Hydrographic data from the CTD mounted on the trace metal rosette (TMR) aboard R/V Falkor cruise (160115) during the ProteOMZ expedition in the Central Pacific in 2016.

Website: https://www.bco-dmo.org/dataset/734608 Data Type: Cruise Results Version: 1 Version Date: 2018-05-01

Project

» <u>The ProteOMZ Expedition: Investigating Life Without Oxygen in the Pacific Ocean</u> (ProteOMZ (Proteomics in an Oxygen Minimum Zone))

| Contributors | Affiliation | Role |
|---------------------------|---|---------------------------------|
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Coverage

Spatial Extent: N:10.544984 **E**:-158.320979 **S**:-26.364655 **W**:-179.289931 **Temporal Extent**: 2016-01-16 - 2016-02-11

Dataset Description

Hydrographic data files from the SeaBird SBE19 CTD mounted on the trace metal rosette (TMR).

Methods & Sampling

Data were collected using the Trace Metal Rosette (TMR, Sea-Bird SEACAT 19+), equipped standard conductivity, temperature and pressure sensors, as well as an added optional SBE 43 dissolved oxygen sensor. All four sensors were factory refurbished/calibrated immediately prior to the expedition in November of 2015 by Sea-Bird Electronics (Bellevue WA).

Notes on CTD/O2 data acquisition and processing using Sea-Bird hardware and software. The GO-SHIP Repeat Hydrography Manual: A Collection of Expert Reports and Guidelines. IOCCP Perort No. 14, ICPO Publication Series No. 134, v. 1. 2010.

Location: Tropical/equatorial Pacific along 150º W; Honolulu, Hawai'i to Pape'ete, French Polynesia

Data Processing Description

Data from the SBE19Plus were processed using Sea-Bird's SBE Data Processing software, v. 7.23.2.

SBE processing modules were applied in the following order: *Data Conversion, Filter, Align CTD, Cell Thermal Mass,* and *Loop Edit* were applied to the input variables using the parameters identified in the .cnv file header shown below. Oxygen data were first processed using the raw sensor voltage, then converted to units of umol/kg using the *Derive* module. Finally, *Wild Edit* was used to remove extraneous values and data were binned by depth into 1 m bins using *Bin Average* and converted to ASCII format using *ASCII Out*.

Sea state during the cruise and issues with the block used to deploy the TMR did not allow full in-water equilibration of the CTD sensors and pumping system prior to each cast. As a result, we recommend using data from the upcasts (designated with the prefix 'u' in the filename).

```
# datcnv date = May 19 2016 15:57:33, 7.23.2 [datcnv vars = 5]
# datcnv in = C:\Users\Santoro\Desktop\Falkor 2016\TMR\160131TMR21CTDdata.hex
C:\Users\Santoro\Desktop\Falkor_2016\TMR\SBE19plusV2_6801.xmlcon
# datcnv skipover = 0
# datcnv ox hysteresis correction = yes
# filter date = May 19 2016 15:58:16, 7.23.2
# filter in = C:\Users\Santoro\Desktop\Falkor 2016\tmr process\160131TMR21CTDdata.cnv
# filter low pass to A = 0.500
# filter low pass to B = 0.150
# filter low pass A vars = depSM tv290C c0mS/cm sbeox0V
# filter low pass B vars = prdM
# alignctd date = May 19 2016 15:58:49, 7.23.2
# alignctd in = C:\Users\Santoro\Desktop\Falkor 2016\tmr process\160131TMR21CTDdata.cnv
# alignctd adv = c0mS/cm 0.073 # celltm date = May 19 2016 15:59:14, 7.23.2
# celltm in = C:\Users\Santoro\Desktop\Falkor 2016\tmr process\160131TMR21CTDdata.cnv
# celltm alpha = 0.0300, 0.0000
# celltm tau = 7.0000, 0.0000
# celltm_temp_sensor_use_for_cond = primary,
# loopedit date = May 19 2016 16:00:13, 7.23.2
# loopedit in = C:\Users\Santoro\Desktop\Falkor 2016\tmr process\160131TMR21CTDdata.cnv
# loopedit minVelocity = 0.250
# loopedit surfaceSoak: minDepth = 5.0, maxDepth = 20, useDeckPress = 1
# loopedit excl bad scans = yes
# Derive date = May 19 2016 16:03:09, 7.23.2 [derive vars = 5]
# Derive in = C:\Users\Santoro\Desktop\Falkor 2016\tmr process\160131TMR21CTDdata.cnv
C:\Users\Santoro\Desktop\Falkor 2016\TMR\SBE19plusV2 6801.xmlcon
# derive time window docdt = seconds: 2
# derive ox tau correction = yes
# wildedit date = May 19 2016 16:03:51, 7.23.2
# wildedit in = C:\Users\Santoro\Desktop\Falkor 2016\tmr process\160131TMR21CTDdata.cnv
# wildedit pass1 nstd = 2.0
# wildedit pass2 nstd = 20.0
# wildedit pass2 mindelta = 0.000e+000
# wildedit npoint = 100
# wildedit vars = prdM depSM tv290C c0mS/cm sbeox0V sal00 potemp090C density00 sigma-È00
sbeox0Mm/Kg
# wildedit excl bad scans = yes
# file type = ascii
```

BCO-DMO Data Processing Notes:

- Files were originally grouped in separate zip files, but were compressed into one to serve.

```
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```

Related Datasets

IsRelatedTo

Saito, M. A., Saunders, J. (2022) **Relative protein abundance from scaled and corrected exclusive peptide spectral counts from the ProteOMZ R/V Falkor expedition cruise FK160115 in the Pelagic central Pacific Ocean in 2016.** Biological and Chemical Oceanography Data Management Office (BCO-DMO). (Version 1) Version Date 2022-01-13 doi:10.26008/1912/bco-dmo.868030.1 [view at BCO-DMO] *Relationship Description: This dataset was collected asynchronously using another instrument at the same stations during the expedition.*

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Parameters

| Parameter | Description | Units |
|-------------|--------------------------------|--------------------------|
| PrdM | Pressure, Strain Gauge | db |
| DepSM | Depth of salt water | meters |
| Tv290C | Temperature [ITS-90] | Celsius |
| C0mS/cm | Conductivity | mS per cm |
| Sal00 | Salinity | Practical Salinity Units |
| Potemp090C | Potential Temperature [ITS-90] | Celsius |
| Density00 | Density | kilogram per meter cubed |
| Sigma-E00 | Density [sigma-theta] | kilogram per meter cubed |
| Sbeox0Mm/Kg | Oxygen, SBE 43 | umol per kilogram |
| Flag | Flag | unitless |

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Instruments

| Dataset-specific Instrument Name | Trace metal rosette |
|-------------------------------------|---|
| Generic Instrument Name | Trace Metal Bottle |
| Dataset-specific Description | Used to collect samples |
| Generic Instrument Description | Trace metal (TM) clean rosette bottle used for collecting trace metal clean seawater samples. |

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Deployments

FK160115

| Website | https://www.bco-dmo.org/deployment/708387 |
|-------------|---|
| Platform | R/V Falkor |
| Report | https://service.rvdata.us/data/cruise/FK160115/doc/FK160115_OfficialCruiseReport_Saito_v3.pdf |
| Start Date | 2016-01-16 |
| End Date | 2016-02-11 |
| Description | Project: Using Proteomics to Understand Oxygen Minimum Zones (ProteOMZ) More information is available from the ship operator at <u>https://schmidtocean.org/cruise/investigating-life-without-oxygen-in-the</u> Additional cruise information is available from the Rolling Deck to Repository (R2R): <u>https://www.rvdata.us/search/cruise/FK160115</u> |

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Project Information

The ProteOMZ Expedition: Investigating Life Without Oxygen in the Pacific Ocean (ProteOMZ (Proteomics in an Oxygen Minimum Zone))

Website: https://schmidtocean.org/cruise/investigating-life-without-oxygen-in-the-tropical-pacific/#team

Coverage: Central Pacific Ocean (Hawaii to Tahiti)

From Schmidt Ocean Institute's ProteOMZ Project page:

Rising temperatures, ocean acidification, and overfishing have now gained widespread notoriety as humancaused phenomena that are changing our seas. In recent years, scientists have increasingly recognized that there is yet another ingredient in that deleterious mix: a process called deoxygenation that results in less oxygen available in our seas.

Large-scale ocean circulation naturally results in low-oxygen areas of the ocean called oxygen deficient zones (ODZs). The cycling of carbon and nutrients – the foundation of marine life, called biogeochemistry – is fundamentally different in ODZs than in oxygen-rich areas. Because researchers think deoxygenation will greatly expand the total area of ODZs over the next 100 years, studying how these areas function now is important in predicting and understanding the oceans of the future. This first expedition of 2016 led by Dr. Mak Saito from the Woods Hole Oceanographic Institution (WHOI) along with scientists from University of Maryland Center for Environmental Science, University of California Santa Cruz, and University of Washington aimed to do just that, investigate ODZs.

During the 28 day voyage named "ProteOMZ," researchers aboard R/V *Falkor* traveled from Honolulu, Hawaii to Tahiti to describe the biogeochemical processes that occur within this particular swath of the ocean's ODZs. By doing so, they contributed to our greater understanding of ODZs, gathered a database of baseline measurements to which future measurements can be compared, and established a new methodology that could be used in future research on these expanding ODZs.

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Funding

| Funding Source | Award |
|---|--|
| Gordon and Betty Moore Foundation: Marine Microbiology Initiative (MMI) | <u>GBMF3782</u> |
| Alfred P. Sloan Foundation (Sloan) | Unknown ProteOMZ Sloan Foundation |
| Schmidt Ocean Institute (SOI) | R/V Falkor 160115 SOI ProteOMZ Expedition |

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