

Survival Ensembles

Torsten Hothorn^{1,*}, Peter Bühlmann², Sandrine Dudoit³,
Annette Molinaro⁴ and Mark J. van der Laan³

¹Institut für Medizininformatik, Biometrie und Epidemiologie
Friedrich-Alexander-Universität Erlangen-Nürnberg
Waldstraße 6, D-91054 Erlangen, Germany
Tel: ++49-9131-8522707
Fax: ++49-9131-8525740
Torsten.Hothorn@R-project.org

²Seminar für Statistik, ETH Zürich, CH-8032 Zürich, Switzerland
buhlmann@stat.math.ethz.ch

³Division of Biostatistics, University of California, Berkeley
140 Earl Warren Hall, #7360, Berkeley, CA 94720-7360, USA
sandrine@stat.Berkeley.EDU
laan@stat.Berkeley.EDU

⁴Division of Biostatistics, Epidemiology and Public Health
Yale University School of Medicine, 206 LEPH
60 College Street PO Box 208034, New Haven CT 06520-8034
annette.molinaro@yale.edu

1 Illustrations and Applications

This document reproduces the data analyses presented in [Hothorn et al. \(2006\)](#).
For a description of the theory behind applications shown here we refer to the
original manuscript.

1.1 Acute myeloid leukemia

Data preprocessing Compute IPC weights, define risk score and set up
learning sample:

```
R> AMLw <- IPCweights(Surv(clinical$time, clinical$event))  
R> risk <- rep(0, nrow(clinical))  
R> rlev <- levels(clinical[, "Cytogenetic.group"])
```

```

R> risk[clinical[, "Cytogenetic.group"] %in% rlev[c(7,
8, 4)]] <- "low"
R> risk[clinical[, "Cytogenetic.group"] %in% rlev[c(5,
9)]] <- "intermediate"
R> risk[clinical[, "Cytogenetic.group"] %in% rlev[-c(4,
5, 7, 8, 9)]] <- "high"
R> risk <- as.factor(risk)
R> AMLlearn <- cbind(clinical[, c("time", "Sex",
"Age", "LDH", "WBC", "FLT3.aberration.", "MLL.PTD",
"Tx.Group.")], risk = risk, iexpressions[,
colnames(iexpressions) %in% selgenes[["Clone.ID"]]])
R> cc <- complete.cases(AMLlearn)
R> AMLlearn <- AMLlearn[AMLw > 0 & cc, ]
R> AMLw <- AMLw[AMLw > 0 & cc]

```

Model fitting Fit random forest for censored data

```

R> ctrl <- cforest_control(mincriterion = 0.1, mtry = 5,
minsplitt = 5, ntree = 250)
R> AMLrf <- cforest(I(log(time)) ~ ., data = AMLlearn,
control = ctrl, weights = AMLw)

```

and L_2 Boosting for censored data

```

R> AML12b <- glmboost(I(log(time)) ~ ., data = AMLlearn,
weights = AMLw, control = boost_control(mstop = 5000))

```

Compute fitted values

```

R> AML12b <- AML12b[mstop(aic)]
R> cAML <- coef(AML12b)
R> cAML[abs(cAML) > 0]

```

	Age	WBC	
(Intercept)	0.03094981	0.00854937	-0.00364371
MLL.PTDyes	Tx.Group.AUTO	Tx.Group.IC	
-0.50709786	0.90185340	0.04037578	
Tx.Group.Ind	riskintermediate	`IMAGE:145643`	
-1.86134842	0.11825619	0.19788355	
`IMAGE:2542486`	`IMAGE:345601`	`IMAGE:377560`	
0.00442375	0.02935101	0.11000322	
`IMAGE:428782`	`IMAGE:2043415`	`IMAGE:1584563`	
0.01010658	0.05911671	-0.17883619	
`IMAGE:347035`	`IMAGE:262695`	`IMAGE:950479`	
-0.03307600	0.00080156	0.09049309	
`IMAGE:898305`	`IMAGE:1472689`	`IMAGE:150702`	
0.00523016	0.03498572	0.01367553	
`IMAGE:1526826`	`IMAGE:66507`	`IMAGE:786302`	
-0.01805326	0.00399127	0.08941300	
`IMAGE:243614`	`IMAGE:417884`	`IMAGE:1592006`	
-0.05776062	-0.04890054	-0.02269622	
`IMAGE:1917063`	`IMAGE:884333`	`IMAGE:133273`	

```
R> plot(aic <- AIC(AML12b))
```

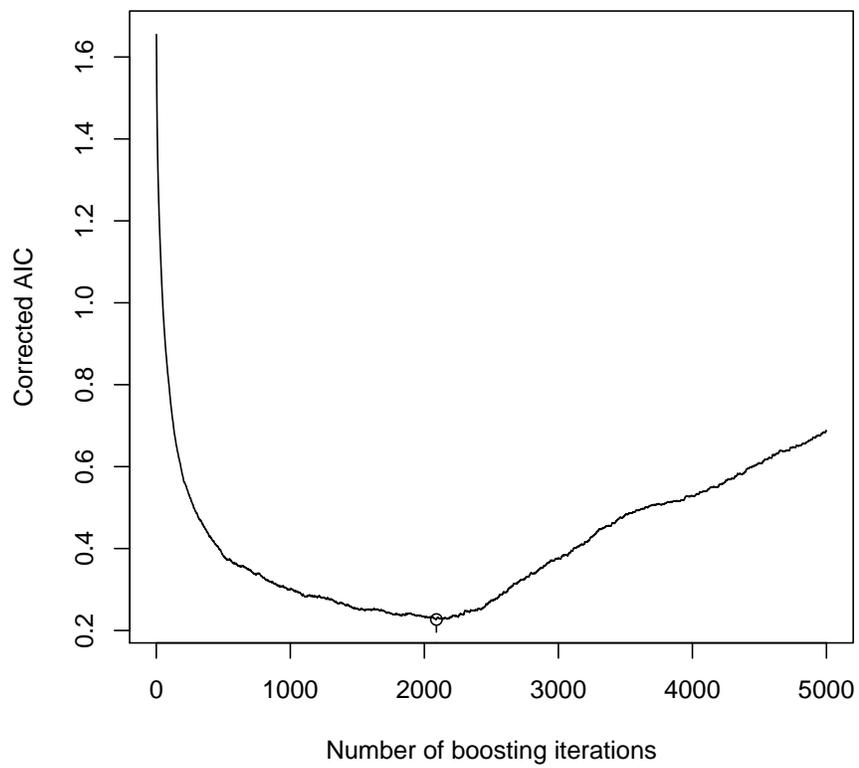


Figure 1: AIC criterion for AML data.

```

-0.06536720      0.04189990      0.06594787
`IMAGE:950888` `IMAGE:809533` `IMAGE:49389`
  0.02027810      -0.15986981      0.06352703
`IMAGE:789357` `IMAGE:142139` `IMAGE:1558053`
 -0.01252187      0.00089307      0.07795515
`IMAGE:856174` `IMAGE:504421` `IMAGE:435036`
  0.01115234      0.06861766      0.06094620
`IMAGE:491751` `IMAGE:782835` `IMAGE:52930`
  0.04336285      -0.17924185      -0.03503330
`IMAGE:2545705` `IMAGE:756405` `IMAGE:502664`
 -0.09886616      0.07713650      0.03620466
`IMAGE:129032` `IMAGE:1610168` `IMAGE:327676`
 -0.31322459      0.01260374      -0.02117310
`IMAGE:69002` `IMAGE:121551` `IMAGE:2019101`
 -0.41671336      -0.08107446      -0.06531175
`IMAGE:1456160` `IMAGE:430318` `IMAGE:2566064`
 -0.10208684      -0.07297586      0.06126683
`IMAGE:74537` `IMAGE:1606557` `IMAGE:306812`
  0.04523784      0.14243526      0.03504441
`IMAGE:565083` `IMAGE:843028` `IMAGE:68794`
  0.29555347      0.05619983      0.23722775
`IMAGE:488505` `IMAGE:167205` `IMAGE:291756`
  0.33464829      0.00217136      0.04973319
`IMAGE:810801` `IMAGE:1702742` `IMAGE:380462`
  0.08725523      -0.04428190      -0.13182519
`IMAGE:154472` `IMAGE:302540` `IMAGE:135221`
 -0.24723347      0.17175129      -0.01972168
`IMAGE:1567220` `IMAGE:594630`
  0.02473376      -0.07396882

```

```

R> AMLprf <- predict(AMLrfl, newdata = AMLlearn)
R> AMLpb <- predict(AMLl2b, newdata = AMLlearn)

```

1.2 Node-positive breast cancer

Data preprocessing Compute IPC weights and set up learning sample:

```

R> data("GBSG2", package = "ipred")
R> GBSG2w <- IPCweights(Surv(GBSG2$time, GBSG2$cens))
R> GBSG2learn <- cbind(GBSG2[, -which(names(GBSG2) %in%
  c("time", "cens"))], ltime = log(GBSG2$time))
R> n <- nrow(GBSG2learn)

```

Model fitting

```

R> LMmod <- lm(ltime ~ ., data = GBSG2learn, weights = GBSG2w)
R> LMerisk <- sum((GBSG2learn$ltime - predict(LMmod))^2 *
  GBSG2w)/n
R> TRmod <- rpart(ltime ~ ., data = GBSG2learn, weights = GBSG2w)
R> TRerisk <- sum((GBSG2learn$ltime - predict(TRmod))^2 *
  GBSG2w)/n
R> ctrl <- cforest_control(mincriterion = qnorm(0.95),

```

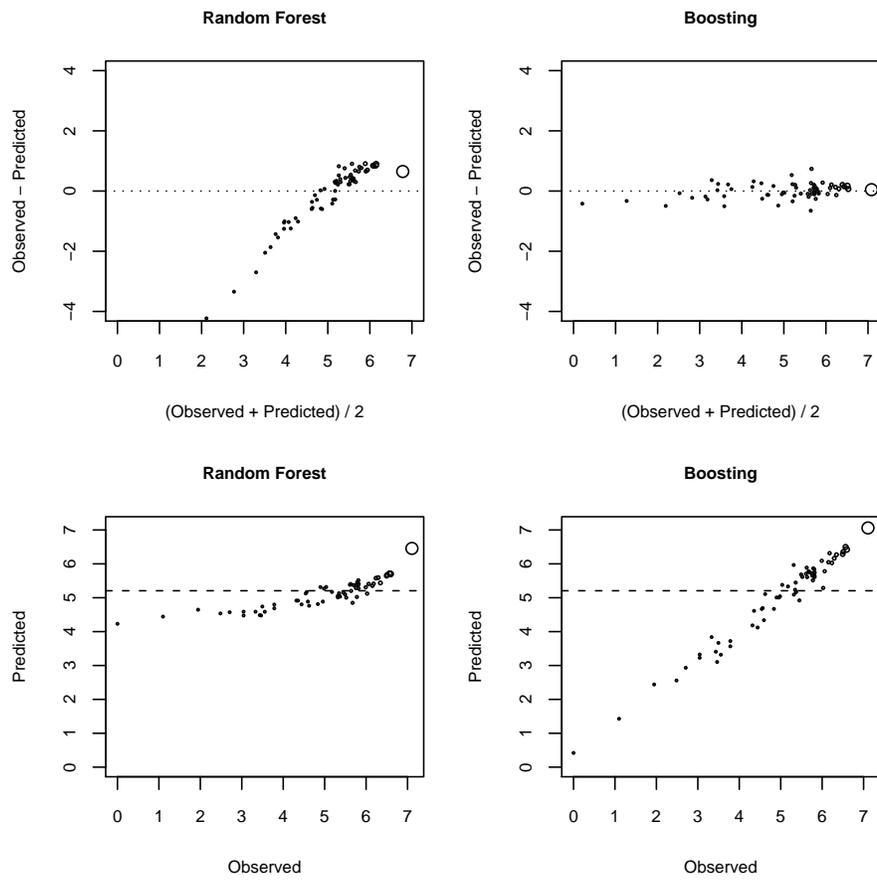


Figure 2: AML data: Reproduction of Figure 1.

```
      mtry = 5, minsplit = 5, ntree = 100)
R> RFmod <- cforest(ltime ~ ., data = GBSG2learn,
  weights = GBSG2w, control = ctrl)
R> L2Bmod <- glmboost(ltime ~ ., data = GBSG2learn,
  weights = GBSG2w, control = boost_control(mstop = 250))
R> L2BHubermod <- glmboost(ltime ~ ., data = GBSG2learn,
  weights = GBSG2w, family = Huber(d = log(2)))
```

Compute fitted values:

```
R> GBSG2Hp <- predict(L2BHubermod, newdata = GBSG2learn)
R> L2Berisk <- sum((GBSG2learn$ltime - predict(L2Bmod,
  newdata = GBSG2learn))^2 * GBSG2w)/n
R> RFerisk <- sum((GBSG2learn$ltime - predict(RFmod,
  newdata = GBSG2learn))^2 * GBSG2w)/n
```

```
R> plot(aic <- AIC(L2Bmod))
```

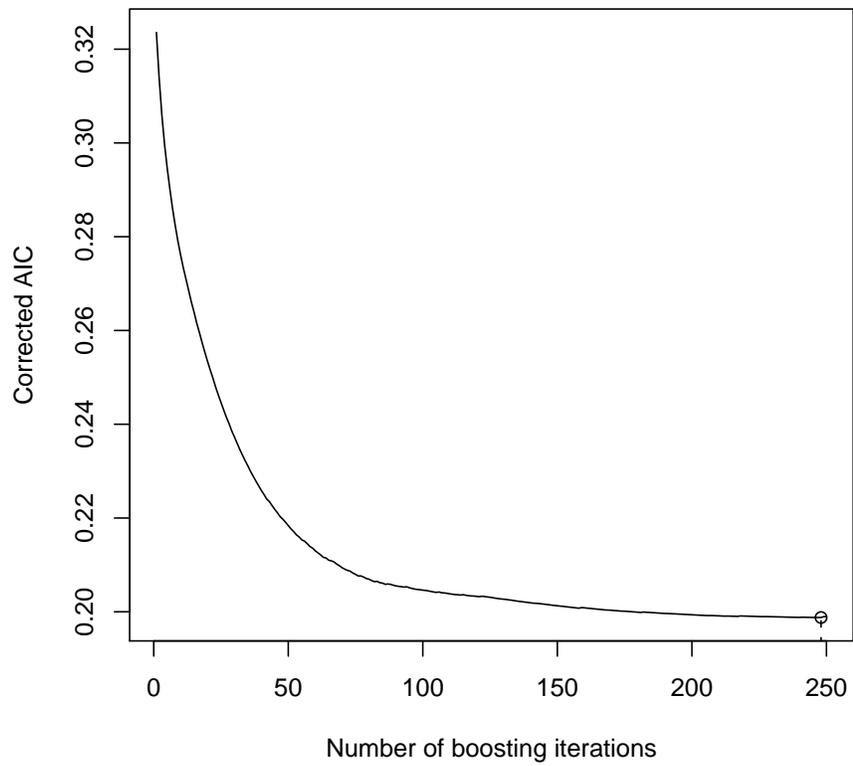


Figure 3: AIC criterion for GBSG2 data.

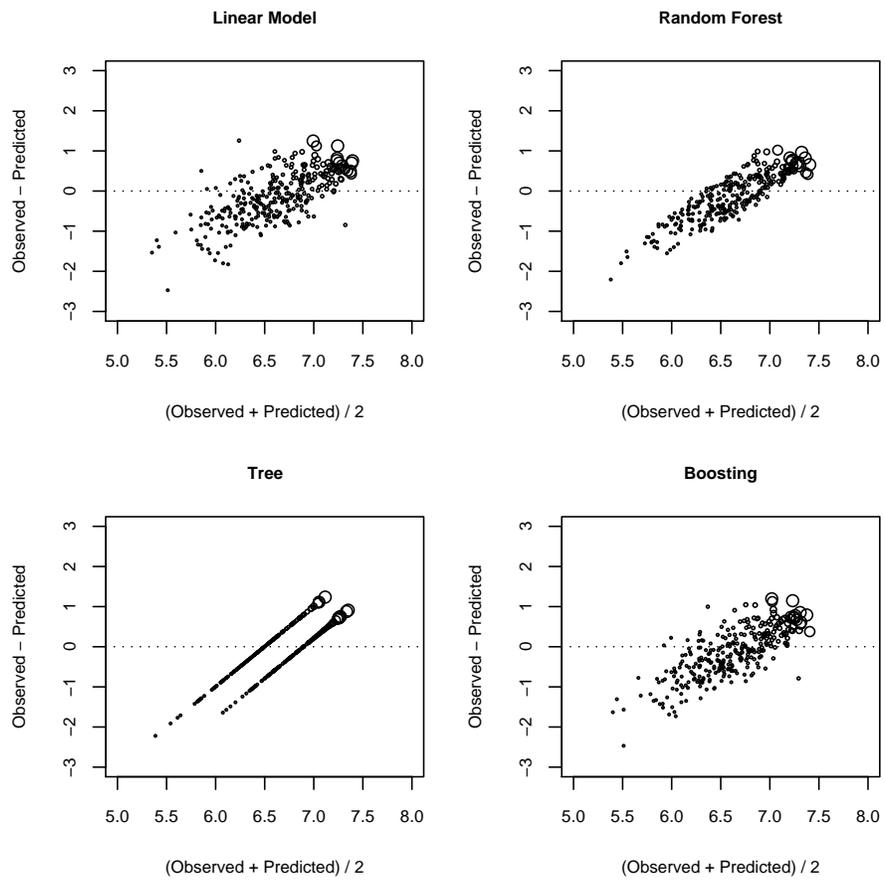


Figure 4: GBSG-2 data: Reproduction of Figure 3.

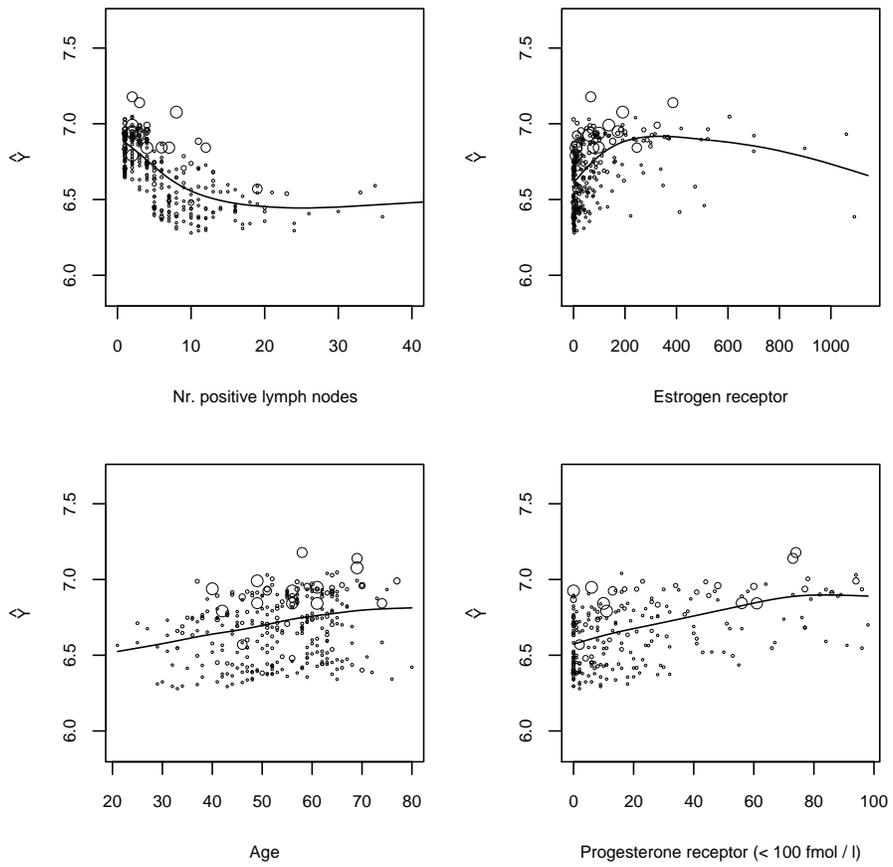


Figure 5: GBSG-2 data: Reproduction of Figure 5.

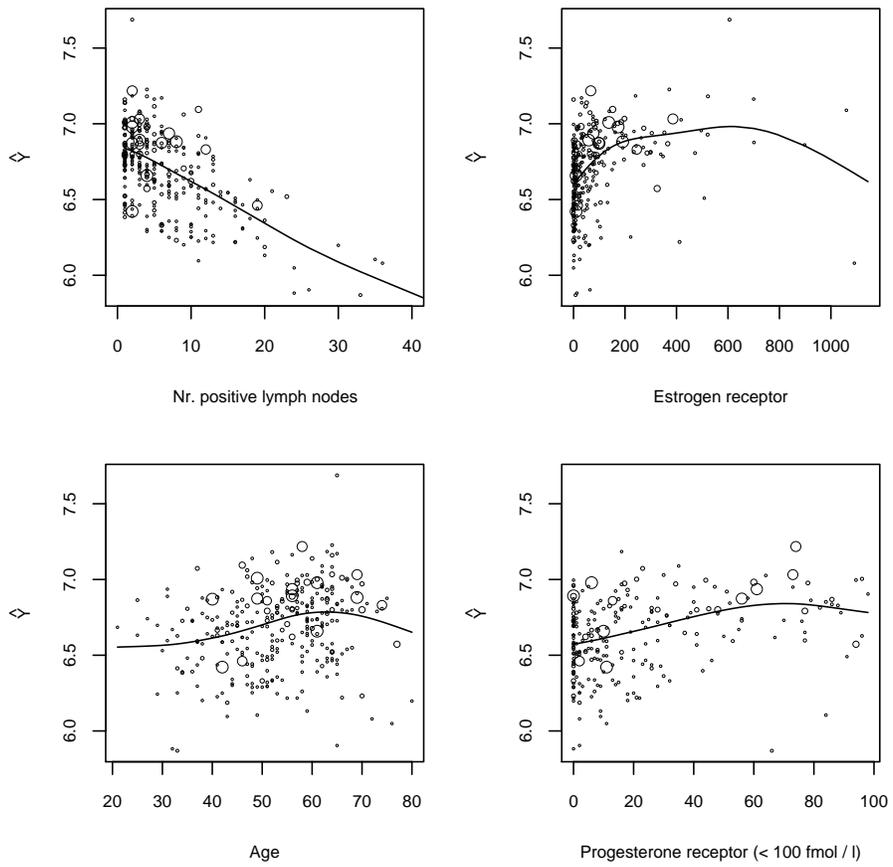


Figure 6: GBSG-2 data: Reproduction of Figure 6.

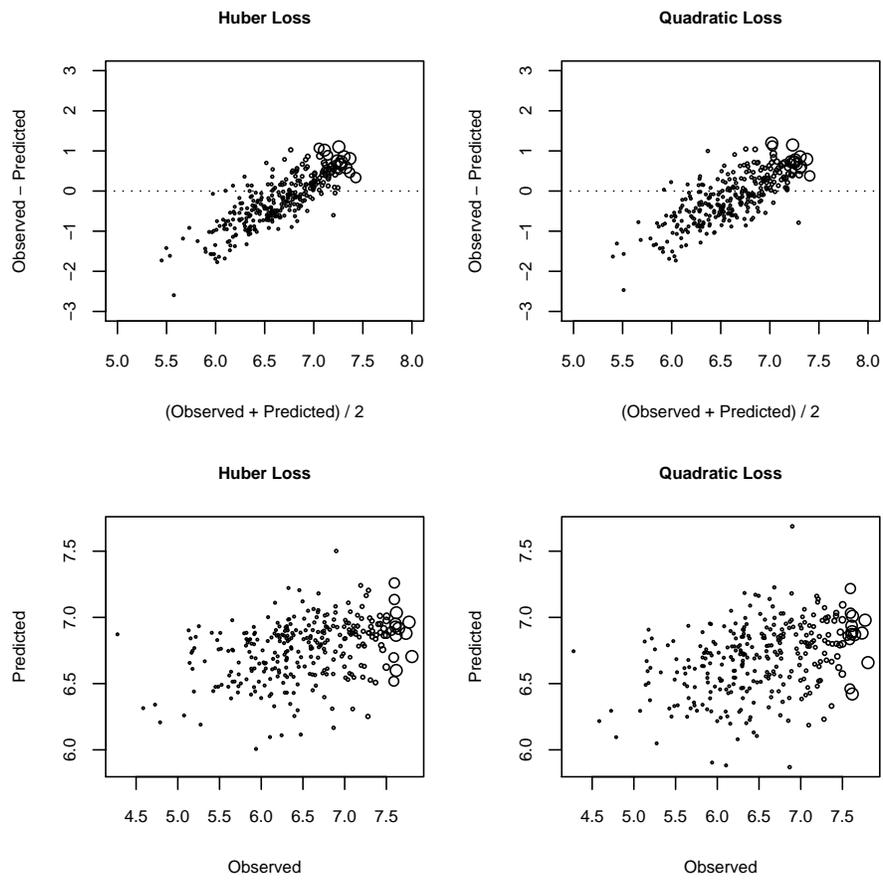


Figure 7: GBSG-2 data: Reproduction of Figure 7.

References

- T. Hothorn, P. Bühlmann, S. Dudoit, A. Molinaro, and M. van der Laan. Survival ensembles. *Biostatistics*, 7:355–373, 2006.