APC-model Poisson with log(Y) offset : ( APC ):\n"

```

Analysis of deviance for Age-Period-Cohort model

	Resid. Df	Resid. Dev	Df	Deviance	Pr(>Chi)
Age	3990	18794.7			
Age-drift	3989	7763.3	1	11031.4	< 2.2e-16 ***
Age-Cohort	3983	6581.4	6	1181.9	< 2.2e-16 ***
Age-Period-Cohort	3979	6568.2	4	13.3	0.01011 *
Age-Period	3985	7709.3	-6	-1141.2	< 2.2e-16 ***
Age-drift	3989	7763.3	-4	-54.0	5.313e-11 ***

```

---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
> nDapcF <- apc.fit( subset(nD,sex=="F"),
+                   ref.p=2000,
+                   parm="APC",
+                   npar=list(A=A.nD.kn,P=P.nD.kn,C=C.nD.kn) )
[1] "ML of APC-model Poisson with log(Y) offset : ( APC ):\n"

Analysis of deviance for Age-Period-Cohort model

              Resid. Df Resid. Dev Df Deviance  Pr(>Chi)
Age                3990    13636.1
Age-drift          3989     7014.9  1   6621.3 < 2.2e-16 ***
Age-Cohort         3983     5469.6  6   1545.3 < 2.2e-16 ***
Age-Period-Cohort  3979     5444.9  4    24.7 5.784e-05 ***
Age-Period         3985     6905.4 -6  -1460.5 < 2.2e-16 ***
Age-drift          3989     7014.9 -4   -109.5 < 2.2e-16 ***
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
> par( mar=c(3,4,1,4), mgp=c(3,1,0)/1.6, las=1 )
> plot ( nDapcF, lty=1, ci=TRUE, col="red", r.txt="Rate per 1000 PY" )

cp.offset      RR.fac
      1790         100

> lines( nDapcM, lty=1, ci=TRUE, col="blue" )

```

3.4.2 Mortality among DM patients

Here we use `D.DM` and `Y.DM` as response variables in the analysis of mortality rates among non-diabetics, and again we first need to define the age and period properly:

```

> DM <- TT[,c("sex","A","P","D.DM","Y.DM")]
> names( DM )[4:5] <- c("D","Y")
> DM <- subset( DM, Y>0 )

```

With this groomed data frame in place we can fit separate models for men and women and plot the estimates together:

```

> ( A.DM.kn <- quantile( rep(DM$A      ,DM$D), probs=(1:8-0.5)/8 ) )
  6.25%  18.75%  31.25%  43.75%  56.25%  68.75%  81.25%  93.75%
56.33333 66.33333 71.66667 75.66667 79.66667 83.33333 86.66667 91.66667
> ( P.DM.kn <- quantile( rep(DM$P      ,DM$D), probs=(1:6-0.5)/6 ) )
8.333333%      25% 41.66667% 58.33333%      75% 91.66667%
1998.333 2002.333 2005.667 2009.333 2011.667 2014.667
> ( C.DM.kn <- quantile( rep(DM$P-DM$A,DM$D), probs=(1:8-0.5)/8 ) )
  6.25%  18.75%  31.25%  43.75%  56.25%  68.75%  81.25%  93.75%
1912.667 1919.333 1923.333 1927.333 1931.667 1936.333 1942.333 1951.667
> DMacpM <- apc.fit( subset(DM,sex=="M"),
+                   ref.c=1950,
+                   parm="ACP",
+                   npar=list(A=A.DM.kn,P=P.DM.kn,C=C.DM.kn) )

```

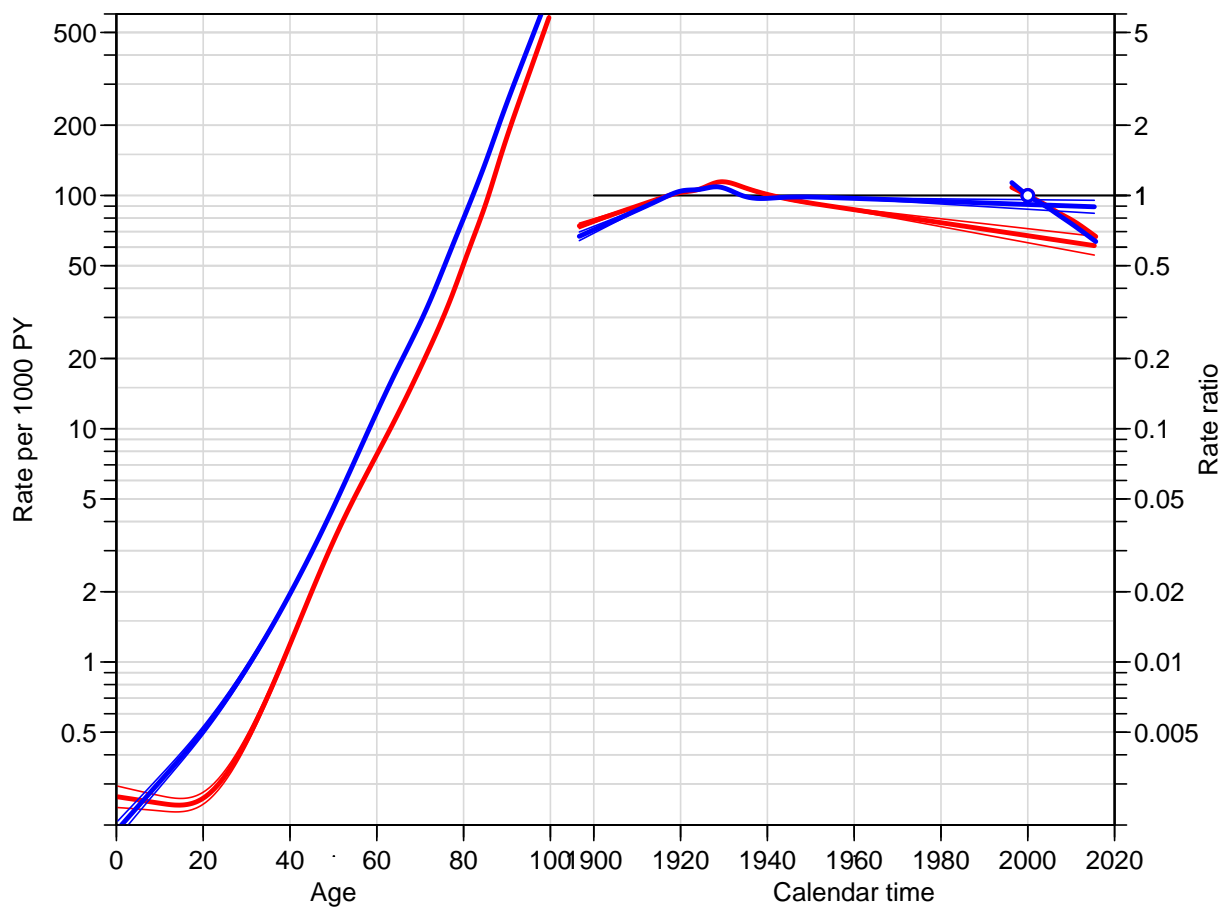


Figure 3.9: Estimates from an APC-model for mortality among non-diabetic individuals in Denmark 1995–2011 (original definition of DM), period constrained to be 1 at 2000, cohort slope to be 0. Blue: Men, red: Women.

```
[1] "ML of APC-model Poisson with log(Y) offset : ( ACP ):\n"
```

Analysis of deviance for Age-Period-Cohort model

	Resid. Df	Resid. Dev	Df	Deviance	Pr(>Chi)
Age	3984	7139.2			
Age-drift	3983	3626.6	1	3512.6	< 2e-16 ***
Age-Cohort	3977	3463.0	6	163.6	< 2e-16 ***
Age-Period-Cohort	3973	3455.1	4	8.0	0.09198 .
Age-Period	3979	3617.6	-6	-162.5	< 2e-16 ***
Age-drift	3983	3626.6	-4	-9.1	0.05967 .

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```
> DMacpF <- apc.fit( subset(DM,sex=="F"),
+                   ref.c=1950,
+                   parm="ACP",
+                   npar=list(A=A.DM.kn,P=P.DM.kn,C=C.DM.kn) )
```

```
[1] "ML of APC-model Poisson with log(Y) offset : ( ACP ):\n"
```

Analysis of deviance for Age-Period-Cohort model

```

                Resid. Df Resid. Dev Df Deviance Pr(>Chi)
Age                3977      5621.6
Age-drift           3976      3499.6  1  2122.00 < 2.2e-16 ***
Age-Cohort          3970      3311.6  6   188.08 < 2.2e-16 ***
Age-Period-Cohort   3966      3296.7  4    14.82  0.005091 **
Age-Period           3972      3485.2 -6  -188.44 < 2.2e-16 ***
Age-drift           3976      3499.6 -4   -14.46  0.005974 **
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
> par( mar=c(3,4,1,4), mgp=c(3,1,0)/1.6, las=1 )
> plot ( DMacpF, lty=1, ci=TRUE, col="red", r.txt="Rate per 1000 PY" )
cp.offset  RR.fac
      1790      10
> lines( DMacpM, lty=1, ci=TRUE, col="blue" )

```

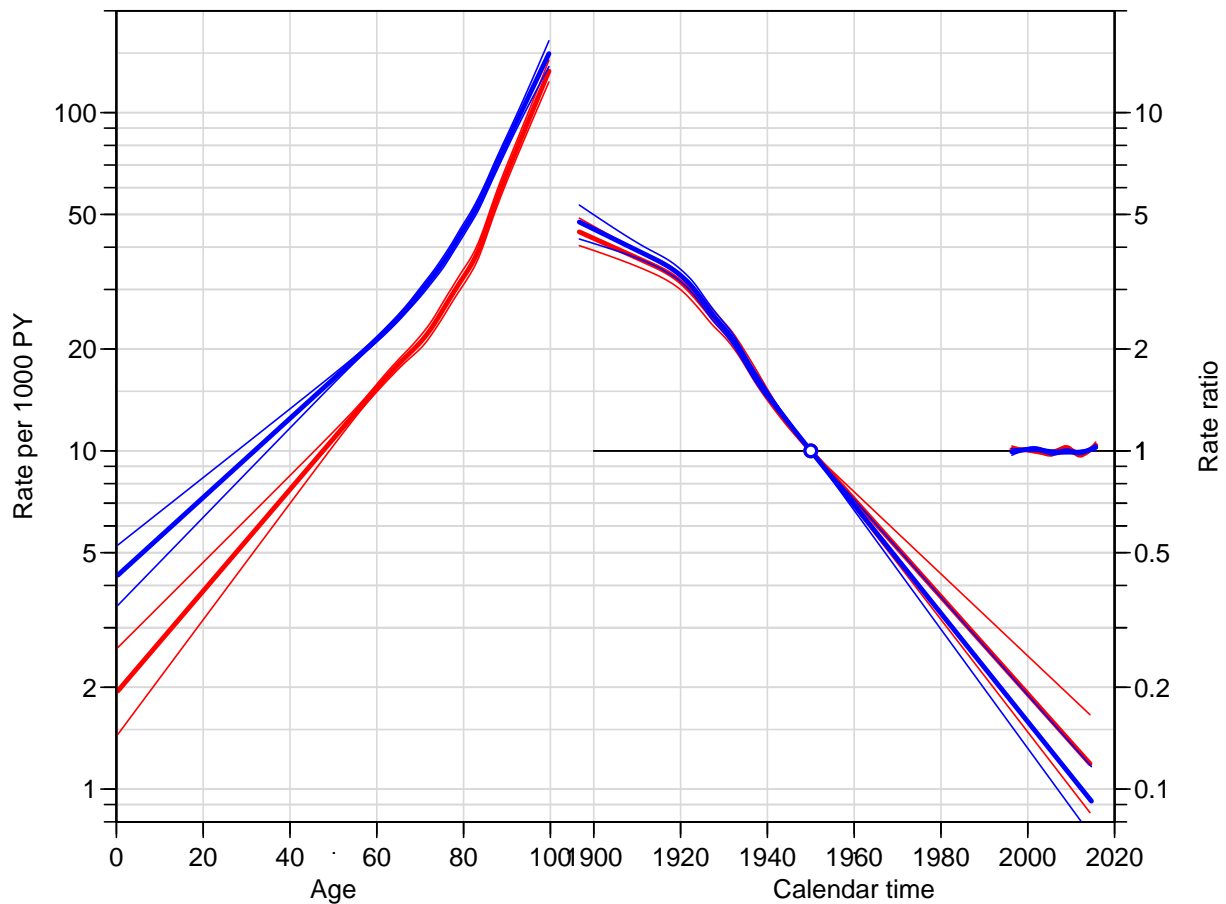


Figure 3.10: *Estimates from an APC-model for mortality among DM patients in Denmark 1995–2011 (original definition), cohort constrained to be 1 at 1950, period slope to be 0. Blue: Men, red: Women.*

We also fit using the period-major parametrization:

```

> DMacpM <- apc.fit( subset(DM,sex=="M"),
+                   ref.p=2000,
+                   parm="APC",
+                   npar=list(A=A.DM.kn,P=P.DM.kn,C=C.DM.kn) )

```

```
[1] "ML of APC-model Poisson with log(Y) offset : ( APC ):\n"
```

```
Analysis of deviance for Age-Period-Cohort model
```

	Resid.	Df	Resid. Dev	Df	Deviance	Pr(>Chi)
Age	3984		7139.2			
Age-drift	3983		3626.6	1	3512.6	< 2e-16 ***
Age-Cohort	3977		3463.0	6	163.6	< 2e-16 ***
Age-Period-Cohort	3973		3455.1	4	8.0	0.09198 .
Age-Period	3979		3617.6	-6	-162.5	< 2e-16 ***
Age-drift	3983		3626.6	-4	-9.1	0.05967 .

```
---
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
> DMapcF <- apc.fit( subset(DM,sex=="F"),
+                   ref.p=2000,
+                   parm="APC",
+                   npar=list(A=A.DM.kn,P=P.DM.kn,C=C.DM.kn) )
```

```
[1] "ML of APC-model Poisson with log(Y) offset : ( APC ):\n"
```

```
Analysis of deviance for Age-Period-Cohort model
```

	Resid.	Df	Resid. Dev	Df	Deviance	Pr(>Chi)
Age	3977		5621.6			
Age-drift	3976		3499.6	1	2122.00	< 2.2e-16 ***
Age-Cohort	3970		3311.6	6	188.08	< 2.2e-16 ***
Age-Period-Cohort	3966		3296.7	4	14.82	0.005091 **
Age-Period	3972		3485.2	-6	-188.44	< 2.2e-16 ***
Age-drift	3976		3499.6	-4	-14.46	0.005974 **

```
---
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

```
> par( mar=c(3,4,1,4), mgp=c(3,1,0)/1.6, las=1 )
> plot ( DMapcF, lty=1, ci=TRUE, col="red", r.txt="Rate per 1000 PY" )
```

```
cp.offset    RR.fac
   1790         100
```

```
> lines( DMapcM, lty=1, ci=TRUE, col="blue" )
```

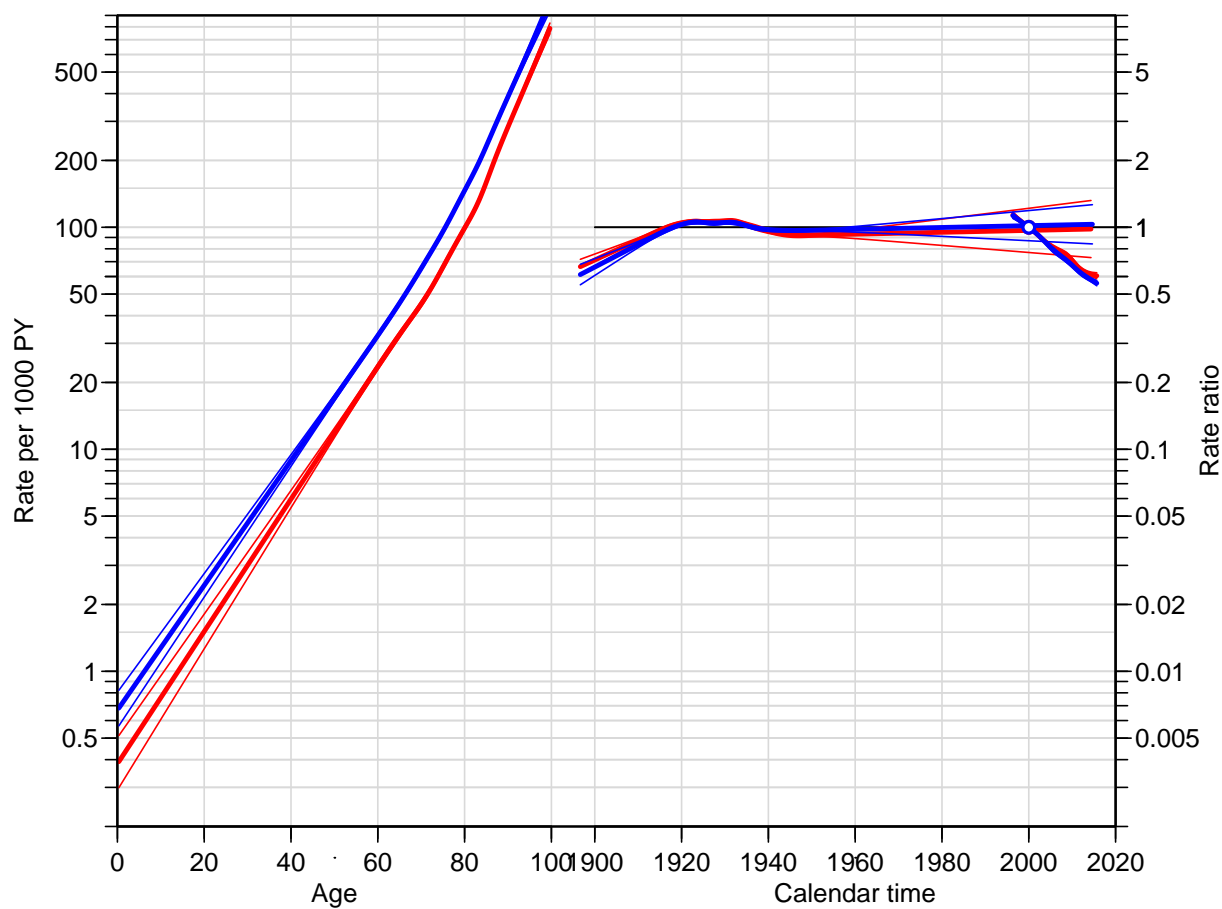


Figure 3.11: *Estimates from an APC-model for mortality among non-diabetic individuals in Denmark 1995–2011 (original definition of DM), period constrained to be 1 at 2000, cohort slope to be 0. Blue: Men, red: Women.*

3.5 Summary of APC models for incidence and mortality

The deviance analysis of the model did not surprisingly show that both cohort and period have non-linear effects, however this formal significance is largely due to the large data base, clearly there is no epidemiologically significant period-effect.

```
> load( file="../data/inc.Rda" )
> par( mfrow=c(3,2), mar=c(0,0,0,0), oma=c(3,4,2,4), mgp=c(3,1,0)/1.6, las=1 )
> frm <- function( side, wh ) {
+   apc.frame( a.lab=seq(0,80,20), a.tic=c(0,seq(10,100,10)),
+             cp.lab=seq(1900,2020,20), cp.tic=seq(1900,2020,10),
+             r.lab=c(outer(c(1,2,5),10^(-1:1)),100), tic.fac=2,
+             r.tic=c(outer(2:9,10^(-1:1)),100,150,200), rr.ref=10,
+             gap=15, r.txt=wh, side=side ) }
> frm( 2, "DM incidence per 1000 PY" )
> lines( acpM, col="blue", ci=TRUE )
> lines( acpF, col="red" , ci=TRUE )
> frm( 4 )
> lines( apcM, col="blue", ci=TRUE )
> lines( apcF, col="red" , ci=TRUE )
> frm( 2, "non-DM mortality per 1000 PY" )
> lines( nDacpM, col="blue", ci=TRUE )
> lines( nDacpF, col="red" , ci=TRUE )
> frm( 4 )
> lines( nDapcM, col="blue", ci=TRUE )
> lines( nDapcF, col="red" , ci=TRUE )
> frm( 1:2, "DM mortality per 1000 PY" )
> lines( DMacpM, col="blue", ci=TRUE )
> lines( DMacpF, col="red" , ci=TRUE )
> frm( c(1,4) )
> lines( DMapcM, col="blue", ci=TRUE )
> lines( DMapcF, col="red" , ci=TRUE )
> mtext( "Cohort major", side=3, line=0.5, at=0.25, outer=TRUE )
> mtext( "Period major", side=3, line=0.5, at=0.75, outer=TRUE )
```

3.5.1 Time-trends in mortality rates

We can extract the timetrends for diabetics and non-diabetics by sex, and print the annual percentwise change:

```
> DA <- NArray( c( list( who = c("non-DM","DM"),
+                         sex = c("M","F") ),
+                 dimnames( nDacpM$Drift ) ) )
> DA["non-DM","M",,] <- nDacpM$Drift
> DA["non-DM","F",,] <- nDacpF$Drift
> DA[  "DM","M",,] <- DMacpM$Drift
> DA[  "DM","F",,] <- DMacpF$Drift
> round( ftable( (DA-1)*100, row.vars=1:2 ), 1 )
              APC (D-weights)                A-d
              exp(Est.) 2.5% 97.5% exp(Est.) 2.5% 97.5%
who  sex
non-DM M      -2.9 -2.9 -2.8          -2.6 -2.7 -2.6
      F      -2.4 -2.5 -2.4          -2.0 -2.0 -1.9
DM    M      -3.7 -3.8 -3.6          -3.6 -3.7 -3.5
      F      -3.4 -3.5 -3.2          -3.1 -3.2 -2.9
```

We see that there is not much difference in the overall trend between man and women, but there seem to be a substantially steeper decrease in mortality among diabetes patients than among persons without diabetes.

```
> library( Epi )  
> library( splines )
```

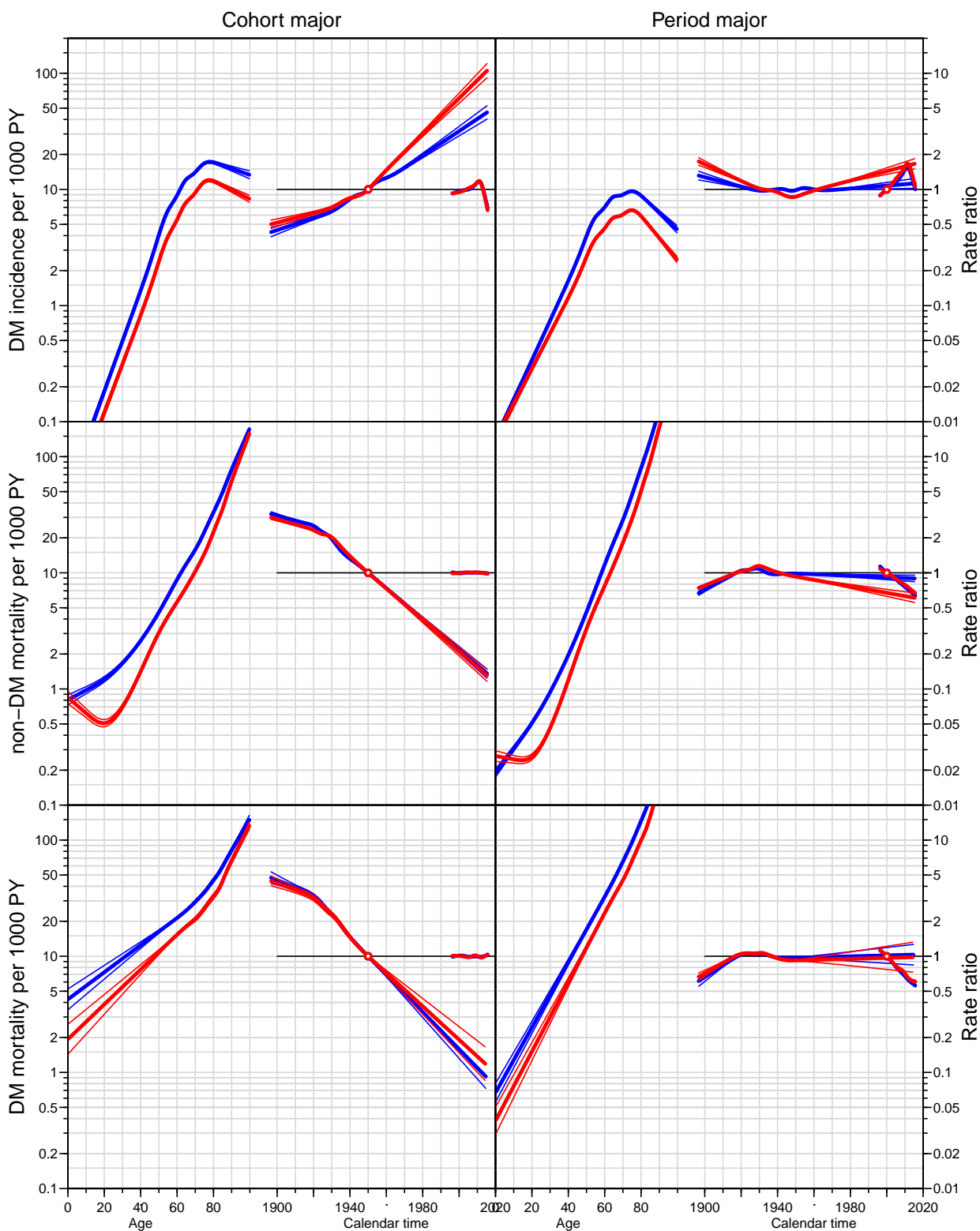



Figure 3.12: Age-Period-Cohort models for mortality among men (blue) and women (red). Top panel is the mortality among non-diabetics and the lower panel is the mortality among diabetes patients.

Chapter 4

Prediction of rates

Despite the previous section's modeling of incidence and mortality rates by APC-models we re-fit the models for prediction purposes.

4.1 Analysis date

We model the incidence and mortality rates from a tabulation of the diabetes register made in conjunction with the annual update of the register. The analytical units are the Lexis triangles, classified by age, period and cohort (∇ and \triangleleft):

```
> load( '../data/inits.Rda' )
> load( '../data/TT.Rda' )
> # convert to person years from person-millenia as used in the SAS-programs
> TT[,c("Y.nD", "Y.DM")] <- TT[,c("Y.nD", "Y.DM")]*1000
> TT <- subset( TT, P > 1996 )
> str( TT )
'data.frame':      8000 obs. of  9 variables:
 $ sex : Factor w/ 2 levels "M","F": 1 1 1 1 1 1 1 1 1 1 ...
 $ A   : num  0.333 0.667 0.333 0.667 0.333 ...
 $ P   : num  1997 1996 1998 1997 1999 ...
 $ U   : num  0 1 0 1 0 1 0 1 0 1 ...
 $ X   : num  0 1 1 1 2 2 2 2 1 3 ...
 $ D.nD: num  0 28 1 18 2 18 0 11 0 21 ...
 $ Y.nD: num  17570 17884 17726 17321 17116 ...
 $ D.DM: num  0 0 0 0 0 0 0 0 0 0 ...
 $ Y.DM: num  0 0.476 0.461 0.17 0.927 ...
> head( TT )
  sex      A      P U X D.nD      Y.nD D.DM      Y.DM
1  M 0.3333333 1996.667 0 0  0 17569.79  0 0.0000000
2  M 0.6666667 1996.333 1 1  28 17883.86  0 0.4757016
3  M 0.3333333 1997.667 0 1  1 17725.66  0 0.4613279
4  M 0.6666667 1997.333 1 1  18 17320.57  0 0.1704312
5  M 0.3333333 1998.667 0 2  2 17116.36  0 0.9267625
6  M 0.6666667 1998.333 1 2  18 17093.26  0 0.7015743
> summary( TT )
sex      A      P      U      X      D.nD
M:4000  Min.   : 0.3333  Min.   :1996  Min.   :0.0  Min.   : 0.0  Min.   : 0
F:4000  1st Qu.:25.1667  1st Qu.:2001  1st Qu.:0.0  1st Qu.: 6.0  1st Qu.: 7
        Median :50.0000  Median :2006  Median :0.5  Median :21.0  Median :51
        Mean   :50.0000  Mean   :2006  Mean   :0.5  Mean   :41.2  Mean   :119
```

```

3rd Qu.:74.8333 3rd Qu.:2011 3rd Qu.:1.0 3rd Qu.: 68.0 3rd Qu.:205
Max. :99.6667 Max. :2016 Max. :1.0 Max. :280.0 Max. :590
Y.nD D.DM Y.DM
Min. : 22.35 Min. : 0.00 Min. : 0.00
1st Qu.: 8509.06 1st Qu.: 0.00 1st Qu.: 67.03
Median :16058.13 Median : 5.00 Median : 243.74
Mean :13225.82 Mean : 18.92 Mean : 419.75
3rd Qu.:17903.57 3rd Qu.: 33.00 3rd Qu.: 638.96
Max. :23342.36 Max. :116.00 Max. :2705.43
> table( TT$Y.DM==0,TT$Y.nD==0 )
FALSE
FALSE 7977
TRUE 23

```

We see from the tabulation that we truly have data in Lexis-triangles:

```

> with( subset( TT, A<5 & P<1999 ),
+ print( table( Age=round(A,2),
+ Per=round(P,2) ),
+ zero.print="." ) )
Per
Age 1996.33 1996.67 1997.33 1997.67 1998.33 1998.67
0.33 . 2 . 2 . 2
0.67 2 . 2 . 2 .
1.33 . 2 . 2 . 2
1.67 2 . 2 . 2 .
2.33 . 2 . 2 . 2
2.67 2 . 2 . 2 .
3.33 . 2 . 2 . 2
3.67 2 . 2 . 2 .
4.33 . 2 . 2 . 2
4.67 2 . 2 . 2 .

```

A brief overview of the number of events and PY:

```

> tt <- xtabs( cbind(X,D.nD,D.DM,Y.nD,Y.DM) ~ sex + floor(P),
+ data = TT )
> round( ftable( addmargins(tt,1:2), row.vars=1:2 ) )
sex floor(P) X D.nD D.DM Y.nD Y.DM
M 1996 6602 26894 2958 2551813 47789
1997 6528 26205 3069 2559636 51223
1998 6991 25607 3124 2565493 54920
1999 7468 25235 3415 2570306 58725
2000 7414 24463 3406 2575532 62791
2001 7491 24655 3504 2582024 66845
2002 7720 24359 3747 2587849 70815
2003 8785 24141 3872 2591456 75133
2004 9061 23316 3871 2593652 80158
2005 8864 22600 3947 2597064 85083
2006 9228 22863 4124 2602379 90126
2007 9772 22646 4242 2611223 95272
2008 10628 22224 4280 2624040 101316
2009 11246 21934 4763 2635748 107637
2010 12188 21689 4788 2644351 114590
2011 13907 20833 4928 2651824 122766
2012 12145 20419 5141 2659128 130855
2013 9267 20523 5290 2670097 136249

```

	2014	8867	19938	5470	2688000	139869
	2015	9912	20296	5532	2689703	142459
	Sum	184084	460840	83471	52251318	1834623
F	1996	5413	27411	2661	2620753	43025
	1997	5277	27274	2728	2628018	45520
	1998	5501	26407	2765	2633905	48237
	1999	5811	27243	2873	2638757	50976
	2000	5862	26260	2971	2644189	53927
	2001	6012	26533	3045	2650333	56871
	2002	6010	26784	3110	2655795	59848
	2003	6939	25872	3253	2658828	62949
	2004	7057	24718	3124	2661464	66811
	2005	7196	24459	3323	2664904	70570
	2006	7276	24431	3424	2669731	74431
	2007	8007	24679	3566	2676849	78479
	2008	8481	23711	3484	2687706	83325
	2009	8610	23783	3710	2699204	88124
	2010	9365	23162	3884	2708667	93337
	2011	11220	22341	3802	2716290	99759
	2012	9804	22023	3837	2722084	106525
	2013	7309	21752	3981	2731402	111039
	2014	6648	21080	4063	2745803	113931
	2015	7681	21276	4271	2740586	115710
	Sum	145479	491199	67875	53555267	1523395
Sum	1996	12015	54305	5619	5172566	90814
	1997	11805	53479	5797	5187655	96743
	1998	12492	52014	5889	5199397	103158
	1999	13279	52478	6288	5209063	109701
	2000	13276	50723	6377	5219720	116717
	2001	13503	51188	6549	5232356	123717
	2002	13730	51143	6857	5243644	130663
	2003	15724	50013	7125	5250284	138082
	2004	16118	48034	6995	5255116	146969
	2005	16060	47059	7270	5261968	155653
	2006	16504	47294	7548	5272110	164557
	2007	17779	47325	7808	5288072	173752
	2008	19109	45935	7764	5311746	184641
	2009	19856	45717	8473	5334951	195761
	2010	21553	44851	8672	5353018	207927
	2011	25127	43174	8730	5368114	222524
	2012	21949	42442	8978	5381212	237380
	2013	16576	42275	9271	5401500	247288
	2014	15515	41018	9533	5433802	253800
	2015	17593	41572	9803	5430289	258170
	Sum	329563	952039	151346	105806585	3358018

```

> tt <- xtabs( cbind(X,D.nD,Y.nD,D.DM,Y.DM) ~ sex + gP,
+             data = transform( subset( TT, P > 1996 ),
+                               gP = factor( (P>=2001)+(P>=2006)+(P>=2011),
+                                           labels=c("1996-2000",
+                                                  "2001-2005",
+                                                  "2006-2010",
+                                                  "2011-2015") ) ) )
> round( ftable( addmargins(tt,1:2), row.vars=1:2 ) )

```

		X	D.nD	Y.nD	D.DM	Y.DM
sex	gP					
M	1996-2000	35003	128404	12822780	15972	275448
	2001-2005	41921	119071	12952045	18941	378035
	2006-2010	53062	111356	13117741	22197	508942

	2011–2015	54098	102009	13358752	26361	672198
	Sum	184084	460840	52251318	83471	1834623
F	1996–2000	27864	134595	13165621	13998	241685
	2001–2005	33214	128366	13291324	15855	317049
	2006–2010	41739	119766	13442157	18068	417696
	2011–2015	42662	108472	13656165	19954	546964
	Sum	145479	491199	53555267	67875	1523395
Sum	1996–2000	62867	262999	25988401	29970	517133
	2001–2005	75135	247437	26243369	34796	695084
	2006–2010	94801	231122	26559898	40265	926638
	2011–2015	96760	210481	27014917	46315	1219162
	Sum	329563	952039	105806585	151346	3358018

We set up arrays to hold the predicted incidence and mortality rates from the different models, separately for the two sexes:

```
> Lambda <- Mu.W <- Mu.DM <- NArray( list( a = a.pt,
+                                       p = p.pt,
+                                       sex = c("M", "F"),
+                                       mod = c("ap", "apc", "att", "i20", "i25", "i30") ) )
> str( Lambda )
logi [1:1200, 1:528, 1:2, 1:6] NA NA NA NA NA NA ...
- attr(*, "dimnames")=List of 4
..$ a : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.2916666666666667"
..$ p : chr [1:528] "1996.041666666667" "1996.125" "1996.208333333333" "1996.291666666667"
..$ sex: chr [1:2] "M" "F"
..$ mod: chr [1:6] "ap" "apc" "att" "i20" ...
> prod( dim(Lambda) )
[1] 7603200
```

Note that we have a level on dimension 4, `att`, designed to hold predictions where we attenuate the linear trajectories from the period and cohort terms.

4.1.1 Datasets for rate modeling

First we construct simple datasets for APC and Lee-Carter analysis separately for the three types of transitions we are going to consider.

We can explore the entire set of models for men and women separately using the `apc.LCa` function:

```
> incdat <- TT[,c("A", "P", "X", "Y.nD", "sex")]
> mnDdat <- TT[,c("A", "P", "D.nD", "Y.nD", "sex")]
> mDMdat <- TT[,c("A", "P", "D.DM", "Y.DM", "sex")]
> names( incdat )[3:4] <-
+ names( mnDdat )[3:4] <-
+ names( mDMdat )[3:4] <-c("D", "Y")
> mDMdat <- subset( mDMdat, Y>0 )
```

4.2 Models for incidence and mortality rates

There are two immediate alternatives to the age-period-cohort models on the log-scale; age-period-cohort models on the additive scale and Lee-Carter models. These may provide a better fit to data (some of the expanded Lee-Carter models necessarily do).

```

> minc <- apc.LCa( subset(incdat,sex=="M"), eps=1e-4 )
> finc <- apc.LCa( subset(incdat,sex=="F"), eps=1e-4 )
> mmnD <- apc.LCa( subset(mnDdat,sex=="M"), eps=1e-4 )
> fmnD <- apc.LCa( subset(mnDdat,sex=="F"), eps=1e-4 )
> mmDM <- apc.LCa( subset(mDMdat,sex=="M"), eps=1e-4 )
> fmDM <- apc.LCa( subset(mDMdat,sex=="F"), eps=1e-4 )
> save( incdat, mnDdat, mDMdat,
+       minc, finc, mmnD, fmnD, mmDM, fmDM,
+       file = "imdat.Rda" )

```

Once we have fitted all models for all transitions we can show their relative merits in terms of deviance, and assess to what extent the APC models would suffice or whether substantial improvements could be obtained by using a Lee-Carter model:

```

> load( file = "imdat.Rda" )
> par( mfcol=c(3,2) )
> show.apc.LCa( minc, col.txt="blue" ); text( 10, 90, "DM\n incidence", cex=2 )
> show.apc.LCa( mmDM, col.txt="blue" ); text( 10, 90, "DM\n mortality", cex=2 )
> show.apc.LCa( mmnD, col.txt="blue" ); text( 10, 90, "non-DM\n mortality", cex=2 )
> show.apc.LCa( finc, col.txt="red" )
> show.apc.LCa( fmDM, col.txt="red" )
> show.apc.LCa( fmnD, col.txt="red" )

```

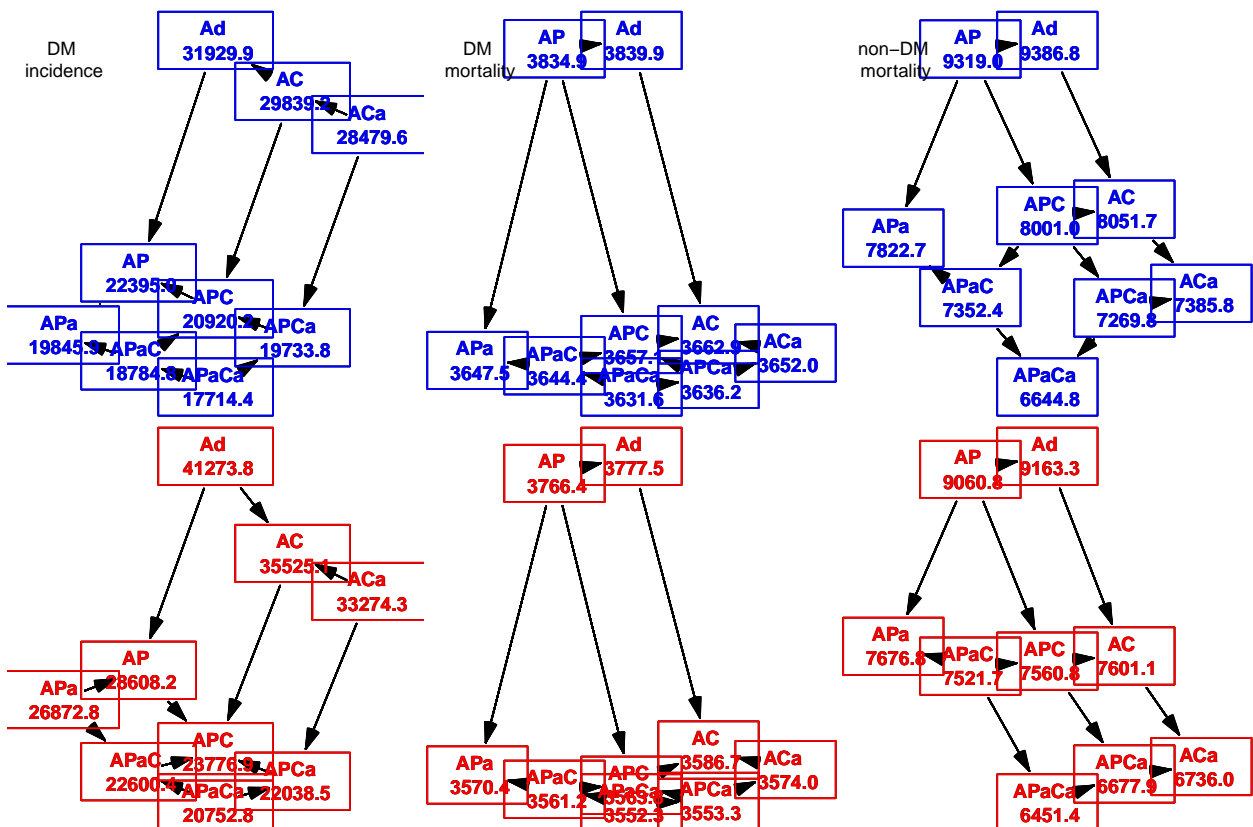


Figure 4.1: Relative fit of APC and Lee-Carter models for incidence and mortality rates for men and women respectively.

From the relative model fits shown in figure 4.1 we see that for diabetes incidence and mortality the APC is quite well fitting relative to the classical Lee-Carter model (APa),

but that the Lee-Carter type models offer some improvement relative to the APC model for non-DM mortality but only if cohort-age effect is included.

Thus, we shall proceed with APC-models for the incidence and mortality rates.

4.3 Incidence rates

Based on this we can now derive the location of the knots for this model (first loading the prerequisites):

```
> load( file="../data/inits.Rda" )
> ( ki.a <- with( incdat, quantile( rep( A,D), qn(nk.a) ) ) )
  6.25%  18.75%  31.25%  43.75%  56.25%  68.75%  81.25%  93.75%
35.33333 47.66667 54.33333 59.66667 64.33333 68.66667 74.33333 82.66667
> ( ki.p <- with( incdat, quantile( rep( P ,D), qn(nk.p) ) ) )
8.333333%      25% 41.66667% 58.33333%      75% 91.66667%
1998.333  2002.333  2005.667  2009.333  2011.667  2014.333
> ( ki.c <- with( incdat, quantile( rep( P-A,D), qn(nk.c) ) ) )
  6.25%  18.75%  31.25%  43.75%  56.25%  68.75%  81.25%  93.75%
1922.333 1931.667 1938.333 1943.333 1947.667 1953.333 1960.333 1972.667
```

The model we set up is an age-period-cohort model with these three terms in it. As we are only going to use the model for predictions we need not bother about parametrization issues, so it is not an issue that the model we fit is formally over-parametrized. However we want to extract the average trend from the APC-model, so we also fit the model with the parametrization that allows us to extract the trend.

```
> m.inc.ap <- glm( D ~ Ns(A,knots=ki.a) + Ns(P,knots=ki.p),
+               offset = log( Y ),
+               family = poisson,
+               data = subset(incdat,sex=="M") )
> m.inc.aPC <- update( m.inc.ap, . ~ . - Ns(P ,kn=ki.p) + I(P) +
+                 detrend( Ns(P ,kn=ki.p), P , D ) +
+                 detrend( Ns(P-A,kn=ki.c), P-A, D ) )
> m.inc.apc <- update( m.inc.ap, . ~ . + Ns(P-A,kn=ki.c) )
> c( m.inc.apc$deviance, m.inc.aPC$deviance )
[1] 11179.21 11179.21
> f.inc.ap <- update( m.inc.ap , data = subset(incdat,sex=="F") )
> f.inc.apc <- update( m.inc.apc, data = subset(incdat,sex=="F") )
> f.inc.aPC <- update( m.inc.aPC, data = subset(incdat,sex=="F") )
```

The average annual trends in incidence from the multiplicative models:

```
> inc.chg <- rbind( ci.exp(m.inc.aPC,subset="I\\(P)"-1,
+                       ci.exp(f.inc.aPC,subset="I\\(P)"-1)*100
> rownames( inc.chg ) <- c("DM incidence change      Men",
+                          "                          Women")
> round( inc.chg, 1 )
                                exp(Est.) 2.5% 97.5%
DM incidence change      Men      2.2  2.1  2.2
                          Women    2.6  2.5  2.7
```

The average increase is higher for women than for men, but the period effect is massively non-linear, so these summary figures are not really informative, see the comparative figure with the mortality rates, ??

4.3.1 Incidence rate predictions

Finally we need the predicted incidence rates at a grid of points suitable for the calculations of predicted prevalences. We make the predictions for all combinations of `a.pt` and `p.pt`.

However, all the predictions should be in units of the interval length chosen for calculations. We note from the calculations above that the quantities that enter the expressions for the transition probabilities are all cumulative rates over the intervals. Thus we use a prediction data frame with the person-years-variables set to `int`,

```
> nd <- data.frame( A = rep(a.pt,      length(p.pt)),
+                  P = rep(p.pt,each=length(a.pt)),
+                  Y = int )
> head( nd )
      A      P      Y
1 0.04166667 1996.042 0.08333333
2 0.12500000 1996.042 0.08333333
3 0.20833333 1996.042 0.08333333
4 0.29166667 1996.042 0.08333333
5 0.37500000 1996.042 0.08333333
6 0.45833333 1996.042 0.08333333
> dim( nd )
[1] 633600      3
```

Note that the prediction data frame was set up with age varying fastest, and the `Lambda` array with age before period, so that the column-major storage of arrays conforms with the predictions form `nd`:

```
> Lambda[,,"M","ap" ] <- predict.glm( m.inc.ap , type="response", newdata=nd )
> Lambda[,,"F","ap" ] <- predict.glm( f.inc.ap , type="response", newdata=nd )
> Lambda[,,"M","apc" ] <- predict.glm( m.inc.apc, type="response", newdata=nd )
> Lambda[,,"F","apc" ] <- predict.glm( f.inc.apc, type="response", newdata=nd )
```

Thus we have the incidence rates that we need for two sexes, and for two different modeling approaches.

4.3.2 A damping extrapolation

Instead of using the naive extrapolation we may dampen the trend derived from the naive application of the natural splines. In the vein of the recommendation by Sasieni [?] we use an attenuation of the trend beyond 2016 for period and beyond 2000 for cohort. The attenuation is traditionally set to 0.92 per year, but we shall be more conservative.

The arithmetic goes as follows: Suppose the slope of the period or cohort effect is β , but that it would be an exaggeration to continue the period effect indefinitely at a slope of β , so we choose a *damping* factor, d , say, such that the slope of the effect at tie t is not β , but rather βd^t . Thus for the effect $f(t)$ we have that:

$$f'(t) = \beta d^t \quad \Leftrightarrow \quad f(t) = k + \beta d^t / \log(d) \quad \Rightarrow \quad f(0) = k + \beta / \log(d)$$

Now, if the original linear extrapolation we have from the natural spline parametrization has the form:

$$f(t) = \alpha + \beta t$$

then at the prediction start (for convenience of notation, $t = 0$):

$$f(0) = \alpha = k + \beta / \log(d) \quad \Leftrightarrow \quad k = \alpha - \beta / \log(d)$$

These formulae are implemented in the function `damp`, which assumes that the predicted effects at times `t` are in `fval`, and that the effects are linear beyond `t0`, but that we attenuate the effect with the damping factor `damp`, in cancer epidemiology by default often set to 0.92, but we shall be a bit more conservative and set it to 0.88 (which also is a figure taken out of thin air):

```
> damp <-
+ function( fval, t, t0, dfac=0.88 )
+   {
+ # where are the earliest an latest prediction point after t0
+ wh <- match( range( t[t>t0] ), t )
+ # slope of predicted curve - we assume linearity
+ beta <- diff( fval[wh] ) / diff( t[wh] )
+ # the attenuated curve - revision beyond first point after t0:
+ kons <- fval[wh[1]] - beta/log(dfac)
+ ifelse( t>t[wh[1]], kons + beta * dfac^(t-t[wh[1]]) / log( dfac ), fval )
+   }
> t <- seq( 0,30,,100 )
> f <- 2.5 + 0.4 * t
> t0 <- 16
> matplot( t, cbind( f, damp(f,t,5) ), lty=1, lwd=c(8,2), type="l" )
> abline( v= 5 )
```

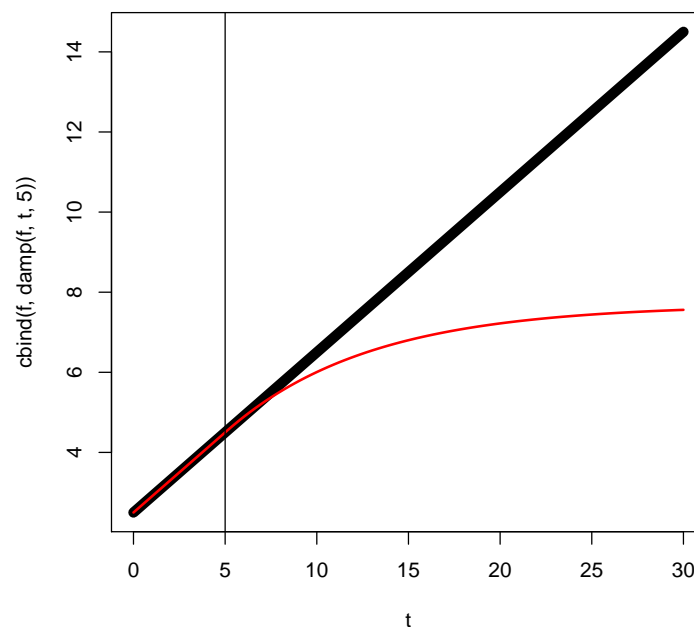


Figure 4.2: *Illustration of the damp function for attenuation of linear effects.*

We can now employ this damping function on the terms-predictions from the apc-models, and show how it pans out in practice for the period and cohort effects. We adjust the

predictions for the period and cohort terms and use these to reconstruct the predicted (cumulative) rates, note in particular that the intercept is not in the terms components, it must be retrieved from the attribute `const` in the `predict.glm` object:

```
> predatt.apc <-
+ function( obj, nd, dfac=0.88, p.end=2016, c.end=2000 )
+ {
+ # predicted terms --- linear beyond outer knots
+ pc <- po <- predict.glm( obj, type="terms", newdata=nd )
+ # replace with the attenuated curves
+ pc[,2] <- damp( pc[,2], nd$P      , p.end, dfac=dfac )
+ pc[,3] <- damp( pc[,3], nd$P-nd$A, c.end, dfac=dfac )
+ # return predicted rates as well as original and revised terms predictions
+ list( pr.resp = exp( apply( pc, 1, sum ) + attr(pc,"const") + log(nd$Y) ),
+       pr.terms = pc,
+       pr.org   = po )
+ }
```

We have observed that the incidence rates show a decreasing tendency over the last few years of observation, hence we may want not only to investigate a scenario where rates are kept constant, but also one where we increase the period effect by 2% per year — this is only going to be used for the incidence rates as a sensitivity analysis, and together with the

```
> predinc.apc <-
+ function( obj, nd, dfac=0.025, p.end=2016, c.end=2000 )
+ {
+ # predicted terms --- linear beyond outer knots
+ pc <- po <- predict.glm( obj, type="terms", newdata=nd )
+ # replace with the revised curves
+ mxp <- match( max( nd$P[nd$P<p.end] ), nd$P )
+ pc[,2] <- ifelse( nd$P>p.end, pc[mxp,2] + pmax(0, (nd$P-p.end)*dfac), pc[,2] )
+ # return predicted rates as well as original and revised terms predictions
+ list( pr.resp = exp( apply( pc, 1, sum ) + attr(pc,"const") + log(nd$Y) ),
+       pr.terms = pc,
+       pr.org   = po )
+ }
```

We can then illustrate the two variants for the terms

```
> zz <- predatt.apc( m.inc.apc, nd )
> ww <- predinc.apc( m.inc.apc, nd )
> str( zz )
```

List of 3

```
$ pr.resp : Named num [1:633600] 5.50e-06 5.54e-06 5.57e-06 5.61e-06 5.64e-06 ...
..- attr(*, "names")= chr [1:633600] "1" "2" "3" "4" ...
$ pr.terms: num [1:633600, 1:3] -3.28 -3.27 -3.27 -3.26 -3.25 ...
..- attr(*, "dimnames")=List of 2
.. ..$ : chr [1:633600] "1" "2" "3" "4" ...
.. ..$ : chr [1:3] "Ns(A, knots = ki.a)" "Ns(P, knots = ki.p)" "Ns(P - A, kn = ki.c)"
..- attr(*, "constant")= num -6.09
$ pr.org : num [1:633600, 1:3] -3.28 -3.27 -3.27 -3.26 -3.25 ...
..- attr(*, "dimnames")=List of 2
.. ..$ : chr [1:633600] "1" "2" "3" "4" ...
.. ..$ : chr [1:3] "Ns(A, knots = ki.a)" "Ns(P, knots = ki.p)" "Ns(P - A, kn = ki.c)"
..- attr(*, "constant")= num -6.09
> str( nd )
```

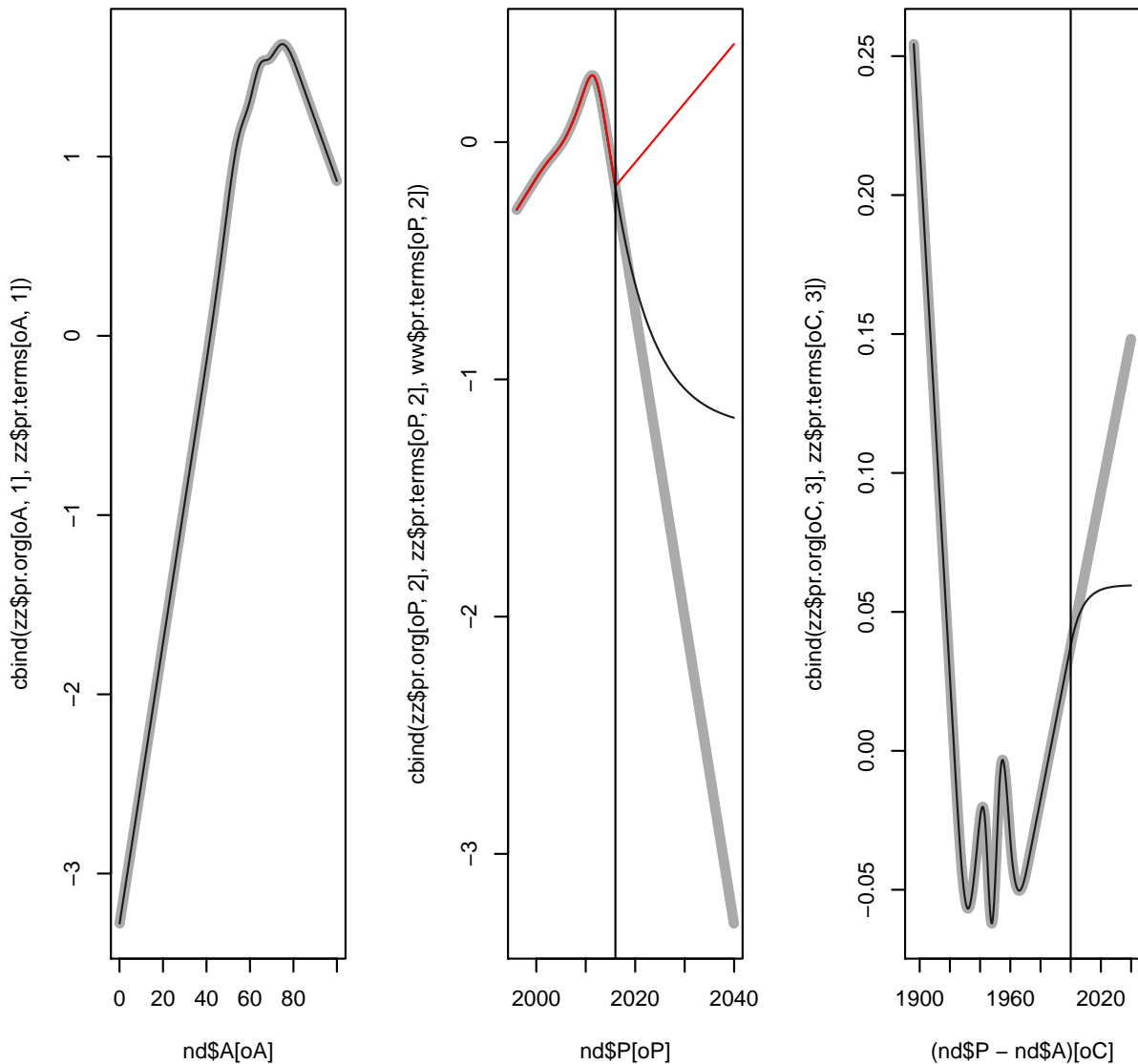



Figure 4.3: *Demonstration of the attenuation of the period and cohort terms from an APC-model. The attenuation effect used is the magic number 0.88 per year; that is, the slope of the effects decreases by a factor 0.88 per year, corresponding to a 47% decrease at 5 years and 72% decrease at 10 years in the slope of the effects. The red line is an annual increase *i* periodo effect of 2.5% intended as a sensitivity analysis tool.*

```
> m.md.apc <- update( m.md.ap, . ~ . + Ns(P-A, kn=kmd.c) )
> f.md.ap <- update( m.md.ap, data = subset( mDMdat, sex=="F" ) )
> f.md.apc <- update( m.md.apc, data = subset( mDMdat, sex=="F" ) )
> f.md.aPC <- update( m.md.aPC, data = subset( mDMdat, sex=="F" ) )
> Mu.DM[, "M", "ap" ] <- predict.glm( m.md.ap, type="response", newdata=nd )
> Mu.DM[, "F", "ap" ] <- predict.glm( f.md.ap, type="response", newdata=nd )
> Mu.DM[, "M", "apc" ] <- predict.glm( m.md.apc, type="response", newdata=nd )
> Mu.DM[, "F", "apc" ] <- predict.glm( f.md.apc, type="response", newdata=nd )
```

```

> Mu.DM[,,"M","att"] <- predatt.apc( m.md.apc, nd )$pr.resp
> Mu.DM[,,"F","att"] <- predatt.apc( f.md.apc, nd )$pr.resp
> Mu.DM[,,"M","i20"] <- predinc.apc( m.md.apc, nd, dfac=0.020 )$pr.resp
> Mu.DM[,,"F","i20"] <- predinc.apc( f.md.apc, nd, dfac=0.020 )$pr.resp
> Mu.DM[,,"M","i25"] <- predinc.apc( m.md.apc, nd, dfac=0.025 )$pr.resp
> Mu.DM[,,"F","i25"] <- predinc.apc( f.md.apc, nd, dfac=0.025 )$pr.resp
> Mu.DM[,,"M","i30"] <- predinc.apc( m.md.apc, nd, dfac=0.030 )$pr.resp
> Mu.DM[,,"F","i30"] <- predinc.apc( f.md.apc, nd, dfac=0.030 )$pr.resp

```

4.4.2 Persons without diabetes

The mortality in the population without diabetes is modeled in exactly the same way, except we put in knots early in age:

```

> ( kmw.a <- with( mnDdat, c( 5, 15,
+                          quantile( rep(A ,D), qn(nk.a) ) ) ) )
      6.25%  18.75%  31.25%  43.75%  56.25%  68.75%  81.25%  93.75%
5.00000 15.00000 50.66667 64.33333 71.66667 77.33333 81.33333 85.33333 88.66667 93.66667
> ( kmw.p <- with( mnDdat, quantile( rep(P ,D), qn(nk.p) ) ) )
8.333333%  25% 41.66667% 58.33333%  75% 91.66667%
1997.333 2000.667 2003.667 2006.667 2010.333 2014.333
> ( kmw.c <- with( mnDdat, quantile( rep(P-A,D), qn(nk.c) ) ) )
      6.25%  18.75%  31.25%  43.75%  56.25%  68.75%  81.25%  93.75%
1909.667 1915.667 1920.333 1924.333 1928.667 1934.333 1942.667 1955.667
> m.mw.ap <- glm( D ~ Ns(A,knots=kmw.a) + Ns(P,knots=kmw.p),
+               offset = log(Y),
+               family = poisson,
+               data = subset( mnDdat, sex=="M" ) )
> m.mw.aPC <- update( m.mw.ap, . ~ . - Ns(P ,kn=kmw.p) + I(P) +
+                   detrend( Ns(P ,kn=kmw.p), P , D ) +
+                   detrend( Ns(P-A,kn=kmw.c), P-A, D ) )
> m.mw.apc <- update( m.mw.ap, . ~ . + Ns(P-A,kn=kmw.c) )
> f.mw.ap <- update( m.mw.ap , data = subset( mnDdat, sex=="F" ) )
> f.mw.apc <- update( m.mw.apc, data = subset( mnDdat, sex=="F" ) )
> f.mw.aPC <- update( m.mw.aPC, data = subset( mnDdat, sex=="F" ) )
> Mu.W[,,"M","ap" ] <- predict.glm( m.mw.ap , type="response", newdata=nd )
> Mu.W[,,"F","ap" ] <- predict.glm( f.mw.ap , type="response", newdata=nd )
> Mu.W[,,"M","apc" ] <- predict.glm( m.mw.apc, type="response", newdata=nd )
> Mu.W[,,"F","apc" ] <- predict.glm( f.mw.apc, type="response", newdata=nd )
> Mu.W[,,"M","att" ] <- predatt.apc( m.mw.apc, nd )$pr.resp
> Mu.W[,,"F","att" ] <- predatt.apc( f.mw.apc, nd )$pr.resp
> Mu.W[,,"M","i20" ] <- predinc.apc( m.mw.apc, nd, dfac=0.020 )$pr.resp
> Mu.W[,,"F","i20" ] <- predinc.apc( f.mw.apc, nd, dfac=0.020 )$pr.resp
> Mu.W[,,"M","i25" ] <- predinc.apc( m.mw.apc, nd, dfac=0.025 )$pr.resp
> Mu.W[,,"F","i25" ] <- predinc.apc( f.mw.apc, nd, dfac=0.025 )$pr.resp
> Mu.W[,,"M","i30" ] <- predinc.apc( m.mw.apc, nd, dfac=0.030 )$pr.resp
> Mu.W[,,"F","i30" ] <- predinc.apc( f.mw.apc, nd, dfac=0.030 )$pr.resp

```

4.5 Average trends

The average annual trends in all of the rates can now be summarized:

```

> mort.chg <- rbind( ci.exp(m.md.aPC,subset="I\\(P")-1,
+                       ci.exp(f.md.aPC,subset="I\\(P")-1,
+                       ci.exp(m.mw.aPC,subset="I\\(P")-1,
+                       ci.exp(f.mw.aPC,subset="I\\(P")-1 )*100
> rownames( mort.chg ) <- c("Mortality change,   DM: Men",
+                           "                               Women",
+                           "Mortality change, Well: Men",
+                           "                               Women")
> round( rbind( inc.chg, mort.chg ), 1 )

```

		exp(Est.)	2.5%	97.5%
DM incidence change	Men	2.2	2.1	2.2
	Women	2.6	2.5	2.7
Mortality change,	DM: Men	-3.7	-3.8	-3.6
	Women	-3.4	-3.5	-3.2
Mortality change, Well:	Men	-2.9	-2.9	-2.8
	Women	-2.4	-2.5	-2.4

Thus it appears that the incidence rates of diabetes are increasing by some 2.2% per year for women but 3.5% per year for men, while mortality rates are decreasing 3.6% per year for persons with diabetes, but only 2.5–2.9% per year for persons without.

For convenience of calculations and for subsequent use, we save the estimated rates and other quantities of interest:

```

> save( Lambda, Mu.W, Mu.DM, a.pt, p.pt, t.pt, int, qn, file="../data/rateEsts.Rda" )
> load( file="../data/rateEsts.Rda" )

```

4.6 Trends in estimated rates

In order to show how the models predict the time trends in diabetes incidence and mortality, we make a graphical display of the estimated rates in ages 20,...,90 versus calendar time:

A brief overview of the mortality and incidence rates over time:

```

> pts <- as.numeric( dimnames(Lambda)[[2]] ) [1:240]
> ( dimnames(Lambda)[[1]][agr <- seq(240,1080,120)] )
[1] "19.95833333333333" "29.95833333333333" "39.95833333333333" "49.95833333333333"
[5] "59.95833333333333" "69.95833333333333" "79.95833333333333" "89.95833333333333"
> par( mfrow=c(3,2), mar=c(1,1,1,1), oma=c(3,3,0,0), mgp=c(3,1,0)/1.6, las=1, bty="n" )
> yticks <- outer( 1:9, 10^(-2:1) ) [1:30]
> rpl <- function( Lambda, sx, yl ){
+   matplot( pts, t(Lambda[agr,1:240,sx,"apc"])*1000,
+           log="y", yaxt="n", ylim=c(0.01,30), ylab="", xaxt="n",
+           lty=1, lwd=5, type="l",
+           col=gray(4:11/13) )#if(sx=="M") "blue" else "red" )
+   mtext( side=2, yl, line=2.5, las=0 )
+ }
> rpl( Lambda, "M", "DM incidence per 1000 PY" )
>   axis( side=2 ) ; axis( side=2, at=yticks, labels=NA )
> text( 1996, 30, "Men", adj=c(0,1), cex=1.5 )
> rpl( Lambda, "F", "" )
> text( 1996, 30, "Women", adj=c(0,1), cex=1.5 )
> rpl( Mu.W , "M", "Population mortality per 1000 PY" )
>   axis( side=2 ) ; axis( side=2, at=yticks, labels=NA )
> rpl( Mu.W , "F", "" )

```

```

> rpl( Mu.DM, "M", "DM mortality per 1000 PY" )
>           axis( side=2 ) ; axis( side=2, at=yticks, labels=NA )
>           axis( side=1 ) ; axis( side=1, at=1996:2016, labels=NA )
> rpl( Mu.DM, "F", "" ) ; axis( side=1 ) ; axis( side=1, at=1996:2016, labels=NA )
> mtext( "Date of follow-up", side=1, line=2, outer=TRUE )

```

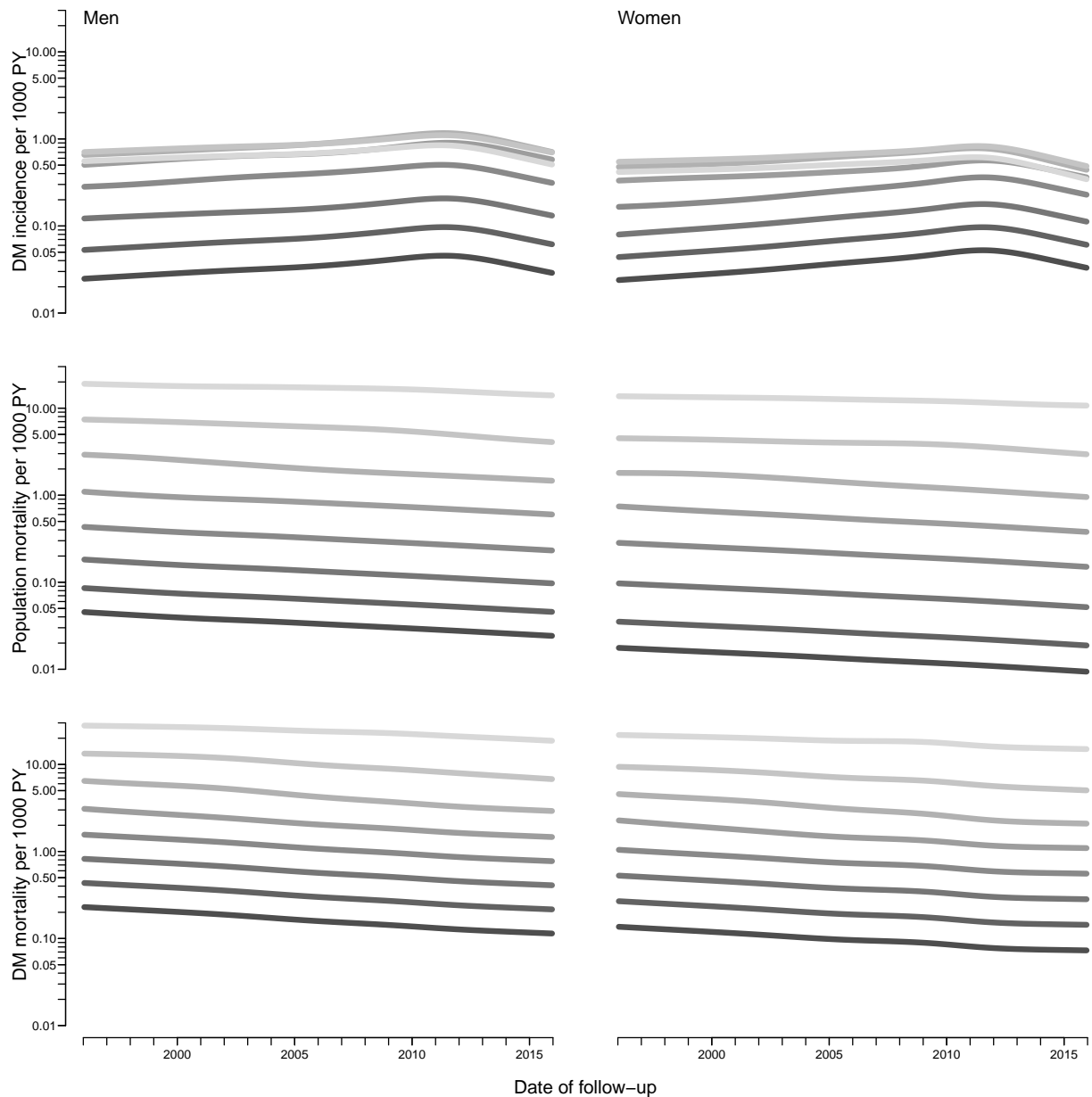


Figure 4.4: Trends in incidence and mortality rates for ages 20 (darkest), 30, ..., 90 (lightest), as estimated from the age-period-cohort models.

4.7 Extrapolation of rates

It is possible to extrapolate the rates beyond the observed dates by simply extending the linear part of the natural splines; in fact this is already done in the rate-objects `Lambda`,

Mu.W and Mu.DM. However, as seen in figure 4.5 the predicted decline in diabetes rates is presumably way too dramatic.

```
> pts <- as.numeric( dimnames(Lambda)[[2]] )
> ( dimnames(Lambda)[[1]][agr <- seq(240,1080,120)] )
[1] "19.95833333333333" "29.95833333333333" "39.95833333333333" "49.95833333333333"
[5] "59.95833333333333" "69.95833333333333" "79.95833333333333" "89.95833333333333"
> par( mfrow=c(3,2), mar=c(1,1,1,1), oma=c(3,3,0,0), mgp=c(3,1,0)/1.6, las=1, bty="n" )
> yticks <- outer( 1:9, 10^(-2:1) )[1:30]
> rpl <- function( Lambda, sx, yl, inc=FALSE ){
+   matplot( pts, t(Lambda[agr,,sx,"apc"])*1000,
+           log="y", yaxt="n", ylim=c(0.01,30), ylab="", xaxt="n",
+           lty=1, lwd=5, type="l", col=gray(4:11/13) )
+   matlines( pts, t(Lambda[agr,,sx,"att"])*1000,
+            lty=1, lwd=1, type="l", col="black" )
+   if( inc )
+     matlines( pts, t(Lambda[agr,,sx,"i25"])*1000,
+              lty=1, lwd=1, type="l", col="red" )
+   abline( v=2016, lty=3, col=gray(0.6) )
+   mtext( side=2, yl, line=2.5, las=0 )
+ }
> rpl( Lambda, "M", "DM incidence per 1000 PY", inc=TRUE )
> axis( side=2 ) ; axis( side=2, at=yticks, labels=NA, tcl=-0.3 )
> text( 1996, 30, "Men", adj=c(0,1), cex=1.5 )
> rpl( Lambda, "F", "", inc=TRUE )
> text( 1996, 30, "Women", adj=c(0,1), cex=1.5 )
> rpl( Mu.W , "M", "Population mortality per 1000 PY" )
> axis( side=2 ) ; axis( side=2, at=yticks, labels=NA, tcl=-0.3 )
> rpl( Mu.W , "F", "" )
> rpl( Mu.DM, "M", "DM mortality per 1000 PY" )
> axis( side=2 ) ; axis( side=2, at=yticks, labels=NA, tcl=-0.3 )
> axis( side=1 ) ; axis( side=1, at=1996:2030, labels=NA, tcl=-0.3 )
> rpl( Mu.DM, "F", "" ) ; axis( side=1 ) ; axis( side=1, at=1996:2030, labels=NA, tcl=-0.3 )
> mtext( "Date of follow-up", side=1, line=2, outer=TRUE )
```

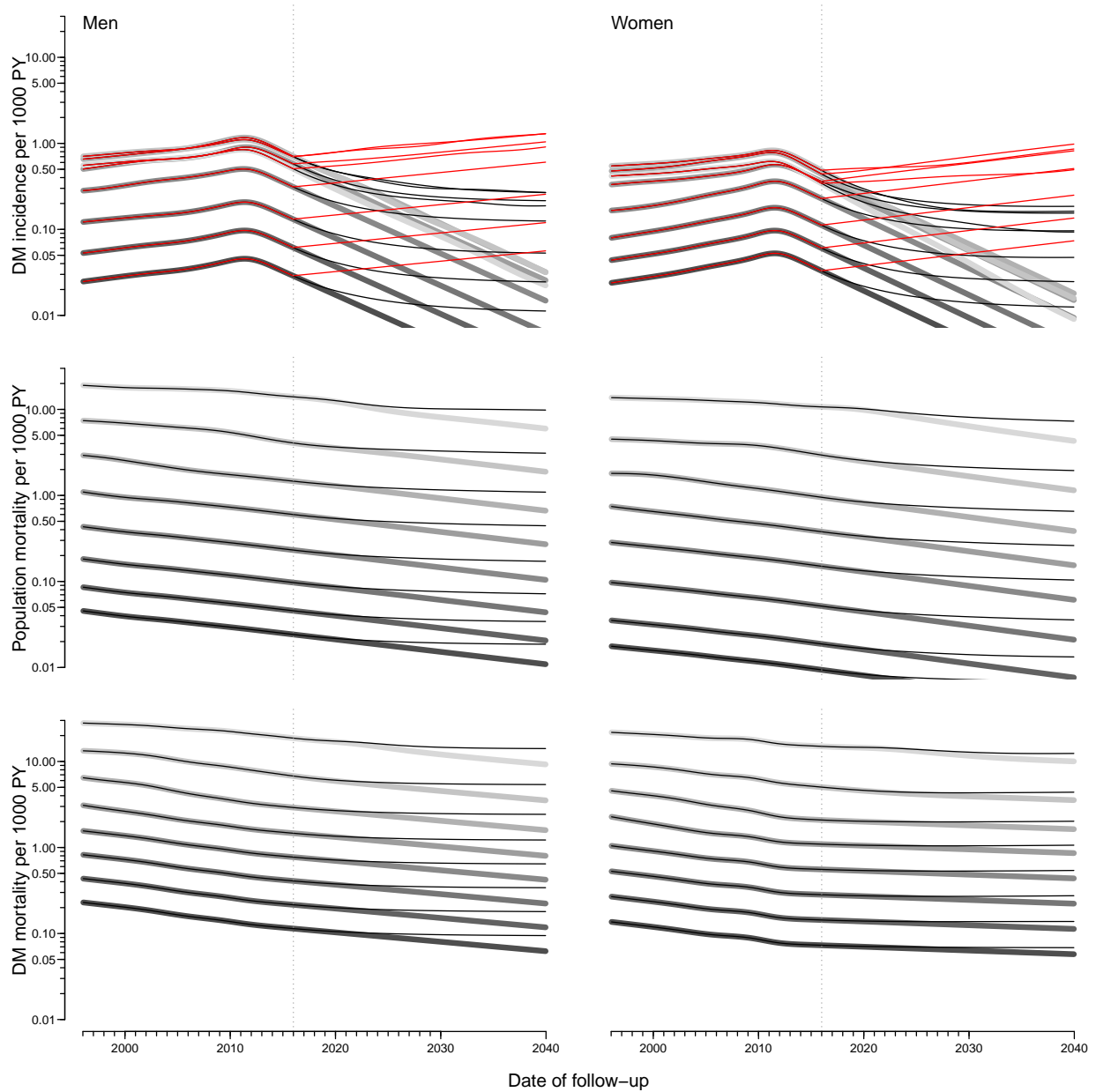



Figure 4.5: Trends in incidence and mortality rates for ages 20 (darkest), 30, . . . , 90 (lightest), as estimated from the age-period-cohort models and predicted by naive extrapolation of the natural splines. The vertical dotted lines indicate the end of available data, and the thin overlaid lines represent the rate predictions based on attenuated period- and cohort-effects. The red lines are predictions assuming an annual increase in period effects of 2.5%/year — only used for the incidence rates.

Chapter 5

Components of prevalence

The purpose of this chapter is to use the estimated transition rates to predict the prevalences at later times. This is in itself not an interesting endeavour, because we have the prevalence data available, but it will serve as an illustration that the rates are adequately modelled and that the degree of approximation is adequate when using an interval length as chosen.

```
> load( file="../data/rateEsts.Rda")
> lls()
  name  mode   class  size
1 a.pt  numeric numeric 1200
2 int   numeric numeric   1
3 Lambda numeric array  1200 528 2 6
4 Mu.DM numeric array  1200 528 2 6
5 Mu.W  numeric array  1200 528 2 6
6 p.pt  numeric numeric  528
7 qn    function function 1
8 t.pt  numeric numeric  529
```

We shall use the simulation scheme to predict the course of DM prevalence development in the population under various scenarios of mortality and incidence development. So we set up various structures to hold results and clarify calculations:

pr.fit — array of empirical age-specific prevalences at 1.1.1996–1.1.2016, smoothed by natural splines.

TR — array of transition probabilities between states Well and DM and Death. Transition probabilities are computed under the 4 different scenarios combining mortality and incidence rates either as they actually developed 1996–2016 or assuming they were constant at the 1996 level. These refer to intervals of length **int** and are therefore labeled on the period dimension by the midpoint of these, a total of $20/\text{int}$.

prv — array of predicted prevalences based on the initial prevalences at 1.1.1996 and the transition probabilities as put in **TR**. The scenario dimension refers to the 4 scenarios: “**obs**”, “**m-fix**”, “**i-fix**” and “**all-f**”, but this dimension in the array is expanded by 3 extra levels “**mort**”, “**inc**” and “**const**” that are to be filled with the part of the prevalences that are attributable to decrease in mortality, increase in incidence and the disequilibrium between rates and prevalence in 1996. Likewise, the period dimension is expanded by one relative to that in **TR** (replacing **p.pt** (period points)

with `t.pt` (time points)), since this refer to points in time and not time intervals (periods).

`prn` — array of predicted *number* of DM patients in one-year age classes at the 1 January each year. So the same structure as `prv`, but with substantially fewer entries.

5.1 Transition probabilities

In order to get the predicted *number* of persons by age, period and prediction type, we need the (1-step) transition matrices at all combinations of age (*a*) and date (*p*), this is put in array.

Note that the structures `Lambda`, `Mu.W` and `Mu.DM`, contain predicted rates up to 2040 under different scenarios, that we do not need:

```
> str( Lambda )
num [1:1200, 1:528, 1:2, 1:6] 5.62e-06 5.65e-06 5.69e-06 5.72e-06 5.76e-06 ...
- attr(*, "dimnames")=List of 4
..$ a : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.291666666666667"
..$ p : chr [1:528] "1996.041666666667" "1996.125" "1996.208333333333" "1996.291666666667"
..$ sex: chr [1:2] "M" "F"
..$ mod: chr [1:6] "ap" "apc" "att" "i20" ...
```

...we only need the dates till 2016, and the models `ap` and `apc`:

```
> # 2nd dimension of rates is all the way to 2040, only need to 2016
> dimnames(Lambda)[[2]][250+1:4]
[1] "2016.875" "2016.958333333333" "2017.041666666667" "2017.125"
> Lambda <- Lambda[,1:252,,1:2]
> Mu.W <- Mu.W [,1:252,,1:2]
> Mu.DM <- Mu.DM [,1:252,,1:2]
> states <- c("Well", "DM")
> TR <- NArray( c( dimnames(Lambda),
+ list( from = states,
+ to = states,
+ scene = c("obs", "m-fix", "i-fix", "all-f" ) ) ) )
> str( TR )
logi [1:1200, 1:252, 1:2, 1:2, 1:2, 1:2, 1:4] NA NA NA NA NA NA ...
- attr(*, "dimnames")=List of 7
..$ a : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.291666666666667"
..$ p : chr [1:252] "1996.041666666667" "1996.125" "1996.208333333333" "1996.291666666667"
..$ sex: chr [1:2] "M" "F"
..$ mod : chr [1:2] "ap" "apc"
..$ from : chr [1:2] "Well" "DM"
..$ to : chr [1:2] "Well" "DM"
..$ scene: chr [1:4] "obs" "m-fix" "i-fix" "all-f"
```

The situation where both the mortality rates and incidence rates are fixed at the 1996 level is trivial, because transition probabilities in that case only depend on age and not on period.

In order to fill `TR`, we need the cumulative incidences over intervals of length `int`. But these were exactly the ones we predicted in the previous sections by setting the person years equal to `int` in the data frame supplied to the `newdata` argument.

So we can now compute the one-`int`-step transition matrices for every combination of `a.pt` and `p.pt`, both in steps of `int` (in this case 0.083 year):

```
> TR[,,,,"Well","Well","obs"] <- exp(-Lambda-Mu.W)
> TR[,,,,"Well","DM" ,"obs"] <- Lambda
> TR[,,,,"DM" ,"Well","obs"] <- 0
> TR[,,,,"DM" ,"DM" ,"obs"] <- exp(-Mu.DM)
```

Note that we have not included the “Dead” state in the calculations, because we only bother about the *fraction* of diabetes patients in each age class at each time-point. So the probabilities we compute do not sum to 1 within the “from” states; we only compute how many of the persons alive that end up being alive at the next timepoint

When we fix the mortality or incidence at the 1996 level we just replace the expressions above with expressions where we replace the date dimension by `rep(1,np)`, (where `np` is the number of periods) for either incidence, mortality or both:

```
> ( np <- dim(Lambda)[2] )
P
252
> TR[,,,,"Well","Well","m-fix"] <- exp(-Lambda-Mu.W[,rep(1,np),,])
> TR[,,,,"Well","DM" ,"m-fix"] <- Lambda
> TR[,,,,"DM" ,"Well","m-fix"] <- 0
> TR[,,,,"DM" ,"DM" ,"m-fix"] <- exp(-Mu.DM[,rep(1,np),,])

> TR[,,,,"Well","Well","i-fix"] <- exp(-Lambda[,rep(1,np),,]-Mu.W)
> TR[,,,,"Well","DM" ,"i-fix"] <- Lambda[,rep(1,np),,]
> TR[,,,,"DM" ,"Well","i-fix"] <- 0
> TR[,,,,"DM" ,"DM" ,"i-fix"] <- exp(-Mu.DM)

> TR[,,,,"Well","Well","all-f"] <- exp(-Lambda[,rep(1,np),,]-Mu.W[,rep(1,np),,])
> TR[,,,,"Well","DM" ,"all-f"] <- Lambda[,rep(1,np),,]
> TR[,,,,"DM" ,"Well","all-f"] <- 0
> TR[,,,,"DM" ,"DM" ,"all-f"] <- exp(-Mu.DM[,rep(1,np),,])
```

We have now collected the transition probabilities between “Well” and “DM” as well as the probabilities of remaining in each of these, all referring to a duration of `int`, a total of about 30 mil. numbers:

```
> str( TR )
num [1:1200, 1:252, 1:2, 1:2, 1:2, 1:2, 1:4] 1 1 1 1 1 ...
- attr(*, "dimnames")=List of 7
 ..$ a : chr [1:1200] "0.0416666666666667" "0.125" "0.208333333333333" "0.291666666666667"
 ..$ p : chr [1:252] "1996.041666666667" "1996.125" "1996.208333333333" "1996.291666666667"
 ..$ sex : chr [1:2] "M" "F"
 ..$ mod : chr [1:2] "ap" "apc"
 ..$ from : chr [1:2] "Well" "DM"
 ..$ to : chr [1:2] "Well" "DM"
 ..$ scene: chr [1:4] "obs" "m-fix" "i-fix" "all-f"

> prod( dim(TR) )
[1] 19353600

> save( TR, file="../data/TR.Rda" )
```

5.2 Prediction of the observed prevalences

Note that we do not need to predict the population size; we can get away with only predicting the prevalences as fractions. When we multiply the fraction of persons in states (Well,DM) with the transition matrix, we get fraction of the persons in the previous state that are in states (Well,DM), which does not sum to 1 (because of the ones dying), so we must rescale to prevalence age in each step.

First we set up an array to hold the predicted prevalences under different scenarios. Later we shall also compute the fraction of the prevalences that are attributable to trends in mortality and incidence as well as to the non-stationarity of the rates/prevalences as of 1996, so we put in three extra levels of the last dimension, and one extra levels of the period dimension because we want to predict to the end of the last period too (or, to put it differently, we need an extra first level to hold the starting prevalences as of 1.1.1996).

```
> dpr <- c( dimnames(Lambda)[1:4],
+          list( c(dimnames(TR)[["scene"]], "mort", "inc", "const") ) )
> names( dpr )[c(2,5)] <- c("t", "what")
> t.pt[250+1:4]
[1] 2016.833 2016.917 2017.000 2017.083
> dpr[["t"]] <- t.pt[1:253]
> prv <- NArray(dpr)
> str( prv )

logi [1:1200, 1:253, 1:2, 1:2, 1:7] NA NA NA NA NA NA ...
- attr(*, "dimnames")=List of 5
..$ a : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.2916666666666666
..$ t : chr [1:253] "1996" "1996.08333333333333" "1996.166666666667" "1996.25" ...
..$ sex : chr [1:2] "M" "F"
..$ mod : chr [1:2] "ap" "apc"
..$ what: chr [1:7] "obs" "m-fix" "i-fix" "all-f" ...

> prod( dim(prv) )
[1] 8500800
```

Then we make a loop that updates the prevalences at 1.1.1996 to those at subsequent times, but first we must initialize the prevalences as modeled at 1996-01-01, as well as the prevalences at ages 0 (the new-born) that we set to 0; note that we exploit the column major storage of arrays to get `prv` filled with identical values across the two last dimensions (model and scenario).

```
> load( file="../data/prFit.Rda" )
> str( pr.fit ); str( prv )

num [1:2, 1:1200, 1:21] 0.000626 0.000579 0.00063 0.000583 0.000635 ...
- attr(*, "dimnames")=List of 3
..$ sex: chr [1:2] "M" "F"
..$ A : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.2916666666666666
..$ P : chr [1:21] "1996" "1997" "1998" "1999" ...

logi [1:1200, 1:253, 1:2, 1:2, 1:7] NA NA NA NA NA NA ...
- attr(*, "dimnames")=List of 5
..$ a : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.2916666666666666
..$ t : chr [1:253] "1996" "1996.08333333333333" "1996.166666666667" "1996.25" ...
..$ sex : chr [1:2] "M" "F"
..$ mod : chr [1:2] "ap" "apc"
..$ what: chr [1:7] "obs" "m-fix" "i-fix" "all-f" ...
```

```

> # Smoothed prevalences at 1.1.1996 - the starting values
> for( sx in c("M","F") ) prv[,1,sx,,] <- pr.fit[sx,,1]
> # Prevalences at age 0
> prv[1,,"M",,] <- 0
> prv[1,,"F",,] <- 0
> tt <- prv[c(1:2,floor(dim(prv)[1]/1.5)+1:2),1:2,,]
> dimnames( tt )[[1]] <- round( as.numeric(dimnames( tt )[[1]]), 3 )
> dimnames( tt )[[2]] <- round( as.numeric(dimnames( tt )[[2]]), 3 )
> round( ftable( tt, row.vars=4:1 )*100, 1 )

```

mod	sex	t	a	what	obs	m-fix	i-fix	all-f	mort	inc	const
ap	M	1996	0.042	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
			0.125	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
			66.708	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0
			66.792	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0
		1996.083	0.042	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
		0.125	NA	NA	NA	NA	NA	NA	NA	NA	NA
	F	1996	0.042	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
			0.125	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
			66.708	3.7	3.7	3.7	3.7	3.7	3.7	3.7	3.7
			66.792	3.7	3.7	3.7	3.7	3.7	3.7	3.7	3.7
		1996.083	0.042	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
		0.125	NA	NA	NA	NA	NA	NA	NA	NA	NA
apc	M	1996	0.042	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
			0.125	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
			66.708	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0
			66.792	5.0	5.0	5.0	5.0	5.0	5.0	5.0	5.0
		1996.083	0.042	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
		0.125	NA	NA	NA	NA	NA	NA	NA	NA	NA
	F	1996	0.042	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
			0.125	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
			66.708	3.7	3.7	3.7	3.7	3.7	3.7	3.7	3.7
			66.792	3.7	3.7	3.7	3.7	3.7	3.7	3.7	3.7
		1996.083	0.042	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0
		0.125	NA	NA	NA	NA	NA	NA	NA	NA	NA

So now we have checked that we have put initial values correctly into `prv`, basically at the period edge (at 1996) and the age edge (at 0). Then we can compute the predicted prevalences under the different scenarios. We take the fraction of the population in age class `ia` at time `ip` that end up as diabetes patients at time `ip+1` (and hence in age class `ia+1`), and divide by the fraction of all that remain alive, which is the diabetes patients (survivors and new), *plus* those who survive free of diabetes:

```

> system.time(
+ for( ip in 1:(dim(prv)[2]-1) )
+ for( ia in 1:(dim(prv)[1]-1) )
+ prv[ia+1,ip+1,,1:4] <-
+ ( prv[ia,ip,,1:4] * TR[ia,ip,, "DM" , "DM" ,]
+ +(1-prv[ia,ip,,1:4]) * TR[ia,ip,, "Well", "DM" ,] ) /

```

```

+ ( prv[ia,ip,,,1:4] * TR[ia,ip,,, "DM" , "DM" ,]
+   +(1-prv[ia,ip,,,1:4]) * TR[ia,ip,,, "Well", "DM" ,]
+   +(1-prv[ia,ip,,,1:4]) * TR[ia,ip,,, "Well", "Well",] )
+
  user  system elapsed
12.53   0.02   12.55

```

Note that the reason that the last dimension, `scene`, is explicitly mentioned in the array `prv` is because the length of this dimension is 7, but the corresponding in `TR` only 4 — recall that `prv` also has three extra levels to provide for the estimated part of the prevalences attributable to mortality change, incidence changes, and non-equilibrium at 1996.

Moreover, note that the code above is particularly simple because we only need to compute the prevalence at the next date and age. If we had had a more elaborate model with, say complications states, the calculations in the loop would have been a matrix-multiplication updating the state-distribution, but this simplification would have been at the expense of another three loop-levels, namely over the three last dimensions of the `prv` array — the matrix machinery only operates on 2-dimensional structures, matrices. (Well, maybe some `apply` construction could be set up).

We can then show a few of the predicted prevalences in (

```

> round( prv[1:4,1:4,1,1,1,drop=F]*100, 3 )
, , sex = M, mod = ap, what = obs

      t
a      1996 1996.083333333333 1996.166666666667 1996.25
0.04166666666666667 0.000          0.000          0.000  0.000
0.125              0.063          0.001          0.001  0.001
0.2083333333333333 0.063          0.064          0.001  0.001
0.2916666666666667 0.064          0.064          0.064  0.002

> save( a.pt, prv, file="../data/prv.Rda" )
> load(          file="../data/prv.Rda" )

```

5.2.1 Checking the prediction

With this initial prediction in place we can now check whether we have made a reasonable approximation to the observed prevalences at 1.1.2016.

In the array `prv` are all the prevalences as predicted from the prevalence in 1996 using the estimated incidences and mortalities; predicted at intervals of `inc` whereas we have the smoothed empirical prevalences in 1996, 2001, 2006, 2011 and 2016 in the array `pr.fit`:

Thus we have the predicted age-specific prevalences for men in say 2000 in `prv[, "2000", "M", "apc", "obs"]`, and the smoothed empirical in `pr.fit[, "M", "2000"]`.

We now plot these in the same plot:

```

> nd <- c( grep("2001",dimnames(prv)[[2]])[1],
+         grep("2006",dimnames(prv)[[2]])[1],
+         grep("2011",dimnames(prv)[[2]])[1],
+         grep("2016",dimnames(prv)[[2]])[1] )
> ( wh <- dimnames(prv)[[2]][nd] )
[1] "2001" "2006" "2011" "2016"

```



```

> par( mfrow=c(1,2), mar=c(0,0,0,0), oma=c(3,4,1,1), mgp=c(3,1,0)/1.6,
+     las=1, bty="n" )
> matplot( a.pt, pr.fit["M",,wh]*100,
+         xlim=c(10,95), ylim=c(0,18), yaxs="i",
+         xlab="Age", ylab="Prevalence (%)",
+         type="l", col="blue", lty=1, lwd=2 )
> axis( side=1, at=1:9*10, labels=NA )
> matlines( a.pt, prv[,wh,"M","apc","obs"]*100,
+         type="l", col="blue", lty="32", lwd=4 )
> matplot( a.pt, pr.fit["F",,wh]*100,
+         xlim=c(10,95), ylim=c(0,18), yaxs="i",
+         xlab="Age", ylab="", yaxt="n",
+         type="l", col="red", lty=1, lwd=2 )
> matlines( a.pt, prv[,wh,"F","apc","obs"]*100,
+         type="l", col="red", lty="32", lwd=3 )
> axis( side=1, at=1:9*10, labels=NA )
> mtext( "Prevalence of DM (%)", side=2, line=2, las=0, outer=TRUE )
> mtext( "Age", side=1, line=2, las=0, outer=TRUE )

```

```

> par( mfrow=c(1,2), mar=c(0,0,0,0), oma=c(3,4,1,1), mgp=c(3,1,0)/1.6,
+     las=1, bty="n" )
> matplot( a.pt, cbind(prv[,wh,"M","apc","obs"],
+                    prv[,wh,"M","ap" ,"obs"],
+                    pr.fit["M",,wh])*100,
+         xlim=c(10,95), ylim=c(0,18), yaxs="i",
+         xlab="Age", ylab="Prevalence (%)",
+         type="l", col="black", lty=rep(c(0,1),c(8,4)), lwd=2 )
> matlines( a.pt, cbind(prv[,wh,"M","ap" ,"obs"],
+                    prv[,wh,"M","apc","obs"])*100,
+         type="l", col="blue", lty=rep(c("12","42"),c(4,4)), lwd=4 )
> axis( side=1, at=1:9*10, labels=NA )
> matplot( a.pt, cbind(prv[,wh,"F","apc","obs"],
+                    prv[,wh,"F","ap" ,"obs"],
+                    pr.fit["F",,wh])*100,
+         xlim=c(10,95), ylim=c(0,18), yaxs="i",
+         xlab="Age", ylab="", yaxt="n",
+         type="l", col="black", lty=rep(c(0,1),c(8,4)), lwd=2 )
> matlines( a.pt, cbind(prv[,wh,"F","ap" ,"obs"],
+                    prv[,wh,"F","apc","obs"])*100,
+         type="l", col="red", lty=rep(c("12","42"),c(4,4)), lwd=4 )
> axis( side=1, at=1:9*10, labels=NA )
> mtext( "Prevalence of DM (%)", side=2, line=2, las=0, outer=TRUE )
> mtext( "Age", side=1, line=2, las=0, outer=TRUE )

```

Since the APC-models for rates clearly provide a better fit (figure 5.1), we shall use these in the reporting of the different scenarios.

We now compare the predicted prevalences under the four scenarios at 1.1.2016:

```

> str( prv )
num [1:1200, 1:253, 1:2, 1:2, 1:7] 0 0.00063 0.000635 0.000639 0.000644 ...
- attr(*, "dimnames")=List of 5
..$ a : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.2916666666666666
..$ t : chr [1:253] "1996" "1996.083333333333" "1996.166666666667" "1996.25" ...
..$ sex : chr [1:2] "M" "F"
..$ mod : chr [1:2] "ap" "apc"
..$ what: chr [1:7] "obs" "m-fix" "i-fix" "all-f" ...
> dimnames(prv)[["t"]][np <- 241]

```

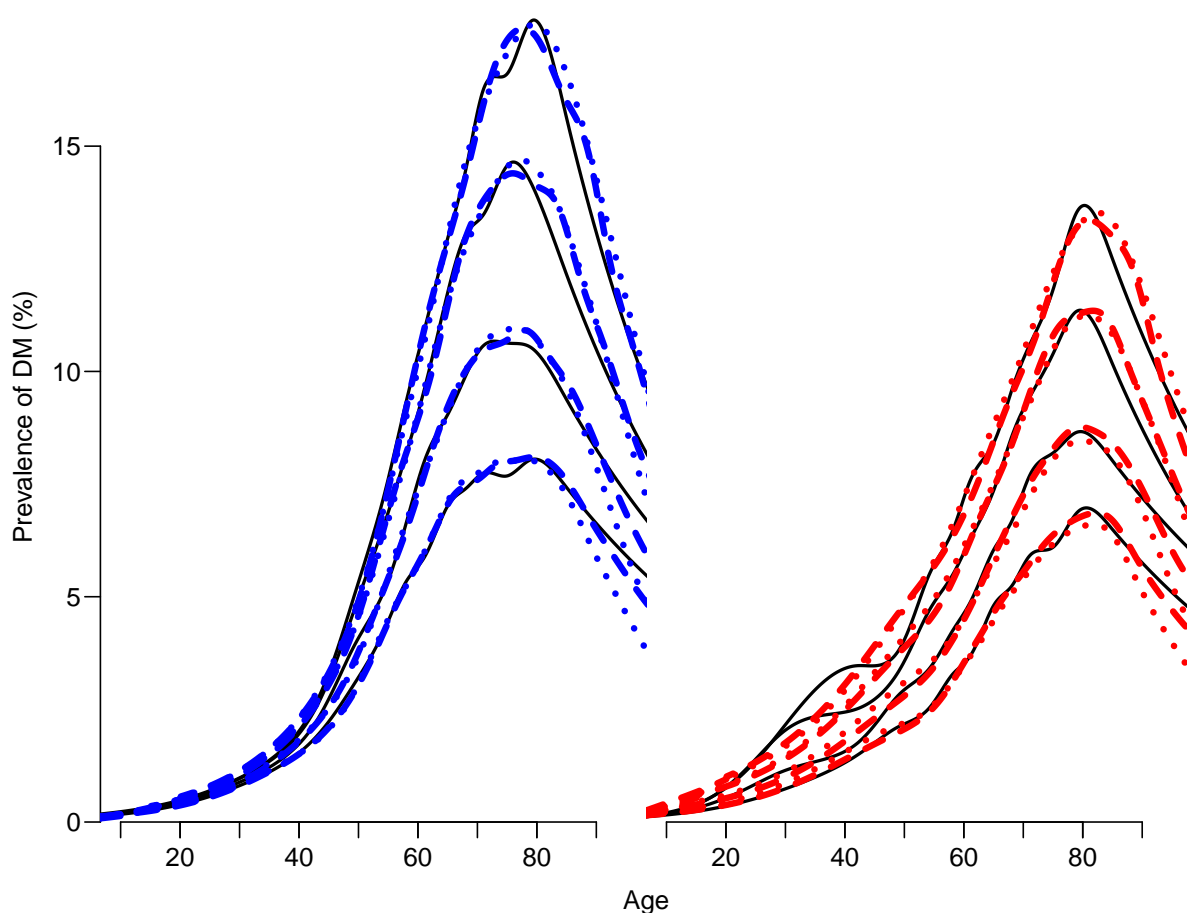



Figure 5.1: Plot of observed (full lines) and predicted prevalences in 2001, using simple age-period-models (dotted lines) or age-period-cohort models (broken lines). Clearly the broken lines gives the better approximation to the smoothed empirical rates (black lines).

[1] "2016"

```
> prv[floor(dim(prv)[1]/1.5)+1:5,np,"M","apc",]*100
```

what

a	obs	m-fix	i-fix	all-f	mort	inc	const
66.7083333333333	13.74514	12.52098	10.45399	9.454924	NA	NA	NA
66.7916666666667	13.78295	12.54914	10.49134	9.483485	NA	NA	NA
66.875	13.82081	12.57731	10.52854	9.511871	NA	NA	NA
66.9583333333333	13.85873	12.60549	10.56559	9.540077	NA	NA	NA
67.0416666666667	13.89672	12.63369	10.60249	9.568100	NA	NA	NA

```
> par( mfrow=c(1,2), mar=c(0,0,0,0), oma=c(3,4,1,1),
+      mgp=c(3,1,0)/1.6, las=1, bty="n" )
> lpl <- function( sx, cl )
+ matplot( a.pt, cbind(prv[,np,sx,"apc",c("obs","m-fix","i-fix","all-f")],
+                    prv[, 1,sx,"apc",1 ])*100,
+          xlim=c(20,90), ylim=c(0,18), yaxs="i", yaxt="n",
+          xlab="Age", ylab="Prevalence (%)",
+          type="l", lty=c(1,2,1,2,1),
+          col=cl, lwd=c(5,5,3,3,1) )
> lpl( "M", "blue" )
> axis( side=2 )
> axis( side=2, at=0:18, labels=NA, tcl=-0.3)
```

```
> lpl( "F", "red" )
> dimnames(prv)[[5]]
[1] "obs"    "m-fix"  "i-fix"  "all-f"  "mort"   "inc"    "const"
```

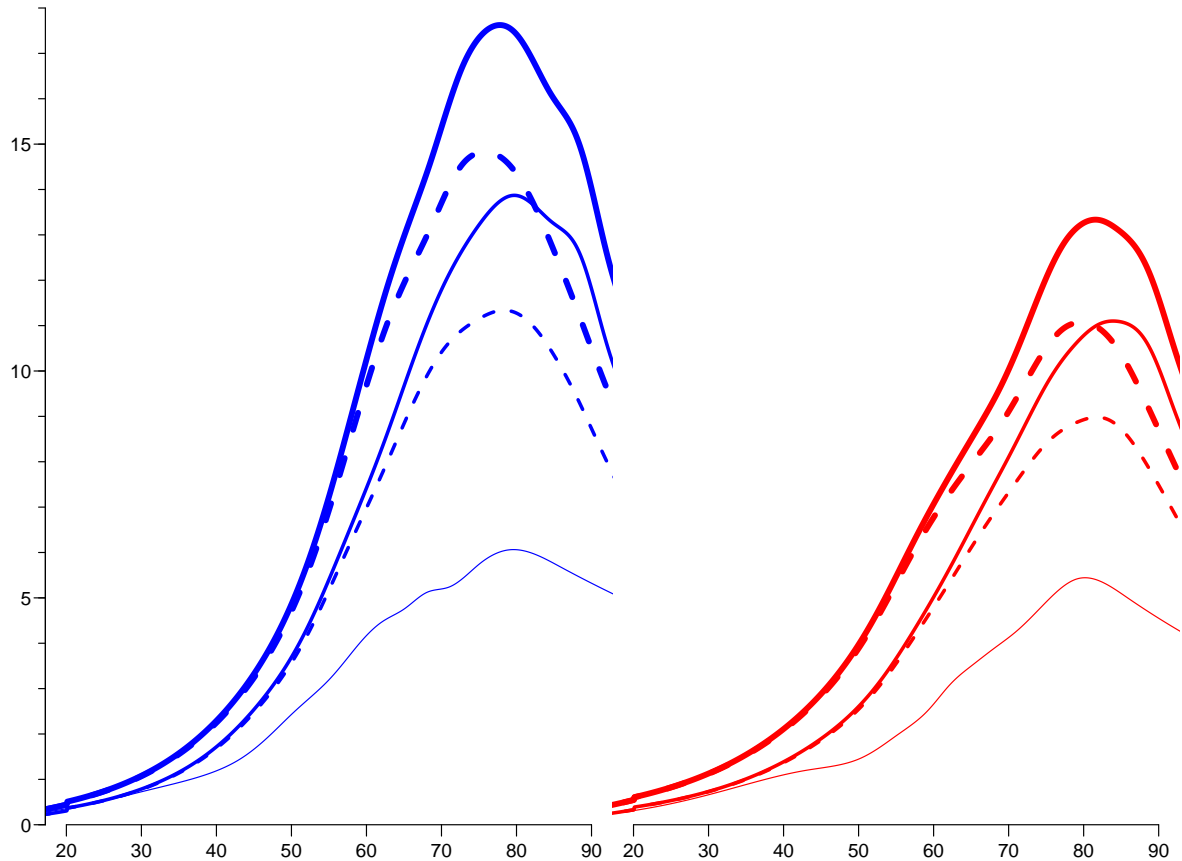


Figure 5.2: *The predicted prevalences under different scenarios:*

Full lines: Mortality rates evolve as observed.

Broken lines: Mortality rates remain as 1996.

Thick lines: Incidence rates evolve as observed.

Thin lines: Incidence rates remain as in 1996.

The very thin lines lowest in the two displays are the observed age-specific prevalences in 1996.

```
> scen <- c("Mort changes, Inc changes",
+          "Mort 1996, Inc changes",
+          "Mort changes, Inc 1996",
+          "Mort 1996, Inc 1996")
> c.a <- dimnames(prv)[[1]][floor(dim(prv)[1]/1.5)]
> n.a <- as.numeric( c.a )
> hts <- prv[c.a,np,"M","apc",1:4]*100
> cau.exp <-
+ function( wh=1:4, fill=FALSE )
+ {
+ pdf( paste( "comp-DMpr-", paste(wh,collapse=""), if( fill ) "F",
+                               ".pdf", sep="" ), height=8, width=11 )
```

```

+ par( mfrow=c(1,2), mar=c(0,0,0,0), oma=c(3,4,1,1),
+       mgp=c(3,1,0)/1.6, las=1, bty="n" )
+ matplot( a.pt, cbind(prv[,np,"M","apc"],prv[,1,"M","apc",1])*100,
+           xlim=c(20,90), ylim=c(0,18), xlab="Age",
+           ylab="Prevalence (%)", yaxs="i",
+           type="l", lty=rep(c(1,0),2), lwd=c(4,4,2,2,0)+1, col="blue" )
+ axis( side=2, at=0:18, labels=NA, tcl=-0.3 )
+ mtext( "Prevalence (%)", side=2, line=2.5, outer=T, las=0 )
+ matlines( a.pt, prv[,np,"M","apc",1]*100,
+           type="l", lty=rep(c("11","22"),2), lwd=c(4,4,2,2)+1, col="blue" )
+ matlines( a.pt, prv[,1,"M","apc",1]*100,
+           type="l", lty=1, lwd=1, col="blue" )
+ text( rep(20,4)[wh], hts[wh], scen[wh], adj=0, col="blue", cex=1.2 )
+ for( i in 1:15 )
+   arrows( (20.20+strwidth(scen,cex=1.2))[wh], hts[wh],
+           rep(n.a,4)[wh], hts[wh],
+           col="blue", angle=i, lwd=2 )
+ if( fill ) polygon( c(a.pt,rev(a.pt)),
+                   c(prv[,np,"M","apc",wh[1]],
+                     rev(prv[,np,"M","apc",wh[2]]))*100,
+                   col=rgb(0,0,1,0.3), border="transparent" )
+ matplot( a.pt, cbind(prv[,np,"F","apc"],prv[,1,"F","apc",1])*100,
+           xlim=c(20,90), ylim=c(0,18), xlab="Age", yaxt="n", yaxs="i",
+           type="l", lty=rep(c(1,0),2), lwd=c(4,4,2,2,0)+1, col="red" )
+ matlines( a.pt, prv[,np,"F","apc",1]*100,
+           type="l", lty=rep(c("11","22"),2), lwd=c(4,4,2,2)+1, col="red" )
+ matlines( a.pt, prv[,1,"F","apc",1]*100, type="l", lty=1, lwd=1, col="red" )
+ if( fill ) polygon( c(a.pt,rev(a.pt)),
+                   c(prv[,np,"F","apc",wh[1]],
+                     rev(prv[,np,"F","apc",wh[2]]))*100,
+                   col=rgb(1,0,0,0.3), border="transparent" )
+ dev.off()
+ }
> cau.exp(1:4)

pdf
2

> for( ff in c(FALSE,TRUE) )
+   {
+   cau.exp(1:2,fill=ff)
+   cau.exp(3:4,fill=ff)
+   cau.exp(c(1,3),fill=ff)
+   cau.exp(c(2,4),fill=ff)
+   }

```

Figure 5.2 shows the predicted prevalences under 4 different scenarios compared to the observed prevalences as of 1.1.1996.

5.3 How much is attributable to what?

We can compute how much of the age-specific prevalences that are attributable to mortality changes and how much to changes in incidence rates.

The effect of mortality decline can be computed either as the difference between “obs” and “m-fix” or as the difference between “i-fix” and “all-f”. But there is no guarantee that these two quantities are the same.

Similarly the effect of incidence increase can be computed either as the difference between “obs” and “i-fix” or as the difference between “m-fix” and “all-f”. And there is no guarantee that these two are the same either.

Hence we explore how different these quantities are:

```
> dimnames( prv )[5]
$what
[1] "obs"    "m-fix"  "i-fix"  "all-f"  "mort"   "inc"    "const"
> par( mfrow=c(1,2), mar=c(3,0,0,0), oma=c(0,4,1,1),
+      mgp=c(3,1,0)/1.6, las=1, bty="n" )
> matplot( a.pt, cbind( prv[,np,"M","apc","obs" ]-
+                      prv[,np,"M","apc","m-fix"],
+                      prv[,np,"M","apc","i-fix"]-
+                      prv[,np,"M","apc","all-f"] )*100,
+          xlim=c(20,90), ylim=c(0,5), xlab="Age", ylab="Prevalence (%)",
+          type="l", lty=1, lwd=c(4,2)+1, col="blue", yaxs="i" )
> axis( side=2, at=0:14/2, labels=NA, tcl=-0.3 )
> mtext( "Prevalence difference (%)", side=2, line=2.5, outer=T, las=0 )
> matlines(a.pt, cbind( prv[,np,"M","apc","obs" ]-
+                      prv[,np,"M","apc","i-fix"],
+                      prv[,np,"M","apc","m-fix"]-
+                      prv[,np,"M","apc","all-f"] )*100,
+          type="l", lty="22", lwd=c(4,2)+1, col="blue" )
> matplot( a.pt, cbind( prv[,np,"F","apc","obs" ]-
+                      prv[,np,"F","apc","m-fix"],
+                      prv[,np,"F","apc","i-fix"]-
+                      prv[,np,"F","apc","all-f"] )*100,
+          xlim=c(20,90), ylim=c(0,5), xlab="Age", yaxt="n", yaxs="i",
+          type="l", lty=1, lwd=c(4,2)+1, col="red" )
> matlines(a.pt, cbind( prv[,np,"F","apc","obs" ]-
+                      prv[,np,"F","apc","i-fix"],
+                      prv[,np,"F","apc","m-fix"]-
+                      prv[,np,"F","apc","all-f"] )*100,
+          type="l", lty="22", lwd=c(4,2)+1, col="red" )
```

From figure 5.3 we see that the two different ways of computing the contribution give pretty much the same results — the differences never exceed some 0.5%. Therefore, if we want to attribute fractions of the prevalence in 2010 to decreasing mortality and increasing incidence respectively, we would want two measures that had a sum equal the the difference between the scenario with observed mortality and incidence rates (“obs”), and the scenario with rates fixed to those from 1996 (“all-f”). This is obtained by taking the average of the two curves in each scenario.

The thin lines at the bottom of figure 5.2 represent the prevalence at 1.1.1996, so it is pretty clear that the incidence and mortality rates as observed by 1996 did not provide for at steady state.

So basically we can subdivide the prevalence at any point in time into 4 components:

1. the “inherited” prevalences from 1996.
2. the prevalence attributable to rates of mortality and incidence as of 1996 — the “epidemiological disequilibrium” as of 1996.
3. the prevalence attributable to the *increase* in the incidence rates.
4. the prevalence attributable to the *decrease* in the mortality rates.

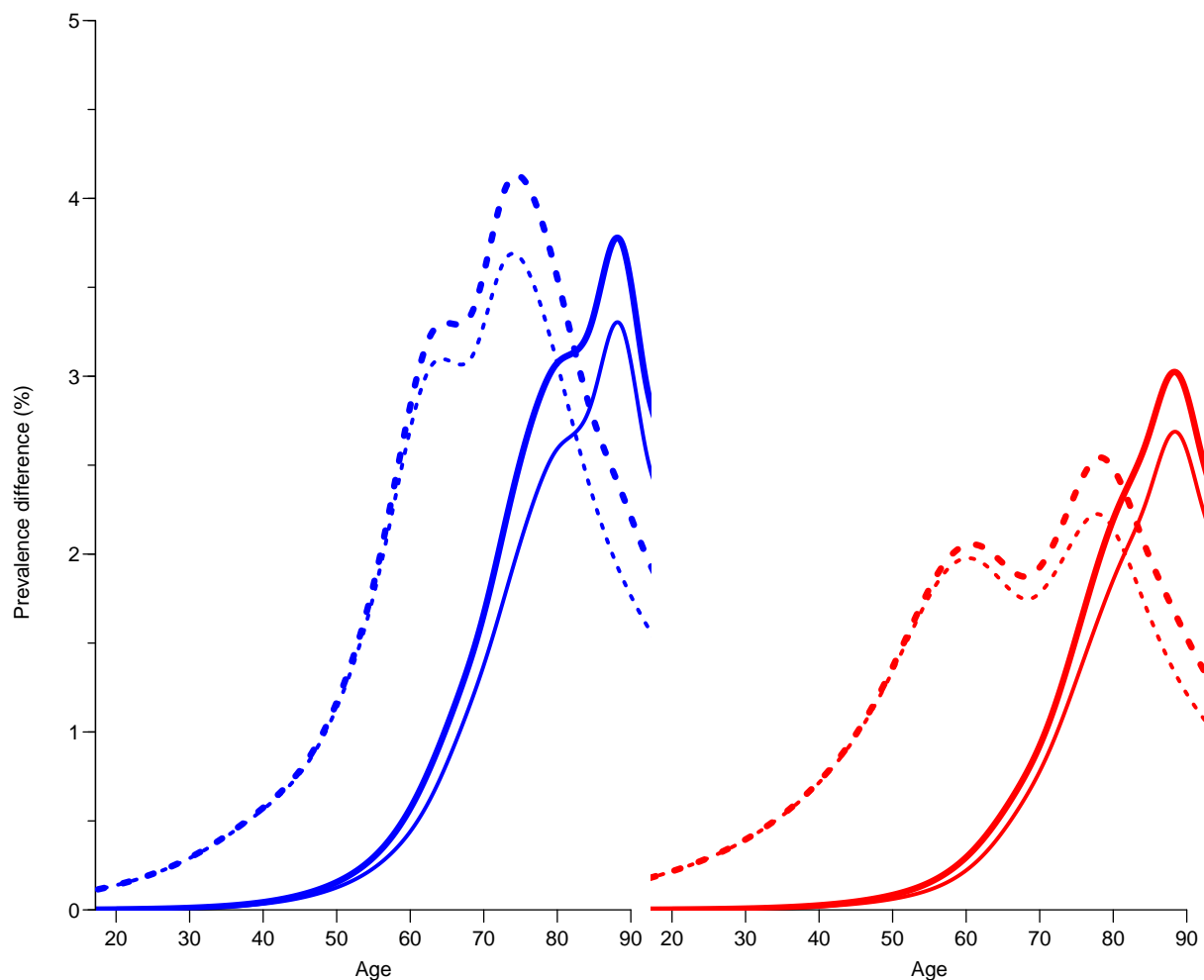


Figure 5.3: *Suggested contributions to age-specific prevalences from increasing mortalities over the period 1996–2016; the thick lines are obtained by subtracting the prediction based on fixing one rate from the one using the observed rates; thin lines based on subtracting the prediction based on fixing both rates from that where only one is fixed. Full lines are for differences attributable to changes in mortality rates, broken lines are for changes attributable to changes in incidence rates.*

So we now fill out the remaining 3 dimensions of `prv`:

```
> prv[,,,, "mort" ] <- ( prv[,,,, "obs" ] - prv[,,,, "m-fix" ] +
+   prv[,,,, "i-fix" ] - prv[,,,, "all-f" ] ) / 2
> prv[,,,, "inc" ] <- ( prv[,,,, "obs" ] - prv[,,,, "i-fix" ] +
+   prv[,,,, "m-fix" ] - prv[,,,, "all-f" ] ) / 2
> prv[,,,, "const" ] <- prv[,,,, "all-f" ] - prv[,rep(1, dim(prv)[2]),, "obs" ]
```

The components `obs`, `const`, `inc` and `mort` now together make up the total prevalence of diabetes for a given combinations of sex, age and date. Thus we can show these for each of the 15 dates 1996, ..., 2010.

First we define a function to make the component plots, and then we can plot the resulting development for men and women, for convenience we also put the latter in a function.

```

> poly.parts <-
+ function( x, crv, col, xlim, ylim, txt="" )
+ {
+ crv <- t(apply(cbind(0,crv),1,cumsum))
+ matplot( x, crv, type="n", xaxt="n", yaxt="n", xlab="", ylab="",
+         xlim=xlim, ylim=ylim, yaxs="i", bty="n" )
+ for( i in 2:ncol(crv) )
+ polygon( c(x,rev(x)), c(crv[,i],rev(crv[,i-1])),
+         col=col[i-1], border=col[i-1])
+ text( par("usr")[1:2]*%c(0.1,0.9),
+       par("usr")[3:4]*%c(0.9,0.1), txt, adj=c(1,0), font=2 )
+ }
> one.comp <-
+ function( sex, clr )
+ {
+ par( mfrow=c(3,6), mar=c(0,0,0,0), oma=c(3,3,1,1), mgp=c(3,1,0)/1.6 )
+ for(j in 1:18)
+ {
+ yn <- round(seq(1,dim(prv)[2],,18))
+ poly.parts( a.pt,
+             cbind(prv[,1 ,sex,"apc","obs"],
+                 prv[,yn[j],sex,"apc",c("const","inc","mort")])*100,
+             col=clr, xlim=c(20,90), ylim=c(0,18),
+             txt=dimnames(prv)[[2]][yn[j]] )
+ abline(h=0)
+ if( j==1 ) text( rep(25,3), c(13,15,17)+0.5,
+                 c("Imbalance","Incidence","Mortality"),
+                 col=clr[2:4], font=2, adj=0, cex=1.2 )
+ if( j %in% c(1,7,13) ) axis( side=2 )
+ if( j %in% 13:18 ) axis( side=1 )
+ mtext( "Age", side=1, outer=TRUE, line=1.5, font=1, las=0 )
+ mtext( "Prevalence of DM", side=2, outer=TRUE, line=1.5, font=1, las=0 )
+ }
+ }

> par( mfrow=c(1,2), mar=c(1,0,1,0), mgp=c(3,1,0)/1.6, las=1,
+       oma=c(2,3,0,1), bty="n" )
> clr <- rgb(c(3,2,1.5,0)/3,c(3,2,1.5,0)/3,1)
> poly.parts( a.pt, cbind(prv[,1 ,"M","apc","obs"],
+                         prv[,np,"M","apc",c("const","inc","mort")])*100,
+             col=clr, xlim=c(20,90), ylim=c(0,18) )
> abline(h=0:22,v=2:9*10,col=gray(0.9))
> axis( side=1 )
> axis( side=2 )
> axis( side=2, at=1:18, labels=NA, tcl=-0.3 )
> text( rep(25,3), 17:19-1.5,
+       c("Imbalance","Incidence","Mortality"),
+       col=clr[2:4], font=2, adj=0, cex=1.0 )
> clr <- rgb(1,c(3,2,1.5,0)/3,c(3,2,1.5,0)/3)
> poly.parts( a.pt, cbind(prv[,1 ,"F","apc","obs"],
+                         prv[,np,"F","apc",c("const","inc","mort")])*100,
+             col=clr, xlim=c(20,90), ylim=c(0,18) )
> abline(h=0:22,v=2:9*10,col=gray(0.9))
> axis( side=1 )
> text( rep(25,3), 17:19-1.5,
+       c("Imbalance","Incidence","Mortality"),
+       col=clr[2:4], font=2, adj=0, cex=1.0 )
> mtext( "Age", side=1, outer=TRUE, line=0.8, font=1, las=0 )
> mtext( "Prevalence of DM (%)", side=2, outer=TRUE, line=2, font=1, las=0 )

```

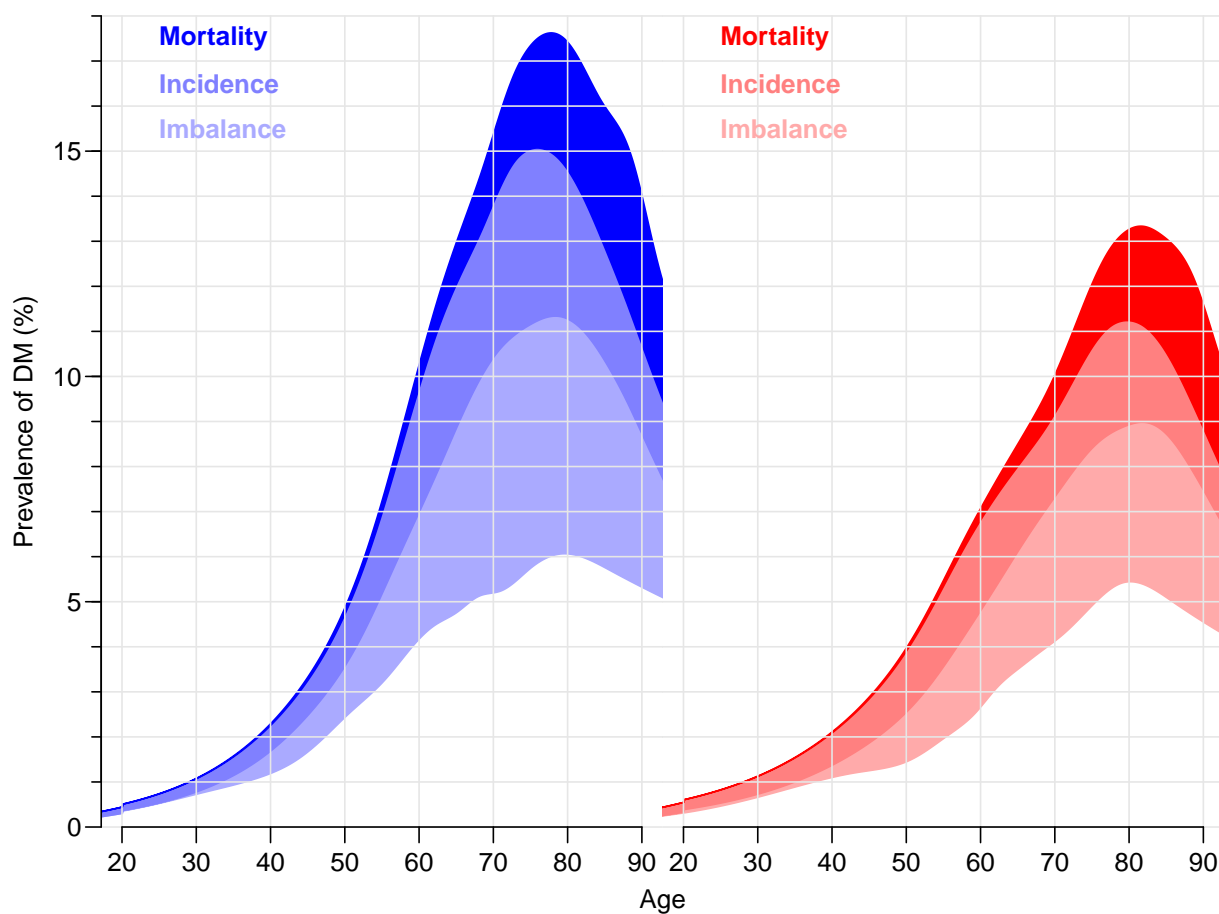


Figure 5.4: Changes in predicted age-specific prevalences of DM in Denmark 2016 among men (blue) and women (red), partitioned by the contribution from rates as they were in 1996 (“Imbalance”), increases in incidence and decrease in mortality, respectively.

5.4 Number of diabetes patients in Denmark

In the previous section we only looked at the age-specific prevalences, because these are the quantities that are driven by the incidence and mortality rates. However, it is also of interest to see how the actual number of diabetes patients would have looked under the different scenarios, specifically how the *number* of the current patients that can be attributed to the various components.

Also note that since the previous calculations were for age-specific prevalences we have a constant reference as the prevalences at 1996, but when we multiply by the population figures we would of course see differences in numbers and age-distribution of the diabetes population even if the age-specific prevalences were unchanged.

To show these effects we set up an array `prn` with `structure` (but not `extent`) as `prv` to hold the number of diabetes patients by category, assuming the age-distribution in the population to be as actually observed (that is as extracted from Statistics Denmark). However `prn` will have 100 age-classes rather than 1200 (`100/int`), and only 18 dates: `prv`.

```
> dn <- dimnames(prv)
> dn[[1]] <- 0:99
> dn[[2]] <- 1996:2016
> dn[[5]] <- dn[[5]][c(5:7,1)]
> prn <- NArray( dn )
> length(prv) ; str( prv ) ; table( prv<0 )
[1] 8500800
num [1:1200, 1:253, 1:2, 1:2, 1:7] 0 0.00063 0.000635 0.000639 0.000644 ...
- attr(*, "dimnames")=List of 5
..$ a : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.2916666666666666
..$ t : chr [1:253] "1996" "1996.083333333333" "1996.166666666667" "1996.25" ...
..$ sex : chr [1:2] "M" "F"
..$ mod : chr [1:2] "ap" "apc"
..$ what: chr [1:7] "obs" "m-fix" "i-fix" "all-f" ...
FALSE TRUE
8321273 179527
> length(prn) ; str( prn )
[1] 33600
logi [1:100, 1:21, 1:2, 1:2, 1:4] NA NA NA NA NA NA ...
- attr(*, "dimnames")=List of 5
..$ a : chr [1:100] "0" "1" "2" "3" ...
..$ t : chr [1:21] "1996" "1997" "1998" "1999" ...
..$ sex : chr [1:2] "M" "F"
..$ mod : chr [1:2] "ap" "apc"
..$ what: chr [1:4] "mort" "inc" "const" "obs"
```

In order to fill in the numbers we use the estimated age-specific prevalences at 1st January each year, that is at the dates 1996-01-01,...,2016-01-01 in the entries along the `t`-dimension of `prv`. Moreover we want the prevalences for a 1 year age class rather than age-classes of length `int`. So we take the average prevalences from `prv` over each one-year age-interval. The vectors `wh.a` and `wh.p` will hold the number of the age and period classes from `prv` which have the desired prevalences (as proportions) that we will use for multiplication with the population figures:


```

> comp <- c("mort","inc","const")
> str( prn[,,,,comp ] )
logi [1:100, 1:21, 1:2, 1:2, 1:3] NA NA NA NA NA NA ...
- attr(*, "dimnames")=List of 5
..$ a : chr [1:100] "0" "1" "2" "3" ...
..$ t : chr [1:21] "1996" "1997" "1998" "1999" ...
..$ sex : chr [1:2] "M" "F"
..$ mod : chr [1:2] "ap" "apc"
..$ what: chr [1:3] "mort" "inc" "const"

> # Find the dates in the predicted prevalences prv that matches the
> # dates in prn where empirical rates are available.
> prv.p <- as.numeric( dimnames(prv)[["t"]] )
> prn.p <- as.numeric( dimnames(prn)[["t"]] )
> wh.p <- match( prn.p, prv.p )
> if( any(is.na(wh.p)) ) # Need to find approximate dates if they do not match
+ for( ip in 1:length(prn.p) )
+ {
+   dd <- abs( prn.p[ip]-prv.p )
+   wh.p[ip] <- (1:length(dd))[dd==min(dd)]
+ }
> wh.p

[1] 1 13 25 37 49 61 73 85 97 109 121 133 145 157 169 181 193 205 217 229 241

> prv <- pmax( prv, 0 )
> # Ages in the two arrays
> prv.a <- as.numeric( dimnames(prv)[["a"]] )
> prn.a <- as.numeric( dimnames(prn)[["a"]] )
> for( ip in 1:length(wh.p) )
+ for( ia in 1:length(prn.a) )
+ {
+ wh.a <- which( prn.a[ia]==floor(prv.a) )
+ prn[ia,ip,,,comp ] <- apply( prv[wh.a,wh.p[ip],,, comp], 2:4, mean )
+ prn[ia,ip,,, "obs"] <- apply( prv[wh.a, 1,,, "obs"], 2:3, mean )
+ }

```

Now `prn` contains the estimated prevalences (as fractions) for 100 age classes and the 21 dates. We need to multiply these prevalences by the population figures at these times. The population figures are in `pr`:

```

> head( pr )
sex A P X N
2 M 0 1996 2 36258
3 M 0 1997 0 34920
4 M 0 1998 1 34936
5 M 0 1999 2 34151
6 M 0 2000 2 34035
7 M 0 2001 1 34432

> pop <- xtabs( N ~ A + P + sex, data=pr )[1:100,,]
> dmp <- xtabs( X ~ A + P + sex, data=pr )[1:100,,]
> str( pop )

table [1:100, 1:21, 1:2] 36258 36077 35003 35344 33830 ...
- attr(*, "dimnames")=List of 3
..$ A : chr [1:100] "0" "1" "2" "3" ...
..$ P : chr [1:21] "1996" "1997" "1998" "1999" ...
..$ sex: chr [1:2] "M" "F"

> str( dmp )

```

```

table [1:100, 1:21, 1:2] 2 8 10 18 14 21 33 38 52 39 ...
- attr(*, "dimnames")=List of 3
..$ A : chr [1:100] "0" "1" "2" "3" ...
..$ P : chr [1:21] "1996" "1997" "1998" "1999" ...
..$ sex: chr [1:2] "M" "F"

> str( prn )

num [1:100, 1:21, 1:2, 1:2, 1:4] 0 0 0 0 0 0 0 0 0 0 ...
- attr(*, "dimnames")=List of 5
..$ a : chr [1:100] "0" "1" "2" "3" ...
..$ t : chr [1:21] "1996" "1997" "1998" "1999" ...
..$ sex: chr [1:2] "M" "F"
..$ mod: chr [1:2] "ap" "apc"
..$ what: chr [1:4] "mort" "inc" "const" "obs"

> for( i in dimnames(prn)[[4]] )
+ for( j in dimnames(prn)[[5]] )
+ prn[,,,i,j] <- prn[,,,i,j] * pop

```

First we draw a simple pyramid of the age-distribution of diabetes patients in Denmark:

```

> # Note: This uses the undocumented feature that if the first
> # number in a column is negative this is taken as the left endpoint of
> # the bar. So c(-m,m,f) is a bar starting at -m, and a division at
> # -m+m(=0) and an upper end at -m+m-f. Coloring is from the top, that is
> # the part stretching from -m+m to -m+m+f get the first color
> pp <- "2016"
> oo <- c("mort","inc","const","obs")
> lim <- 6
> clr <- c("red","blue")
> draw.dmp <-
+ function(pp)
+ {
+   par( mar=c(3,3,3,0), mgp=c(3,1,0)/1.6, las=1 )
+   barplot( height=t( cbind( -dmp[,pp,"M"],
+                             dmp[,pp,"M"],
+                             dmp[,pp,"F"] ) ) / 1000,
+           horiz=TRUE, col=clr,
+           border="transparent",space=0,axes=FALSE,
+           names.arg=rep("",dim(prn)[1]),
+           xlim=c(-1,1)*lim*1.05,
+           xlab="Persons in 1 year class (1000s)",ylab="Age")
+   abline(h=seq(0,100,5),
+          v=seq(-lim,lim,0.5),
+          col="white")
+   axis( side=1, at=seq(-lim,lim,1), labels=abs(seq(-lim,lim,1)) )
+   axis( side=2, at=seq(0,100,20) )
+   axis( side=2, at=seq(0,100,5), labels=NA, tcl=-0.3 )
+   mtext( pp, at=-lim, adj=1.4, cex=1.3, font=1 )
+   mtext( formatC(sum(dmp[,pp,"M"]),0,format="f",big.mark=","),
+          at=-1, col="blue", line=0, cex=0.99 )
+   mtext( formatC(sum(dmp[,pp,"F"]),0,format="f",big.mark=","),
+          at=1, col="red", line=0, cex=0.99 )
+   mtext( "N", at=0, line=0, cex=0.99 )
+ }
> pdf( "comp-obs-film.pdf", width=8, height=6 )
> for( pp in paste(1996:2016) ) draw.dmp(pp)
> dev.off()

```

```
null device
1
```

```
> for( pp in paste(1996:2016) )
+ {
+ pdf( paste("comp-obs-", pp, ".pdf", sep=""), width=8, height=6 )
+ draw.dmp(pp)
+ dev.off()
+ }
```

Now we can also draw a population pyramid of the *predicted* number of DM patients using colors that range from very light to full:

```
> shd <- c(0.0, 1.1, 2.0, 2.8) / 3
> een <- rep(1,4)
> clr <- rgb( c(een,rev(shd)),
+           c(shd,rev(shd)),
+           c(shd, een ) )
> clr
[1] "#FF0000" "#FF5E5E" "#FFAAAA" "#FFEEEE" "#EEEEFF" "#AAAAFF" "#5E5EFF" "#0000FF"
> # Note: This uses the undocumented feature that if the first
> # number in a column is negative this is taken as the left endpoint of
> # the bar. So c(-m,m,f) is a bar starting at -m, and a division at
> # -m+m and an upper end at -m+m+f. Coloring is from the top, that is
> # the part stretching from -m+m to -m+m+f get the first color
> oo <- c("mort","inc","const","obs")
> lim <- 6
> draw.pyr <-
+ function(pp)
+ {
+ par( mar=c(3,3,3,0), mgp=c(3,1,0)/1.6, las=1 )
+ barplot( height=t( cbind( -apply(prn[,pp,"M","apc",      ], 1, sum),
+                             prn[,pp,"M","apc",      oo ],
+                             prn[,pp,"F","apc",rev(oo)] ) ) / 1000,
+         horiz=TRUE, col=clr[c(1,8:2)], border=rep("transparent",8),
+         space=0, axes=FALSE, names.arg=rep("",dim(prn)[1]),
+         xlim=c(-1,1)*lim*1.05,
+         xlab="Persons in 1 year class (1000s)",ylab="Age")
+ abline(h=seq(0,100,5),
+        v=seq(-lim,lim,0.5),
+        col="white")
+ axis( side=1, at=seq(-lim,lim,1), labels=abs(seq(-lim,lim,1)) )
+ axis( side=2, at=seq(0,100,20) )
+ axis( side=2, at=seq(0,100,5), labels=NA, tcl=-0.3 )
+ tt <- addmargins( apply( prn[,pp,,"apc",,2:3, sum ), 2 )
+ nn <- tt / tt[,5] * 100
+ ppos <- seq(1,5.9,,5)-0.1
+ npos <- -rev(ppos)
+ mtext( pp, at=-lim, adj=1.8, line=2, cex=1.2, font=1 )
+ mtext( c(lg<- c("Mort","Inc","Imbal","Org","All"),rev(lg)),
+       at=c(npos,ppos), col="black", cex=0.99, line=2 )
+ mtext( formatC(tt["M",1:5],0,,"f",,,","),
+       at=npow, col="blue", line=1, cex=0.99 )
+ mtext( formatC(tt["F",5:1],0,,"f",,,","),
+       at=ppos, col="red" , line=1, cex=0.99 )
+ mtext( formatC(nn["M",1:4],1,4,"f"),
+       at=npow[1:4], col="blue", line=0, cex=0.99 )
+ mtext( formatC(nn["F",4:1],1,4,"f"),
+       at=ppos[2:5], col="red" , line=0, cex=0.99 )
```

```

+ mtext( "N", at=0, line=1, cex=0.99 )
+ mtext( "%", at=0, line=0, cex=0.99 )
+ }
> pdf( "comp-DMpr-film.pdf", width=9, height=6 )
> for( pp in paste(1996:2016) ) draw.pyr(pp)
> dev.off()

null device
      1

> for( pp in paste(1996:2016) )
+ {
+ pdf( paste("comp-DMpr-", pp, ".pdf", sep=""), width=8, height=6 )
+ draw.pyr(pp)
+ dev.off()
+ }

```

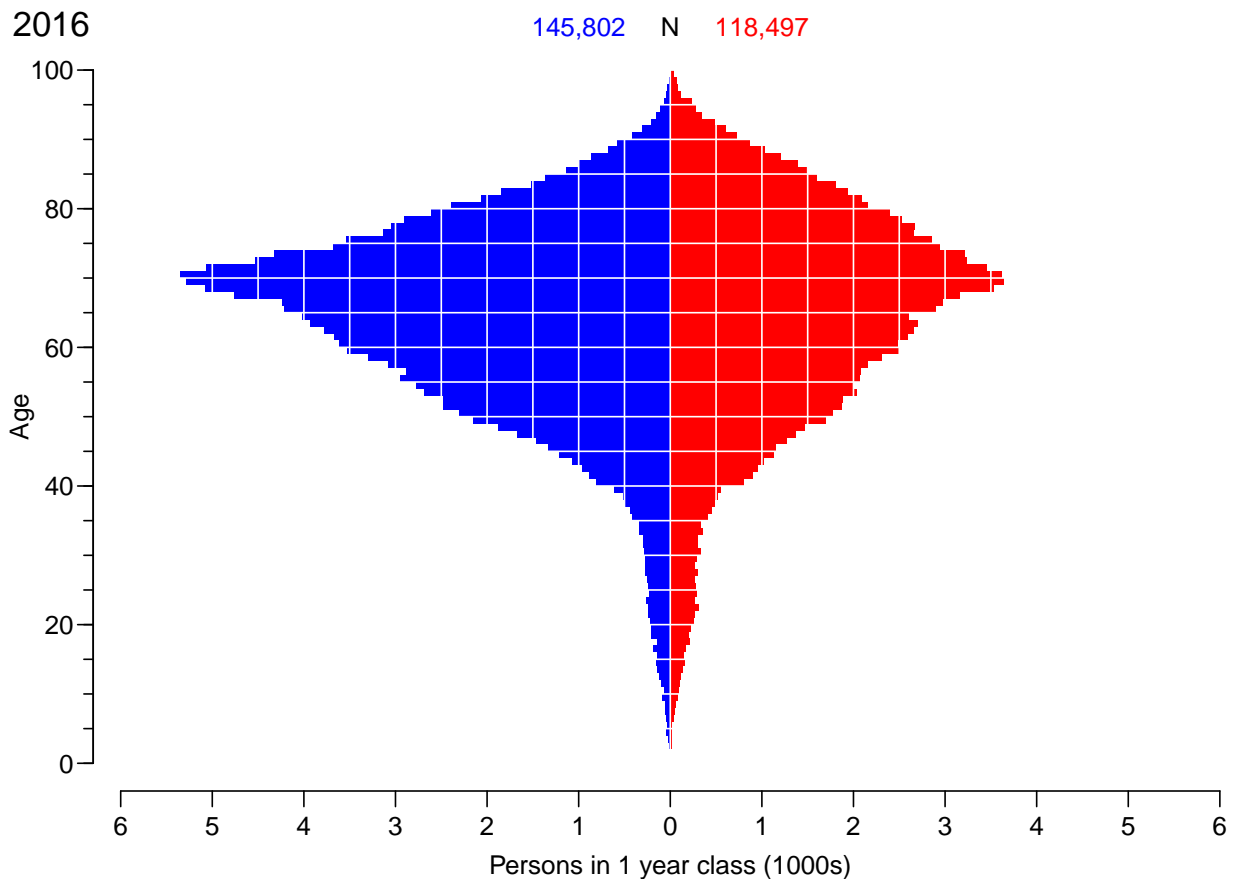


Figure 5.5: *Empirical age-distribution of the diabetes cases in Denmark as of 1.1.2016.*

5.5 Timetrend in the components

It is of course also of interest to see how large a fraction of the DM-patients in various ages that can be attributed to the different components at different times.

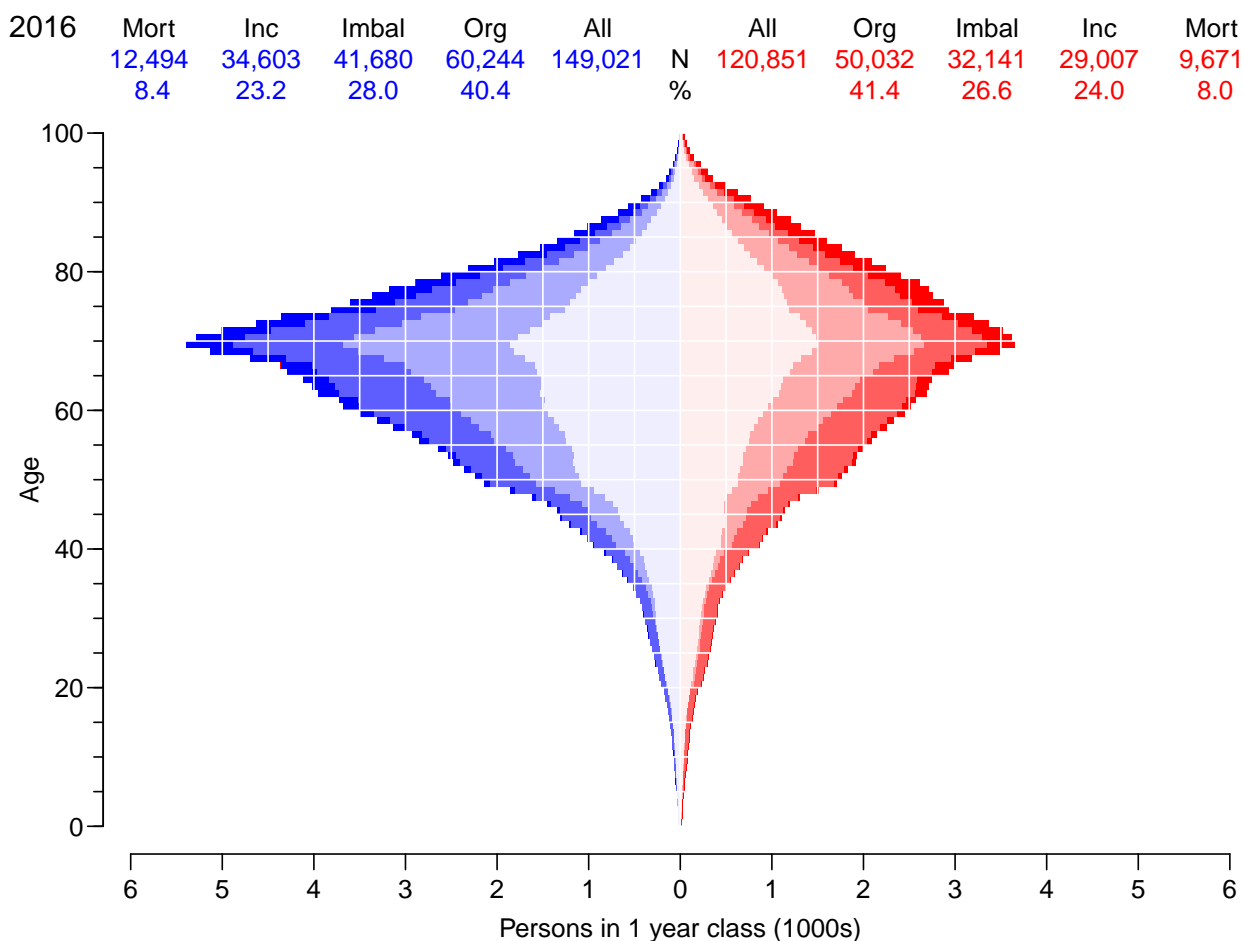


Figure 5.6: Age-distribution of the predicted no. of diabetes cases in Denmark as of 1.1.2016, subdivided by the components of disease prevalence: *Mort*: decrease in mortality, *Inc*: increase in incidence, *Imbal*: constant rates from 1996 (non-steady-state imbalance between incidence and mortality in 1996, *Org*: age-specific prevalence in 1996.

```
> str( prv )
num [1:1200, 1:253, 1:2, 1:2, 1:7] 0 0.00063 0.000635 0.000639 0.000644 ...
- attr(*, "dimnames")=List of 5
..$ a : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.2916666666666666
..$ t : chr [1:253] "1996" "1996.083333333333" "1996.166666666667" "1996.25" ...
..$ sex : chr [1:2] "M" "F"
..$ mod : chr [1:2] "ap" "apc"
..$ what: chr [1:7] "obs" "m-fix" "i-fix" "all-f" ...
> dimnames( prv )[[5]]
[1] "obs" "m-fix" "i-fix" "all-f" "mort" "inc" "const"
```

We extract the prevalences at ages 60, 70 and 80, and compute the fractions of all DM patients attributable to each component:

```
> aloc <- match( 6:8*10, floor(as.numeric(dimnames(prv)[[1]])) )
> ptrend <- ( prv[aloc,, "apc", -(2:4)] + prv[aloc-1,, "apc", -(2:4)] )/2
> str( ptrend )
num [1:3, 1:253, 1:4] 0.0417 0.052 0.0606 0.0419 0.0524 ...
- attr(*, "dimnames")=List of 4
```

```

..$ a    : chr [1:3] "60.0416666666667" "70.0416666666667" "80.0416666666667"
..$ t    : chr [1:253] "1996" "1996.083333333333" "1996.166666666667" "1996.25" ...
..$ sex  : chr [1:2] "M" "F"
..$ what: chr [1:4] "obs" "mort" "inc" "const"

> # Fraction of all DM at each age
> ptrend[,,,] <- ptrend[,,,]/ptrend[,,,rep("obs",4)]
> # Reduce to fraction attributable to org. prevalence
> ptrend[,,,"obs"] <- ptrend[,,,"obs" ] -
+                       ptrend[,,,"mort" ] -
+                       ptrend[,,,"inc" ] -
+                       ptrend[,,,"const"]

```

We can now plot the stacked fractions of the components for ages 60, 70 and 80 separately for men and women:

```

> p.pt <- as.numeric(dimnames(ptrend)[[2]])
> par( mfcol=c(3,2), mar=c(0,0,0,2), oma=c(3,4,1,0), mgp=c(3,1,0)/1.6, las=1, bty="n" )
> for( sx in c("M","F") )
+ for( ag in 1:3 )
+ {
+ plot( NA, xlim=c(1996,2016), ylim=0:1*100,
+       xaxs="i", yaxs="i", xaxt="n", yaxt="n", xlab="", ylab="" )
+ #if( sx=="M" ){
+ #axis( side=2 )
+ #axis( side=2, at=1:9*10, labels=NA ) }
+ if( ag==3 ){
+ axis( side=1 )
+ axis( side=1, at=1:20+1996, labels=NA, tcl=-0.3 ) }
+ polygon( c(p.pt,rev(p.pt)), c( ptrend[ag,,sx,"mort"] *100,
+                               rev(ptrend[ag,,sx,"mort"])* 0),
+         col=clr[if(sx=="M") 8 else 1], border="transparent" )
+ polygon( c(p.pt,rev(p.pt)), c(ptrend[ag,,sx,"mort"],
+                               rev(ptrend[ag,,sx,"mort"]+
+                                   ptrend[ag,,sx,"inc" ]))*100,
+         col=clr[if(sx=="M") 7 else 2], border="transparent" )
+ polygon( c(p.pt,rev(p.pt)), c(ptrend[ag,,sx,"mort"]+
+                               ptrend[ag,,sx,"inc" ],
+                               rev(ptrend[ag,,sx,"mort"]+
+                                   ptrend[ag,,sx,"inc" ]+
+                                   ptrend[ag,,sx,"const"]))*100,
+         col=clr[if(sx=="M") 6 else 3], border="transparent" )
+ abline( v=seq(2000,2015,5), h=1:9*10, col="white" )
+ axis( side=4, at=1:7*10, tcl=-0.5 )
+ axis( side=4, at=0:14*5, tcl=-0.4, labels=NA )
+ axis( side=4, at=0:70 , tcl=-0.2, labels=NA )
+ text( 1997, 75, paste( if(sx=="F") "Women\n" else "Men\n", "age", (6:8*10)[ag] ),
+       cex=1.2, font=2, adj=0 )
+ box(col=gray(0.7),bty="o")
+ }
> mtext( side=1, "Date", line=2, outer=TRUE, cex=0.67 )
> mtext( side=2, "Prevalence component (%)", line=2, outer=TRUE, cex=0.67, las=0 )

```

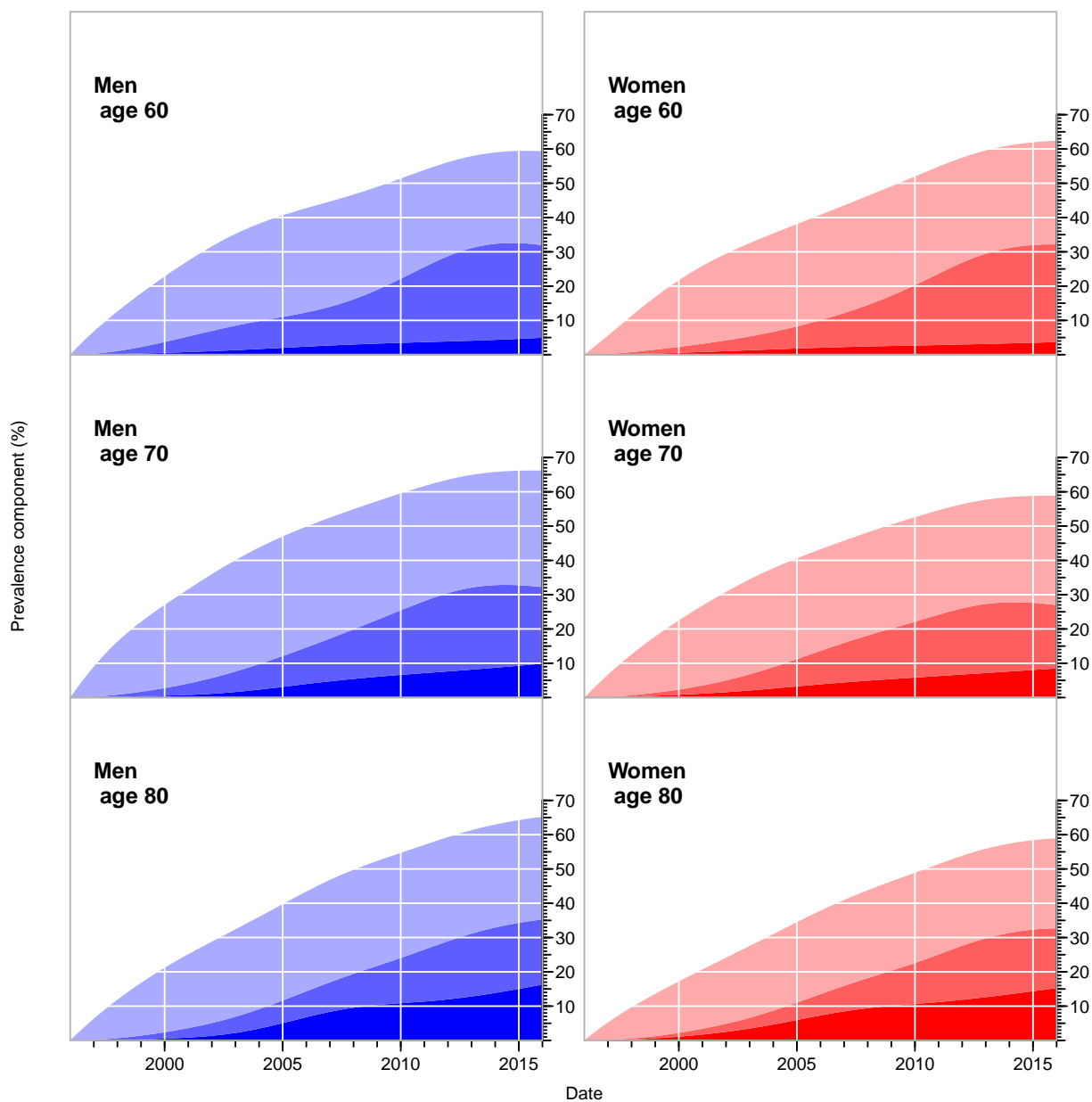


Figure 5.7: Fraction of the prevalent cases at different times attributable to a) declining mortality (bottom, full color), b) increasing incidence (middle, pale color) and c) prevalence/mortality imbalance at 1996 (top, weak color). The white area above the curves corresponds to the fraction of the cases that would have been around if incidence and mortality rates had remained as in 1996.

Chapter 6

Predicting prevalence of diabetes

In the previous chapter we predicted the number of prevalent cases, or more precisely the age-specific prevalences of DM over the period 1996 – 2016 under different scenarios.

In this chapter we shall essentially repeat this for the period 2016–2030, using three different scenarios:

- Use the naively predicted rates from the APC-model with natural splines — the “**apc**” component if the rate-arrays.

This will give a prediction of the numbers which is the least credible.

- Use the attenuated rates — the “**att**” component of the rate-arrays.

```
> library( Epi )
> library( splines )
> clear()
```

6.1 Predicted rates

In the previous chapter we did predictions of the incidence and mortality rates, and used prevalences at 1996-01-01 as starting point to predict prevalences at 2016-01-01.

Here we shall do the same as in the exercise assessing the components of prevalence, except that we will start with the observed (smoothed) age-specific prevalences at 2016-01-01 and then use the three different scenarios laid out above to predict the prevalences each year till 2030.

First we load the estimated / predicted rates

```
> load( file="../data/rateEsts.Rda" )
```

We shall use the simulation scheme to predict the course of DM prevalence development in the population under various scenarios of mortality and incidence development. So we set up various structures to hold results and clarify calculations:

pr.fit — array of predicted age-specific prevalences at 1995-01-01 to 2016-01-01, smoothed by natural splines. This is actually derived in the section on prevalence:

```
> load( file="../data/prFit.Rda" )
> str( pr.fit )
```



```

num [1:2, 1:1200, 1:21] 0.000626 0.000579 0.00063 0.000583 0.000635 ...
- attr(*, "dimnames")=List of 3
..$ sex: chr [1:2] "M" "F"
..$ A : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.2916666666666667" ...
..$ P : chr [1:21] "1996" "1997" "1998" "1999" ...

```

TR — array of transition probabilities between states Well and DM and Death. Transition probabilities are computed under the 3 different scenarios combining mortality and incidence rates either as they actually developed 1996–2015. These refer to intervals of length `int` and are therefore labeled on the period dimension by the midpoint of these, a total of $15/\text{int}$.

prv — array of predicted prevalences based on the initial prevalences at 2016-01-01 and the transition probabilities as put in **TR**. The scenario dimension refers to the 3 scenarios: “lin”, “att” and “fix”. Moreover, the period dimension is expanded by one relative to that in **TR**, since this refer to points in time and not time intervals.

prn — array of predicted *number* of DM patients in one-year age classes at the 1 January each year. So the same structure as **prv**, but with substantially fewer entries.

6.2 Transition probabilities

In order to get the predicted *number* of persons by age, period and prediction type, we need the (1-step) transition matrices at all combinations of age (a) and date (p), this is put in array. But we will only need the rates from 2016 and onward, so we restrict the arrays with the rates to this period:

```

> dimnames( Lambda )[[2]][240:241]
[1] "2015.9583333333333" "2016.041666666667"
> rLambda <- Lambda[,-(1:240),,]
> rMu.W <- Mu.W [,-(1:240),,]
> rMu.DM <- Mu.DM[,-(1:240),,]
> states <- c("Well", "DM")
> TR <- NArray( c( dimnames( rLambda ),
+                 list( from = states,
+                       to = states ) ) )
> dimnames( TR )[[4]][1] <- "fix"
> str( TR )

logi [1:1200, 1:288, 1:2, 1:6, 1:2, 1:2] NA NA NA NA NA NA ...
- attr(*, "dimnames")=List of 6
..$ a : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.2916666666666667" ...
..$ p : chr [1:288] "2016.041666666667" "2016.125" "2016.208333333333" "2016.291666666667" ...
..$ sex: chr [1:2] "M" "F"
..$ mod: chr [1:6] "fix" "apc" "att" "i20" ...
..$ from: chr [1:2] "Well" "DM"
..$ to : chr [1:2] "Well" "DM"

```

The situation where both the mortality rates and incidence rates are fixed at the 2016 level (“fix”) is trivial, because transition probabilities in that case only depend on age and not on period.

In order to fill `TR`, we need the cumulative incidences over intervals of length `int`. But these were exactly the ones we predicted in the previous sections by setting the person years equal to `int` in the data frame supplied to the `newdata` argument.

So we can now compute the one-`int`-step transition matrices for every combination of `a.pt` and `p.pt`, in steps of `int` (in this case 0.0833 year).

```
> TR[,,,,"Well","Well"] <- exp(-rLambda-rMu.W)
> TR[,,,,"Well","DM" ] <- 1 - exp(-rLambda)
> TR[,,,,"DM" ,"Well"] <- 0
> TR[,,,,"DM" ,"DM" ] <- exp(-rMu.DM)
```

Note that we here fill in the transition probabilities from the age-period model in the `fix` slot if the 4th dimension of `TR`, but we overwrite this with the rates as of 2016:

```
> TR[,,,,"fix","Well","Well"] <- exp(-rLambda[,rep(1,dim(TR)[2]),,"apc"]-rMu.W[,rep(1,dim(TR)[2]),,"apc"])
> TR[,,,,"fix","Well","DM" ] <- 1 - exp(-rLambda[,rep(1,dim(TR)[2]),,"apc"])
> TR[,,,,"fix","DM" ,"Well"] <- 0
> TR[,,,,"fix","DM" ,"DM" ] <- exp(- rMu.DM[,rep(1,dim(TR)[2]),,"apc"])
```

Likewise we fill in the entries referring to the three scenarios of increasing incidence rates — note that it is only for the incidence rates we impose an increase in rates:

```
> TR[,,,,"i20","Well","Well"] <- exp(-rLambda[,,,,"i20"]-rMu.W[,,,,"att"])
> TR[,,,,"i20","Well","DM" ] <- 1 - exp(-rLambda[,,,,"i20"])
> TR[,,,,"i20","DM" ,"Well"] <- 0
> TR[,,,,"i20","DM" ,"DM" ] <- exp(- rMu.DM[,,,,"att"])
> TR[,,,,"i25","Well","Well"] <- exp(-rLambda[,,,,"i25"]-rMu.W[,,,,"att"])
> TR[,,,,"i25","Well","DM" ] <- 1 - exp(-rLambda[,,,,"i25"])
> TR[,,,,"i25","DM" ,"Well"] <- 0
> TR[,,,,"i25","DM" ,"DM" ] <- exp(- rMu.DM[,,,,"att"])
> TR[,,,,"i30","Well","Well"] <- exp(-rLambda[,,,,"i30"]-rMu.W[,,,,"att"])
> TR[,,,,"i30","Well","DM" ] <- 1 - exp(-rLambda[,,,,"i30"])
> TR[,,,,"i30","DM" ,"Well"] <- 0
> TR[,,,,"i30","DM" ,"DM" ] <- exp(- rMu.DM[,,,,"att"])
```

Finally, note that we have not included the “Dead” state in the calculations, because we only bother about the *fraction* of diabetes patients in each age class at each time-point. So the probabilities we compute do not sum to 1 within the “from” states; we only compute how many of the persons alive that end up being alive at the next time point

We have now collected the transition probabilities between “Well” and “DM” as well as the probabilities of remaining in each of these, all referring to a duration of `int`, a total of almost 5 mil. numbers:

```
> str( TR )
num [1:1200, 1:288, 1:2, 1:6, 1:2, 1:2] 1 1 1 1 1 ...
- attr(*, "dimnames")=List of 6
..$ a : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.2916666666666667"
..$ p : chr [1:288] "2016.041666666667" "2016.125" "2016.208333333333" "2016.291666666667"
..$ sex : chr [1:2] "M" "F"
..$ mod : chr [1:6] "fix" "apc" "att" "i20" ...
..$ from: chr [1:2] "Well" "DM"
..$ to : chr [1:2] "Well" "DM"

> prod( dim(TR) )
[1] 16588800

> save( TR, file="../data/TRf.Rda" )
```

6.3 Prediction of the observed prevalences

Note that we do not need to predict the population size; we can get away with only predicting the prevalences as fractions. When we multiply the fraction of persons in states (Well,DM) with the transition matrix, we get fraction of the persons in the previous state that are in states (Well,DM), which does not sum to 1 (because of the ones dying), so we must rescale to prevalence age in each step.

First we set up an array to hold the predicted prevalences under different scenarios. Later we shall also compute the fraction of the prevalences that are attributable to trends in mortality and incidence as well as to the non-stationarity of the rates/prevalences as of 1995, so we put in three extra levels of the last dimension, and one extra levels of the period dimension because we want to predict to the end of the last period too (or, to put it differently, we need an extra first level to hold the starting prevalences as of 2016-01-01).

```
> dpr <- dimnames(TR)[1:4]
> names(dpr)[2] <- "t"
> dpr[["t"]] <- t.pt[-(1:240)]
> prv <- NArray(dpr)
> str(prv)
logi [1:1200, 1:289, 1:2, 1:6] NA NA NA NA NA NA ...
- attr(*, "dimnames")=List of 4
..$ a : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.2916666666666667"
..$ t : chr [1:289] "2016" "2016.083333333333333" "2016.166666666667" "2016.25" ...
..$ sex: chr [1:2] "M" "F"
..$ mod: chr [1:6] "fix" "apc" "att" "i20" ...
> prod(dim(prv))
[1] 4161600
```

Thus we must make a loop that updates the prevalences at 2016-01-01 to those at subsequent times, but first we must initialize the prevalences as modeled on 2016-01-01, as well as the prevalences at ages 0 (the new-born) that we set to 0; note that we exploit the column major storage of arrays to get prv filled with identical values across the two last dimensions (model and scenario).

```
> load(file="../data/prFit.Rda")
> str(pr.fit)
num [1:2, 1:1200, 1:21] 0.000626 0.000579 0.00063 0.000583 0.000635 ...
- attr(*, "dimnames")=List of 3
..$ sex: chr [1:2] "M" "F"
..$ A : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.2916666666666667"
..$ P : chr [1:21] "1996" "1997" "1998" "1999" ...
> # Smoothed prevalences at 2016-01-01 - the starting values
> for(sx in c("M", "F")) prv[, "2016", sx, ] <- pr.fit[sx, "2016"]
> # Prevalences at age 0
> prv[1, ,, ] <- 0
> tt <- prv[c(1:2, floor(dim(prv)[1]/1.5)+1:3), 1:3, , ]
> dimnames(tt)[[1]] <- round(as.numeric(dimnames(tt)[[1]]), 2)
> dimnames(tt)[[2]] <- round(as.numeric(dimnames(tt)[[2]]), 2)
> round(ftable(tt, col.vars=3:4)*100, 1)
      sex      M
      mod fix  apc  att  i20  i25  i30  sex      F
      mod fix  apc  att  i20  i25  i30  mod fix  apc  att  i20  i25  i30
a      t
0.04  2016      0.0  0.0  0.0  0.0  0.0  0.0  0.0  0.0  0.0  0.0  0.0  0.0  0.0
```

```

      2016.08      0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0
      2016.17      0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0 0.0
0.12 2016         0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1 0.1
      2016.08      NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA
      2016.17      NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA
66.71 2016       13.8 13.8 13.8 13.8 13.8 13.8 9.0 9.0 9.0 9.0 9.0
      2016.08      NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA
      2016.17      NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA
66.79 2016       13.8 13.8 13.8 13.8 13.8 13.8 9.1 9.1 9.1 9.1 9.1
      2016.08      NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA
      2016.17      NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA
66.88 2016       13.9 13.9 13.9 13.9 13.9 13.9 9.1 9.1 9.1 9.1 9.1
      2016.08      NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA
      2016.17      NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA  NA

```

So now it is checked that we have put the initial values correctly into `prv`. Then we can compute the predicted prevalences under the different scenarios. We take the fraction of the population in age class `ia` at time `ip` that end up as diabetes patients at time `ip+1` (and hence in age class `ia+1`), and divide by the fraction of all that remain alive, which is the diabetes patients, *plus* those who survive free of diabetes:

```

> system.time(
+ for( ip in 1:(dim(prv)[2]-1) )
+ for( ia in 1:(dim(prv)[1]-1) )
+ prv[ia+1,ip+1,,] <-
+ (   prv[ia,ip,,] * TR[ia,ip,, "DM" , "DM" ]
+   +(1-prv[ia,ip,,] * TR[ia,ip,, "Well", "DM" ] ) ) /
+ (   prv[ia,ip,,] * TR[ia,ip,, "DM" , "DM" ]
+   +(1-prv[ia,ip,,] * TR[ia,ip,, "Well", "DM" ]
+   +(1-prv[ia,ip,,] * TR[ia,ip,, "Well", "Well" ] )
+ )
      user  system elapsed
      12.94    0.00   12.94

```

Note that the code above is particularly simple because we only need to compute the prevalence at the next date and age. If we had had a more elaborate model with, say complications states, the calculations in the loop would have been a matrix-multiplication updating the state-distribution, but this simplification would have been at the expense of another three loop-levels, namely over the the three last dimensions of the `prv` array.

We can then show a few of the predicted prevalences in (

```

> round( prv[1:4,1:2,1,1,drop=F]*100, 3 )
, , sex = M, mod = fix

      t
a      2016 2016.083333333333
0.04166666666666667 0.000      0.000
0.125      0.121      0.001
0.2083333333333333 0.121      0.121
0.29166666666666667 0.122      0.122

> save( a.pt, prv, file="../data/prv-comp.Rda" )
> load(      file="../data/prv-comp.Rda" )

```

6.4 The actual numbers of diabetes patients in Denmark

In the previous section we only looked at the age-specific prevalences, because these are the quantities that are driven by the incidence and mortality rates. However, it is also of interest to see how the actual number of diabetes patients would have looked under the different scenarios.

To show the number of patients we set up an array `prn` with `structure` (but not `extent`) as `prv` to hold the number of diabetes patients by category, assuming the age-distribution in the population to be as actually observed (that is as extracted from Statistics Denmark). However `prn` will have 100 age-classes rather than 1200 ($100/\text{int}$), and only 15 dates (2016–2030): `prv`.

```
> dn <- dimnames(prv)
> dn[[1]] <- 0:99
> dn[[2]] <- 2016:2030
> prn <- NArray( dn )
> table( (prv>0) + (prv>=0) )
      1      2
3468 4158132
> str( prv )
num [1:1200, 1:289, 1:2, 1:6] 0 0.00121 0.00121 0.00122 0.00123 ...
- attr(*, "dimnames")=List of 4
..$ a : chr [1:1200] "0.0416666666666667" "0.125" "0.2083333333333333" "0.291666666666667" ...
..$ t : chr [1:289] "2016" "2016.083333333333" "2016.16666666667" "2016.25" ...
..$ sex: chr [1:2] "M" "F"
..$ mod: chr [1:6] "fix" "apc" "att" "i20" ...
> str( prn )
logi [1:100, 1:15, 1:2, 1:6] NA NA NA NA NA NA NA ...
- attr(*, "dimnames")=List of 4
..$ a : chr [1:100] "0" "1" "2" "3" ...
..$ t : chr [1:15] "2016" "2017" "2018" "2019" ...
..$ sex: chr [1:2] "M" "F"
..$ mod: chr [1:6] "fix" "apc" "att" "i20" ...
```

In order to fill in the numbers we use the estimates age-specific prevalences at 1st January each year, that is at the dates 2016-01-01,...,2030-01-01 in the entries along the `t`-dimension of `prv`. Moreover we want the prevalences for a 1 year age class rather than age-classes of length `int`. So we take the average prevalences from `prv` over each one-year age-interval. The vectors `wh.a` and `wh.p` will hold the number of the age and period classes from `prv` which have the desired prevalences (as proportions) that we will use for multiplication with the population figures:

```
> # Find the dates in the predicted prevalences prv that matches the
> # dates in prn where empirical rates are available.
> prv.p <- as.numeric( dimnames(prv)[["t"]] )
> prn.p <- as.numeric( dimnames(prn)[["t"]] )
> wh.p <- match( prn.p, prv.p )
> if( any(is.na(wh.p)) ) # Need to find approximate dates if they do not match
+ for( ip in 1:length(prn.p) )
+ {
+   dd <- abs( prn.p[ip]-prv.p )
+   wh.p[ip] <- (1:length(dd))[dd==min(dd)]
+ }
> wh.p
```

```

[1] 1 13 25 37 49 61 73 85 97 109 121 133 145 157 169
> prv <- pmax( prv, 0 )
> # Ages in the two arrays
> prv.a <- as.numeric( dimnames(prv)[["a"]] )
> prn.a <- as.numeric( dimnames(prn)[["a"]] )
> for( ip in 1:length(wh.p) )
+ for( ia in 1:length(prn.a) )
+ {
+ wh.a <- which( prn.a[ia]==floor(prv.a) )
+ prn[ia,ip,,] <- apply( prv[wh.a,wh.p[ip],,], 2:3, mean )
+ }
> range( prv )
[1] 0.000000 0.275464
> range( prn )
[1] 6.336452e-06 2.246479e-01

```

Now `prn` contains the prevalences (as fractions) for 100 age classes and the 15 dates. We need to multiply these prevalences by the population figures at these times.

6.5 Population forecast from DST

One prerequisite for the prediction is that we have some knowledge about the future population size of Denmark, so we have acquired this from the data bank of Statistics Denmark:

```

> bef <- read.csv2( "../data/bef2040.csv", header=TRUE )
> bef$sex <- ifelse( bef$sex==" ", NA, bef$sex )
> for( j in 1:3 )
+ for( i in 2:nrow(bef) )
+ if( is.na(bef[i,j]) ) bef[i,j] <- bef[i-1,j]
> table( bef$sex )
  2  3
3277 3277
> bef$sex <- factor( bef$sex, levels=3:2, labels=c("M","F") )
> bef <- subset( bef, !is.na(N) )
> addmargins( xtabs( N ~ P + sex, data = bef ), 2 )

```

	sex		
P	M	F	Sum
2016	2837887	2869364	5707251
2017	2863749	2891251	5755000
2018	2887236	2911306	5798542
2019	2906801	2928091	5834892
2020	2923339	2942471	5865810
2021	2938021	2955444	5893465
2022	2951822	2967966	5919788
2023	2965270	2980437	5945707
2024	2978491	2992896	5971387
2025	2991402	3005238	5996640
2026	3003925	3017277	6021202
2027	3016021	3028944	6044965
2028	3027617	3040154	6067771
2029	3038690	3050838	6089528
2030	3049243	3060972	6110215
2031	3059248	3070527	6129775

```

2032 3068731 3079506 6148237
2033 3077703 3087895 6165598
2034 3086217 3095728 6181945
2035 3094266 3102972 6197238
2036 3101872 3109674 6211546
2037 3109101 3115831 6224932
2038 3115961 3121489 6237450
2039 3122496 3126666 6249162
2040 3128698 3131429 6260127

```

```
> str( bef )
```

```

'data.frame':      6300 obs. of  4 variables:
 $ sex: Factor w/ 2 levels "M","F": 1 1 1 1 1 1 1 1 1 1 ...
 $ A  : int  0 0 0 0 0 0 0 0 0 0 ...
 $ P  : int  2016 2017 2018 2019 2020 2021 2022 2023 2024 2025 ...
 $ N  : int  30049 30897 31660 32444 33259 34197 35259 36232 36895 37240 ...

```

We need the population figures in an array of the same shape as (some of the dimensions of) `prv`

```

> pop <- xtabs( N ~ A + P + sex,
+             data = subset( bef, A<100 & P<2031 ) )
> str( pop )
int [1:100, 1:15, 1:2] 30049 29750 29459 30834 31038 33643 33495 34918 34180 34688 ...
- attr(*, "dimnames")=List of 3
..$ A : chr [1:100] "0" "1" "2" "3" ...
..$ P : chr [1:15] "2016" "2017" "2018" "2019" ...
..$ sex: chr [1:2] "M" "F"
- attr(*, "class")= chr [1:2] "xtabs" "table"
- attr(*, "call")= language xtabs(formula = N ~ A + P + sex, data = subset(bef, A < 100 &
> str( prn )
num [1:100, 1:15, 1:2, 1:6] 0.00114 0.00133 0.00142 0.00152 0.00162 ...
- attr(*, "dimnames")=List of 4
..$ a : chr [1:100] "0" "1" "2" "3" ...
..$ t : chr [1:15] "2016" "2017" "2018" "2019" ...
..$ sex: chr [1:2] "M" "F"
..$ mod: chr [1:6] "fix" "apc" "att" "i20" ...
> dmp <- prn
> for( im in dimnames(dmp)[[4]] ) dmp[,,,im] <- prn[,,,im] * pop
> save( dmp, file="dmp.Rda" )
> load( file="dmp.Rda" )

```

First we draw simple population pyramids of the age-distribution of the diabetes patients in Denmark, as predicted under different scenarios:

```

> # Note: This uses the undocumented feature that if the first
> # number in a column is negative this is taken as the left endpoint of
> # the bar. So c(-m,m,f) is a bar starting at -m, and a division at
> # -m+m(=0) and an upper end at -m+m-f. Coloring is from the top, that is
> # the part stretching from -m+m to -m+m+f get the first color
> pp <- "2016"
> lim <- 6
> clr <- c("red","blue")
> draw.dmp <-
+ function(pp,wh)
+ {
+ par( mar=c(3,3,3,0), mgp=c(3,1,0)/1.6, las=1 )

```



```

+ barplot( height=t( cbind( -dmp[,pp,"M",wh],
+                          dmp[,pp,"M",wh],
+                          dmp[,pp,"F",wh] ) )/ 1000,
+         horiz=TRUE, col=clr,
+         border="transparent",space=0,axes=FALSE,
+         names.arg=rep("",dim(prn)[1]),
+         xlim=c(-1,1)*lim*1.05,
+         xlab="Persons in 1 year class (1000s)",ylab="Age")
+ abline(h=seq(0,100,5),
+        v=seq(-lim,lim,0.5),
+        col="white")
+ axis( side=1, at=seq(-lim,lim,1), labels=abs(seq(-lim,lim,1)) )
+ axis( side=1, at=seq(-lim,lim,0.5), labels=NA, tcl=-0.3 )
+ axis( side=2, at=seq(0,100,20) )
+ axis( side=2, at=seq(0,100,5), labels=NA, tcl=-0.3 )
+ mtext( pp, at=-lim, adj=1.2, cex=1.0, font=2 )
+ mtext( formatC(sum(dmp[,pp,"M",wh]),0,format="f",big.mark=","),
+        at=-1, col="blue", line=0, cex=0.99, adj=1 )
+ mtext( formatC(sum(dmp[,pp,"F",wh]),0,format="f",big.mark=","),
+        at= 1, col="red" , line=0, cex=0.99, adj=0 )
+ mtext( "N", at=0, line=0, cex=0.99 )
+ }
> pdf( "pred-inc-film.pdf", width=8, height=6 )
> for( pp in paste(2016:2030) ) draw.dmp(pp,"fix")
> dev.off()
null device
  1

> pdf( "pred-apc-film.pdf", width=8, height=6 )
> for( pp in paste(2016:2030) ) draw.dmp(pp,"apc")
> dev.off()
null device
  1

> pdf( "pred-att-film.pdf", width=8, height=6 )
> for( pp in paste(2016:2030) ) draw.dmp(pp,"att")
> dev.off()
null device
  1

> for( pp in paste(2016:2030) )
+ {
+ pdf( paste("pred-att-", pp, ".pdf", sep=""), width=8, height=6 )
+ draw.dmp(pp,"att")
+ dev.off()
+ }

> par( mfrow=c(2,2), mar=c(3,3,0,0), oma=c(0,0,2,0), mgp=c(3,1,0)/1.6, las=1 )
> draw.dmp("2016","att")
> draw.dmp("2020","att")
> draw.dmp("2025","att")
> draw.dmp("2030","att")
> mtext( "Incidence rate decrease attenuates from 2016", side=3, line=0, outer=TRUE )

> par( mfrow=c(2,2), mar=c(3,3,0,0), oma=c(0,0,2,0), mgp=c(3,1,0)/1.6, las=1 )
> draw.dmp("2016","fix")
> draw.dmp("2020","fix")
> draw.dmp("2025","fix")
> draw.dmp("2030","fix")
> mtext( "Incidence and mortality rates constant from 2016", side=3, line=0, outer=TRUE )

```



```

> par( mfrow=c(2,2), mar=c(3,3,0,0), oma=c(0,0,2,0), mgp=c(3,1,0)/1.6, las=1 )
> draw.dmp("2016","apc")
> draw.dmp("2020","apc")
> draw.dmp("2025","apc")
> draw.dmp("2030","apc")
> mtext( "Naive linear prediction from 2016", side=3, line=0, outer=TRUE )

> par( mfrow=c(2,2), mar=c(3,3,0,0), oma=c(0,0,2,0), mgp=c(3,1,0)/1.6, las=1 )
> draw.dmp("2016","i20")
> draw.dmp("2020","i20")
> draw.dmp("2025","i20")
> draw.dmp("2030","i20")
> mtext( "Incidence rates increase 2.0%/y from 2016, mortality decrease", side=3, line=0, o

> par( mfrow=c(2,2), mar=c(3,3,0,0), oma=c(0,0,2,0), mgp=c(3,1,0)/1.6, las=1 )
> draw.dmp("2016","i25")
> draw.dmp("2020","i25")
> draw.dmp("2025","i25")
> draw.dmp("2030","i25")
> mtext( "Incidence rates increase 2.5%/y from 2016, mortality decrease", side=3, line=0, o

> par( mfrow=c(2,2), mar=c(3,3,0,0), oma=c(0,0,2,0), mgp=c(3,1,0)/1.6, las=1 )
> draw.dmp("2016","i30")
> draw.dmp("2020","i30")
> draw.dmp("2025","i30")
> draw.dmp("2030","i30")
> mtext( "Incidence rates increase 3.0%/y from 2016, mortality decrease", side=3, line=0, o

```

6.6 Timetrend in prevalent number of DM patients

We would like to see the overall change in the number of diabetes patients, as recorded in the structure `dmp`

```

> DMall <- dmp[,,"M",] + dmp[,,"F",]
> DMcum <- apply( DMall, 2:3, cumsum )
> DMcum <- DMcum[c(1,1:100),,]
> DMcum[1,,] <- 0
> DMcum <- DMcum/1000
> str( DMcum )
num [1:101, 1:15, 1:6] 0 0.0663 0.1427 0.2242 0.3151 ...
- attr(*, "dimnames")=List of 3
..$ a : chr [1:101] "0" "0" "1" "2" ...
..$ t : chr [1:15] "2016" "2017" "2018" "2019" ...
..$ mod: chr [1:6] "fix" "apc" "att" "i20" ...

```

Finally we can plot the predicted numbers from the different scenarios:

```

> range( DMcum )
[1] 0.000 369.521

```

```

> ryr <- c(2016:2030,2030:2016)
> leg <- c("All rates fixed at 2016 level",
+         "Linear projection from 2016",
+         "Attenuated linear projection",
+         "DM incidence increasing 2.0%/y",
+         "DM incidence increasing 2.5%/y",
+         "DM incidence increasing 3.0%/y")
> par( mfrow=c(3,2), mar=c(2,1,0,3), mgp=c(3,1,0)/1.6, las=1, bty="n" )
> for( j in c(2,3,1,4:6) )
+ {
+ plot( NA,
+       xlim=c(2015.5,2030), xlab="", xaxt="n", xaxs="i",
+       ylim=c(0,420), yaxs="i", yaxt="n", ylab="" )
+ axis( side=4, at=0:4*100 )
+ axis( side=4, at=seq(0,4,1/4)*100, labels=NA, tcl=-0.3 )
+ axis( side=1, at=2015+1:3*5 )
+ axis( side=1, at=2016:2030, labels=NA, tcl=-0.3 )
+ for( i in 1:10 ) polygon( ryr, c( DMcum[1+(i-1)*10,,j],
+                                 rev( DMcum[1+ i *10,,j] ) ),
+                          col=gray( (17-i)/18 ), border=gray(0.8) ) #"transparent" )
+ abline( h=seq(50,400,50), v=c(2020,2025), col=gray(1), lty="16" )
+ for( i in seq(55,85,10) ) text( 2029, DMcum[paste(i),"2029",j],
+                               paste( i-5,"-",i+4,sep="" ) )
+ text( 2016, 400, paste( dimnames(DMcum)[[3]][j], ":", leg[j], sep="" ), adj=c(0,1) )
+ }

```

From figure 6.7 it appears that it is the decreasing incidence rates of diabetes that carries the major differences of more than 100,000 patients in 2030. The decrease in the number of incident cases is very recent; during the period 2012–2014 there was a drop and a very slight pick-up during 2015. Tabulations in the SAS-programs 08-mkFU (p. ??) and 09-mkPr (p. ??), show that there is no particular break in data where the decrease occur.

Thus the prediction of the number of future patients is crucially dependent on the tiny amount of information available about future diabetes incidence rates in the rather odd behaviour of the rates in 2012–2015.

However, even the quite brutal assumption of a pick up of increasing DM incidence rates by 2.5% per year will not bring the predicted number of patients over 400,000 in 2030. So to say that the number of diabetes patients is less than this in 2030 seems to be a fairly safe bet.

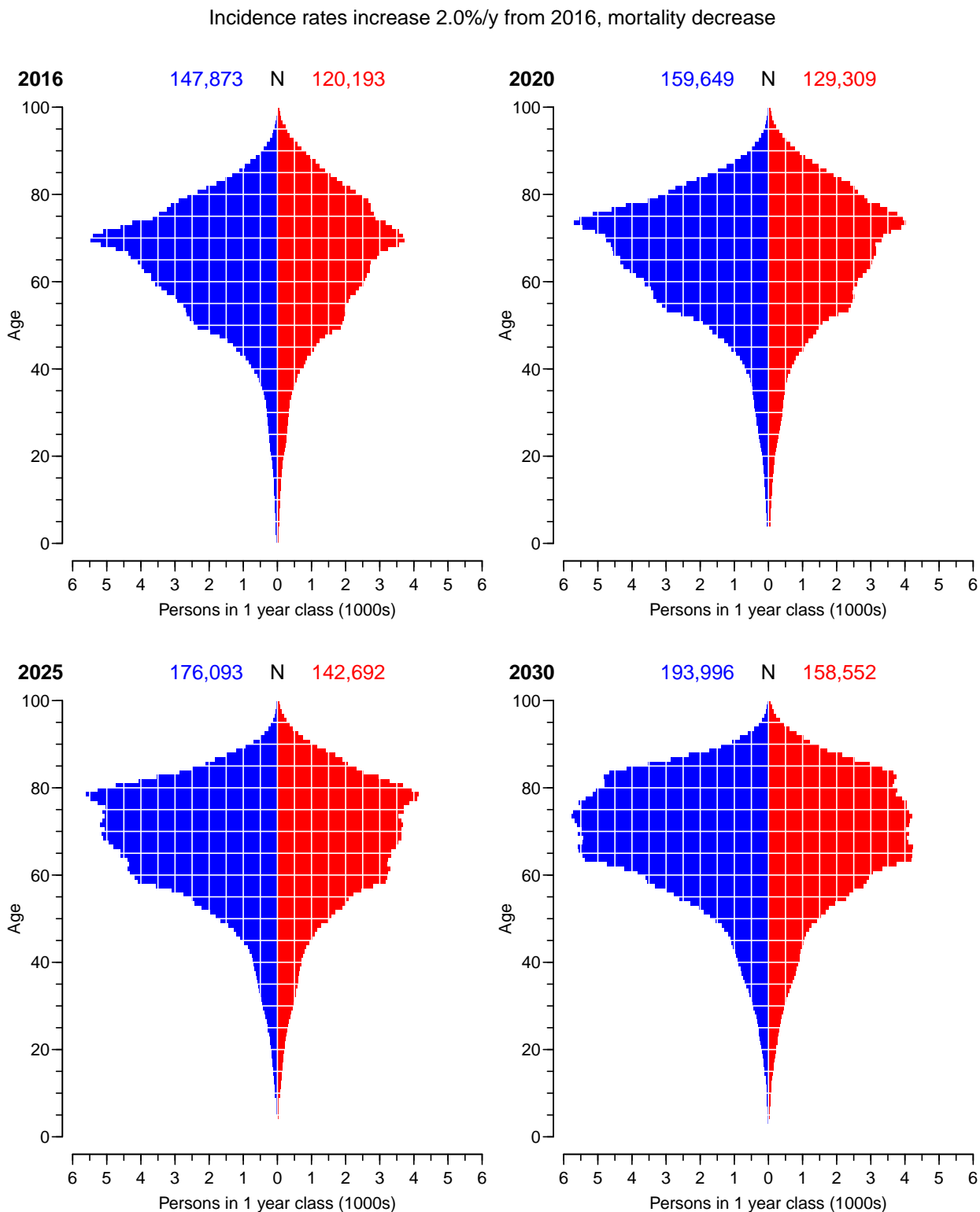


Figure 6.1: Empirical age-distribution of the diabetes cases in Denmark based on a unchanged mortality and increasing incidence rates (2.0%/y) from 2016.

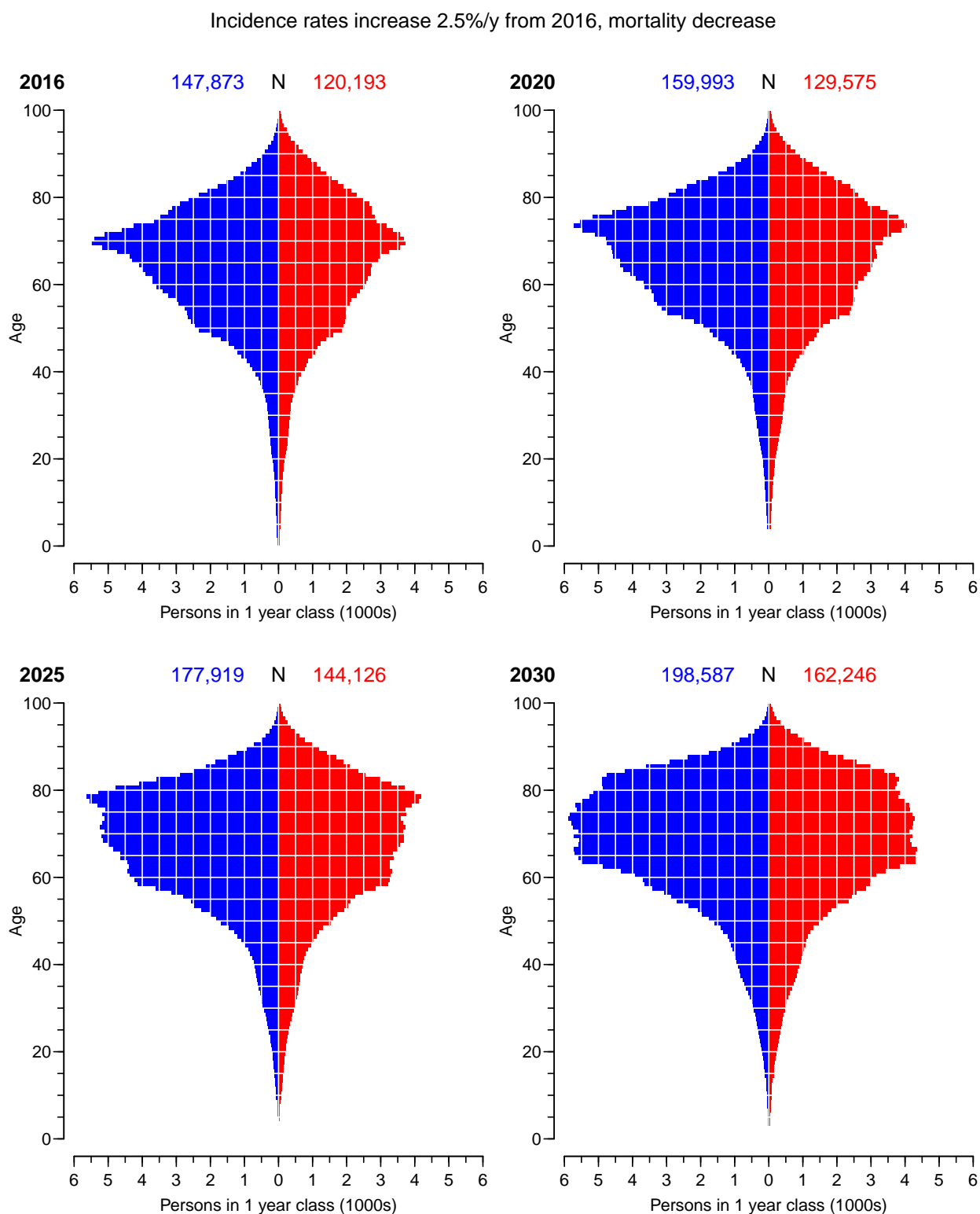


Figure 6.2: Empirical age-distribution of the diabetes cases in Denmark based on a unchanged mortality and increasing incidence rates (2.5%/y) from 2016.

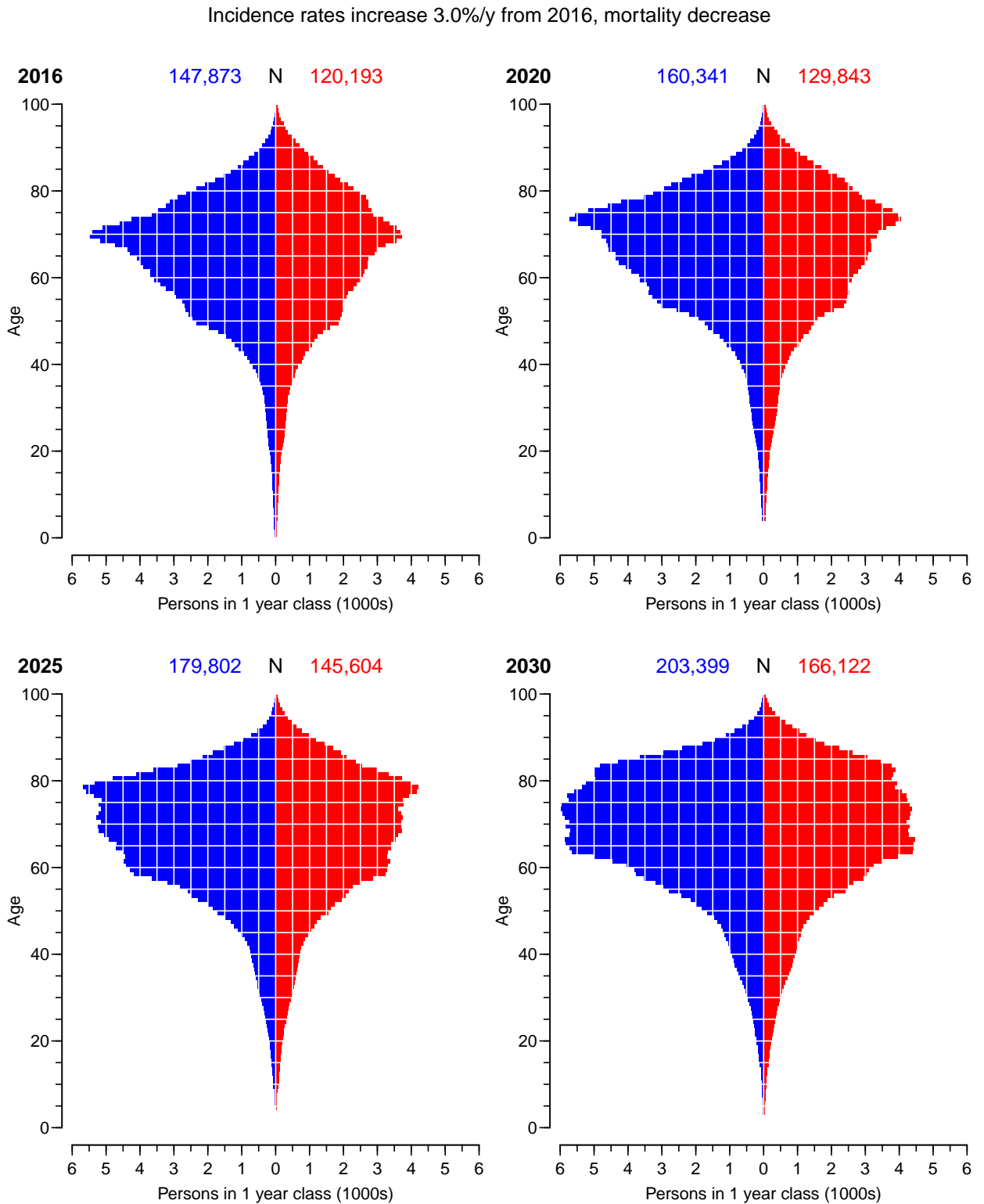


Figure 6.3: Empirical age-distribution of the diabetes cases in Denmark based on a unchanged mortality and increasing incidence rates (3.0%/y) from 2016.

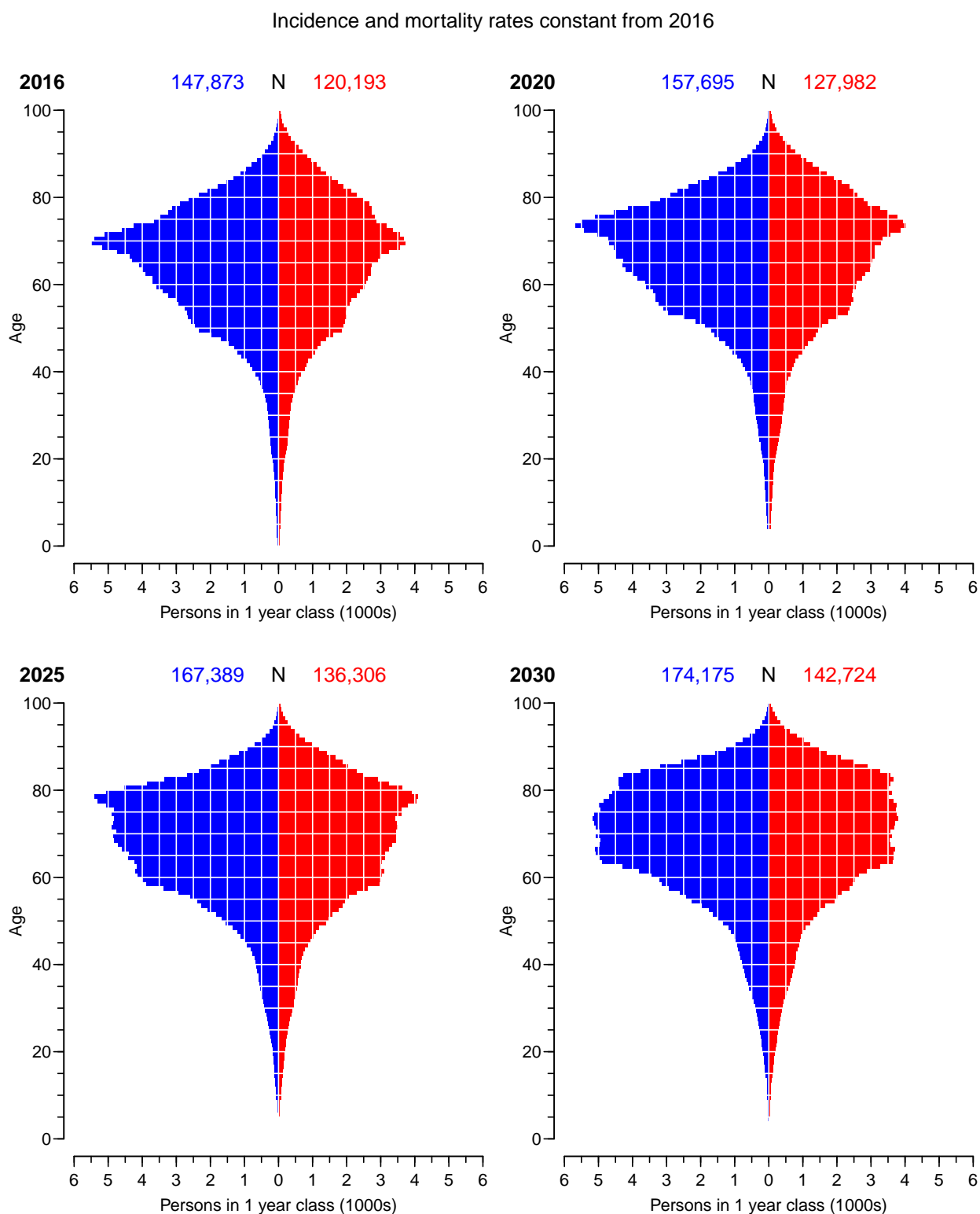


Figure 6.4: Empirical age-distribution of the diabetes cases in Denmark based on a unchanged mortality and incidence rates from 2016.

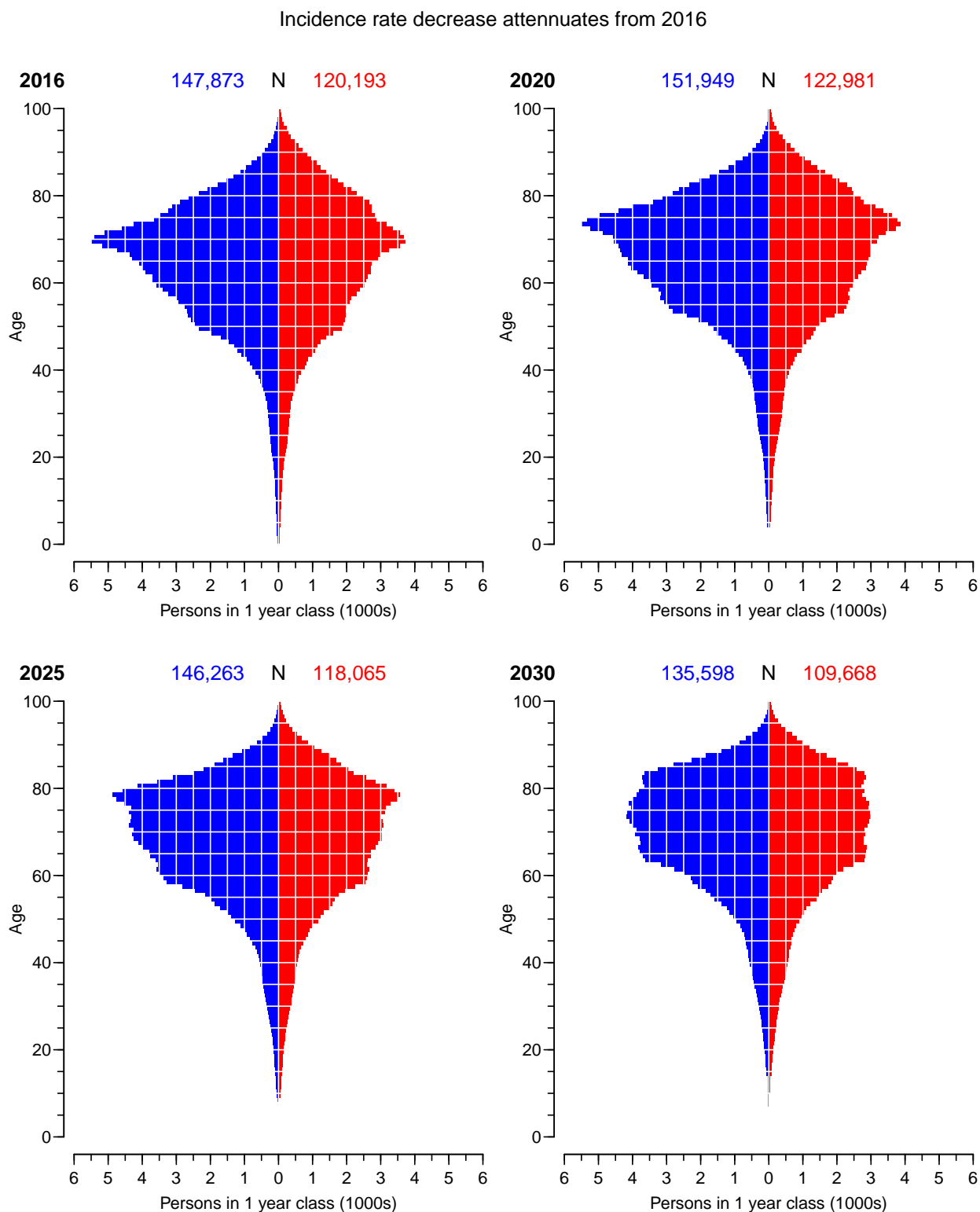


Figure 6.5: Empirical age-distribution of the diabetes cases in Denmark based on an attenuated linear prediction of mortality and incidence rates.

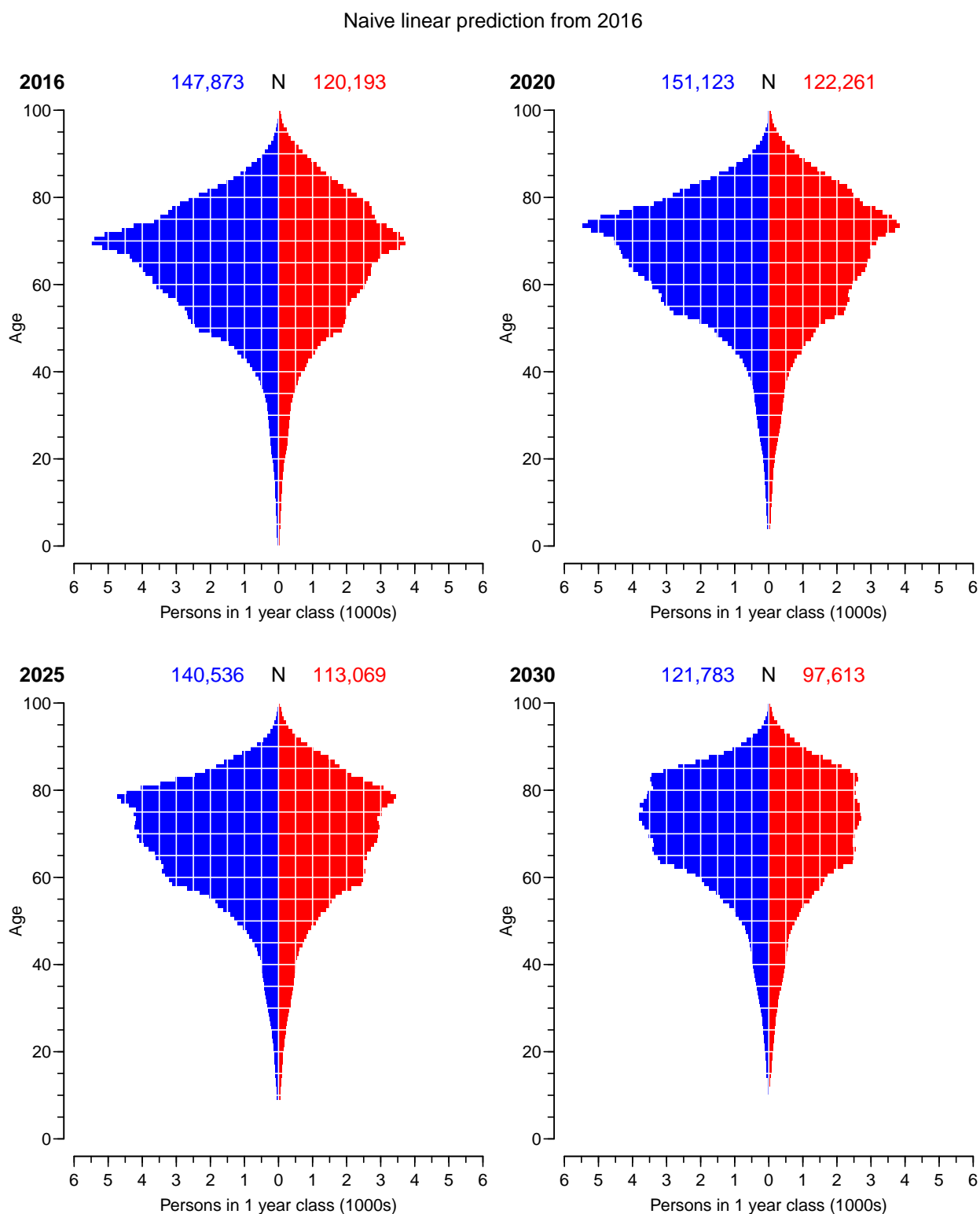


Figure 6.6: Empirical age-distribution of the diabetes cases in Denmark based on a linear prediction of mortality and incidence rates — mainly decreasing.

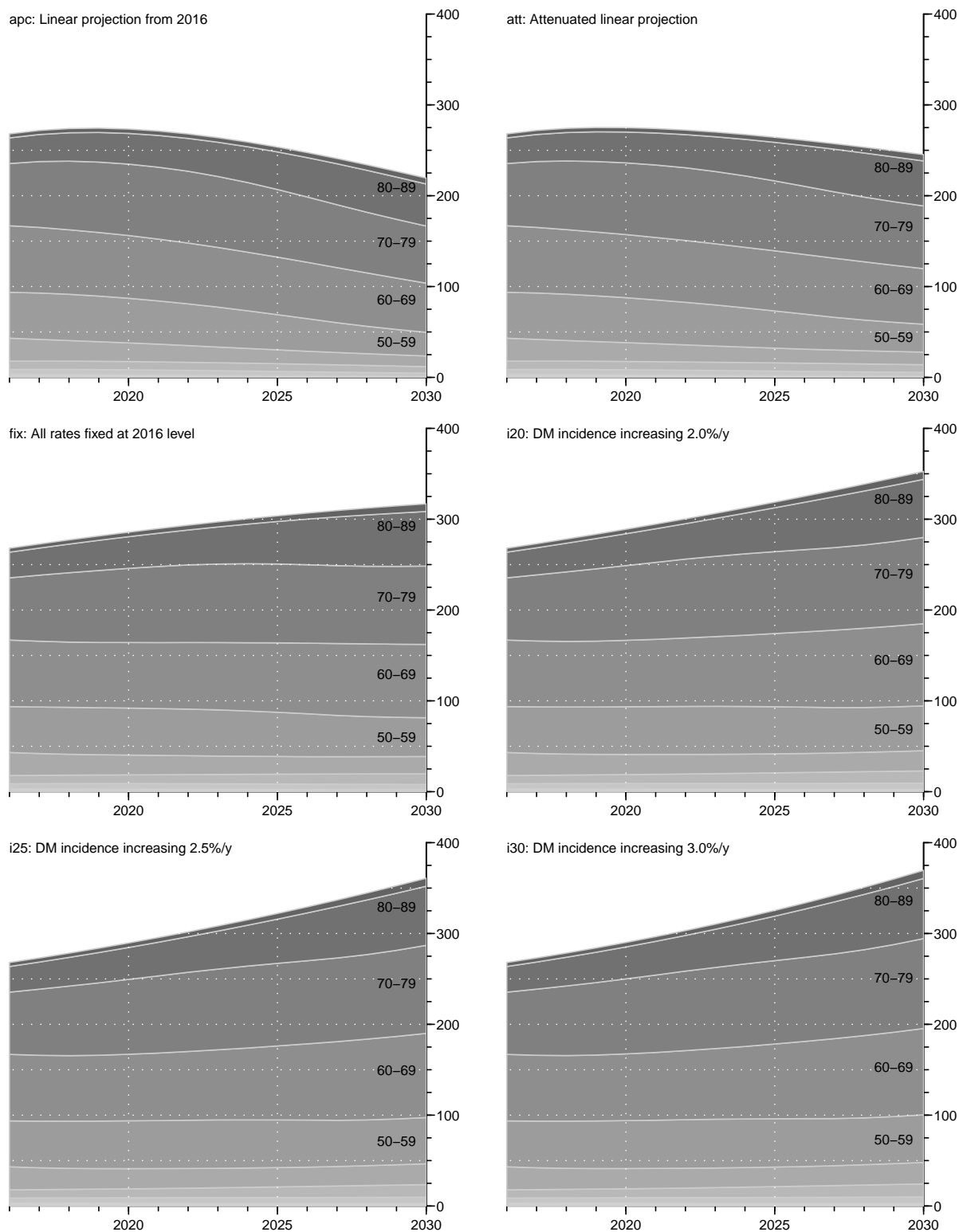


Figure 6.7: Predicted number of diabetes patients in Denmark under different scenarios.

References

- [1] B Carstensen. Age-Period-Cohort models for the Lexis diagram. *Statistics in Medicine*, 26(15):3018–3045, July 2007.
- [2] B. Carstensen, Christensen J.K., Marcussen M.M., and Borch-Johnsen K. The National Diabetes Register. *Scandinavian Journal of Public Health*, 39(7 suppl):58–61, 2011.
- [3] Anders Green, Camilla Sorts, Peter Bjdstrup Jensen, and Martha Emneus. Validation of the Danish National Diabetes Register. *Clinical Epidemiology*, 7:5–15, 2015.

Chapter 7

SAS programs

7.1 Rationale and overview

The following documented programs sequentially construct datasets with dates of diagnosis of DM according to different criteria, then merge these to pick the earliest.

All created data will be in the data folder as SAS-datasets, and

We have the following programs:

00-base Collects the base information on all persons in Denmark, that is sex, date of birth and date of death, and stores it in the dataset **bef**.

01-npr Uses the national patient register (NPR) to generate three datasets, all with **pnr** as key:

- a dataset **npr** with the earliest date of DM diagnosis in the NPR, **doNPR**, as well as a variable **nprtyp** with values T1 (ICD10: E10) or T2 (ICD10: E11) or NA (anything else), based on whether E10 or E11 or neither is recorded on more than half of the person's NPR entries. Thus formally some of the follow-up will be based on type-information from future recording.
- a dataset **pcos** with the earliest date of registered PCOS, **doPCOS**
- a dataset **gdm** with recorded dates of GDM that are at least 200 days apart, **doGDM1**, **doGDM2**, ...

02-dvdd Uses the DADD to identify persons from outpatient clinics (and in due course from GPs) and to seek out persons deemed to be T1D patients. It creates a dataset, **dvdd** with key **pnr** and a variable for type of diabetes **dvdtyp**, based on whether T1 resp. T2 is recorded on more than half of the available clinical records. Thus formally some of the follow-up will be based on type-information from future recording.

03-ndr Uses the NDR to define data of diagnosis ignoring the originally implemented blood-glucose criteria and foot-therapy, and creates a dataset **ndr** with a revised version of the "old" NDR.

04-rmps Generates a dataset with **pnr** as key with one record per person, with dates of first dispensation of each of a number of drugs, **donlMet**, **danyMet** etc. as well as **doOAD** and **doIns**, which are the two criteria dates that are carried on to the register. Note that this is at variance with the original NDR that used the *second* dispensation date.

06-define Collects data from the previously created datasets and defines date of diagnosis and type of diabetes (T1/T2), and thus generates a DM-register with sex, date of birth, date of death, date of inclusion, the latter being the smaller of date of OAD, date of insulin, date of DADD recording and date of NPR recording.

For persons with a record from DADD with type of diabetes defined, this is used. For other persons with a record from NPR the type is taken as that derived from the NPR if T1 or T2. For persons with no classification from DADD or from NPR, the person is classified as T1 if any type of medication has been taken out before age 15 or if insulin has been taken out before age 30, otherwise as T2.

06a-comp Compares the prevalent cases as of 1996, 2004 and 2012 from the olde revised NDR and from the reconstructed DMreg.

7.2 Program execution

All data analyses are run on the servers at Statistics Denmark. In order to have a thorough documentation of the data processing all SAS-programs have been run in sequence as batch jobs from the command prompt (`cmd`), where the program in the file `xxx.sas`, say, is run and produces the files `xxx.log` and `xxx.lst`. Since the code from `xxx.sas` is contained in `xxx.log`, it suffices to show the files `xxx.log` and `xxx.lst` to provide full documentation of the data acquisition process.

The practical execution of the SAS-programs is done using the `cmd`-script `sj.bat` which reads:

```
start "sas job" /min sjx %~n1
```

The running of the program `xxx.sas` is started by issuing “`sj xxx`” at the command prompt.

The script `sj.bat` just starts a new process which in turn runs the script `sjx.bat`, which reads:

```
"C:\Program Files\SASHome\SASFoundation\9.4\sas.exe" ^
-COFIG "C:\Program Files\SASHome\SASFoundation\9.4\nls\en\sasv9.cfg" ^
-$lognote1 "Program %~n1.sas" -autoexec optslibs.sas -sysin %~n1.sas
copy %~n1.log + ctrl1 + %~n1.lst %~n1.yt
exit
```

The second last line in the script simply copies the two result-files from SAS into one for convenience of inspection. It is the two result files that are transferred from DST to a local computer for inclusion in a documentation report.

This way there is a reasonable documentation that the results are actually produced by the listed code (in the `.log` file). Hopefully the program code is written in a reasonably human-readable way.

7.3 Program documentation

The following is a listing of the SAS-programs and -results (that is the `.log` and `.lst` files) used to generate the base datasets. Each is preceded by a very brief description; main technical points are included as comments in the program code, found in the `.log` files.

Note that according to rules of DST, all table entries of 5 or less in `.lst` file are masked as a “*”.

7.3.1 optslibs.sas

This is common set of declarative commands that defines a couple of options, the location of the raw and the derived datasets and some global macro variables. It is included as autoexec file in all runs, with `options nonotes` for brevity of output:

```
* options used throughout ;
options nocenter nonotes nomprint nosource2
      ps = 10000 /* 105 */
      ls = 90    /* 160 */
      obs = max
      formchar = '          '
/* format libraries we use */
fmtsearch = ( dsfmt.times_personstatistik
              dsfmt.brancher
              dsfmt.uddannelser
              dsfmt.geokoder
              daffmt.drugfmt
              ttfmt.ttformat00
              ttfmt.ttformat10
              ttfmt.ttformat20 ) ;

* data libraries ;
libname DELPOP15 'E:\rawdata\705093\Delpopulationer\' ;
libname EKST15  'E:\rawdata\705093\Eksterne data\' ;
libname EKST16  'E:\rawdata\705093\Eksterne data\Diabasen 28-11-2016' ;
libname GRUND15 'E:\rawdata\705093\Grunddata\' ;
libname POPUL15 'E:\rawdata\705093\Population\' ;
libname TTDATA  'E:\workdata\705093\BxC\demoDM\DATA\' ;

* format libraries ;
libname TTfmt 'E:\workdata\705093\QSN_MLiJ\' ;
libname daffmt 'E:\workdata\705093\BxC\daffodil\data\' ;
libname DSfmt 'E:\Formater\SAS formater i Danmarks Statistik\FORMATKATALOG';

* useful constants ;
%let primo    = '01JAN1995'd ;
%let cutdate  = '01JAN2016'd ; * we have data including 2015 ;
%let gdmint   = 200 ; * minimal distance between GDM dates to constitute 2 GDM events ;

* macro to exclude observations with dates in GDM grace period ;
%macro xgdm( xdate, gdmwin = 365 ) ;
%do n = 1 %to 12 ;
    if doGDM&n. < &xdate. < ( doGDM&n. + &gdmwin. ) then delete ;
%end ;
%mend ;

* page ;
options notes ;
```

7.3.2 xgdm.sas

Note that the `optslibs.sas` also contains the definition of the `xgdm` macro: For each of the criteria it is necessary to exclude dates of meeting the criterion which fall within a grace period after a diagnosis of GDM. This is what the macro `xgdm` is for; it relies on the structure of the GDM dataset constructed in the `01-npr` program, which has the GDM dates in the wide form for person with at least one date of GDM. It iterates up to 12 in order to produce a note from the SAS system, that documents that only 11 GDM dataes are needed.

7.4 00-base

Reads the files with all person ids (**pnr**), for each calendar year of data, and forms a total roster of all **pnr** with demographic information (sex, date of birth, date of death).

7.4.1 00-base.lst

7.5 01-npr

Processes the records from the NPR, and produces records with GDM diagnoses and PCOS diagnoses.

Persons cannot enter on any criterion in a 365 days grace period after each GDM diagnosis. GDM diagnoses occurring within 200 days of another one is not counted, though. Thus all GDM diagnoses in the same person are at least 200 days apart.

Outputs the earliest NPR diagnosis clear of GDM and PCOS, and derives a tentative T1/T2 classification in the variable **nprtyp**.

```

1                                     "Program 01-npr.sas"

NOTE: Copyright (c) 2002-2012 by SAS Institute Inc., Cary, NC, USA.
NOTE: SAS (r) Proprietary Software 9.4 (TS1M3)
      Licensed to FORSKNING 2, Site 50800723.
NOTE: This session is executing on the X64_SRV12 platform.

NOTE: Updated analytical products:
      SAS/STAT 14.1

NOTE: Additional host information:
      X64_SRV12 WIN 6.2.9200 Server

NOTE: SAS initialization used:
      real time          0.06 seconds
      cpu time           0.07 seconds

NOTE: AUTOEXEC processing beginning; file is E:\workdata\705093\BXC\demoDM\sas\optslibs.sas.

NOTE: AUTOEXEC processing completed.

1      * read the NPR datasets in the two different formats and combine them ;
2
3      %macro mold ;
4      data all_npr1977_93 ;
5          set %do i = 1977 %to 1993 ;
6              GRUND15.LPRHEL&i. (keep = pnr recnum c_adiag d_inddto )
7          %end ; ;
8      * the ICD-8 codes incl. GDM / PCOS ;
9      if c_adiag in('24900','24901','24902','24903','24904',
10                 '24905','24906','24907','24908','24909',
11                 '25000','25001','25002','25003','25004',
12                 '25005','25006','25007','25008','25009',
13                 '63474','Y6449','61520','61521') ;
14      if substr(c_adiag,1,3) eq '249' then nprtyp = 'T1' ;
15      if substr(c_adiag,1,3) eq '250' then nprtyp = 'T2' ;
16      run ;
17      %mend ;
18      %mold ;

```

```

NOTE: There were 536220 observations read from the data set GRUND15.LPRHEL1977.
NOTE: There were 576400 observations read from the data set GRUND15.LPRHEL1978.
NOTE: There were 593234 observations read from the data set GRUND15.LPRHEL1979.
NOTE: There were 603085 observations read from the data set GRUND15.LPRHEL1980.
NOTE: There were 602068 observations read from the data set GRUND15.LPRHEL1981.
NOTE: There were 625449 observations read from the data set GRUND15.LPRHEL1982.
NOTE: There were 651393 observations read from the data set GRUND15.LPRHEL1983.
NOTE: There were 669830 observations read from the data set GRUND15.LPRHEL1984.
NOTE: There were 694317 observations read from the data set GRUND15.LPRHEL1985.
NOTE: There were 727270 observations read from the data set GRUND15.LPRHEL1986.
NOTE: There were 754514 observations read from the data set GRUND15.LPRHEL1987.
NOTE: There were 792682 observations read from the data set GRUND15.LPRHEL1988.
NOTE: There were 821261 observations read from the data set GRUND15.LPRHEL1989.
NOTE: There were 852349 observations read from the data set GRUND15.LPRHEL1990.
NOTE: There were 874069 observations read from the data set GRUND15.LPRHEL1991.
NOTE: There were 922350 observations read from the data set GRUND15.LPRHEL1992.
NOTE: There were 989322 observations read from the data set GRUND15.LPRHEL1993.
NOTE: The data set WORK.ALL_NPR1977_93 has 239128 observations and 5 variables.
NOTE: DATA statement used (Total process time):
      real time          4.79 seconds
      cpu time           1.21 seconds

```

```

19
20      %macro mnew ;
21      data all_npr1994_15 ;
22          set %do i = 1994 %to 2015 ;
23              GRUND15.LPRPOP&i. (keep = pnr recnum c_adiag d_inddto )
24          %end ;
25          %do i = 14 %to 15 ;
26              GRUND15.LPRPOP_UAFAMB&i. (keep = pnr recnum c_adiag d_inddto )
27          %end ; ;
28      * the ICD-10 codes incl GDM / PCOS ;
29      if substr(c_adiag,2,3) in ('E10','E11','E12','E13','E14','O24') or
30      substr(c_adiag,2,4) in ('H360','E748','E891','R730','R819','E282' ) ;
31      if substr(c_adiag,2,3) eq 'E10' then nprtyp = 'T1' ;
32      if substr(c_adiag,2,3) eq 'E11' then nprtyp = 'T2' ;
33      run ;
34      %mend ;
35      %mnew ;

```

WARNING: Multiple lengths were specified for the variable C_ADIAG by input data set(s).
This can cause truncation of data.

```

NOTE: There were 2257703 observations read from the data set GRUND15.LPRPOP1994.
NOTE: There were 3099164 observations read from the data set GRUND15.LPRPOP1995.
NOTE: There were 3254161 observations read from the data set GRUND15.LPRPOP1996.
NOTE: There were 3348359 observations read from the data set GRUND15.LPRPOP1997.
NOTE: There were 3438023 observations read from the data set GRUND15.LPRPOP1998.
NOTE: There were 3544774 observations read from the data set GRUND15.LPRPOP1999.
NOTE: There were 3554711 observations read from the data set GRUND15.LPRPOP2000.
NOTE: There were 3676892 observations read from the data set GRUND15.LPRPOP2001.
NOTE: There were 3734587 observations read from the data set GRUND15.LPRPOP2002.
NOTE: There were 3714168 observations read from the data set GRUND15.LPRPOP2003.
NOTE: There were 4771057 observations read from the data set GRUND15.LPRPOP2004.
NOTE: There were 4970581 observations read from the data set GRUND15.LPRPOP2005.
NOTE: There were 5146887 observations read from the data set GRUND15.LPRPOP2006.
NOTE: There were 5169427 observations read from the data set GRUND15.LPRPOP2007.
NOTE: There were 5544338 observations read from the data set GRUND15.LPRPOP2008.
NOTE: There were 5835150 observations read from the data set GRUND15.LPRPOP2009.
NOTE: There were 7287169 observations read from the data set GRUND15.LPRPOP2010.
NOTE: There were 7664856 observations read from the data set GRUND15.LPRPOP2011.
NOTE: There were 6069159 observations read from the data set GRUND15.LPRPOP2012.
NOTE: There were 6227306 observations read from the data set GRUND15.LPRPOP2013.
NOTE: There were 6386162 observations read from the data set GRUND15.LPRPOP2014.
NOTE: There were 6798419 observations read from the data set GRUND15.LPRPOP2015.
NOTE: There were 317697 observations read from the data set GRUND15.LPRPOP_UAFAMB14.
NOTE: There were 1860412 observations read from the data set GRUND15.LPRPOP_UAFAMB15.
NOTE: The data set WORK.ALL_NPR1994_15 has 1333078 observations and 5 variables.
NOTE: DATA statement used (Total process time):
      real time          38.53 seconds

```

```
cpu time          12.46 seconds
```

```
36
37     data all_npr ;
38     set all_npr1977_93
39     all_npr1994_15 ;
40     run ;
```

NOTE: There were 239128 observations read from the data set WORK.ALL_NPR1977_93.
 NOTE: There were 1333078 observations read from the data set WORK.ALL_NPR1994_15.
 NOTE: The data set WORK.ALL_NPR has 1572206 observations and 5 variables.
 NOTE: DATA statement used (Total process time):
 real time 0.17 seconds
 cpu time 0.17 seconds

```
41
42     proc sort data = all_npr ; by pnr d_inddto ; run ;
```

NOTE: There were 1572206 observations read from the data set WORK.ALL_NPR.
 NOTE: The data set WORK.ALL_NPR has 1572206 observations and 5 variables.
 NOTE: PROCEDURE SORT used (Total process time):
 real time 0.29 seconds
 cpu time 0.67 seconds

```
43
44     * only records from persons in the base population -
45     GDM & PCOS diagnoses are put in separate file ;
46     data DM
47     gdm
48     pcos ;
49     merge all_npr      ( in = npr )
50     TTDATA.pop ( in = pop ) ;
51     by pnr ;
52     length diaggr $ 4 ;
53     if npr and pop ;
54     * GDM / PCOS (excluding men) or 'real' DM diagnoses ;
55     if substr(c_adiag,2,3) in('024'          ) or
56     c_adiag          in('63474','Y6449')
57     then do ;
58     if sex eq 2 then diaggr = 'GDM' ; else delete ;
59     end ;
60     else
61     if substr(c_adiag,2,4) in('E282'          ) or
62     c_adiag          in('61520','61521')
63     then do ;
64     if sex eq 2 then diaggr = 'PCOS' ; else delete ;
65     end ;
66     else
67     diaggr = 'DM' ;
68     if diaggr eq 'DM' then output DM ;
69     if diaggr eq 'GDM' then output gdm ;
70     if diaggr eq 'PCOS' then output pcos ;
71     run ;
```

NOTE: There were 1572206 observations read from the data set WORK.ALL_NPR.
 NOTE: There were 7361669 observations read from the data set TTDATA.POP.
 NOTE: The data set WORK.DM has 1371306 observations and 9 variables.
 NOTE: The data set WORK.GDM has 53671 observations and 9 variables.
 NOTE: The data set WORK.PCOS has 32167 observations and 9 variables.
 NOTE: DATA statement used (Total process time):
 real time 1.78 seconds
 cpu time 1.56 seconds

```
72
73     title1 'PCOS records - id and first date of PCOS' ;
74     proc sort data = pcos ; by pnr d_inddto ; run ;
```


NOTE: There were 32167 observations read from the data set WORK.PCOS.

NOTE: The data set WORK.PCOS has 32167 observations and 9 variables.

NOTE: PROCEDURE SORT used (Total process time):

real time	0.01 seconds
cpu time	0.01 seconds

```
75      data TTdata.pcos ( keep = pnr doPCOS ) ;
76          set pcos ;
77          by pnr d_inddto ;
78          if first.pnr ;
79          doPCOS = d_inddto ;
80      run ;
```

NOTE: There were 32167 observations read from the data set WORK.PCOS.

NOTE: The data set TTDATA.PCOS has 20773 observations and 2 variables.

NOTE: DATA statement used (Total process time):

real time	0.03 seconds
cpu time	0.00 seconds

```
81
82      proc contents data = TTdata.pcos ; run ;
```

NOTE: PROCEDURE CONTENTS used (Total process time):

real time	0.01 seconds
cpu time	0.01 seconds

NOTE: The PROCEDURE CONTENTS printed page 1.

```
83      proc tabulate data = TTdata.pcos missing noseps ;
84          class doPCOS ;
85          table doPCOS, n * f=comma10.
86              / rts = 8 ;
87          format doPCOS year4. ;
88      run ;
```

NOTE: There were 20773 observations read from the data set TTDATA.PCOS.

NOTE: The PROCEDURE TABULATE printed page 2.

NOTE: PROCEDURE TABULATE used (Total process time):

real time	0.01 seconds
cpu time	0.01 seconds

```
89      proc tabulate data = TTdata.pcos missing noseps ;
90          where doPCOS ge '01JAN2013'd ;
91          class doPCOS ;
92          table doPCOS, n * f=comma10.
93              / rts = 10 ;
94          format doPCOS yymms8. ;
95      run ;
```

NOTE: There were 2354 observations read from the data set TTDATA.PCOS.
WHERE doPCOS>='01JAN2013'D;

NOTE: The PROCEDURE TABULATE printed page 3.

NOTE: PROCEDURE TABULATE used (Total process time):

real time	0.01 seconds
cpu time	0.01 seconds

```
96
97      title1 'GDM records - id and any date of GDM except if too close' ;
98      proc sort data = gdm ; by pnr d_inddto ; run ;
```

NOTE: There were 53671 observations read from the data set WORK.GDM.

NOTE: The data set WORK.GDM has 53671 observations and 9 variables.

NOTE: PROCEDURE SORT used (Total process time):

real time	0.01 seconds
cpu time	0.01 seconds

```

99
100     data gdm ( keep = pnr doGDM dno ) ;
101         set gdm ( rename = ( d_inddto = doGDM ) ) ;
102     by pnr doGDM ;
103     retain prevGDM ;
104     if first.pnr then do ;
105         dno = 1 ;
106         prevGDM = doGDM ;
107         output ;
108         end ;
109     if ^first.pnr and ( doGDM - prevGDM ) gt &gdmint. then do ;
110         dno + 1 ;
111         output ;
112         prevGDM = doGDM ;
113         end ;
114     run ;

```

NOTE: There were 53671 observations read from the data set WORK.GDM.

NOTE: The data set WORK.GDM has 27882 observations and 3 variables.

NOTE: DATA statement used (Total process time):

```

real time      0.01 seconds
cpu time       0.01 seconds

```

```

115
116     proc transpose  data = gdm
117                   out = TTdata.gdm ( drop = _NAME_ _LABEL_ )
118                   prefix = doGDM ;
119     by pnr ;
120     var doGDM ;
121     id dno ;
122     run ;

```

NOTE: There were 27882 observations read from the data set WORK.GDM.

NOTE: The data set TTDATA.GDM has 21146 observations and 12 variables.

NOTE: PROCEDURE TRANSPOSE used (Total process time):

```

real time      0.06 seconds
cpu time       0.06 seconds

```

```

123
124     %let doGDMn = doGDM2 doGDM3 doGDM4 doGDM5 doGDM6 doGDM7 doGDM8 doGDM9 doGDM10
124     ! doGDM11 ;
125     title 'The recorded dates of Gestational diabetes' ;
126     proc contents  data = TTdata.gdm ; run ;

```

NOTE: PROCEDURE CONTENTS used (Total process time):

```

real time      0.00 seconds
cpu time       0.00 seconds

```

NOTE: The PROCEDURE CONTENTS printed page 4.

```

127     proc tabulate  data = TTdata.gdm missing noseps ;
128         class doGDM1 ;
129         var &doGDMn. ;
130         table doGDM1 all &doGDMn.,
131             n * f=comma10. / rts=9 ;
132         format doGDM1 year4. ;
133     run ;

```

NOTE: There were 21146 observations read from the data set TTDATA.GDM.

NOTE: The PROCEDURE TABULATE printed page 5.

NOTE: PROCEDURE TABULATE used (Total process time):

```

real time      0.01 seconds
cpu time       0.01 seconds

```

```

134     proc tabulate  data = TTdata.gdm missing noseps ;

```

```

135     where doGDM1 ge '01JAN2013'd ;
136     class doGDM1 ;
137     var &doGDMn. ;
138     table doGDM1 all &doGDMn.,
139           n * f=comma10. / rts=9 ;
140     format doGDM1 yymms7. ;
141     run ;

```

NOTE: There were 4312 observations read from the data set TTDATA.GDM.
WHERE doGDM1>='01JAN2013'D;

NOTE: The PROCEDURE TABULATE printed page 6.

NOTE: PROCEDURE TABULATE used (Total process time):

```

real time      0.01 seconds
cpu time       0.01 seconds

```

```

142     title1 ;
143
144     options mprint ;
145     data npr ( keep = pnr sex d_inddto nprtyp c_adiag ) ;
146         merge DM ( in = DM )
147             TTdata.gdm ;
148     by pnr ;
149     if DM ;
150     %xgdm( d_inddto ) ;
MPRINT(XGDM):  if ( doGDM1 - 30 ) < d_inddto < ( doGDM1 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM2 - 30 ) < d_inddto < ( doGDM2 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM3 - 30 ) < d_inddto < ( doGDM3 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM4 - 30 ) < d_inddto < ( doGDM4 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM5 - 30 ) < d_inddto < ( doGDM5 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM6 - 30 ) < d_inddto < ( doGDM6 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM7 - 30 ) < d_inddto < ( doGDM7 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM8 - 30 ) < d_inddto < ( doGDM8 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM9 - 30 ) < d_inddto < ( doGDM9 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM10 - 30 ) < d_inddto < ( doGDM10 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM11 - 30 ) < d_inddto < ( doGDM11 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM12 - 30 ) < d_inddto < ( doGDM12 + 365 ) then delete ;
151     run ;

```

NOTE: Variable doGDM12 is uninitialized.

NOTE: Missing values were generated as a result of performing an operation on missing values.

Each place is given by: (Number of times) at (Line):(Column).

```

1300548 at 150:18  1300548 at 150:54  1332935 at 150:20  1332935 at 150:56
1352536 at 150:20  1352536 at 150:56  1359343 at 150:20  1359343 at 150:56
1360953 at 150:20  1360953 at 150:56  1361516 at 150:20  1361516 at 150:56
1361597 at 150:20  1361597 at 150:56  1361626 at 150:20  1361626 at 150:56
1361626 at 150:20  1361626 at 150:56  1361626 at 150:20  1361626 at 150:56
1361626 at 150:20  1361626 at 150:56  1361627 at 150:20  1361627 at 150:56

```

NOTE: There were 1371306 observations read from the data set WORK.DM.

NOTE: There were 21146 observations read from the data set TTDATA.GDM.

NOTE: The data set WORK.NPR has 1361627 observations and 5 variables.

NOTE: DATA statement used (Total process time):

```

real time      3.62 seconds
cpu time       3.62 seconds

```

```

152     options nomprint ;
153
154     title1 'NPR diagnoses used - records' ;
155     proc tabulate data = npr missing noseps ;
156         class c_adiag nprtyp d_inddto ;
157         table all c_adiag d_inddto,
158               nprtyp * f=comma10.
159               / rts = 10 ;
160     format d_inddto year4. ;
161     title1 ;
162

```

NOTE: There were 1361627 observations read from the data set WORK.NPR.

NOTE: The PROCEDURE TABULATE printed page 7.
 NOTE: PROCEDURE TABULATE used (Total process time):
 real time 0.29 seconds
 cpu time 0.71 seconds

```
163     data npr1 ;
164     set npr ( keep = pnr d_inddto
165             rename = ( d_inddto = doNPR ) ) ;
166     by pnr ;
167     if first.pnr ;
168     run ;
```

NOTE: There were 1361627 observations read from the data set WORK.NPR.
 NOTE: The data set WORK.NPR1 has 223402 observations and 2 variables.
 NOTE: DATA statement used (Total process time):
 real time 0.12 seconds
 cpu time 0.12 seconds

```
169
170     * Classifiy persons according to the most Frequently occurring type ;
171     data TTdata.npr ;
172     merge npr npr1 ;
173     by pnr ;
174     retain nT1 nT2 ;
175     if first.pnr then do ;
176         nT1 = 0 ;
177         nT2 = 0 ;
178         nRc = 0 ;
179     end ;
180     nT1 + ( nprtyp eq 'T1' ) ;
181     nT2 + ( nprtyp eq 'T2' ) ;
182     nRc + 1 ;
183     * If more than half of records agree on one type ;
184     if last.pnr then do ;
185         nprtyp = 'NA' ;
186         if nT1 > nRc/2 then nprtyp = 'T1' ;
187         if nT2 > nRc/2 then nprtyp = 'T2' ;
188     output ;
189     end ;
190     label doNPR = ' ' ;
191     run ;
```

NOTE: There were 1361627 observations read from the data set WORK.NPR.
 NOTE: There were 223402 observations read from the data set WORK.NPR1.
 NOTE: The data set TTDATA.NPR has 223402 observations and 9 variables.
 NOTE: DATA statement used (Total process time):
 real time 0.26 seconds
 cpu time 0.23 seconds

```
192
193     title1 'Diagnoses of DM accepted from NPR' ;
194     proc contents data = TTdata.npr ; run ;
```

NOTE: PROCEDURE CONTENTS used (Total process time):
 real time 0.00 seconds
 cpu time 0.00 seconds

NOTE: The PROCEDURE CONTENTS printed page 8.

```
195
196     proc tabulate data = TTdata.npr missing noseps ;
197     class doNPR nprtyp sex ;
198     table all doNPR,
199           ( all sex nprtyp ) * f=comma10.
200           / rts = 7 ;
201     format doNPR year4.
202           sex koen_t. ;
```

```
203          run ;
```

NOTE: There were 223402 observations read from the data set TTDATA.NPR.

NOTE: The PROCEDURE TABULATE printed page 9.

NOTE: PROCEDURE TABULATE used (Total process time):

real time	0.12 seconds
cpu time	0.12 seconds

```
204
205          title2 '- only from 1 January 2013 - checking seasonality' ;
206          proc tabulate data = TTdata.npr missing noseps ;
207              where doNPR ge '01JAN2013'd ;
208              class doNPR nprtyp sex ;
209              table all doNPR,
210                  ( all sex nprtyp ) * f=comma10.
211                  / rts = 10 ;
212              format doNPR yymms8.
213                  sex koen_t. ;
214          run ;
```

NOTE: There were 24630 observations read from the data set TTDATA.NPR.

WHERE doNPR>='01JAN2013'D;

NOTE: The PROCEDURE TABULATE printed page 10.

NOTE: PROCEDURE TABULATE used (Total process time):

real time	0.03 seconds
cpu time	0.03 seconds

```
215          title1 ;
```

NOTE: SAS Institute Inc., SAS Campus Drive, Cary, NC USA 27513-2414

NOTE: The SAS System used:

real time	50.45 seconds
cpu time	21.26 seconds

7.5.1 01-npr.lst

PCOS records - id and first date of PCOS

17:15 Thursday, January 12, 2017 1

The CONTENTS Procedure

Data Set Name	TTDATA.PCOS	Observations	20773
Member Type	DATA	Variables	*
Engine	V9	Indexes	0
Created	12/01/2017 17:15:49	Observation Length	24
Last Modified	12/01/2017 17:15:49	Deleted Observations	0
Protection		Compressed	NO
Data Set Type		Sorted	NO
Label			
Data Representation	WINDOWS_64		
Encoding	wlatin1 Western (Windows)		

Engine/Host Dependent Information

Data Set Page Size	65536
Number of Data Set Pages	8
First Data Page	*
Max Obs per Page	2715
Obs in First Data Page	2664
Number of Data Set Repairs	0
ExtendObsCounter	YES
Filename	E:\workdata\705093\BxC\demoDM\DATA\pcos.sas7bd
Release Created	9.0401M3
Host Created	X64_SRV12

Alphabetic List of Variables and Attributes

#	Variable	Type	Len	Format	Informat	Label
2	doPCOS	Num	8			
1	pnr	Char	12	\$12.	\$10.	Personnummer

N

doPCOS

1976	*
1977	577
1978	607
1979	590
1980	648
1981	741
1982	682
1983	757
1984	717
1985	757
1986	742
1987	610
1988	637
1989	630
1990	589
1991	574
1992	535
1993	459
1994	109
1995	108
1996	153
1997	153
1998	188
1999	229
2000	193
2001	212
2002	272
2003	378
2004	479
2005	532
2006	538
2007	631
2008	566
2009	695
2010	732
2011	705
2012	691
2013	789
2014	832
2015	733

N

doPCOS

2013/01	73
2013/02	54
2013/03	58
2013/04	66
2013/05	68
2013/06	65
2013/07	45
2013/08	67
2013/09	74
2013/10	85
2013/11	67

2013/12	67
2014/01	82
2014/02	68
2014/03	60
2014/04	61
2014/05	71
2014/06	69
2014/07	49
2014/08	68
2014/09	82
2014/10	78
2014/11	74
2014/12	70
2015/01	71
2015/02	49
2015/03	79
2015/04	64
2015/05	72
2015/06	79
2015/07	38
2015/08	58
2015/09	67
2015/10	61
2015/11	54
2015/12	41

The CONTENTS Procedure

Data Set Name	TTDATA.GDM	Observations	21146
Member Type	DATA	Variables	12
Engine	V9	Indexes	0
Created	12/01/2017 17:15:49	Observation Length	104
Last Modified	12/01/2017 17:15:49	Deleted Observations	0
Protection		Compressed	NO
Data Set Type		Sorted	NO
Label			
Data Representation	WINDOWS_64		
Encoding	wlatin1 Western (Windows)		

Engine/Host Dependent Information

Data Set Page Size	65536
Number of Data Set Pages	34
First Data Page	*
Max Obs per Page	629
Obs in First Data Page	607
Number of Data Set Repairs	0
ExtendObsCounter	YES
Filename	E:\workdata\705093\BxC\demoDM\DATA\gdm.sas7bdat
Release Created	9.0401M3
Host Created	X64_SRV12

Alphabetic List of Variables and Attributes

#	Variable	Type	Len	Format	Informat	Label
*	doGDM1	Num	8	DDMMYY10.		
*	doGDM2	Num	8	DDMMYY10.		
*	doGDM3	Num	8	DDMMYY10.		
*	doGDM4	Num	8	DDMMYY10.		
6	doGDM5	Num	8	DDMMYY10.		
7	doGDM6	Num	8	DDMMYY10.		
8	doGDM7	Num	8	DDMMYY10.		
9	doGDM8	Num	8	DDMMYY10.		
10	doGDM9	Num	8	DDMMYY10.		
11	doGDM10	Num	8	DDMMYY10.		

12	doGDM11	Num	8	DDMMYY10.		
*	pnr	Char	12	\$12.	\$10.	Personnummer

N

doGDM1	
1980	*
1987	51
1988	82
1989	108
1990	96
1991	84
1992	89
1993	160
1994	474
1995	435
1996	604
1997	591
1998	681
1999	546
2000	583
2001	585
2002	590
2003	774
2004	949
2005	1,020
2006	1,041
2007	1,112
2008	1,250
2009	1,354
2010	1,231
2011	1,245
2012	1,098
2013	1,347
2014	1,478
2015	1,487
All	21,146
doGDM2	5,108
doGDM3	1,218
doGDM4	292
doGDM5	82
doGDM6	23
doGDM7	8
doGDM8	*
doGDM9	*
doGDM10	*
doGDM11	*

N

doGDM1	
2013/01	110
2013/02	108
2013/03	100
2013/04	128
2013/05	104
2013/06	126
2013/07	101
2013/08	123
2013/09	107
2013/10	104
2013/11	113
2013/12	123
2014/01	143
2014/02	110

2014/03	133
2014/04	133
2014/05	150
2014/06	128
2014/07	123
2014/08	121
2014/09	116
2014/10	101
2014/11	109
2014/12	111
2015/01	164
2015/02	131
2015/03	139
2015/04	103
2015/05	122
2015/06	148
2015/07	136
2015/08	140
2015/09	109
2015/10	107
2015/11	118
2015/12	70
All	4,312
doGDM2	240
doGDM3	12
doGDM4	0
doGDM5	0
doGDM6	0
doGDM7	0
doGDM8	0
doGDM9	0
doGDM10	0
doGDM11	0

nprtyp			
T1 T2			
	N	N	N
All	217,741	482,966	660,920
Aktions-			
diagnose			
24900	.	5,914	.
24901	.	2,260	.
24902	.	366	.
24903	.	224	.
24904	.	96	.
24905	.	715	.
24906	.	70	.
24907	.	1,325	.
24908	.	2,512	.
24909	.	21,160	.
25000	.	.	19,663
25001	.	.	4,017
25002	.	.	320
25003	.	.	642
25004	.	.	86
25005	.	.	919
25006	.	.	113
25007	.	.	1,879
25008	.	.	4,768
25009	.	.	46,640
DE10	.	1,314	.
DE100	.	3,842	.
DE100A	.	25	.
DE100B	.	402	.

DE100C	.	34	.
DE100D	.	87	.
DE100E	.	504	.
DE100F	.	236	.
DE101	.	20,757	.
DE102	.	16,513	.
DE103	.	29,194	.
DE104	.	9,241	.
DE105	.	14,990	.
DE105A	.	258	.
DE105B	.	3,631	.
DE105C	.	1,186	.
DE105D	.	27	.
DE106	.	3,622	.
DE107	.	40,553	.
DE108	.	42,135	.
DE109	.	252,402	.
DE109A	.	7,371	.
DE11	.	.	836
DE110	.	.	3,014
DE110A	.	.	21
DE110B	.	.	167
DE110C	.	.	54
DE110D	.	.	46
DE110E	.	.	45
DE111	.	.	1,668
DE112	.	.	34,802
DE113	.	.	15,172
DE114	.	.	20,538
DE115	.	.	21,169
DE115A	.	.	446
DE115B	.	.	10,122
DE115C	.	.	1,387
DE115D	.	.	61
DE116	.	.	5,600
DE117	.	.	44,029
DE118	.	.	68,759
DE119	.	.	341,690
DE119A	.	.	12,247
DE12	8	.	.
DE120	225	.	.
DE120A	10	.	.
DE120B	10	.	.
DE120C	*	.	.
DE121	156	.	.
DE122	111	.	.
DE123	87	.	.
DE124	80	.	.
DE125	404	.	.
DE125A	*	.	.
DE125B	40	.	.
DE125C	42	.	.
DE126	39	.	.
DE127	91	.	.
DE128	134	.	.
DE129	298	.	.
DE13	45	.	.
DE130	77	.	.
DE131	510	.	.
DE132	328	.	.
DE133	1,667	.	.
DE134	309	.	.
DE135	213	.	.
DE135A	*	.	.
DE135B	141	.	.
DE135C	35	.	.
DE135D	*	.	.
DE136	191	.	.
DE137	521	.	.
DE138	869	.	.

DE139	5,543	.	.
DE14	378	.	.
DE140	726	.	.
DE140A	53	.	.
DE140B	12	.	.
DE140C	20	.	.
DE140D	20	.	.
DE141	2,262	.	.
DE142	898	.	.
DE143	3,458	.	.
DE144	3,909	.	.
DE145	5,941	.	.
DE145A	30	.	.
DE145B	2,437	.	.
DE145C	449	.	.
DE145D	17	.	.
DE146	476	.	.
DE147	1,375	.	.
DE148	5,738	.	.
DE149	39,751	.	.
DE748	213	.	.
DE748A	6	.	.
DE748B	41	.	.
DE748C	9	.	.
DE748E	9	.	.
DE748G	*	.	.
DE891	310	.	.
DE891A	80	.	.
DE891B	*	.	.
DH360	113,008	.	.
DH360A	1,506	.	.
DH360B	2,877	.	.
DH360C	1,724	.	.
DH360D	1,471	.	.
DH360E	1,271	.	.
DH360F	2,543	.	.
DH360H	3,230	.	.
DH360J	3,150	.	.
DH360K	4,501	.	.
DR730	1,048	.	.
DR730A	342	.	.
DR819	253	.	.
Indl- elsesda- to			
1941	.	*	.
1968	.	*	.
1970	.	9	.
1971	.	9	.
1972	.	14	*
1973	.	46	*
1974	.	37	*
1975	.	28	*
1976	.	28	48
1977	.	10	3,228
1978	.	63	4,094
1979	.	83	4,398
1980	*	104	4,700
1981	.	118	5,039
1982	.	90	5,598
1983	*	147	5,566
1984	*	107	6,381
1985	*	145	7,411
1986	6	281	7,777
1987	*	4,347	3,592
1988	57	5,194	3,509
1989	69	5,717	3,498
1990	59	5,774	3,459
1991	63	6,570	3,941
1992	117	6,952	4,265

1993	651	13,835	7,484
1994	2,475	13,872	9,790
1995	3,624	14,623	11,248
1996	4,006	14,301	13,100
1997	5,041	15,576	14,039
1998	7,681	15,301	16,039
1999	7,572	17,561	18,330
2000	6,537	15,504	18,510
2001	7,167	16,213	19,139
2002	7,495	16,556	19,620
2003	9,580	19,142	23,752
2004	10,608	19,287	22,988
2005	11,653	26,729	28,494
2006	11,939	21,130	30,991
2007	12,545	17,671	26,508
2008	19,695	43,773	59,929
2009	18,985	34,241	53,635
2010	13,674	19,625	33,534
2011	13,707	23,212	33,566
2012	11,736	15,259	24,028
2013	12,108	23,470	42,929
2014	10,824	13,700	27,644
2015	8,052	16,507	29,110

The CONTENTS Procedure

Data Set Name	TTDATA.NPR	Observations	223402
Member Type	DATA	Variables	9
Engine	V9	Indexes	0
Created	12/01/2017 17:15:53	Observation Length	72
Last Modified	12/01/2017 17:15:53	Deleted Observations	0
Protection		Compressed	NO
Data Set Type		Sorted	NO
Label			
Data Representation	WINDOWS_64		
Encoding	wlatin1 Western (Windows)		

Engine/Host Dependent Information

Data Set Page Size	65536
Number of Data Set Pages	247
First Data Page	*
Max Obs per Page	908
Obs in First Data Page	881
Number of Data Set Repairs	0
ExtendObsCounter	YES
Filename	E:\workdata\705093\BxC\demoDM\DATA\npr.sas7bdat
Release Created	9.0401M3
Host Created	X64_SRV12

Alphabetic List of Variables and Attributes

#	Variable	Type	Len	Format	Informat	Label
2	C_ADIAG	Char	6	\$20.	\$20.	Aktionsdiagnose
3	D_INDDTO	Num	8	DDMMYY10.	DATE9.	Indlelsesdato
6	doNPR	Num	8	DDMMYY10.	DATE9.	
9	nRc	Num	8			
7	nT1	Num	8			
8	nT2	Num	8			
4	nprtyp	Char	*			
1	pnr	Char	12	\$12.	\$10.	Personnummer
5	sex	Num	8			

	sex				nprtyp	
	All	Mand	Kvinde	NA	T1	T2
	N	N	N	N	N	N
All	223,402	126,834	96,568	37,839	44,535	141,028
doNPR						
1941	*	.	*	.	*	.
1968	*	*	.	.	*	.
1970	*	*	*	.	*	.
1971	*	.	*	*	*	.
1972	9	*	7	*	7	*
1973	19	12	7	*	15	*
1974	18	9	9	*	12	*
1975	16	10	6	8	7	*
1976	69	33	36	14	41	14
1977	2,414	1,170	1,244	465	1,241	708
1978	2,289	1,145	1,144	448	1,148	693
1979	2,066	1,045	1,021	362	1,017	687
1980	1,987	1,041	946	381	896	710
1981	1,835	972	863	320	805	710
1982	2,014	1,071	943	311	839	864
1983	1,881	950	931	321	745	815
1984	2,084	992	1,092	327	847	910
1985	2,166	1,145	1,021	342	827	997
1986	2,265	1,218	1,047	336	943	986
1987	2,314	1,206	1,108	312	975	1,027
1988	2,430	1,340	1,090	319	1,040	1,071
1989	2,605	1,362	1,243	352	1,048	1,205
1990	2,760	1,508	1,252	351	1,049	1,360
1991	3,062	1,644	1,418	393	1,139	1,530
1992	3,240	1,723	1,517	423	1,113	1,704
1993	5,127	2,927	2,200	774	1,706	2,647
1994	5,996	3,288	2,708	1,027	1,520	3,449
1995	6,457	3,627	2,830	1,120	1,466	3,871
1996	6,791	3,832	2,959	1,137	1,403	4,251
1997	7,038	3,925	3,113	1,119	1,403	4,516
1998	7,899	4,406	3,493	1,316	1,466	5,117
1999	8,021	4,439	3,582	1,398	1,340	5,283
2000	8,109	4,525	3,584	1,278	1,313	5,518
2001	8,094	4,580	3,514	1,440	1,290	5,364
2002	8,307	4,706	3,601	1,607	1,271	5,429
2003	9,356	5,378	3,978	1,732	1,245	6,379
2004	9,276	5,333	3,943	1,723	1,296	6,257
2005	9,041	5,108	3,933	1,604	1,236	6,201
2006	9,341	5,413	3,928	1,797	1,259	6,285
2007	9,210	5,317	3,893	1,667	1,210	6,333
2008	8,742	5,167	3,575	1,486	1,196	6,060
2009	8,624	5,158	3,466	1,469	1,178	5,977
2010	8,635	5,202	3,433	1,314	1,181	6,140
2011	8,860	5,286	3,574	1,399	1,129	6,332
2012	8,297	4,871	3,426	1,462	916	5,919
2013	8,314	4,958	3,356	1,429	935	5,950
2014	8,114	4,873	3,241	1,396	902	5,816
2015	8,202	4,914	3,288	1,352	913	5,937

- only from * January 2013 - checking seasonality

	sex				nprtyp	
	All	Mand	Kvinde	NA	T1	T2
	N	N	N	N	N	N
All	24,630	14,745	9,885	4,177	2,750	17,703
doNPR						
2013/01	823	501	322	139	107	577

2013/02	678	409	269	116	89	473
2013/03	730	452	278	129	91	510
2013/04	774	478	296	145	87	542
2013/05	775	451	324	137	74	564
2013/06	658	395	263	110	74	474
2013/07	535	296	239	72	57	406
2013/08	641	372	269	107	73	461
2013/09	671	389	282	111	58	502
2013/10	730	430	300	130	71	529
2013/11	712	416	296	124	85	503
2013/12	587	369	218	109	69	409
2014/01	825	507	318	155	91	579
2014/02	629	403	226	117	79	433
2014/03	790	470	320	148	89	553
2014/04	622	390	232	105	74	443
2014/05	671	375	296	112	92	467
2014/06	618	348	270	108	50	460
2014/07	532	326	206	101	58	373
2014/08	567	356	211	84	72	411
2014/09	734	427	307	137	79	518
2014/10	760	471	289	136	70	554
2014/11	757	437	320	101	87	569
2014/12	609	363	246	92	61	456
2015/01	743	481	262	93	84	566
2015/02	699	438	261	110	82	507
2015/03	774	463	311	133	77	564
2015/04	628	388	240	106	65	457
2015/05	708	412	296	121	81	506
2015/06	729	424	305	116	74	539
2015/07	525	303	222	93	63	369
2015/08	610	339	271	108	61	441
2015/09	792	483	309	131	82	579
2015/10	719	434	285	136	87	496
2015/11	727	424	303	132	75	520
2015/12	548	325	223	73	82	393

7.6 02-dvdd

The DVDD contains annual records for diabetes patients, mostly from out-patient clinics, but (eventually) also from GPs. These records contain type and date of diagnosis. The program chooses the earliest reported date of diagnosis and the type of diabetes reported more than half of the times (`dvdtyp`).

Uses the GDM dates to exclude possible inclusion dates in GDM grace periods.

```
1                                     "Program 02-dvdd.sas"
```

```
NOTE: Copyright (c) 2002-2012 by SAS Institute Inc., Cary, NC, USA.
```

```
NOTE: SAS (r) Proprietary Software 9.4 (TS1M3)
```

```
      Licensed to FORSKNING 1, Site 50800722.
```

```
NOTE: This session is executing on the X64_SRV12 platform.
```

```
NOTE: Updated analytical products:
```

```
      SAS/STAT 14.1
```

```
NOTE: Additional host information:
```

```
      X64_SRV12 WIN 6.2.9200 Server
```

```
NOTE: SAS initialization used:
```

```
      real time          0.06 seconds
```

```
      cpu time           0.09 seconds
```

NOTE: AUTOEXEC processing beginning; file is E:\workdata\705093\BXC\demoDM\sas\optslibs.sas.

NOTE: AUTOEXEC processing completed.

```
1      proc sort data = EKST15.DVDD out = dvdd ;
2          by pnr status_dato diag_dato diag_type ;
3      run ;
```

NOTE: There were 349662 observations read from the data set EKST15.DVDD.

NOTE: The data set WORK.DVDD has 349662 observations and 53 variables.

NOTE: PROCEDURE SORT used (Total process time):

```
real time      10.89 seconds
cpu time       0.70 seconds
```

```
4
5      * check number of *persons* in the data set ;
6      proc sort data = dvdd out = pers nodupkey ;
7          by pnr ;
8      run ;
```

NOTE: There were 349662 observations read from the data set WORK.DVDD.

NOTE: 260882 observations with duplicate key values were deleted.

NOTE: The data set WORK.PERS has 88780 observations and 53 variables.

NOTE: PROCEDURE SORT used (Total process time):

```
real time      0.32 seconds
cpu time       0.40 seconds
```

```
9
10     options mprint ;
11     * only persons in base and included before 1.1.2015 ;
12     data dvdd ;
13         merge dvdd ( in = dvdd )
14             TTDATA.pop ( in = pop )
15             TTdata.gdm ;
16     by pnr ;
17     if pop and dvdd ;
18     * remove status records after the cut date ;
19     if status_dato > &cutdate. then delete ;
20     * Do not count diagnosis or status dates in GDM grace period ;
21     %xgdm( diag_dato ) ;
MPRINT(XGDM):  if ( doGDM1 - 30 ) < diag_dato < ( doGDM1 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM2 - 30 ) < diag_dato < ( doGDM2 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM3 - 30 ) < diag_dato < ( doGDM3 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM4 - 30 ) < diag_dato < ( doGDM4 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM5 - 30 ) < diag_dato < ( doGDM5 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM6 - 30 ) < diag_dato < ( doGDM6 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM7 - 30 ) < diag_dato < ( doGDM7 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM8 - 30 ) < diag_dato < ( doGDM8 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM9 - 30 ) < diag_dato < ( doGDM9 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM10 - 30 ) < diag_dato < ( doGDM10 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM11 - 30 ) < diag_dato < ( doGDM11 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM12 - 30 ) < diag_dato < ( doGDM12 + 365 ) then delete ;
22     %xgdm( status_dato ) ;
MPRINT(XGDM):  if ( doGDM1 - 30 ) < status_dato < ( doGDM1 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM2 - 30 ) < status_dato < ( doGDM2 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM3 - 30 ) < status_dato < ( doGDM3 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM4 - 30 ) < status_dato < ( doGDM4 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM5 - 30 ) < status_dato < ( doGDM5 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM6 - 30 ) < status_dato < ( doGDM6 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM7 - 30 ) < status_dato < ( doGDM7 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM8 - 30 ) < status_dato < ( doGDM8 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM9 - 30 ) < status_dato < ( doGDM9 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM10 - 30 ) < status_dato < ( doGDM10 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM11 - 30 ) < status_dato < ( doGDM11 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM12 - 30 ) < status_dato < ( doGDM12 + 365 ) then delete ;
```



```

48      * DVDD will provide classification of follow-up as T1 / *not* T1 (=T2) ;
49      * tabulation of the sequences of type classifications occurring ;
50      proc sort data = dvdd out = dvdd_type ;
51          by pnr status_dato ;
52      run ;

```

NOTE: There were 344543 observations read from the data set WORK.DVDD.
NOTE: The data set WORK.DVDD_TYPE has 344543 observations and 7 variables.
NOTE: PROCEDURE SORT used (Total process time):
real time 0.07 seconds
cpu time 0.07 seconds

```

53
54      data dvdd_type ( keep = pnr doDVDD status_dato typ )
55          dvdd_hist ( keep = pnr hist ) ;
56          set dvdd_type ;
57          by pnr ;
58          length typ $ 4 hist $ 80 ;
59          retain hist ;
60          typ = substr( diag_type, 1, 2 ) ;
61          if typ eq "Ty" then typ = "T" || substr( diag_type, 6, 1 ) ;
62          if first.pnr then hist = typ ;
63          if ^first.pnr and ( diag_type ne lag(diag_type) )
64              then hist = trim(hist) || " " || typ ;
65          output dvdd_type ;
66          if last.pnr then output dvdd_hist ;
67      run ;

```

NOTE: There were 344543 observations read from the data set WORK.DVDD_TYPE.
NOTE: The data set WORK.DVDD_TYPE has 344543 observations and 4 variables.
NOTE: The data set WORK.DVDD_HIST has 88398 observations and 2 variables.
NOTE: DATA statement used (Total process time):
real time 0.14 seconds
cpu time 0.11 seconds

```

68
69      * classification rule: if more than half of registrations T1 then T1 ;
70      *                          if more than half of registrations T2 then T2 ;
71      data dvdd ( keep = pnr doDVDD dvdtyp nT1 nT2 nRc ) ;
72          set dvdd_type ;
73          by pnr status_dato ;
74          retain nT1 nT2 ;
75          if first.pnr then do ;
76              nT1 = 0 ;
77              nT2 = 0 ;
78              nRc = 0 ;
79          end ;
80          nT1 + ( typ eq "T1" ) ;
81          nT2 + ( typ eq "T2" ) ;
82          nRc + 1 ;
83          * If most and more than half of records agree on one type ;
84          if last.pnr then do ;
85              if nT1 > nRc/2 then dvdtyp = 'T1' ; else
86              if nT2 > nRc/2 then dvdtyp = 'T2' ; else dvdtyp = 'NA' ;
87          output ;
88          end ;
89      run ;

```

NOTE: There were 344543 observations read from the data set WORK.DVDD_TYPE.
NOTE: The data set WORK.DVDD has 88398 observations and 6 variables.
NOTE: DATA statement used (Total process time):
real time 0.07 seconds
cpu time 0.07 seconds

```

90
91      data TTdata.dvdd ;
92          merge dvdd

```

```

93             dvdd_hist ( keep = pnr hist ) ;
94             by pnr ;
95             run ;

```

NOTE: There were 88398 observations read from the data set WORK.DVDD.
NOTE: There were 88398 observations read from the data set WORK.DVDD_HIST.
NOTE: The data set TTDATA.DVDD has 88398 observations and 7 variables.
NOTE: DATA statement used (Total process time):

real time	0.20 seconds
cpu time	0.03 seconds

```

96
97             title1 'Dates and types from DVDD' ;
98             proc contents data = TTdata.dvdd ; run ;

```

NOTE: PROCEDURE CONTENTS used (Total process time):

real time	0.03 seconds
cpu time	0.03 seconds

NOTE: The PROCEDURE CONTENTS printed page 1.

```

99             title2 'Classification based on most frequent type recorded in DVDD - persons'
99             ! ;
100            proc tabulate data = TTdata.dvdd missing noseps ;
101                class dvdtyp doDVDD nT1 nT2 NRc ;
102                table all doDVDD,
103                    ( all dvdtyp ) * f=comma9.
104                    / rts = 8 ;
105                table nRc * nT1,
106                    nT2 * f=5.
107                    / rts = 5 indent = 1 ;
108                format doDVDD year4. ;
109            run ;

```

NOTE: There were 88398 observations read from the data set TTDATA.DVDD.
NOTE: The PROCEDURE TABULATE printed pages 2-3.
NOTE: PROCEDURE TABULATE used (Total process time):

real time	0.07 seconds
cpu time	0.03 seconds

```

110
111            proc tabulate data = TTdata.dvdd missing noseps order = freq ;
112                class dvdtyp hist ;
113                table all hist="sequence of different types",
114                    ( all dvdtyp ) * f=comma7.
115                    / rts = 30 ;
116            run ;

```

NOTE: There were 88398 observations read from the data set TTDATA.DVDD.
NOTE: The PROCEDURE TABULATE printed page 4.
NOTE: PROCEDURE TABULATE used (Total process time):

real time	0.01 seconds
cpu time	0.04 seconds

```

117            title1 ;

```

NOTE: SAS Institute Inc., SAS Campus Drive, Cary, NC USA 27513-2414
NOTE: The SAS System used:

real time	19.64 seconds
cpu time	6.39 seconds

7.6.1 02-dvdd.lst

Dates and types from DVDD

14:17 Tuesday, January 24, 2017 1

The CONTENTS Procedure

Data Set Name	TTDATA.DVDD	Observations	88398
Member Type	DATA	Variables	7
Engine	V9	Indexes	0
Created	24/01/2017 14:18:15	Observation Length	128
Last Modified	24/01/2017 14:18:15	Deleted Observations	0
Protection		Compressed	NO
Data Set Type		Sorted	NO
Label			
Data Representation	WINDOWS_64		
Encoding	wlatin1 Western (Windows)		

Engine/Host Dependent Information

Data Set Page Size	65536
Number of Data Set Pages	174
First Data Page	*
Max Obs per Page	511
Obs in First Data Page	498
Number of Data Set Repairs	0
ExtendObsCounter	YES
Filename	E:\workdata\705093\BxC\demoDM\DATA\dvdd.sas7bdat
Release Created	9.0401M3
Host Created	X64_SRV12

Alphabetic List of Variables and Attributes

#	Variable	Type	Len	Format	Informat	Label
2	doDVDD	Num	8			
6	dvdtyp	Char	*			
7	hist	Char	80			
5	nRc	Num	8			
3	nT1	Num	8			
4	nT2	Num	8			
1	pnr	Char	12	\$12.	\$10.	Personnummer

Dates and types from DVDD 14:17 Tuesday, January 24, 2017 2
 Classification based on most frequent type recorded in DVDD - persons

	dvdtyp			
	All	NA	T1	T2
	N	N	N	N
All	88,398	3,555	23,947	60,896
doDVDD				
1890	*	.	.	*
1899	*	.	.	*
1900	57	*	11	45
1901	*	.	.	*
1909	*	*	.	.
1919	*	.	.	*
1920	13	.	*	9
1923	*	*	.	.
1927	*	.	*	.
1931	*	.	.	*
1933	*	.	*	.
1934	*	.	*	.
1936	*	.	*	.
1937	*	.	*	.
1938	*	.	*	.
1939	*	.	*	*
1940	6	.	6	.
1941	*	.	*	.

1942	8	.	7	*
1943	*	.	*	.
1944	9	.	9	.
1945	16	.	15	*
1946	13	.	12	*
1947	21	.	21	.
1948	28	*	26	*
1949	26	.	26	.
1950	37	.	34	*
1951	34	.	34	.
1952	36	.	31	*
1953	40	*	38	*
1954	51	*	47	*
1955	80	*	74	*
1956	117	.	114	*
1957	86	.	84	*
1958	109	.	104	*
1959	101	*	94	*
1960	176	*	163	12
1961	173	*	162	6
1962	182	*	169	11
1963	191	*	179	10
1964	182	*	165	15
1965	185	*	170	13
1966	200	*	179	17
1967	211	*	188	22
1968	213	*	191	21
1969	237	*	207	28
1970	350	*	283	65
1971	301	*	261	36
1972	386	*	309	74
1973	352	6	291	55
1974	390	6	313	71
1975	430	7	316	107
1976	434	*	343	86
1977	468	7	351	110
1978	567	6	414	147
1979	545	9	392	144
1980	884	19	460	405
1981	590	7	411	172
1982	714	18	410	286
1983	711	8	417	286
1984	755	13	417	325
1985	1,058	19	420	619
1986	970	12	467	491
1987	1,055	22	476	557
1988	1,144	24	470	650
1989	1,170	30	499	641
1990	2,015	41	590	1,384
1991	1,407	36	543	828
1992	1,860	31	535	1,294
1993	1,708	30	501	1,177
1994	1,967	45	599	1,323
1995	2,664	56	594	2,014
1996	2,381	55	588	1,738
1997	2,499	65	634	1,800
1998	3,079	53	635	2,391
1999	2,825	62	556	2,207
2000	3,847	86	627	3,134
2001	3,143	79	668	2,396
2002	3,267	85	588	2,594
2003	3,317	96	544	2,677
2004	3,646	114	584	2,948
2005	3,889	104	566	3,219
2006	4,061	181	613	3,267
2007	3,989	177	576	3,236
2008	4,037	218	614	3,205
2009	3,649	290	533	2,826
2010	3,503	231	527	2,745
2011	2,999	231	432	2,336

2012	2,533	286	401	1,846
2013	2,180	324	341	1,515
2014	1,639	282	237	1,120
2015	154	38	18	98

Dates and types from DVDD 14:17 Tuesday, January 24, 2017 3
 Classification based on most frequent type recorded in DVDD - persons

		nT2													
		0	*	*	*	*	*	6	7	8	9	10	11	12	13
		N	N	N	N	N	N	N	N	N	N	N	N	N	N
*	0	1288	20100
*	2259
*	0	445	220	10634
*	58	173
*	1934
*	0	251	79	160	7239
*	24	7	133
*	29	107
*	1992
*	0	109	32	53	104	5229
*	8	*	10	146
*	9	8	79
*	37	116
*	1819
*	0	68	14	28	22	72	4614
*	*	*	*	*	*	100
*	8	*	*	85
*	17	*	82
*	30	119
*	2194
6	0	51	14	*	18	26	66	3534
*	*	*	*	*	*	*	122
*	10	*	.	*	96
*	7	.	*	80
*	9	*	59
*	35	84
6	2472
7	0	60	10	9	7	6	16	55	2822
*	6	.	*	*	*	*	*	101
*	7	.	*	*	*	*	104
*	6	*	*	*	65
*	*	*	*	66
*	9	*	83
6	27	101
7	3143
8	0	40	9	*	*	6	16	9	25	2382
*	*	.	.	*	*	.	*	*	53
*	.	*	.	*	*	.	*	50
*	*	*	*	*	*	57
*	*	.	.	*	64
*	6	*	*	79
6	12	*	60
7	24	67
8	2963
9	0	46	*	*	*	*	*	*	7	17	2103

*	*	.	.	*	*	.	.	*	59
*	*	.	*	.	.	*	*	34
*	*	.	.	*	*	*	49
*	*	.	.	*	*	41
*	*	*	*	*	55
6	6	*	.	39
7	7	*	33
8	24	70
9	3255
10														
0	*	*	*	.	*	257	.	.	.
*	*	.	7
*	.	*	*	.	*
*	*
*	*
6	.	.	.	*	*
7	.	.	*	*
8	*	*	6
9	11	8
10	319
11														
0	*	*	20	.	.
*	*	.	.	.
*	.	.	*	*
9	.	.	*
10	*	*
11	30
12														
0	*	.
10	.	.	*
12	*
13														
0	*
13	*

Dates and types from DVDD 14:17 Tuesday, January 24, 2017 4
 Classification based on most frequent type recorded in DVDD - persons

	dvdtyp			
	All	T2	T1	NA
	N	N	N	N
All	88,398	60,896	23,947	3,555
sequence of different types				
T2	58,940	58,940	.	.
T1	22,384	.	22,384	.
An	2,308	.	.	2,308
T2 T1	1,287	404	693	190
T1 T2	933	502	262	169
An T2	585	328	.	257
T2 T1 T2	369	342	15	12
T2 An	344	133	.	211
T1 T2 T1	274	18	248	8
An T1	192	.	129	63
T1 An	159	.	79	80
T2 An T2	111	107	.	*
T1 An T1	55	.	48	7
T2 T1 T2 T1	44	20	13	11
An T2 An	35	*	.	32
-I T2	35	30	.	*
T1 T2 T1 T2	28	13	6	9
-I	27	.	.	27
-I An	25	.	.	25
-I T1	21	.	19	*
An T2 T1	17	*	*	13

T1 T2 An	15	*	*	11
An T1 T2	15	*	*	9
T2 An T1	14	*	*	9
An T1 An	14	.	*	13
T2 T1 An	14	*	.	13
T2 T1 T2 T1 T2	13	12	.	*
T1 An T2	11	*	.	7
T2 -I T2	10	10	.	.
T1 -I T1	9	.	9	.
T2 T1 An T1	7	.	*	*
T2 T1 T2 An	6	*	*	*
An T2 An T2	6	*	.	*
-I An T2	6	.	.	6
T1 T2 An T1	*	.	*	*
T1 T2 T1 T2 T1	*	.	*	*
T2 T1 T2 T1 T2 T1	*	*	*	.
An T1 An T1	*	.	*	*
T2 -I	*	*	.	*
An T2 T1 T2	*	*	.	*
T1 An T2 T1	*	.	*	*
An -I An	*	.	.	*
-I T2 T1	*	.	*	*
T2 An T2 An	*	.	.	*
T1 An T1 An	*	.	*	*
An T2 An T1	*	.	.	*
An T1 T2 T1	*	.	*	*
-I T1 T2 T1	*	.	*	.
T1 T2 T1 An	*	.	*	*
-I An T1	*	.	*	*
T1 An T1 An T1	*	.	*	*
T2 T1 An T2	*	*	.	*
An T2 T1 An	*	*	.	*
-I T1 T2	*	*	*	.
T2 An T1 T2	*	*	.	.
An T1 An T2	*	.	.	*
T1 An T2 An	*	.	.	*
-I T2 T1 T2	*	.	.	*
-I An -I An T2	*	.	.	*
T1 -I An	*	.	.	*
T1 T2 An T2	*	.	.	*
An T2 An T2 An	*	.	.	*
An T1 T2 An	*	.	.	*
T1 -I An T1 T2	*	.	.	*
-I An T2 An	*	.	.	*
T1 T2 T1 T2 T1 T2 T1 T2	*	*	.	.
T1 An T1 T2 An	*	.	.	*
T2 An T1 T2 T1 An T1	*	.	*	.
T2 -I T1 T2 T1	*	.	.	*
An -I An T2 An	*	.	.	*
T2 -I An T2 An T2 T1	*	.	.	*
T1 -I An T2	*	.	.	*
An T2 T1 T2 T1	*	.	*	.
T1 T2 An T2 An T1	*	.	.	*
T1 T2 T1 T2 T1 T2 T1	*	.	*	.
-I T2 T1 T2 T1	*	*	.	.
T2 An T1 T2 T1	*	*	.	.
-I An T1 An	*	.	.	*
T2 An T1 An	*	.	.	*

7.7 03-ndr

Constructs a version of the old NDR without using the blood-glucose criteria and the foot-therapy criterion.

NOTE: Copyright (c) 2002-2012 by SAS Institute Inc., Cary, NC, USA.
 NOTE: SAS (r) Proprietary Software 9.4 (TS1M3)
 Licensed to FORSKNING 1, Site 50800722.
 NOTE: This session is executing on the X64_SRV12 platform.

NOTE: Updated analytical products:
 SAS/STAT 14.1

NOTE: Additional host information:
 X64_SRV12 WIN 6.2.9200 Server

NOTE: SAS initialization used:
 real time 0.07 seconds
 cpu time 0.06 seconds

NOTE: AUTOEXEC processing beginning; file is E:\workdata\705093\BXC\demoDM\sas\optslibs.sas.

NOTE: AUTOEXEC processing completed.

```
1 *****
2 * NDR ;
3 proc sort data = EKST15.ndr out = ndr ;
4   by pnr ;
5   run ;
```

NOTE: Input data set is already sorted; it has been copied to the output data set.
 NOTE: There were 523770 observations read from the data set EKST15.NDR.
 NOTE: The data set WORK.NDR has 523770 observations and 9 variables.
 NOTE: PROCEDURE SORT used (Total process time):
 real time 0.54 seconds
 cpu time 0.07 seconds

```
6
7   * Only persons present in the base (TTDATA.bef_grund) ;
8   data TTDATA.NDR ( keep = pnr sex doNDR datNDR doBth doDth inklaars ) ;
9     merge ndr ( in = ndr
10              rename = ( d_inkldto = DoNDR ) )
11           TTDATA.pop ( in = pop ) ;
12   by pnr ;
13   if ndr and pop ;
14   * revised date of inclusion (ignoring blood glucose criteria) ;
15   doNDR = min( d_lpr, /*d_fodt,*/ d_ins, d_oad ) ;
16   if doNDR eq d_lpr then inklaars = 'lpr' ;
17   * if doNDR eq d_fodt then inklaars = 'fod' ;
18   if doNDR eq d_oad then inklaars = 'oad' ;
19   if doNDR eq d_ins then inklaars = 'ins' ;
20   * To avoid small cell entries in overview table ;
21   datNDR = max( doNDR, '01JAN1990'd ) ;
22   label datNDR="doDM" ;
23   if doNDR gt .z ;
24   run ;
```

NOTE: Missing values were generated as a result of performing an operation on missing values.
 Each place is given by: (Number of times) at (Line):(Column).
 106885 at 15:12

NOTE: There were 523770 observations read from the data set WORK.NDR.
 NOTE: There were 7361669 observations read from the data set TTDATA.POP.
 NOTE: The data set TTDATA.NDR has 389304 observations and 7 variables.
 NOTE: DATA statement used (Total process time):
 real time 2.67 seconds
 cpu time 1.28 seconds


```

25
26      title1 'Persons from NDR with valid corrected DM dates' ;
27      title2 'datNDR is the doNDR left censored at 1.1.1990' ;
28      proc tabulate data = TTdata.NDR missing noseps formchar='      ' ;
29          class datNDR inklaars ;
30          table all datNDR,
31              ( all inklaars ) * f=comma7.
32              ( all * f= 8.1
33                inklaars * f=4.1 ) * pctn< all inklaars >
34              / rts = 8 ;
35          format datNDR year4. ;
36          keylabel n = " " ;
37      run ;

```

NOTE: There were 389304 observations read from the data set TTDATA.NDR.

NOTE: At least one W.D format was too small for the number to be printed. The decimal may be shifted by the "BEST" format.

NOTE: The PROCEDURE TABULATE printed page 1.

NOTE: PROCEDURE TABULATE used (Total process time):

```

real time      0.15 seconds
cpu time       0.15 seconds

```

```

38      title2 ;
39
40      proc contents data = TTdata.ndr ;
41      run ;

```

NOTE: PROCEDURE CONTENTS used (Total process time):

```

real time      0.00 seconds
cpu time       0.00 seconds

```

NOTE: The PROCEDURE CONTENTS printed page 2.

NOTE: SAS Institute Inc., SAS Campus Drive, Cary, NC USA 27513-2414

NOTE: The SAS System used:

```

real time      3.53 seconds
cpu time       1.59 seconds

```

7.7.1 03-ndr.lst

Persons from NDR with valid corrected DM dates
datNDR is the doNDR left censored at 1.1.1990

10:42 Tuesday, January 24, 2017 1

	inklaars							
	inklaars				All	ins	lpr	oad
	All	ins	lpr	oad	PctN	PctN	PctN	PctN
All	389,304	10,163	181,868	197,273	100.0	100	100	100
doDM								
1990	12,842	.	12,842	.	100.0	.	100	.
1991	8,854	.	8,854	.	100.0	.	100	.
1992	8,236	.	8,236	.	100.0	.	100	.
1993	12,362	.	12,362	.	100.0	.	100	.
1994	37,349	6,094	7,037	24,218	100.0	16.3	18.8	64.8
1995	12,458	598	5,877	5,983	100.0	4.8	47.2	48.0
1996	12,476	317	6,013	6,146	100.0	2.5	48.2	49.3
1997	12,027	187	5,964	5,876	100.0	1.6	49.6	48.9
1998	13,175	177	6,468	6,530	100.0	1.3	49.1	49.6
1999	13,636	191	6,723	6,722	100.0	1.4	49.3	49.3
2000	14,123	148	7,338	6,637	100.0	1.0	52.0	47.0
2001	14,996	139	7,646	7,211	100.0	0.9	51.0	48.1
2002	15,386	163	8,042	7,181	100.0	1.1	52.3	46.7
2003	17,208	148	8,659	8,401	100.0	0.9	50.3	48.8

2004	17,749	156	8,440	9,153	100.0	0.9	47.6	51.6
2005	17,439	163	8,236	9,040	100.0	0.9	47.2	51.8
2006	18,062	177	8,313	9,572	100.0	1.0	46.0	53.0
2007	18,966	241	8,153	10,572	100.0	1.3	43.0	55.7
2008	20,088	264	7,738	12,086	100.0	1.3	38.5	60.2
2009	20,951	262	7,652	13,037	100.0	1.3	36.5	62.2
2010	22,396	236	7,485	14,675	100.0	1.1	33.4	65.5
2011	24,997	265	7,192	17,540	100.0	1.1	28.8	70.2
2012	23,528	237	6,598	16,693	100.0	1.0	28.0	70.9

Persons from NDR with valid corrected DM dates

10:42 Tuesday, January 24, 2017 2

The CONTENTS Procedure

Data Set Name	TTDATA.NDR	Observations	389304
Member Type	DATA	Variables	7
Engine	V9	Indexes	0
Created	24/01/2017 10:42:45	Observation Length	56
Last Modified	24/01/2017 10:42:45	Deleted Observations	0
Protection		Compressed	NO
Data Set Type		Sorted	NO
Label			
Data Representation	WINDOWS_64		
Encoding	wlatin1 Western (Windows)		

Engine/Host Dependent Information

Data Set Page Size	65536
Number of Data Set Pages	334
First Data Page	*
Max Obs per Page	1167
Obs in First Data Page	1136
Number of Data Set Repairs	0
ExtendObsCounter	YES
Filename	E:\workdata\705093\BxC\demoDM\DATA\ndr.sas7bdat
Release Created	9.0401M3
Host Created	X64_SRV12

Alphabetic List of Variables and Attributes

#	Variable	Type	Len	Format	Informat	Label
2	DoNDR	Num	8	DATE9.	DATE9.	D_INKLDTO
7	datNDR	Num	8			doDM
4	doBth	Num	8			
5	doDth	Num	8			
6	inklaars	Char	*			
1	pnr	Char	12	\$12.	\$10.	V_CPR
3	sex	Num	8			

7.8 04-rmps

Processes the records from the RMPS with other target medications and creates a file (pRMPS) with one record per person with at least one prescription of either OAD or insulin.

Uses the GDM dates to exclude possible inclusion dates in GDM grace periods.

1

"Program 04-rmps.sas"

NOTE: Copyright (c) 2002-2012 by SAS Institute Inc., Cary, NC, USA.

NOTE: SAS (r) Proprietary Software 9.4 (TS1M3)

Licensed to FORSKNING 1, Site 50800722.

NOTE: This session is executing on the X64_SRV12 platform.

NOTE: Updated analytical products:

SAS/STAT 14.1

NOTE: Additional host information:

X64_SRV12 WIN 6.2.9200 Server

NOTE: SAS initialization used:

real time 0.07 seconds
cpu time 0.09 seconds

NOTE: AUTOEXEC processing beginning; file is E:\workdata\705093\BXC\demoDM\sas\optslibs.sas.

NOTE: AUTOEXEC processing completed.

```

1      %macro med ;
2      data rmps ;
3          length druggr $ 7 ;
4          set %do i = 1995 %to 2015 ;
5              GRUND15.lmdb&i. ( keep = pnr ATC eksd apk volume packsize doso
6                  where = ( substr(atc,1,3) eq 'A10' ) )
7          %end ; ;
8          * Grouping of drugs as a character ;
9              druggr = put(          atc          , $atc5grp. ) ;
10         if ( druggr eq "Other" ) then druggr = put( substr(atc,1,5), $atc4grp. ) ;
11     run ;
12     %mend ;
13     %med ;

```

NOTE: There were 561599 observations read from the data set GRUND15.LMDB1995.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 626172 observations read from the data set GRUND15.LMDB1996.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 670759 observations read from the data set GRUND15.LMDB1997.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 732363 observations read from the data set GRUND15.LMDB1998.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 795049 observations read from the data set GRUND15.LMDB1999.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 826889 observations read from the data set GRUND15.LMDB2000.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 906757 observations read from the data set GRUND15.LMDB2001.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 985120 observations read from the data set GRUND15.LMDB2002.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 1066934 observations read from the data set GRUND15.LMDB2003.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 1179915 observations read from the data set GRUND15.LMDB2004.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 1290865 observations read from the data set GRUND15.LMDB2005.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 1407844 observations read from the data set GRUND15.LMDB2006.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 1519880 observations read from the data set GRUND15.LMDB2007.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 1659373 observations read from the data set GRUND15.LMDB2008.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 1751002 observations read from the data set GRUND15.LMDB2009.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 1875270 observations read from the data set GRUND15.LMDB2010.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 2003071 observations read from the data set GRUND15.LMDB2011.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 2101642 observations read from the data set GRUND15.LMDB2012.

WHERE SUBSTR(atc, 1, 3)='A10';

NOTE: There were 2130915 observations read from the data set GRUND15.LMDB2013.

```

WHERE SUBSTR(atc, 1, 3)='A10';
NOTE: There were 2152130 observations read from the data set GRUND15.LMDB2014.
WHERE SUBSTR(atc, 1, 3)='A10';
NOTE: There were 2164414 observations read from the data set GRUND15.LMDB2015.
WHERE SUBSTR(atc, 1, 3)='A10';
NOTE: The data set WORK.RMPS has 28407963 observations and 8 variables.
NOTE: DATA statement used (Total process time):
      real time          4:43.26
      cpu time           43.21 seconds

```

```

14
15      proc sort  data = rmps ;  by pnr druggr eksd ;  run ;

```

```

NOTE: There were 28407963 observations read from the data set WORK.RMPS.
NOTE: The data set WORK.RMPS has 28407963 observations and 8 variables.
NOTE: PROCEDURE SORT used (Total process time):
      real time          6.21 seconds
      cpu time           12.67 seconds

```

```

16
17      * Generate one record per person with a drug dispensation
18      but excluding dispensations in the time after GDM and PCOS ;
19      options mprint ;
20      data rmps
21          pospcos ( keep = pnr pospcos ) ;
22      merge rmps      ( in = r )
23          TTdata.GDM
24          TTdata.PCOS ( in = p )
25          TTdata.pop  ( in = b  keep = pnr doBth ) ;
26      by pnr ;
27      if r and b ;
28      pcos = p ;
29      retain pospcos ;
30      * exclude dispensations in GDM-window ;
31      %xgdm( eksd ) ;
MPRINT(XGDM):  if ( doGDM1 - 30 ) < eksd < ( doGDM1 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM2 - 30 ) < eksd < ( doGDM2 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM3 - 30 ) < eksd < ( doGDM3 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM4 - 30 ) < eksd < ( doGDM4 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM5 - 30 ) < eksd < ( doGDM5 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM6 - 30 ) < eksd < ( doGDM6 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM7 - 30 ) < eksd < ( doGDM7 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM8 - 30 ) < eksd < ( doGDM8 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM9 - 30 ) < eksd < ( doGDM9 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM10 - 30 ) < eksd < ( doGDM10 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM11 - 30 ) < eksd < ( doGDM11 + 365 ) then delete ;
MPRINT(XGDM):  if ( doGDM12 - 30 ) < eksd < ( doGDM12 + 365 ) then delete ;
32      * exclude metformin dispensations after PCOS if before age 40 ;
33      if pcos and
34          druggr eq '11' and
35          ( doPCOS - 30 ) < eksd < ( doBth + 365.25*40 ) then delete ;
36      output rmps ;
37      * if metformin (druggr=11) is the only dispensation between age 20 and 40,
38      it is possibly PCOS so the person is flagged by pospcos ;
39      if first.pnr then pospcos =
40          ( doBth + 365.25*20 ) < eksd < ( doBth + 365.25*40 ) and druggr eq '11' ;
41      if ( doBth + 365.25*20 ) < eksd < ( doBth + 365.25*40 ) and druggr ne '11'
42          then pospcos = 0 ;
43      if last.pnr and pospcos then output pospcos ;
44      run ;

```

NOTE: Variable doGDM12 is uninitialized.

NOTE: Missing values were generated as a result of performing an operation on missing values.

Each place is given by: (Number of times) at (Line):(Column).

```

27625444 at 31:18  27625444 at 31:54  28022030 at 31:20  28022030 at 31:56
28236064 at 31:20  28236064 at 31:56  28309326 at 31:20  28309326 at 31:56
28328127 at 31:20  28328127 at 31:56  28333628 at 31:20  28333628 at 31:56

```

```

28334561 at 31:20 28334561 at 31:56 28334796 at 31:20 28334796 at 31:56
28334831 at 31:20 28334831 at 31:56 28334831 at 31:20 28334831 at 31:56
28334831 at 31:20 28334831 at 31:56 28334831 at 31:20 28334831 at 31:56

```

NOTE: There were 28407963 observations read from the data set WORK.RMPS.

NOTE: There were 21146 observations read from the data set TTDATA.GDM.

NOTE: There were 20773 observations read from the data set TTDATA.PCOS.

NOTE: There were 7361669 observations read from the data set TTDATA.POP.

NOTE: The data set WORK.RMPS has 28295912 observations and 24 variables.

NOTE: The data set WORK.POSPCOS has 19885 observations and 2 variables.

NOTE: DATA statement used (Total process time):

```

real time          1:28.60
cpu time           1:22.62

```

```

45     options nomprint ;
46
47     * Exclude metformin dispensation between 20 and 40 for those who start
48     ! metformin
49     after 20 oand only fill metformin before age 40 ;
49     data rmps ;
50     merge rmps pospcos ;
51     by pnr ;
52     drop pospcos ;
53     if pospcos and
54     ( doBth + 365.25*20 ) < eksd < ( doBth + 365.25*40 )
55     then delete ;
56     run ;

```

NOTE: There were 28295912 observations read from the data set WORK.RMPS.

NOTE: There were 19885 observations read from the data set WORK.POSPCOS.

NOTE: The data set WORK.RMPS has 27943194 observations and 23 variables.

NOTE: DATA statement used (Total process time):

```

real time          16.07 seconds
cpu time           7.70 seconds

```

```

57
58     * Naming convention for date variables :
59     "donlXX" is first date of single drug medication with XX
60     "danyXX" is first date of *any* medication with drug XX
61     Thus "donl" is for selection on mono-users, otherwise "dany" is used.
62     We put the names of these data variables into a macro variable and
63     later in an array --- note that insulins are the last 5 entries in the array
63     ! ;
64     %let drdates = donlMet    danyMet
65                     donlSU    danySU
66                     donlTZD   danyTZD
67                     donlDPP4  danyDPP4
68                     danyGLP1
69                     donlSGLT2 danySGLT2
70                     donlAca   danyAca
71                     danyMeg
72                     dofIns doiIns domIns dolIns doIns ;
73
74     data TTdata.pRMPS ( keep = pnr &drdates. doOAD doRMPS ) ;
75     set rmps ;
76     by pnr druggr eksd ;
77     retain          &drdates. ;
78     array drd [19] &drdates. ;
79     * initiate all drugdates to missing ;
80     if first.pnr then do i = 1 to dim(drd) ; drd(i) = . ; end ;
81     * dates of first exposure ;
82     if first.druggr then do ;
83         if druggr eq '11' then donlMet = eksd ;
84         if index(substr(druggr,2,2),'1') then danyMet = min(eksd,danyMet) ;
85         if druggr eq '12' then donlSU = eksd ;
86         if index(substr(druggr,2,2),'2') then danySU = min(eksd,danySU) ;
87         if druggr eq '13' then donlTZD = eksd ;
88         if index(substr(druggr,2,2),'3') then danyTZD = min(eksd,danyTZD) ;
89         if druggr eq '14' then donlDPP4 = eksd ;

```

```

90         if index(substr(druggr,2,2),'4') then danyDPP4 = min(eksd,danyDPP4) ;
91         if druggr eq '15' then danyGLP1 = eksd ;
92         if druggr eq '16' then donlSGLT2= eksd ;
93         if index(substr(druggr,2,2),'6') then danySGLT2= min(eksd,danySGLT2) ;
94         if druggr eq '18' then donlAca = eksd ;
95         if index(substr(druggr,2,2),'8') then danyAca = min(eksd,danyAca) ;
96         if druggr eq '19' then danyMeg = eksd ;
97         if druggr eq 'fastIns' then dofIns = eksd ;
98         if druggr eq 'intIns' then doiIns = eksd ;
99         if druggr eq 'mixIns' then domIns = eksd ;
100        if druggr eq 'longIns' then dolIns = eksd ;
101        end ;
102        * Date of any drugexposure / resp. any OAD (i.e. non-ins) exposure ;
103        if last.pnr then do ; doRMPS = min( of drd[*] ) ;
104        do i = 1 to dim(drd)-5 ; doOAD = min( doOAD, drd[i] ) ; end ;
105        doIns = min( dofIns, doiIns, domIns, dolIns ) ;
106        output ;
107        end ;
108        run ;

```

NOTE: Missing values were generated as a result of performing an operation on missing values.

Each place is given by: (Number of times) at (Line):(Column).

881239 at 104:40 260604 at 105:14

NOTE: There were 27943194 observations read from the data set WORK.RMPS.

NOTE: The data set TTDATA.PRMPMS has 409844 observations and 22 variables.

NOTE: DATA statement used (Total process time):

real time 10.76 seconds
cpu time 6.65 seconds

```

109
110        title1 "Persons from the RPMS" ;
111        proc contents data = TTdata.pRMPS ; run ;

```

NOTE: PROCEDURE CONTENTS used (Total process time):

real time 0.03 seconds
cpu time 0.03 seconds

NOTE: The PROCEDURE CONTENTS printed page 1.

```

112        proc tabulate data = TTdata.pRMPS missing noseps ;
113        class &drdates. doOAD doRMPS ;
114        table all &drdates. doOAD doRMPS,
115        n * f=comma10.
116        / rts = 15 ;
117        table all doOAD,
118        ( all doINS ) * f=comma7.
119        / rts = 7 ;
120        format &drdates. doOAD doRMPS year4. ;
121        keylabel n = ' ' ;
122        run ;

```

NOTE: There were 409844 observations read from the data set TTDATA.PRMPMS.

NOTE: The PROCEDURE TABULATE printed pages 2-5.

NOTE: PROCEDURE TABULATE used (Total process time):

real time 0.42 seconds
cpu time 1.11 seconds

```

123        title1 ;
124
125

```

NOTE: SAS Institute Inc., SAS Campus Drive, Cary, NC USA 27513-2414

NOTE: The SAS System used:

real time 6:45.54
cpu time 2:34.10

7.8.1 04-rmps.lst

Persons from the RPMS

14:04 Tuesday, January 24, 2017 1

The CONTENTS Procedure

Data Set Name	TTDATA.PRMP5	Observations	409844
Member Type	DATA	Variables	22
Engine	V9	Indexes	0
Created	24/01/2017 14:10:53	Observation Length	184
Last Modified	24/01/2017 14:10:53	Deleted Observations	0
Protection		Compressed	NO
Data Set Type		Sorted	NO
Label			
Data Representation	WINDOWS_64		
Encoding	wlatin1 Western (Windows)		

Engine/Host Dependent Information

Data Set Page Size	65536
Number of Data Set Pages	1155
First Data Page	*
Max Obs per Page	355
Obs in First Data Page	338
Number of Data Set Repairs	0
ExtendObsCounter	YES
Filename	E:\workdata\705093\BxC\demoDM\DATA\prmps.sas7bdat
Release Created	9.0401M3
Host Created	X64_SRV12

Alphabetic List of Variables and Attributes

#	Variable	Type	Len	Format	Informat	Label
14	danyAca	Num	8			
9	danyDPP4	Num	8			
10	danyGLP1	Num	8			
15	danyMeg	Num	8			
*	danyMet	Num	8			
12	danySGLT2	Num	8			
*	danySU	Num	8			
7	danyTZD	Num	8			
20	doIns	Num	8			
22	doOAD	Num	8			
21	doRMPS	Num	8			
16	dofIns	Num	8			
17	doiIns	Num	8			
19	dolIns	Num	8			
18	domIns	Num	8			
13	donlAca	Num	8			
8	donLDPP4	Num	8			
*	donlMet	Num	8			
11	donLSGLT2	Num	8			
*	donLSU	Num	8			
6	donlTZD	Num	8			
*	pnr	Char	12	\$12.	\$10.	Personnummer

Persons from the RPMS

14:04 Tuesday, January 24, 2017 2

All	409,844
donlMet	
.	117,064
1995	13,332
1996	4,358
1997	4,065
1998	4,948
1999	6,813
2000	8,906

2001	9,388
2002	9,974
2003	11,755
2004	13,278
2005	13,319
2006	14,407
2007	16,728
2008	19,263
2009	21,005
2010	22,782
2011	27,064
2012	23,187
2013	16,641
2014	15,096
2015	16,471
danyMet	
.	115,985
1995	13,332
1996	4,358
1997	4,065
1998	4,948
1999	6,813
2000	8,906
2001	9,388
2002	9,974
2003	11,755
2004	13,339
2005	13,382
2006	14,554
2007	16,856
2008	19,372
2009	21,139
2010	22,897
2011	27,106
2012	23,267
2013	16,709
2014	15,149
2015	16,550
donlsu	
.	214,656
1995	45,851
1996	10,198
1997	9,615
1998	10,402
1999	10,093
2000	9,110
2001	9,389
2002	8,772
2003	10,016
2004	9,589
2005	9,099
2006	8,298
2007	8,183
2008	7,272
2009	6,004
2010	5,907
2011	4,633
2012	3,637
2013	3,255
2014	2,911
2015	2,954
danySU	
.	214,655
1995	45,851
1996	10,198
1997	9,615
1998	10,402
1999	10,093
2000	9,110
2001	9,389

2002	8,772
2003	10,016
2004	9,589
2005	9,099
2006	8,298
2007	8,184
2008	7,273
2009	6,005
2010	5,906
2011	4,632
2012	3,637
2013	3,255
2014	2,911
2015	2,954
donlTZD	
.	405,171
2000	100
2001	652
2002	424
2003	333
2004	426
2005	468
2006	584
2007	573
2008	390
2009	303
2010	178
2011	90
2012	52
2013	39
2014	36
2015	25
danyTZD	
.	399,747
2000	100
2001	652
2002	424
2003	334
2004	1,114
2005	1,422
2006	2,126
2007	1,847
2008	952
2009	617
2010	312
2011	56
2012	43
2013	36
2014	37
2015	25
donlDPP4	
.	369,776
2007	2,290
2008	4,452
2009	3,239
2010	4,425
2011	4,669
2012	4,471
2013	5,055
2014	5,418
2015	6,049
danyDPP4	
.	349,498
2007	2,290
2008	5,153
2009	5,464
2010	7,541
2011	7,764
2012	7,453
2013	7,830

2014	8,125
2015	8,726
danyGLP1	
.	377,256
2007	176
2008	1,030
2009	2,217
2010	7,256
2011	6,471
2012	5,235
2013	3,566
2014	2,969
2015	3,668
donlSGLT2	
.	400,423
2012	20
2013	1,948
2014	2,958
2015	4,495
danySGLT2	
.	400,183
2012	20
2013	1,948
2014	3,041
2015	4,652
donlAca	
.	402,362
1995	685
1996	1,232
1997	1,174
1998	1,172
1999	712
2000	453
2001	427
2002	246
2003	221
2004	193
2005	173
2006	134
2007	115
2008	111
2009	72
2010	95
2011	57
2012	75
2013	55
2014	43
2015	37
danyAca	
.	402,362
1995	685
1996	1,232
1997	1,174
1998	1,172
1999	712
2000	453
2001	427
2002	246
2003	221
2004	193
2005	173
2006	134
2007	115
2008	111
2009	72
2010	95
2011	57
2012	75
2013	55
2014	43

2015	37
danyMeg	
.	409,844
dofIns	
.	333,788
1995	17,490
1996	3,203
1997	2,427
1998	2,234
1999	2,276
2000	2,240
2001	2,206
2002	2,311
2003	2,325
2004	2,536
2005	2,610
2006	2,697
2007	2,832
2008	2,983
2009	3,189
2010	3,308
2011	3,407
2012	3,531
2013	3,856
2014	4,065
2015	4,330
doiIns	
.	308,325
1995	29,190
1996	4,509
1997	3,616
1998	3,742
1999	4,027
2000	4,188
2001	4,102
2002	4,421
2003	4,844
2004	4,684
2005	4,400
2006	4,206
2007	4,143
2008	3,011
2009	2,590
2010	2,568
2011	2,579
2012	2,535
2013	3,075
2014	2,740
2015	2,349
domIns	
.	348,584
1995	7,828
1996	2,766
1997	2,215
1998	2,610
1999	2,753
2000	2,688
2001	2,746
2002	2,705
2003	3,533
2004	4,116
2005	4,025
2006	4,321
2007	4,448
2008	3,152
2009	2,582
2010	1,934
2011	1,761
2012	1,539
2013	1,406

2014	1,191
2015	941
dolIns	
.	349,929
1996	10
1997	6
1998	*
1999	*
2000	*
2004	2,590
2005	4,688
2006	3,149
2007	2,943
2008	6,758
2009	6,778
2010	5,630
2011	5,239
2012	5,210
2013	4,747
2014	5,486
2015	6,671
doIns	
.	260,604
1995	33,463
1996	4,865
1997	4,009
1998	4,347
1999	4,804
2000	4,977
2001	4,895
2002	5,204
2003	5,947
2004	6,395
2005	6,231
2006	6,305
2007	6,552
2008	6,194
2009	6,257
2010	5,957
2011	6,282
2012	6,303
2013	6,616
2014	6,641
2015	6,996
doOAD	
.	52,998
1995	48,996
1996	10,882
1997	10,141
1998	11,113
1999	11,421
2000	11,548
2001	12,223
2002	12,126
2003	14,339
2004	14,722
2005	14,646
2006	15,150
2007	16,588
2008	18,298
2009	19,160
2010	21,329
2011	25,032
2012	21,853
2013	15,984
2014	14,747
2015	16,548
doRMPS	
1995	79,747
1996	12,916

1997	11,321
1998	12,245
1999	12,435
2000	12,605
2001	13,250
2002	13,132
2003	15,361
2004	15,909
2005	15,757
2006	16,377
2007	17,715
2008	19,427
2009	20,194
2010	22,314
2011	25,900
2012	22,673
2013	17,089
2014	15,865
2015	17,612

Persons from the RPMS

14:04 Tuesday, January 24, 2017 3

		doIns									
		All	.	1995	1996	1997	1998	1999	2000	2001	2002
All	409,844	260,604	33,463	4,865	4,009	4,347	4,804	4,804	4,977	4,895	5,204
doOAD											
.	52,998	.	27,298	2,044	1,214	1,151	1,137	1,137	1,172	1,147	1,169
1995	48,996	26,300	2,712	2,170	1,970	1,993	2,069	2,069	1,855	1,415	1,416
1996	10,882	5,917	375	286	259	277	329	329	365	329	338
1997	10,141	5,587	273	48	279	252	282	282	274	337	307
1998	11,113	6,070	278	26	64	325	298	298	299	336	333
1999	11,421	6,354	293	33	30	75	381	381	284	289	327
2000	11,548	6,609	275	29	24	41	67	67	407	292	276
2001	12,223	7,163	241	17	22	26	27	27	77	460	300
2002	12,126	7,449	226	24	19	23	36	36	40	63	470
2003	14,339	9,283	179	19	20	20	32	32	26	35	61
2004	14,722	10,191	130	17	7	16	21	21	23	29	32
2005	14,646	10,402	122	9	9	14	18	18	19	29	31
2006	15,150	11,280	101	15	12	15	16	16	13	18	23
2007	16,588	12,824	127	17	15	24	17	17	14	17	21
2008	18,298	14,886	110	15	9	11	11	11	23	12	15
2009	19,160	16,022	135	17	9	17	11	11	18	10	18
2010	21,329	18,556	120	10	8	12	16	16	11	19	18
2011	25,032	22,582	127	14	9	16	7	7	17	19	17
2012	21,853	19,831	127	12	6	16	8	8	17	12	*
2013	15,984	14,429	91	8	8	14	8	8	9	9	11
2014	14,747	13,451	63	20	9	*	*	*	*	14	7
2015	16,548	15,418	60	15	7	*	9	9	9	*	9

(Continued)

Persons from the RPMS

14:04 Tuesday, January 24, 2017 4

		doIns									
		2003	2004	2005	2006	2007	2008	2009	2010	2011	2012
All	5,947	6,395	6,231	6,305	6,552	6,194	6,257	6,257	5,957	6,282	6,303
doOAD											
.	1,147	1,227	1,210	1,287	1,287	1,316	1,306	1,306	1,291	1,239	1,188
1995	1,403	1,201	931	772	629	438	364	364	311	290	239
1996	357	363	282	277	243	176	151	151	121	118	91
1997	377	351	318	263	233	187	139	139	141	139	114
1998	425	407	360	314	319	240	199	199	179	168	146

1999	407	419	351	338	341	279	235	188	184	173
2000	385	381	381	350	362	280	271	212	217	198
2001	377	394	402	365	414	328	300	259	243	242
2002	322	382	362	352	341	306	322	251	261	226
2003	480	413	408	424	447	368	375	308	332	292
2004	63	559	383	348	346	388	353	295	343	300
2005	22	64	605	370	371	323	336	310	325	354
2006	26	31	60	575	355	333	373	311	336	334
2007	19	34	41	61	617	338	340	288	347	328
2008	29	31	21	32	73	689	317	293	284	342
2009	19	28	23	31	36	64	712	312	303	312
2010	21	26	24	40	47	37	75	709	321	277
2011	23	23	25	27	25	27	31	73	723	320
2012	15	20	18	28	17	23	31	59	54	727
2013	12	13	10	20	11	18	14	15	22	61
2014	11	10	7	19	19	21	*	20	16	16
2015	7	18	9	12	19	15	8	11	17	23

(Continued)

Persons from the RPMS

14:04 Tuesday, January 24, 2017 5

	doIns		
	2013	2014	2015
All	6,616	6,641	6,996
doOAD			
.	1,380	1,380	1,408
1995	200	175	143
1996	83	76	69
1997	80	82	78
1998	144	92	91
1999	164	143	133
2000	160	168	163
2001	229	176	161
2002	229	223	199
2003	281	276	260
2004	320	278	280
2005	317	284	312
2006	312	307	304
2007	358	381	360
2008	359	361	375
2009	346	344	373
2010	283	310	389
2011	288	313	326
2012	299	262	266
2013	705	235	261
2014	57	709	259
2015	22	66	786

7.9 05-diab

Uses eye-screening dates from the national eye-screening database to supplement persons with diabetes and update dates of diabetes diagnosis. Uses the clinically recorded type of diabetes to define the status of the patients.

Uses the dataset with GDM dates to exclude inclusion dates in GDM grace periods.

1

"Program 05-diab.sas"

NOTE: Copyright (c) 2002-2012 by SAS Institute Inc., Cary, NC, USA.

NOTE: SAS (r) Proprietary Software 9.4 (TS1M3)

Licensed to FORSKNING 2, Site 50800723.

NOTE: This session is executing on the X64_SRV12 platform.

NOTE: Updated analytical products:

SAS/STAT 14.1

NOTE: Additional host information:

X64_SRV12 WIN 6.2.9200 Server

NOTE: SAS initialization used:

real time 0.07 seconds
cpu time 0.09 seconds

NOTE: AUTOEXEC processing beginning; file is E:\workdata\705093\BXC\demoDM\sas\optslibs.sas.

NOTE: AUTOEXEC processing completed.

```
1          title1 'Original Diabase' ;
2          proc contents data = ekst15.diabasen ; run ;
```

NOTE: PROCEDURE CONTENTS used (Total process time):

real time 0.03 seconds
cpu time 0.01 seconds

NOTE: The PROCEDURE CONTENTS printed page 1.

```
3          title1 ;
4
5          proc sort data = ekst15.diabasen
6                  out = diab ;
7          by pnr Report_EyeScreeningDate ;
8          run ;
```

NOTE: There were 98929 observations read from the data set EKST15.DIABASEN.

NOTE: The data set WORK.DIAB has 98929 observations and 13 variables.

NOTE: PROCEDURE SORT used (Total process time):

real time 0.15 seconds
cpu time 0.10 seconds

```
9
10         options mprint ;
11         data diab ;
12         merge diab ( in = d )
13             TTdata.GDM ;
14         by pnr ;
15         * exclude visits in GDM grace period ;
16         %xgdm( Report_EyeScreeningDate ) ;
MPRINT(XGDM):  if ( doGDM1 - 30 ) < Report_EyeScreeningDate < ( doGDM1 + 365 ) then
delete ;
MPRINT(XGDM):  if ( doGDM2 - 30 ) < Report_EyeScreeningDate < ( doGDM2 + 365 ) then
delete ;
MPRINT(XGDM):  if ( doGDM3 - 30 ) < Report_EyeScreeningDate < ( doGDM3 + 365 ) then
delete ;
MPRINT(XGDM):  if ( doGDM4 - 30 ) < Report_EyeScreeningDate < ( doGDM4 + 365 ) then
delete ;
MPRINT(XGDM):  if ( doGDM5 - 30 ) < Report_EyeScreeningDate < ( doGDM5 + 365 ) then
delete ;
MPRINT(XGDM):  if ( doGDM6 - 30 ) < Report_EyeScreeningDate < ( doGDM6 + 365 ) then
delete ;
MPRINT(XGDM):  if ( doGDM7 - 30 ) < Report_EyeScreeningDate < ( doGDM7 + 365 ) then
delete ;
MPRINT(XGDM):  if ( doGDM8 - 30 ) < Report_EyeScreeningDate < ( doGDM8 + 365 ) then
delete ;
MPRINT(XGDM):  if ( doGDM9 - 30 ) < Report_EyeScreeningDate < ( doGDM9 + 365 ) then
```

```

delete ;
MPRINT(XGDM):  if ( doGDM10 - 30 ) < Report_EyeScreeningDate < ( doGDM10 + 365 ) then
delete ;
MPRINT(XGDM):  if ( doGDM11 - 30 ) < Report_EyeScreeningDate < ( doGDM11 + 365 ) then
delete ;
MPRINT(XGDM):  if ( doGDM12 - 30 ) < Report_EyeScreeningDate < ( doGDM12 + 365 ) then
delete ;
17          run ;

```

NOTE: Variable doGDM12 is uninitialized.

NOTE: Missing values were generated as a result of performing an operation on missing values.

Each place is given by: (Number of times) at (Line):(Column).

```

89780 at 16:18      89780 at 16:54      107772 at 16:20      107772 at 16:56
112844 at 16:20      112844 at 16:56      114221 at 16:20      114221 at 16:56
114499 at 16:20      114499 at 16:56      114577 at 16:20      114577 at 16:56
114616 at 16:20      114616 at 16:56      114625 at 16:20      114625 at 16:56
114626 at 16:20      114626 at 16:56      114626 at 16:20      114626 at 16:56
114626 at 16:20      114626 at 16:56      114627 at 16:20      114627 at 16:56

```

NOTE: There were 98929 observations read from the data set WORK.DIAB.

NOTE: There were 21146 observations read from the data set TTDATA.GDM.

NOTE: The data set WORK.DIAB has 114627 observations and 25 variables.

NOTE: DATA statement used (Total process time):

```

real time          0.35 seconds
cpu time           0.34 seconds

```

```

18
19          data TTdata.diab ( keep = pnr doDIAB )
20              diab ;
21          set diab ;
22          by pnr Report_EyeScreeningDate ;
23          doDiab = Report_EyeScreeningDate ;
24          visit = "later" ;
25          if first.pnr then do ;
26              output TTdata.diab ;
27              visit = "first" ;
28          end ;
29          output diab ;
30          run ;

```

NOTE: There were 114627 observations read from the data set WORK.DIAB.

NOTE: The data set TTDATA.DIAB has 57676 observations and 2 variables.

NOTE: The data set WORK.DIAB has 114627 observations and 27 variables.

NOTE: DATA statement used (Total process time):

```

real time          0.09 seconds
cpu time           0.06 seconds

```

```

31
32          title1 'Distribution of first and later visits' ;
33          proc tabulate data = diab missing noseps ;
34              class doDIAB visit ;
35              table doDIAB,
36                  visit * f=comma9.
37                  / rts = 20 ;
38              format doDIAB yymms7. ;
39          run ;

```

NOTE: There were 114627 observations read from the data set WORK.DIAB.

NOTE: The PROCEDURE TABULATE printed page 2.

NOTE: PROCEDURE TABULATE used (Total process time):

```

real time          0.04 seconds
cpu time           0.07 seconds

```

```

40          title1 ;
41
42          title1 'First visits from Diabasen' ;
43          proc contents data = TTdata.diab ; run ;

```


NOTE: PROCEDURE CONTENTS used (Total process time):

real time	0.00 seconds
cpu time	0.00 seconds

NOTE: The PROCEDURE CONTENTS printed page 3.

NOTE: SAS Institute Inc., SAS Campus Drive, Cary, NC USA 27513-2414

NOTE: The SAS System used:

real time	0.82 seconds
cpu time	0.73 seconds

7.9.1 05-diab.lst

Original Diabase

17:36 Thursday, January 12, 2017 1

The CONTENTS Procedure

Data Set Name	EKST15.DIABASEN	Observations	98929
Member Type	DATA	Variables	13
Engine	V9	Indexes	0
Created	27/11/2015 14:29:35	Observation Length	336
Last Modified	27/11/2015 14:29:35	Deleted Observations	0
Protection		Compressed	NO
Data Set Type		Sorted	YES
Label	Diabasen		
Data Representation	WINDOWS_64		
Encoding	wlatin1 Western (Windows)		

Engine/Host Dependent Information

Data Set Page Size	32768
Number of Data Set Pages	1020
First Data Page	*
Max Obs per Page	97
Obs in First Data Page	88
Number of Data Set Repairs	0
Filename	E:\rawdata\705093\Eksterne data\diabasen.sas7bdat
Release Created	9.0401M3
Host Created	X64_ES08R2

Alphabetic List of Variables and Attributes

#	Variable	Type	Len	Format	Informat	Label
13	BlurredMediaLeftEyeCode	Char	25		\$25.	Slrede medier venstre je
12	BlurredMediaRightEyeCode	Char	25		\$25.	Slrede medier hjre je
*	EyeScreeningIndication Code	Char	10		\$10.	enscreeningsindikation
10	MaculopatiStatus LeftEyeCode	Char	53		\$53.	Maculopati-status venstre je
11	MaculopatiStatus RightEyeCode	Char	53		\$53.	Maculopati-status hjre je
*	PriorEyeSurgery LeftEyeCode	Char	23		\$23.	Tidligere jenkirurgi venstre je
*	PriorEyeSurgery RightEyeCode	Char	23		\$23.	Tidligere jenkirurgi hjre je
*	Report_EyeScreeningDate	Num	8	IS8601DA10.	IS8601DA10.	enscreeningsdato
8	RetinaStatusLeftEyeCode	Char	23		\$23.	Retina status venstre je
9	RetinaStatusRightEyeCode	Char	23		\$23.	Retina status hjre je
6	VisualAcuityLeftEyeCode	Char	28		\$28.	Synsstyrke venstre je
7	VisualAcuityRightEyeCode	Char	28		\$28.	Synsstyrke hjre je
*	pnr	Char	12	\$12.	\$10.	Personnummer

Sort Information

Sortedby pnr
 Validated YES
 Character Set ANSI

Distribution of first and later visits

17:36 Thursday, January 12, 2017 2

	visit	
	first	later
	N	N
doDiab		
.	18,197	.
2009/01	385	.
2009/02	409	.
2009/03	389	*
2009/04	354	*
2009/05	282	7
2009/06	337	19
2009/07	149	15
2009/08	328	47
2009/09	359	64
2009/10	357	53
2009/11	354	51
2009/12	302	54
2010/01	625	121
2010/02	431	159
2010/03	543	287
2010/04	412	262
2010/05	444	255
2010/06	695	342
2010/07	604	219
2010/08	980	434
2010/09	1,246	498
2010/10	1,304	488
2010/11	1,269	538
2010/12	779	449
2011/01	1,251	645
2011/02	1,247	601
2011/03	1,475	798
2011/04	628	548
2011/05	993	748
2011/06	1,062	726
2011/07	454	369
2011/08	724	936
2011/09	877	1,076
2011/10	794	1,058
2011/11	885	1,264
2011/12	554	961
2012/01	752	1,293
2012/02	560	1,001
2012/03	766	1,307
2012/04	619	1,002
2012/05	624	1,113
2012/06	632	1,223
2012/07	295	602
2012/08	439	1,003
2012/09	399	1,149
2012/10	397	1,219
2012/11	357	1,157
2012/12	177	693
2013/01	507	1,252
2013/02	325	966
2013/03	332	929
2013/04	416	1,192
2013/05	331	1,157
2013/06	348	1,099

2013/07	100	412
2013/08	293	910
2013/09	250	1,108
2013/10	267	1,209
2013/11	238	1,252
2013/12	178	778
2014/01	234	1,390
2014/02	393	1,327
2014/03	548	1,501
2014/04	486	1,285
2014/05	425	1,324
2014/06	409	1,303
2014/07	268	589
2014/08	359	1,128
2014/09	509	1,418
2014/10	414	1,451
2014/11	469	1,360
2014/12	384	1,259
2015/01	362	1,359
2015/02	336	1,134

First visits from Diabasen

17:36 Thursday, January 12, 2017 3

The CONTENTS Procedure

Data Set Name	TTDATA.DIAB	Observations	57676
Member Type	DATA	Variables	*
Engine	V9	Indexes	0
Created	12/01/2017 17:36:50	Observation Length	24
Last Modified	12/01/2017 17:36:50	Deleted Observations	0
Protection		Compressed	NO
Data Set Type		Sorted	NO
Label			
Data Representation	WINDOWS_64		
Encoding	wlatin1 Western (Windows)		

Engine/Host Dependent Information

Data Set Page Size	65536
Number of Data Set Pages	22
First Data Page	*
Max Obs per Page	2715
Obs in First Data Page	2664
Number of Data Set Repairs	0
ExtendObsCounter	YES
Filename	E:\workdata\705093\BxC\demoDM\DATA\diab.sas7bdat
Release Created	9.0401M3
Host Created	X64_SRV12

Alphabetic List of Variables and Attributes

#	Variable	Type	Len	Format	Informat	Label
2	doDiab	Num	8			
1	pnr	Char	12	\$12.	\$10.	Personnummer

7.10 06-define

Collects records from the processed registers and defines a diabetes register and the relevant dates in it.

The inclusion date will be the smaller of the earliest dates from NPR, RMPS, DADD and DIAB, and the inclusion criterion will be the the one that triggered the inclusion.

Diabetes type is derived as described above.

Also derives a diabetes register exclusively based on drug information only.

```

1                                     "Program 06-define.sas"

NOTE: Copyright (c) 2002-2012 by SAS Institute Inc., Cary, NC, USA.
NOTE: SAS (r) Proprietary Software 9.4 (TS1M3)
      Licensed to FORSKNING 1, Site 50800722.
NOTE: This session is executing on the X64_SRV12 platform.

NOTE: Updated analytical products:
      SAS/STAT 14.1

NOTE: Additional host information:
      X64_SRV12 WIN 6.2.9200 Server

NOTE: SAS initialization used:
      real time          0.06 seconds
      cpu time           0.07 seconds

NOTE: AUTOEXEC processing beginning; file is E:\workdata\705093\BXC\demoDM\sas\optslibs.sas.

NOTE: AUTOEXEC processing completed.

1      /* get the TTdata.gdm, and exclude all registration in the year after the gdm
1      ! */
2      /* not here but in all other programs generating dates of inclusion */
3
4      data DMreg ;
5          merge TTdata.npr    ( in = npr    keep = pnr doNPR  nprtyp )
6                  TTdata.DVDD ( in = dvdd   keep = pnr doDVDD dvttyp )
7                  TTdata.pRMPS ( in = rmps   keep = pnr doRMPS doOAD doIns )
8                  TTdata.DIAB ( in = diab   keep = pnr doDIAB )
9                  TTdata.pop   ( in = pop   );
10         by pnr ;
11         format doBth doDM doDth doNPR doDVDD doOAD doIns doDIAB ddmmyy10. ;
12         if pop and ( npr or dvdd or rmps or diab ) ;
13         * Date of diagnosis - GDM and PCOS - taken care of in RMPS / DVDD ;
14         doDM = min( doNPR, doDVDD, doOAD, doIns, doDIAB ) ;
15         if doDM > doBth ;
16         if doDM eq doNPR then inCr = "NPR" ;
17         if doDM eq doDVDD then inCr = "DVDD" ;
18         if doDM eq doOAD then inCr = "OAD" ;
19         if doDM eq doIns then inCr = "Ins" ;
20         if doDM eq doDIAB then inCr = "DIAB" ;
21         * Type of diabetes is T1 if known from DVDD or NPR or
22           if any dispensation before age 15 or insulin before 30 ;
23         if ( nprtyp eq 'T1' and dvttyp ne 'T2' ) or
24             dvttyp eq 'T1' or
25             .z < (doOAD - doBth) < ( 365.25 * 15 ) or
26             .z < (doIns - doBth) < ( 365.25 * 30 ) then DMtp = 'T1' ;
27         else DMtp = 'T2' ;
28         * never override DVDD verdict of T2 ;
29         if dvttyp eq 'T2' then DMtp = 'T2' ;
30         * for simplification of tables ;
31         doDMx = max( doDM, '31DEC1994'd ) ;
32     run ;

NOTE: Missing values were generated as a result of performing an operation on missing
      values.
      Each place is given by: (Number of times) at (Line):(Column).
      14385 at 14:10 36528 at 25:18 274757 at 26:18
NOTE: There were 223402 observations read from the data set TTDATA.NPR.
NOTE: There were 88398 observations read from the data set TTDATA.DVDD.
NOTE: There were 409844 observations read from the data set TTDATA.PRMP.
NOTE: There were 57676 observations read from the data set TTDATA.DIAB.

```

NOTE: There were 7361669 observations read from the data set TTDATA.POP.
 NOTE: The data set WORK.DMREG has 428251 observations and 16 variables.
 NOTE: DATA statement used (Total process time):
 real time 3.93 seconds
 cpu time 2.57 seconds

```
33
34     title1 'Overview of typing of DM' ;
35     proc tabulate data = DMreg missing noseps ;
36         class dvdtyp nprtyp DMtp inCr doDMx ;
37         table ( all dmTP ) *
38             ( all nprtyp ),
39             ( all dvdtyp ) * f=comma7.
40         / rts = 15 ;
41         keylabel n = ' ' ;
42         format doDMx year4. ;
43     run ;
```

NOTE: There were 428251 observations read from the data set WORK.DMREG.
 NOTE: The PROCEDURE TABULATE printed page 1.
 NOTE: PROCEDURE TABULATE used (Total process time):
 real time 0.12 seconds
 cpu time 0.17 seconds

```
44
45     title1 'The reconstructed diabetes register' ;
46     data TTdata.DMreg ;
47     set DMreg ;
48     keep pnr sex DMtp inCr /* inklaars */
49     doBth doDM doDth doNPR doDVDD doOAD doIns doDIAB ;
50     label DMtp = 'Type of DM'
51     inCr = 'Inclusion criterion' ;
52     run ;
```

NOTE: There were 428251 observations read from the data set WORK.DMREG.
 NOTE: The data set TTDATA.DMREG has 428251 observations and 12 variables.
 NOTE: DATA statement used (Total process time):
 real time 0.20 seconds
 cpu time 0.09 seconds

```
53     proc contents data = TTdata.DMreg ; run ;
```

NOTE: PROCEDURE CONTENTS used (Total process time):
 real time 0.00 seconds
 cpu time 0.00 seconds

NOTE: The PROCEDURE CONTENTS printed page 2.

```
54
55     * add temporary variables for the tabulation ;
56     data a ;
57     set TTdata.DMreg ;
58     a1 = floor( ( doDM - doBth ) / 365.25 ) ;
59     a5 = floor( ( doDM - doBth ) / (365.25*5) ) * 5 ;
60     doDM = max( doDM, '31DEC1994'd ) ;
61     run ;
```

NOTE: There were 428251 observations read from the data set TTDATA.DMREG.
 NOTE: The data set WORK.A has 428251 observations and 14 variables.
 NOTE: DATA statement used (Total process time):
 real time 0.25 seconds
 cpu time 0.07 seconds

```
62
63     proc tabulate data = a missing noseps ;
64         class sex doDM DMtp a1 a5 inCr ; * inklaars ;
```

```

65     table all doDM,
66         ( all DMtp ) * f = comma8.
67         all * pctn< DMtp all > * f = 5.1
68         DMtp * pctn< DMtp all > * f = 4.1
69     / rts = 9 condense ;
70     table all doDM,
71         ( all inCr ) * f = comma7.
72         all * pctn< inCr all > * f = 5.1
73         inCr * pctn< inCr all > * f = 4.1
74     / rts = 9 condense ;
75     table all a1,
76         sex * (
77             all * f = comma7.
78             all * pctn< inCr all > * f = 5.1
79             inCr * pctn< inCr all > * f = 4.1 )
80     / rts = 9 condense ;
81     table sex, all a1 = "DM age",
82         ( all inCr ) * f = comma7.
83         all * pctn< inCr all > * f = 5.1
84         inCr * pctn< inCr all > * f = 4.1
85     / rts = 9 condense ;
86     table all DMtp,
87         all a5,
88         all * f = comma7.
89         doDM * f = comma6.
90     / rts = 6 condense ;
91     format doDM year4.
92         sex koen_t. ;
93     run ;

```

NOTE: There were 428251 observations read from the data set WORK.A.

NOTE: At least one W.D format was too small for the number to be printed. The decimal may be shifted by the "BEST" format.

NOTE: The PROCEDURE TABULATE printed pages 3-7.

NOTE: PROCEDURE TABULATE used (Total process time):

```

real time      0.15 seconds
cpu time       0.31 seconds

```

```

94
95     title2 'Diagnoses 2012 ff. only' ;
96     proc tabulate data = TTdata.DMreg missing noseps ;
97         where doDM > '31DEC2011'd ;
98         class sex doDM inCr DMtp ;
99         table all doDM,
100             ( all DMtp ) * f = comma8.
101             all * pctn< DMtp all > * f = 5.1
102             DMtp * pctn< DMtp all > * f = 4.1
103         / rts = 9 condense ;
104         table all doDM,
105             ( all inCr ) * f = comma7.
106             all * pctn< inCr all > * f = 5.1
107             inCr * pctn< inCr all > * f = 4.1
108         / rts = 9 condense ;
109         format doDM yymms7.
110             sex koen_t. ;
111     run ;

```

NOTE: There were 73169 observations read from the data set TTDATA.DMREG.
WHERE doDM>'31DEC2011'D;

NOTE: The PROCEDURE TABULATE printed pages 8-9.

NOTE: PROCEDURE TABULATE used (Total process time):

```

real time      0.04 seconds
cpu time       0.10 seconds

```

```

112     title1 ;
113
114     title1 'The reconstructed diabetes *drug* register' ;
115     data TTdata.DMreg ;

```

```

116      merge TTdata.pRMPS ( in = rmps  keep = pnr doOAD doIns )
117          TTdata.pop    ( in = pop  ) ;
118      by pnr ;
119      keep pnr sex DMtp inCr
120          doBth doDM doDth doOAD doIns ;
121      format doBth doDM doDth doOAD doIns  ddmmyy10. ;
122      if pop and rmps ;
123      * Date of diagnosis - GDM and PCOS - taken care of in RMPS / DVDD ;
124      doDM = min( doOAD, doIns ) ;
125      if doDM > doBth ;
126      * Type of diabetes is T1 if known from DVDD or
127        if any dispensation before age 15 or insulin before 30 ;
128      if .z < doOAD - doBth < ( 365.25 * 15 ) or
129        .z < doIns - doBth < ( 365.25 * 30 ) then DMtp = 'T1' ;
130      if DMtp ne 'T1' then DMtp = 'T2' ;
131      if doDM eq doOAD then inCr = "OAD " ;
132      if doDM eq doIns then inCr = "Ins " ;
133      run ;

```

NOTE: Missing values were generated as a result of performing an operation on missing values.

Each place is given by: (Number of times) at (Line):(Column).
52995 at 128:17 260453 at 129:17

NOTE: There were 409844 observations read from the data set TTDATA.PRMPS.

NOTE: There were 7361669 observations read from the data set TTDATA.POP.

NOTE: The data set TTDATA.DMDREG has 409841 observations and 9 variables.

NOTE: DATA statement used (Total process time):

real time	1.45 seconds
cpu time	1.31 seconds

```

134
135      proc contents data = TTdata.Dmdreg ; run ;

```

NOTE: PROCEDURE CONTENTS used (Total process time):

real time	0.01 seconds
cpu time	0.01 seconds

NOTE: The PROCEDURE CONTENTS printed page 10.

```

136      proc tabulate data = TTdata.Dmdreg missing noseps ;
137          class sex doDM DMtp inCr ;
138          table all doDM,
139              ( all DMtp inCr ) * f = comma7.
140              all * pctn< all > * f = 6.1
141              ( DMtp * pctn< DMtp >
142              inCr * pctn< inCr > ) * f = 4.1
143              / rts = 6 ;
144          keylabel n = ' ' ;
145          format doDM year4.
146              sex koen_t. ;
147      run ;

```

NOTE: There were 409841 observations read from the data set TTDATA.DMDREG.

NOTE: The PROCEDURE TABULATE printed page 11.

NOTE: PROCEDURE TABULATE used (Total process time):

real time	0.17 seconds
cpu time	0.23 seconds

NOTE: SAS Institute Inc., SAS Campus Drive, Cary, NC USA 27513-2414

NOTE: The SAS System used:

real time	6.51 seconds
cpu time	4.99 seconds

7.10.1 06-define.lst

		dvdtyp				
		All	NA	T1	T2	
All	All	428,251	339,982	3,552	23,886	60,831
	nprtyp					
		204,986	204,207	66	25	688
	NA	37,821	25,998	1,174	2,687	7,962
	T1	44,479	21,086	702	19,833	2,858
	T2	140,965	88,691	1,610	1,341	49,323
DMtp	All	47,339	22,602	851	23,886	.
T1	nprtyp					
		665	638	*	25	.
	NA	3,434	638	109	2,687	.
	T1	41,621	21,086	702	19,833	.
	T2	1,619	240	38	1,341	.
T2	All	380,912	317,380	2,701	.	60,831
	nprtyp					
		204,321	203,569	64	.	688
	NA	34,387	25,360	1,065	.	7,962
	T1	2,858	.	.	.	2,858
	T2	139,346	88,451	1,572	.	49,323

The reconstructed diabetes register

14:29 Tuesday, January 24, 2017 2

The CONTENTS Procedure

Data Set Name	TTDATA.DMREG	Observations	428251
Member Type	DATA	Variables	12
Engine	V9	Indexes	0
Created	24/01/2017 14:29:14	Observation Length	96
Last Modified	24/01/2017 14:29:14	Deleted Observations	0
Protection		Compressed	NO
Data Set Type		Sorted	NO
Label			
Data Representation	WINDOWS_64		
Encoding	wlatin1 Western (Windows)		

Engine/Host Dependent Information

Data Set Page Size	65536
Number of Data Set Pages	629
First Data Page	*
Max Obs per Page	681
Obs in First Data Page	658
Number of Data Set Repairs	0
ExtendObsCounter	YES
Filename	E:\workdata\705093\BxC\demoDM\DATA\dmreg.sas7bdat
Release Created	9.0401M3
Host Created	X64_SRV12

Alphabetic List of Variables and Attributes

#	Variable	Type	Len	Format	Informat	Label
12	DMtp	Char	*			Type of DM
8	doBth	Num	8	DDMMYY10.		
10	doDM	Num	8	DDMMYY10.		
*	doDVDD	Num	8	DDMMYY10.		
6	doDiab	Num	8	DDMMYY10.		
9	doDth	Num	8	DDMMYY10.		
*	doIns	Num	8	DDMMYY10.		
*	doNPR	Num	8	DDMMYY10.	DATE9.	
*	doOAD	Num	8	DDMMYY10.		
11	inCr	Char	*			Inclusion criterion

* pnr Char 12 \$12. \$10. Personnummer
 7 sex Num 8

The reconstructed diabetes register

14:29 Tuesday, January 24, 2017 3

	Type of DM			Type of DM		
	All	T1	T2	All	T1	T2
	N	N	N	PctN	PctN	PctN
All	428,251	47,339	380,912	100.0	11.1	88.9
doDM						
1994	56,578	21,640	34,938	100.0	38.2	61.8
1995	38,055	3,151	34,904	100.0	8.3	91.7
1996	12,117	1,323	10,794	100.0	10.9	89.1
1997	11,589	1,306	10,283	100.0	11.3	88.7
1998	12,883	1,335	11,548	100.0	10.4	89.6
1999	12,788	1,183	11,605	100.0	9.3	90.7
2000	13,784	1,243	12,541	100.0	9.0	91.0
2001	13,685	1,260	12,425	100.0	9.2	90.8
2002	13,882	1,181	12,701	100.0	8.5	91.5
2003	15,817	1,129	14,688	100.0	7.1	92.9
2004	16,286	1,154	15,132	100.0	7.1	92.9
2005	16,124	1,094	15,030	100.0	6.8	93.2
2006	16,719	1,170	15,549	100.0	7.0	93.0
2007	17,861	1,126	16,735	100.0	6.3	93.7
2008	19,553	1,167	18,386	100.0	6.0	94.0
2009	20,034	1,097	18,937	100.0	5.5	94.5
2010	21,853	1,083	20,770	100.0	5.0	95.0
2011	25,474	1,015	24,459	100.0	4.0	96.0
2012	22,346	936	21,410	100.0	4.2	95.8
2013	17,049	935	16,114	100.0	5.5	94.5
2014	16,012	927	15,085	100.0	5.8	94.2
2015	17,762	884	16,878	100.0	5.0	95.0

The reconstructed diabetes register

14:29 Tuesday, January 24, 2017 4

	Inclusion criterion						Inclusion criterion					
	All	DIA	DVD	Ins	NPR	OAD	All	DIA	DVD	Ins	NPR	OAD
	N	N	N	N	N	N	PctN	PctN	PctN	PctN	PctN	PctN
All	428,251	509	69,363	12,080	84,063	262,236	100.0	0.1	16.2	2.8	19.6	61.2
doDM												
1994	56,578	.	23,636	.	32,942	.	100.0	.	41.8	.	58.2	.
1995	38,055	.	2,358	3,493	2,561	29,643	100.0	.	6.2	9.2	6.7	77.9
1996	12,117	.	1,995	478	2,389	7,255	100.0	.	16.5	3.9	19.7	59.9
1997	11,589	.	2,102	364	2,373	6,750	100.0	.	18.1	3.1	20.5	58.2
1998	12,883	.	2,535	387	2,643	7,318	100.0	.	19.7	3.0	20.5	56.8
1999	12,788	.	2,265	394	2,638	7,491	100.0	.	17.7	3.1	20.6	58.6
2000	13,784	.	3,086	396	2,714	7,588	100.0	.	22.4	2.9	19.7	55.0
2001	13,685	.	2,541	358	2,667	8,119	100.0	.	18.6	2.6	19.5	59.3
2002	13,882	.	2,573	345	2,851	8,113	100.0	.	18.5	2.5	20.5	58.4
2003	15,817	.	2,587	361	3,116	9,753	100.0	.	16.4	2.3	19.7	61.7
2004	16,286	.	2,671	384	2,826	10,405	100.0	.	16.4	2.4	17.4	63.9
2005	16,124	.	2,856	389	2,572	10,307	100.0	.	17.7	2.4	16.0	63.9
2006	16,719	.	2,858	397	2,523	10,941	100.0	.	17.1	2.4	15.1	65.4
2007	17,861	.	2,724	425	2,544	12,168	100.0	.	15.3	2.4	14.2	68.1
2008	19,553	.	2,623	477	2,366	14,087	100.0	.	13.4	2.4	12.1	72.0
2009	20,034	6	2,306	473	2,246	15,003	100.0	0.0	11.5	2.4	11.2	74.9
2010	21,853	48	2,265	431	2,005	17,104	100.0	0.2	10.4	2.0	9.2	78.3
2011	25,474	266	1,936	419	1,924	20,929	100.0	1.0	7.6	1.6	7.6	82.2
2012	22,346	80	1,522	440	1,791	18,513	100.0	0.4	6.8	2.0	8.0	82.8
2013	17,049	37	1,167	544	1,809	13,492	100.0	0.2	6.8	3.2	10.6	79.1

2014	16,012	63	733	539	2,074	12,603	100.0	0.4	4.6	3.4	13.0	78.7
2015	17,762	9	24	586	2,489	14,654	100.0	0.1	0.1	3.3	14.0	82.5

The reconstructed diabetes register

14:29 Tuesday, January 24, 2017 5

	sex													
	Mand							Kvinde						
	Inclusion criterion													
	All	All	DIA	DVD	Ins	NPR	OAD	All	All	DIA	DVD	Ins	NPR	OAD
N	PctN	PctN	PctN	PctN	PctN	PctN	N	PctN	PctN	PctN	PctN	PctN	PctN	
All	236,401	100.0	0.1	17.3	2.5	19.5	60.5	191,850	100.0	0.1	14.8	3.2	19.7	62.1
a1														
0	124	100.0	.	51.6	4.0	42.7	1.6	126	100.0	.	60.3	2.4	34.1	3.2
*	280	100.0	.	50.7	1.1	47.1	1.1	235	100.0	.	47.2	2.6	48.9	1.3
*	298	100.0	.	49.7	2.0	47.3	1.0	226	100.0	.	53.5	0.4	45.1	0.9
*	303	100.0	.	51.5	0.7	46.9	1.0	291	100.0	.	50.5	1.4	47.4	0.7
*	319	100.0	.	52.0	1.6	45.8	0.6	297	100.0	.	50.8	0.7	47.8	0.7
*	334	100.0	.	55.4	0.9	42.8	0.9	368	100.0	.	56.0	0.5	43.5	.
6	387	100.0	.	54.5	0.8	44.2	0.5	367	100.0	.	61.0	0.8	37.9	0.3
7	366	100.0	.	54.4	0.8	44.8	.	374	100.0	.	60.2	0.3	39.3	0.3
8	384	100.0	.	53.4	0.5	45.6	0.5	435	100.0	.	62.8	0.5	35.9	0.9
9	479	100.0	.	58.7	0.6	40.3	0.4	507	100.0	.	60.7	0.6	38.3	0.4
10	450	100.0	.	60.0	0.9	39.1	.	526	100.0	.	62.5	0.8	35.6	1.1
11	563	100.0	.	65.2	1.1	33.2	0.5	585	100.0	.	59.8	1.4	37.4	1.4
12	649	100.0	.	60.4	0.6	37.8	1.2	559	100.0	.	62.8	0.7	35.1	1.4
13	635	100.0	.	61.7	1.3	35.7	1.3	453	100.0	.	60.0	1.5	34.7	3.8
14	604	100.0	.	61.8	0.7	35.6	2.0	409	100.0	.	62.3	1.0	31.5	5.1
15	519	100.0	.	64.0	0.8	33.9	1.3	437	100.0	.	54.9	2.7	27.2	15.1
16	475	100.0	.	64.8	2.1	30.7	2.3	438	100.0	.	56.4	1.8	20.8	21.0
17	483	100.0	.	72.9	1.9	23.0	2.3	522	100.0	.	47.7	1.5	21.6	29.1
18	488	100.0	.	63.9	1.8	30.3	3.9	555	100.0	.	41.6	1.6	17.7	39.1
19	518	100.0	.	66.8	2.3	26.3	4.6	598	100.0	.	37.3	1.5	18.6	42.6
20	449	100.0	.	62.4	3.3	32.1	2.2	436	100.0	.	55.3	4.8	28.2	11.7
21	509	100.0	.	67.0	3.9	25.9	3.1	472	100.0	0.4	49.6	3.4	31.6	15.0
22	575	100.0	.	63.8	3.1	29.6	3.5	522	100.0	0.2	48.5	4.0	30.7	16.7
23	542	100.0	.	64.2	2.2	29.0	4.6	567	100.0	.	51.3	4.1	30.3	14.3
24	633	100.0	.	64.3	2.8	29.2	3.6	607	100.0	.	46.5	4.6	29.2	19.8
25	594	100.0	.	58.4	3.9	32.7	5.1	644	100.0	.	43.5	4.2	30.6	21.7
26	722	100.0	.	60.2	3.7	29.4	6.6	703	100.0	0.1	41.8	6.8	33.1	18.1
27	653	100.0	0.2	58.3	2.8	28.3	10.4	705	100.0	0.3	42.4	4.5	27.0	25.8
28	798	100.0	.	53.9	4.0	32.6	9.5	744	100.0	0.1	40.6	6.3	29.6	23.4
29	838	100.0	0.1	55.3	4.3	28.4	11.9	777	100.0	0.3	41.7	6.3	29.1	22.7
30	862	100.0	0.1	52.7	3.7	29.9	13.6	826	100.0	0.2	39.5	7.4	29.5	23.4
31	923	100.0	0.2	49.6	3.0	31.4	15.7	844	100.0	0.4	40.2	8.3	28.6	22.6
32	1,039	100.0	0.1	48.4	3.6	31.9	16.1	880	100.0	0.2	39.2	9.0	29.4	22.2
33	1,149	100.0	0.1	49.1	3.1	31.4	16.3	879	100.0	.	37.7	9.9	30.3	22.2
34	1,177	100.0	0.1	46.9	3.7	32.0	17.3	974	100.0	0.3	39.0	10.0	28.6	22.1
35	1,302	100.0	0.2	45.1	3.1	31.3	20.4	981	100.0	0.1	36.8	9.5	30.3	23.3
36	1,451	100.0	0.1	43.7	3.6	33.8	18.8	994	100.0	0.2	40.2	9.1	29.1	21.4
37	1,524	100.0	0.1	45.0	2.9	32.8	19.2	1,077	100.0	0.2	38.0	9.5	28.6	23.8
38	1,593	100.0	0.1	44.6	3.3	33.7	18.3	1,146	100.0	0.3	40.8	9.2	29.4	20.2
39	1,819	100.0	.	47.2	3.1	33.2	16.5	1,270	100.0	0.4	44.7	7.8	29.4	17.7
40	3,216	100.0	0.0	25.4	1.6	17.0	56.0	3,457	100.0	0.1	14.4	3.5	9.1	72.9
41	2,656	100.0	0.1	32.3	3.0	23.1	41.5	2,035	100.0	0.1	24.9	5.3	15.0	54.7
42	2,725	100.0	0.1	31.0	2.5	22.6	43.9	2,039	100.0	.	25.6	5.0	16.2	53.3
43	3,044	100.0	0.0	30.8	2.2	21.6	45.4	2,006	100.0	0.1	26.3	3.5	18.5	51.6
44	3,264	100.0	.	29.8	2.4	22.1	45.7	2,083	100.0	0.0	26.2	3.6	18.7	51.4
45	3,436	100.0	0.1	27.6	2.2	22.3	47.8	2,244	100.0	0.1	26.0	3.6	17.9	52.5
46	3,723	100.0	0.1	26.3	2.2	21.2	50.3	2,346	100.0	0.1	25.5	3.7	17.6	53.1
47	4,008	100.0	0.2	25.8	2.1	20.3	51.6	2,380	100.0	.	25.0	3.4	17.1	54.5
48	4,201	100.0	0.1	24.5	1.9	20.4	53.2	2,611	100.0	0.1	23.9	3.3	17.8	55.0
49	4,522	100.0	0.1	25.0	2.6	19.3	53.1	2,752	100.0	.	23.2	3.1	17.4	56.3
50	4,789	100.0	0.0	23.6	1.8	18.5	56.1	2,952	100.0	0.2	20.4	2.8	16.4	60.3

51	4,999	100.0	0.0	22.0	1.7	19.1	57.2	3,143	100.0	0.1	21.4	2.7	17.9	57.9
52	5,053	100.0	.	19.6	1.7	19.4	59.3	3,307	100.0	0.1	20.5	2.0	18.1	59.4
53	5,137	100.0	0.1	18.8	2.2	18.1	60.8	3,450	100.0	0.1	18.8	2.4	18.8	59.9
54	5,517	100.0	0.1	17.7	2.1	18.9	61.2	3,417	100.0	0.2	16.9	1.8	18.9	62.2
55	5,616	100.0	0.1	17.2	1.8	19.6	61.3	3,590	100.0	0.1	16.4	1.9	19.5	62.1
56	5,727	100.0	0.1	16.2	1.9	18.1	63.8	3,770	100.0	0.1	16.1	2.0	19.6	62.3
57	6,042	100.0	0.1	15.2	1.9	17.7	65.2	3,802	100.0	0.2	14.4	1.7	20.6	63.1
58	6,058	100.0	0.1	14.0	1.8	18.5	65.5	3,981	100.0	0.1	14.8	1.7	18.4	65.0
59	6,097	100.0	0.2	13.9	2.0	18.1	65.8	4,067	100.0	0.1	15.4	1.8	18.5	64.2
60	6,370	100.0	0.2	12.7	2.0	16.8	68.4	4,234	100.0	0.1	12.2	2.0	19.4	66.2
61	6,547	100.0	0.2	10.6	2.0	17.1	70.2	4,276	100.0	0.1	12.2	2.2	18.7	66.7
62	6,400	100.0	0.1	10.3	2.0	16.6	70.9	4,324	100.0	0.2	10.2	2.1	19.3	68.2
63	6,342	100.0	0.2	9.1	2.3	16.7	71.7	4,465	100.0	0.2	10.2	2.2	19.4	68.0
64	6,450	100.0	0.2	8.6	2.2	16.7	72.4	4,610	100.0	0.1	8.9	2.3	18.0	70.7
65	6,170	100.0	0.1	7.6	2.5	16.3	73.5	4,622	100.0	0.0	7.7	2.6	18.5	71.2
66	5,983	100.0	0.1	7.2	2.2	16.7	73.8	4,582	100.0	0.1	7.5	2.7	19.1	70.5
67	5,869	100.0	0.2	6.5	2.3	16.0	75.0	4,549	100.0	0.3	7.2	2.4	18.7	71.4
68	5,662	100.0	0.2	6.1	2.8	17.0	74.0	4,582	100.0	0.2	5.5	2.3	20.3	71.8
69	5,614	100.0	0.2	5.1	2.9	16.1	75.8	4,367	100.0	0.1	5.5	2.5	19.9	72.0
70	5,189	100.0	0.2	5.1	2.5	16.3	75.9	4,375	100.0	0.2	5.1	2.5	19.3	72.9
71	4,941	100.0	0.2	4.4	2.8	15.8	76.7	4,299	100.0	0.1	3.8	2.9	19.3	73.9
72	4,669	100.0	0.1	4.1	2.9	17.8	75.0	4,315	100.0	0.1	4.3	3.0	17.8	74.8
73	4,645	100.0	0.2	3.6	3.1	15.9	77.2	4,353	100.0	0.2	3.0	3.1	18.6	75.2
74	4,273	100.0	0.3	3.1	3.1	16.2	77.3	4,151	100.0	0.1	3.4	2.9	17.8	75.8
75	3,910	100.0	0.1	3.2	3.2	16.4	77.0	3,902	100.0	0.2	3.0	3.4	18.6	74.7
76	3,885	100.0	0.2	2.6	3.3	17.2	76.8	3,860	100.0	0.1	2.1	2.7	19.4	75.7
77	3,445	100.0	0.1	1.9	2.8	17.3	77.8	3,633	100.0	0.1	2.4	2.7	18.9	75.9
78	3,190	100.0	0.3	1.9	3.2	16.6	78.0	3,479	100.0	0.2	1.5	3.2	18.1	77.0
79	2,924	100.0	0.2	1.9	3.6	15.6	78.6	3,340	100.0	0.0	1.8	3.8	18.6	75.8
80	2,651	100.0	0.1	1.8	3.5	15.5	79.0	3,134	100.0	0.1	1.2	3.8	17.2	77.7
81	2,315	100.0	0.2	1.5	3.5	15.5	79.4	2,899	100.0	0.1	1.1	4.1	16.5	78.1
82	2,068	100.0	0.1	1.9	4.2	14.4	79.4	2,695	100.0	0.1	0.7	3.8	16.2	79.1
83	1,810	100.0	.	1.3	4.5	15.4	78.8	2,440	100.0	0.2	1.1	4.3	17.5	77.0
84	1,501	100.0	0.1	1.1	3.5	15.8	79.5	2,165	100.0	0.1	0.8	4.3	18.2	76.6
85	1,352	100.0	0.1	1.0	4.6	15.4	78.9	1,997	100.0	0.1	0.7	3.8	15.7	79.8
86	1,111	100.0	0.2	0.3	4.3	16.1	79.1	1,708	100.0	0.1	0.6	4.1	17.4	77.7
87	890	100.0	0.1	0.4	4.6	14.3	80.6	1,514	100.0	0.1	0.2	4.2	15.1	80.4
88	701	100.0	0.1	0.7	4.7	15.3	79.2	1,236	100.0	0.2	0.4	4.5	16.5	78.4
89	514	100.0	.	0.2	3.7	14.0	82.1	974	100.0	.	0.5	4.9	17.2	77.3
90	404	100.0	.	.	5.4	14.6	80.0	815	100.0	.	0.5	4.0	14.0	81.5
91	297	100.0	.	0.3	5.1	19.5	75.1	604	100.0	.	.	5.3	13.2	81.5
92	230	100.0	.	0.4	7.0	17.0	75.7	456	100.0	.	0.2	7.7	11.0	81.1
93	138	100.0	.	0.7	3.6	16.7	79.0	364	100.0	.	0.3	6.3	13.7	79.7
94	112	100.0	.	.	5.4	18.8	75.9	267	100.0	0.4	0.4	3.7	13.9	81.6
95	87	100.0	.	.	5.7	11.5	82.8	169	100.0	.	.	11.2	13.0	75.7
96	38	100.0	.	.	5.3	5.3	89.5	126	100.0	.	.	7.1	15.1	77.8
97	29	100.0	.	.	3.4	10.3	86.2	74	100.0	.	.	4.1	24.3	71.6
98	17	100.0	.	.	5.9	5.9	88.2	54	100.0	.	.	3.7	24.1	72.2
99	11	100.0	.	.	18.2	18.2	63.6	24	100.0	.	.	12.5	12.5	75.0
100	*	100.0	100	15	100.0	.	.	6.7	20.0	73.3
101	*	100.0	100	9	100.0	.	.	11.1	11.1	77.8
102	*	100.0	.	.	50.0	.	50.0	*	100.0	100
104	*	100.0	100

The reconstructed diabetes register

14:29 Tuesday, January 24, 2017 6

sex Mand

	Inclusion criterion						Inclusion criterion					
	All	DIA	DVD	Ins	NPR	OAD	All	DIA	DVD	Ins	NPR	OAD
	N	N	N	N	N	N	PctN	PctN	PctN	PctN	PctN	PctN
All	236,401	277	40,999	5,896	46,213	143,016	100.0	0.1	17.3	2.5	19.5	60.5
DM age												
0	124	.	64	*	53	*	100.0	.	51.6	4.0	42.7	1.6
*	280	.	142	*	132	*	100.0	.	50.7	1.1	47.1	1.1
*	298	.	148	6	141	*	100.0	.	49.7	2.0	47.3	1.0

*	303	.	156	*	142	* 100.0	.	51.5	0.7	46.9	1.0
*	319	.	166	*	146	* 100.0	.	52.0	1.6	45.8	0.6
*	334	.	185	*	143	* 100.0	.	55.4	0.9	42.8	0.9
6	387	.	211	*	171	* 100.0	.	54.5	0.8	44.2	0.5
7	366	.	199	*	164	. 100.0	.	54.4	0.8	44.8	.
8	384	.	205	*	175	* 100.0	.	53.4	0.5	45.6	0.5
9	479	.	281	*	193	* 100.0	.	58.7	0.6	40.3	0.4
10	450	.	270	*	176	. 100.0	.	60.0	0.9	39.1	.
11	563	.	367	6	187	* 100.0	.	65.2	1.1	33.2	0.5
12	649	.	392	*	245	8 100.0	.	60.4	0.6	37.8	1.2
13	635	.	392	8	227	8 100.0	.	61.7	1.3	35.7	1.3
14	604	.	373	*	215	12 100.0	.	61.8	0.7	35.6	2.0
15	519	.	332	*	176	7 100.0	.	64.0	0.8	33.9	1.3
16	475	.	308	10	146	11 100.0	.	64.8	2.1	30.7	2.3
17	483	.	352	9	111	11 100.0	.	72.9	1.9	23.0	2.3
18	488	.	312	9	148	19 100.0	.	63.9	1.8	30.3	3.9
19	518	.	346	12	136	24 100.0	.	66.8	2.3	26.3	4.6
20	449	.	280	15	144	10 100.0	.	62.4	3.3	32.1	2.2
21	509	.	341	20	132	16 100.0	.	67.0	3.9	25.9	3.1
22	575	.	367	18	170	20 100.0	.	63.8	3.1	29.6	3.5
23	542	.	348	12	157	25 100.0	.	64.2	2.2	29.0	4.6
24	633	.	407	18	185	23 100.0	.	64.3	2.8	29.2	3.6
25	594	.	347	23	194	30 100.0	.	58.4	3.9	32.7	5.1
26	722	.	435	27	212	48 100.0	.	60.2	3.7	29.4	6.6
27	653	*	381	18	185	68 100.0	0.2	58.3	2.8	28.3	10.4
28	798	.	430	32	260	76 100.0	.	53.9	4.0	32.6	9.5
29	838	*	463	36	238	100 100.0	0.1	55.3	4.3	28.4	11.9
30	862	*	454	32	258	117 100.0	0.1	52.7	3.7	29.9	13.6
31	923	*	458	28	290	145 100.0	0.2	49.6	3.0	31.4	15.7
32	1,039	*	503	37	331	167 100.0	0.1	48.4	3.6	31.9	16.1
33	1,149	*	564	36	361	187 100.0	0.1	49.1	3.1	31.4	16.3
34	1,177	*	552	43	377	204 100.0	0.1	46.9	3.7	32.0	17.3
35	1,302	*	587	40	408	265 100.0	0.2	45.1	3.1	31.3	20.4
36	1,451	*	634	52	490	273 100.0	0.1	43.7	3.6	33.8	18.8
37	1,524	*	686	44	500	292 100.0	0.1	45.0	2.9	32.8	19.2
38	1,593	*	711	53	537	291 100.0	0.1	44.6	3.3	33.7	18.3
39	1,819	.	859	56	603	301 100.0	.	47.2	3.1	33.2	16.5
40	3,216	*	818	50	547	1,800 100.0	0.0	25.4	1.6	17.0	56.0
41	2,656	*	857	81	613	1,102 100.0	0.1	32.3	3.0	23.1	41.5
42	2,725	*	845	67	616	1,195 100.0	0.1	31.0	2.5	22.6	43.9
43	3,044	*	937	68	657	1,381 100.0	0.0	30.8	2.2	21.6	45.4
44	3,264	.	972	79	721	1,492 100.0	.	29.8	2.4	22.1	45.7
45	3,436	*	950	74	765	1,643 100.0	0.1	27.6	2.2	22.3	47.8
46	3,723	*	978	82	789	1,871 100.0	0.1	26.3	2.2	21.2	50.3
47	4,008	7	1,036	83	815	2,067 100.0	0.2	25.8	2.1	20.3	51.6
48	4,201	*	1,030	80	855	2,233 100.0	0.1	24.5	1.9	20.4	53.2
49	4,522	*	1,129	118	871	2,401 100.0	0.1	25.0	2.6	19.3	53.1
50	4,789	*	1,129	85	887	2,687 100.0	0.0	23.6	1.8	18.5	56.1
51	4,999	*	1,100	83	957	2,857 100.0	0.0	22.0	1.7	19.1	57.2
52	5,053	.	989	86	980	2,998 100.0	.	19.6	1.7	19.4	59.3
53	5,137	*	968	115	928	3,123 100.0	0.1	18.8	2.2	18.1	60.8
54	5,517	*	975	118	1,045	3,374 100.0	0.1	17.7	2.1	18.9	61.2
55	5,616	7	966	99	1,101	3,443 100.0	0.1	17.2	1.8	19.6	61.3
56	5,727	6	927	107	1,036	3,651 100.0	0.1	16.2	1.9	18.1	63.8
57	6,042	6	917	114	1,067	3,938 100.0	0.1	15.2	1.9	17.7	65.2
58	6,058	8	849	111	1,121	3,969 100.0	0.1	14.0	1.8	18.5	65.5
59	6,097	10	847	120	1,106	4,014 100.0	0.2	13.9	2.0	18.1	65.8
60	6,370	11	806	125	1,073	4,355 100.0	0.2	12.7	2.0	16.8	68.4
61	6,547	10	692	132	1,117	4,596 100.0	0.2	10.6	2.0	17.1	70.2
62	6,400	7	662	130	1,062	4,539 100.0	0.1	10.3	2.0	16.6	70.9
63	6,342	11	578	146	1,059	4,548 100.0	0.2	9.1	2.3	16.7	71.7
64	6,450	10	553	143	1,074	4,670 100.0	0.2	8.6	2.2	16.7	72.4
65	6,170	9	467	153	1,004	4,537 100.0	0.1	7.6	2.5	16.3	73.5
66	5,983	7	428	133	1,002	4,413 100.0	0.1	7.2	2.2	16.7	73.8
67	5,869	11	380	137	938	4,403 100.0	0.2	6.5	2.3	16.0	75.0
68	5,662	10	344	156	962	4,190 100.0	0.2	6.1	2.8	17.0	74.0
69	5,614	11	286	160	902	4,255 100.0	0.2	5.1	2.9	16.1	75.8
70	5,189	8	264	132	848	3,937 100.0	0.2	5.1	2.5	16.3	75.9
71	4,941	9	219	140	783	3,790 100.0	0.2	4.4	2.8	15.8	76.7
72	4,669	*	193	136	832	3,504 100.0	0.1	4.1	2.9	17.8	75.0

73	4,645	8	169	144	740	3,584	100.0	0.2	3.6	3.1	15.9	77.2
74	4,273	13	133	131	691	3,305	100.0	0.3	3.1	3.1	16.2	77.3
75	3,910	*	125	125	643	3,012	100.0	0.1	3.2	3.2	16.4	77.0
76	3,885	6	100	128	667	2,984	100.0	0.2	2.6	3.3	17.2	76.8
77	3,445	*	64	98	597	2,681	100.0	0.1	1.9	2.8	17.3	77.8
78	3,190	8	62	103	530	2,487	100.0	0.3	1.9	3.2	16.6	78.0
79	2,924	6	57	106	457	2,298	100.0	0.2	1.9	3.6	15.6	78.6
80	2,651	*	48	94	412	2,094	100.0	0.1	1.8	3.5	15.5	79.0
81	2,315	*	34	80	359	1,838	100.0	0.2	1.5	3.5	15.5	79.4
82	2,068	*	40	86	298	1,641	100.0	0.1	1.9	4.2	14.4	79.4
83	1,810	.	23	82	279	1,426	100.0	.	1.3	4.5	15.4	78.8
84	1,501	*	16	52	237	1,194	100.0	0.1	1.1	3.5	15.8	79.5
85	1,352	*	13	62	208	1,067	100.0	0.1	1.0	4.6	15.4	78.9
86	1,111	*	*	48	179	879	100.0	0.2	0.3	4.3	16.1	79.1
87	890	*	*	41	127	717	100.0	0.1	0.4	4.6	14.3	80.6
88	701	*	*	33	107	555	100.0	0.1	0.7	4.7	15.3	79.2
89	514	.	*	19	72	422	100.0	.	0.2	3.7	14.0	82.1
90	404	.	.	22	59	323	100.0	.	.	5.4	14.6	80.0
91	297	.	*	15	58	223	100.0	.	0.3	5.1	19.5	75.1
92	230	.	*	16	39	174	100.0	.	0.4	7.0	17.0	75.7
93	138	.	*	*	23	109	100.0	.	0.7	3.6	16.7	79.0
94	112	.	.	6	21	85	100.0	.	.	5.4	18.8	75.9
95	87	.	.	*	10	72	100.0	.	.	5.7	11.5	82.8
96	38	.	.	*	*	34	100.0	.	.	5.3	5.3	89.5
97	29	.	.	*	*	25	100.0	.	.	3.4	10.3	86.2
98	17	.	.	*	*	15	100.0	.	.	5.9	5.9	88.2
99	11	.	.	*	*	7	100.0	.	.	18.2	18.2	63.6
100	*	*	100.0	100
101	*	*	100.0	100
102	*	.	.	*	.	*	100.0	.	.	50.0	.	50.0

sex Kvinde

	Inclusion criterion						Inclusion criterion					
	All	DIA	DVD	Ins	NPR	OAD	All	DIA	DVD	Ins	NPR	OAD
	N	N	N	N	N	N	PctN	PctN	PctN	PctN	PctN	PctN
All	191,850	232	28,364	6,184	37,850	119,220	100.0	0.1	14.8	3.2	19.7	62.1
DM age												
0	126	.	76	*	43	*	100.0	.	60.3	2.4	34.1	3.2
*	235	.	111	6	115	*	100.0	.	47.2	2.6	48.9	1.3
*	226	.	121	*	102	*	100.0	.	53.5	0.4	45.1	0.9
*	291	.	147	*	138	*	100.0	.	50.5	1.4	47.4	0.7
*	297	.	151	*	142	*	100.0	.	50.8	0.7	47.8	0.7
*	368	.	206	*	160	.	100.0	.	56.0	0.5	43.5	.
6	367	.	224	*	139	*	100.0	.	61.0	0.8	37.9	0.3
7	374	.	225	*	147	*	100.0	.	60.2	0.3	39.3	0.3
8	435	.	273	*	156	*	100.0	.	62.8	0.5	35.9	0.9
9	507	.	308	*	194	*	100.0	.	60.7	0.6	38.3	0.4
10	526	.	329	*	187	6	100.0	.	62.5	0.8	35.6	1.1
11	585	.	350	8	219	8	100.0	.	59.8	1.4	37.4	1.4
12	559	.	351	*	196	8	100.0	.	62.8	0.7	35.1	1.4
13	453	.	272	7	157	17	100.0	.	60.0	1.5	34.7	3.8
14	409	.	255	*	129	21	100.0	.	62.3	1.0	31.5	5.1
15	437	.	240	12	119	66	100.0	.	54.9	2.7	27.2	15.1
16	438	.	247	8	91	92	100.0	.	56.4	1.8	20.8	21.0
17	522	.	249	8	113	152	100.0	.	47.7	1.5	21.6	29.1
18	555	.	231	9	98	217	100.0	.	41.6	1.6	17.7	39.1
19	598	.	223	9	111	255	100.0	.	37.3	1.5	18.6	42.6
20	436	.	241	21	123	51	100.0	.	55.3	4.8	28.2	11.7
21	472	*	234	16	149	71	100.0	0.4	49.6	3.4	31.6	15.0
22	522	*	253	21	160	87	100.0	0.2	48.5	4.0	30.7	16.7
23	567	.	291	23	172	81	100.0	.	51.3	4.1	30.3	14.3
24	607	.	282	28	177	120	100.0	.	46.5	4.6	29.2	19.8

25	644	.	280	27	197	140	100.0	.	43.5	4.2	30.6	21.7
26	703	*	294	48	233	127	100.0	0.1	41.8	6.8	33.1	18.1
27	705	*	299	32	190	182	100.0	0.3	42.4	4.5	27.0	25.8
28	744	*	302	47	220	174	100.0	0.1	40.6	6.3	29.6	23.4
29	777	*	324	49	226	176	100.0	0.3	41.7	6.3	29.1	22.7
30	826	*	326	61	244	193	100.0	0.2	39.5	7.4	29.5	23.4
31	844	*	339	70	241	191	100.0	0.4	40.2	8.3	28.6	22.6
32	880	*	345	79	259	195	100.0	0.2	39.2	9.0	29.4	22.2
33	879	.	331	87	266	195	100.0	.	37.7	9.9	30.3	22.2
34	974	*	380	97	279	215	100.0	0.3	39.0	10.0	28.6	22.1
35	981	*	361	93	297	229	100.0	0.1	36.8	9.5	30.3	23.3
36	994	*	400	90	289	213	100.0	0.2	40.2	9.1	29.1	21.4
37	1,077	*	409	102	308	256	100.0	0.2	38.0	9.5	28.6	23.8
38	1,146	*	468	106	337	232	100.0	0.3	40.8	9.2	29.4	20.2
39	1,270	*	568	99	373	225	100.0	0.4	44.7	7.8	29.4	17.7
40	3,457	*	497	121	316	2,521	100.0	0.1	14.4	3.5	9.1	72.9
41	2,035	*	506	107	306	1,114	100.0	0.1	24.9	5.3	15.0	54.7
42	2,039	.	521	102	330	1,086	100.0	.	25.6	5.0	16.2	53.3
43	2,006	*	527	70	371	1,035	100.0	0.1	26.3	3.5	18.5	51.6
44	2,083	*	546	76	390	1,070	100.0	0.0	26.2	3.6	18.7	51.4
45	2,244	*	584	80	401	1,177	100.0	0.1	26.0	3.6	17.9	52.5
46	2,346	*	598	87	412	1,246	100.0	0.1	25.5	3.7	17.6	53.1
47	2,380	.	595	81	408	1,296	100.0	.	25.0	3.4	17.1	54.5
48	2,611	*	623	87	464	1,435	100.0	0.1	23.9	3.3	17.8	55.0
49	2,752	.	638	86	478	1,550	100.0	.	23.2	3.1	17.4	56.3
50	2,952	6	601	83	483	1,779	100.0	0.2	20.4	2.8	16.4	60.3
51	3,143	*	673	85	564	1,819	100.0	0.1	21.4	2.7	17.9	57.9
52	3,307	*	679	65	597	1,964	100.0	0.1	20.5	2.0	18.1	59.4
53	3,450	*	649	83	650	2,065	100.0	0.1	18.8	2.4	18.8	59.9
54	3,417	6	579	60	645	2,127	100.0	0.2	16.9	1.8	18.9	62.2
55	3,590	*	587	70	699	2,230	100.0	0.1	16.4	1.9	19.5	62.1
56	3,770	*	607	74	738	2,347	100.0	0.1	16.1	2.0	19.6	62.3
57	3,802	7	546	66	783	2,400	100.0	0.2	14.4	1.7	20.6	63.1
58	3,981	*	588	68	734	2,587	100.0	0.1	14.8	1.7	18.4	65.0
59	4,067	*	626	75	751	2,611	100.0	0.1	15.4	1.8	18.5	64.2
60	4,234	6	517	86	821	2,804	100.0	0.1	12.2	2.0	19.4	66.2
61	4,276	*	523	96	798	2,854	100.0	0.1	12.2	2.2	18.7	66.7
62	4,324	9	440	92	834	2,949	100.0	0.2	10.2	2.1	19.3	68.2
63	4,465	8	455	99	865	3,038	100.0	0.2	10.2	2.2	19.4	68.0
64	4,610	*	410	106	832	3,257	100.0	0.1	8.9	2.3	18.0	70.7
65	4,622	*	355	122	854	3,289	100.0	0.0	7.7	2.6	18.5	71.2
66	4,582	6	344	123	877	3,232	100.0	0.1	7.5	2.7	19.1	70.5
67	4,549	12	326	111	850	3,250	100.0	0.3	7.2	2.4	18.7	71.4
68	4,582	7	251	104	931	3,289	100.0	0.2	5.5	2.3	20.3	71.8
69	4,367	*	242	107	871	3,143	100.0	0.1	5.5	2.5	19.9	72.0
70	4,375	9	223	111	844	3,188	100.0	0.2	5.1	2.5	19.3	72.9
71	4,299	6	163	123	830	3,177	100.0	0.1	3.8	2.9	19.3	73.9
72	4,315	*	184	128	770	3,228	100.0	0.1	4.3	3.0	17.8	74.8
73	4,353	7	129	135	809	3,273	100.0	0.2	3.0	3.1	18.6	75.2
74	4,151	*	141	119	740	3,148	100.0	0.1	3.4	2.9	17.8	75.8
75	3,902	6	119	134	727	2,916	100.0	0.2	3.0	3.4	18.6	74.7
76	3,860	*	82	105	747	2,921	100.0	0.1	2.1	2.7	19.4	75.7
77	3,633	*	86	97	688	2,757	100.0	0.1	2.4	2.7	18.9	75.9
78	3,479	8	51	113	628	2,679	100.0	0.2	1.5	3.2	18.1	77.0
79	3,340	*	59	127	621	2,532	100.0	0.0	1.8	3.8	18.6	75.8
80	3,134	*	37	120	540	2,435	100.0	0.1	1.2	3.8	17.2	77.7
81	2,899	*	33	120	478	2,264	100.0	0.1	1.1	4.1	16.5	78.1
82	2,695	*	20	102	437	2,132	100.0	0.1	0.7	3.8	16.2	79.1
83	2,440	*	26	104	426	1,880	100.0	0.2	1.1	4.3	17.5	77.0
84	2,165	*	18	94	393	1,658	100.0	0.1	0.8	4.3	18.2	76.6
85	1,997	*	13	76	314	1,593	100.0	0.1	0.7	3.8	15.7	79.8
86	1,708	*	11	70	298	1,327	100.0	0.1	0.6	4.1	17.4	77.7
87	1,514	*	*	63	229	1,217	100.0	0.1	0.2	4.2	15.1	80.4
88	1,236	*	*	56	204	969	100.0	0.2	0.4	4.5	16.5	78.4
89	974	.	*	48	168	753	100.0	.	0.5	4.9	17.2	77.3
90	815	.	*	33	114	664	100.0	.	0.5	4.0	14.0	81.5
91	604	.	.	32	80	492	100.0	.	.	5.3	13.2	81.5
92	456	.	*	35	50	370	100.0	.	0.2	7.7	11.0	81.1
93	364	.	*	23	50	290	100.0	.	0.3	6.3	13.7	79.7
94	267	*	*	10	37	218	100.0	0.4	0.4	3.7	13.9	81.6

95	169	.	.	19	22	128	100.0	.	.	11.2	13.0	75.7
96	126	.	.	9	19	98	100.0	.	.	7.1	15.1	77.8
97	74	.	.	*	18	53	100.0	.	.	4.1	24.3	71.6
98	54	.	.	*	13	39	100.0	.	.	3.7	24.1	72.2
99	24	.	.	*	*	18	100.0	.	.	12.5	12.5	75.0
100	15	.	.	*	*	11	100.0	.	.	6.7	20.0	73.3
101	9	.	.	*	*	7	100.0	.	.	11.1	11.1	77.8
102	*	*	100.0	100
104	*	*	100.0	100

The reconstructed diabetes register

14:29 Tuesday, January 24, 2017 7

All

doDM											
All	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	
N	N	N	N	N	N	N	N	N	N	N	N
All	428,251	56,578	38,055	12,117	11,589	12,883	12,788	13,784	13,685	13,882	15,817
a5											
0	2,499	1,322	60	58	62	49	67	59	57	47	56
*	4,001	2,012	71	55	80	74	68	98	100	100	95
10	5,433	2,750	81	81	86	82	103	95	126	106	101
15	5,033	2,253	78	98	93	91	76	81	85	106	91
20	5,312	2,511	145	120	114	121	133	103	134	125	132
25	7,178	2,857	255	189	183	203	176	212	189	188	185
30	9,553	3,168	540	300	279	319	303	311	306	264	322
35	13,157	3,717	831	412	405	425	419	537	494	485	529
40	26,525	4,483	1,144	588	567	657	676	810	727	683	861
45	32,223	5,089	1,801	913	820	946	875	1,046	974	1,059	1,164
50	41,764	5,049	2,459	1,312	1,289	1,503	1,444	1,595	1,446	1,434	1,552
55	48,750	4,894	2,980	1,200	1,262	1,449	1,551	1,688	1,848	1,947	2,181
60	54,018	4,475	4,067	1,367	1,310	1,455	1,499	1,660	1,626	1,768	2,129
65	52,000	4,096	5,127	1,418	1,275	1,492	1,459	1,502	1,527	1,605	1,856
70	45,210	3,505	6,017	1,421	1,300	1,406	1,341	1,419	1,431	1,425	1,623
75	35,568	2,517	5,461	1,213	1,215	1,235	1,193	1,191	1,153	1,116	1,356
80	23,678	1,299	4,119	855	741	800	834	786	835	848	926
85	11,997	478	2,101	407	384	439	420	437	464	411	465
90	3,687	95	624	89	106	120	133	129	140	139	161
95	629	8	87	21	17	16	18	24	22	23	29
100	36	.	7	.	*	*	.	*	*	*	*

(Continued)

doDM												
	2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015
	N	N	N	N	N	N	N	N	N	N	N	N
All	16,286	16,124	16,719	17,861	19,553	20,034	21,853	25,474	22,346	17,049	16,012	17,762
a5												
0	42	58	54	49	64	64	49	62	49	64	50	57
*	105	117	81	92	91	110	112	117	88	116	115	104
10	139	123	163	143	159	146	179	143	156	150	154	167
15	91	137	143	137	193	167	187	189	212	175	183	167
20	95	113	140	135	149	164	159	141	152	132	136	158
25	223	246	225	212	242	222	209	207	207	168	193	187
30	295	304	320	351	327	341	296	304	256	221	218	208
35	484	512	462	446	449	411	421	399	339	350	322	308
40	985	977	1,134	1,252	1,269	1,315	1,479	1,528	1,438	1,288	1,329	1,335
45	1,152	1,137	1,202	1,283	1,477	1,612	1,662	1,900	1,767	1,448	1,391	1,505
50	1,585	1,493	1,683	1,738	1,841	1,998	2,176	2,454	2,201	1,711	1,749	2,052
55	2,298	2,223	2,034	2,215	2,326	2,390	2,628	3,020	2,607	2,048	1,837	2,124

60	2,210	2,287	2,549	2,704	3,124	3,077	3,424	3,785	3,172	2,141	1,993	2,196
65	1,884	1,913	2,015	2,181	2,563	2,672	3,046	4,014	3,385	2,406	2,163	2,401
70	1,727	1,626	1,675	1,902	2,011	2,145	2,340	2,919	2,529	1,786	1,672	1,990
75	1,358	1,252	1,299	1,377	1,553	1,489	1,680	2,128	1,876	1,346	1,186	1,374
80	962	944	910	937	997	1,000	1,067	1,296	1,107	864	739	812
85	451	480	431	504	511	528	541	623	588	456	438	440
90	169	154	166	162	177	154	153	212	184	155	118	147
95	29	28	32	39	29	26	43	31	31	21	26	29
100	*	.	*	*	*	*	*	*	*	*	.	*

Type of DM T1

		doDM										
		All	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003
		N	N	N	N	N	N	N	N	N	N	N
All		47,339	21,640	3,151	1,323	1,306	1,335	1,183	1,243	1,260	1,181	1,129
a5												
0		2,338	1,231	54	54	58	47	63	58	52	42	52
*		3,878	1,945	71	53	79	73	66	97	97	100	92
10		5,215	2,626	80	79	81	79	100	91	123	103	100
15		3,603	1,949	65	86	77	76	58	65	67	71	63
20		3,519	1,865	114	82	81	85	83	64	93	86	75
25		3,677	1,803	164	103	114	114	83	101	100	87	72
30		3,061	1,565	125	89	86	104	95	83	87	81	64
35		2,805	1,317	119	82	88	75	61	102	86	78	89
40		2,724	1,202	108	90	77	100	86	79	78	70	65
45		2,526	1,103	173	88	76	102	59	68	67	63	56
50		2,461	975	199	99	94	89	87	90	79	68	58
55		2,359	920	256	73	80	69	78	82	69	82	71
60		2,244	840	289	86	70	75	58	57	66	45	75
65		2,132	819	371	75	61	66	67	60	74	60	41
70		1,919	690	403	71	77	71	58	46	49	50	55
75		1,484	495	286	61	55	52	41	48	39	41	53
80		895	216	200	32	33	39	25	24	19	35	27
85		367	63	63	18	15	15	9	23	13	12	16
90		111	14	10	*	*	*	6	*	*	*	*
95		20	*	*	.	*	.	.	.	*	*	.
100		*

(Continued)

		doDM											
		2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015
		N	N	N	N	N	N	N	N	N	N	N	N
All		1,154	1,094	1,170	1,126	1,167	1,097	1,083	1,015	936	935	927	884
a5													
0		40	55	53	48	60	61	46	56	48	61	46	53
*		103	107	79	91	91	108	105	112	83	112	110	104
10		131	115	159	138	150	143	171	136	153	148	151	158
15		57	80	71	61	94	90	83	94	108	85	97	106
20		57	57	61	66	78	82	76	74	86	77	82	95
25		70	69	95	80	76	72	76	67	65	72	85	109
30		65	66	74	70	57	62	64	51	46	44	54	29
35		63	80	81	59	65	73	75	52	52	43	39	26
40		71	56	79	87	88	65	72	65	47	55	40	44
45		74	58	55	68	66	60	52	57	51	56	37	37
50		71	52	71	75	50	51	48	39	35	45	56	30

55	88	71	61	68	45	44	46	47	35	28	24	22
60	69	60	73	60	76	55	47	44	35	26	27	11
65	54	45	39	40	55	34	35	42	36	21	22	15
70	46	35	37	39	35	30	30	26	21	19	20	11
75	40	32	36	37	33	26	24	18	15	18	19	15
80	27	32	35	22	32	19	17	21	11	12	8	9
85	20	14	9	12	9	13	8	11	7	6	6	*
90	6	9	*	*	6	8	6	*	*	7	*	*
95	*	*	*	*	*	.	*	*	.	.	.	*
100	*

Type of DM T2

												doDM											
												All	1994	1995	1996	1997	1998	1999	2000	2001	2002	2003	
												N	N	N	N	N	N	N	N	N	N	N	
All	380,912	34,938	34,904	10,794	10,283	11,548	11,605	12,541	12,425	12,701	14,688												
a5																							
0	161	91	6	*	*	*	*	*	*	*	*											*	
*	123	67	.	*	*	*	*	*	*	*	*											*	
10	218	124	*	*	*	*	*	*	*	*	*											*	
15	1,430	304	13	12	16	15	18	16	18	35	28												
20	1,793	646	31	38	33	36	50	39	41	39	57												
25	3,501	1,054	91	86	69	89	93	111	89	101	113												
30	6,492	1,603	415	211	193	215	208	228	219	183	258												
35	10,352	2,400	712	330	317	350	358	435	408	407	440												
40	23,801	3,281	1,036	498	490	557	590	731	649	613	796												
45	29,697	3,986	1,628	825	744	844	816	978	907	996	1,108												
50	39,303	4,074	2,260	1,213	1,195	1,414	1,357	1,505	1,367	1,366	1,494												
55	46,391	3,974	2,724	1,127	1,182	1,380	1,473	1,606	1,779	1,865	2,110												
60	51,774	3,635	3,778	1,281	1,240	1,380	1,441	1,603	1,560	1,723	2,054												
65	49,868	3,277	4,756	1,343	1,214	1,426	1,392	1,442	1,453	1,545	1,815												
70	43,291	2,815	5,614	1,350	1,223	1,335	1,283	1,373	1,382	1,375	1,568												
75	34,084	2,022	5,175	1,152	1,160	1,183	1,152	1,143	1,114	1,075	1,303												
80	22,783	1,083	3,919	823	708	761	809	762	816	813	899												
85	11,630	415	2,038	389	369	424	411	414	451	399	449												
90	3,576	81	614	87	103	116	127	124	139	134	156												
95	609	6	86	21	16	16	18	24	21	21	29												
100	35	.	7	.	*	*	.	*	*	*	*												

(Continued)

												doDM											
												2004	2005	2006	2007	2008	2009	2010	2011	2012	2013	2014	2015
												N	N	N	N	N	N	N	N	N	N	N	N
All	15,132	15,030	15,549	16,735	18,386	18,937	20,770	24,459	21,410	16,114	15,085	16,878											
a5																							
0	*	*	*	*	*	*	*	6	*	*	*	*											
*	*	10	*	*	.	*	7	*	*	*	*	*											
10	8	8	*	*	9	*	8	7	*	*	*	9											
15	34	57	72	76	99	77	104	95	104	90	86	61											
20	38	56	79	69	71	82	83	67	66	55	54	63											
25	153	177	130	132	166	150	133	140	142	96	108	78											
30	230	238	246	281	270	279	232	253	210	177	164	179											
35	421	432	381	387	384	338	346	347	287	307	283	282											
40	914	921	1,055	1,165	1,181	1,250	1,407	1,463	1,391	1,233	1,289	1,291											
45	1,078	1,079	1,147	1,215	1,411	1,552	1,610	1,843	1,716	1,392	1,354	1,468											

50	1,514	1,441	1,612	1,663	1,791	1,947	2,128	2,415	2,166	1,666	1,693	2,022
55	2,210	2,152	1,973	2,147	2,281	2,346	2,582	2,973	2,572	2,020	1,813	2,102
60	2,141	2,227	2,476	2,644	3,048	3,022	3,377	3,741	3,137	2,115	1,966	2,185
65	1,830	1,868	1,976	2,141	2,508	2,638	3,011	3,972	3,349	2,385	2,141	2,386
70	1,681	1,591	1,638	1,863	1,976	2,115	2,310	2,893	2,508	1,767	1,652	1,979
75	1,318	1,220	1,263	1,340	1,520	1,463	1,656	2,110	1,861	1,328	1,167	1,359
80	935	912	875	915	965	981	1,050	1,275	1,096	852	731	803
85	431	466	422	492	502	515	533	612	581	450	432	435
90	163	145	165	160	171	146	147	210	182	148	114	144
95	27	27	31	36	28	26	41	30	31	21	26	27
100	*	.	*	*	*	*	*	*	*	*	.	*

The reconstructed diabetes register
Diagnoses 2012 ff. only

14:29 Tuesday, January 24, 2017 8

	Type of DM			Type of DM		
	All	T1	T2	All	T1	T2
	N	N	N	PctN	PctN	PctN
All	73,169	3,682	69,487	100.0	5.0	95.0
doDM						
2012/01	2,955	298	2,657	100.0	10.1	89.9
2012/02	1,851	47	1,804	100.0	2.5	97.5
2012/03	2,211	57	2,154	100.0	2.6	97.4
2012/04	1,583	42	1,541	100.0	2.7	97.3
2012/05	1,950	66	1,884	100.0	3.4	96.6
2012/06	2,185	75	2,110	100.0	3.4	96.6
2012/07	1,271	51	1,220	100.0	4.0	96.0
2012/08	1,526	63	1,463	100.0	4.1	95.9
2012/09	1,695	63	1,632	100.0	3.7	96.3
2012/10	1,827	72	1,755	100.0	3.9	96.1
2012/11	1,813	61	1,752	100.0	3.4	96.6
2012/12	1,479	41	1,438	100.0	2.8	97.2
2013/01	2,435	250	2,185	100.0	10.3	89.7
2013/02	1,494	80	1,414	100.0	5.4	94.6
2013/03	1,468	72	1,396	100.0	4.9	95.1
2013/04	1,511	66	1,445	100.0	4.4	95.6
2013/05	1,363	52	1,311	100.0	3.8	96.2
2013/06	1,369	63	1,306	100.0	4.6	95.4
2013/07	1,029	41	988	100.0	4.0	96.0
2013/08	1,170	60	1,110	100.0	5.1	94.9
2013/09	1,342	51	1,291	100.0	3.8	96.2
2013/10	1,318	61	1,257	100.0	4.6	95.4
2013/11	1,320	73	1,247	100.0	5.5	94.5
2013/12	1,230	66	1,164	100.0	5.4	94.6
2014/01	1,840	174	1,666	100.0	9.5	90.5
2014/02	1,277	67	1,210	100.0	5.2	94.8
2014/03	1,475	75	1,400	100.0	5.1	94.9
2014/04	1,162	62	1,100	100.0	5.3	94.7
2014/05	1,304	80	1,224	100.0	6.1	93.9
2014/06	1,306	47	1,259	100.0	3.6	96.4
2014/07	1,049	62	987	100.0	5.9	94.1
2014/08	1,079	71	1,008	100.0	6.6	93.4
2014/09	1,414	76	1,338	100.0	5.4	94.6
2014/10	1,410	61	1,349	100.0	4.3	95.7
2014/11	1,342	81	1,261	100.0	6.0	94.0
2014/12	1,354	71	1,283	100.0	5.2	94.8
2015/01	1,421	87	1,334	100.0	6.1	93.9
2015/02	1,423	80	1,343	100.0	5.6	94.4
2015/03	1,670	82	1,588	100.0	4.9	95.1
2015/04	1,432	70	1,362	100.0	4.9	95.1
2015/05	1,387	71	1,316	100.0	5.1	94.9
2015/06	1,629	69	1,560	100.0	4.2	95.8
2015/07	1,168	65	1,103	100.0	5.6	94.4
2015/08	1,310	66	1,244	100.0	5.0	95.0

2015/09	1,601	83	1,518	100.0	5.2	94.8
2015/10	1,589	82	1,507	100.0	5.2	94.8
2015/11	1,515	65	1,450	100.0	4.3	95.7
2015/12	1,617	64	1,553	100.0	4.0	96.0

The reconstructed diabetes register
Diagnoses 2012 ff. only

14:29 Tuesday, January 24, 2017 9

	Inclusion criterion						Inclusion criterion					
	All	DIA	DVD	Ins	NPR	OAD	All	DIA	DVD	Ins	NPR	OAD
	N	N	N	N	N	N	PctN	PctN	PctN	PctN	PctN	PctN
All	73,169	189	3,446	2,109	8,163	59,262	100.0	0.3	4.7	2.9	11.2	81.0
doDM												
2012/01	2,955	18	965	34	180	1,758	100.0	0.6	32.7	1.2	6.1	59.5
2012/02	1,851	10	46	40	134	1,621	100.0	0.5	2.5	2.2	7.2	87.6
2012/03	2,211	8	59	35	155	1,954	100.0	0.4	2.7	1.6	7.0	88.4
2012/04	1,583	6	39	36	125	1,377	100.0	0.4	2.5	2.3	7.9	87.0
2012/05	1,950	*	58	30	172	1,685	100.0	0.3	3.0	1.5	8.8	86.4
2012/06	2,185	9	135	44	159	1,838	100.0	0.4	6.2	2.0	7.3	84.1
2012/07	1,271	*	26	31	114	1,096	100.0	0.3	2.0	2.4	9.0	86.2
2012/08	1,526	*	47	35	169	1,271	100.0	0.3	3.1	2.3	11.1	83.3
2012/09	1,695	6	34	41	140	1,474	100.0	0.4	2.0	2.4	8.3	87.0
2012/10	1,827	*	38	39	184	1,564	100.0	0.1	2.1	2.1	10.1	85.6
2012/11	1,813	7	34	40	159	1,573	100.0	0.4	1.9	2.2	8.8	86.8
2012/12	1,479	*	41	35	100	1,302	100.0	0.1	2.8	2.4	6.8	88.0
2013/01	2,435	*	776	35	168	1,452	100.0	0.2	31.9	1.4	6.9	59.6
2013/02	1,494	*	40	41	166	1,244	100.0	0.2	2.7	2.7	11.1	83.3
2013/03	1,468	*	40	44	166	1,214	100.0	0.3	2.7	3.0	11.3	82.7
2013/04	1,511	*	32	33	170	1,272	100.0	0.3	2.1	2.2	11.3	84.2
2013/05	1,363	*	29	46	165	1,122	100.0	0.1	2.1	3.4	12.1	82.3
2013/06	1,369	*	70	49	136	1,111	100.0	0.2	5.1	3.6	9.9	81.2
2013/07	1,029	.	20	64	117	828	100.0	.	1.9	6.2	11.4	80.5
2013/08	1,170	*	36	52	128	952	100.0	0.2	3.1	4.4	10.9	81.4
2013/09	1,342	*	25	35	141	1,138	100.0	0.2	1.9	2.6	10.5	84.8
2013/10	1,318	*	35	58	161	1,059	100.0	0.4	2.7	4.4	12.2	80.3
2013/11	1,320	*	36	39	152	1,088	100.0	0.4	2.7	3.0	11.5	82.4
2013/12	1,230	*	28	48	139	1,012	100.0	0.2	2.3	3.9	11.3	82.3
2014/01	1,840	*	411	55	182	1,187	100.0	0.3	22.3	3.0	9.9	64.5
2014/02	1,277	*	38	47	139	1,049	100.0	0.3	3.0	3.7	10.9	82.1
2014/03	1,475	6	34	34	185	1,216	100.0	0.4	2.3	2.3	12.5	82.4
2014/04	1,162	*	28	39	157	934	100.0	0.3	2.4	3.4	13.5	80.4
2014/05	1,304	*	33	41	161	1,064	100.0	0.4	2.5	3.1	12.3	81.6
2014/06	1,306	8	32	31	146	1,089	100.0	0.6	2.5	2.4	11.2	83.4
2014/07	1,049	*	29	36	145	837	100.0	0.2	2.8	3.4	13.8	79.8
2014/08	1,079	*	31	39	151	854	100.0	0.4	2.9	3.6	14.0	79.1
2014/09	1,414	7	25	45	198	1,139	100.0	0.5	1.8	3.2	14.0	80.6
2014/10	1,410	10	34	59	209	1,098	100.0	0.7	2.4	4.2	14.8	77.9
2014/11	1,342	6	27	49	218	1,042	100.0	0.4	2.0	3.7	16.2	77.6
2014/12	1,354	*	11	64	183	1,094	100.0	0.1	0.8	4.7	13.5	80.8
2015/01	1,421	*	21	49	204	1,145	100.0	0.1	1.5	3.4	14.4	80.6
2015/02	1,423	7	*	52	206	1,155	100.0	0.5	0.2	3.7	14.5	81.2
2015/03	1,670	.	.	55	236	1,379	100.0	.	.	3.3	14.1	82.6
2015/04	1,432	.	.	44	208	1,180	100.0	.	.	3.1	14.5	82.4
2015/05	1,387	.	.	50	207	1,130	100.0	.	.	3.6	14.9	81.5
2015/06	1,629	.	.	54	218	1,357	100.0	.	.	3.3	13.4	83.3
2015/07	1,168	.	.	41	168	959	100.0	.	.	3.5	14.4	82.1
2015/08	1,310	.	.	44	202	1,064	100.0	.	.	3.4	15.4	81.2
2015/09	1,601	.	.	58	219	1,324	100.0	.	.	3.6	13.7	82.7
2015/10	1,589	.	.	40	231	1,318	100.0	.	.	2.5	14.5	82.9
2015/11	1,515	.	.	49	205	1,261	100.0	.	.	3.2	13.5	83.2
2015/12	1,617	.	.	50	185	1,382	100.0	.	.	3.1	11.4	85.5

The reconstructed diabetes *drug* register

14:29 Tuesday, January 24, 2017 10

The CONTENTS Procedure

```

Data Set Name      TTDATA.DMDREG      Observations      409841
Member Type       DATA              Variables         9
Engine            V9                 Indexes           0
Created            24/01/2017 14:29:15  Observation Length 72
Last Modified     24/01/2017 14:29:15  Deleted Observations 0
Protection        Compressed         NO
Data Set Type     Sorted             NO
Label
Data Representation WINDOWS_64
Encoding          wlatin1 Western (Windows)
    
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Engine/Host Dependent Information

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Data Set Page Size      65536
Number of Data Set Pages 452
First Data Page         *
Max Obs per Page        908
Obs in First Data Page  881
Number of Data Set Repairs 0
ExtendObsCounter        YES
Filename                 E:\workdata\705093\BxC\demoDM\DATA\dmdreg.sas7bdat
Release Created          9.0401M3
Host Created             X64_SRV12
    
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Alphabetic List of Variables and Attributes

#	Variable	Type	Len	Format	Informat	Label
8	DMtp	Char	*			
5	doBth	Num	8	DDMMYY10.		
7	doDM	Num	8	DDMMYY10.		
6	doDth	Num	8	DDMMYY10.		
2	doIns	Num	8	DDMMYY10.		
3	doOAD	Num	8	DDMMYY10.		
9	inCr	Char	*			
1	pnr	Char	12	\$12.	\$10.	Personnummer
4	sex	Num	8			

The reconstructed diabetes *drug* register

14:29 Tuesday, January 24, 2017 11

	DMtp		inCr		All	T1	T2	Ins	OAD	PctN	PctN	PctN	PctN	PctN			
	DMtp	inCr	All	T1											T2	Ins	OAD
	All	T1	T2	Ins											OAD	PctN	PctN
All	409,841	16,029	393,812	66,056	343,785	100.0	3.9	96.1	16.1	83.9							
doDM																	
1995	79,746	4,629	75,117	31,393	48,353	100.0	5.8	94.2	39.4	60.6							
1996	12,915	1,268	11,647	2,519	10,396	100.0	9.8	90.2	19.5	80.5							
1997	11,321	530	10,791	1,597	9,724	100.0	4.7	95.3	14.1	85.9							
1998	12,245	475	11,770	1,605	10,640	100.0	3.9	96.1	13.1	86.9							
1999	12,434	498	11,936	1,587	10,847	100.0	4.0	96.0	12.8	87.2							
2000	12,605	472	12,133	1,660	10,945	100.0	3.7	96.3	13.2	86.8							
2001	13,250	505	12,745	1,610	11,640	100.0	3.8	96.2	12.2	87.8							
2002	13,132	507	12,625	1,601	11,531	100.0	3.9	96.1	12.2	87.8							
2003	15,361	480	14,881	1,567	13,794	100.0	3.1	96.9	10.2	89.8							
2004	15,909	506	15,403	1,714	14,195	100.0	3.2	96.8	10.8	89.2							
2005	15,757	497	15,260	1,685	14,072	100.0	3.2	96.8	10.7	89.3							
2006	16,377	527	15,850	1,759	14,618	100.0	3.2	96.8	10.7	89.3							
2007	17,715	508	17,207	1,782	15,933	100.0	2.9	97.1	10.1	89.9							
2008	19,427	550	18,877	1,793	17,634	100.0	2.8	97.2	9.2	90.8							
2009	20,194	583	19,611	1,777	18,417	100.0	2.9	97.1	8.8	91.2							
2010	22,314	583	21,731	1,792	20,522	100.0	2.6	97.4	8.0	92.0							
2011	25,900	561	25,339	1,657	24,243	100.0	2.2	97.8	6.4	93.6							

2012	22,673	580	22,093	1,620	21,053	100.0	2.6	97.4	7.1	92.9
2013	17,089	574	16,515	1,779	15,310	100.0	3.4	96.6	10.4	89.6
2014	15,865	585	15,280	1,781	14,084	100.0	3.7	96.3	11.2	88.8
2015	17,612	611	17,001	1,778	15,834	100.0	3.5	96.5	10.1	89.9

7.11 06a-comp

Compares the prevalence of DM as derived from the revised (shrunk) version of the NDR and the newly constructed DMreg, at 1 Jan 1996, 2004 and 2012.

```

1                                     "Program 06a-comp.sas"

NOTE: Copyright (c) 2002-2012 by SAS Institute Inc., Cary, NC, USA.
NOTE: SAS (r) Proprietary Software 9.4 (TS1M3)
      Licensed to FORSKNING 1, Site 50800722.
NOTE: This session is executing on the X64_SRV12 platform.

NOTE: Updated analytical products:
      SAS/STAT 14.1

NOTE: Additional host information:
      X64_SRV12 WIN 6.2.9200 Server

NOTE: SAS initialization used:
      real time          0.07 seconds
      cpu time           0.09 seconds

NOTE: AUTOEXEC processing beginning; file is E:\workdata\705093\BXC\demoDM\sas\optslibs.sas.

NOTE: AUTOEXEC processing completed.

1      data prev ;
2          merge TTdata.ndr      ( in = nn )
3              TTdata.dmreg     ( in = dd ) ;
4          by pnr ;
5          ndr = nn ;
6          dmreg = dd ;
7          mdif = (doNDR - doDM )/30 ;
8          do pdat = 1996, 2004, 2012 ;
9              a5 = floor( (mdy(1,1,pdat)-doBth )/(365.25*5) ) * 5 ;
10             if ( ndr      and doNDR < mdy(1,1,pdat) < min(doDth,mdy(1,2,pdat)) ) or
11                 ( dmreg and doDM < mdy(1,1,pdat) < min(doDth,mdy(1,2,pdat)) ) then
11         ! output ;
12         end ;
13     run ;

NOTE: Missing values were generated as a result of performing an operation on missing
      values.
      Each place is given by: (Number of times) at (Line):(Column).
      95279 at 7:17
NOTE: There were 389304 observations read from the data set TTDATA.NDR.
NOTE: There were 428251 observations read from the data set TTDATA.DMREG.
NOTE: The data set WORK.PREV has 499300 observations and 20 variables.
NOTE: DATA statement used (Total process time):
      real time          1.03 seconds
      cpu time           0.64 seconds

14
15     proc tabulate data = prev noseps missing ;

```

```

16      class a5 ndr dmreg pdat sex inklaars inCr ;
17      table all a5,
18          pdat * ( all * f=comma9.
19              ndr * dmreg * pctn<ndr*dmreg> * f=5.1 )
20          / rts = 5 ;
21      table pdat * ( all a5 ),
22          ( all * f=comma7.
23              ndr * dmreg * inCr="DMreg" * pctn<ndr*dmreg*inCr> * f=5.1 )
24          / rts = 11 indent = 1 ;
25      table pdat * ( all sex ) *
26          ( all inklaars="NDR" ),
27          ( all inCr ) * f=comma7.
28          / rts = 19 ;
29      keylabel n = ' ' ;
30      run ;
    
```

NOTE: There were 499300 observations read from the data set WORK.PREV.

NOTE: The PROCEDURE TABULATE printed pages 1-3.

NOTE: PROCEDURE TABULATE used (Total process time):

```

real time      0.09 seconds
cpu time       0.26 seconds
    
```

NOTE: SAS Institute Inc., SAS Campus Drive, Cary, NC USA 27513-2414

NOTE: The SAS System used:

```

real time      1.34 seconds
cpu time       0.99 seconds
    
```

7.11.1 06a-comp.lst

The SAS System

14:32 Tuesday, January 24, 2017 1

	pdat											
	1996				2004				2012			
	ndr				ndr				ndr			
	0	*			0	*			0	*		
dmreg	dmreg			dmreg	dmreg			dmreg	dmreg			
*	0	*		*	0	*		*	0	*		
All	PctN	PctN	PctN	All	PctN	PctN	PctN	All	PctN	PctN	PctN	
All	95,310	2.0	3.3	94.7	154,703	2.6	3.8	93.7	249,287	3.7	4.7	91.6
a5	*	.	100.0
-10												
0	102	9.8	10.8	79.4	121	4.1	7.4	88.4	122	13.9	5.7	80.3
*	368	11.4	2.4	86.1	498	6.4	2.6	91.0	553	3.6	4.2	92.2
10	571	8.4	0.5	91.1	915	4.7	2.5	92.8	1,129	3.9	2.1	94.0
15	1,153	5.6	1.1	93.2	1,207	6.4	2.6	91.1	1,951	6.1	3.4	90.5
20	1,658	3.6	1.7	94.7	1,515	6.4	2.2	91.4	2,288	11.1	4.7	84.2
25	2,356	3.8	2.9	93.3	2,527	5.9	3.6	90.5	2,642	13.9	7.1	79.0
30	3,187	3.7	2.1	94.2	3,721	6.0	6.0	87.9	4,002	12.6	9.1	78.2
35	4,100	4.7	2.2	93.1	5,828	4.7	5.5	89.8	6,280	7.8	8.8	83.4
40	5,113	3.3	2.2	94.5	7,776	4.9	3.7	91.4	10,526	6.1	6.1	87.8
45	6,908	2.3	2.3	95.4	10,400	3.7	2.7	93.6	16,189	5.2	4.0	90.8
50	8,551	1.6	1.5	96.9	13,254	2.7	2.6	94.8	20,269	4.2	3.1	92.6
55	8,599	1.2	1.3	97.5	19,605	1.9	2.0	96.1	26,043	3.6	3.0	93.5
60	9,492	1.1	1.9	97.1	19,253	1.7	2.4	95.8	32,840	2.7	3.1	94.2
65	10,369	1.1	2.8	96.1	18,258	1.5	2.9	95.6	39,434	2.4	3.4	94.1
70	10,800	1.3	3.6	95.1	16,585	1.8	3.8	94.4	30,531	2.6	4.0	93.4
75	9,716	1.1	5.1	93.8	14,492	1.7	5.0	93.3	24,249	2.4	5.4	92.1
80	7,129	1.6	6.5	91.9	10,724	1.9	6.6	91.5	16,804	2.9	7.0	90.2
85	3,823	2.7	9.1	88.2	5,598	2.3	8.4	89.3	9,239	3.2	10.2	86.6

90	1,139	4.3	10.4	85.3	2,030	2.8	10.7	86.5	3,459	3.2	13.1	83.7
95	159	8.2	11.3	80.5	363	3.3	11.0	85.7	673	4.9	15.9	79.2
100	15	13.3	6.7	80.0	31	3.2	12.9	83.9	60	3.3	18.3	78.3
105	*	100.0	.	.	*	50.0	.	50.0	*	.	25.0	75.0

The SAS System

14:32 Tuesday, January 24, 2017 2

ndr												
0						*						
dmreg						dmreg						
*						0						
DMreg						DMreg						
	DIA	DVD	Ins	NPR	OAD		DIA	DVD	Ins	NPR	OAD	
All	PctN	PctN	PctN	PctN	PctN	PctN	PctN	PctN	PctN	PctN	PctN	
1996												
All	95,310	.	0.0	0.2	1.5	0.4	3.3	.	27.4	3.4	32.9	31.1
-10	*	100.0
0	102	.	.	.	9.8	.	10.8	.	51.0	.	28.4	.
*	368	.	0.5	.	10.9	.	2.4	.	70.4	.	15.5	0.3
10	571	.	0.7	.	7.7	.	0.5	.	76.0	0.2	14.9	.
15	1,153	.	0.3	0.3	4.8	0.3	1.1	.	72.3	0.9	19.9	0.1
20	1,658	.	0.1	0.8	2.6	0.1	1.7	.	71.5	1.3	21.4	0.5
25	2,356	.	0.1	0.5	2.8	0.4	2.9	.	65.7	2.9	23.0	1.6
30	3,187	.	0.2	0.4	2.9	0.3	2.1	.	61.2	6.0	23.8	3.2
35	4,100	.	0.1	0.5	3.6	0.5	2.2	.	54.3	9.0	25.8	4.0
40	5,113	.	0.0	0.4	2.7	0.2	2.2	.	51.6	7.7	28.0	7.1
45	6,908	.	0.0	0.3	1.7	0.3	2.3	.	50.9	4.4	27.9	12.2
50	8,551	.	0.0	0.2	1.1	0.2	1.5	.	44.2	2.3	30.4	19.9
55	8,599	.	0.0	0.1	0.7	0.4	1.3	.	35.5	2.1	33.7	26.2
60	9,492	.	.	0.1	0.7	0.3	1.9	.	25.4	2.1	35.4	34.2
65	10,369	.	.	0.1	0.6	0.4	2.8	.	13.8	2.6	37.0	42.6
70	10,800	.	.	0.1	0.7	0.5	3.6	.	5.7	3.0	38.6	47.8
75	9,716	.	.	0.1	0.7	0.4	5.1	.	1.2	3.0	38.0	51.5
80	7,129	.	.	0.1	1.1	0.4	6.5	.	0.3	3.6	35.7	52.4
85	3,823	.	.	0.0	2.1	0.6	9.1	.	.	2.9	34.8	50.5
90	1,139	.	.	0.1	4.0	0.3	10.4	.	.	3.0	32.7	49.6
95	159	.	.	.	6.3	1.9	11.3	.	.	1.9	30.2	48.4
100	15	.	.	.	13.3	.	6.7	.	.	13.3	13.3	53.3
105	*	.	.	.	100.0
2004												
All	154,703	.	0.0	0.5	0.8	1.3	3.8	0.0	29.6	2.3	20.8	40.9
0	121	.	.	.	2.5	1.7	7.4	.	.	4.1	83.5	0.8
*	498	.	0.4	1.6	3.0	1.4	2.6	.	29.5	1.8	59.6	.
10	915	.	0.3	0.8	3.3	0.3	2.5	.	73.1	0.7	18.9	0.1
15	1,207	.	0.4	0.7	4.1	1.2	2.6	.	72.3	1.1	16.1	1.6
20	1,515	.	0.4	0.9	3.0	2.1	2.2	.	67.7	2.1	19.3	2.2
25	2,527	.	0.2	0.9	2.9	1.9	3.6	.	63.9	2.3	20.6	3.8
30	3,721	.	0.3	1.6	1.9	2.2	6.0	.	56.6	3.3	20.3	7.8
35	5,828	.	0.1	1.0	1.6	2.0	5.5	.	52.2	3.5	23.0	11.1
40	7,776	.	0.1	1.1	1.8	1.9	3.7	.	46.1	4.8	21.6	18.9
45	10,400	.	0.1	0.6	1.5	1.6	2.7	.	43.7	5.0	21.4	23.5
50	13,254	.	0.0	0.6	1.1	1.0	2.6	.	40.9	3.3	19.6	30.9
55	19,605	.	0.0	0.4	0.5	0.9	2.0	0.0	36.6	2.0	20.0	37.4
60	19,253	.	0.0	0.3	0.4	0.9	2.4	.	30.9	1.4	20.3	43.4
65	18,258	.	0.0	0.2	0.2	1.0	2.9	0.0	25.2	1.5	20.4	48.4
70	16,585	.	0.0	0.3	0.3	1.3	3.8	0.0	17.5	1.5	20.7	54.7
75	14,492	.	.	0.2	0.3	1.3	5.0	.	10.6	2.2	21.1	59.4
80	10,724	.	.	0.3	0.3	1.3	6.6	0.0	5.0	1.7	21.1	63.6
85	5,598	.	.	0.3	0.4	1.5	8.4	.	1.3	2.1	21.0	64.9
90	2,030	.	.	0.3	0.6	1.9	10.7	.	0.5	2.1	19.2	64.6
95	363	.	.	0.3	1.4	1.7	11.0	.	.	0.8	17.1	67.8

100	31	.	.	.	3.2	.	12.9	.	.	6.5	12.9	64.5
105	*	50.0	50.0
2012												
All	249,287	0.1	0.1	0.4	0.6	2.5	4.7	0.1	24.5	1.5	13.7	51.8
0	122	.	.	1.6	7.4	4.9	5.7	.	.	4.1	76.2	.
*	553	.	.	0.5	1.4	1.6	4.2	.	.	0.7	91.1	0.4
10	1,129	.	.	0.8	1.8	1.3	2.1	.	1.1	1.4	89.9	1.6
15	1,951	.	0.7	0.6	1.4	3.4	3.4	.	47.6	1.1	36.2	5.6
20	2,288	.	0.5	0.8	2.1	7.7	4.7	0.0	58.0	1.3	14.6	10.4
25	2,642	.	0.6	0.9	2.5	10.0	7.1	0.1	54.2	2.0	15.0	7.7
30	4,002	0.0	0.5	1.1	2.1	8.9	9.1	0.1	47.3	2.2	17.6	11.1
35	6,280	0.0	0.5	1.4	1.6	4.3	8.8	0.1	45.8	2.0	18.0	17.5
40	10,526	0.1	0.2	0.8	1.0	3.9	6.1	0.0	36.6	1.8	15.6	33.7
45	16,189	0.0	0.2	0.6	0.9	3.5	4.0	0.0	32.4	2.2	14.2	42.0
50	20,269	0.0	0.2	0.5	0.7	2.7	3.1	0.0	29.6	2.4	13.3	47.2
55	26,043	0.0	0.1	0.5	0.7	2.3	3.0	0.1	27.0	2.4	12.7	51.3
60	32,840	0.1	0.1	0.3	0.4	1.8	3.1	0.1	25.5	1.5	12.4	54.7
65	39,434	0.1	0.1	0.3	0.3	1.7	3.4	0.1	22.7	1.1	12.2	58.1
70	30,531	0.1	0.1	0.2	0.3	2.0	4.0	0.1	19.6	0.9	11.9	60.9
75	24,249	0.1	0.1	0.2	0.2	1.9	5.4	0.1	17.1	0.9	12.2	61.8
80	16,804	0.1	0.0	0.2	0.2	2.4	7.0	0.0	12.5	1.2	12.6	63.8
85	9,239	0.1	.	0.3	0.3	2.5	10.2	0.1	8.5	1.5	12.4	64.2
90	3,459	.	.	0.2	0.2	2.7	13.1	.	4.9	1.6	12.9	64.3
95	673	.	.	.	0.7	4.2	15.9	.	1.9	2.2	12.0	63.0
100	60	.	.	.	3.3	.	18.3	.	.	3.3	11.7	63.3
105	*	25.0	.	.	.	25.0	50.0

			Inclusion criterion						
			All	DIA	DVD	Ins	NPR	OAD	
pdat 1996	All	All	95,310	3,116	.	26,109	3,388	32,728	29,969
		NDR	1,930	.	30	160	1,397	343	
		ins	6,757	268	.	2,301	2,172	1,970	46
		lpr	53,238	2,611	.	16,966	818	25,177	7,666
	oad	33,385	237	.	6,812	238	4,184	21,914	
	sex *	All	49,701	1,275	.	14,714	1,341	17,189	15,182
		NDR	710	.	.	18	74	459	159
		ins	3,406	84	.	1,446	727	1,124	25
		lpr	28,427	1,094	.	9,412	421	13,534	3,966
	oad	17,158	97	.	3,838	119	2,072	11,032	
	* NDR	All	45,609	1,841	.	11,395	2,047	15,539	14,787
		ins	1,220	.	.	12	86	938	184
lpr		3,351	184	.	855	1,445	846	21	
oad		24,811	1,517	.	7,554	397	11,643	3,700	
2004	All	All	154,703	5,835	*	45,915	4,328	33,318	65,303
		NDR	3,947	.	.	74	712	1,203	1,958
		ins	6,487	258	.	2,517	2,387	1,196	129
		lpr	78,037	5,439	*	28,964	1,065	28,704	13,861
	oad	66,232	138	.	14,360	164	2,215	49,355	
	sex *	All	83,439	2,340	*	26,477	1,790	18,350	34,480
		NDR	1,542	.	.	48	336	352	806
		ins	3,279	77	.	1,564	850	725	63
		lpr	42,821	2,196	*	16,538	531	16,159	7,395
	oad	35,797	67	.	8,327	73	1,114	26,216	
	* NDR	All	71,264	3,495	*	19,438	2,538	14,968	30,823
		ins	2,405	.	.	26	376	851	1,152
lpr		3,208	181	.	953	1,537	471	66	
oad		35,216	3,243	*	12,426	534	12,545	6,466	

		oad	30,435	71	.	6,033	91	1,101	23,139
2012	All	All	249,287	11,609	321	61,435	4,846	35,535	135,541
		NDR							
			9,246	.	150	324	1,016	1,405	6,351
		ins	6,470	259	.	2,579	2,566	737	329
		lpr	103,985	10,925	117	39,835	1,099	32,206	19,803
		oad	129,586	425	54	18,697	165	1,187	109,058
	sex	All	136,043	4,685	180	35,929	2,094	19,447	73,708
	*	NDR							
			3,565	.	87	162	462	430	2,424
		ins	3,269	76	.	1,579	975	458	181
		lpr	56,796	4,333	67	23,163	584	17,946	10,703
		oad	72,413	276	26	11,025	73	613	60,400
	*	All	113,244	6,924	141	25,506	2,752	16,088	61,833
		NDR							
			5,681	.	63	162	554	975	3,927
		ins	3,201	183	.	1,000	1,591	279	148
		lpr	47,189	6,592	50	16,672	515	14,260	9,100
		oad	57,173	149	28	7,672	92	574	48,658

7.12 Tabulation of follow-up and prevalence

For analysis of incidence rates of diabetes and mortality rates for persons with and without diabetes we tabulate the events and person-years for the entire Danish population by sex, state (T1, T2 and no DM), age and date of follow-up and date of birth, the latter three in 1-year intervals. Since we have individual records for the entire population (with sex, date of birth, migration and death) we could in principle choose smaller intervals of say half a year or 3 months.

7.13 08-mkFU

Defines follow-up intervals for all persons in the population, taking emigration and immigration records into account. These intervals are subdivided in pre- and post-DM diagnosis, the latter marked as T1 or T2. This follow-up is then split by age and subsequently by date using the `Lexis` macro.

Subsequently the number of deaths and DM (T1 or T2) events are tabulated by sex, age and date of follow-up and date of birth too.

Finally all is merged to a dataset, `FUtab`, classified by sex, age and calendar time and upper/lower Lexis triangle (corresponding to the two possible periods of date of birth for a given combination of period of age and date of follow-up) [1].

```
1 "Program 08-mkFU.sas"
```

```
NOTE: Copyright (c) 2002-2012 by SAS Institute Inc., Cary, NC, USA.
```

```
NOTE: SAS (r) Proprietary Software 9.4 (TS1M3)
```

```
      Licensed to FORSKNING 1, Site 50800722.
```

```
NOTE: This session is executing on the X64_SRV12 platform.
```

```
NOTE: Updated analytical products:
```

```
      SAS/STAT 14.1
```

```
NOTE: Additional host information:
```

```
      X64_SRV12 WIN 6.2.9200 Server
```

NOTE: SAS initialization used:
 real time 0.06 seconds
 cpu time 0.10 seconds

NOTE: AUTOEXEC processing beginning; file is E:\workdata\705093\BXC\demoDM\sas\optslibs.sas.

NOTE: AUTOEXEC processing completed.

```

1      * Begin and end of study period, respectively ;
2      %let bos = '01jan1995'd ;
3      %let eos = '31dec2015'd ;
4
5      proc sort  data = TTdata.pop    out = pop    ; by pnr    ; run ;

```

NOTE: There were 7361669 observations read from the data set TTDATA.POP.

NOTE: The data set WORK.POP has 7361669 observations and 4 variables.

NOTE: PROCEDURE SORT used (Total process time):
 real time 2.96 seconds
 cpu time 2.23 seconds

```

6      proc sort  data = TTdata.xdk    out = xdk    ; by pnr doEm ; run ;

```

NOTE: There were 1825733 observations read from the data set TTDATA.XDK.

NOTE: The data set WORK.XDK has 1825733 observations and 3 variables.

NOTE: PROCEDURE SORT used (Total process time):
 real time 0.51 seconds
 cpu time 0.59 seconds

```

7      proc sort  data = TTdata.DMreg  out = dmreg  ; by pnr    ; run ;

```

NOTE: There were 428251 observations read from the data set TTDATA.DMREG.

NOTE: The data set WORK.DMREG has 428251 observations and 12 variables.

NOTE: PROCEDURE SORT used (Total process time):
 real time 0.64 seconds
 cpu time 0.25 seconds

```

8
9      data FU ;
10     merge pop ( in = p )
11           xdk ( in = x ) ;
12     by pnr ;
13     format doBth doDth doEm doIm entry exit ddmmyy10. ;
14     if ^p then delete ;
15     retain entry exit ;
16     if first.pnr then do ;
17         entry = max( &bos., doBth ) ;
18         exit = min( &eos., doDth ) ;
19     end ;
20     * if no migration records just output ;
21     if ^x then output ;
22     else do ;
23         if doEm > .z then do ;
24             exit = min( exit, doEm ) ;
25             output ;
26             firstout = 1 ;
27         end ;
28         if doIm > .z then do ;
29             entry = max( entry, doIm ) ;
30             exit = min( &eos., doDth ) ;
31             firstout = 0 ;
32         end ;
33         if last.pnr and ^firstout then output ;
34     end ;
35     run ;

```

NOTE: There were 7361669 observations read from the data set WORK.POP.
 NOTE: There were 1825733 observations read from the data set WORK.XDK.
 NOTE: The data set WORK.FU has 7825474 observations and 9 variables.
 NOTE: DATA statement used (Total process time):
 real time 2.46 seconds
 cpu time 2.10 seconds

```

36
37     data TTdata.FU ;
38         merge FU DMreg ( in = dm
39                         keep = pnr doDM DMtp ) ;
40     by pnr ;
41     state = "Well" ;
42     if ^dm then output ;
43     if dm then do ;
44         if exit < doDM          then output ;
45         if doDM < entry then do ;
46             state = DMtp ;
47             output ;
48             end ;
49         if entry < doDM < exit then do ;
50             oldex = exit ;
51             exit = doDM ;
52             output ;
53             entry = doDM ;
54             exit = oldex ;
55             state = DMtp ;
56             output ;
57             end ;
58     end ;
59     run ;

```

NOTE: There were 7825474 observations read from the data set WORK.FU.
 NOTE: There were 428251 observations read from the data set WORK.DMREG.
 NOTE: The data set TTDATA.FU has 8191903 observations and 13 variables.
 NOTE: DATA statement used (Total process time):
 real time 5.86 seconds
 cpu time 2.09 seconds

```

60
61     data FU ;
62         set TTdata.FU ;
63         fail = ( doDth eq exit ) ;
64     run ;

```

NOTE: There were 8191903 observations read from the data set TTDATA.FU.
 NOTE: The data set WORK.FU has 8191903 observations and 14 variables.
 NOTE: DATA statement used (Total process time):
 real time 5.67 seconds
 cpu time 1.28 seconds

```

65
66     options nosource2 ;
67     %inc '../..../sas/lexis.sas' ;
224
225     * tabulation interval for risk time and events ;
226     %let int = 1 ;
227
228     %lexis( data = FU,
229            out = FUa,
230            breaks = 0 to 100 by &int.,
231            origin = doBth,
232            scale = 365.25,
233            left = A ) ;

```

NOTE: There were 8191903 observations read from the data set WORK.FU.

NOTE: The data set WORK.DISCRD has 91 observations and 17 variables.
 NOTE: The data set WORK.FUA has 121849450 observations and 17 variables.
 NOTE: DATA statement used (Total process time):
 real time 55.10 seconds
 cpu time 39.75 seconds

```
234
235     %lexis( data = FUA,
236             out = FUap,
237             breaks = 190 to 220 by &int.,
238             origin = '01jan1800'd,
239             scale = 365.25,
240             left = P ) ;
```

NOTE: There were 121849450 observations read from the data set WORK.FUA.
 NOTE: The data set WORK.DISCRD has 0 observations and 18 variables.
 NOTE: The data set WORK.FUAP has 235985275 observations and 18 variables.
 NOTE: DATA statement used (Total process time):
 real time 3:27.73
 cpu time 2:47.21

```
241
242     data FU ;
243     set FUap ;
244     * set the counters of death and DM incidence ;
245     D = ( doDth eq exit ) ;
246     T1 = 0 ;
247     T2 = 0 ;
248     if ( doDM eq exit ) then do ;
249         if ( DMtp eq 'T1' ) then T1 = 1 ;
250         if ( DMtp eq 'T2' ) then T2 = 1 ;
251     end ;
252     C = floor( (doBth-'01jan1800'd)/(365.25*&int.) ) * &int. + 1800 ;
253     P = P + 1800 ;
254     Y = risk/1000 ;
255     label A = "Age at FU"
256           P = "Date of FU" ;
257     run ;
```

NOTE: There were 235985275 observations read from the data set WORK.FUAP.
 NOTE: The data set WORK.FU has 235985275 observations and 23 variables.
 NOTE: DATA statement used (Total process time):
 real time 1:45.43
 cpu time 41.15 seconds

```
258
259     * The dataset FU now contains the follow-up classified by sex and state ;
260     * The deaths are those where doDth equals exit ;
261     * The dm diagnoses are those where doDM eq exit ;
262
263     proc summary data = FU nway ;
264         class sex state A P C ;
265         var D T1 T2 Y ;
266         output out = FUtab ( drop = _FREQ_ _TYPE_ )
267             sum = ;
268     run ;
```

NOTE: There were 235985275 observations read from the data set WORK.FU.
 NOTE: The data set WORK.FUTAB has 25723 observations and 9 variables.
 NOTE: PROCEDURE SUMMARY used (Total process time):
 real time 1:11.65
 cpu time 2:37.65

```
269
270     data TTdata.FUtab ;
271     set FUtab ;
```



```

First Data Page          1
Max Obs per Page        817
Obs in First Data Page  793
Number of Data Set Repairs 0
ExtendObsCounter        YES
Filename                 E:\workdata\705093\BxC\demoDM\DATA\futab.sas7bdat
Release Created          9.0401M3
Host Created             X64_SRV12

```

Alphabetic List of Variables and Attributes

#	Variable	Type	Len	Label
*	A	Num	8	Age at FU
*	C	Num	8	
6	D	Num	8	
*	P	Num	8	Date of FU
7	T1	Num	8	
8	T2	Num	8	
10	U	Num	8	
9	Y	Num	8	
*	sex	Num	8	
*	state	Char	4	

Tabulation of the population follow-up in DK

14:39 Tuesday, January 24, 2017 2

	All				state			
					T1			
	D	T1	T2	Y	D	T1	T2	Y
All	1,166,535	24,842	342,446	114,424.04	15,668	0	0	568.89
sex								
Mand	575,622	14,247	188,969	56,682.35	9,061	0	0	321.98
Kvinde	590,913	10,595	153,477	57,741.69	6,607	0	0	246.91
U								
0	587,188	14,971	174,206	58,540.14	7,936	0	0	292.31
*	579,347	9,871	168,240	55,883.90	7,732	0	0	276.58
Date of FU								
1994	215	38	461	25.02	*	0	0	0.10
1995	62,935	3,064	34,162	5,234.42	779	0	0	22.94
1996	59,924	1,336	10,679	5,263.38	697	0	0	23.72
1997	59,276	1,295	10,510	5,284.40	749	0	0	24.32
1998	57,903	1,236	11,256	5,302.56	757	0	0	24.84
1999	58,766	1,193	12,086	5,318.76	782	0	0	25.23
2000	57,100	1,259	12,017	5,336.44	813	0	0	25.66
2001	57,737	1,165	12,338	5,356.07	782	0	0	26.07
2002	58,000	1,123	12,607	5,374.31	828	0	0	26.43
2003	57,138	1,108	14,616	5,388.37	851	0	0	26.65
2004	55,029	1,107	15,011	5,402.09	778	0	0	26.95
2005	54,329	1,133	14,927	5,417.62	810	0	0	27.21
2006	54,842	1,107	15,397	5,436.67	805	0	0	27.55
2007	55,133	1,085	16,694	5,461.82	784	0	0	27.83
2008	53,699	1,095	18,014	5,496.39	754	0	0	28.21
2009	54,190	1,053	18,803	5,530.71	787	0	0	28.52
2010	53,523	1,057	20,496	5,560.95	777	0	0	28.82
2011	51,904	977	24,150	5,590.64	712	0	0	29.09
2012	51,420	866	21,083	5,618.59	631	0	0	29.36
2013	51,546	859	15,717	5,648.79	632	0	0	29.61
2014	50,551	812	14,703	5,687.60	585	0	0	29.88
2015	51,375	874	16,719	5,688.46	572	0	0	29.90

(Continued)

	state							
	T2				Well			
	D	T1	T2	Y	D	T1	T2	Y
All	141,075	0	0	2,870.52	1,009,792	24,842	342,446	110,984.63
sex								
Mand	77,164	0	0	1,555.30	489,397	14,247	188,969	54,805.06
Kvinde	63,911	0	0	1,315.22	520,395	10,595	153,477	56,179.56
U								
0	71,554	0	0	1,485.06	507,698	14,971	174,206	56,762.77
*	69,521	0	0	1,385.46	502,094	9,871	168,240	54,221.86
Date of FU								
1994	8	0	0	0.17	204	38	461	24.75
1995	4,607	0	0	58.19	57,549	3,064	34,162	5,153.29
1996	4,922	0	0	67.10	54,305	1,336	10,679	5,172.57
1997	5,048	0	0	72.42	53,479	1,295	10,510	5,187.65
1998	5,132	0	0	78.32	52,014	1,236	11,256	5,199.40
1999	5,506	0	0	84.47	52,478	1,193	12,086	5,209.06
2000	5,564	0	0	91.06	50,723	1,259	12,017	5,219.72
2001	5,767	0	0	97.64	51,188	1,165	12,338	5,232.36
2002	6,029	0	0	104.23	51,143	1,123	12,607	5,243.64
2003	6,274	0	0	111.44	50,013	1,108	14,616	5,250.28
2004	6,217	0	0	120.02	48,034	1,107	15,011	5,255.12
2005	6,460	0	0	128.44	47,059	1,133	14,927	5,261.97
2006	6,743	0	0	137.01	47,294	1,107	15,397	5,272.11
2007	7,024	0	0	145.92	47,325	1,085	16,694	5,288.07
2008	7,010	0	0	156.43	45,935	1,095	18,014	5,311.75
2009	7,686	0	0	167.24	45,717	1,053	18,803	5,334.95
2010	7,895	0	0	179.10	44,851	1,057	20,496	5,353.02
2011	8,018	0	0	193.43	43,174	977	24,150	5,368.11
2012	8,347	0	0	208.02	42,442	866	21,083	5,381.21
2013	8,639	0	0	217.68	42,275	859	15,717	5,401.50
2014	8,948	0	0	223.92	41,018	812	14,703	5,433.80
2015	9,231	0	0	228.27	41,572	874	16,719	5,430.29

7.14 09-mkPr

Uses the follow-up data to count the persons that are alive at the 1 January 1995,...,2016, classified by sex, state (T1/T2/noDM) and age in 1-year classes. The resulting dataset, prv is classified by sex, state, age and date has only the extra variable n with the counts

```
1                                     "Program 09-mkPr.sas"
```

NOTE: Copyright (c) 2002-2012 by SAS Institute Inc., Cary, NC, USA.

NOTE: SAS (r) Proprietary Software 9.4 (TS1M3)

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NOTE: This session is executing on the X64_SRV12 platform.

NOTE: Updated analytical products:

SAS/STAT 14.1

NOTE: Additional host information:

X64_SRV12 WIN 6.2.9200 Server

NOTE: SAS initialization used:

real time 0.09 seconds

cpu time 0.10 seconds

NOTE: AUTOEXEC processing beginning; file is E:\workdata\705093\BXC\demoDM\sas\optslibs.sas.

NOTE: AUTOEXEC processing completed.

```

1      data prv ( keep = state sex age pdat ) ;
2      set TTdata.FU ;
3      * To make sure we get the last minute exiters in too ;
4      if exit ge '31DEC2015'd then exit = '02JAN2016'd ;
5      do pdat = 1995 to 2016 ;
6          if entry <= mdy(1,1,pdat) < exit then do ;
7              age = floor( (mdy(1,1,pdat)-doBth)/365.25 ) ;
8              output ;
9          end ;
10     end ;
11     run ;

```

NOTE: Missing values were generated as a result of performing an operation on missing values.

Each place is given by: (Number of times) at (Line):(Column).
 2002 at 7:15 2002 at 7:36 2002 at 7:43

NOTE: There were 8191903 observations read from the data set TTDATA.FU.

NOTE: The data set WORK.PRIV has 120001859 observations and 4 variables.

NOTE: DATA statement used (Total process time):

real time	28.21 seconds
cpu time	21.85 seconds

```

12
13     proc summary data = prv nway ;
14         class pdat state sex age ;
15         output out = TTdata.priv ( keep = pdat state sex age _freq_
16                                 rename = ( _freq_ = n ) ) ;
17     run ;

```

NOTE: There were 120001859 observations read from the data set WORK.PRIV.

NOTE: The data set TTDATA.PRIV has 13545 observations and 5 variables.

NOTE: PROCEDURE SUMMARY used (Total process time):

real time	17.36 seconds
cpu time	33.96 seconds

```

18
19     proc contents data = TTdata.priv ; run ;

```

NOTE: PROCEDURE CONTENTS used (Total process time):

real time	0.03 seconds
cpu time	0.03 seconds

NOTE: The PROCEDURE CONTENTS printed page 1.

```

20
21     proc tabulate data = TTdata.priv ( rename = ( n = x ) ) missing noseps ;
22         class pdat state ;
23         var x ;
24         table pdat,
25             ( x = ' ' * sum = 'no. persons alive' * f=comma9.
26               x = ' ' * n = 'no. records in data' * f=comma6. ) * ( all state )
27             / rts = 6 ;
28     run ;

```

NOTE: There were 13545 observations read from the data set TTDATA.PRIV.

NOTE: The PROCEDURE TABULATE printed page 2.

NOTE: PROCEDURE TABULATE used (Total process time):

real time	0.03 seconds
cpu time	0.03 seconds

NOTE: SAS Institute Inc., SAS Campus Drive, Cary, NC USA 27513-2414

NOTE: The SAS System used:

real time	45.82 seconds
-----------	---------------

cpu time 56.04 seconds

7.14.1 09-mkPr.lst

The SAS System 14:51 Tuesday, January 24, 2017 1

The CONTENTS Procedure

Data Set Name	TTDATA.PRV	Observations	13545
Member Type	DATA	Variables	5
Engine	V9	Indexes	0
Created	24/01/2017 14:52:19	Observation Length	40
Last Modified	24/01/2017 14:52:19	Deleted Observations	0
Protection		Compressed	NO
Data Set Type		Sorted	NO
Label			
Data Representation	WINDOWS_64		
Encoding	wlatin1 Western (Windows)		

Engine/Host Dependent Information

Data Set Page Size	65536
Number of Data Set Pages	9
First Data Page	1
Max Obs per Page	1632
Obs in First Data Page	1595
Number of Data Set Repairs	0
ExtendObsCounter	YES
Filename	E:\workdata\705093\BxC\demoDM\DATA\prv.sas7bdat
Release Created	9.0401M3
Host Created	X64_SRV12

Alphabetic List of Variables and Attributes

#	Variable	Type	Len
4	age	Num	8
5	n	Num	8
1	pdat	Num	8
3	sex	Num	8
2	state	Char	4

The SAS System 14:51 Tuesday, January 24, 2017 2

	no. persons alive				no. records in data			
	state				state			
	All	T1	T2	Well	All	T1	T2	Well
pdat								
1995	5,222,317	21,263	34,435	5,166,619	605	192	198	215
1996	5,253,550	23,600	64,444	5,165,506	612	194	203	215
1997	5,276,124	24,251	70,243	5,181,630	609	193	200	216
1998	5,294,854	24,792	75,672	5,194,390	611	194	199	218
1999	5,312,778	25,254	81,831	5,205,693	613	194	201	218
2000	5,328,484	25,668	88,379	5,214,437	608	194	199	215
2001	5,348,316	26,095	94,799	5,227,422	608	195	199	214
2002	5,367,470	26,459	101,325	5,239,686	612	196	200	216
2003	5,383,257	26,726	107,841	5,248,690	612	196	200	216
2004	5,396,881	26,958	116,142	5,253,781	618	200	202	216
2005	5,411,220	27,261	124,841	5,259,118	619	199	202	218
2006	5,428,440	27,564	133,148	5,267,728	615	197	201	217
2007	5,449,469	27,865	141,703	5,279,901	614	198	201	215
2008	5,480,236	28,175	151,253	5,300,808	614	198	200	216

2009	5,518,953	28,533	162,218	5,328,202	613	197	199	217
2010	5,548,845	28,816	173,245	5,346,784	614	198	200	216
2011	5,579,763	29,079	185,682	5,365,002	617	197	202	218
2012	5,608,449	29,340	201,528	5,377,581	623	199	204	220
2013	5,637,470	29,561	214,139	5,393,770	624	197	205	222
2014	5,670,648	29,784	221,087	5,419,777	625	197	204	224
2015	5,713,553	30,005	226,703	5,456,845	628	197	205	226
2016	5,768,780	30,278	234,098	5,504,404	631	198	206	227