

Analysis of base-line follow-up data

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Outline

- ▶ Observations / measurements for each individual, i :
 - ▶ Baseline y_{0i}
 - ▶ Follow-up y_{1i}
 - ▶ treatment group
 - ▶ covariates
- ▶ Topic of interest:
 - ▶ How much is the change from baseline to follow-up
 - ▶ How much does this depend on treatment / covariates

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The change from baseline to follow-up

- ▶ Baseline is subject to random error
- ▶ If the random error at baseline is large **positive**:
 - ▶ baseline measurement “artificially” **large**
 - ▶ **change** to follow-up **smaller**
- ▶ If the random error at baseline is large **negative**:
 - ▶ baseline measurement “artificially” **small**
 - ▶ **change** to follow-up **larger**
- ▶ $\Rightarrow$ the change depends on the baseline **measurement**.

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Example from Vickers et al.

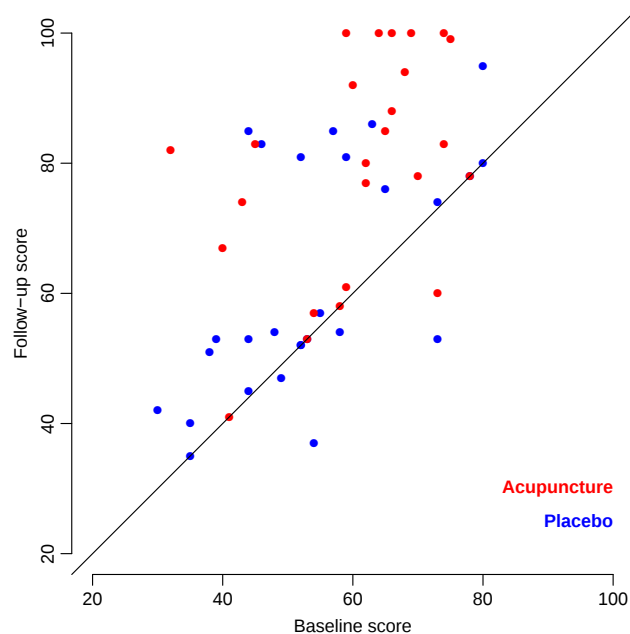
```
> library( Epi )
> library( foreign )
> acp <- read.dta( "./data/sportsmen.dta" )[, -4]
> names( acp ) <- c("bl", "fu", "gr")
> acp$gr <- factor( acp$gr, labels=c("Placebo", "Acupuncture") )
> str( acp )
'data.frame': 54 obs. of 3 variables:
 $ bl: num  59 53 46 38 52 63 30 73 44 48 ...
 $ fu: num  81 53 83 51 81 86 42 74 45 54 ...
 $ gr: Factor w/ 2 levels "Placebo","Acupuncture": 1 1 1 1 1 1 1 1 1 1 ...
> head( acp )
   bl fu   gr
1  59 81 Placebo
2  53 53 Placebo
3  46 83 Placebo
4  38 51 Placebo
5  52 81 Placebo
6  63 86 Placebo
```

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Example data from
Vickers *et al.*:

Randomization to
acupuncture /
placebo

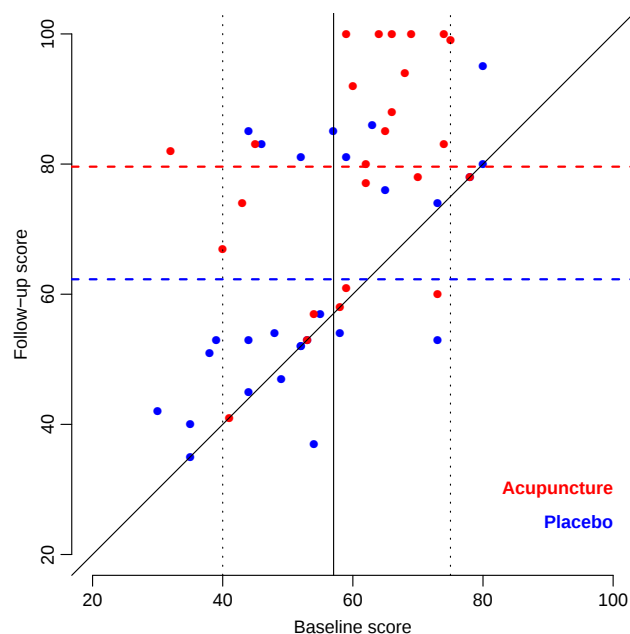
Outcome:
Pain/function rating
of shoulder pain
(0–100).



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Follow-up analysis

If the study is
randomized,
analysis of follow-up
is in principle
unbiased, because
baseline distribution
is the same in
randomization
groups.



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Analysis of follow-up

```
> # Follow-up
> fu <- with( acp, tapply( fu , gr, mean ) )
> c( fu, diff( fu ) )
      Placebo Acupuncture Acupuncture
      62.2963    79.6000    17.3037
> mf <- lm( fu ~ gr, data=acp )
> round( ci.lin( mf ), 3 )
              Estimate StdErr      z P    2.5% 97.5%
(Intercept)    62.296   3.378 18.440 0 55.675 68.918
grAcupuncture   17.304   4.872  3.551 0  7.754 26.853
```

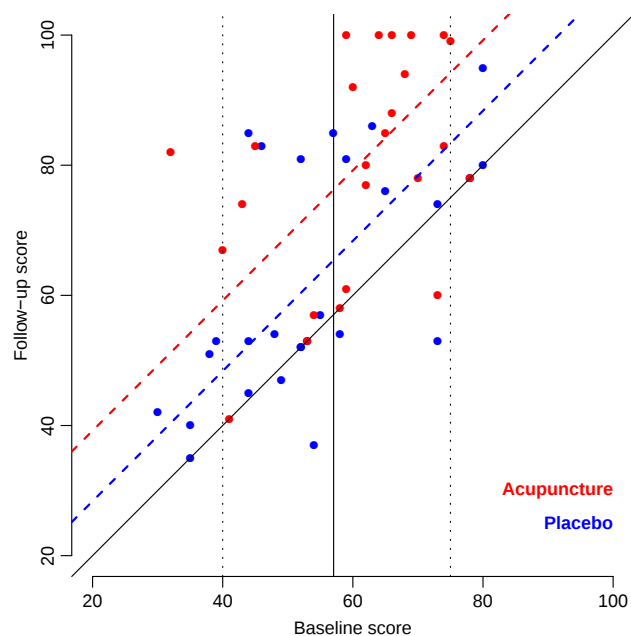
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Analysis of change scores

$$y_1 - y_0$$

If not randomized
this is also biased by
baseline differences

The change scores
are found as the
distance to the 45°
line.



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Analysis of change scores

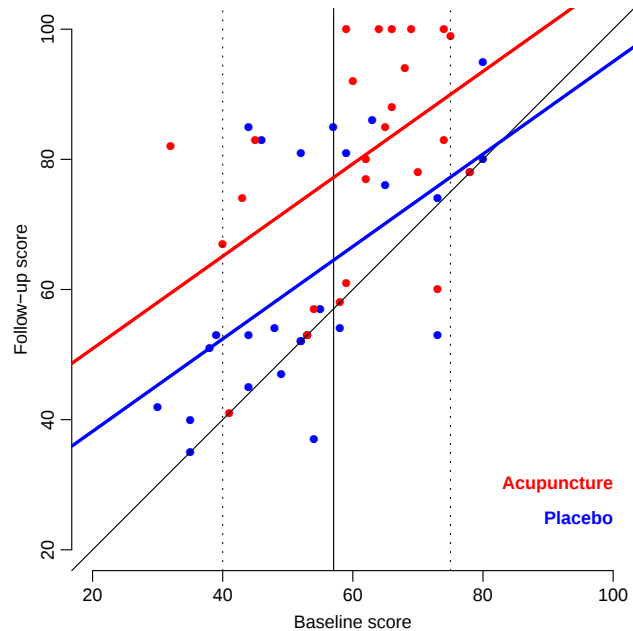
```
> df <- with( acp, tapply( fu-bl, gr, mean ) )
> c( df, diff( df ) )
      Placebo Acupuncture Acupuncture
      8.37037    19.20000    10.82963
> md <- lm( fu-bl ~ gr, data=acp )
> round( ci.lin( md ), 3 )
              Estimate StdErr      z      P    2.5% 97.5%
(Intercept)     8.37    2.948  2.839 0.005  2.592 14.148
grAcupuncture   10.83    4.252  2.547 0.011  2.497 19.163
```

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Conditioning on baseline $y_1|y_0$

Accounts for possible imbalances in baseline distribution.

Separates treatment effect and baseline effect on outcome.



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Conditioning on baseline

```
> mc <- lm( fu ~ bl + gr, data=acp )
> round( ci.lin( mc ), 4 )

              Estimate StdErr      z      P    2.5%   97.5%
(Intercept)   23.9973  9.1092  2.6344 0.0084  6.1435 41.8511
bl             0.7102  0.1602  4.4323 0.0000  0.3962  1.0243
grAcupuncture  12.7057  4.2857  2.9647 0.0030  4.3059 21.1056
```

- ▶ $y_{i1} = M + B y_{i0} + D_g$
- ▶ treatment effect (D_g) is 12.7 points:
 - ▶ change in placebo:

$$M + (B - 1)y_{i0} = 23.997 - 0.290 \times y_{0i}$$
 - ▶ change in acupuncture:

$$M + (B - 1)y_{i0} + D_g = 23.997 - 0.290 \times y_{0i} + 12.706$$
- ▶ change from baseline to FU depend on baseline
- ▶ treatment effect is **difference** in changes

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Comparing the three approaches

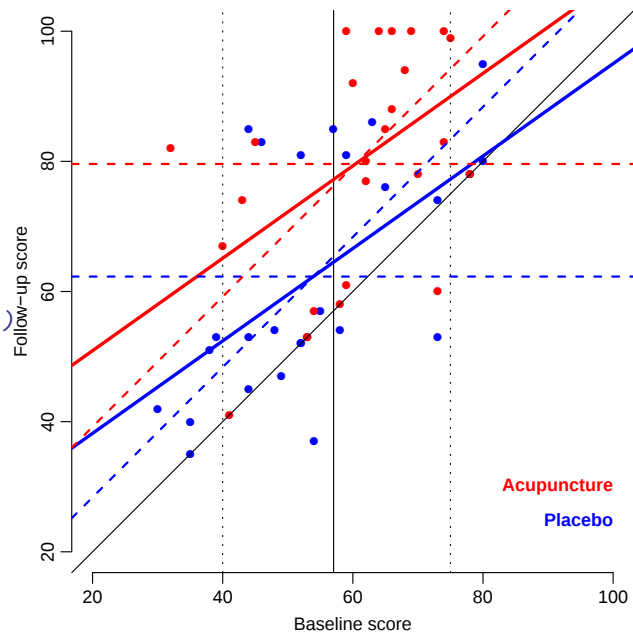
```
> cmp.cf <- rbind( ci.lin( mf, subset="Acu" ),
+                 ci.lin( md, subset="Acu" ),
+                 ci.lin( mc, subset="Acu" ) )
> rownames( cmp.cf ) <- c("FU", "Chg-sc", "Cond")
> round( cmp.cf, 4 )

              Estimate StdErr      z      P    2.5%   97.5%
FU           17.3037  4.8723  3.5515 0.0004  7.7542 26.8532
Chg-sc       10.8296  4.2516  2.5472 0.0109  2.4966 19.1627
Cond         12.7057  4.2857  2.9647 0.0030  4.3059 21.1056
```

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Comparing the three approaches

```
> round( cmp.cf[,c(1,2,4)], 3 )
      Estimate StdErr      P
FU      17.304   4.872 0.000
Chg-sc   10.830   4.252 0.011
Cond     12.706   4.286 0.003
```



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Changes from baseline to FU

```
> ( cf <- coef(mc) )
      (Intercept)          bl grAcupuncture
      23.9973054      0.7102148      12.7057205
> ( mb <- mean( acp$bl ) )
[1] 57.04259
> y0 <- c(40,mb,75)
> p.ch <- cf[1] - (cf[2]-1)*y0
> a.ch <- cf[1] - (cf[2]-1)*y0 + cf[3]
> chg <- cbind( p.ch, a.ch, a.ch-p.ch )
> colnames( chg ) <- c( levels( acp$gr ), "Diff" )
> rownames( chg ) <- round(y0,2)
> round( chg, 2 )

      Placebo Acupuncture Diff
40      35.59      48.29 12.71
57.04    40.53      53.23 12.71
75      45.73      58.44 12.71
```

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Constant treatment effect?

- Depends on baseline & treatment
- ... but treatment **difference** in change does not
- But **does** the treatment effect depend on baseline?

```
> mi <- lm( fu ~ gr + gr:bl, data=acp )
> round( ci.lin( mi ), 4 )

      Estimate StdErr      z      P      2.5% 97.5%
(Intercept)   20.3488 11.7437 1.7327 0.0831  -2.6685 43.3661
grAcupuncture  22.1307 19.4070 1.1403 0.2541 -15.9062 60.1677
grPlacebo:bl    0.7779  0.2110 3.6865 0.0002   0.3643  1.1914
grAcupuncture:bl 0.6146  0.2509 2.4498 0.0143   0.1229  1.1063
> anova( mi, mc )
Analysis of Variance Table

Model 1: fu ~ gr + gr:bl
Model 2: fu ~ bl + gr
      Res.Df  RSS Df Sum of Sq    F Pr(>F)
1         48 10942
2         49 10998 -1    -56.565 0.2481 0.6207
```

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