Case Study: MAP Prior with Binary Endpoint

Hongtao Zhang (github/squallteo)

Historical Data

The historical data used in this case study are from four studies of rheumatoid arthritis (RA) in patients who were treated with methotrexate (MTX). The primary binary outcome is the rate of patients satisfying American College of Rheumatology 50 criteria (ACR50) at week 12.

```
library(tidyverse)
library(knitr)
library(kableExtra)
histdt <- tibble(study = 1:4, r = c(33, 98, 3, 36), n = c(221, 651, 20, 214)) %>%
    mutate(ACR50 = round(r/n,3))
histdt %>% kbl(caption="Historical Data") %>%
    kable_classic(full_width = F, html_font = "Cambria", latex_options = "HOLD_position")
```

Table 1: Historical Data

study	r	n	ACR50
1	33	221	0.149
2	98	651	0.151
3	3	20	0.150
4	36	214	0.168

The ACR50 rates in the four historical trials are fairly consistent. The random effect meta-analysis yields a point estimate of 0.154 and an SE of 0.0108. The predictive CI for current control is (0.11, 0.21).

```
with(histdt, meta::metaprop(event = r, n = n, method = "Inverse", prediction = T))
```

```
proportion
                           95%-CI %W(fixed) %W(random)
## 1
         0.1493 [0.1051; 0.2033]
                                       19.5
                                                   19.5
## 2
         0.1505 [0.1239; 0.1804]
                                       57.9
                                                   57.9
## 3
         0.1500 [0.0321; 0.3789]
                                        1.8
                                                    1.8
         0.1682 [0.1207; 0.2252]
                                       20.8
                                                   20.8
##
## Number of studies combined: k = 4
##
##
                                               95%-CI
                         proportion
                             0.1538 [0.1337; 0.1763]
## Fixed effect model
## Random effects model
                             0.1538 [0.1337; 0.1763]
## Prediction interval
                                    [0.1127; 0.2065]
## Quantifying heterogeneity:
  tau^2 = 0 [0.0000; 0.0150]; tau = 0 [0.0000; 0.1225]
   I^2 = 0.0\% [0.0\%; 84.7\%]; H = 1.00 [1.00; 2.56]
##
```

```
## Test of heterogeneity:
## Q d.f. p-value
## 0.43   3  0.9337
##
## Details on meta-analytical method:
## - Inverse variance method
## - DerSimonian-Laird estimator for tau^2
## - Jackson method for confidence interval of tau^2 and tau
## - Logit transformation
## - Clopper-Pearson confidence interval for individual studies
```

Borrowing with Conjugate Beta Prior

We could conduct fixed borrowing via the Beta-Binomial conjugate model. Recall that the point estimate and SE of historical data are 0.154 and 0.0108, respectively. The Beta(a, b) prior for ACR50 rate can be derived to have the same mean and SE, that is, to "match the moments". Specifically, we solve a, b from the equation system

$$\frac{a}{a+b} = 0.154$$
 and $\frac{ab}{(a+b)(a+b+1)^2} = 0.0108^2$.

The solution is Beta(170, 935). Once Y_C ACR50 responses are observed among n_C subjects in the current trial, the posterior distribution of ACR50 rate, which is also a Beta distribution due to conjugacy, is $Beta(170 + Y_C, 935 + n_C - Y_C)$. Further estimations and inferences are drawn based on this known posterior Beta distribution.

MAP Prior

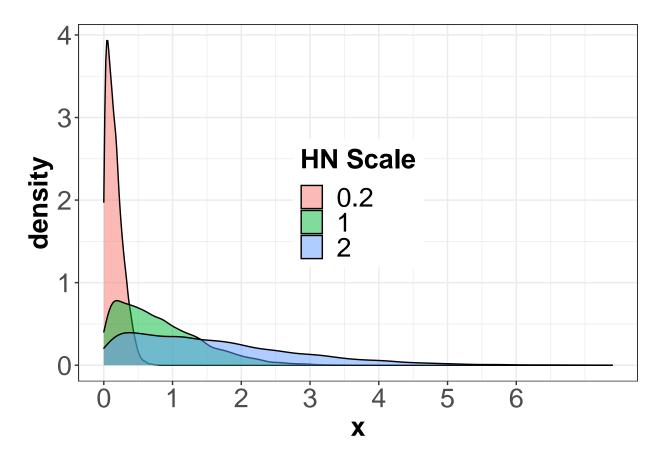
The original MAP prior assumes a common normal distribution for log-odds $\theta_i = \log \frac{p_i}{1-p_i}$ where subscript $i = 1, \dots, 4$ indices the historical trials:

$$\theta_1, \cdots, \theta_4 \sim N(\mu_C, \sigma_C^2),$$

where the hyper-priors are

$$\mu_C \sim N(0, 2^2)$$
 and $\sigma_C \sim HN(s)$.

A standard deviation of 2 for μ_C is fairly large on the logit scale so that the hyper-prior is vague. The half-normal prior regulates the extent of borrowing and thus the scale s should be chosen carefully. One way is to examine the densities corresponding to various scales (0.2, 1, 2 in the plot). We will use HN(s=1) from hereafter.



Below is the command to derive the original MAP prior.

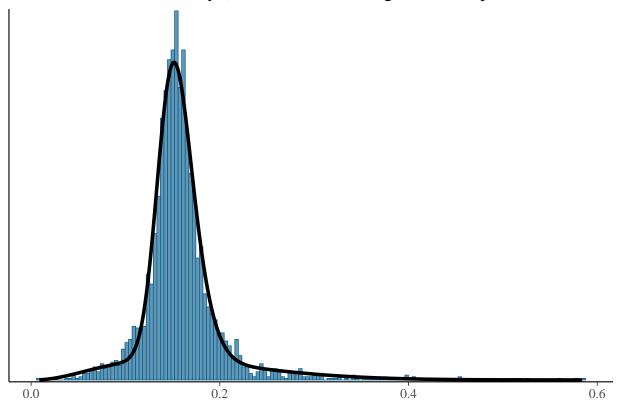
The distribution of the MAP prior is unknown. Approximation is needed to convert it to a "workable" form. With binomial data, the MAP prior is approximated by a weighted mixture of Beta distributions. Three Beta components are used and the approximation is satisfactory as shown both numerically and visually.

```
map_c_hat = mixfit(map_c_mcmc, Nc=3)
print(map_c_hat)
```

```
## EM for Beta Mixture Model
## Log-Likelihood = 5969.913
##
## Univariate beta mixture
## Mixture Components:
## comp1 comp2 comp3
## w 0.3893370 0.3880054 0.2226576
## a 46.5702699 72.0148367 3.5054258
## b 243.4141931 408.0689941 16.2800457
```

```
summary(map_c_mcmc)$theta.pred
                                     sd
                                               2.5%
                                                          50%
                                                                   97.5%
                        mean
## theta_resp_pred 0.1599893 0.04597546 0.08825363 0.1544131 0.2759806
summary(map_c_hat)
##
                               2.5%
                                          50.0%
                                                     97.5%
         mean
                      sd
## 0.16017713 0.04415167 0.08156843 0.15476832 0.28404156
plot(map_c_hat)$mix
```

Parametric Mixture Density (black line) and Histogram of Sample



Robust MAP Prior

The vague prior f_V to construct rMAP prior is Beta(1,1) and we form a 50-50 mixture of original MAP prior and the vague prior, that is,

$$f_{rMAP} = 0.5 * \hat{f}_{MAP} + 0.5 * f_V.$$

```
#RBesT package automatically bumps up n by 1
#see the help document of robustify command for more details
rmap_c <- robustify(map_c_hat, weight=0.5, mean=0.5, n=1)
print(rmap_c)
## Univariate beta mixture</pre>
```

```
## Univariate beta mixture
## Mixture Components:
## comp1 comp2 comp3 robust
## w 0.1946685 0.1940027 0.1113288 0.5000000
```

```
## a 46.5702699 72.0148367 3.5054258 1.0000000
## b 243.4141931 408.0689941 16.2800457 1.0000000
```

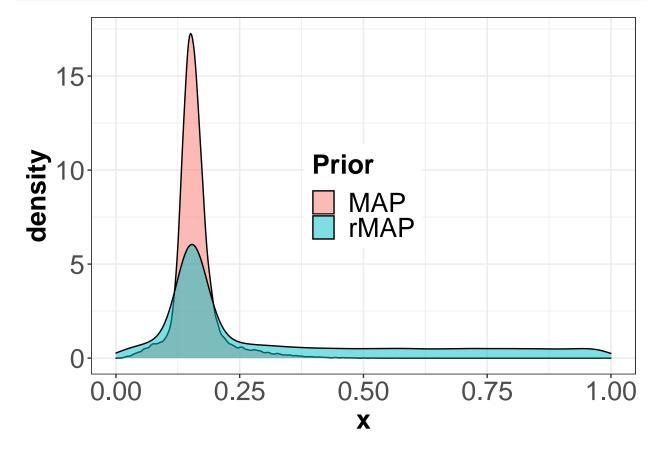
```
summary(rmap_c)
```

```
## mean sd 2.5% 50.0% 97.5%
## 0.33008857 0.26741587 0.04515052 0.18031091 0.95001226
```

We could compare the density functions of original and robust MAP prior. It can be seen that both priors have roughly the same mode. The density of rMAP prior has heavier tails, and therefore places more density on more extreme rates.

```
sample_MAP <- tibble(x=rmix(map_c_hat,100000), Prior = "MAP")
sample_rMAP <- tibble(x=rmix(rmap_c,100000), Prior = "rMAP")

rbind(sample_MAP, sample_rMAP) %>%
    ggplot(aes(x=x, fill=Prior)) + geom_density(alpha=0.5) +
    theme_bw() +
    theme(axis.title = element_text(face="bold",size=20),
        axis.text = element_text(size=20),
        legend.title=element_text(size=20,face="bold"),
        legend.text=element_text(size=20),
        legend.position = c(0.5, 0.5)
)
```



New Trial Design

Suppose that we are designing a new randomized two-arm proof-of-concept study in the same RA population. The endpoint remains the same, that is, ACR50 at week 12. Randomziation ratio is 2:1 - 60 in treatment and 30 in control. The robust MAP prior derived in previous section is used to augment the current control arm. We typically don't augment the treatment arm, and therefore assign a flat prior Beta(1,1) to p_T .

```
(prior_t = mixbeta(c(1,1,1)))
```

```
## Univariate beta mixture
## Mixture Components:
## comp1
## w 1
## a 1
## b 1
```

The operating characteristics are evaluated for current control rates between 0.11 and 0.21, guided by the predictive interval from meta-analysis. Assuming an improvement of 0.25 in ACR50 rate in the treatment arm, we may calculate the frequentist power that does not borrow historical data as a reference.

The trial is claimed a success if the single-criterion decision rule is met:

$$Pr(p_T - p_C > 0|Y_1, ..., Y_4, Y_C, Y_T) > 0.975.$$

The type I error and power can be calculated in RBesT package. To do that, we first define the decision rule then the design

```
(rule_single <- decision2S(pc = c(0.975), qc = c(0), lower.tail = F))

## 2 sample decision function
## Conditions for acceptance:
## P(x1 - x2 > 0) > 0.975

## Link: identity
design <- oc2S(prior_t, rmap_c, 60, 30, rule_single)</pre>
```

On a side note, a dual-criteria decision rule can also be defined, e.g.:

$$Pr(p_T - p_C > 0|Y_1, ..., Y_4, Y_C, Y_T) > 0.975$$

and

$$Pr(p_T - p_C > 0.25|Y_1, ..., Y_4, Y_C, Y_T) > 0.6.$$

```
decision2S(pc = c(0.975, 0.6), qc = c(0, 0.25), lower.tail = F)
```

```
## 2 sample decision function

## Conditions for acceptance:

## P(x1 - x2 > 0) > 0.975

## P(x1 - x2 > 0.25) > 0.6

## Link: identity
```

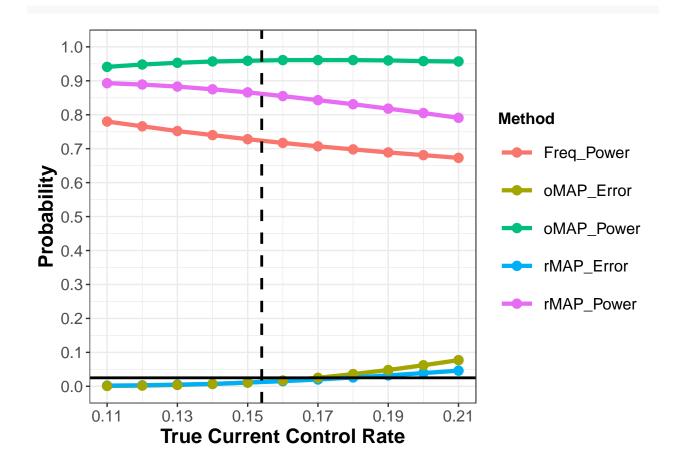
The "design" object is a function that computes operating characteristics given control and treatment rates. In the results shown below, type I error corresponds to an effect size of 0, while power corresponds to an effect size of 0.25. The operating characteristics corresponding to original MAP is also computed for comparison purpose.

Table 2: Operating Characteristics

CtrlRate	Freq_Error	rMAP_Error	oMAP_Error	Freq_Power	rMAP_Power	oMAP_Power
0.11	0.025	0.002	0.001	0.780	0.893	0.941
0.12	0.025	0.003	0.002	0.766	0.889	0.948
0.13	0.025	0.005	0.004	0.752	0.883	0.953
0.14	0.025	0.007	0.007	0.740	0.875	0.957
0.15	0.025	0.011	0.011	0.728	0.866	0.959
0.16	0.025	0.015	0.017	0.717	0.855	0.961
0.17	0.025	0.020	0.025	0.707	0.843	0.961
0.18	0.025	0.026	0.036	0.698	0.831	0.961
0.19	0.025	0.032	0.048	0.689	0.818	0.960
0.20	0.025	0.039	0.062	0.681	0.805	0.958
0.21	0.025	0.046	0.077	0.673	0.791	0.957

We also generate a plot that overlays the OCs. The horizontal solid line indicates the constant frequentist type I error $\alpha=0.025$. The dashed vertical line represents the meta-analysis point estimate of historical control data. Within the range of true current control rate, the power using robust MAP prior is about 10% higher than no borrowing. In the meantime, the type I error rate is very well contained. Robust MAP prior also demonstrates its robustness especially when current control rate is large.

```
plotdt <-
  ocdt %>% pivot_longer(!CtrlRate, names_to = "Method", values_to = "Prob") %>%
  filter(Method != "Freq Error")
plotdt %>%
  ggplot(aes(x=CtrlRate,y=Prob, color=Method)) +
  geom_line(size = 1.5) + geom_point(size = 3) +
  scale_x_continuous(breaks = seq(0.11, 0.21, 0.02), name = "True Current Control Rate") +
  scale_y_continuous(limits = c(0,1), breaks = seq(0,1,0.1), name = "Probability") +
  geom_vline(xintercept = 0.154, linetype=2, size=1) +
  geom_hline(yintercept = 0.025, linetype=1, size=1) +
  theme bw() +
  theme(axis.title = element_text(face="bold",size=15),
        axis.text = element_text(size=12),
        legend.title=element_text(size=12,face="bold"),
        legend.text=element text(size=12),
        legend.key.size = unit(1, "cm")
```



Analysis with Observed Current Trial Data

After the new trial is unblinded and analyzed, the robust MAP prior is updated with observed current control data. The posterior distribution is also a mixture of Beta components. Comparing with the prior mixture, the parameters of each Beta component is updated, and so is the weight w.

```
y_c <- 6
(post_c <- postmix(rmap_c, r=y_c, n=30))</pre>
## Univariate beta mixture
## Mixture Components:
##
     comp1
                 comp2
                              comp3
                                           robust
       0.3454043
                   0.3155126
                                0.1433055
                                             0.1957776
## a 52.5702699 78.0148367
                                9.5054258
                                             7.0000000
## b 267.4141931 432.0689941
                               40.2800457
                                            25.0000000
rmap_c
## Univariate beta mixture
## Mixture Components:
##
     comp1
                                           robust
                 comp2
                              comp3
## W
       0.1946685
                    0.1940027
                                0.1113288
                                             0.5000000
## a 46.5702699
                  72.0148367
                                3.5054258
                                             1.0000000
## b 243.4141931 408.0689941
                               16.2800457
                                             1.0000000
```

Likewise, the vague prior for p_T is updated with observed current treatment data.

```
y_t <- 30
(post_t <- postmix(prior_t, r=y_t, n=60))

## Univariate beta mixture
## Mixture Components:
## comp1</pre>
```

w 1 ## a 31 ## b 31

Thanks to the RBesT package, the mixture distribution (and their difference) in R works in the same way as other probability distributions, in the sense that d/p/q/r functions can be used. Therefore, we can use the *pmixdiff* command to evaluate the single-criterion decision rule.

```
pmixdiff(post_t, post_c, 0, lower.tail=FALSE)
```

```
## [1] 0.9993733
```

To verify, we can draw samples from respective posterior distributions and calculate the probability. The probability matches that calculated using pmixdiff. Either way, the current trial can be claimed successful.

```
sample_t <- rmix(post_t, 100000)
sample_c <- rmix(post_c, 100000)
mean((sample_t - sample_c) > 0)
```

[1] 0.99943

The point estimate and 95% credible interval for the control rate can be extracted from the summary statistics of the posterior mixture distribution.

```
## mean sd 2.5% 50.0% 97.5%
## 0.17518995 0.04776647 0.11497612 0.16329924 0.31042833
```