Hypothesis Testing

Slides

Chapter slides here. (To convert html to pdf, press $E \to Print \to Destination: Save to pdf)$

R code

```
# This code generates the numerical results in chapter 2
# load the survival package
library(survival)
# install and load the WR package for
# 1. dataset hfaction_cpx9;
# 2. function WRrec() for win ratio test (of recurrent events and death)
# 3. functions base() and WRSS() for sample size calculation
# install.packages("WR")
library(WR)
library(tidyverse) # for data wrangling (dplyr, ggplot2, etc.)
##### Read in HF-ACTION DATA #######
# same as rmt::hfaction used in chap 1
# (except for status coding)
data(hfaction_cpx9)
hfaction <- hfaction_cpx9
head(hfaction)
#> Shows the first few rows of hfaction_cpx9 dataset
# count unique patients in each arm
hfaction |>
group_by(trt_ab) |>
```

```
distinct(patid) |>
  count(trt_ab)
#> This gives the number of unique patients (patid) by treatment arm (trt_ab)
#### demo ##########
# WRrec() fits the recurrent event plus death model (Win Ratio approach)
obj <- WRrec(
  ID = hfaction$patid,
 time = hfaction$time,
  status = hfaction$status,
 trt = hfaction$trt_ab,
 strata = hfaction$age60,
 naive = TRUE
# summary results
#> Displays the main results, including win ratio estimates for each method.
# LWR
beta <- obj$log.WR # log-win ratio for LWR
                # standard error for log-win ratio (LWR)
se <- obj$se
# test
pval <- 2 * (1 - pnorm(abs(beta / se)))</pre>
#> Two-sided p-value for LWR
# NWR.
beta.naive <- obj$log.WR.naive # log-win ratio for naive WR (NWR)
                               # its standard error
se.naive <- obj$se.naive</pre>
# test
pval.naive <- 2 * (1 - pnorm(abs(beta.naive / se.naive)))</pre>
pval.naive
#> Two-sided p-value for NWR
beta.FI <- obj$log.WR.FI # log-win ratio for Fisher's information-based WR (FWR)
se.FI <- obj$se.FI
                         # standard error for log(FWR)
pval.FI <- 2 * (1 - pnorm(abs(beta.FI / se.FI)))</pre>
pval.FI
#> Two-sided p-value for FWR
```

```
####################################
# Win ratio analyses: tabulate #
####################################
data <- hfaction
### create a dataset with only the first hospitalization -> data.H1
# hospitalization data
tmpH <- data[data$status == 2, ]</pre>
# get the first record of each id
o <- order(tmpH$patid, tmpH$time)</pre>
tmpH <- tmpH[o, ]</pre>
tmpFH <- tmpH[!duplicated(tmpH$patid), ]</pre>
# combine it with mortality data
data.H1 <- rbind(tmpFH, data[data$status != 2, ])</pre>
o <- order(data.H1$patid, data.H1$time)
data.H1 <- data.H1[o, ]</pre>
# Function to create a summary table for
# PWR, NWR, FWR, and LWR with their 95% CI and p-values
# ind: index (logical) for rows in the main 'data'
# ind1: index (logical) for rows in 'data.H1'
       number of decimals for rounding in the output
gwr.fun = function(ind, ind1, r = 2) {
  # fit NWR, FWR, and LWR to original data (multiple events)
  obj <- WRrec(
   ID = data$patid[ind],
   time = data$time[ind],
   status = data$status[ind],
   trt = data$trt_ab[ind],
   strata = data$age60[ind],
   naive = TRUE
  # fit sWR (PWR) to dataset with first hospitalization only
  # This typically addresses "semi-competing" event structure
  obj1 <- WRrec(
    ID = data.H1$patid[ind1],
   time = data.H1$time[ind1],
    status = data.H1$status[ind1],
```

```
trt = data.H1$trt_ab[ind1],
 strata = data.H1$age60[ind1],
 naive = FALSE
# critical value for a 95% confidence interval
za \leftarrow qnorm(0.975)
## LWR results
beta <- obj$log.WR
se <- obj$se
theta <- obj$theta # proportion of pairwise comparisons that are wins/losses
# Format: percentage of wins, percentage of losses,
         win ratio & 95% CI, p-value
r4 <- c(
 paste0(round(100 * theta[1], 1), "%"), # Win
 paste0(round(100 * theta[2], 1), "%"), # Loss
 paste0(
   round(exp(beta), r), " (",
   round(exp(beta - za * se), r), ", ",
   round(exp(beta + za * se), r), ")"
 ),
 round(1 - pchisq((beta / se)^2, 1), 3)
## PWR results
beta1 <- obj1$log.WR
se1 <- obj1$se
theta1 <- obj1$theta
r1 <- c(
 paste0(round(100 * theta1[1], 1), "%"),
 paste0(round(100 * theta1[2], 1), "%"),
 paste0(
   round(exp(beta1), r), " (",
   round(exp(beta1 - za * se1), r), ", ",
   round(exp(beta1 + za * se1), r), ")"
 ),
 round(1 - pchisq((beta1 / se1)^2, 1), 3)
)
```

```
## NWR results
 beta.naive <- obj$log.WR.naive</pre>
  se.naive <- obj$se.naive
 theta.naive <- obj$theta.naive</pre>
 r2 <- c(
   paste0(round(100 * theta.naive[1], 1), "%"),
   paste0(round(100 * theta.naive[2], 1), "%"),
   paste0(
     round(exp(beta.naive), r), " (",
     round(exp(beta.naive - za * se.naive), r), ", ",
     round(exp(beta.naive + za * se.naive), r), ")"
   ),
   round(1 - pchisq((beta.naive / se.naive)^2, 1), 3)
  ## FWR results
 beta.FI <- obj$log.WR.FI
 se.FI <- obj$se.FI
 theta.FI <- obj$theta.FI
 r3 <- c(
   paste0(round(100 * theta.FI[1], 1), "%"),
   paste0(round(100 * theta.FI[2], 1), "%"),
   paste0(
     round(exp(beta.FI), r), " (",
     round(exp(beta.FI - za * se.FI), r), ", ",
     round(exp(beta.FI + za * se.FI), r), ")"
   ),
   round(1 - pchisq((beta.FI / se.FI)^2, 1), 3)
 )
 # Combine rows into a single table
 result <- rbind(r1, r2, r3, r4)
 rownames(result) <- c("PWR", "NWR", "FWR", "LWR")</pre>
 return(result)
# Create table
## Age <= 60 years
ind <- (data\$age60 == 0)
```

```
ind1 \leftarrow (data.H1\$age60 == 0)
result.lt60 <- gwr.fun(ind, ind1, r = 2)
## Age > 60 years
ind \leftarrow (data\$age60 == 1)
ind1 <- (data.H1$age60 == 1)
result.ge60 <- gwr.fun(ind, ind1, r = 2)
## overall
ind <- rep(TRUE, nrow(data))</pre>
ind1 <- rep(TRUE, nrow(data.H1))</pre>
result.all <- gwr.fun(ind, ind1, r = 2)
# combine results
results <- rbind(result.lt60, result.ge60, result.all)</pre>
colnames(results) <- c("Win", "Loss", "Win ratio (95% CI)", "p-value")
noquote(results)
#> Final table of all 4 measures (PWR, NWR, FWR, LWR) across strata.
Sample size calculation
# get training arm data
pilot <- hfaction |>
 filter(trt_ab == 1)
# number of subjects
pilot |>
 distinct(patid) |>
  count()
#> This indicates how many unique subjects were in the training arm
############ estimate parameters ############
# Get the variables from pilot dataset
# to estimate baseline parameters
# lambda_D, lambda_H, kappa
outcome_base <- gumbel.est(pilot$patid, pilot$time / 12, pilot$status)
lambda_D <- outcome_base$lambda_D</pre>
lambda_H <- outcome_base$lambda_H</pre>
kappa <- outcome_base$kappa
```

```
lambda_D
lambda_H
kappa
#> Baseline hazards for death/hospitalization and the gumbel 'kappa' parameter
## Kendall's rank correlation
1 - 1/kappa
#> [1] 0.360812
#> This measures correlation between timing of repeated events
### demo ################
# set design parameters
tau_b <- 3  # time from baseline to start of follow-up for base() computation
           # total follow-up (in years)
lambda_L < -0.01 # Additional parameter used in the base() function
# use base() function to compute zeta2 and delta
## may take up to 30s
bparam <- base(lambda_D, lambda_H, kappa, tau_b, tau, lambda_L)
#> bparam includes the baseline rates and distribution shape
#> used for sample size calculations
# compute sample size under HRs 0.8 and 0.9
# for death and nonfatal event, respectively
obj <- WRSS(
 xi = log(c(0.9, 0.8)),
 bparam = bparam,
  q = 0.5,
 alpha = 0.05,
 power = 0.8
obj$n
#> The required sample size for the given HRs at 80% power
## effect size specification
thetaD <- seq(0.6, 0.95, by = 0.05) # hazard ratio for death
thetaH <- seq(0.6, 0.95, by = 0.05) # hazard ratio for hospitalization
## create a matrix "SS08" for sample size powered at 80%
## under each combination of thetaD and thetaH
mD <- length(thetaD)</pre>
mH <- length(thetaH)
```

```
SSO8 <- matrix(NA, mD, mH)
rownames(SS08) <- thetaD
colnames(SS08) <- thetaH
## fill in the computed sample size values
for (i in 1:mD) {
  for (j in 1:mH) {
    ## sample size under hazard ratios thetaD[i] for death
    ## and thetaH[j] for hospitalization
    SS08[i, j] <- WRSS(
      xi = log(c(thetaD[i], thetaH[j])),
      bparam = bparam,
      q = 0.5,
      alpha = 0.05,
      power = 0.8
    )$n
  }
}
## print the calculated sample sizes
print(SS08)
#> Shows how sample size changes under different hazard ratios for death/hosp
## repeating the same calculation for power = 90%
SS09 <- matrix(NA, mD, mH)
rownames(SS09) <- thetaD
colnames(SSO9) < -thetaH # As in original code; sets the colnames to the sequence of thetaH
## fill in the computed sample size values
for (i in 1:mD) {
  for (j in 1:mH) {
    ## sample size under hazard ratios thetaD[i] for death
    ## and thetaH[j] for hospitalization
    SS09[i, j] <- WRSS(
      xi = log(c(thetaD[i], thetaH[j])),
      bparam = bparam,
      q = 0.5,
      alpha = 0.05,
      power = 0.9
    )$n
  }
## print the calculated sample sizes
```

```
print(SS09)
#> Sample sizes under 90% power requirements
oldpar <- par(mfrow = par("mfrow"))</pre>
par(mfrow = c(1, 2))
persp(
  thetaD, thetaH, SS08 / 1000,
  theta = 50, phi = 15, expand = 0.8, col = "gray",
  ltheta = 180, lphi = 180, shade = 0.75,
  ticktype = "detailed",
  xlab = "\n HR on Death", ylab = "\n HR on Hospitalization",
  zlab = paste0("\n Sample Size (10e3)"),
  main = "Power = 80%",
  zlim = c(0, 26)
)
#> 3D perspective plot of sample size (in thousands) for power=80%
#> over various hazard ratios for death/hosp
persp(
  thetaD, thetaH, SS09 / 1000,
  theta = 50, phi = 15, expand = 0.8, col = "gray",
  ltheta = 180, lphi = 180, shade = 0.75,
  ticktype = "detailed",
  xlab = "\nHR on Death", ylab = "\nHR on Hospitalization",
  zlab = paste0("\n Sample Size (10e3)"),
  main = "Power = 90%",
  zlim = c(0, 26)
#> Similar 3D perspective for power=90%
```

Win Ratio: Definitions and Properties

The win ratio (WR) is a pairwise, nonparametric method for comparing composite outcomes in a prioritized manner. Each subject in the treatment arm is compared to each subject in the control arm over a shared follow-up window. A win is declared if the treated subject experiences a better outcome—typically defined as longer survival or, if tied on survival, fewer or later nonfatal events.