

# Misclassification Activity

## JSM 2024 Short Course

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## Recall this example

```
data.tbl <- matrix(c(45,94,257,945),  
  dimnames = list(c("CHD+", "CHD-"),c("Resin+", "Resin-")),  
  nrow = 2, byrow = TRUE)
```

```
data.tbl
```

##	Resin+	Resin-
## CHD+	45	94
## CHD-	257	945

# Naive analysis presuming correct exposure classification

Inference for exposure-disease odds-ratio

```
logOR.hat <- sum(c(1,-1,-1,1)*log(as.vector(data.tbl)))
```

```
logOR.SE <- sqrt(sum(1/as.vector(data.tbl)))
```

```
exp(logOR.hat + c(0, -1.96, 1.96)*logOR.SE)
```

```
## [1] 1.76 1.20 2.58
```

## Assuming nondifferential exposure misclassification with 90% sensitivity and 80% specificity

Again, recall from slides:

```
require(episensr)

ft <- misclassification(data.tbl,
  type="exposure", bias_parms=c(0.9, 0.9, 0.8, 0.8))

# point and interval estimation of OR
ft$adj.measures[2,]
```

```
##          2.5% 97.5%
## 10.67    1.64 69.55
```

## Activity A

Check you can reproduce one of the **differential** classification adjustments given in the slides (i.e., one of the off-diagonal table entries on slide 151).

For instance, try presuming 90% specificity for all subjects, but sensitivity of 90% for controls, compared to 80% for cases.

Might help:

```
help(misclassification)
```

## Activity B: Uncertainty about misclassification rates

Say the investigator is confident that the misclassification is nondifferential.

Has 85% sensitivity and 85% specificity as “best guesses.”

But thinks either guess could be off by as much as five percentage points.

Can you look at

```
help(probsens)
```

and then provide an appropriate analysis?

HINT: First example in the help gives a template.

HINT: For simplicity, maybe “triangular” or “uniform” instead of “trapezoidal”

## Activity C - Role of data

We have useful heuristics in statistics, such as the primal role of  $\sqrt{n}$ .

If I want interval estimates *twice* as narrow, I likely need about *four times* as much data.

Repeat Activity B, but with four times as much data. (Simplest to just keep cell *proportions* fixed in the 2 by 2 data table).

Reflect on what you find.

## Activity D - Bayesian approach

Factor the joint dist. of  $(X, X^*, Y)$  in terms of  $(Y)$ ,  $(X|Y)$ ,  $(X^*|X, Y)$ .

Leave  $(Y)$  as unmodeled [since  $Y$  observed, and given case-control design, parameter of most interest is determined by  $X|Y$ ].

Parameterize as  $Pr(X = 1|Y = y) = r_y$ , and

$$Pr(X^* = X|X = x, Y = y) = \begin{cases} Sp & \text{if } x = 0, \\ Sn & \text{if } x = 1. \end{cases}$$

Parameter of most interest:  $\psi = \log OR(X, Y) = \text{logit}(r_1) - \text{logit}(r_0)$ .

Priors  $r_j \sim \text{Unif}(0, 1)$ ,  $Sn \sim \text{beta}(a_{sn}, b_{sn})$ ,  $Sp \sim \text{beta}(a_{sp}, b_{sp})$ .



## Activity D, continued

A bit too much overhead with trying to get JAGS/rJAGS going in a matter of minutes (unless you have experience. . .)

Here (meaning sitting in the .Rmd file generating these slides) is a bespoke R function (called **bespoke()**) to do MCMC for this model/prior only:

## bespoke()

```
bespoke <- function(n.0, n.1, mstr.0, mstr.1,
                   a.sn, b.sn, a.sp, b.sp,
                   N.REP=50000, N.BURN=100) {

  ### INPUTS
  ### n.j is size of control (j=0) and case (j=1) samples
  ### mstr.j is number (out of n.j) of apparently exposed
  ### a.sn, b.sn, a.sp, b.sp are hyperparameters

  ### LATENTS
  ### m.j is number (out of n.j) actually exposed
  ### t.j is number (out of mstr.j) actually exposed
  ###      amongst the apparents

  ### OUTPUT output will be matrix, MC sample from posterior
  ans <- matrix(NA, N.REP, 8)
  colnames(ans) <- c("r0", "r1", "sn", "sp", "m0", "m1", "t0", "t1")
}
```

## Example

Say I am pretty sure that the exposure classification is very good (though probably not perfect). I encode this with priors  $S_n \sim \text{Beta}(140, 10)$ ,  $S_p \sim \text{Beta}(140, 10)$ .

*Sidenote:* As a thought experiment, this would formally be the evidence had we done an external validation of 150 truly unexposed and 150 truly exposed individuals, and found 10 misclassifications in each group.

```
mc.opt <- bespoke(n.0=257+945, n.1=45+94,  
                  mstr.0=257, mstr.1=45,  
                  a.sn=140, b.sn=10, a.sp=140, b.sp=10)
```

## Example, continued

```
### focus on target parameter
```

```
mc.trg <- logit(mc.opt[, "r1"]) - logit(mc.opt[, "r0"])  
summary(mc.trg)
```

```
##      Min. 1st Qu.  Median    Mean 3rd Qu.    Max.  
## -0.513   0.570   0.736   0.739   0.905   2.150
```

```
### estimate OR
```

```
exp(mean(mc.trg))
```

```
## [1] 2.09
```

```
### corresponding 95% credible interval
```

```
exp(quantile(mc.trg, c(0.025, 0.975)))
```

```
## 2.5% 97.5%
```

```
## 1.28 3.46
```

## Activity D - you try

Can you carry out a Bayesian analysis with *about* the same sort of uncertainty about misclassification parameters as in Activity B.

## Activity D - further thinking/doing points

Depending on your background and interests, you could:

- Look at the code for **bespoke()** to confirm how the MCMC algorithm (in this case the Gibbs sampler) bounces back and forth between sampling complete data given parameters, and sampling parameters given complete data.
- Take a closer look at the Monte Carlo output to confirm that you do not get the luxury of *iid* draws from the posterior distribution, but rather have to live with serially autocorrelated draws.
- Take a closer look at the Monte Carlo output to consider what the posterior distribution of  $S_n$  and  $S_p$  looks like, compared to the prior.