

Study Design and Data Collection

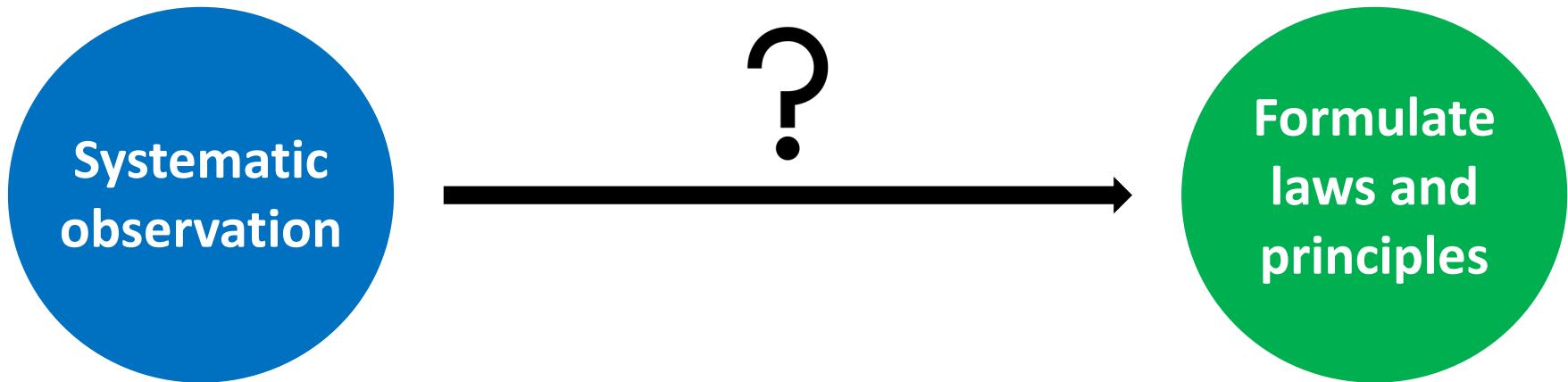
Ecological and Epidemiological Modeling in Madagascar

Centre ValBio, Ranomafana, January 2020

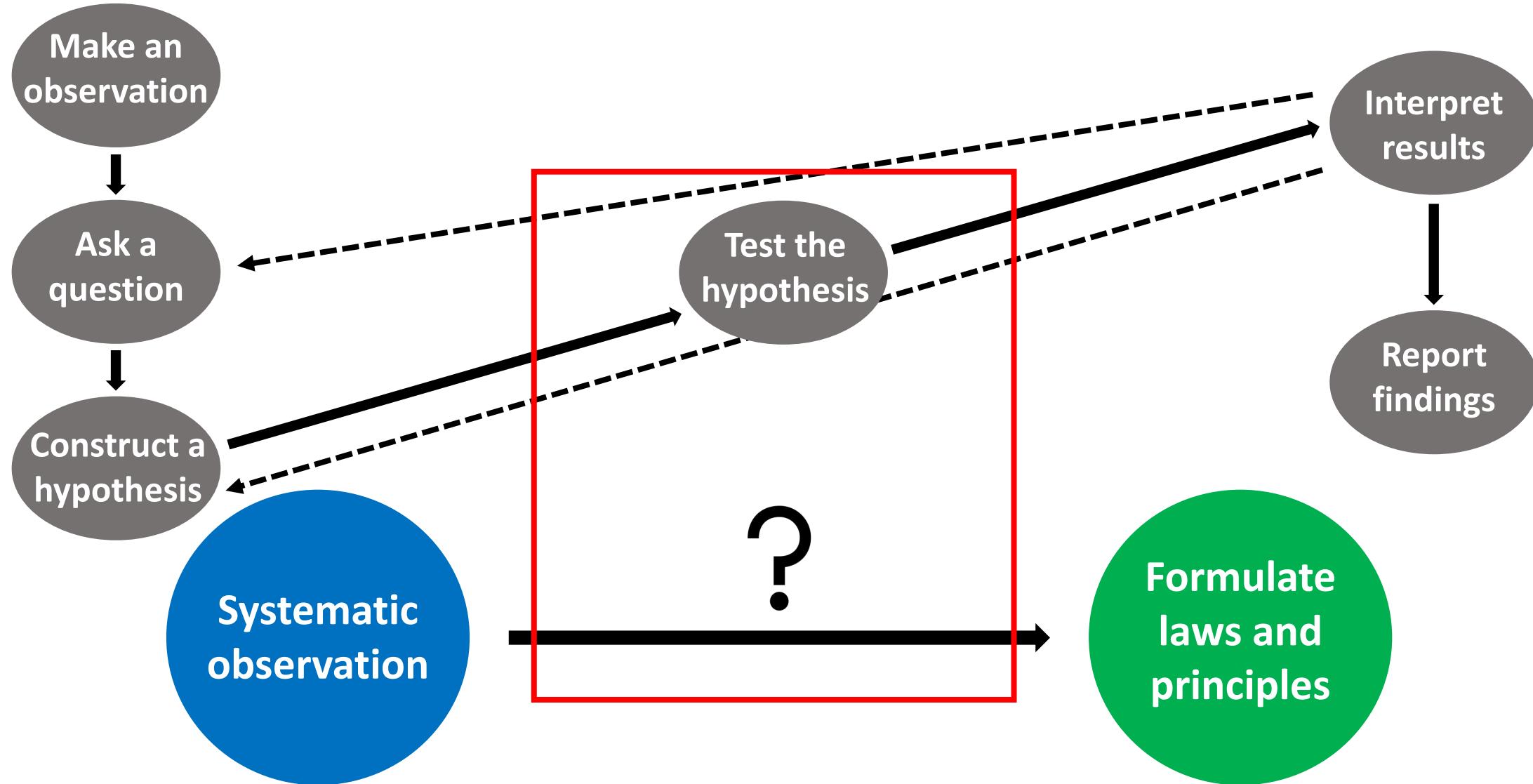
What *is* science?

the **systematic observation** of natural events and conditions in order to **discover facts** about them and to **formulate laws and principles** based on these facts

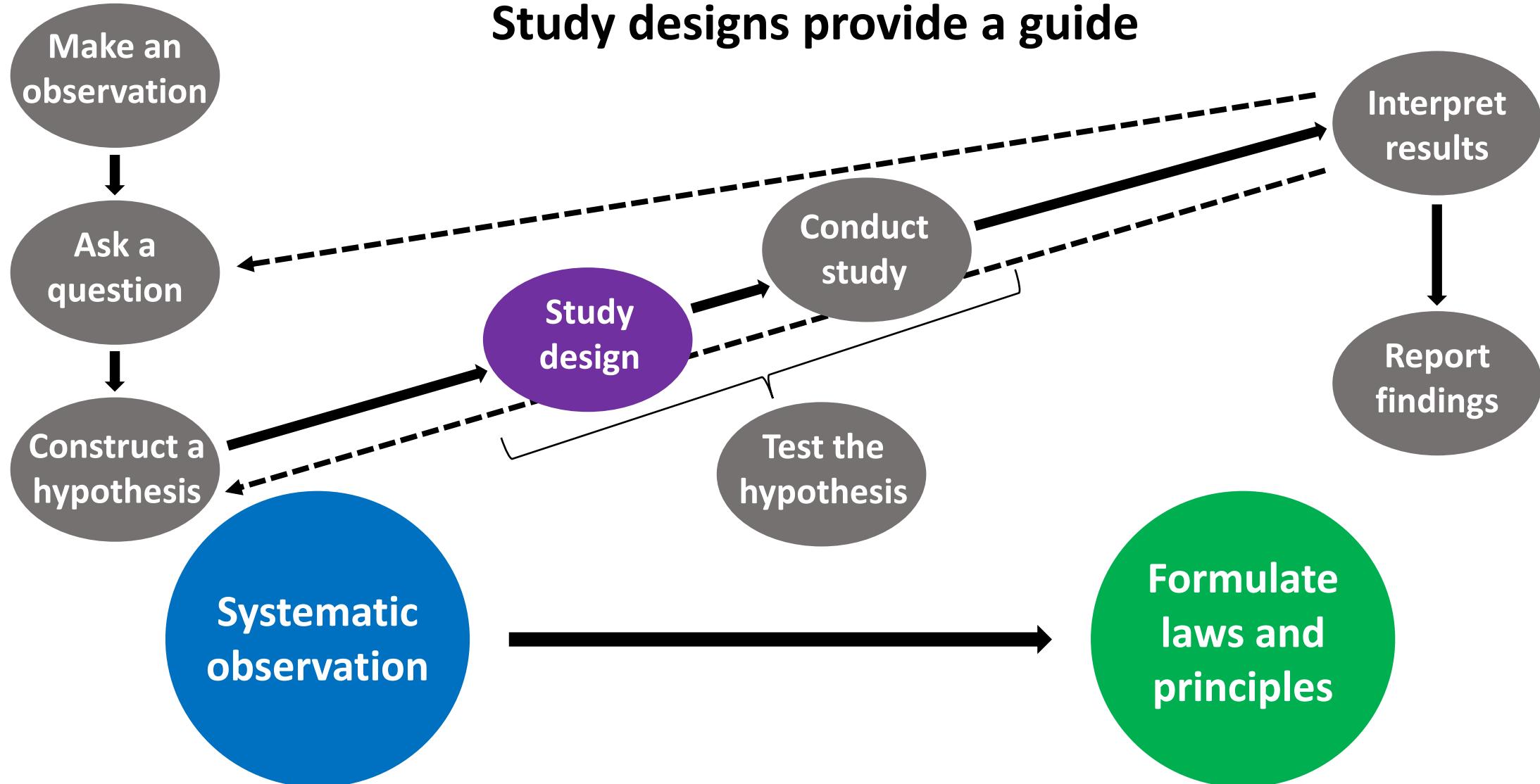
-- Academic Press Dictionary of Science and Technology



How to do science?



How to do science?



Goals for this lecture

- Outline the study design process
- Understand why study design helps us “do” science
- R tutorial: sample and data organization with *Ekipa Fanihy*

Study Design

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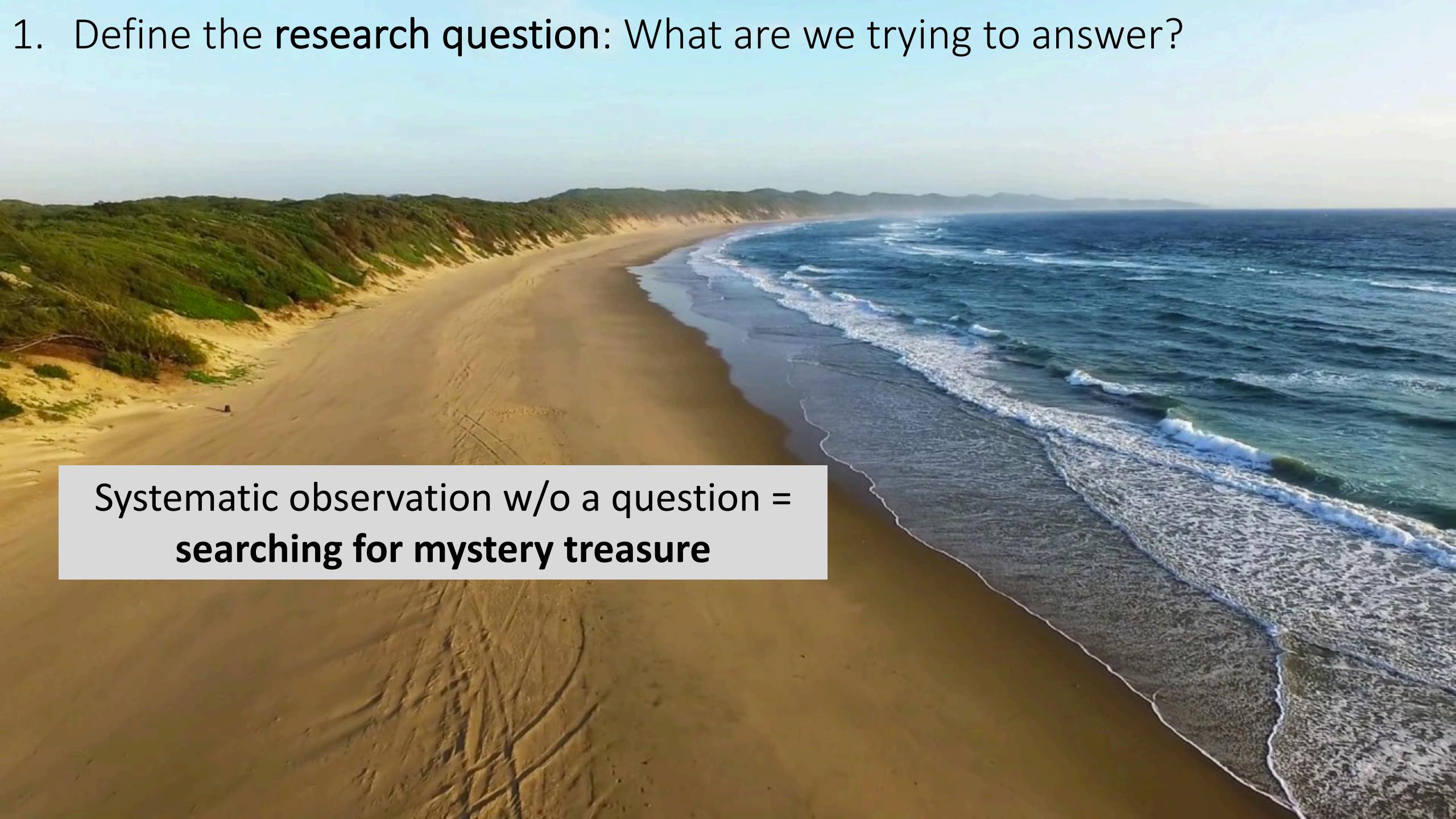
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6. Outline a **data organization plan**: How should we organize our data?
7. Be **flexible**: How can we prepare for potential/unanticipated challenges?

1. Define the research question: What are we trying to answer?



Mahaliana
IT ALWAYS STARTS WITH A QUESTION

1. Define the research question: What are we trying to answer?

An aerial photograph of a long, sandy beach stretching towards the horizon. The beach is bordered by green dunes on the left and the ocean on the right. The water is a deep blue with white-capped waves breaking near the shore. The sky is clear and light blue.

Systematic observation w/o a question =
searching for mystery treasure

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Every component of study design hinges on the research question

- **Research question** = the target
- **Study design** = an effective and efficient route to answering the question



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Good research questions are:

- Clear
- Focused and testable
- Original
- Based on previous observations



1. Define the research question: What are we trying to answer?

- Formulate a **hypothesis**
- Develop a **model** to demonstrate your hypothesis

Every component of study design hinges on the research question

- **Research question** = the target
- **Study design** = an effective and efficient route to answering the question

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1. Define the research question: What are we trying to answer?

How do bat populations maintain
virulent human-infection viruses?



Photo by Cara Brook

1. Define the research question: What are we trying to answer?

How do bat populations maintain
virulent human-infection viruses?

broad

↓
Focused/
testable

What is the **force of infection** of
henipaviruses in Madagascar
Eidolon dupreanum populations?



2. Define the sample type: What data do we need to answer our question?

What is **data**?

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What is data?

= **evidence** to support a **claim**

Force of infection =
Rate at which bats
become infected

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Data:

- Age data
- Serology



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Sample:

- Teeth
- Serum



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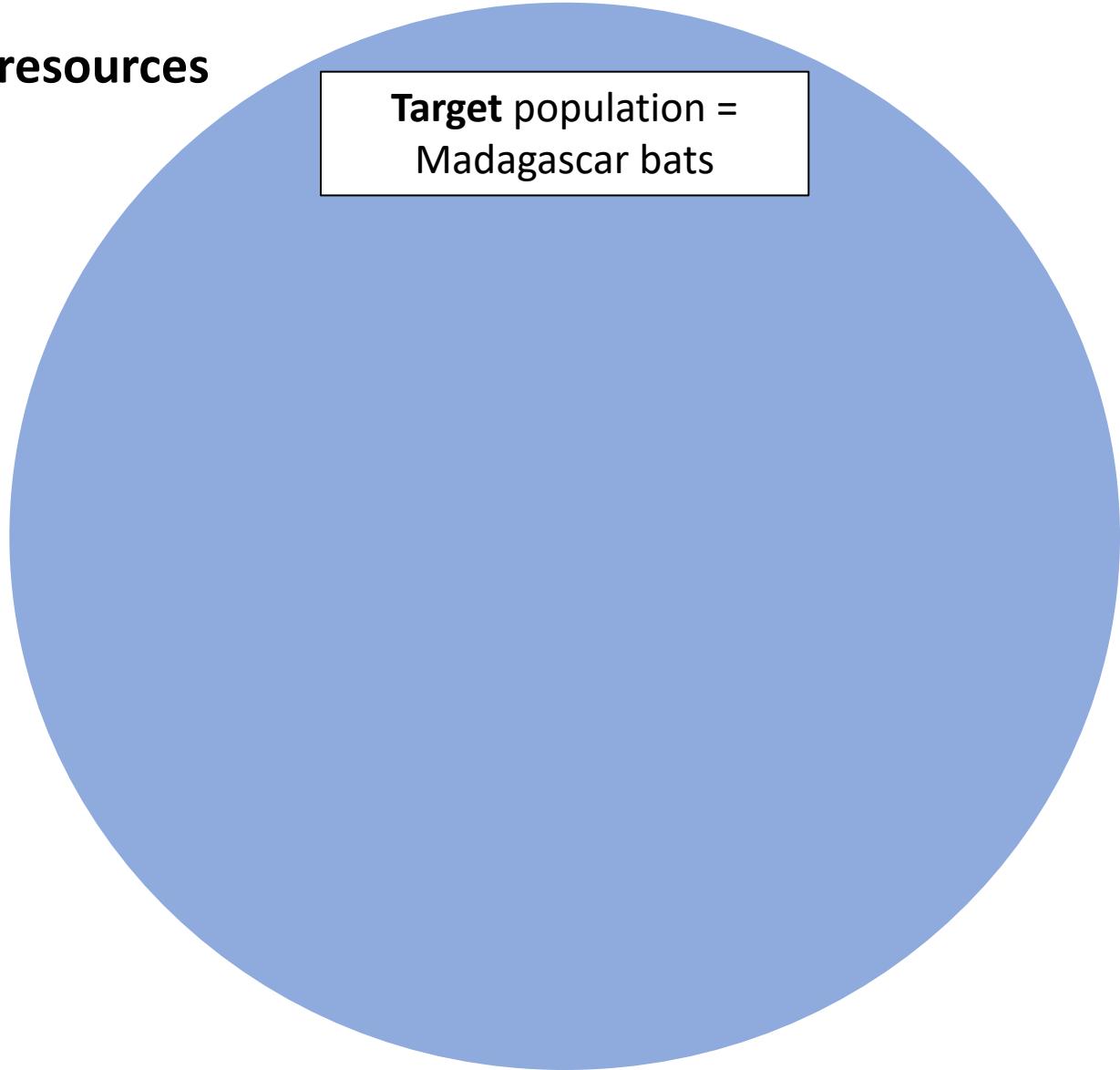
3. Identify a system: Where can we collect our data?

Choosing a study population that allows you to **answer your research question**:

- **effectively**
- **tractably**—time, money, and effort are **limited resources**

Target population:

Want to make inferences about



Target population =
Madagascar bats

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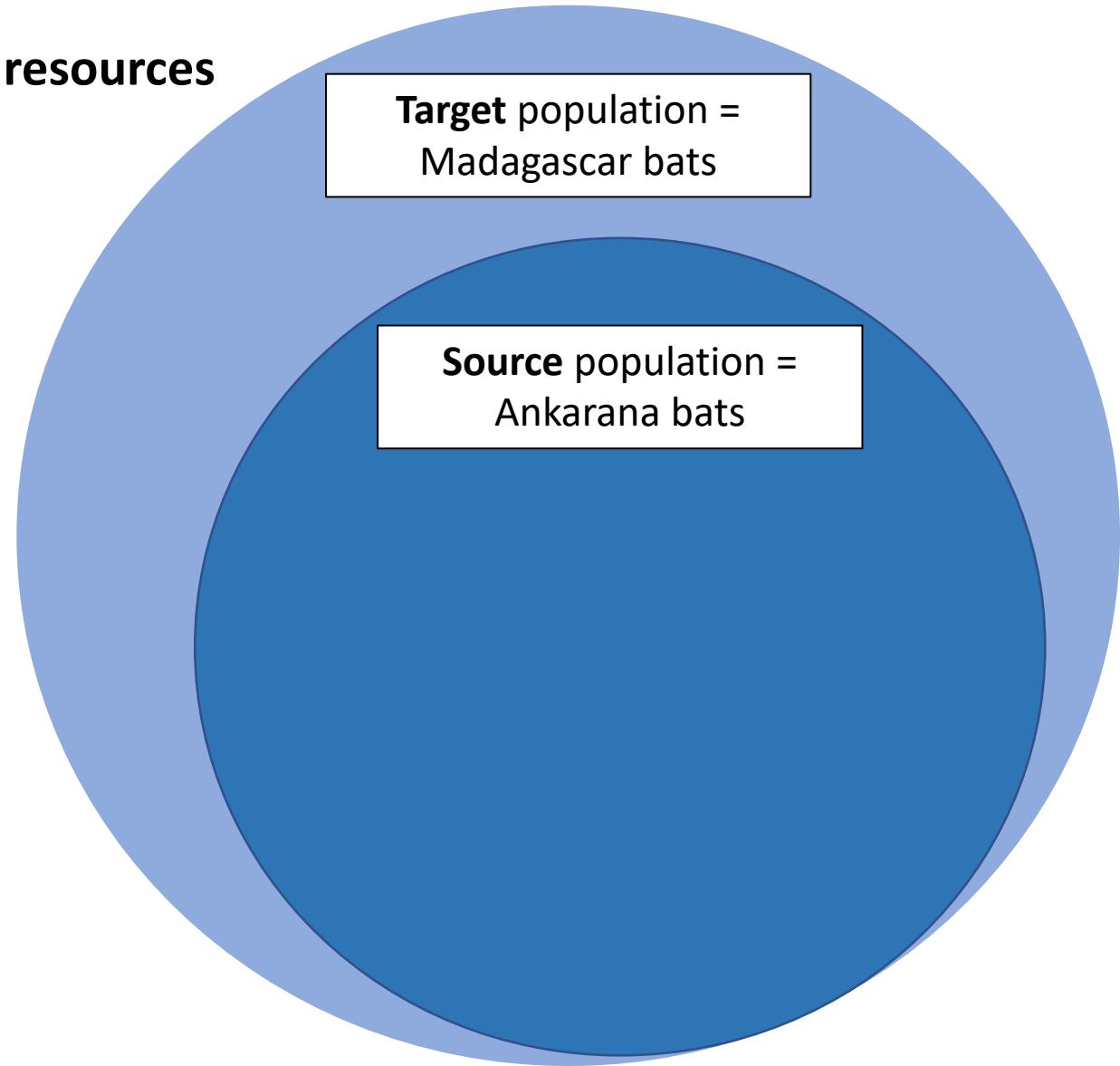
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Source population:

Choosing study population



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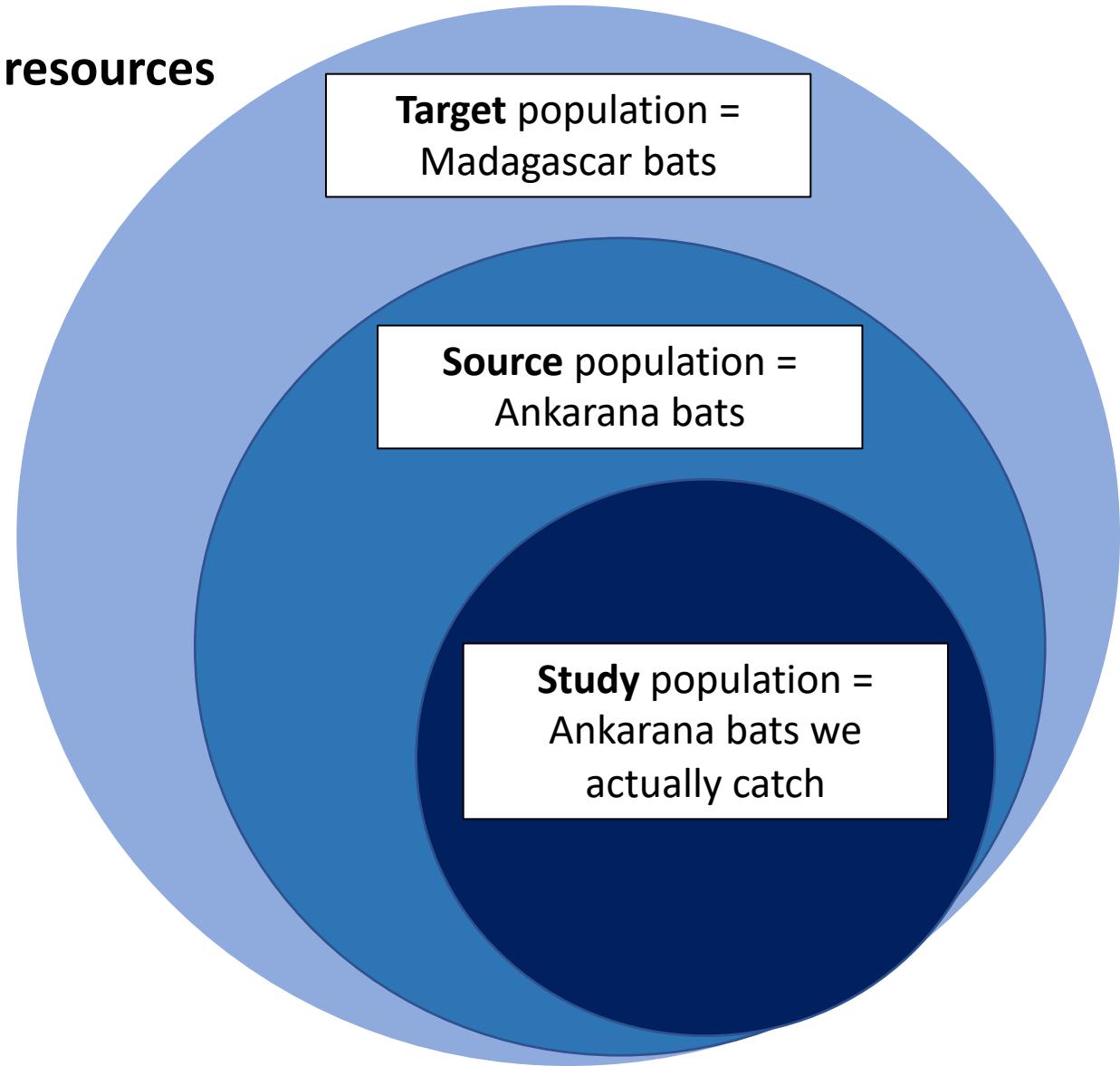
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Source population:

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Study population:

Sampled individuals



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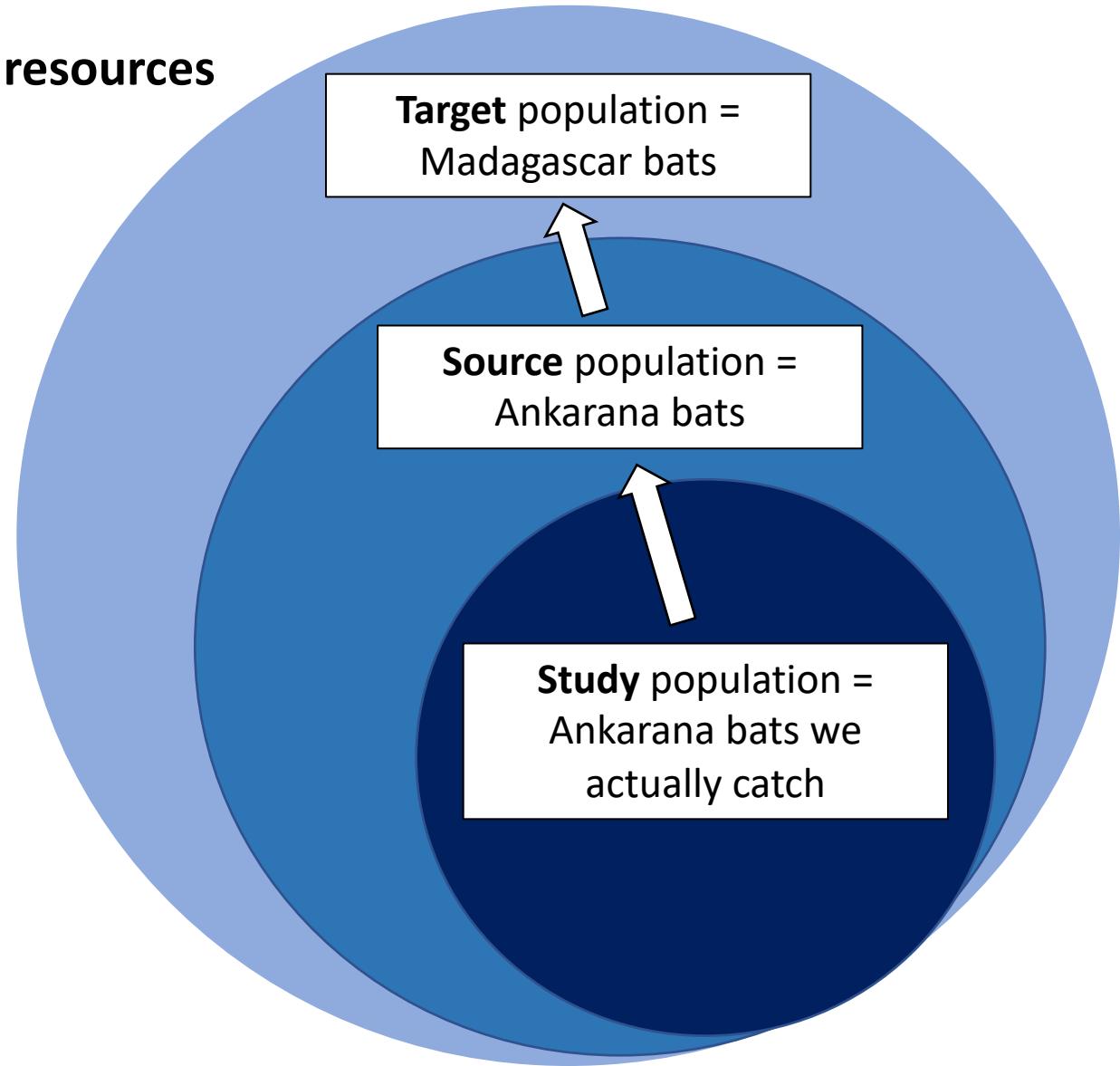
Source population:

Choosing study population

Study population:

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Well designed studies allow us to make inference about the target population



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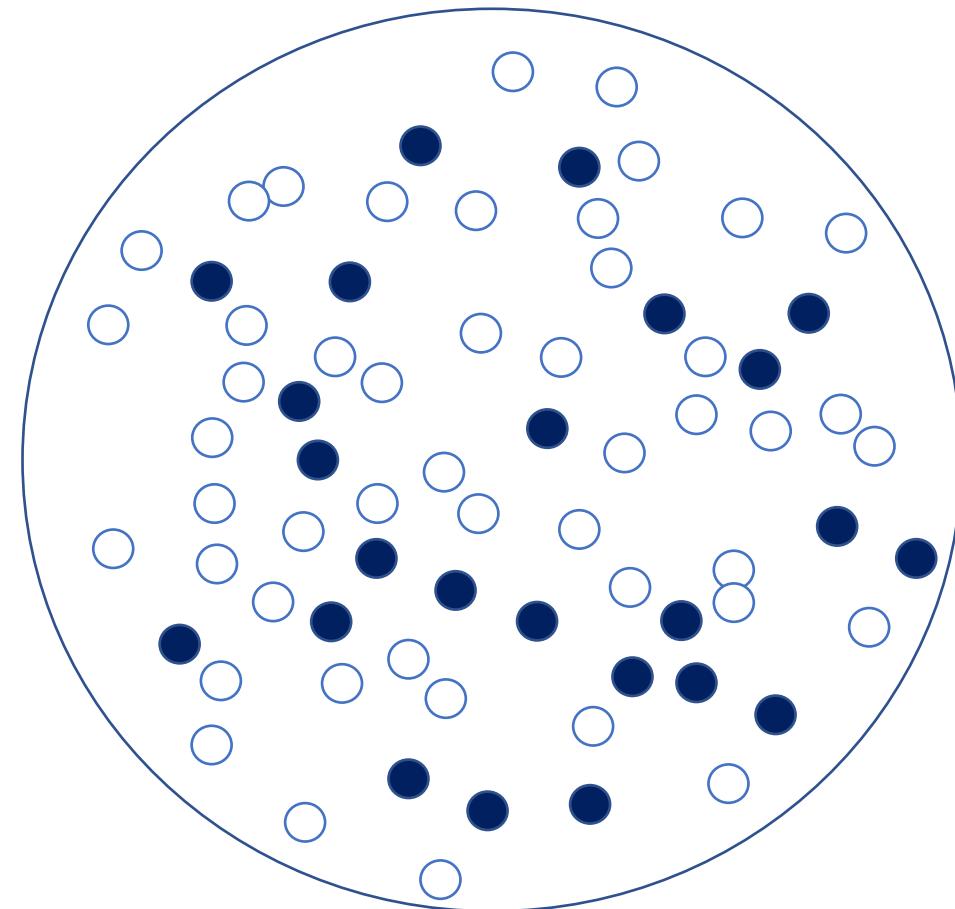
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4. Choose a sampling scheme: How should we collect our data?

The **study** population should be **randomly selected** from the **source** population



Choose a sampling scheme: How should we collect our data?

Observational

- Descriptive
- Cross-sectional
- Longitudinal
- Ecological

There are several **study types** to choose from...
but not all types will be able to answer your question

Experimental

- Experimental Ecology
- Randomized Control Trial (RCT)

Choose a sampling scheme: How should we collect our data?

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Observational vs. Experimental?

Experimental

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- Randomized Control Trial (RCT)

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Descriptive studies = observational research that describes the **characteristics** of a population

- focus on the **what** instead of the **why**

Good for **generating hypotheses**, especially when data is limited

- **Bat survey:** we need to find the bats before we can study their viruses

Experimental

- Randomized Control Trial (RCT)

Not all studies are hypothesis-driven...
but research really does always start with a question

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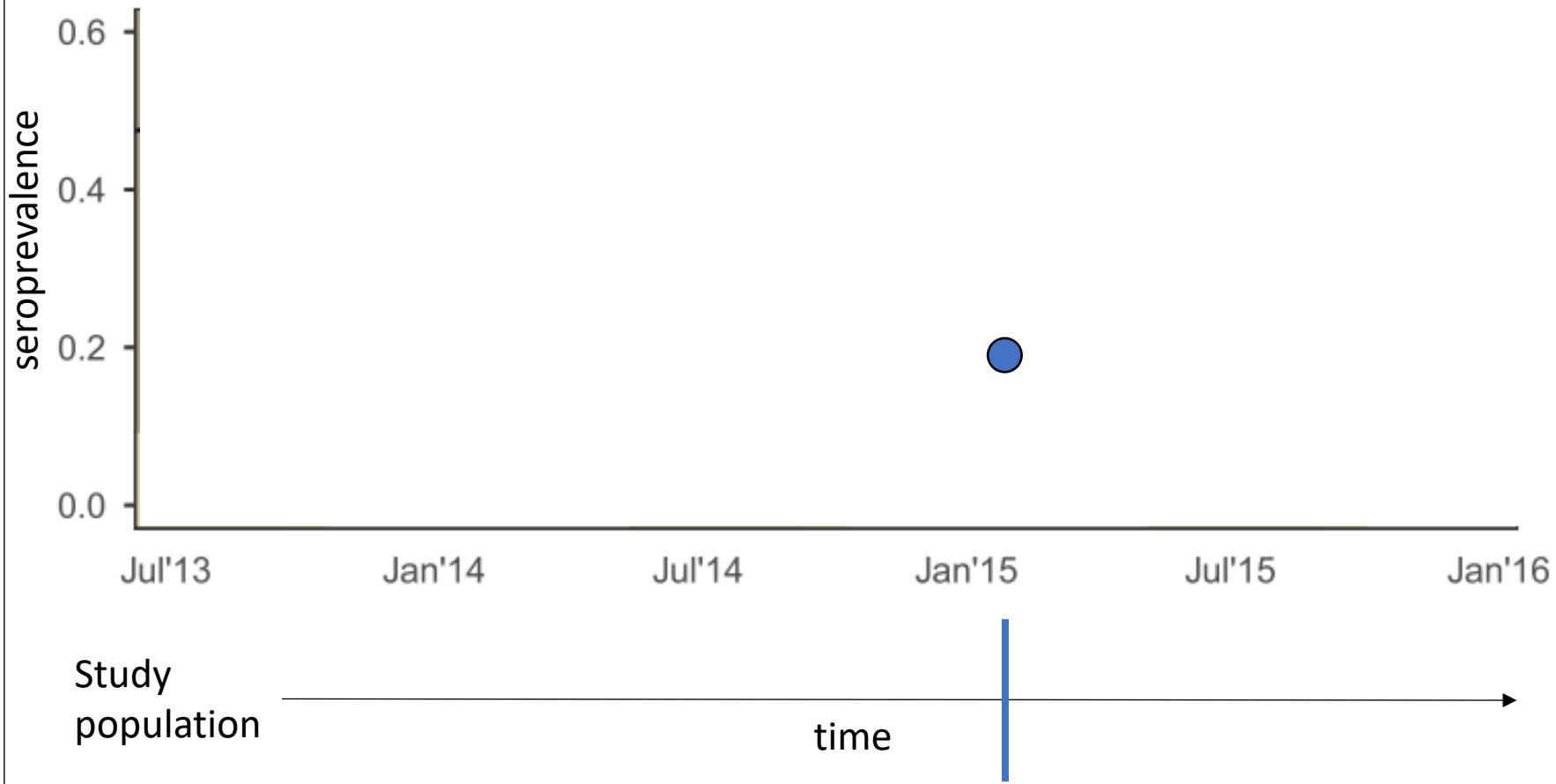
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Cross-sectional studies = snapshot of the population at a particular point in time



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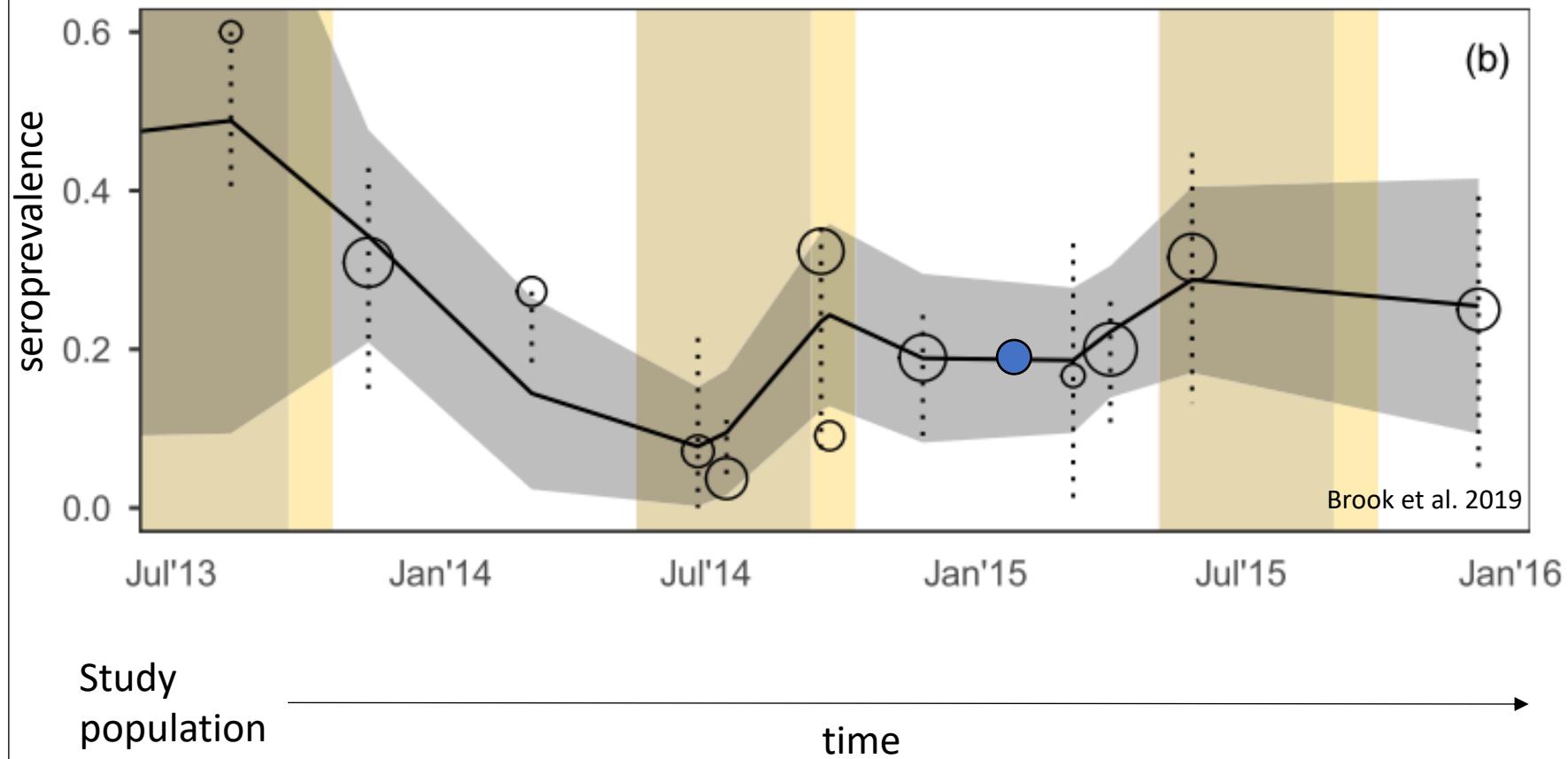
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- **Longitudinal**
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Longitudinal studies = follow a population over a **period of time**



Experimental

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➤ **Cohort** studies = follow a group of individuals over a **period of time**



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Unit of comparison = populations instead of individuals



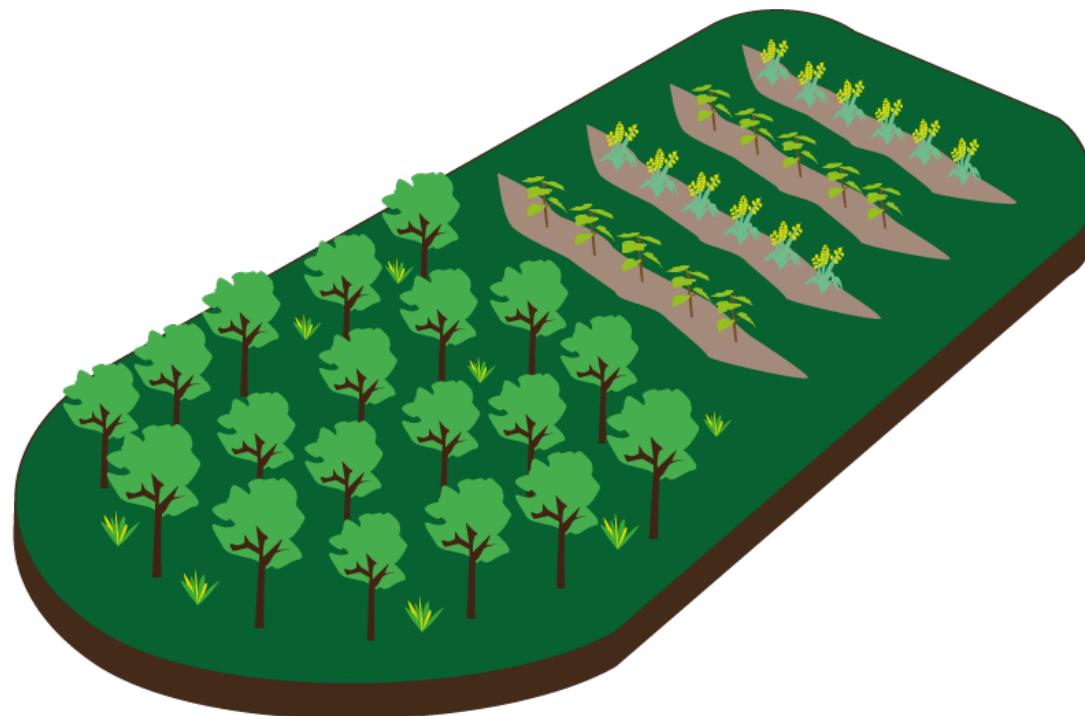
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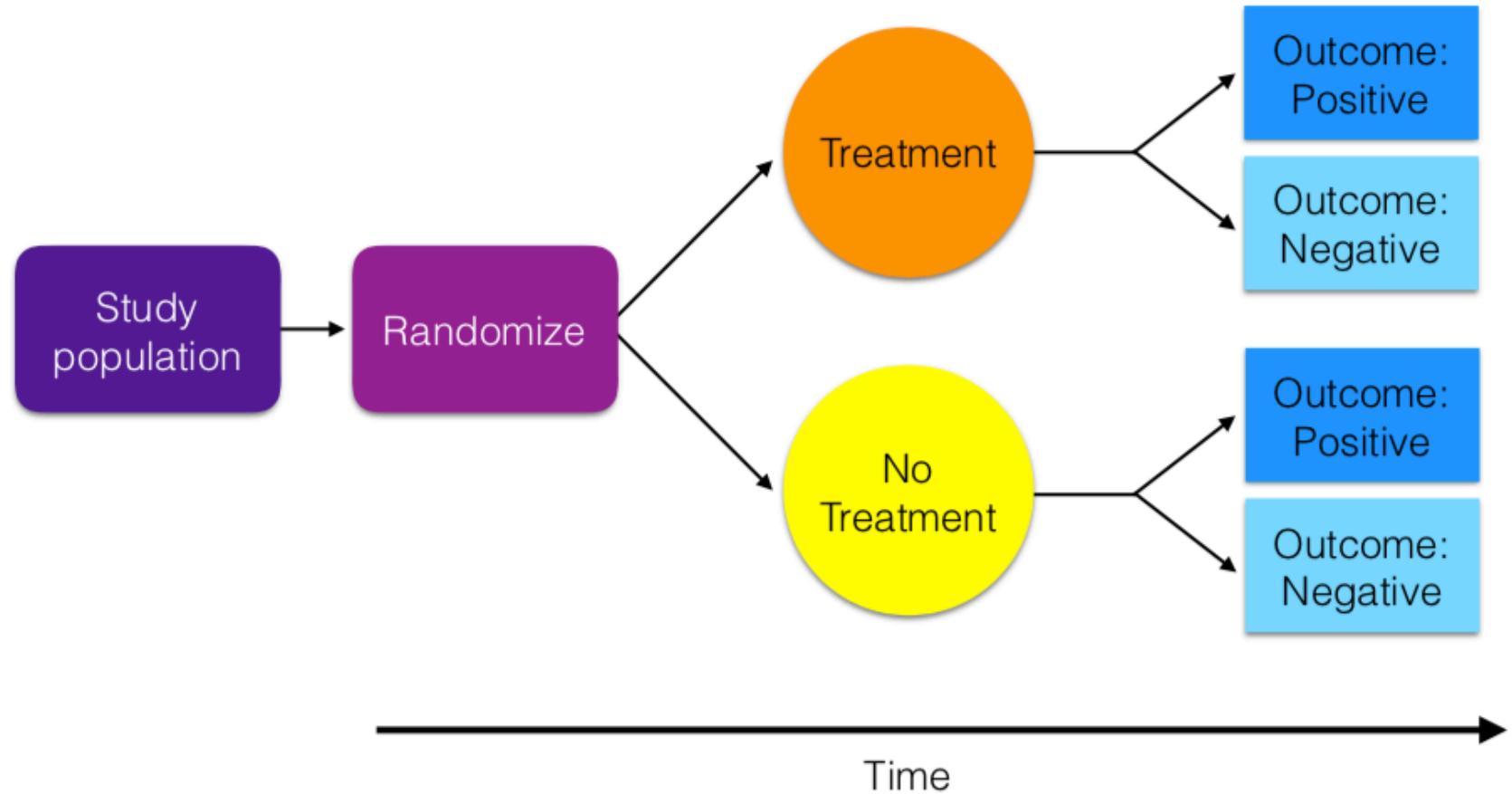
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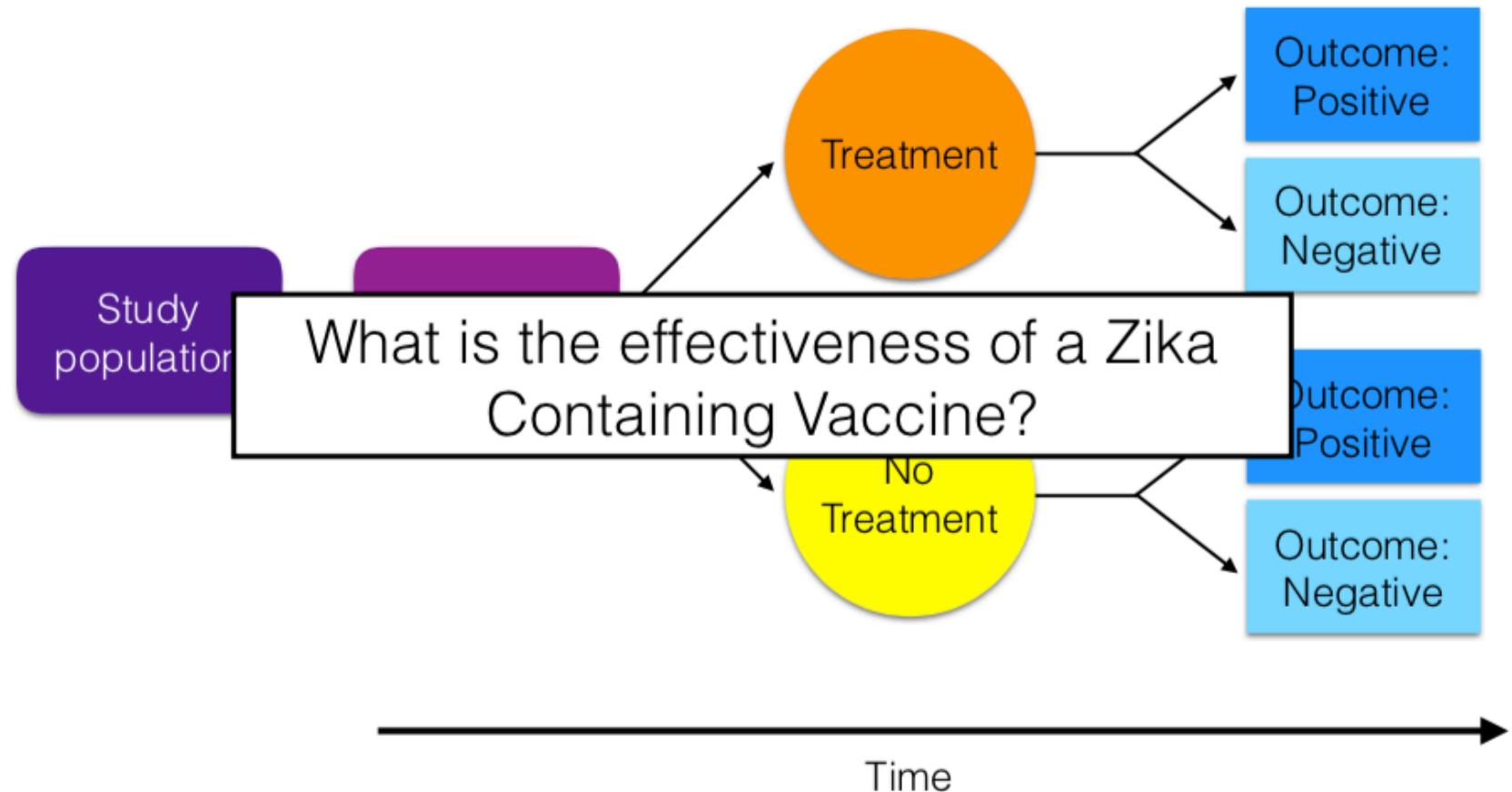
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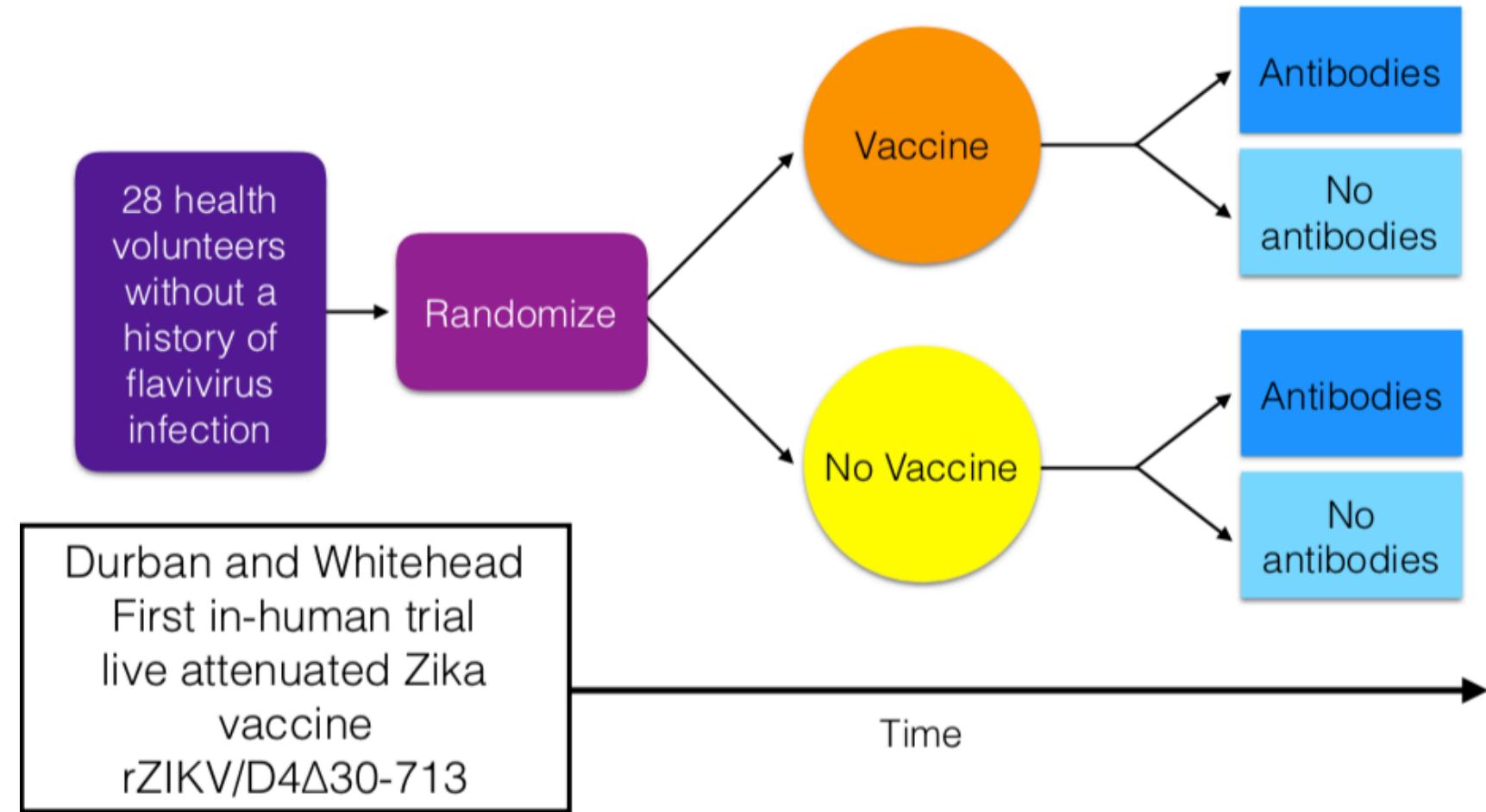
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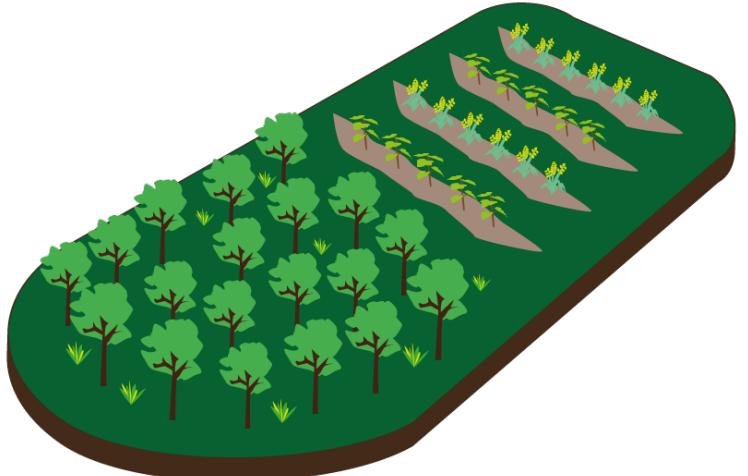
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6. Outline a data organization plan



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Designing your 1) field datasheet

| IDENTITY | | | | MEASUREMENTS | | | | | | MISC. SAMPLES | | SWABS | | BLOOD | | | | AGE | Temp Track Write as: Temp (time) |
|-----------|-------------------------------------|---|--------------------|----------------|----------------|---------------|------------|----------|----------------------|------------------|----------------------------------|---------------------------|-----------------------------|-----------------------------|--------------------------|--------------------------------|------------------|------------------|--|
| Sample ID | Tag # | M / F? | Mom/baby pair? Y/N | Weight (g) | Body (cm) | Fore-arm (mm) | Tibia (mm) | Ear (mm) | Tes/Mam (L) (W) | Hair Y/N | Ectos (# BF, MS, M, FL, T) | UR (#) | Amt plain eppendorf (serum) | Amt EDTA eppendorf (plasma) | Amt RNA-tube (ul) | # slides thick | Fltr? | Tooth? Y/N | |
| | recap? Y/N | class (M: J/A: F: J/ NL/P/L) *if young of the yr | ID # of mom/baby | | | | | | | FEC (#) | Wing Punch in EtOH (#) | THR (#) | Target: Eid Pier: >500ul | Target: Eid Pier: >200ul | Target: Eid Pier: >200ul | Assess (new, mild, mod, heavy) | | | |
| | | S: F 956000005601848 Rou160 N | C: NL | | 86 34 52 | 112.1 | 70.9 | 33.8 | 14.6 | L: 304 W: 24 | Hair: Y WP: 1 1 | BF: 2 THR: 1 M: 4 | UR: 1 THR: 1 FEC: 1 | 350 | 150 | 100 | TK: 1 TN: 2 | T: N A: Mild. | |
| | S: M 956000005670812 Rou161 N | C: A | | 85 34 51 | 111.8 | 71.0 | 34.7 | 12.9 | L: small W: small | Hair: Y WP: 1 | BF: 1 THR: 1 FL: 1 T: 1 | UR: 1 THR: 1 FEC: 1 | 350 | 150 | 100 | TK: 1 TN: 2 | T: M A: Mild. | | |
| | S: F 956000006449656 Rou163 N | C: NL | | 81 34 47 | 110.7 | 71.3 | 31.9 | 12.9 | L: small W: small | Hair: Y WP: 1 | BF: 3 THR: 1 M: 6 T: 1 | UR: 1 THR: 1 FEC: 1 | 400 | 150 | 80 | TK: 1 TN: 2 | T: N A: Mild. | | |
| | S: F 956000005605949 Rou164 N | C: NL | | 89 34 56 | 113.1 | 73.4 | 34.6 | 12.8 | L: 1 W: 32 | Hair: Y WP: 1 | BF: 7 M: 6 FL: 2 | UR: 1 THR: 1 FEC: 1 | 350 | 150 | 100 | TK: 1 TN: 2 | T: N A: Mild. | | |
| | S: M 956000005605949 Rou164 N | C: NL | | 82 32 56 | 110.8 | 72.1 | 32.1 | 12.1 | L: small W: small | Hair: Y WP: 1 | BF: 6 | UR: 1 | | | | TK: 1 | T: N | | |

6. Outline a data organization plan

Designing your 1) field datasheet
2) database structure

Compiling data the “long way”
➤ identifying information stored
in columns

| SITE DATA | | | | | Identification | |
|-------------|--------------|----------|-----------|---------------|----------------|--|
| Roost Site | Researchers | Date | Net Night | Bat Species | Sample ID | |
| Lakato | CB, CR, LA | 8/22/13 | 2 | Eidolon dupr | LKT-001 | |
| Lakato | CB, CR, LA | 8/22/13 | 2 | Eidolon dupr | LKT-002 | |
| Lakato | CB, CR, LA | 8/22/13 | 2 | Eidolon dupr | LKT-003 | |
| Lakato | CB, CR, LA | 8/22/13 | 2 | Eidolon dupr | LKT-004 | |
| Lakato | CB, CR, LA | 8/22/13 | 2 | Eidolon dupr | LKT-005 | |
| Marovitsika | CB, CR, AR | 11/4/13 | 1 | Pteropus rufi | MAR01 | |
| Marovitsika | CB, CR, AR | 11/4/13 | 1 | Pteropus rufi | MAR02 | |
| Marovitsika | CB, CR, AR | 11/4/13 | 1 | Pteropus rufi | MAR03 | |
| Marovitsika | CB, CR, AR | 11/4/13 | 1 | Pteropus rufi | MAR04 | |
| Marovitsika | CB, CR, AR | 11/4/13 | 1 | Pteropus rufi | MAR05 | |
| Marovitsika | CB, CR, AR | 11/6/13 | 3 | Pteropus rufi | MAR06 | |
| Marovitsika | CB, CR, AR | 11/7/13 | 4 | Pteropus rufi | MAR07 | |
| Marovitsika | CB, CR, AR | 11/7/13 | 4 | Pteropus rufi | MAR08 | |
| Marovitsika | CB, CR, AR | 11/7/13 | 4 | Pteropus rufi | MAR09 | |
| Marovitsika | CB, CR, AR | 11/8/13 | 5 | Pteropus rufi | MAR010 | |
| Marovitsika | CB, CR, AR | 11/8/13 | 5 | Pteropus rufi | MAR011 | |
| Marovitsika | CB, CR, AR | 11/8/13 | 5 | Pteropus rufi | MAR012 | |
| Marovitsika | CB, CR, AR | 11/8/13 | 5 | Pteropus rufi | MAR013 | |
| Ambakoana | CB,CR, AR, R | 11/15/13 | 1 | Pteropus rufi | AMB1 | |
| Ambakoana | CB,CR, AR, R | 11/15/13 | 1 | Pteropus rufi | AMB2 | |
| Ambakoana | CB,CR, AR, R | 11/15/13 | 1 | Pteropus rufi | AMB3 | |
| Ambakoana | CB,CR, AR, R | 11/15/13 | 1 | Pteropus rufi | AMB4 | |
| Ambakoana | CB,CR, AR, R | 11/15/13 | 1 | Pteropus rufi | AMB5 | |
| Ambakoana | CB,CR, AR, R | 11/15/13 | 1 | Pteropus rufi | AMB6 | |

6. Outline a data organization plan

Designing your 1) field datasheet
2) database structure
3) sample storage system

Freezer mgp

Necropsies
(columns by species)
in back: Caras old phd samples

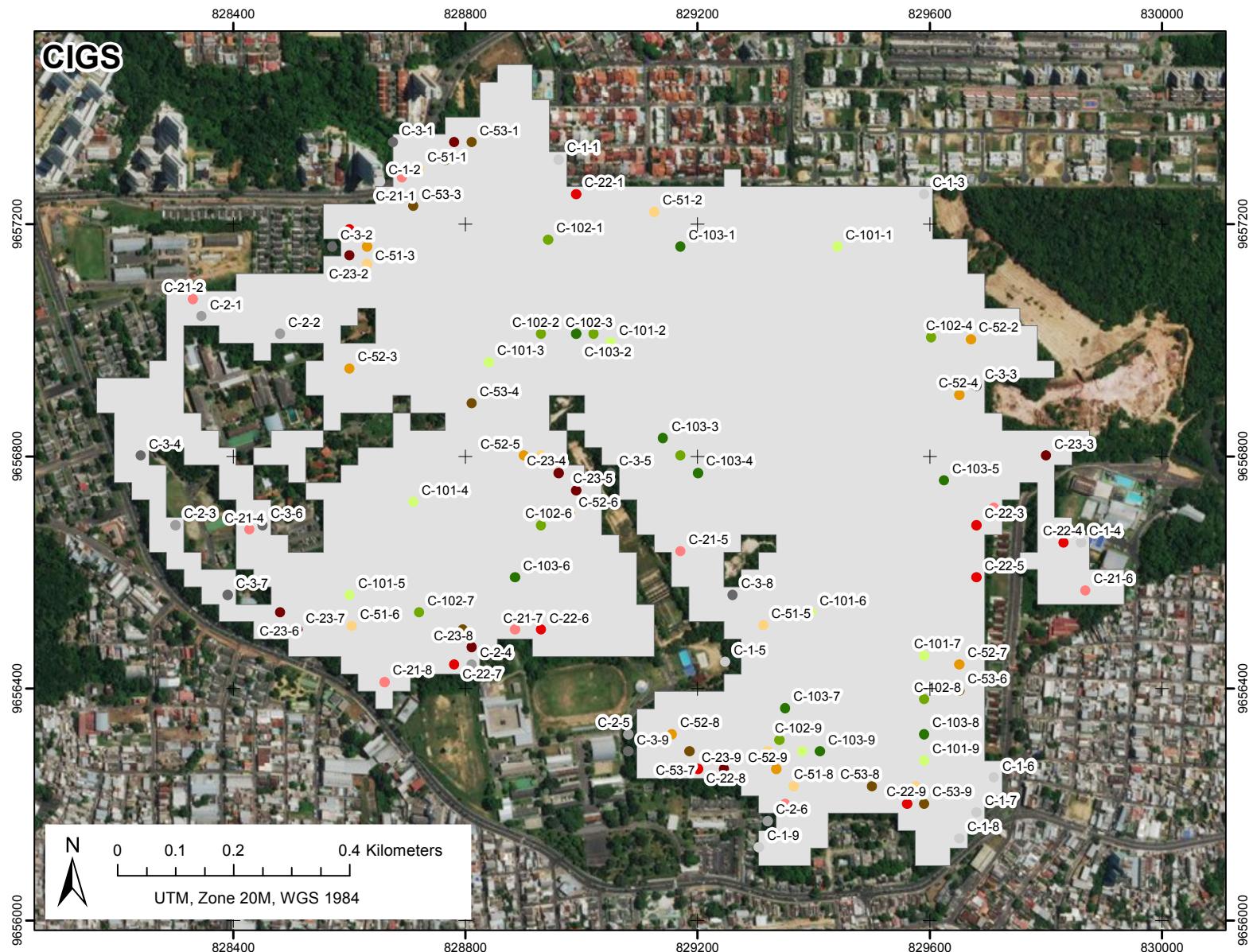
| FEC | UR | THR | SERA | SERB |
|------------------|------|--------------------|--|------|
| EDTA PEL/plas | PELB | RNA whole blood | | |
| PELA | WP | | MB (microbiome) -not collecting anymore | |

| Box Number | Sample Type | Species | Reagent | SampleID | Date Stocked | Notes |
|------------|--------------|-------------|-------------|----------|--------------|-------|
| PR-FEC-1 | Feces | Asio madaga | UTM | AMAK_001 | 4/25/18 | |
| PR-RNA-BLD | Whole Blood | Asio madaga | RNA-protect | AMAK_001 | 4/25/18 | |
| PR-THR-1 | Throat | Asio madaga | UTM | AMAK_001 | 4/25/18 | |
| PR-SER-A-1 | Serum A | Asio madaga | raw | AMAK_001 | 4/25/18 | |
| PR-SER-B-1 | Serum B | Asio madaga | raw | AMAK_001 | 4/25/18 | |
| PR-PEL-A-1 | Blood Pellet | Asio madaga | raw | AMAK_001 | 4/25/18 | |
| PR-PEL-B-1 | Blood Pellet | Asio madaga | raw | AMAK_001 | 4/25/18 | |

7. Be flexible



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Study Design: Examples from E2M2?

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