

phia_lca2_jsg2145

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```
# Read public-release datasets
#Biomarker dataset
biomarker <- read_csv(file = "./data/Shims22016adultbio.csv",
                      col_names = TRUE,
                      col_types = NULL,
                      locale = default_locale(),
                      na = c("", "NA"),
                      quoted_na = TRUE,
                      quote = "\"",
                      comment = "#",
                      trim_ws = TRUE,
                      skip = 0,
                      n_max = Inf,
                      progress = show_progress(),
                      skip_empty_rows = TRUE)
```

```
bio_dat <- biomarker %>%
  filter(hiv1statusfinalsurvey == 1) %>%
  select(personid,
         hiv1statusfinalsurvey,
         awareselfreported,
         gender,
         arvstatus,
         artselfreported,
         resultvlc,
         vls,
         tri90,
         tri90aware,
         tri90art,
         tri90vls) %>%
  mutate(resultvlc = recode(resultvlc,
                            "< LLOD" = "1",
                            "< LLOQ: 20" = "20",
                            "< LLOQ: 40" = "40",
                            "< LLOQ: 400" = "400",
                            "< LLOQ: 839" = "839",
                            "> ULOQ 10000000" = "10000000"),
         resultvlc = as.numeric(resultvlc),
         vlunder200 = if_else(resultvlc < 200, 1, 2),
         across(.funs = as_factor))
```

```
str(bio_dat)
```

```
## tibble [3,055 x 13] (S3: tbl_df/tbl/data.frame)
## $ personid      : chr [1:3055] "SW0000000000000502" "SW0000000000000803" "SW0000000000000806" "SW0000000000000807" ...
## $ hiv1statusfinalsurvey: chr [1:3055] "1" "1" "1" "1" ...
## $ awareselfreported  : num [1:3055] 1 2 1 1 1 1 1 1 1 1 ...
## $ gender            : num [1:3055] 2 2 2 2 1 2 2 2 2 1 ...
## $ arvstatus          : num [1:3055] 1 2 2 1 1 1 1 2 1 2 ...
## $ artselfreported    : num [1:3055] 1 99 99 1 1 1 1 2 1 2 ...
## $ resultvlc          : num [1:3055] 20 99137 1468618 1 20 ...
## $ vls                : num [1:3055] 1 2 2 1 1 1 1 2 1 2 ...
## $ tri90              : num [1:3055] 1 1 2 1 1 1 1 1 1 1 ...
## $ tri90aware         : num [1:3055] 1 2 99 1 1 1 1 1 1 1 ...
## $ tri90art           : num [1:3055] 1 3 99 1 1 1 1 2 1 2 ...
## $ tri90vls           : num [1:3055] 1 2 99 1 1 1 1 2 1 2 ...
## $ vlunder200         : num [1:3055] 1 2 2 1 1 1 1 2 1 2 ...
```

```
# aware:      1 - Aware or considered aware because ARVs detectable
#              2 - Unaware and ARVs not detectable, or unaware and ARV testing results missing
#              99 - Missing
#
# art:        1 - ARVs detectable, self-reported on ART, or both ARVs detectable and self-reported on ART
#              2 - Unaware or aware, ARVs not detectable and self-reported not on ART, or aware and self-reported not on ART
#              99 - Missing
# awareselfreported: 1 - Self-report aware of HIV + status
#                   2 - Self-report not aware of HIV + status
#                   99 - missing
# artselfreported:  1 - On ART
#                   2 - Not on ART
#                   99 - Missing
# awareartselfreported: 1 - Self-report as not previously diagnosed
#                       2 - Self-report as previously diagnosed, not on ART
#                       3 - Previously diagnosed, on ART
#                       99 - Missing, including incomplete tri90 information
# arvstatus:      1 - ARV detected
#                 2 - ARV not detected
#                 99 - Missing
# resultvlc:      > ULOQ 10000000 - Upper limit of quantification 10000000
#                 < LLOD - less than lower limit of detection
#                 < LLOQ: 839 - less than lower limit of quantification of 839
#                 < LLOQ: 400 - less than lower limit of quantification of 400
#                 < LLOQ: 40 - less than lower limit of quantification of 40
#                 < LLOQ: 20 - less than lower limit of quantification of 20
# hivselfreport:  1 - Self-reported positive
#                 2 - Self-reported negative
#                 3 - Self-reported never tested or never received test result
#                 99 - Missing
# tri90art        1 - ARVs detectable, self-reported on ART, or both ARVs detectable and self-reported on ART
#                 2 - ARVs not detectable and self-reported not on ART or missing ARV data and self-reported not on ART
#                 3 - Recoded as not on ART (unaware and ARVs not detectable or unaware and ARVs not detectable and self-reported not on ART)
#                 99 - Incomplete Tri90 information
#
```

```

bio_dat_lca = bio_dat %>%
  select(awareselfreported,
         arvstatus,
         artselfreported,
         gender,
         vls,
         tri90,
         tri90aware,
         tri90art,
         tri90vls,
         vlunder200) %>%
  mutate(awareselfreported = recode(awareselfreported, "1" = "1", "2" = "0", "99" = NULL),
         arvstatus = recode(arvstatus, "1" = "1", "2" = "0", "99" = NULL),
         artselfreported = recode(artselfreported, "1" = "1", "2" = "0", "99" = NULL),
         gender = recode(gender, "1" = "1", "2" = "0", "99" = NULL),
         vls = recode(vls, "1" = "1", "2" = "0", "99" = NULL),
         tri90 = recode(tri90, "1" = "1", "2" = "0", "99" = NULL),
         tri90aware = recode(tri90aware, "1" = "1", "2" = "0", "99" = NULL),
         tri90art = recode(tri90art, "1" = "1", "2" = "0", "99" = NULL),
         tri90vls = recode(tri90vls, "1" = "1", "2" = "0", "99" = NULL),
         vlunder200 = recode(vlunder200, "1" = "1", "2" = "0", "99" = NULL))

```

```

bio_dat_lca_full = bio_dat %>%
  select(awareselfreported,
         arvstatus,
         gender,
         vlunder200) %>%
  mutate(across(everything(), ~recode(., "1" = "1", "2" = "0", "99" = NULL)))

```

```

bio_dat1 = bio_dat %>%
  mutate(awareselfreported = recode(awareselfreported, "1" = "1", "2" = "0", "99" = NULL))

lca_prepper = function(.) {
  mutate(across(.funs = recode(., "1" = "1", "2" = "0", "99" = NULL))
}

```

“Full Model”

```

# full model
bio_lca_full = bio_dat_lca %>%
  select(awareselfreported, arvstatus, vlunder200)

set.seed(22)
bio_lca = randomLCA::randomLCA(bio_lca_full, calcSE = TRUE)
summary(bio_lca)

```

```

##   Classes      AIC      BIC      AIC3    logLik penlogLik
##           2 6777.542 6819.714 6784.542 -3381.771 -3381.818

```

```
## Class probabilities
## Class 1 Class 2
## 0.2347 0.7653
## Outcome probabilities
##      awareselfreported arvstatus vlunder200
## Class 1      0.5015      0.0251      0.0464
## Class 2      0.9773      0.9872      0.9118
```

```
probs = outcomeProbs(bio_lca)
postClassProbs(bio_lca)
```

	awareselfreported	arvstatus	vlunder200	Freq	Class 1	Class 2
## 1	0	0	0	329	0.9998197716	0.0001802284
## 2	0	0	1	16	0.9630959474	0.0369040526
## 3	0	1	0	13	0.6488982251	0.3511017749
## 4	0	1	1	46	0.0086194775	0.9913805225
## 5	0	NA	0	3	0.9864386751	0.0135613249
## 6	0	NA	1	3	0.2549479739	0.7450520261
## 7	1	0	0	332	0.9923495506	0.0076504494
## 8	1	0	1	42	0.3789605022	0.6210394978
## 9	1	1	0	204	0.0414237541	0.9585762459
## 10	1	1	1	2015	0.0002032504	0.9997967496
## 11	1	NA	0	7	0.6297360897	0.3702639103
## 12	1	NA	1	43	0.0079374908	0.9920625092
## 13	NA	0	0	1	0.9960598053	0.0039401947
## 14	NA	1	0	1	0.0776775598	0.9223224402

```
BIC(bio_lca)
```

```
## [1] 6819.714
```

```
diseased <- ifelse(probs[[1]]$Outcome[1] < probs[[2]]$Outcome[1], 2, 1)
notdiseased <- 3 - diseased
sens <- apply(probs[[diseased]], 1, function(x)

  sprintf("%3.2f (%3.2f, %3.2f)",
    x[1],
    x[2],
    x[3])
)
spec <- apply(probs[[notdiseased]], 1, function(x)
  sprintf("%3.2f (%3.2f, %3.2f)",
    1 - x[1],
    1 - x[3],
    1 - x[2])
)

stable <- data.frame(sens, spec)
names(stable) <- c("Sensitivity", "Specificity")
print(stable, row.names = TRUE)
```

```

stable = function(probs = probs){

  diseased <- ifelse(probs[[1]]$Outcome[1] < probs[[2]]$Outcome[1], 2, 1)

  notdiseased <- 3 - diseased

  sens <- apply(probs[[diseased]], 1, function(x)
    sprintf("%3.2f (%3.2f, %3.2f)",
            x[1],
            x[2],
            x[3])
  )

  spec <- apply(probs[[notdiseased]], 1, function(x)
    sprintf("%3.2f (%3.2f, %3.2f)",
            1 - x[1],
            1 - x[3],
            1 - x[2])
  )

  stable <- data.frame(sens, spec)

  names(stable) <- c("Sensitivity", "Specificity")

  return(print(stable, row.names = TRUE))
}

# stable(probs2)

```

“Fuller model”

```

# full model
set.seed(22)
bio_lca_fuller = randomLCA::randomLCA(bio_dat_lca_full, random = FALSE, calcSE = TRUE)

probs_full = outcomeProbs(bio_lca_fuller)

stable(probs_full)

```

	Sensitivity	Specificity
## awareselfreported	0.98 (0.97, 0.98)	0.50 (0.46, 0.54)
## arvstatus	0.99 (0.98, 0.99)	0.97 (0.94, 0.99)
## gender	0.31 (0.29, 0.33)	0.62 (0.58, 0.66)
## vlunder200	0.91 (0.90, 0.92)	0.96 (0.93, 0.97)

“Simple Model”

A model of sensitivity and specificity for just one variable should be compared to a true (known) outcome. As such, a contingency table should be sufficient to calculate the sensitivity and specificity of each variable.

awareselfreported

```
xtabs(~ awareselfreported + tri90aware, data = bio_dat_lca)
```

```
##           tri90aware
## awareselfreported  0    1
##                0 346   59
##                1    0 2591
```

Assuming the final classifications are true for tri90aware, there are 59 false negatives and no false positives. This corresponds to a sensitivity of $2591/2650$, or 0.978, and a specificity of $346/346$, or 1.00. The prevalence (of awareness of HIV status) is therefore 0.885. The PPV can be calculated as $2591/2591$, or 1.00, and the NPV is $346/405 = 0.854$.

arvstatus

```
xtabs(~ arvstatus + tri90aware, data = bio_dat_lca)
```

```
##           tri90aware
## arvstatus  0    1
##          0 345 369
##          1    0 2279
```

Assuming the final classifications are true for tri90aware, there are 369 false negatives and no false positives. This corresponds to a sensitivity of $2279/2648$, or 0.861, and a specificity of $345/345$, or 1.00. The prevalence (of awareness of HIV status) is therefore 0.885. The PPV can be calculated as $2279/2279$, or 1.00, and the NPV is $345/714 = 0.483$.

vlunder200

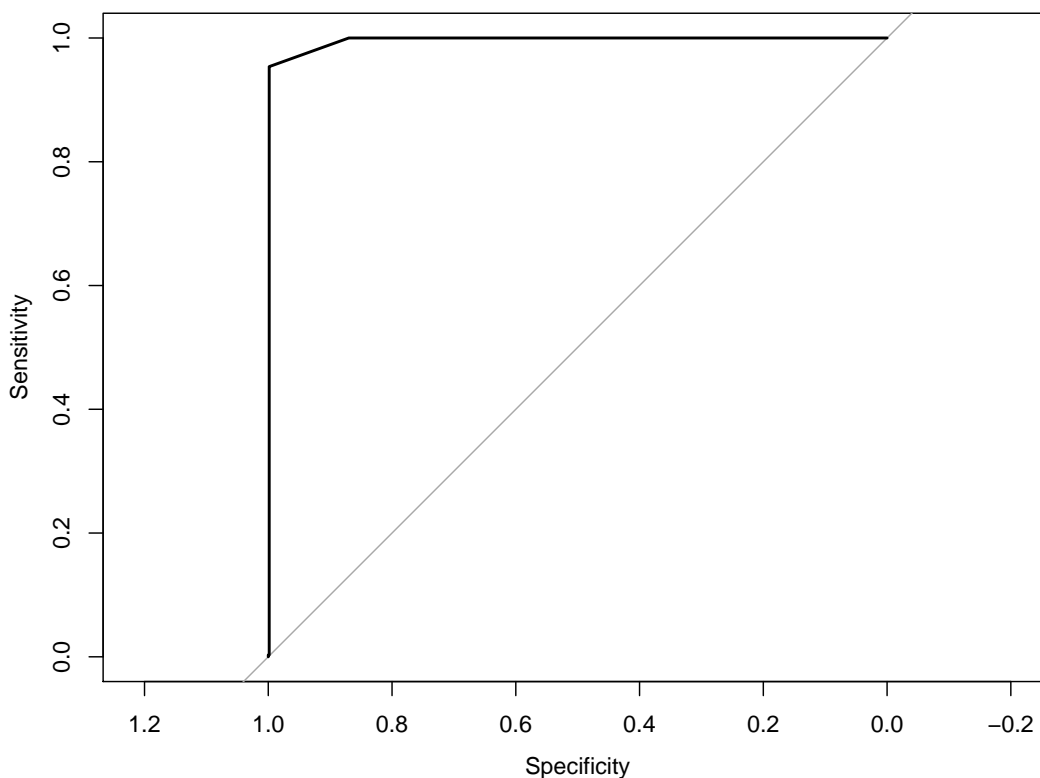
```
xtabs(~ vlunder200 + tri90aware, data = bio_dat_lca)
```

```
##           tri90aware
## vlunder200  0    1
##          0 330 546
##          1   16 2105
```

Assuming the final classifications are true for tri90aware, there are 546 false negatives and 16 false positives. This corresponds to a sensitivity of $2105/2651$, or 1, and a specificity of $330/346$, or 0.954. The prevalence (of awareness of HIV status) is therefore 0.885. The PPV can be calculated as $2105/2121$, or 0.992, and the NPV is $330/876 = 0.377$.

ROC

```
roc(bio_dat$tri90aware, c(bio_dat$awareselfreported + bio_dat$arvstatus + bio_dat$vlunder200), data = b
plot())
```



```
test1 = roc(tri90aware ~ awareselfreported, bio_dat)
test2 = roc(tri90aware ~ arvstatus, bio_dat)
roc.test(test1, test2)
```

```
##
## DeLong's test for two correlated ROC curves
##
## data: test1 and test2
## Z = 15.526, p-value < 2.2e-16
## alternative hypothesis: true difference in AUC is not equal to 0
## sample estimates:
## AUC of roc1 AUC of roc2
## 0.9884949 0.9294748
```

```
roc.test(bio_dat$tri90aware, bio_dat$awareselfreported, bio_dat$arvstatus)
```

```
##
## DeLong's test for two correlated ROC curves
##
## data: bio_dat$awareselfreported and bio_dat$arvstatus by bio_dat$tri90aware (1, 2)
## Z = 15.526, p-value < 2.2e-16
```

```
## alternative hypothesis: true difference in AUC is not equal to 0
## sample estimates:
## AUC of roc1 AUC of roc2
## 0.9884949 0.9294748
```