#### Preregistration

# Preregistration Practice for Living Data Project 2025 Productivity and Reproducibility Course

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## **Study Information**

This is a practice preregistration using the "palmerpenguins" dataset from the R package of the same name.

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#### Description For this project, the raw data was extracted from the "palmerpenguins" R package,

and saved as a .csv to a raw data folder. That is the full extent of data use in this project, and therefore no further elements of this preregistration template will be used.

#### Hypotheses

Enter your response here.

## Design Plan

#### Study type

**Experiment**. A researcher randomly assigns treatments to study subjects, this includes field or lab experiments. This is also known as an intervention experiment and includes randomized controlled trials.

**Observational Study**. Data is collected from study subjects that are not randomly assigned to a treatment. This includes surveys, natural experiments, and regression discontinuity designs.

Meta-Analysis. A systematic review of published studies.

Other. Please explain.

#### Blinding

No blinding is involved in this study.

For studies that involve human subjects, they will not know the treatment group to which they have been assigned.

Personnel who interact directly with the study subjects (either human or non-human subjects) will not be aware of the assigned treatments.

Personnel who analyze the data collected from the study are not aware of the treatment applied to any given group.

#### Study design

Enter your response here.

#### Randomization

Enter your response here.

### Sampling Plan

#### Existing data

Registration prior to creation of data. As of the date of submission of this research plan for preregistration, the data have not yet been collected, created, or realized.

Registration prior to any human observation of the data. As of the date of submission, the data exist but have not yet been quantified, constructed, observed, or reported by anyone - including individuals that are not associated with the proposed study. Examples include museum specimens that have not been measured and data that have been collected by non-human collectors and are inaccessible.

Registration prior to accessing the data. As of the date of submission, the data exist, but have not been accessed by you or your collaborators. Commonly, this includes data that has been collected by another researcher or institution.

Registration prior to analysis of the data. As of the date of submission, the data exist and you have accessed it, though no analysis has been conducted related to the research plan (including calculation of summary statistics). A common situation for this scenario when a large dataset exists that is used for many different studies over time, or when a data set is randomly split into a sample for exploratory analyses, and the other section of data is reserved for later confirmatory data analysis.

Registration following analysis of the data. As of the date of submission, you have accessed and analyzed some of the data relevant to the research plan. This includes preliminary analysis of variables, calculation of descriptive statistics, and observation of data distributions. Please see <a href="https://cos.io/prereg">https://cos.io/prereg</a> for more information.

Explanation of existing data

Enter your response here.

Data collection procedures

Enter your response here.

Sample size	Enter your response here.
Sample size rationale	Enter your response here.
Stopping rule	Enter your response here.
	Variables
Manipulated variables	Enter your response here.
Measured variables	Enter your response here.
Indices	Enter your response here.
	Analysis Plan
Statistical models	Analysis Plan  Enter your response here.
Statistical models  Transformations	
	Enter your response here.
Transformations	Enter your response here.
Transformations Inference criteria	Enter your response here.  Enter your response here.

# Other

Other (Optional) Enter your response here.

# References