

# Protocol for compiling raw data from multiple sources and deduplicating records

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## Goal

Compile raw data from multiple sources and deduplicate records to create an initial dataset that will be enhanced throughout the course of this project.

## Relevant fields in the dataset

### Enhanced data recorded in

- gbif
- idigbio

## Process & Parties Responsible

This process is completed by the Digitization Specialist using the iDigBio and GBIF websites and R.

1. Acquire specimen occurrence data from iDigBio by searching in the Portal (<https://www.idigbio.org/portal/search>) for *Basis of record* = “preservedspecimen” and *Order* = “chiroptera” and downloading the results.
2. Acquire specimen occurrence data from GBIF by searching in the user interface (<https://www.gbif.org/occurrence/search>) for *Basis of record* = “Preserved specimen” and *Family* = “Hipposideridae” or *Family* = “Rhinolophidae” or *Family* = “Rhinonycteridae” and downloading the results.
3. Prepare iDigBio data in R:
  - a. Import both the raw occurrence data and the occurrence data processed by iDigBio during ingestion.
  - b. Add a suffix to field names to indicate data source (“\_idbR” for raw and “\_idbP” for processed).
  - c. Subset data to include only records belonging to one of the families of interest (Hipposideridae, Rhinolophidae, or Rhinonycteridae).
  - d. Split the field *idigbio\_geoPoint\_idbP* into two new fields, *decimalLatitude\_idbP* and *decimalLongitude\_idbP* to match the format of the iDigBio raw, GBIF raw, and GBIF processed data.

This protocol was created as part of [NSF DBI 2033973](https://www.nsf.gov/awardsearch/showAward?AWD_ID=2033973), RAPID Grant: Rapid Creation of a Data Product for the World's Specimens of Horseshoe Bats and Relatives, a Known Reservoir for Coronaviruses. Documents associated with this grant are archived at <https://doi.org/10.5281/zenodo.3974999>.

4. Prepare GBIF data in R:
  - a. Import both the raw occurrence data and the occurrence data processed by GBIF during ingestion.
  - b. Add a suffix to field names to indicate data source (“\_gbifR” for raw and “\_gbifP” for processed).
  - c. Exclude records published as part of a “checklist” dataset. These are typically based on literature referring to specimens and are not specimen occurrences themselves.
5. Compile data from iDigBio and from GBIF into a single dataset, deduplicating specimen records based on matches where the combination of *institutionCode*, *collectionCode*, *catalogNumber*, and *occurrenceID* is the same in both the GBIF raw and iDigBio raw data.
6. Record the provenance of each record in the fields *idigbio* (“1” = record was present in iDigBio data; “0” = record was absent in iDigBio data) and *gbif* (“1” = record was present in GBIF data; “0” = record was absent in GBIF data).
7. Save the dataset as a CSV file.

## Communication

Communication for this task will be via weekly team meetings.

## Results

This task was completed by Digitization Specialist Erica Krimmel on 2020-09-23 and took a total of approximately 31 hours, including the initial data exploration required to write effective code. The iDigBio search (Step #1) retrieved 829,063 records; the decision to search based on order versus the specific families of interest to this project was due to concern that iDigBio taxonomic data may be slightly out of date. GBIF searches (Step #2) resulted in 40,101 Hipposideridae and 40,404 Rhinolophidae records. Although Rhinonycteridae is also a family of interest, there are no records in GBIF in this family. The raw data from Steps #1 and #2 were archived on Zenodo in the form of three Darwin Core Archives, one from iDigBio and two from GBIF. Step #3c reduced the number of iDigBio records to 59,122. Step #4c reduced the number of GBIF records from 80,505 to 79,855 by excluding 650 records from 25 GBIF “checklist” datasets. Step #5 resulted in a dataset of 89,837 records, of which 9,982 were only present in iDigBio, 30,715 were only present in GBIF, and 49,140 were present in both iDigBio and GBIF (Step #6). There are almost certainly instances where our deduplication criteria of *institutionCode* + *collectionCode* + *catalogNumber* + *occurrenceID* were too strict and two records representing the same physical specimen were not merged. Given the potential complexity of solving this problem (e.g., truly unique specimens of the same taxon collected at the same place and time could be erroneously labeled as duplicates and removed from the dataset), we determined to be conservative in asserting that two records were exact duplicates.

Code associated with this protocol can be found in ‘RAPID-code\_compile-deduplicate.R’ (archived at <https://doi.org/10.5281/zenodo.3974999>).

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