

Bulk stable isotopes from siphonophores collected during four research cruises on the R/V Wester Flyer in the California Current Ecosystem between 2019 and 2021

Website: <https://www.bco-dmo.org/dataset/916958>

Data Type: Cruise Results

Version: 1

Version Date: 2024-01-11

Project

» [Collaborative research: The effects of predator traits on the structure of oceanic food webs](#) (SiphWeb)

Contributors	Affiliation	Role
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Abstract

Samples of siphonophores (Cnidaria, Hydrozoa) were collected using blue-water diving, midwater trawls, and remotely operated vehicles in the California Current Ecosystem, from 0 to 3,000 meters depth. Siphonophore samples were collected on four research cruises on the R/V Wester Flyer between 2019-2021. To remove potential biases associated with tissue-specific variability in stable isotope values, the gelatinous swimming bells (nectophores) of siphonophores were sampled. This approach was possible for most specimens, except for physonect species that are extremely fragile or have nectosomes that are a small fraction of the colony length and are often not collected. For these species (e.g., *Apolemia* spp.), the gelatinous bracts and pieces of the siphosome, excluding gastrozooids, were used. For small individuals (*Diphyes dispar*, *Nanomia bijuga*, and *Sphaeronectes koellikeri*), nectophores from several colonies that were captured at the same time and sampling location were pooled to obtain an adequate mass for isotope analyses. A subset of samples was selected for compound-specific isotope analysis of amino acids. These specific taxa were selected as representatives of different depth habitats, suborders, and hypothesized diets. Bulk and compound-specific isotope analyses were performed at the University of Hawaii's Biogeochemistry Stable Isotope Facility. This dataset includes the bulk stable isotope measurements along with metadata for specimens when possible (collection month and year, latitude, longitude, depth).

Table of Contents

- [Coverage](#)
- [Dataset Description](#)
 - [Methods & Sampling](#)
 - [BCO-DMO Processing Description](#)
- [Data Files](#)
- [Related Publications](#)
- [Related Datasets](#)
- [Parameters](#)
- [Instruments](#)
- [Deployments](#)
- [Project Information](#)
- [Funding](#)

Coverage

Spatial Extent: N:37.1967 E:-117.717 S:32.72 W:-125.038

Temporal Extent: 2019-03 - 2021-07

Methods & Sampling

Samples were collected in the central and southern California Current. Most samples were collected in the Monterey Bay region, but a subset of samples were collected in southern California in 2020 and 2021. Samples were collected from between 0 to 3,000 meters depth. Samples were collected on four cruises across three years. All cruises were on the R/V Western Flyer. Dr. Steven Haddock (haddock@mbari.org) was the Chief Scientist on all cruises. Cruises occurred in March 2019 (Cruise ID: WF0319), January 2020 (Cruise ID: WF0120), July 2020 (Cruise ID: WF0720), and July 2021 (Cruise ID: WF0721). Sample locations and dates are provided as columns in the data file.

Siphonophores were collected using three methods: (1) a remotely operated vehicle, (2) blue water diving, and (3) a midwater trawl.

(1) We used the Remotely Operated Vehicle (ROV) Doc Ricketts (<https://www.mbari.org/technology/rov-doc-ricketts/>) to collect siphonophores, which is an electro-hydraulic vehicle that operates between 200 and 4000 meters. The vehicle was fitted with high-definition video cameras, environmental data instrumentation (e.g. depth, temperature, salinity, and oxygen sensors), and suction and detritus samples to collect in-tact siphonophore specimens. ROV collections occurred during daylight hours.

(2) Siphonophores were collected by blue water diving between 0 and 20 meters during daylight hours. Blue water diving techniques followed the guidelines in the following publication: Haddock, Steven HD, and John N. Heine. "Scientific blue-water diving." (2005). From Haddock and Heine (2005): "In a typical blue-water dive, working divers are connected to a surface platform (and indirectly to each other) by tethers attached to a central hub, which is tended by a safety diver. This hub is connected to a down-line, providing a vertical point of reference. A surface float allows the divers to drift freely through the upper water-column, focusing on their work while the safety diver acts as a buddy for everyone."

(3) A Tucker Trawl with a frame area: 2 square meters (m²), mesh size: 500 micrometers (µm) was towed obliquely for ~2 hours between 900 meters and the surface at night.

Upon collection, siphonophores were identified to the finest taxonomic level, which was either genus or species. For some genera, there are likely undescribed and/or cryptic species (e.g., *Apolemia*) and for these taxa, genera-level identifications were used. All siphonophores were rinsed with DI water and frozen at -80°C until further processing. Siphonophore tissues were weighed, lyophilized, packaged into tin capsules for bulk isotope analysis, and analyzed at the University of Hawaii's Isotope Geochemistry Facility.

For bulk stable isotope analysis, siphonophore samples were analyzed using a Costech (Valencia, CA, USA) elemental combustion system coupled to a Thermo-Finnigan Delta XP isotope ratio mass spectrometer with N2 standard for nitrogen and Vienna Pee Dee Belemnite for carbon.

A subset of samples was selected for compound-specific isotope analysis of amino acids (CSIA-AA). CSIA-AA was also conducted at the University of Hawaii's Isotope Geochemistry Facility using acid hydrolysis followed by derivatization (see Popp et al. (2007) and Hannides et al. (2013) for details). The CSIA-AA data are available in a separate BCO-DMO dataset (see 'Related Datasets').

Related Resources:

Some of the siphonophores collected in this dataset were also used for metabarcoding. Those data are published Damian-Serrano, et al. (2022) (doi: [10.1371/journal.pone.0267761](https://doi.org/10.1371/journal.pone.0267761))

Illumina sequencing data files can be found in the NCBI BioProject PRJNA733192 (<https://www.ncbi.nlm.nih.gov/bioproject/733192>)

Prey 18S reference database enhancement sequences are available in NCBI (accession numbers between [MZ333540](#) - [MZ333629](#))

Other data, intermediary files, and all code can be found in the GitHub repository: https://github.com/dunnlab/siphweb_metabarcoding (see DOI: doi.org/10.1371/journal.pone.0267761)

BCO-DMO Processing Description

- Imported original file "Siphonophore bulk stable isotope data (NSF Project Number 1829812).csv" into the BCO-DMO system.

- Marked 'NA' as a missing data value (missing data are blank/empty in the final CSV file).
- Renamed fields to comply with BCO-DMO naming conventions.
- Saved the final file as "916958_v1_siphonophore_bulk_stable_isotopes.csv".

[[table of contents](#) | [back to top](#)]

Data Files

File
916958_v1_siphonophore_bulk_stable_isotopes.csv (Comma Separated Values (.csv), 14.15 KB) MD5:b7e7d4e4f065cc22490ea53f6c437d0f
Primary data file for dataset ID 916958, version 1

[[table of contents](#) | [back to top](#)]

Related Publications

Damian-Serrano, A., Hetherington, E. D., Choy, C. A., Haddock, S. H. D., Lapidés, A., & Dunn, C. W. (2022). Characterizing the secret diets of siphonophores (Cnidaria: Hydrozoa) using DNA metabarcoding. PLOS ONE, 17(5), e0267761. <https://doi.org/10.1371/journal.pone.0267761>

Related Research

Haddock, S. H. D., Heine, J. N., United States. National Oceanic and Atmospheric Administration, California Sea Grant College Program, & National Sea Grant College Program (U.S.). (2005). Scientific blue-water diving. California Sea Grant College Program. <https://isbnsearch.org/isbn/978-1-888691-13-9>

Methods

Hannides, C. C. S., Popp, B. N., Choy, C. A., & Drazen, J. C. (2013). Midwater zooplankton and suspended particle dynamics in the North Pacific Subtropical Gyre: A stable isotope perspective. Limnology and Oceanography, 58(6), 1931–1946. doi:[10.4319/lo.2013.58.6.1931](https://doi.org/10.4319/lo.2013.58.6.1931)

Methods

Hetherington, ED, Close, H, Haddock, SHD, Damian-Serrano, A, Dunn, CW, Wallsgrove, N, Doherty, S, and Choy, CA. Vertical trophic structure and niche partitioning of gelatinous predators in a pelagic food web: insights from stable isotopes of siphonophores. Under Review at Limnology and Oceanography.

Results

Popp, B. N., Graham, B. S., Olson, R. J., Hannides, C. C. S., Lott, M. J., López-Ibarra, G. A., ... Fry, B. (2007). Insight into the Trophic Ecology of Yellowfin Tuna, *Thunnus albacares*, from Compound-Specific Nitrogen Isotope Analysis of Proteinaceous Amino Acids. Terrestrial Ecology, 173–190. doi:[10.1016/s1936-7961\(07\)01012-3](https://doi.org/10.1016/s1936-7961(07)01012-3)

Methods

[[table of contents](#) | [back to top](#)]

Related Datasets

IsRelatedTo

Hetherington, E. D., Choy, C. A. (2024) **Compound-specific isotope analysis of amino acids (CSIA-AA) from a subset of siphonophore samples collected during four research cruises on the R/V Wester Flyer in the California Current Ecosystem between 2019 and 2021.** Biological and Chemical Oceanography Data Management Office (BCO-DMO). (Version 1) Version Date 2023-12-19 doi:10.26008/1912/bco-dmo.917239.1 [[view at BCO-DMO](#)]

Yale University. DNA Metabarcoding of Siphonophore Gut Contents. 2021/05. In: BioProject [Internet]. Bethesda, MD: National Library of Medicine (US), National Center for Biotechnology Information; 2011-. Available from: <http://www.ncbi.nlm.nih.gov/bioproject/PRJNA733192>. NCBI:BioProject: PRJNA733192.

Parameters

Parameter	Description	Units
Ship	Ship Name	unitless
Chief_Scientist	Chief Scientist Name	unitless
Cruise_Name	Cruise name, where the format is the ship's initials (e.g., WF for Western Flyer) followed by MMY (month and year)	unitless
Year	Year of sample collection	unitless
Month	Month of sample collection	unitless
Family	Taxonomic family of specimen	unitless
Genus	Taxonomic genus of specimen	unitless
Best_Taxonomic_ID	Best taxonomic identification of specimen, usually at the genus or species level	unitless
Depth_m	collection depth; trawl sample depths were the midpoint depths of the trawl, all blue water diving samples were assigned a depth of 10m, which was the average of dive depths; specimens collected with the ROV have discrete collection depths	meters (m)
Latitude	latitude of sample collection (positive values = North)	decimal degrees
Longitude	longitude of sample collection (negative values = West)	decimal degrees
EAweight_mg	mass of the sample analyzed on the Elemental Analyzer	milligrams (mg)
ug_C	mass of carbon detected in the sample by the elemental analyzer	micrograms (ug)
d13C	carbon isotope values relative to the international standard Vienna Pee Dee Belemnite	parts per thousand
ug_N	mass of nitrogen detected in the sample by the elemental analyzer	micrograms (ug)
d15N	nitrogen isotope values relative to the international standard N2	parts per thousand

Instruments

Dataset-specific Instrument Name	Costech (Valencia, CA, USA) elemental combustion system
Generic Instrument Name	Elemental Analyzer
Dataset-specific Description	For bulk stable isotope analysis, siphonophore samples were analyzed using a Costech (Valencia, CA, USA) elemental combustion system coupled to a Thermo-Finnigan Delta XP isotope ratio mass spectrometer with N2 standard for nitrogen and Vienna Pee Dee Belemnite for carbon.
Generic Instrument Description	Instruments that quantify carbon, nitrogen and sometimes other elements by combusting the sample at very high temperature and assaying the resulting gaseous oxides. Usually used for samples including organic material.

Dataset-specific Instrument Name	Thermo-Finnigan Delta XP isotope ratio mass spectrometer
Generic Instrument Name	Isotope-ratio Mass Spectrometer
Dataset-specific Description	For bulk stable isotope analysis, siphonophore samples were analyzed using a Costech (Valencia, CA, USA) elemental combustion system coupled to a Thermo-Finnigan Delta XP isotope ratio mass spectrometer with N2 standard for nitrogen and Vienna Pee Dee Belemnite for carbon.
Generic Instrument Description	The Isotope-ratio Mass Spectrometer is a particular type of mass spectrometer used to measure the relative abundance of isotopes in a given sample (e.g. VG Prism II Isotope Ratio Mass-Spectrometer).

Dataset-specific Instrument Name	blue water diving
Generic Instrument Name	Manual Biota Sampler
Dataset-specific Description	Siphonophores were collected by blue water diving between 0-20 meters during daylight hours. Blue water diving techniques following the guidelines in the following publication: Haddock, Steven HD, and John N. Heine. "Scientific blue-water diving." (2005).
Generic Instrument Description	"Manual Biota Sampler" indicates that a sample was collected in situ by a person, possibly using a hand-held collection device such as a jar, a net, or their hands. This term could also refer to a simple tool like a hammer, saw, or other hand-held tool.

Dataset-specific Instrument Name	Remotely Operated Vehicle Doc Ricketts
Generic Instrument Name	ROV Doc Ricketts
Dataset-specific Description	We used the Remotely Operated Vehicle Doc Ricketts (https://www.mbari.org/technology/rov-doc-ricketts/) to collect siphonophores, which is an electro-hydraulic vehicle that operates between 200 and 4000 meters. The vehicle was fitted with high-definition video cameras, environmental data instrumentation (e.g. depth, temperature, salinity, and oxygen sensors), and suction and detritus samples to collect in-tact siphonophore specimens. ROV collections occurred during daylight hours.
Generic Instrument Description	The remotely operated vehicle (ROV) Doc Ricketts is operated by the Monterey Bay Aquarium Research Institute (MBARI). ROV Doc Ricketts is capable of diving to 4000 meters (about 2.5 miles). The R/V Western Flyer is the support vessel for Doc Ricketts and was designed with a center well whose floor can be opened to allow Doc Ricketts to be launched from within the ship into the water below. For a complete description, see: https://www.mbari.org/at-sea/vehicles/remotely-operated-vehicles/rov-doc...

Dataset-specific Instrument Name	Tucker Trawl
Generic Instrument Name	Tucker Trawl
Dataset-specific Description	A Tucker Trawl with a frame area of 2 square meters and a mesh size of 500 micrometers was towed obliquely for ~2h hours between 900 meters and the surface at night.
Generic Instrument Description	The original Tucker Trawl, a net with a rectangular mouth opening first built in 1951 by G.H. Tucker, was not an opening/closing system, but shortly thereafter it was modified so that it could be opened and closed. The original had a 183 cm by 183 cm flexible rectangular mouth opening 914 cm long net with 1.8 cm stretched mesh for the first 457 cm and 1.3 cm mesh for last 457 cm. 152 cm of coarse plankton or muslin netting lined the end of the net. Tucker designed the net to collect animals associated with the deep scattering layers, principally euphausiids, siphonophores, and midwater fish. (from Wiebe and Benfield, 2003). Currently used Tucker Trawls usually have 1-m2 openings and can have a single net or multiple nets on the frame.

[[table of contents](#) | [back to top](#)]

Deployments

WF0319

Website	https://www.bco-dmo.org/deployment/917024
Platform	R/V Western Flyer
Description	Cruise occurred in March 2019

WF0120

Website	https://www.bco-dmo.org/deployment/917025
Platform	R/V Western Flyer
Description	Cruise occurred in January 2020

WF0720

Website	https://www.bco-dmo.org/deployment/917030
Platform	R/V Western Flyer
Description	Cruise occurred in July 2020

WF0721

Website	https://www.bco-dmo.org/deployment/917032
Platform	R/V Western Flyer
Description	Cruise occurred in July 2021

[[table of contents](#) | [back to top](#)]

Project Information

Collaborative research: The effects of predator traits on the structure of oceanic food webs (SiphWeb)

Coverage: North Pacific

Food webs describe who eats whom, tracing the flow of energy from plants up to large animals. While many connections in food webs on land are quite familiar (lions eat antelope and antelope eat grass, for example), there are large gaps in our understanding of ocean food webs. Closing these gaps is critical to understanding how nutrients and energy move through ocean ecosystems, how organisms interact in the ocean, and how best to manage ocean resources. This project will study ocean food web structure with a focus on siphonophores, an abundant group of predators in the open ocean that range in length from less than an inch to more than one hundred feet. Siphonophores are closely related to corals and many jellyfish. They are known to be important predators within ocean food webs, but they are difficult to study because they live across great ocean depths and are gelatinous and fragile. The details of what they eat, as well as many other features of their biology, remain poorly known. This project will combine direct observations of feeding, genetic analysis of siphonophore gut contents, and stable isotope analyses to identify what different species of siphonophores eat. The team will also examine why they eat what they do. This will provide a new understanding of how the structure of food webs arise, aiding in our ability to predict future changes to food webs as the global climate shifts. Siphonophores feed in a very unique manner--they have highly specialized tentacles that are used solely for capturing prey--thus, the prey captured is determined largely by the anatomy and function of these tentacles. The project will describe these tentacles, reconstruct their evolutionary history, and investigate how evolutionary shifts in tentacle structure have led to changes in diet. This project will train one PhD student, one Master's student, a postdoc, and undergraduate students, including individuals of underrepresented groups. This project will support the production of scientifically rigorous yet engaging videos, foster the expansion of a citizen-science program, and create K-12 teaching modules.

This project will advance three scientific aims: First, it will identify the diet of a diverse range of siphonophores using DNA metabarcoding of gut contents and prey field, remotely operated vehicle (ROV) video of prey encounters, and stable isotope analysis. These approaches are highly complementary and allow for extensive cross validation. Second, the project will characterize the selectivity of siphonophore diets by comparing them to the relative prey abundances in the habitats of each of these species. Third, the project will characterize the structure of the siphonophore prey capture apparatus across species through detailed morphological analysis of their tentacles and nematocysts. These data will be integrated in an ecological and evolutionary framework to identify predator features associated with prey specialization. In a larger context, addressing these questions will advance our understanding of oceanic predation by revealing how evolutionary changes in predator selectivity correspond to evolutionary changes in habitat and feeding apparatus and how these changes shape current food web structure in the open ocean. We will test and refine an integrated approach to describing the structure and origin of food web topology, and evaluate the potential for phylogenetic

relationships to explain prey selectivity.

This award reflects NSF's statutory mission and has been deemed worthy of support through evaluation using the Foundation's intellectual merit and broader impacts review criteria.

[[table of contents](#) | [back to top](#)]

Funding

Funding Source	Award
NSF Division of Ocean Sciences (NSF OCE)	OCE-1829812

[[table of contents](#) | [back to top](#)]