## Appendix II - R Code for Enhancing Patient Engagement in Preventative Care: A Field Experiment on the Effectiveness of Personalized and Generic Digital Outreach Strategies

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```
# Load and preview the first few rows of the data
# This gives a quick view of variables like variant_assignment, unsubscribed, opened, click, and visit
print(head(data))
##
        id variant_assignment last_seen unsubscribed opened click visit
##
     <int>
                       <char>
                                   <char> <int> <int> <int> <int> <int>
## 1:
      21
                                1/17/2025
                      placebo
                                                            1
                                                    1
## 2: 41 treatment_generic a while ago
                                                    1
## 3:
       44
                      placebo
                                9/13/2023
                                                     1
                                                            1
       78 treatment_generic a while ago
## 4:
                                                     1
## 5:
       105 treatment_generic a while ago
                                                    1
                                                                        Λ
                      placebo a while ago
d <- data
# Count the number of patients assigned to each variant group
# Helps confirm balance of randomization across control and treatment groups
data %>%
 count(variant_assignment)
##
         variant_assignment
##
                     <char> <int>
## 1:
                    control
                              108
## 2:
                    placebo
                              132
                              120
          treatment_generic
## 4: treatment_personalized
                              109
# Summarize click behavior by treatment arm
# Outputs total clicks and average click rate per group
d %>%
 group_by(variant_assignment) %>%
 summarise(n = n(), clicks = sum(click, na.rm = TRUE), click_rate = mean(click, na.rm = TRUE))
## # A tibble: 4 x 4
    variant_assignment
                               n clicks click_rate
```

<dbl>

<int> <int>

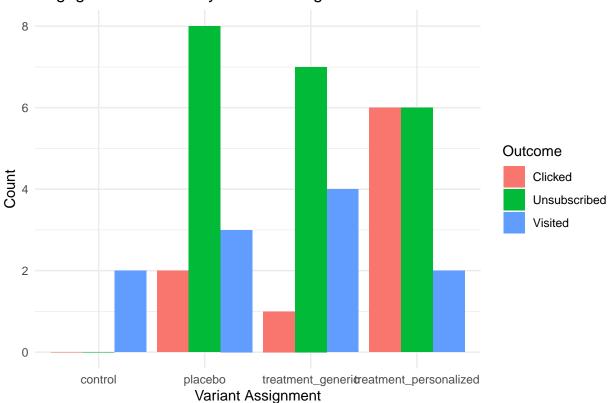
##

<chr>>

```
## 1 control 108 0 0
## 2 placebo 132 2 0.0152
## 3 treatment_generic 120 1 0.00833
## 4 treatment_personalized 109 6 0.0550
```

```
# Summarize key engagement outcomes by treatment group
# Includes counts of clicks, visits, and unsubscribes for plotting
summary counts <- d %>%
  group_by(variant_assignment) %>%
  summarise(
   Clicked = sum(click),
   Visited = sum(visit),
   Unsubscribed = sum(unsubscribed)
  ) %>%
 pivot_longer(
   cols = c(Clicked, Visited, Unsubscribed),
   names_to = "Outcome",
   values_to = "Count"
  )
# Plot bar chart comparing counts of engagement metrics by variant group
# Visualizes Clicked, Visited, and Unsubscribed across Control, Placebo, and Treatment groups
ggplot(summary_counts, aes(x = variant_assignment, y = Count, fill = Outcome)) +
  geom_bar(stat = "identity", position = position_dodge()) +
  labs(title = "Engagement Metrics by Variant Assignment", x = "Variant Assignment", y = "Count") +
 theme_minimal()
```

## **Engagement Metrics by Variant Assignment**



```
# Filter data to include only treatment-related groups (excluding control)
# Used for comparing email open rates across different outreach strategies
data_subset_1 <- data %>%
  filter(variant_assignment %in% c("placebo", "treatment_generic", "treatment_personalized"))
# Estimate linear model to compare open rates across variant_assignment
# Robust standard errors account for heteroskedasticity
model opened <- lm(opened ~ variant assignment, data = data subset 1)
robust_se <- coeftest(model_opened, vcov = vcovHC(model_opened, type = "HC1"))</pre>
print(robust se)
##
## t test of coefficients:
##
##
                                             Estimate Std. Error t value Pr(>|t|)
## (Intercept)
                                            0.9924242 0.0075786 130.9515
## variant assignmenttreatment generic
                                                                            0.3182
                                            0.0075758 0.0075786 0.9996
## variant_assignmenttreatment_personalized 0.0075758 0.0075786
                                                                   0.9996 0.3182
## (Intercept)
                                            ***
## variant_assignmenttreatment_generic
## variant_assignmenttreatment_personalized
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
# Intention-to-treat analysis: calculate open rate and sample size per variant
# Used to assess compliance and attrition
itt_open <- data_subset_1 %>%
  group_by(variant_assignment) %>%
  summarise(open_rate = mean(opened == 1), count = n())
print(itt_open)
## # A tibble: 3 x 3
   variant assignment
                            open rate count
    <chr>
##
                                <dbl> <int>
## 1 placebo
                                0.992
                                        132
## 2 treatment_generic
                                        120
## 3 treatment personalized
                                        109
# Filter to just treatment_generic and treatment_personalized groups
# Fit model to compare unsubscribe rates between personalized and generic emails
data_subset_2 <- data %>%
  filter(variant assignment %in% c("treatment generic", "treatment personalized"))
model_unsubscribed <- lm(unsubscribed ~ variant_assignment, data = data_subset_2)</pre>
# Output summary with robust standard errors (HC1)
coeftest(model_unsubscribed, vcov = vcovHC(model_unsubscribed, type = "HC1"))
## t test of coefficients:
##
```

```
##
                                             Estimate Std. Error t value Pr(>|t|)
## (Intercept)
                                            ## variant assignmenttreatment personalized -0.0032875 0.0307116 -0.1070 0.914849
## (Intercept)
## variant assignmenttreatment personalized
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
# Filter to just treatment_generic and treatment_personalized groups
# Fit model to compare click rates between personalized and generic emails
data subset 3 <- data %>%
 filter(variant_assignment %in% c("treatment_generic", "treatment_personalized"))
model_click <- lm(click ~ variant_assignment, data = data_subset_3)</pre>
# Output summary with robust standard errors (HC1)
coeftest(model_click, vcov = vcovHC(model_click, type = "HC1"))
## t test of coefficients:
##
##
                                            Estimate Std. Error t value Pr(>|t|)
## (Intercept)
                                           0.0083333 0.0083350 0.9998 0.31847
## variant_assignmenttreatment_personalized 0.0467125 0.0234710 1.9902 0.04777
##
## (Intercept)
## variant_assignmenttreatment_personalized *
## ---
## Signif. codes: 0 '***' 0.001 '**' 0.05 '.' 0.1 ' ' 1
# Filter to just control, treatment_qeneric and treatment_personalized groups
# Fit model to compare visit rates between personalized and generic emails
data_subset_4 <- data %>%
 filter(variant_assignment %in% c("control", "treatment_generic", "treatment_personalized"))
model_visit <- lm(visit ~ variant_assignment, data = data_subset_4)</pre>
# Output summary with robust standard errors (HC1)
coeftest(model_visit, vcov = vcovHC(model_visit, type = "HC1"))
##
## t test of coefficients:
##
##
                                              Estimate Std. Error t value
                                            0.01851852 0.01303089 1.4211
## (Intercept)
## variant_assignmenttreatment_generic
                                            0.01481481 0.02099368 0.7057
## variant_assignmenttreatment_personalized -0.00016989 0.01834491 -0.0093
##
                                           Pr(>|t|)
## (Intercept)
                                             0.1562
## variant_assignmenttreatment_generic
                                             0.4809
## variant_assignmenttreatment_personalized
                                             0.9926
```

```
# Add covariate for recency of healthcare interaction, and generate interaction term with treatment.
# Enables HTE analysis to see if treatment works differently for patients not recently seen.
today <- as.Date("2025-04-15")</pre>
d <- d %>%
 mutate(last_seen = ifelse(last_seen == "a while ago", '04/01/2024', last_seen))
d <- d %>%
  mutate(
    # Step 2: Convert from m/d/Y to Date
   last_seen_clean = mdy(last_seen),
    # Step 3: Compute days since seen
   days_since_seen = as.numeric(Sys.Date() - last_seen_clean),
   # Step 4: Define seen_recently as 365 days
   seen_recently = ifelse(days_since_seen <= 365, 1, 0)</pre>
 )
# Fit a model including an interaction between treatment assignment and not_seen_recently.
# Helps test whether timing of last engagement modifies treatment effects.
# Set 'not seen recently' (0) as the reference level
# Create a new variable: not seen recently
d$not_seen_recently <- as.numeric(d$seen_recently == 0)</pre>
# Filter out the placebo group
d_filtered <- d %>% filter(variant_assignment %in% c("control", "treatment_generic", "treatment_persona
# Create a new variable: not_seen_recently
d_filtered$not_seen_recently <- as.numeric(d_filtered$seen_recently == 0)</pre>
# Fit the model using the new variable
model_HTE <- lm(visit ~ variant_assignment * not_seen_recently, data = d_filtered)</pre>
robust_se <- vcovHC(model_HTE, type = "HC1")</pre>
robust_results <- coeftest(model_HTE, vcov = robust_se)</pre>
print(robust_results)
##
## t test of coefficients:
##
                                                                Estimate Std. Error
##
                                                                0.083333 0.056926
## (Intercept)
## variant_assignmenttreatment_generic
                                                               -0.018817 0.072269
                                                               -0.011905 0.075182
## variant_assignmenttreatment_personalized
## not_seen_recently
                                                               -0.083333 0.056926
## variant_assignmenttreatment_generic:not_seen_recently
                                                                0.041289 0.073987
## variant_assignmenttreatment_personalized:not_seen_recently 0.011905 0.075182
                                                               t value Pr(>|t|)
##
## (Intercept)
                                                                1.4639 0.1442
                                                               -0.2604 0.7947
## variant_assignmenttreatment_generic
## variant_assignmenttreatment_personalized
                                                               -0.1583 0.8743
```

```
## not seen recently
                                                                 -1.4639
                                                                           0.1442
## variant_assignmenttreatment_generic:not_seen_recently
                                                                  0.5581
                                                                           0.5772
## variant assignmenttreatment personalized:not seen recently 0.1583
                                                                           0.8743
# Simulate statistical power at different sample sizes for three effect sizes: 0.005, 0.01, 0.02.
# Helps visualize sensitivity of the study to detect effects under realistic assumptions.
set.seed(3)
power_test_t <- function(</pre>
 mean_control = 0.02,
 mean_treat = 0.03,
  sd_control = sqrt(0.02 * 0.98),
  sd_treat = sqrt(0.02 * 0.98),
  number_per_condition = 40,
  power_loops = 1000,
  verbose = FALSE) {
    p_values <- numeric(power_loops)</pre>
    for (i in 1:power_loops) {
        control_group <- rnorm(number_per_condition, mean = mean_control, sd = sd_control)</pre>
        treatment group <- rnorm(number per condition, mean = mean treat, sd = sd treat)
        test_result <- t.test(control_group, treatment_group, var.equal = FALSE)</pre>
        p_values[i] <- test_result$p.value</pre>
    }
    power_estimate <- mean(p_values < 0.05)</pre>
    return(list('p_values' = p_values, 'power' = power_estimate))
# Generate line plot showing how sample size impacts power for each effect size.
# A horizontal line at 0.8 indicates the standard threshold for acceptable power.
# Sample sizes
samples_per_condition <- c(100, 200, 300, 300, 400, 500, 600, 700, 800, 900, 1000, 1200, 1500, 1750, 20
# Calculate power for treatment effects of +0.005, +0.01, +0.02
power_005 <- numeric(length(samples_per_condition))</pre>
power_01 <- numeric(length(samples_per_condition))</pre>
power_02 <- numeric(length(samples_per_condition))</pre>
for(i in seq_along(samples_per_condition)) {
  power_005[i] <- power_test_t(mean_control = 0.02, mean_treat = 0.025,</pre>
                                 number_per_condition = samples_per_condition[i])$power
 power_01[i] <- power_test_t(mean_control = 0.02, mean_treat = 0.03,</pre>
                                number_per_condition = samples_per_condition[i])$power
 power_02[i] <- power_test_t(mean_control = 0.02, mean_treat = 0.04,</pre>
                                number_per_condition = samples_per_condition[i])$power
}
# Assemble power table
power_table <- data.frame(</pre>
 "Sample Size" = samples per condition,
 "ATE 0.005" = power 005,
```

## Power Curves for ATEs of 0.005, 0.01, 0.02

