Unwrapping Customer Delight

Milestone #3 Meeting: The Analysis Phase

The Estée Lauder Companies







• What is an RCT?

- Randomized Controlled Trial = split customers into two groups at random
- Control group (0): does not get the surprise gift
- Treatment group (1): does get the surprise gift

Why randomization matters:

- Makes groups statistically comparable on average (age, tenure, loyalty, spending history)
- Prevents bias that could occur if we "cherry-picked" who gets gifts

Measuring the effect:

- Average Treatment Effect (ATE) = difference in mean outcomes between treatment and control
- \circ Formula: Δ = mean(revenue | treatment=1) mean(revenue | treatment=0)
- \circ Example: If treated customers spend \$110 on average and control spend \$100, then ATE = +\$10
- **Big challenge**: Real data is noisy → harder to detect real differences without large enough samples





Recap: Why Statistical Power Matters

- Statistical power = probability that our experiment correctly detects a real effect
- If power is too low → risk of false negatives (we conclude "no effect" when there actually is one)
- If sample size is too big → wasted resources, longer experiments, possible negative customer experiences
- The 4 levers of power analysis:
 - Minimum Detectable Effect (MDE): The smallest change we care about finding (e.g., +1% revenue lift)
 - Significance level (α): Chance of false positive. Standard = 5%
 - \circ **Power (1–\beta)**: Chance of catching a real effect. Standard = 80–90%
 - o Sample size (N): How many customers we need in the experiment

How they interact:

- Smaller MDE → need larger N
- Higher power (90% vs. 80%) → need larger N
- Lower variance → smaller N is enough
- Takeaway: Designing experiments = balancing these levers to be efficient but still scientifically valid





Recap: Variance Reduction w/ MLy RATE Hatters:

- Outcome data (like revenue) has high natural variability
- This "noise" makes it harder to see the treatment effect

• Solution: Regression adjustment (MLRATE)

- Step 1: Use ML to predict expected revenue based on pre-experiment covariates (tenure, loyalty, prior spend, etc.)
- \circ Step 2: Use those predictions (G(x)) as control variables in the treatment effect regression
- Step 3: Estimate treatment effect on the "leftover" variation (residuals) after accounting for predictable patterns

• Key assumptions:

- G(x) is strongly correlated with revenue
- \circ G(x) is independent of treatment (since assignment is random)

Benefit:

- Reduces unexplained variance → increases recall
- Means fewer customers are needed to detect the same effect
 In practice: MLRATE is like turning down the "noise" so the gift signal comes through more clearly





What You Completed for Milestone #3

You ran two types of power analysis:

1. Standard t-test method

- Effect size = (MDE) ÷ (std. dev. of revenue).
- Sample size based on all variance in the outcome

2. MLRATE-adjusted method

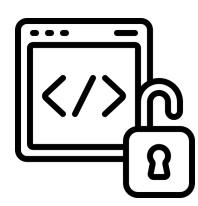
- Effect size = (MDE) ÷ (std. dev. of residuals).
- Sample size based on reduced variance (after regression adjustment)

• Expected outcome:

- MLRATE gave you a smaller required sample size
- \circ Your residual distribution plot shows how much variance was explained by G(x)

Big takeaway:

- Power analysis tells us how many customers we need
- MLRATE lets us run experiments more efficiently (same power, smaller N)







1. Execution

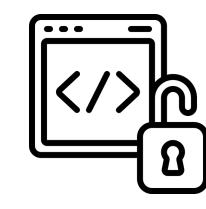
- We've determined the optimal sample size, given the data and business requirements
- Now, we apply our intervention. For example:
 - 1. For each user on the checkout page, randomly assign them into the treatment or control group.
 - 2. Offer a free gift to each customer in the treatment group.
 - 3. Repeat until we have reached the required sample size.
 - 4. Wait until t time has passed
 - 5. Collect the experiment data, wait until t time has passed:
 - a. Revenue: during the experiment period (i.e. time t).
 - b. Covariates: The same ones we used in the power analysis, during the *pre-experiment* period (i.e. time t-1).

We will simulate the experiment execution phase and provide you with the above experiment data.

2. Analysis

Now that we have the experiment data:

- 1. Conduct EDA to validate the data and get an initial, visual read of the treatment effect.
- 2. Estimate the treatment effect using MLRATE.





Milestone #4: Moving to Experiment Data

- What data you'll get (via email):
 - File: experiment_results.parquet
 - Each row = a simulated customer in our experiment (aka gift giving strategy)

Columns included:

- customer_id → unique customer identifier
- aov → average order value before the intervention
- o days_since_last_purchase → recency of last purchase (days)
- tenure_in_days → how long the customer has been active
- loyalty_membership → categorical/boolean flag for loyalty program
- assignment → treatment assignment indicator (0 = control, 1 = treatment)
- revenue (t) → post-intervention revenue (outcome variable of interest)





EDA on Experiment Data

• EDA goals:

- Data quality checks: row counts, datatypes, any missing values or duplicates?
- o **Univariate exploration**: histograms & boxplots for revenue (t), aov, tenure_in_days, days_since_last_purchase.
- Treatment vs. control comparisons:
 - Verify covariate balance across assignment
 - Expectation: Covariates should look *statistically similar*, since assignment was random
 - Compare distributions of revenue (t) between groups
 - This should give you an early visual sense of the treatment effect
- **Relationships**: scatterplots (e.g., aov vs. revenue (t) colored by assignment), grouped box/violin plots (e.g., revenue by loyalty_membership × assignment to see heterogeneity of effects)
- Skew/outliers: check for long-tail distributions

Meeting will also cover:

A lecture on how to estimate the treatment effect (theory)



Project milestones and timeline

These are the milestones for your Challenge Project. They include the <u>CRISP-DM</u> process steps you learned about in your ML Foundations course. In addition, there is an educational component in the front-end.

- EDA on pre-experiment data
- Present EDA findings and data preprocessing

MILESTONE 2

- You will have received the experiment data
- Perform EDA on experimental data
- Present EDA findings and data preprocessing

MILESTONE 4

Estimate MLRATE

- Compare to:
 - Standard t-test ATE
 - Pre-experiment revenue controlled ATE

MILESTONE 6

MILESTONE 1

- Review material from "Helpful resources" slide
- Q&A with challenge advisors

MILESTONE 3

- Power analysis
 - Determine the variance-reduced sample size for the upcoming RCT
 - Compare to standard t-test sample size

MILESTONE 5

- Estimate ATE using two approaches:
 - First by using the standard t-test
 - Then by controlling for pre-experimental revenue

MILESTONE 7

- Deliver final project presentation:
 RCT, MLRATE, key findings
- Present quantified gift impact and campaign recommendations
- Final Q&A and project recap