Examples of focused model comparison: skew-normal models

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Abstract

The following example (from ?) illustrates how focused model comparison can be performed using the **fic** package in a situation where:

- a novel class of models is defined and fitted by custom R functions
- a simple model is extended in two different directions to define the models being compared

Keywords: models.

An outcome y_i and a covariate x_i are observed for individuals i = 1, ..., n. Four different models are compared: two normal models with a constant variance, one without (1) and one with (2) a linear regression term, and two "skew-normal" models without (3) and with (4) the linear regression term. The skew-normal model is defined by an error term $\sigma \epsilon_i$, where the ϵ_i are independently distributed with a density $f(u|\lambda) = \lambda \Phi(u)^{\lambda-1} \phi(u)$. All four models are nested in the "wide" model (4).

- (1) $y_i \sim N(\beta_0, \sigma^2)$
- (2) $y_i \sim N(\beta_0 + \beta_1 x_i, \sigma^2)$
- (3) $y_i = \beta_0 + \sigma \epsilon_i$
- (4) $y_i = \beta_0 + \beta_1 x_i + \sigma \epsilon_i$

1. Fitting the models

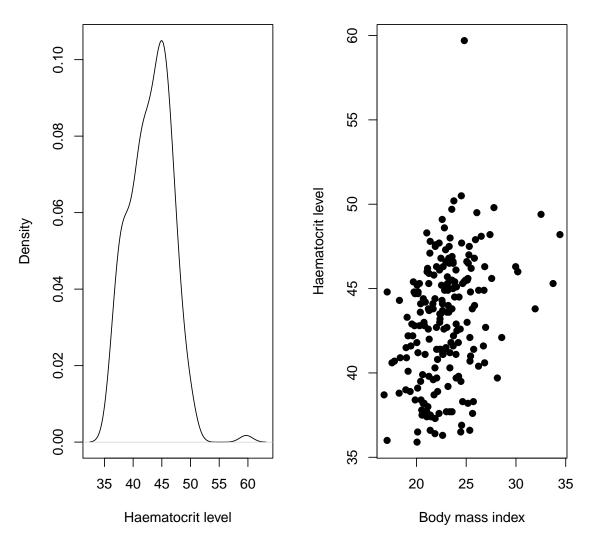
To implement this class of models in R, firstly we define the log density function of the general skew-normal model with mean, scale and skewness parameters μ , σ , λ indicated by arguments mean, sigma and lambda. This defines the distribution of y_i in model (3) with mean $\mu_i = \beta_0$, and in (4) with $\mu_i = \beta_0 + \beta_1 x_i$. Models (1) and (2) are defined by setting $\lambda = 1$ in models (3) and (4) respectively.

```
ldsnorm <- function(x, mean, sd, lambda){
    log(lambda) + (lambda-1)*pnorm(x, mean, sd, log=TRUE) +
        dnorm(x, mean, sd, log=TRUE)
}</pre>
```

The models are fitted to data from the Australian Institute of Sports (Cook and Weisberg, 1994 REF), available as **ais** from the **sn** package REF. The outcome y_i is haematocrit level Hc, and the covariate is body mass index BMI. Figure ?? illustrates the skewed distribution of the outcome and a mild association between the variables.

```
if (!require("sn"))
    stop("The `sn` package should be installed to run code in this vignette")
data(ais)
par(mfrow=c(1,2))
plot(density(ais$Hc), xlab="Haematocrit level")
plot(ais$BMI, ais$Hc, pch=19,
    xlab="Body mass index", ylab="Haematocrit level")
```

density.default(x = ais\$Hc)



The following defines the minus log likelihood for these data as a function of four parameters

 $\beta_0, \beta_1, \sigma \text{ and } \lambda.$

```
mloglik <- function(b0, b1, sd, lambda){
    -sum(ldsnorm(ais$Hc, b0 + b1*ais$BMI, sd, lambda))
}</pre>
```

Then to obtain the maximum likelihood estimates under models 1 to 4, mloglik is rewritten as a function of a single vector par, containing either 2, 3 or 4 parameters to be minimised over, depending on the model. Note here that models 1 and 3 have β_2 fixed at 0, and models 1 and 2 have λ fixed at 1. The positive parameters σ and λ will be estimated by unconstrained maximisation on the log scale. These functions can be passed to the nlm function for optimisation.

```
fn1 <- function(par) mloglik(par[1], 0, exp(par[2]), 1)
fn2 <- function(par) mloglik(par[1], par[2], exp(par[3]), 1)
fn3 <- function(par) mloglik(par[1], 0, exp(par[2]), exp(par[3]))
fn4 <- function(par) mloglik(par[1], par[2], exp(par[3]), exp(par[4]))</pre>
```

nlm also requires plausible initial values for the parameters. A vector of these (ini) is obtained as follows by fitting model (2) with lm and extracting the coefficients with coef (for β_0 and β_1) and the residual standard deviation for $\log(\sigma)$. An initial value of 0 is used for $\log(\lambda)$.

The appropriate objective function (fn1-fn4) is then optimised for each model, starting from the given initial values².

```
opt1 <- nlm(fn1, ini[c("beta0","logsigma")], hessian=TRUE)
opt2 <- nlm(fn2, ini[c("beta0","beta1","logsigma")], hessian=TRUE)
opt3 <- nlm(fn3, ini[c("beta0","logsigma","loglambda")], hessian=TRUE)
opt4 <- nlm(fn4, ini, hessian=TRUE)</pre>
```

Finally, the information required by the fic function (the estimates and covariance matrices) is extracted from the nlm results and arranged into a list, for each of the four models.

¹Note these are the exact maximum likelihood estimates for model 2. The added value of nlm in fitting models 1 and 2, compared to simply using lm, is to conveniently provide the covariance matrix at the maximum likelihood estimates, which is required for focused model comparison

²A warning message of NA/Inf replaced by maximum positive value can be ignored and is the result of nlm trying out extreme and implausible values on the way to finding the maximum likelihood.

```
mod1 <- list(est=opt1$estimate, vcov=solve(opt1$hessian) )
mod2 <- list(est=opt2$estimate, vcov=solve(opt2$hessian) )
mod3 <- list(est=opt3$estimate, vcov=solve(opt3$hessian) )
mod4 <- list(est=opt4$estimate, vcov=solve(opt4$hessian) )</pre>
```

dsn also provided with sn package

2. Focused model comparison

We now perform a focused comparison of the four models. Two alternative focuses are investigated: the mean and median outcome at a covariate value of interest. Expressions for the mean and median of the skew normal, in the parameterisation used here, are given by ? and encoded in the following R functions:

```
mean_snorm <- function(mu, sigma, lambda){
    f <- function(u){u*exp(ldsnorm(u, 0, 1, lambda))}
    mu + sigma * integrate(f, -Inf, Inf)$value
}
median_snorm <- function(mu, sigma, lambda){
    mu + sigma * qnorm(0.5^(1/lambda))
}</pre>
```

As described in the main package vignette (Section ??) the focus function supplied to fic should have arguments defined by a vector par of parameters of the biggest model (in this case the four-parameter model 4), and a matrix of covariate values X, and return the corresponding focus quantity. In this example, the two focus functions are

```
focus1 <- function(par, X){
    mean_snorm(mu = X %*% par[1:2], sigma=exp(par[3]), lambda=exp(par[4]))
}
focus2 <- function(par, X){
    median_snorm(mu = X %*% par[1:2], sigma = exp(par[3]), lambda = exp(par[4]))
}</pre>
```

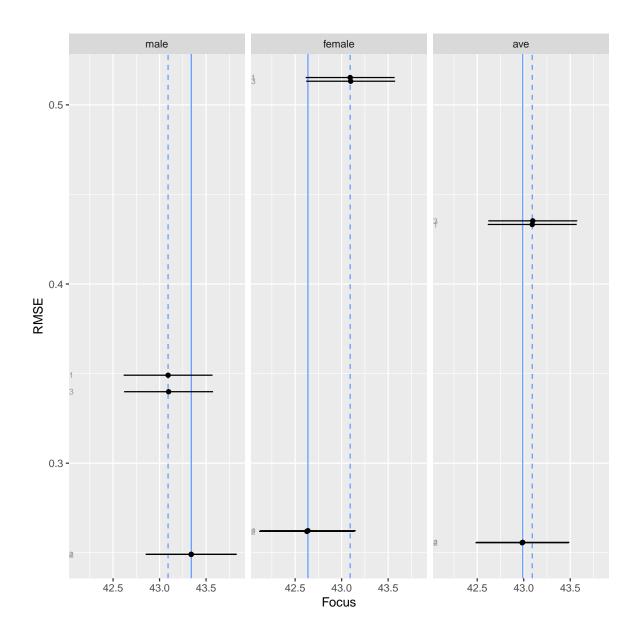
The inds matrix, required by fic is now constructed. Recall this indicates which parameters (columns) are included in each of the models (rows) being compared. The narrow model is in the first row, and the wide model in the last. The functions fns required to extract the estimates and covariance matrix from the fitted model objects are then defined.

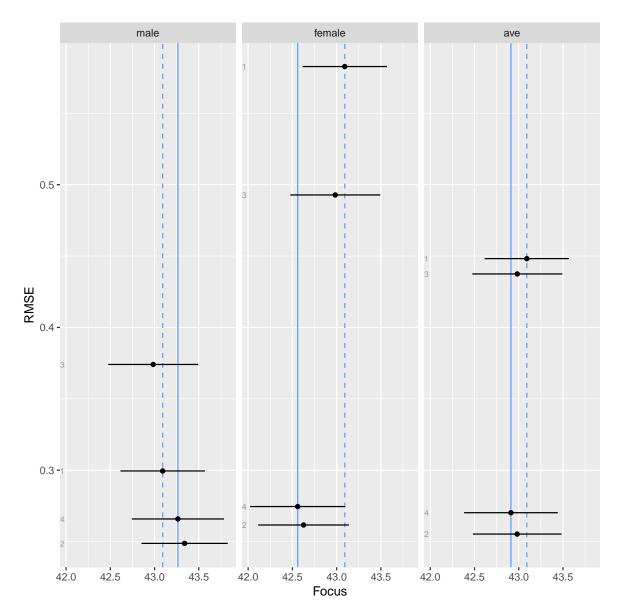
Finally fic is called to compare the four models for focuses defined by the mean and median for average men and women, with covariate values defined by med.bmi

TODO do we really need an intercept? Shouldn't

```
c(1,1,1,1))
fns <- list(coef=function(x)x$est,</pre>
           vcov=function(x)x$vcov,
           nobs=function(x)nrow(ais))
med.bmi <- rbind(male=c(1, 23.56),</pre>
                female=c(1, 21.82))
focus0 <- function(par){X %*% par[1:2]}</pre>
fic(mod2, inds=rbind(c(1,0,1),c(1,1,1)), fns=fns, focus=focus0, X=med.bmi, FIC=TRUE,
   sub=list(mod1, mod2))
##
      vals mods rmse rmse.adj
                                bias
                                        se
                                             FIC focus
             1 0.347
## 1
      male
                         0.347 -0.248 0.244 12.39 43.1
## 4
      male
              2 0.249
                         0.249 0.000 0.249 1.07
                                                  43.3
## 2 female
             1 0.525
                         0.525  0.466  0.244  43.80  43.1
             2 0.262
                        0.262 0.000 0.262 3.79 42.6
## 5 female
                        0.439 0.365 0.244 26.99 43.1
## 3
             1 0.439
       ave
## 6
              2 0.256
                        0.256 0.000 0.256 1.32 43.0
       ave
fmean <- fic(mod4, inds=inds, fns=fns, focus=focus1, X=med.bmi, FIC=TRUE,</pre>
   sub=list(mod1, mod2, mod3, mod4))
fmean
##
       vals mods rmse rmse.adj
                                 bias
                                              FIC focus
                                         se
## 1
       male
             1 0.349
                          0.349 -0.250 0.244 12.61
                                                  43.1
## 4
       male
               2 0.249
                          0.249 0.000 0.249 1.05 43.3
             3 0.340
## 7
                        0.340 -0.237 0.244 11.85 43.1
       male
## 10
       male 4 0.249
                       0.249 0.000 0.249 1.05
                                                  43.3
## 2 female 1 0.515
                       0.515 0.454 0.244 41.62 43.1
     female
               2 0.262
                         0.262 0.000 0.262 3.73 42.6
## 5
## 8 female 3 0.513
                       0.513 0.452 0.244 43.09 43.1
             4 0.262
## 11 female
                        0.262 0.000 0.262 3.78 42.6
## 3
             1 0.433
                        0.433 0.358 0.244 26.02 43.1
        ave
             2 0.256
## 6
                        0.256 0.000 0.256 1.30 43.0
        ave
## 9
               3 0.435
                          0.435 0.360 0.244 26.38 43.1
        ave
## 12
               4 0.256
                          0.256 0.000 0.256 1.32 43.0
        ave
fmed <- fic(mod4, inds=inds, fns=fns, focus=focus2, X=med.bmi,</pre>
   sub=list(mod1, mod2, mod3, mod4))
fmed
##
       vals mods rmse rmse.adj
                                bias
                                         se focus
       male
             1 0.299
                          0.299 -0.174 0.244 43.1
## 1
## 4 male 2 0.242 0.249 0.000 0.249 43.3
```

```
## 7
      male 3 0.374
                    0.374 -0.269 0.260 43.0
## 10
      male 4 0.266
                      0.266 0.000 0.266 43.3
## 2 female 1 0.583
                      0.583 0.529 0.244 43.1
## 5 female 2 0.256
                    0.262 0.000 0.262 42.6
                     0.493 0.419 0.260 43.0
## 8 female 3 0.493
## 11 female 4 0.275
                    0.275 0.000 0.275 42.6
       ave 1 0.448
## 3
                    0.448 0.376 0.244 43.1
       ave 2 0.249 0.255 0.000 0.255 43.0
## 6
## 9
      ave 3 0.437 0.437 0.352 0.260 43.0
                    0.270 0.000 0.270 42.9
## 12 ave 4 0.270
library(ggplot2)
ggplot_fic(fmean)
```





TODO name the models TODO shrink the plots TODO if nobs is supplied then set FIC true mean: we need the covariate, but not necessarily the skewness as well. these conclusions don't depend on cov "For estimating the mean there is no award in involving the Îż aspects of the data, as the added complexity does not alter the large-sample performance of estimators." fic values close to book, not exactly, for mean when wide excludes skew median: we need the covariate again, but skew model has worse rmse makes sense for measures of central tendency. median more robust to skew? todo check vs book, hmm book selects wide model for median refer to algebraic results in book

Very different FICs from book for median can we check the omegas against book? TODO teach people to make a S3 method for their new class TODO can't refit submodels - only works for standard cov selection problems