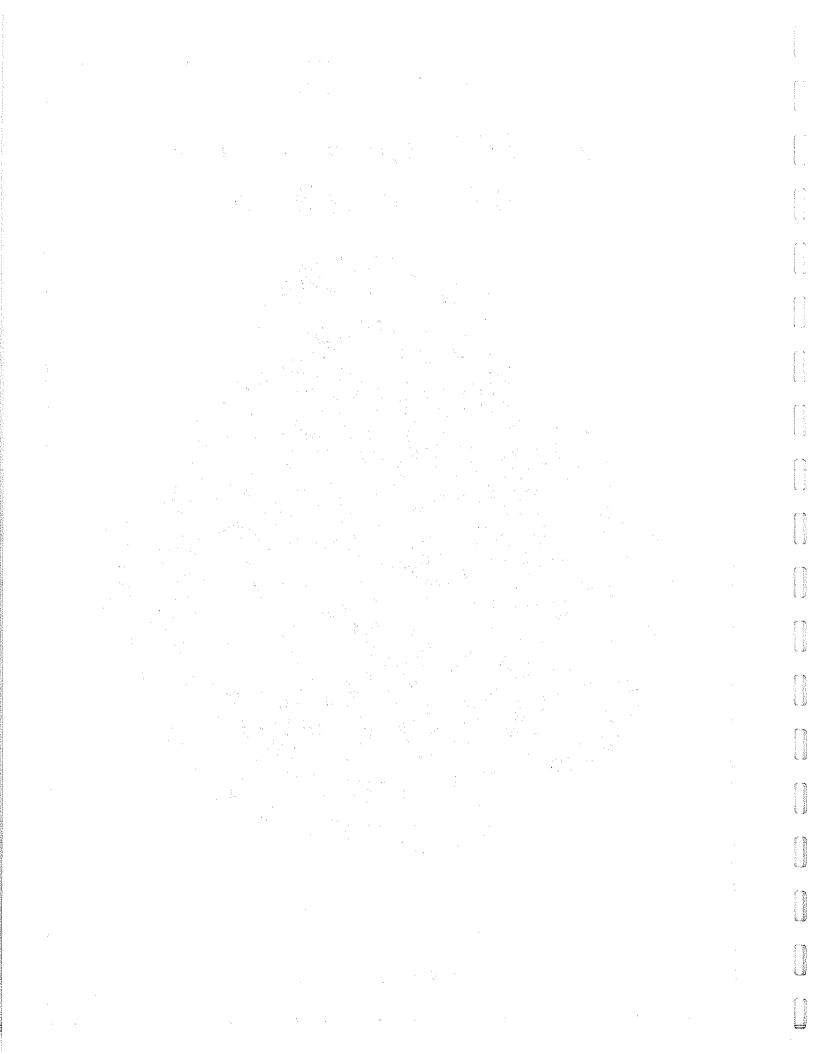
Cruise Report

RV/ISELIN Cruise C19407 to Georges Bank



25 May - 16 June 1994



Cruise Report

RV COLUMBUS ISELIN CRUISE 9407 to Georges Bank

Acknowledgements

This report was prepared by the Chief Scientist, with input from all Principal Investigators. The contributions of Maria Bemis, Bob Campbell, Peter Garrahan and Charlie Miller are greatly appreciated.

We are grateful for the excellent support provided by the Captain and crew of the RV *Columbus Iselin*. The professional atmosphere enabled our research efforts to run smoothly and allowed the scientific party to achieve all of its goals and objectives.

This cruise was sponsored by the National Science Foundation and the National Oceanographic and Atmospheric Administration. All data contained in this report are to be considered preliminary.

TABLE OF CONTENTS

Objectives of the cruise	3
Cruise Narrative	5
Individual PI Reports	14
Hydrography	
CTD data Biological and chemical data	
Zooplankton process studies	
Growth rate studies	
Egg laying rate studies Feeding rate studies	
Population process studies	
Larval fish studies	
Video Plankton recorder studies	32
Acoustic studies (TAPS)	33
Airborne LIDAR fluorescence measurements	34
Personnel list	43
Appendix 1: Event log	
Appendix 2: PI address and e-mail list	,

Objectives of the cruise

Field work for the U.S.-GLOBEC Georges Bank Program began in the spring of 1994. This cruise aboard the RV Columbus Iselin was one of six separate seagoing efforts scheduled for 1994 (Table 1). Two earlier cruises on RV Albatross IV were devoted to process work on the southern flank of Georges Bank and adjacent well mixed areas. A cruise aboard the RV Albatross IV was conducting bank-wide acoustic studies during the period of this cruise.

Our efforts were focused on measuring vital physiological rates of target species on the Bank, primarily the calanoid copepods, *Calanus finmarchicus* and *Pseudocalanus* spp., and larval cod, *Gadus morhua*. A major objective of field activities in 1994 was to shakedown methods and protocols for the intensive 1995 field season and to integrate the various program components. Specific objectives were:

- (1) To conduct a rapid broad-scale survey of U.S. GLOBEC target zooplankton species in order to determine their distribution and abundance.
- (2) To conduct a hydrographic survey of the Bank, including CTD data, in vivo fluorescence, and transmissometry with ancillary measurements of size-fractionated chlorophyll, nutrients, and microplankton.
- (3) To perform experiments measuring vital physiological rates (including growth, feeding, lipid deposition, and egg laying) of target zooplankton and larval fish species under stratifed and well-mixed water column conditions.
- (4) To examine target species distribution and abundance on sub-Bankwide scales using the video plankton recorder (VPR) and acoustic (TAPS) technologies.

The work was a combination of underway activities and station-keeping. Underway activities included CTD casts, stratified MOCNESS sampling for zooplankton, and VPR deployments. Hydrographic data included CTD, fluorometry and transmissometry, with bulk water samples collected for analysis of nutrients, size-fractionated chlorophyll (total, $<20~\mu m$ and $<5~\mu m$), and microplankton. Microplankton samples were collected for analysis by flow cytometry, automated epifluorescence microscopy, and inverted microscopy in order to describe the entire suite of nano- and microplankton including phytoplankton and protozoa. MOCNESS (150 μm mesh) samples were collected on Georges Bank from three depth strata at stations < 100m depth and four strata at stations > 100m depth. MOCNESS samples were collected from 9 strata in two deep basins in the southern Gulf of Maine. Other underway activities included 2 onbank-offbank hydrographic sections of 17 and 10 stations respectively, and a 27-hour VPR transect extending from slopewater off the southern flank of Georges Bank through the well-mixed area and into Georges Basin.

Station-keeping activities consisted of experimental work to measure vital rates in conjunction with twice daily (1200 and 2400) hydrocasts, MOCNESS and pump sampling, and multiple deployments of TAPS and VPR. Station-keeping was conducted in stratified and well-mixed areas of the Bank. Sites were selected on the basis of information obtained during the initial bank-wide survey. A drogued ARGOS drifter was deployed at each site, and tracked by the ship's bridge for the duration of station-keeping operations.

Ancillary activities not funded by the U.S.-GLOBEC program included three overflights by a group from M.I.T.'s Lincoln Laboratory to collect airborne LIDAR fluorescence measurements, phytoplankton net collections for Drs. P.E. Hargraves and J.E. Rines of the University of Rhode Island, and protozoan collections for Mr. R. Pierce of the University of Rhode Island.

Table 1. U.SGLOBEC cruises to Georges Bank during 1994.					
Dates	Vessel	Chief Scientist	Activities		
3-6 May	R.V. Columbus Iselin	J. Irish	Test moorings		
2-13 May	R.V. Albatross IV	L. Madin	Predator vital rates and distributions		
16-27 May	R.V. Albatross IV	G. Lough	Larval fish vital rates and distributions		
24 May-16 June	R.V. Endeavor	D. Gifford	Zooplankton vital rates and distributions		
31 May-10 June	R.V. <i>Albatross IV</i>	P. Wiebe	Acoustic survey		
28 June-1 July	R.V. Endeavor	J. Irish	Test moorings		
19-26 October	R.V. Endeavor	J. Irish	Mooring deployment		

The cruise was an unqualified success. Not only did we shake-down methods and gear for GLOBEC's intensive 1995 field season, a great deal of interesting science was done. Overall, Pls accomplished more than their stated goals. Despite the enormous amount of equipment and scientific personnel crowded into the *Iselin*'s small dry lab, morale was high, and we were all still speaking to each other when we left the ship at the end of the cruise. The *Iselin*'s Captain and crew were competent and extremely helpful. Their skill in locating our sometimes dysfunctional ARGOS drifters under conditions of fog and darkness was truly impressive.

Cruise Narrative

Background. Sea surface temperature maps processed and analyzed from AVHRR data by James Bisagni and associates at the National Marine Fisheries Service Narragansett Laboratories greatly improved our ability to plan our sampling program, particularly with respect to sites where we did NOT want to be because of the presence of streamers from Gulf Stream rings. The images show that sea surface temperature increased from ~7°C to ~9°C during the duration of cruise Cl9407. Sea surface temperature maps from March 1994 showed a large Gulf Stream ring located in slope water off the northeast peak of Georges Bank. As the season progressed, the ring moved south and west along the Bank, and a second large ring moved off the northeast peak. By the time cruise Cl9407 was on the Bank, both rings were pressed against the Bank's southern flank and a number of jets and streamers from the two rings had crossed the 100 meter isobath (Figures 1, 2 and 3).

Initial broadscale survey. 25 - 29 May 1994. The R.V. Columbus Iselin departed Narragansett at 0900 (EDT), sailing to our first station north of the Great South Channel. The initial, rapid bankwide survey sampled 18 stations. The cruise track began north of the Great South Channel, proceeded east across the Bank crest, then southeast to the 100 m isobath (Figure 4). The track proceeded toward the northeast peak between the 60m and 100m isobaths to approximately 41.5 °N, then went north to a station in Georges Basin in the southern Gulf of Maine. From here, we steamed to a station in slopewater east of the northeast peak, then sampled at three more stations in a southwesterly direction to conclude the survey. The original cruise plan called for two stations in the slope water at the end of each of the first two transect legs, but these were cancelled due to inclement weather on 26-27 May.

CTD casts and stratified MOCNESS sampling for zooplankton were done at each station. Hydrographic data collected included CTD, fluorometry and transmissometry, with bulk water samples collected for analysis of major nutrients, size-fractionated chlorophyli (total, $<\!20~\mu\mathrm{m}$ and $<\!5~\mu\mathrm{m}$), and microplankton (=phytoplankton + protozoa). Microplankton samples were collected for analysis by flow cytometry, automated epifluorescence microscopy, and inverted microscopy in order to describe the entire suite of nano- and microplankton prey potentially available to target copepods and larval fish. MOCNESS (150 $\mu\mathrm{m}$ mesh) samples were collected on Georges Bank from three depth strata at stations < 100m depth and four strata at stations > 100m depth. MOCNESS samples were collected from 9 strata in deep basins in the southern Gulf of Maine.

Stratified site. 29 - 31 May 1994; Station 21. Although cruises working on the Bank immediately prior to Cl9407 reported well developed stratification on the southern flank, by the time we arrived, a series of storms had completely mixed the water column at every on-bank station sampled in the initial survey. Thus, the initial "stratified site", located in ~90 m of water, was chosen because the bankwide survey observed abundant Calanus finmarchicus in the area, primarily stages CV and adult females, with some

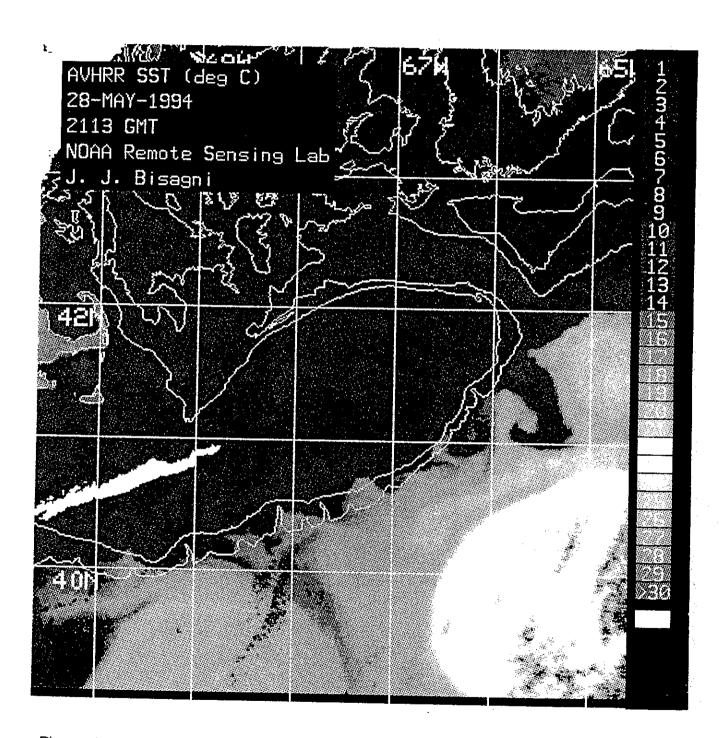


Figure 1. Sea surface temperature on 28 May 1994. 100 m and 200 m isobaths are shown.

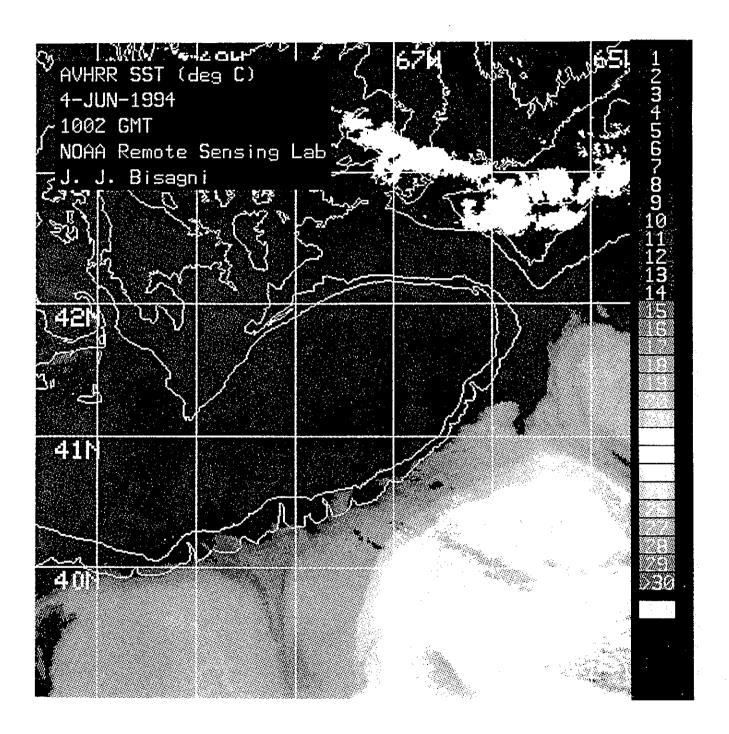


Figure 2. Sea surface temperature on 4 June 1994. 100 m and 200 m isobaths are shown.

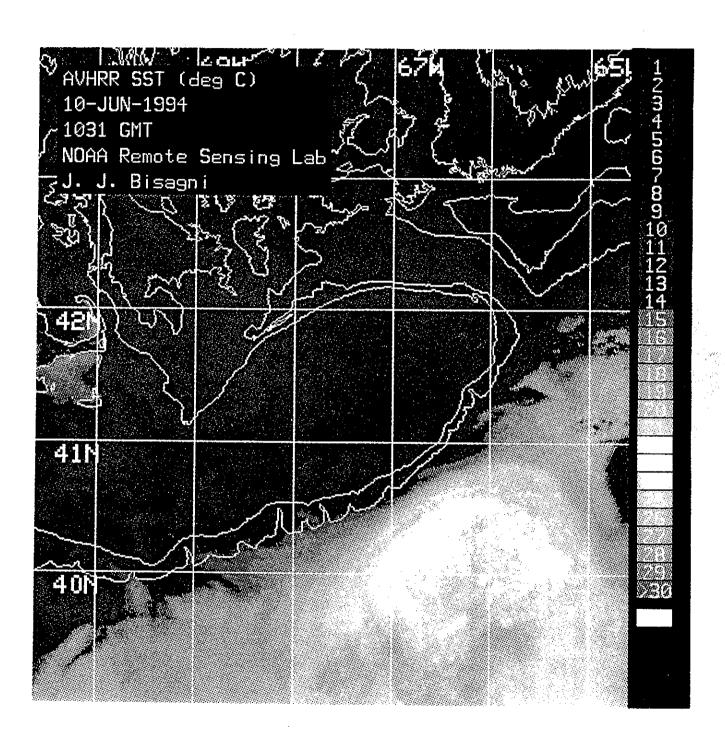
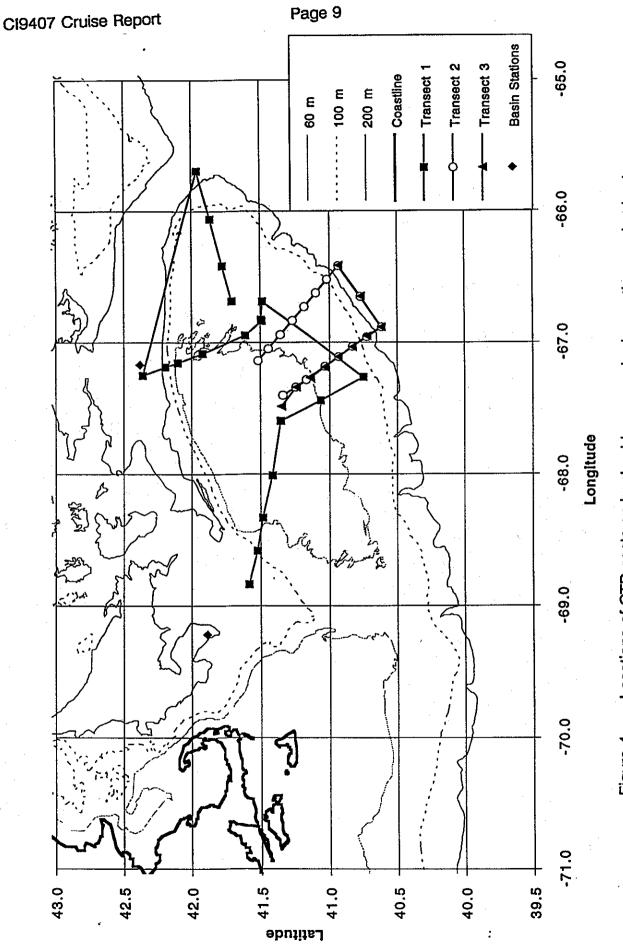


Figure 3. Sea surface temperature on 10 June 1994. 100 m and 200 m isobaths are shown.



Locations of CTD casts on bank-wide survey, hydro sections and at basin stations. Figure 4.

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CIVs. An ARGOS drifter drogued at 10 meters was deployed at the site (initial location 41°49.94'N, 66°12.57'W) and tracked by the ship's bridge for the 48-hour duration of station-keeping operations. Station-keeping activities began with a CTD cast and ring-net tows to characterize the water column and the zooplankton, followed by rosette casts and diaphragm pump deployments to collect water for experiments. A series of net tows, usually with a 150 μ m mesh 3/4m ring net, were then done to collect live zooplankton for experiments. Vital rates measured were growth, feeding and egg-laying of target copepods, and growth, condition and feeding of larval cod.

CTD, MOCNESS and zooplankton pump deployments were done at approximately 1200 and 2400 each day to collect data on hydrography and zooplankton distribution and abundance. Discrete water samples were collected from each noon CTD cast as described for the bankwide survey; in the midnight casts only the CTD was deployed. The TAPS and VPR were deployed at regular intervals throughout each day/night cycle.

The drifter followed the tidal ellipse, moving slowly southward during the 48 hours the station was occupied (Figure 5). The water column remained well mixed during this time. Upon completion of operations at Station 21, the drifter was left in place, and we proceeded to a station located in the mixed area of the Bank.

Mixed site. 31 May - 2 June 1994, Station 22. Our initial mixed site was chosen to lie within the 60 meter isobath on the southern flank of the Bank, but well away from shoal areas. The protocol at this station was as described for the stratified station. In contrast to the stratified station, Calanus finmarchicus were not abundant, and the zooplankton biomass was dominated by the pelagic hydroids, later identified as Clytia cylindrica, also observed during the predation cruise. The copepod assemblage at this station consisted of Centropages in the upper water column, and Temora and Pseudocalanus throughout the water column. The drifter at this station (initial location 41°19.66'n, 67°15.52'W) also followed the tidal ellipse, moving slowly along the 60 meter isobath in a southerly direction during the 48-hour duration of station-keeping (Figure 5). Failure of the VPR's one and only strobe early in station-keeping operations rendered the instrument unusable for the remainder of Leg A. Upon completion of operations at Station 22, the drifter was left in place, and we began a hydrographic section to collect data for GLOBEC investigator, Bob Beardsley.

Hydrosection 1. 2-3 June 1994. CTD casts were done at 11 stations of a hydrosection consisting of two cross-bank and one along-bank transects (open circles in Figure 4). The cross-bank transect consisted of 8 stations running from the well-mixed area near the Bank crest (41° 20.38'N, 67° 24.18'W) to the edge of a warm-core ring at the shelf break (40°36.93'N, 66°52.99'W), the along-bank transect consisted of 3 stations oriented parallel to the shelf break (40°36.93'N, 66°52.99'W to 40°56.08'N, 66°25.00'W), and the second 8-section transect extended from the ring water (40°56.08'N, 66°25.00'W) back onto the crest of the bank (41°30.96', 67°08.43'W). MOCNESS casts were done at 3 stations on each transect. The airborne LIDAR group's first fly-by occurred on 3 June

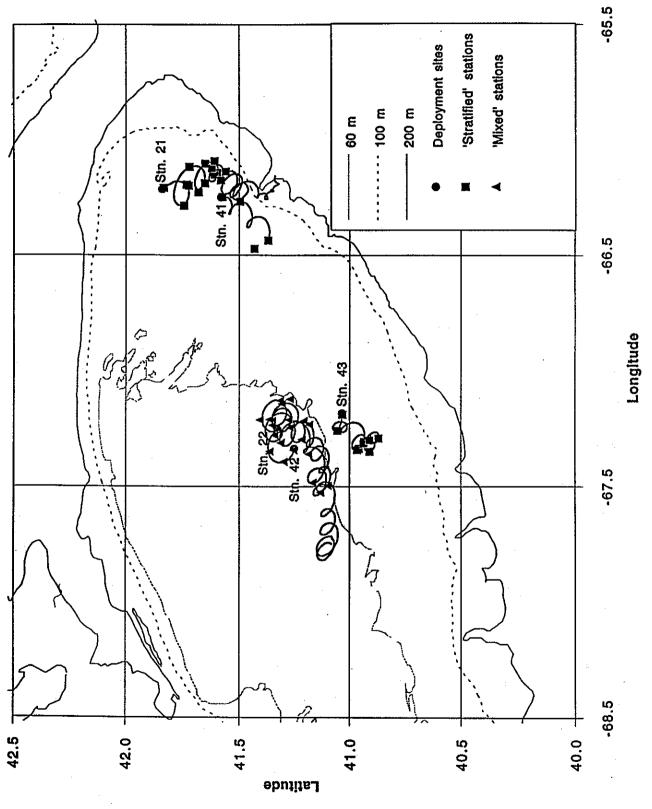


Figure 5. Locations of CTD casts during drifter stations.

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at 1424 (GMT).

Return to stratified site. 3-5 June 1994. After completing the hydrosection, we returned to the first drifter at the "stratified" site. Although we arrived in the vicinity of the drifter in the dark, and the drifter's strobe had failed, the bridge, ably assisted by Peter Garrahan, was able to home in on the drifter's ARGOS signal and pick the drifter's high flyer out of a veritable potpourri of noise on the ship's radar. Upon arrival at station, the drifter was retrieved for a minor repair recommended by its manufacturer, then redeployed. We occupied the station for 24-hours, following a truncated version of the previous 48-hour routine. By the time station-keeping activities were concluded, the water column had begun to stratify weakly. Upon termination of station-keeping activities, the drifter was retrieved because it had become entrained in a streamer from the northernmost Gulf Stream ring, and had begun to spiral off-bank (Figure 5). The ship then steamed to Woods Hole for a scheduled personnel change.

Port stop. 5-6 June 1994. Flying at record speed (15 knots on a 14-knot vessel), the *Iselin* arrived in Woods Hole exactly 3 hours before the bars closed, allowing scientists and crew a pleasant, not-too-rowdy, night on the town. On 6 June, the TAPS (Mark Berman and Joe Kane) and larval fish (Ione Hunt von Herbing and Linda Davis) contingents as well as VPR jockeys Frederika Norrrbin and Marty Marra, zooplankton biologist Jeff Runge, hydrographer Ari Epstein and Durbin assistant Jonathan Hopkins left us and were replaced by hydrographer Ed Dever, Durbin assistant Dave Avery, VPR personnel Andy Girard and Phil Alatalo, and WHOI summer student Lucie Blanchard. The VPR group spent June 6th coaxing their instrument back to life. At one point it was declared brain dead, and the group began offloading their gear. Moments later, Al Morton called from North Falmouth to say he'd located the problem. The gear was reloaded, the VPR was tested off the side of the ship for an hour, and CI9407-Leg B got underway only a few hours late at 1500 (EDT) beneath gray skies with a brisk wind.

Return to mixed site. 7-10 June 1994, Station 42. We returned to the mixed site, with the bridge and Peter Garrahan locating the drifter without problems in the fog. Station-keeping protocol was as before, with the exception that the station was occupied for 72 hours. The increase in station-keeping time was arrived at as a group decision in order to accomplish more experimental work at each station while allowing the experimenters to get more than a few hours of sleep. The drifter continued to follow the tidal ellipse moving southwest along the 60 meter isobath (Figure 5). It was retrieved at the end of station-keeping activities. The airborne LIDAR group's second fly-by occurred on 8 June at 1428 (GMT).

Return to stratified site. 10-13 June 1994, Station 43. We steamed to the spot where our hydrographer's best guess said the drifter would have gone during the time elapsed had it not been entrained in the streamer (Figure 5), and deployed the drifter. Station-keeping protocol was as before, with the exception that the station was occupied for 72 hours. The drifter followed the tidal ellipse while moving south and west between the 60

and 100 meter isobaths. Stratification evolved during the 72 hours we occupied the station. A chlorophyll maximum developed at ~20m and slowly moved deeper over time. The phytoplankton flora in the maximum appeared to be dominated by diatoms contained in a gelatinous matrix, later identified as *Chaetoceros socialis*. An assemblage of abundant microheterotrophs including heterotrophic flagellates, heterotrophic dinoflagellates and aloricate ciliates was associated with the chlorophyll maximum. Upon completion of operations, the drifter was retrieved, and we began a hydrographic section to collect data for GLOBEC investigator, Bob Beardsley. The airborne LIDAR group's third and last fly-by occurred on 10 June at 1650 (GMT).

Hydrosection 2. 13 June 1994. The hydrosection done during Leg A was repeated during Leg B. The first cross-bank transect was repeated as a CTD transect (solid triangles in Figure 4), with MOCNESS casts done at 3 stations. The second was done as a VPR tow-yo transect, described below.

VPR transect. 13-15 June 1994. One long (approximately 200 km and 26 hrs) tow-yo transect was done across Georges Bank from slope water (40°54.62'N, 66°23.39'W), across the mixed area and into Georges Basin (42°38.25'N, 67°16.81'W). Data collected by the VPR included CTD, fluorometry, transmissometry and zooplankton counts.

Deep basin sampling. 15 June 1994. Georges Basin and Wilkinson Basin (solid diamonds on Figure 4) in the southern Gulf of Maine were sampled to collect critical information concerning *Calanus* diapause strategies for Charlie Miller. CTD casts with full hydrographic sampling were done and MOCNESS samples were collected from 9 depth strata at each station. An additional MOCNESS cast was done at the Wilkinson Basin station to collect and freeze *Calanus* for use in a mackerel feeding study being done by GLOBEC investigator Ann Durbin.

Upon completion of sampling, we began steaming to Narragansett. After some delay due to fog, we arrived at the GSO dock at 1000 (EDT) hours on 16 June 1994.

Individual PI Reports

A. Hydrography (Ari Epstein and Edward Dever)

Leg A: 25 May-5 June 1994

Sixty-seven CTD casts were made on CI94-07A, including two hydrographic transects: one 8-station transect running from the well-mixed area near the top of Georges Bank, across the Bank's southern flank and off the shelf break into the edge of a warm-core ring, and another 8-station transect from the ring water back onto the top of the Bank. (These sections were repeated in leg B; the first transect was repeated as a CTD transect, and the second as a VPR tow-yo transect.) The CTD was also used during the bank-wide survey made at the beginning of Leg A and during various 24- and 48-hour stations. The CTD was a Neil Brown Mark III CTD. Data were acquired using the EG&G Data-acquisition model (version 5.1 revision 9) and processed using the EG&G Post-processing module (version 3.0). Most CTD casts (except for "TAPS yo-yo's") were made while the hydrowire was paying out at 30 meters/minute. 150-kHz and 600-kHz RDI ADCPs were running nearly continuously throughout the leg.

The bank-wide survey was done between May 25 and May 29, and included CTD casts 1-19. Casts 1-5 were made during a transect from the northern Great South Channel to a point just south of Georges Shoal, near the center of Georges Bank. Casts 5-7 were made on a transect that extended southward across the shelf-slope front; two casts that had been planned for this transect and the next one were cancelled due to weather. Fog and heavy weather, with some thunderstorms, were present for much of the bank-wide survey. Casts 8-15 were made on a transect from the Bank's southern flank across the northern flank and into Georges Basin. Cast 16 was made in the Northeast Channel, and casts 16-19 were made during a transect from the Northeast Channel to a point near the Hague Line, about halfway between the northern and southern edges of the Bank. During the survey, very little stratification was found in waters shallower than about 80 meters. (The deeper casts, casts 15 and 16, showed the expected degree of stratification, and T-S diagrams there indicated the presence of the expected water masses.)

Casts 20-33 were made during Leg A's first 48-hour station (May 29-31), which was located near the northeast corner of the Bank in about 90 meters of water. This site was chosen in lieu of the expected "stratified site". Casts 34-46 were made during the Leg A's second 48-hour station (May 31 - June 2), located at a site south of Georges Shoal. Many of the CTD casts were "TAPS yo-yo's", in which the CTD was raised and lowered about 5 times at a wire speed of 10 meters/minute. To ensure good CTD data, TAPS yo-yo's were followed by regular CTD casts, made at 30 meters/minute.

Casts 47-54 were made during the hydrographic transect from the top of the Bank

across the southern flank and into the edge of a warm-core ring. Casts 54-56 were made while cruising northward along the 500-fathom isobath, and casts 56-63 were made during the transect from the 500-meter isobath back to the top of the Bank.

Casts 64-67 were made during a 24-hour station, located in about 90 meters of water, at a site south of the site of the second 48-hour station.

Leg B: 7 - 16 June 1994

CTD operations for Leg B of Cl9407 were carried out using a Neil Brown Mk III CTD with EG&G Data Acquisition Module version 5.1 rev 9 and EG&G Post-Processing Module version 3.0. This CTD was attached to a 12 bottle tone fire rosette, unlike the CTD used for Leg A, which was attached to a conventional rosette. The CTD was also equipped with a fluorometer and transmissometer. All casts in water depths of less than 200 m were made at a descent/ascent rate of 30 m/minute. In depths deeper than 200 m a descent/ascent rate of 45 m /minute was used in most cases. CTD casts were taken upon arrival at a mixed layer site within the 60m isobath (casts 68--76), at a stratified site between the tidal front and the shelf/slope front (casts 77--86), in a cross-bank (87--95) and along-bank (95--97) section, in Georges Basin (98), and on the southern edge of the Wilkinson Basin (99). All CTD casts are listed with station numbers in the GLOBEC event log for CI9407 as well as the CTD log. Casts 68 and 69 were subject to severe noise problems. This was not due to an inherent problem with the CTD, but instead may have been a problem in the termination of the conducting cable or in the deck box or power supply. Unfortunately, the data from casts 68 and 69 are for most purposes unusable. The first usable cast, 70, was taken approximately 6 hrs after arrival at Station 42, and indicated top to bottom uniformity at this site. Therefore, it should be possible to use the top and bottom bottle salinities from cast 68 and the ship's sea surface temperature measurements to get a fairly good idea of hydrographic conditions upon arrival at station 42. Also, comparison of casts 70 and 71 indicate initially slow temporal evolution of temperature and salinity at Station 42.

Subsequent CTD casts were not subject to the severe noise problems of Stations 68 and 69. Some spiking (which should be removed in post-processing) was evident. Minor problems with bottle misfires, operator errors, etc are noted in the CTD log. Three casts (80, 82, and 89) were immediately repeated when larger problems were encountered.

CTD casts at the mixed site, Station 42, indicated the drifter could have been moving toward the tidal mixing front. The first successful cast, 70, was uniform in both temperature (~ 0.02 deg C) and salinity (~ 0.01 ppt). As the drifter moved south and west, subsequent stations showed a thin near surface layer of warm water (top to bottom differences up to 1.5 °C), and especially in casts 75 and 76, a slight salinity increase toward bottom (~ 0.1 ppt). This is consistent with Station 42 VPR tows which indicate the mixing front was nearby. It should also be noted that after June 8, seas were

calm, with little wind stress, so that wind induced vertical mixing was probably unimportant.

Stratified site casts (Station 43) showed top to bottom temperature differences of roughly 1.6 ° C and above. Near surface warming in the upper five meters or so probably occurred, as wind speeds were light during this time and skies were nearly clear. This can be checked since all data necessary to make surface heat flux calculations (except incoming long wave radiation) were recorded by the ship. Though temperature stratification (below the near surface layer) was not much greater than at Station 42, far greater salinity stratification existed than at station 42. Top to bottom differences of at least 0.1 ppt and up to 0.5 ppt existed. Near bottom salinity varied from 33.05 to 33.28, while surface salinity tended to be from 32.7 to 32.8 ppt. The strongest stratification existed in the upper 25--30 m. A maximum in fluoresence and minimum in transmission was coincident with the base of the stratification at 25--30m.

The cross bank CTD transect was a repeat of the transect (47--54) performed during Leg A of Cl9407. Both showed the tidal mixing front at the 60 m isobath and the shelf slope front. However, there were some differences. During Leg A, the 33 ppt salinity contour was nearly vertical; during leg B, it had a wedge shape and penetrated further onshelf below 30 m. Higher salinity warm core ring water evident during leg A retreated approximately one station spacing (7 nautical miles) further off-shelf, confirmed by satellite sea surface temperature imagery. Overall warming also occurred on the Bank and the cold band was less evident during leg B.

B. Ancillary Hydrographic Data: chlorophyll, nutrients, phytoplankton and microzooplankton (Dian Gifford, Michael Sieracki and Terrance Cucci)

Objective: To describe quantitatively and qualitatively the nano- and micro- plankton prey fields potentially available to copeods and early stage larval fish on Georges Bank. The data are to be used in interpreting experiments done with target copepods and larval fish described in sections E. and G. below.

Bulk water samples for chlorophyll, major nutrients and phytoplankton and microzooplankton identification were collected from all CTD/rosette casts during the initial bank-wide survey and from each noon CTD/rosette cast during station-keeping operations. At stations where the water column was mixed, samples were collected from the top, middle, and bottom of the water column. At stations where the water column was stratified, samples were collected from 5-9 depths, determined by the particular combination of water column depth and hydrographic features present. For example, if a chlorophyll maximum or density discontinuity was present, samples were collected from above, within and below the feature.

Chlorophyll. One-liter of water was drained through silicone tubing into dark collecting bottles. Samples to assay 3 size fractions of chlorophyll (total, <20 μm and <5 μm) were collected by gentle vacuum filtration of 50 ml onto GF/F filters, which were frozen in liquid nitrogen, then transferred to a freezer until analysis. Size fractionation was done by passing 50 ml of sample through sieves constructed of Nitex mesh of the appropriate porosity. All chlorophyll samples were collected in triplicate. Chlorophyll analysis was by fluorometry.

Nutrients. ~ 50 ml of water from the chlorophyll bottles was poured into precleaned plastic bottles from the chlorophyll bottles following three rinses of ~ 50 ml each with the same water. The samples were frozen until analysis with a nutrient autoanalyzer.

Flow cytometry. 50 ml of water was collected from each Niskin bottle into plastic vials. Subsamples were analyzed immediately using a Beckton-Dickson FACscan flow cytometer. This yielded counts of phytoplankton in size classes of 1-5 μ m, 5-10 μ m, and greater than 10 μ m.

Image analysis-Epifluorescence microscopy. 50 ml of water was collected from each chlorophyll bottle and preserved with glutaraldehyde. Subsamples were post-stained with a mixture of proflavine and DAPI, filtered onto black 0.2 μ m Nucelopore filters, mounted on glass slides in a drop of non-fluorescent oil, and frozen pending subsequent analysis of nano- and micro- phytoplankton and nano-zooplankton by automated epifluorescence microscopy. Some samples were analyzed aboard ship but the majority will be analyzed back at the lab. They will provide numerical abundance, biomass and size distributions for both the phototrophic and heterotrophic nanoplankton which comprise part of the *Calanus* prey field.

Inverted microscopy. Two hundred and fifty ml of water was drained from each Niskin bottle into sample bottles containing an amount of acid Lugol's solution appropriate for a final preservative concentration of 10% (vol:vol). Samples were stored in the dark pending analysis using an inverted microscope. These samples will provide numerical abundance, biomass, and size distributions of the micro-phytoplankton and nano- and microzooplankton which comprise part of the *Calanus* prey field.

Preliminary results: Preliminary observations of the survey samples indicate an extremely diverse assemblage of nano- and microplankton on the Bank including diatoms (chains and single cells), phototrophic dinoflagellates, Prymnesiophytes, Cryptophytes, mixotrophic ciliates, and a diversity of heterotrophic (colorless) protists including ciliates, dinoflagellates, choanoflagellates, and undistinctive flagellates from 2 μ m to 15 μ m in size.

C. Zooplankton Process Studies: Growth and development rates of Calanus finmarchicus and Pseudocalanus sp. (The Durbin Group: Ann Durbin, Edward Durbin and the Planktoneers: Robert Campbell, Maria Bemis, Peter Garrahan, David Avery and Jonathan Hopkins)

Objectives:

- (1) To estimate the recruitment of *Calanus* and *Pseudocalanus* on Georges Bank during the study period.
- (2) To determine the production rates of Calanus and Pseudocalanus at the mixed and stratified sites.

Methods:

MOCNESS and Pump sampling. An initial MOCNESS (150 μ m mesh nets) survey, and several cross-bank transects later in the study (Figure C-1), were conducted to determine zooplankton abundances throughout the region . At the drifter sites both MOCNESS and pump (50 μ m mesh nets) samples were taken at 12 hour intervals for a period of 48 to 72 hours (Figure C-2). These samples will be counted to determine abundance and stage distributions of the target species Calanus and Pseudocalanus as well as other important species such as, Centropages and Temora. It should also be possible to determine recruitment and development rates of the target species at the drifter locations if distinct cohorts are present.

Mesocosm experiments. Development rates of *Calanus* and *Pseudocalanus* were determined in 30-50 gallon mesocosms at the drifter sites. Artificial cohorts were created by screening (Kimmerer and McKinnon 1987) and incubated on deck, in tanks containing ambient water and ambient water enriched with a mixed phytoplankton culture. The mesocosms were sampled every 12 hours for 2 to 3 days and the animals preserved for later stage enumeration. In addition, initial and final samples of copepods from the dominant stages in each size fraction were taken for length, carbon and nitrogen content. The results from the mesocosm experiments will provide estimates of development rate that can be compared to in situ estimates of development rate from MOCNESS and pump samples, as well as determine if development rates were food limited by comparing development rates in ambient and enriched treatments.

Growth experiments. The growth in weight of one selected copepodite stage (C4 - Calanus or C5 Pseudocalanus) was determined at the drifter locations. Copepods were sorted into bottles (2 or 8 liters) containing ambient water from predetermined depths or enriched water, and incubated in a water bath for 1 to 2 days. An initial subsample of copepods was videotaped for length determination and placed in tin boats and dried over desicant for later carbon and nitrogen analysis. At the end of the incubation animals were also collected for length, carbon and nitrogen analysis. The results from these experiments will provide estimates of growth in terms of length, carbon

and nitrogen, as well as determine if growth is food limited.

General observations. During most of the cruise there was an absence of early copepodite stages (C1-C3) of Calanus, so most of our experimental work focused on stage C4 Calanus and Pseudocalanus copepodites. However, we did observe what appeared to be Calanus nauplii in the small size fraction mesocosm experiments, and in the final mesocosm experiment at Station 43, early Calanus copepodites (C1 and C2) were found at the end of the incubation. So, it appears that we were too late in the season to obtain good estimates of growth and development rates on the first generation of Calanus, but perhaps we will be able to determine development rates on the second generation through the pump samples and mesocosm experiments.

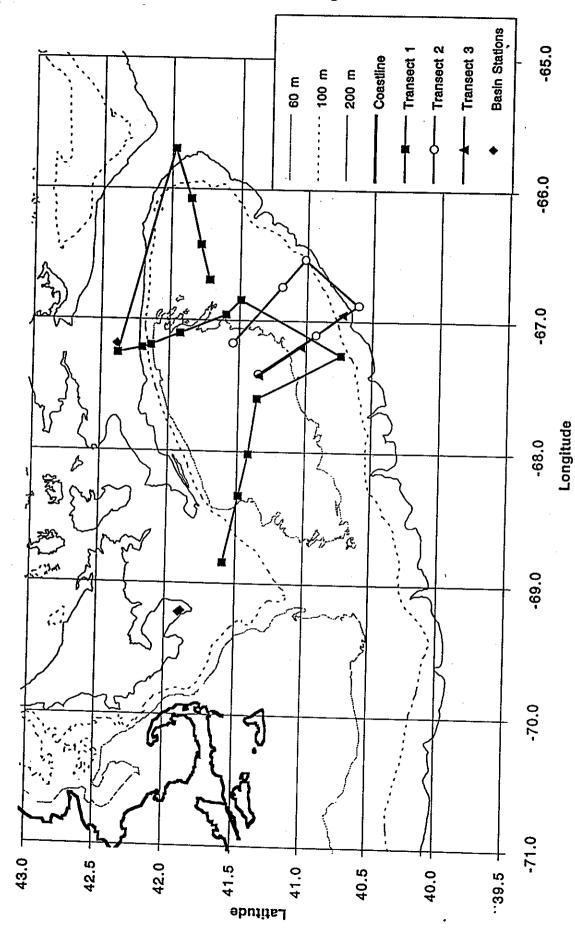
D. Zooplankton Process Studies: Egg production and recruitment rates of Calanus finmarchicus and Pseudocalanus sp. (Jeffrey Runge and Stephane Plourde)

Objectives:

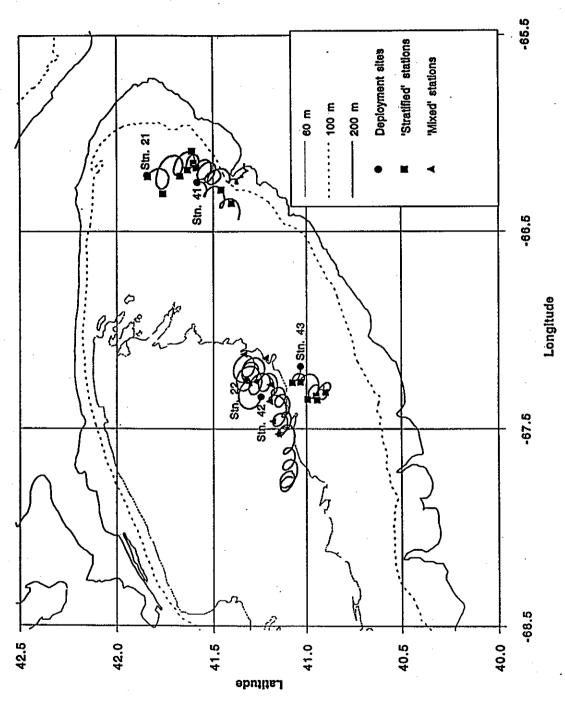
- (1) Examine the hypothesis that the daily production of eggs of *C. finmarchicus* and *Pseudocalanus* sp. in the water column (eggs m² d¹) is different between stratified and mixed regions on Georges Bank, due to variations in both the specific egg laying rate (eggs female⁻¹d⁻¹) and the abundance of females (females m²).
- (2) Conduct preliminary studies of magnitude and causes of mortality of eggs, an estimate of which is needed to determine recruitment rates, where recruitment is defined as the number of eggs hatching m²d¹.

Methods:

C. finmarchicus. Egg laying rates were measured during the first leg at Sta. 17 (in the northeast Channel), Sta. 21 (first drifter station), Sta. 22 (second drifter station), Sta. 41 (third drifter station) and during the secong leg at Sta. 42 (fourth drifter station, well-misted region), Sta 43 (fifth dirfter station, statified region) and Sta. 54 (Georges Basin). At all stations, a standard method was employed to estimate egg production. Forty females were sorted from the catch of an oblique plankton tow (bottom to surface) taken in the afternoon. Individual females were placed into filtered seawater in petri dishes, which were incubated at 8-9 °C for 24 h on a 12 h cycle of darkness and dim light. The dishes were inspected at approximately 8 h intervals and the numbers of eggs released were recorded. At some stations, additional females were sorted and variations of the standard method, including incubations in beakers containing seawater enriched with a high concentration of Gymnodinium and incubations in 45 ml. culture flasks (as a



positions during the initial Bank-wide survey (25-28 May). Transect 2 (2-3 June) and transect 3 (13-14 June) are hydrographic sections 1 and 2. The deep basin stations (15 June) are also shown. Location of MOCNESS stations during transects. Transect 1 shows station Figure C-1.



Positions of initial drifter deployments and each of the Stations 22 (31 May-2 June) and 42 (7-10 June) are located at the Mixed Site. MOCNESS and zooplankton pump stations are shown. Stations 21 (29-31 May), 41 (4-5 June) and 43 (10-13 June) are located at the Startified Site. Drifter paths. Figure C-2.

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simpler alternative to petri dishes), were carried out. Pump and MOCNESS samples taken at these stations will be analyzed for water column abundance of eggs and females, respectively.

Pseudocalanus sp. Egg production rates (i.e. rate of production of clutches of eggs, which are carried by the females until hatching) were estimated at the three drifter stations. Non-bearing females (50-100) were sorted from the catch of the net tow and incubated in ambient seawater (5-15m) at 8-9 °C for 24 h. Another batch of females bearing a complete (approx. 20 eggs) or partial (typically 1-5 eggs) clutch were incubated individually in petri dishes for 24 h. The females were preserved for later analysis. To estimate birth rate (i.e., recruitment rates), females and eggs were collected from the water column with a pump, the outflow from which was filtered through a 50 μm mesh plankton net. At the third drifter station, Pseudocalanus was also collected with a 50 μm plankton net towed vertically from the bottom to the surface.

Egg production of *Temora* and *Centropages*. At the second drifter station, egg production rates of these two species (identified as *T. Iongicornis* and *C. hamatus*) were measured by the standard incubation technique described above for *C. finmarchicus*.

Predation on copepod eggs by polyps of hydroids. At the well-mixed site (second drifter station), a predation experiment was carried out, in order to obtain a preliminary estimate of feeding rates of hydroids on the eggs of dominant copepods at this site. A revised protocol (based on experiences from the preliminary experiment) is attached.

Results and discussion:

C. finmarchicus. Egg production rates ranged between 15 eggs female data at Sta. 17 to 35 eggs female data the third drifter station. Egg production rates of females fed Gymnodinium were not significantly different from the standard estimate (three separate experiments). The number of eggs released during a spawning event averaged 50, which is consistent with previous observations of clutch size of females of equivalent body size (Runge and Plourde 1994). During the secong leg, it wasn't possible to measure egg production rate of C. finmarchicus at the well-mixted station due to low fermales density. Howerver, egg production rate of the species at Sta. 43 (startified region) was >60 eggs female day at the station. Finally, at the Georges Basin station, preliminary observations showed that the majority of the females in the population at this location were reproductively active at the start of the egg laying experiment, suggesting that the egg production rate should be of the same magnitude as rates measured previously at the stratified station.

Assuming (until actual measurements are made at URI) carbon weights of females and eggs of 112 μ gC and 0.23 μ gC, respectively, levels observed during the first leg

represent an egg laying rate of 3%-7% of body carbon per day. Laboratory experiments indicate that maximum egg production rate of *C. finmarchicus* at the measured ambient surface layer temperatures (8-9 °C) is 6-7% per day (Runge and Plourde 1994). This implies that the *Calanus* populations on the Bank were laying eggs at approximately one half maximum level at Station 17 and at near maximum levels at the second and third drifter stations. Examination of the state of reproductive maturity of females in the catch indicated that a small but significant fraction (perhaps up to 25% at some stations) either carried a spermataphore (these females are typically not ready to spawn) or were filled with lipid and appeared to be in diapause mode. During the second leg, egg laying rates represented 8%-14% of body carbon per day. It has to be noted that incubation temperatures ranged between 9-11° C and that surface temperature were ranged between 10-12° C at the stratified station during experiments. These high temparature combined with the high spawning frequency observed (>85%) might explain these high egg production rates.

Pseudocalanus. There appear to be at least two species of Pseudocalanus at the study sites ($P.\ moultoni$ and $P.\ newmani$). Most females were not bearing eggs; a relatively small fraction (estimated to be about 15-20%) were carrying small clusters of 1-5 eggs. Rarely, females were observed with full clutches (18-22 eggs), although some fraction produced full clutches during the 24 h incubation. It appears that egg clusters are easily separated from females during net tows, and accurate estimates of egg ratio will require careful counts of free Pseudocalanus eggs (120 μ m diameter on average) in sample jars.

Temora and Centropages. Temora and Centropages females were abundant at the well-mixed drifter station (in contrast to *C. finmarchicus*, which was relatively uncommon). All Temora females spawned, averaging 71 eggs per clutch, during two separate 24 h incubations. Sixty percent of Centropages females released an average of 46 eggs, yielding a population egg production rate of 28 eggs female d'. Another experiment was done during the secong leg at the well-mixed site (Sta. 42), with similar egg production rates anticipated for Temora and Centropages.

Predation on copepod eggs at the well-mixed station. During the counting of *Calanus* eggs at Sta.22, it was observed that a hydroid polyp, accidentally transferred to a petri dish, had ingested 11 recently spawned eggs. Ten hours later, seven partially-digested eggs were still visible. Subsequent observations showed that hydroid polyps are adept at sweeping up and feeding on substantial numbers of eggs clustered at the bottom of the dish and that the digestive tracts of some hydroids captured in net tows from the station were packed with copepod eggs.

Two prey-disappearance experiments were designed to estimate feeding rates of polyps on suspended eggs. Between 10 and 40 hydroid polyps were introduced into 1 liter bottles containing 100 Temora or Calanus eggs (first experiment) and 150 Temora eggs (second experiment). Replicate bottles were incubated with controls for 3, 13, and

17 h intervals (first experiment) and 3, 6, 9 h intervals (second experiment). The Frost equations were used to calculate clearance rate. During the first experiment, in all replicates, eggs were found in the digestive tract of at least some polyps; the maximum number of eggs observed in a single polyp was 5. Maximum (based on numbers of eggs remaining at the end of the experiment, corrected for controls) and minimum (based on the actual number of eggs found in polyp digestive tracts) clearance rates were 2.2 and 0.6 ml polyp h.1, respectively, on *Calanus* eggs and 2.8 and 1.1 ml polyp h.1 on *Temora* eggs. Preliminary examination of data from the second experiment seems to confirm these results despite a problem with the last control bottle (9h interval, lower number of eggs recovered).

Copepod recruitment rates on Georges Bank between 24 May and 17 June 94: first impressions. There is the potential for large, meso-scale spatial and temporal variation in copepod recruitment rates on Georges Bank during this period (*Pseudocalanus* species may be the exception). Specific egg production rates of *Calanus* varied by a factor of two, probably due in part to food limitation, but also due to spatial variability in the proportion of reproductively active females. Egg production rates were highest at the third drifter station, probably reflecting the higher primary production associated with the onset of stratification. At the same time, there is the potential for great variability in the number of females. Bank-wide, the abundance of females is probably growing as some proportion of the first generation CV's molt into reproductively active adults as suggested by higher egg production rates and spawing frequency measured at the stratified station during the second leg. On the shallow center of the Bank females are probably being eaten as fast as they drift in, so that, even though specific egg production rates are high, recruitment rate, in terms of number per m², is low.

In addition to variablity in numbers of eggs produced, there are clear indications of strong meso-scale variations in the abundance and composition of predators on copepod eggs and nauplii. The presence of hydroids on the central part of the Bank is potentially devasting to recruitment of *Calanus*, *Temora*, and *Centropages*, all species that spawn their eggs into the water. The preliminary experiments (assuming for the moment that encounter rates in botlles on a grazing wheel accurately reflect encounter rates of eggs with hydroids in the water column) suggest that a concentration of between 15 and 70 polyps (not colonies, but individual tentacular polyps) are sufficient to clear the water daily of eggs. *Pseudocalanus* eggs, which are borne by their parent, may be able to avoid the tentacular clutches of hydroids. In any case, the qualitative impression is that the central Bank is relatively devoid of early life stages of copepods, despite the relative high production rates of dominant species (like *Temora*, and *Centropages*) and that the source of exceptionally high early mortality is the invertebrate predator field. For related reasons, the central part of the Bank does not appear to be conducive to growth and survival of fish larvae.

Reference: Runge, J. A. and S. Plourde. 1994. Fecundity of Calanus finmarchicus in coastal waters of eastern Canada. ICES Workshop on the trans-latitudinal study of

Calanus finmarchicus in the North Atlantic. Oslo, 6-8 April, 1994.

E. Zooplankton Process Studies: Feeding rates and diet of Calanus finmarchicus. (Dian Gifford, Michael Sieracki and Terrance Cucci)

Objective: To document the contributions of nano- and micro- zooplankton and phytoplankton prey to the diets of all copepodid stages of *Calanus finmarchicus* under conditions of water column mixing and stratification and to measure *Calanus*' ingestion rates on the prey items.

Methods:

Twelve experiments were done with *Calanus finmarchicus*, summarized in Table E-1. Copepods from live net tows were sorted into 1-liter polycarbonate incubation bottles containing the natural prey assemblage, and the disappearance of prey was followed over 24-hour experimental duration. Control treatments consisted of the natural assemblage alone; experimental treatments were the natural assemblage with copepods added. The bottles were incubated on-deck on a slowly rotating plankton wheel whose temperature was controlled by flowing seawater. Experiments were done with adult females and copepodid stages C4 and C5. Two categories of stage C5 were recognized: "fat" copepods with full oil sacs and "thin" copepods with partly full oil sacs.

The numerical abundance and biomass of phytoplankton prey were quantified in 3 size fractions of chlorophyll \underline{a} (total, <20 μm and < 5 μm), cells counted by flow cytometry, cells counted by image analysis-epifluorescence microscopy, and cells counted by inverted microscopy. The numerical abundance and biomass of nano- and microzooplankton prey were quantified in cells counted by image analysis-epifluorescence microscopy, and cells counted by inverted microscopy. All copepods were collected from the incubation bottles at the end of each experiment for CHN analysis.

Microscopic analyses will be performed in our home laboratories, and final results will not be available for several months. However, preliminary results from flow cytometry indicated that the 3 life history stages of Calanus examined consumed cells > 5 μ m, especially cells > 10 μ m. "Fat" stage CVs did not have a resolvable feeding signal, presumably because they were about to enter diapause and had ceased feeding. In contrast, "thin" CVs consumed the phytoplankton prey resolved by the flow cytometer at rates expected for that life history stage. The results will ultimately be interpreted in the context of growth, production, and lipid deposition rates measured by other GLLOBEC investigators.

Table E-1. Calanus finmarchicus ingestion experiments.

Date	Location	Depth	Life history stage	Number copeods/ bottle
5-29-94	Stratified Station 21	Middle mixed layer	Female	1
5-29-94	Stratified Station 21	Middle mixed layer	C5 "fat"	2
5-29-94	Stratified Station 21	Middle mixed layer	C4	3
5-31-94	Mixed Station 22	Middle mixed layer	Female	1
5-31-94	Mixed Station 22	Middle Mixed Layer	C5 "fat"	2
5-31-94	Mixed Station 22	Middle mixed layer	C5 "thin"	2
6-07-94	Mixed Station 42	Middle mixed layer	Female	1
6-07-94	Mixed Station 42	Middle mixed layer	C5 "thin"	2
6-10-94	Stratified Station 43	Chl max	Female	1
6-10-94	Stratified Station 43	Middle mixed layer	Female	1
6-11-94	Stratified Station 43	Chl max	C5 "thin"	2
6-11-94	Stratified Station 43	Middle mixed layer	C5 "thin"	2

F. Zooplankton Process Studies: Copepod Population Processes (Charles Miller and Cheryl Morgan)

Our interest in this expedition is to test methods for study of *Calanus finmarchicus* and *Pseudocalanus* spp. over Georges Bank. Our goals included:

- (1) Extensive preserved collections of plankton from many sites over and adjacent to Georges Bank for analysis of stage composition. The goal is to determine the field development rates implied by advance of the stock through copepodite stages. The collections are also intended for examination of the morphological correlates of life history transitions. In particular, we intend to examine tooth and gonad development of individuals approaching both maturation and entry to diapause.
- (2) Collection and storage of *Calanus* and *Pseudocalanus* of various sizes and stages for analysis of storage lipid content and composition. This is coupled with video imaging of small groups of individuals in order to compare projection estimates of storage oil content with analytical results. Live vs. preserved size comparisons will also be derived from the video files.
- (3) A general reconnaissance of the planktology of Georges Bank and surrounding waters for late May-early June of 1994.

All of these goals were accomplished well beyond our likely ability to deal with all of the resulting samples by the beginning of next year's Broad Scale Survey cruises. Sampling was mostly with the URI MOCNESS system, which worked well throughout, largely thanks to careful maintenance by David Nelson. We expect a completed cast series of 50 tows, most of them to three depths, some to four, two to eight. The total product is in excess of 200 samples. Sorting for lipid samples frozen in liquid nitrogen produced almost 400 small groups of matched individuals, all with associated video images. These include late life stages of Calanus, Pseudocalanus, Centropages hamatus, Centropages typicus, and Metridia lucens. Detailed analysis will be done ashore. We will need to study this large collection selectively.

Most of the Calanus finmarchicus population was in the fifth copepodite stage (C5) when we arrived in the western Gulf of Maine on 25 May. There were still modest numbers of C4. There were very few adults. As the cruise progressed, shallow samples in deeper areas showed fewer and fewer C4 and many more adults. Deep samples mostly contained C5 obviously in diapause (see below). In the last few days of the cruise we began to see Calanus in late naupliar and C1-C2 stages. The composition of the Calanus stock over the Bank proper was much the same, but its numbers were very low. Sampling over the Bank throughout the cruise shows that the plankton of the shallower, well-mixed areas are not dominated by Calanus. There are more Calanus over the northeast peak, but basically the Bank is not Calanus habitat. The dominant copepod over Georges Bank is Centropages hamatus mixed with substantial numbers

of both *Pseudocalanus moultoni* and *Pseudocalanus newmani*. There were also some *Temora* sp. With practice we were able to distinguish the *Pseudocalanus* species without difficulty, but only in the adult and C5 stages. Both *Centropages* and *Pseudocalanus* were reproductively active (egg bearing, young of all stages present) at all locations on the Bank. It seems unlikely that any cohort structure will exist in this season by which to trace population progress in these species. It is my opinion that despite earlier characterizations of winter plankton over the Bank, we will find relatively few *Calanus* in the well-mixed areas and early starts of *Centropages* and *Pseudocalanus*. Population processes leading to the present composition remain to be revealed by the GLOBEC sampling of next winter and beyond.

Sampling over Georges Bank consistently produced large volumes of hydroids, possibly of the species referred to as *Clytia cylindrica* by H. B. Bigelow. These small polyps on branched stalks are clearly, as Bigelow pointed out, not simply live flotsam ripped from the benthos. Their stalk ends are all healed, usually with formation of polyps at all free points. We watched these polyps eat *Calanus* eggs, whole *Pseudocalanus*, *Sagitta* end on, worms, etc. The polyps are nearly neutrally buoyant and are dispersed through the whole water column. The MOCNESS revealed no particular concentration of them at any level. It was sampling well, since *Centropages* were mostly (not totally) confined to the upper 15 m and abundant pelagic larvae of some polychaete worms were confined to the bottom stratum. *Sagitta elegans* was also very abundant over the Bank, occasionally making a plankton sample look like a pot of saimin.

The distinctiveness of the plankton of the well-mixed region over the Bank was the strongest lesson for me as a West Coast oceanographer. You can read forever about the current loop around Georges Bank isolating a special habitat without quite getting the idea. The intensity of the distinction between the plankton over the Bank and that in the apparently seamlessly adjacent waters is astounding.

The sequence of events in the *Calanus* stock during Cl9407 leads to an hypothesis about at least one aspect of its life history. During Leg B we have seen progressively larger numbers of adult males and females. Experiments by Jeff Runge and Stephane Plourde showed that these were spawning very actively indeed. We have also seen the near disappearance of C4 and substantial accumulations near the bottom over deeper sections of the bank and offshore of C5 clearly in diapause (massive lipid storage, no gonadal development, the motionless hanging posture of resting *Calanus*). There are few or no C4 in this resting group.

The hypothesis expands slightly on the long recognized fact that C. finmarchicus has two stratgies in June. One strategy is to enter diapause immediately, a strategy undertaken as C5. We have added a visit to one of the basins of the Gulf of Maine to check whether the early accumulation of diapausing individuals is all or nearly all C5. That station remains to sample at this writing, but the prediction is that it will be nearly all C5. The other strategy is to mature at about year day 150, reproduce, develop rapidly

in warmed spring waters and enter diapause as C4, probably in early to mid-July. This would account for the recurringly large fraction (30 to 50%) of C4 in the October resting stock in the Gulf of Maine. We will somehow obtain autumn 1994 samples from the Gulf for comparison.

The exact nature of the immediately diapausing and maturing substocks remains obscure. They could be genotypes (or equivalently distinct groups) competing for fractional representation with outcomes depending upon relative survival over the year as whole. The variability of outcomes among years would insure the continuance of both types. Alternatively, the diapause vs. mature decision might depend in some way upon individual experience. This agrees with experimental results (Miller, unpublished) from the Gulf of Maine where all spring and summer collections of C3 maintained in the laboratory matured regardless of photoperiod and temperature treatment. There was no sign in that work of an obligate diapause.

Performance of Dian Gifford as chief scientist on Cl9407 was excellent. She kept good order among competing operations, marshalled people to do the general work without needing a whip, and still had energy to do her own science. She designed smart, effective watch schedules of different kinds for different types of operation. She even managed to make the transitions painless. The Iselin and crew performed well in all respects, although the crew were rather cold at the outset. That was inevitable given the high impact loading of the ship on 24 May. Over twenty people and an astounding mass of gear went aboard and were assembled in eight hours. It's understandably hard to be properly welcoming in the midst of swarming new faces, not all of them sailing. Still, it took substantial work to warm the crew up to having interlopers on their ship, but I think we have slowly done it. A very well spent port stop in Woods Hole helped a lot. It also helps that most of the science team are obviously old hands and know their stuff. Those that aren't have been taken in hand and shown the ropes. The crew can see they are working with professionals of their own caliber, and they've grown appreciative. The ship itself is not particularly seakindly, forcing us to shut down work in very modest seas (State 3+). Fortunately, we have had plenty of excellent weather and only two blows that stopped our work for about a day each.

G. Zooplankton Process Studies: Larval Cod Feeding and Growth (Scott Gallager and Ione Hunt Von Herbing)

The objectives of this study were to examine grazing by newly hatched cod larvae on natural assemblages of microzooplankton and determine growth and survival rates of larvae fed assemblages collected at two different depths, thermocline and surface. The hypothesis was the following: the microzooplankton assemblage will be both quantitatively and qualitatively different between the these locations and thus will influence larval cod grazing and growth differentially.

To test this hypothesis, we conducted numerous experiments on ship board to determine grazing rates and growth and survival of cod larvae exposed to water collected nondestructively from various locations in the water column. Cod embryos were spawned in the laboratory weeks ahead of the cruise and incubated between 2 and 10 °C. Embryonic development was timed so that larvae would be hatching throughout the 25 day cruise period. In addition, a second batch of embryos were to be flown down from Newfoundland (Dr. Joe Brown, Memorial University) and placed on the ship during the layover in Woods Hole on June 6 for experiments during the second leg of the cruise. During the first leg, larvae hatched from four batches each containing about 5,000 to 10,000 embryos. The largest batch was due to hatch about June 4 and would have supplied larvae for two to three more experiments. Unfortunately, all embryos died at the developmental stage of late primitive streak/early tail bud stage. Presumably, this was due to overcrowding and/or lack of oxygen, although no definite reason can be given. The batch of eggs due to arrive in Woods Hole for pick- up on the second leg did not arrive, due to miscomunications with Memorial University confounded by a US and Canadian holiday. Therefore, we were not able to conduct further experiments with new larvae on the second leg, although we did complete a number of experiments initiated on the first leg during the latter half of the cruise.

Long-term Experiments. Two long-term growth and survival experiments were established using larvae hatching on a particular day: one on May 29 with water collected in the region of the stratified drifter (Station 21, LT1) and another on May 31 in the area of the mixed site drifter (Station 22, LT2). LT1 was established with 200 larvae per 12 liter tank. Each tank received water from either the thermocline or surface which was passed through 333 μ m mesh to remove large organisms. There were six treatments, one of each of the following from both the thermocline and surface:

- 1. Natural: all organisms smaller than 333 μ m;
- 2. Large: organisms between 333 and 75 μ m;
- 3. Small: organisms less than 75 μ m.

A second growth experiment was established at Station 22 where samples were taken only at the surface. Since there was some evidence that sampling through the orifice of a Niskin bottle disrupted fragile forms, all samples were siphoned from the top of the Niskin or collected at the surface with a bucket, as indicated below. Buckets supplied all water for LT2 and size fractionation experiments.

Interpretation of the results of these experiments awaits comparison with data from the natural zooplankton assemblages collected simultaneously and fixed in Lugols. In general, yolk sac absorption was least, and growth and survival were greatest in treatments from the mixed water column. However, prey concentrations (microzooplankton and copepod nauplii) were very low at both stations tested, so it is

difficult to establish a solid relationship between prey availability and growth.

Short-Term Grazing Experiments. Throughout the two long-term experiments described above and at approximately daily intervals, subsamples were removed from each treatment and exposed to six prey treatments for a period of 3 hours:

- 1. Natural: <333 μ m (fractions between 333 μ m and 75 μ m were separated and stained with calcien blue (CB) while the <75 μ m fraction was stained with acridine orange (AO); the small and large fractions were then combined);
- 2. Large: 75-333 µm fraction stained with calcien blue;
- 3. Small: $<75 \mu m$ stained with acridine orange.
- 4. Natural + Enhanced: natural treatment plus 0.5/ml stained Balanion and 0.05/ml stained nauplii (Pseudodiomtomus sp.)
- 5. Large + Enhanced: Large fraction plus nauplii
- 6. Small+Enhanced: Small fraction plus Balanion

After grazing for three hours in an illuminated incubator, larvae were removed, mounted on slides and examined under epifluorescence microscopy using either blue or UV excitation for AO or CB, respectively. Fluorescent images of larval guts were captured and stored digitally for quantification of gut fluorescence post cruise. Standard morphological measurements were also made on the stored image (length, height, yolk sac area, myotomal height, eye diameter, etc).

Other studies. In addition to the scheduled grazing experiments outlined above, several ideas concerning the grazing process were also tested. Ship vibrations were though to interfere with feeding by larvae. Therefore we conducted an experiment in which the incubation vessels were suspended in the incubatior using bungle cord. Results suggest that, indeed, larval feeding may be reduced by the ship's vibrations. Future experiment will have to take this into account.

Results of the grazing experiments on various size fractions showed that newly hatched cod larvae feed directly on natural assemblages of microplankton, including protozoans. No copepod nauplii were ingested before day 5 following hatching. Compared with larvae fed enhanced levels of the aloricate ciliate Balanion, however, rates were low, suggesting prey concentration was limiting. A major conclusion from these experiments is that larval growth and survival in the water column at this time of the year would be very poor compared with the prey density we would expect to see earlier in the year when cod and haddock larvae are present. This supports the match-mismatch hypothesis of larval timing in the water column coinciding with maximal prey levels.

Prey Motility Experiments. Fourteen hours of video information were recorded on motility patterns and particle size composition in the water column at various depths following collection both invasively (Niskin port sample) and non-invasively (bucket from the surface or siphon from Niskin bottle). Prey motility patterns will be digitized and calculated with our Motion Analysis System back at Woods Hole and compared with larval capture success from the natural assemblages.

Ancillary Observations. Many hours of recordings were also made of the diatom balls (*Chaetoceros* sp.) collected at various sites and depths where the CTD and VPR fluorometer pegged. Macro views showed regularly organized structures of varying spherical morphologies. High mag (630x) showed chains with spines meshed into a matrix of fibers. Few nano- or picoplankton were associated with these colonies. Further interpretation awaits SEM and culture experiments at the lab.

General Comments. Cl9407 was a great success both in terms of data collection and learning valuable lessons of culturing cod larvae and their prey at sea. We thank the whole crew of the *Columbus Iselin* for providing a pleasant and efficient working environment.

H. Video Plankton Recorder Studies (Cabell Davis and Scott Gallager)

The goal of the VPR sampling was to determine the fine-scale (centimeters to meters) vertical and horizontal distribution of the planktonic community in relation to hydrography in the vicinity of the stratified and well-mixed drifter sites. The main purpose was to measure the degree to which zooplankton are aggregated at the pycnocline. A second purpose was to sample the 3-dimensional distribution of the plankton community in within a 1-km radius of the drifter. The latter sampling will enable us to determine the extent to which "single-point-sampling" (eg. a single double oblique VPR haul, net haul, CTD cast, or pump deployment) at the drifter itself is representative of the the local (1-km) pelagic environment.

Twenty VPR towyos were made during the cruise. After our initial test deployments at the north edge of the bank (VPR 1) and in the Northeast channel on May 28 (VPR 2), we conducted a series of day/night grids at the first drifter site. The first (VPR 3) consisted of 3 parallel transects each about 4 km long. The remaining grids (VPR 4 to 9) consisted of left-handed square inward spirals having an outside dimension of about 2 km (actually 1 nm). All grids were centered on the drifter and thus had a moving coordinate frame. The left-handed grids were used so that the ship could always turn to port, keeping the wire away from the ship (the VPR was towed from the main crane off the port side). The strobe on the VPR malfunctioned at the end of the first leg, and, during the remaining time, we analyzed the video and were able to create 3-D maps of several planktonic groups from the last grid VPR-9. The strobe was repaired in port on May 6 and we obtained a backup strobe as well.

On the second leg of the cruise we conducted 3 grids while following the mixed-area drifter (VPR 11-13) and two while following the stratified drifter (VPR 14-15). We also conducted several short transects (VPR 16-19) in the stratified area and one very long (approx. 200 km and 26 hrs) transect (VPR 20) across Georges Bank from the Slope Water to the mixed area and into the Gulf of Maine (Georges Basin). Prior to the first grid, the VPR winch hydraulic motor failed and the ships hydraulic system was used to run the VPR winch. This changeover actually proved quite beneficial to us, as we were then able to run the winch remotely, in the comfort of the ship's electronics lab. The strobe again failed during VPR 17 and was replaced by the backup which functioned properly for the remainder of the cruise.

We found the crew of the Iselin to be friendly, interested in the science, and helpful. In particular, the engineers (Bob, Jack, and Ralph) helped greatly in connecting the VPR winch to the ship's hydraulic system. The bridge (Mike, Chris, and Rhett) were also very helpful and flexible and were adept at following the drifter along the grid track. The boatswain Steve on the first leg was superb; he was willing to listen, was easy to work with, and developed an efficient method for safe deployment and recovery of the VPR. Other crew were also helpful and competent including Erik, Bob, and John. The cooks Susan and Tori were superb and extremely hard working, and the good food helped keep everyone happy. Chip helped us with electronic hookups and e-lab instrumentation and Dave Nelson was also helpful in loaning us parts (conductivity cell, pinger, and reed-switch).

In all, the cruise was highly successful for us in terms of data acquired and will provide us with unique new insights into the fine-scale distributions of the planktonic community in relation to hydrographic features and circulation. We believe the problem with the VPR strobe was due to adverse effects of bulb failure on the electronics. As expected, the problems encountered during this pilot cruise will enable us to be well-prepared for the full GLOBEC study in 1995.

Acoustic Studies (TAPS) (Mark Berman and Joseph Kane)

The purpose of our research was to further develop techniques for assessing the spatial and temporal distribution of the copepod community on Georges Bank using the Tracor Acoustic Profiling System. Two types of deployment were used, both with TAPS mounted on the CTD frame. The first technique was for quickly surveying the vertical distribution of the copepods. In this mode TAPS collected data during a normal CTD cast, integrating data over 3 m vertical bins. Twenty-five TAPS deployments were made in this mode, during the transects and at the 48 hr stations.

The second mode was used only at the long stations, while the ship kept station with the drifter. In this mode the CTD and TAPS were yoyoed up and down through the top 50 m (water depth permitting) at 10 m/min continuously for 1 hour. This allowed TAPS to

assess the size distribution and concentration of the copepods with a 1 m vertical resolution to characterize the size, position, and frequency of plankton patches. During the latter part of the cruise, the modes were combined to minimize sampling time. For this the TAPS/CTD was yoyoed slowly for 50 min, with the last 10 minutes of the hour going to a normal CTD cast. A total of 11 yoyo casts were completed during this cruise.

A hardware problem within TAPS was discovered during the latter portion of Leg A. It is not yet clear how much of the data collected can be corrected through post-processing. However, the deployment techniques developed here appear to be satisfactory, and will be used on upcoming GLOBEC stratification studies

J. Airborne Lidar Fluorescence Measurements (Charles Primmerman)

On the mornings of 3,8, and 10 June 1994, personnel from the Massachusetts Institute of Technology's Lincoln Laboratory conducted airborne surveys over Georges Bank. These surveys consisted of airborne-lidar fluorescense monitoring of phytoplankton population distribution and abundance. In addition, on the 10 June survey, sea-surface temperature was monitored using an airborne infrared radiometer. The primary objectives of these flights were to provide a 'snapshot' of phytoplankton distribution over a broad region of Georges Bank and to determine the level of 'patchiness' in the phytoplankton distribution. Excellent data were collected on all three flights, and the flight objectives appear to have been accomplished.

The airborne-lidar measurements were coordinated with GLOBEC cruise Cl9407 on the research vessel *Columbus Iselin*. The airborne-lidar measurements were not directly funded by the GLOBEC program, but it is expected that the results will be useful to GLOBEC. Conversely, data collected by the *Iselin* are expected to be useful as comparison points for the airborne measurements.

The MIT Lincoln Laboratory airborne-monitoring system is described below, and the important system parameters are listed. The flights paths and data associated with them are indicated in terms of their latitude, longitude, and GMT times. Pictorial representations of these flight paths are also provided. Finally, an overview of fluorescence monitoring results is presented, and some selected IR data are provided.

Airborne monitoring system. MIT Lincoln Laboratory's fluorescence monitoring system was mounted on a Gulfstream G-1/59 flown out of the MIT Lincoln Laboratory Flight Facility at Hanscom Field in Bedford, Massachusetts. The aircraft was equipped with a Global Positioning System (GPS) receiver and a LORAN system to assist in course heading determination. During the Georges Bank monitoring missions, the aircraft was flown at a speed of approximately 140 kts, at an altitude of approximately 300 ft. Chlorophyll fluorescence was excited using a 20-mJ doubled Nd:YAG laser operating at a wavelength of 532 nm. The laser-beam transmitter produced an 80-cm spot on the

ocean's surface. The laser was operated at a 20-Hz repetition rate so that the horizontal separation between samples was approximately 3.5 meters. Fluorescence radiation in a 30-nm band centered at 685 nm, water Raman radiation at 647 nm, and laser backscatter at 532 nm were collected by an 8-inch Celestron telescope and reimaged onto three avalanche photodiodes through the appropriate filters. Detector output was digitized in 2 nsec increments, to give depth-resolved information, and recorded on a PC-based data acquisition system. A separate 5-cm aperture was used to collect infrared radiation to determine sea-surface temperature. Radiation in the 3-5 micron band was imaged through a germanium filter onto a liquid-nitrogen-cooled indium-antimonide detector. Detector input was chopped at 300 Hz, and its output sampled using a lock-in amplifier. The amplifier's signal was digitized and recorded along with LIDAR data. Sunlight levels were monitored and recorded using a silicon-PIN detector mounted on top of the aircraft. Some important system parameters and hardware specifications are listed in Table J-1.

Flight profiles. The flight paths for 3, 8, and 10 June overlaid on the 50-fathom isobath at Georges Bank are shown in Figures J-1 to J-3. For each mission, the approximate flight path of the aircraft is indicated by the line overlayed on a chart of Georges Bank. The fluorescence data recorded along the indicated flight paths are presented as a normalized ratio, computed by dividing the depth-integrated fluorescence signal by the water-Raman signal. The color bar on the right of each plot relates this ratio to a color. These particular paths were chosen based upon discussions with members of the Woods Hole GLOBEC cruise team. Of particular interest were the northeastern region of the Bank, and the Bank's southern edge, as defined by the 50-fathom isobath.

3 June Mission:

Take-off from Hanscom Field was at 0730 hours (EDT). The airplane was re-fueled at Nantucket airport and departed for the northeastern region of Georges Bank at approximately 0900 hrs. The conditions at the Bank were clear and sunny. The first descent to 300 ft. was made at 41 degrees 17.6 minutes latitude, 67 degrees 35.5 minutes longitude, at time 13:40:23 (GMT). A 300 ft. altitude was maintained for the next 4 hours, and the flight path indicated in Figure J-1 was flown. This flight path brought the airplane over the *Iselin*'s location at approximately 14:24 (GMT), at coordinates 41 degrees 3.1 minutes latitude, 66 degrees 36.8 minutes longitude. The flight path was completed at approximately 17:59 (GMT), and the airplane began the return trip to Hanscom Field.

8 June Mission:

The *IseIin* was contacted by INMARSAT telephone at 0730 hours (EDT), and its location and cruise plans for the day were provided to us. A flight plan to cover the southern edge of the Bank and overlap with the ship was filed before departing Hanscom field at approximately 0814 hours (EDT). The flight path for the 8 June mission is shown in Figure J-2. Descent to 300 ft. altitude was made at time 13:27 (GMT) at location 41 degrees 51.4 minutes latitude, 66 degrees 32.5 minutes longitude. Data collection began

at this time. The flight path intersected the ship's location at approximately 14:28 (GMT), at location 41 degrees 17 minutes latitude, 67 degrees 19.8 minutes longitude. Data collection continued along the indicated flight path until 17:13:37 (GMT), at which time the airplane began the return trip to Hanscom Field.

10 June Mission:

The airplane departed Hanscom Field at approximately 0830 hours (EDT) on route to the planned starting location in the northeast region of the Bank. Descent to 300 ft. altitude was made at location 42 degrees 11.0 minutes latitude, 66 degrees 54.8 minutes longitude, at time 13:36 (GMT). Data collection was begun, and continued along the flight path shown in Figure J-3. This flight path took the airplane within a few kilometers of the *Iselin*'s location at approximately 16:50 (GMT) at location 40 degrees 52 minutes latitude, and 67 degrees 30 minutes longitude. The IR sensor collected data along with the lidar during this flight. Near the end of the data collection, a test was made using the IR sensor without the laser propagating. No change in the IR data stream was noted. Data collection was completed at approximately 17:14 (GMT) at location 40 degrees 44.8 minutes latitude, 68 degrees 30.3 minutes longitude.

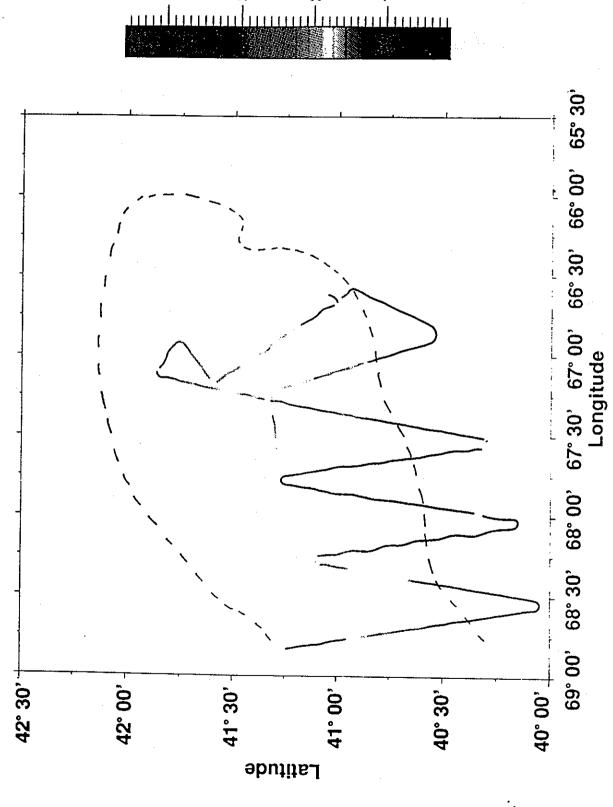
Overview of Results. In addition to the chlorophyll fluorescence data provided in Figures 1-3, an example of depth-integrated fluorescence (normalized to the water Raman signal) is plotted versus distance in Figure J-4. These particular data were collected during the 8 June mission, along a 10 mile stretch over the southeast region of the Bank. The starting and stopping coordinates are given in the figure. The data shown in Figure 4 are representative of the level of structure and spatial scales present in the phytoplankton population distribution for all three missions. Similar spatial scales are seen in data from the infrared radiometer. A sample is presented in Figure J-5. Preliminary analysis of the depth-resolved lidar returns suggests that under certain conditions sufficient signal-to-noise is present to permit measurements of chlorophyll concentration to a depth of 10 meters or more.

Table J-1. COMPONENT PARAMETERS.

AIRFRAME	
Model	Gulfstream G-1/59
Airspeed	140 kts (approximately)
Altitude	300 ft.
Navigation System	GPS and LORAN
LASER AND TRANSMITTER GEOMETRY	
Configuration	x2 Nd:YAG, diode pumped
Wavelength	532 nm
Energy	20 mJ
Pulse Length	10 ns
Repetition Rate	20 Hz
Fluence (at ocean surface)	4 mJ/cm2
Spot Size	80-cm Diameter
Spot-to-Spot Separation	3.5 meters (approximately)
RECEIVER	
Contiguration	Cassegrain. f/11
Apenure Diameter	8 inches
Imaging to Detectors	173
SEA-SURFACE TEMPERATURE SENSOR	
Туре	InSb (at 77 K)
Spectral Band	3 - 4.5 microns (approximately)
Field-of-View	2 degrees
Sampling Bandwidth	1 Hz
Spatial Resolution	3 x 70 meters
Noise-equivalent Temperature	0.5 Kelvin
DETECTORS	
Type	Avalanche photodiodes
Configuration:	532 nm (backscatter)
	647 nm (water Raman)
	685 ± 15 nm (fluorescence)
DIGITIZING AND DATA RECORDING	
Contiguration	4-channel, EISA intertace
Sample Period	2 ns
(All data is IRIG time-stamped, and recorded	
with GPS Coordinates.)	



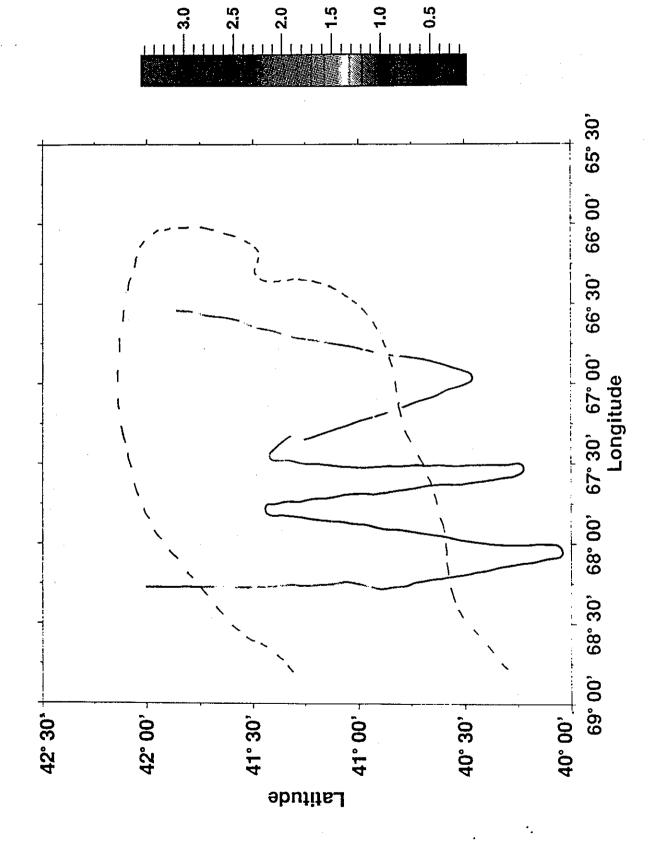




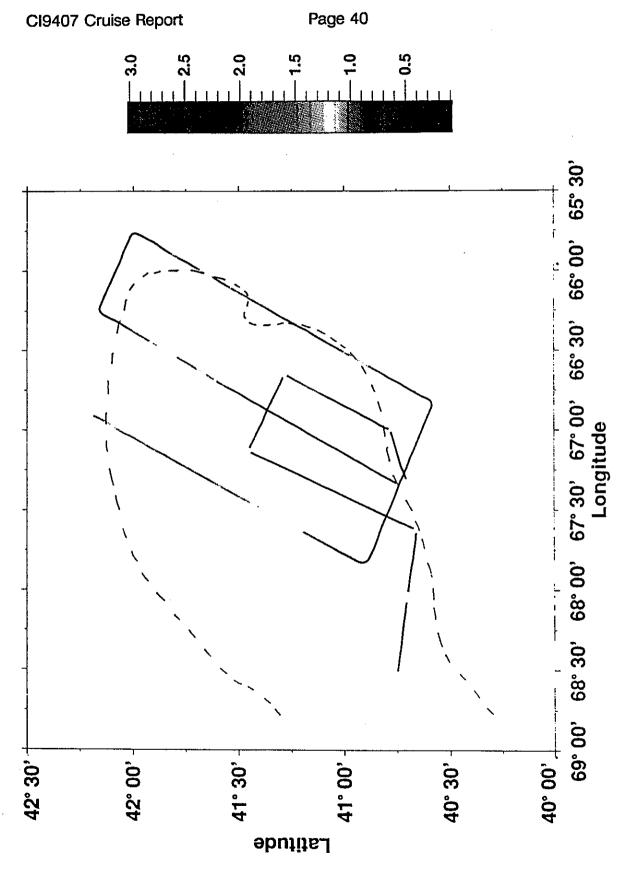
Flight path for June 3, superimposed on 50-fathom isobath of Georges Bank. (Scale indicates normalized fluorescence returns). Figure J-1.



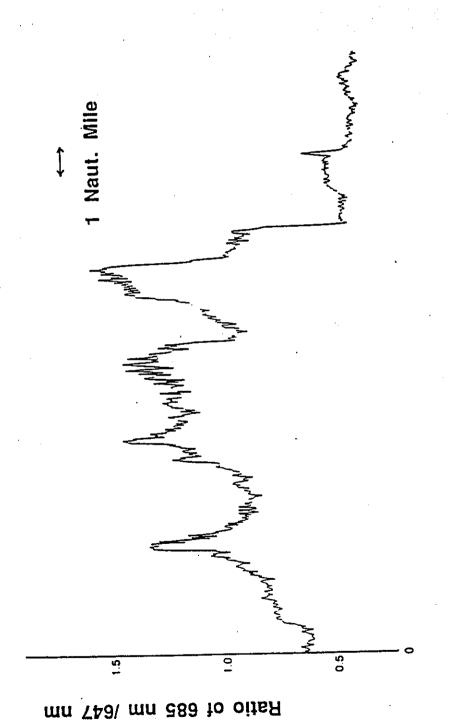




Flight path for June 8, superimposed on 50-fathom isobath of Georges bank. (Scale indicates normalized fluorescence returns). Figure J-2.

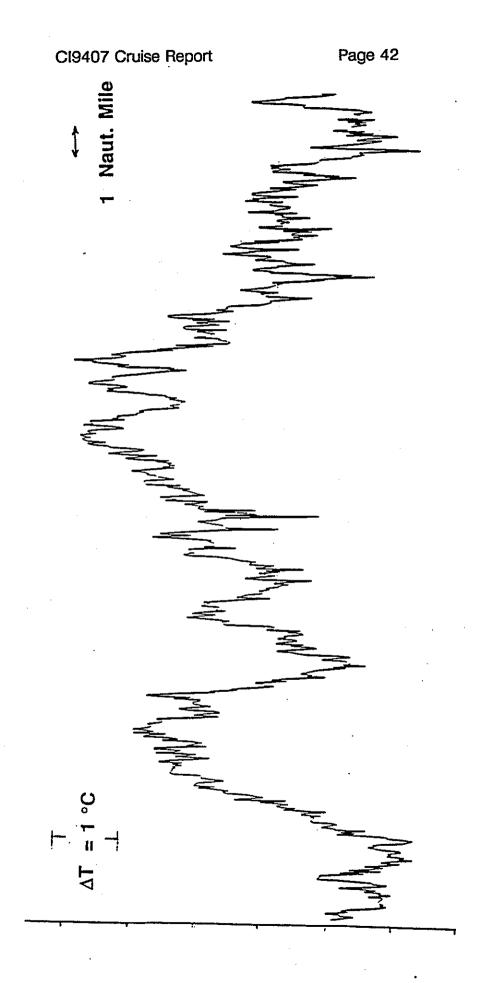


Flight path for June 10, superimposed on 50-fathom isobath of Georges bank. (Scale indicates normalized fluorescence returns). Figure J-3.



Distance along track (from 41° 2', 66° 46' to 40° 40', 66° 51'

Figure J-4. Integrated LIDAR returns (June 8, File 1349).



Distance along track (from 41° 12', 67° 38' to 40° 49', 67° 38'

Figure J-5. IR radiometer signals (June 10, File 1402).

Cl9407 Cruise Report

Page 43

Scientific Personnel Leg A

Name	•	Title	Organization
1.	Dian Gifford	Chief Scientist	University of Rhode Island
2.	Robert Campbell	Post-doctoral Fellow	University of Rhode Island
3.	Maria Bemis	Scientist	University of Rhode Island
4.	Peter Garrahan	Scientist	University of Rhode Island
5.	Richard Bohrer	Scientist	University of Rhode Island
6.	Jonathan Hopkins	Marine Technician	University of Rhode Island
7 .	David Nelson	Marine Technician	University of Rhode Island
8.	Charles Miller	Scientist	Oregon State University
9.	Cheryl Morgan	Scientist	Oregon State University
10.	Ari Epstein	Student	Woods Hole Oceanographic Institution
11.	Michael Sieracki	Scientist	Bigelow Laboratory for Ocean Science
12.	Terrance Cucci	Scientist	Bigelow Laboratory for Ocean Science
13.	Scott Gailager	Scientist	Woods Hole Oceanographic Institution
14.	Ione Hunt von Herbing	Post-doctoral Fellow	Woods Hole Oceanographic Institution
15.	Linda Davis	Scientist	Woods Hole Oceanographic Institution
16.	Cabell Davis	Scientist	Woods Hole Oceanographic Institution
17.	Martin Marra	Scientist	Woods Hole Oceanographic Institution
18.	Frederika Norrbin	Scientist	Woods Hole Oceanographic Institution
19.	Jeffrey Runge	Scientist	Institute Maurice Lamontaigne
20.	Stephane Plourde	Scientist	Institute Maurice Lamontaigne
21.	Mark Berman	Scientist	National Marine Fisheries Service
22.	Joseph Kane	Scientist	National Marine Fisheries Service
23.	Chip Maxwell	Marine Technician	University of Miami
24.	Donald Cucchiara	Marine Technician	University of Miami

RV Columbus Iselin Officers and Crew Leg A

Name	•	Title
1.	Mike Dick	Captain
2.	Chris Vogel	Chief Mate
3.	Rhett McMunn	Second Mate
4.	Steve Vetra	Bosum
5.	Jack Crawford	Chief Engineer
6.	Bob Lapsley	First Engineer
7.	Ralph Harvey	Second Engineer
8.	John Cawley	Seaman
9.	Erik Hutchinson	Seaman
10.	Bob Loos	Seaman
11.	Susan Rafferty	Steward
12.	Torii Young	Cook

Scientific Personnel Leg B

Name	•	Title	Organization
1.	Dian Gifford	Chief Scientist	University of Rhode Island
2.	Robert Campbell	Post-doctoral Fellow	University of Rhode Island
3.	Maria Bemis	Scientist	University of Rhode Island
4.	Peter Garrahan	Scientist	University of Rhode Island
5.	Richard Bohrer	Scientist	University of Rhode Island
6.	David Avery	Scientist	University of Rhode Island
7 .	David Nelson	Marine Technician	University of Rhode Island
8.	Charles Miller	Scientist	Oregon State University
9.	Cheryl Morgan	Scientist	Oregon State University
10.	Edward Dever	Student	Woods Hole Oceanographic Institution
11.	Michael Sieracki	Scientist	Bigelow Laboratory for Ocean Science
12.	Terrance Cucci	Scientist	Bigelow Laboratory for Ocean Science
13.	Scott Gallager	Scientist	Woods Hole Oceanographic Institution
14.	Andrew Girard	Student	Woods Hole Oceanographic Institution
15.	Lucie Blanchard	Student	Woods Hole Oceanographic Institution
16.	Cabell Davis	Scientist	Woods Hole Oceanographic Institution
17.	Philip Alatalo	Scientist	Woods Hole Oceanographic Institution
18.	Stephane Plourde	Scientist	Institute Maurice Lamontaigne
19.	Chip Maxwell	Marine Technician	University of Miami
20.	Donald Cucchiara	Marine Technician	University of Miami

RV Columbus Iselin Officers and Crew Leg B

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APPPENDIX 1: CI4907 EVENT LOG

INSTRUMENT ABBREVIATIONS:

CTD: CTD: Conductivity/tempersture/depth sensing package DFT: ARGOS drifter

MOC: MOCNESS: Multiple opening-closing net and environmental sensing system

PPN: Phytoplankton net

TAP: Tracor acoustic profiling system

VPR: Videoplankton recorder

ZPN: Zooplankton net

ZPP: Zooplankton pump

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	マラマ		Gifford/Durbln		Miller/Durbin		Runge		Davis		Gifford/Durbin		Miller/Durbin			Gifford/Durbin	Glfford/Durbin	Gifford/Durbin Miller/Durbin	Glfford/Durbin Miller/Durbin	Gifford/Durbin Miller/Durbin Gifford/Durbin	Glfford/Durbin Miller/Durbin Glfford/Durbin	Gifford/Durbin Miller/Durbin Gifford/Durbin Miller/Durbin	Gifford/Durbin Miller/Durbin Gifford/Durbin Miller/Durbin	Gifford/Durbin Miller/Durbin Gifford/Durbin Miller/Durbin Gifford, Durbin	Gifford/Durbin Miller/Durbin Gifford/Durbin Miller/Durbin Gifford, Durbin Funge	Gifford/Durbin Miller/Durbin Gifford/Durbin Miller/Durbin Gifford, Durbin Runge	Gifford/Durbin Miller/Durbin Gifford/Durbin Miller/Durbin Gifford, Durbin Hunge Durbin	Gifford/Durbin Gifford/Durbin Miller/Durbin Gifford, Durbin Aunge Durbin Gifford/Durbin
1	ppp	CCCC	0000	0235	0000	0249	0000	0150	NA	NA	0000	0800	8		0085								╶╶┪╼╃╼╃┈╬═╬╌╬╼┾═╏		▗ ▗ ▊▃▋▃▋┈┋▃┞▃╎╶╏╼┠╾╎╶┿╌╸	╶╶┩┈┩┈┩┈╄┈	╒╒╃╒╃╒╃╒╃╒╃╒╇╒╇	╶╶┩┈┩┈┩┈┩┈┩┈╎┈╏┈╏┈╏┈╏┈╏┈ ╏
WatDe	adda	adad bbbb	0250		0259	0263	0347		0458		0094		900			0079												
Longitude WatDe insDep	DOWNER	domin III	6541.93	6541.22	6541.27	6541.32	6541.12	6540.52	6540.14	6548.96	6603.50	6603.91	6603.89	-	6603.92			- 	- 1111	- 1111	- - - - - - - - - - 	- - - - - - - - - - 	- - - - - - - - - - 	╸┤╍┥╸┥┉┈╎╶╏╶┫╼╶╎┈ ┪┯╌╴		▔▐▗▔▐▗▔▋ ┉ ▊ ▗▐ ▗▊▄▐▗▐┉ ▀	- - - - - - - - - - 	▔▗▗▔▗▗ ▃▃
Locitine Latitude	J-IMMOO	ddmm.ff	4157.36	4157.73	4157.90	4158.67	4158.62	4158.91	4158.42	4155.72	4151.81	4151.26	4151.34		4151.47	4151.47	4151.47 4146.47 4146.81	4151.47 4146.47 4146.81 4146.69	4146.47 4146.81 4146.69 4147.05	4151.47 4146.47 4146.81 4146.69 4147.05	4146.47 4146.81 4146.89 4147.05 4142.21 4142.62	_ [
Loctime	SSMALL	hhrimss	140200	142600	143800	151700	153400	155300	162400	192100	202830	204600	210656		212200	212200 241540	212200 241540 243100	212200 241540 243100 234200	241540 241540 243100 234200 235900	241540 241540 243100 234200 235900 012211	212200 241540 243100 234200 235900 012211 013800	241540 241540 243100 234200 235900 012211 013800	212200 241540 243100 234200 235900 012211 013800 015000	241540 241540 243100 234200 235900 012211 013800 015000 021000	241540 243100 234200 235900 012211 013800 015000 053900 NA	241540 243100 234200 234200 012211 013800 015000 021000 053900 NA	212200 241540 234200 235900 012211 013800 015000 021000 053900 NA	241540 243100 234200 235900 012211 013800 015000 021000 053900 NA 061300 NA
Locdate	COMMUN	yymmdd	940528		940528		940528		940528		940528		940528			950528	950528	950528 940528	950528	950528 940528 940528	950528 940528 940528	940528 940528 940529	950528 940528 940528	950528 940528 940528 940529	950528 940528 940529 940529	950528 940528 940529 940529	950528 940528 940528 940529 940529	950528 940528 940529 940529 940529
CAVITATE	IDD: HHAMISS: YYMMDD: HHIMMISS DOWNAFF DOMINFF DODO	Minimss	180200	182600	183800	191700	193441	195300	202400	232100	002830	004238	010656		012200	012200	031546 033100	031546 033100 034200	031546 031546 033100 034200 035900	031546 031546 033100 034200 035900 062211	031546 031546 033100 034200 035900 052211 052211	031546 031546 033100 034200 035900 052211 053800	031546 033100 034200 035900 05211 053800 055000	031546 033100 033100 034200 035900 052211 052211 053800 061000	031546 033100 033100 034200 052211 052211 053800 056000 061000	031546 033100 033100 034200 035900 052211 053800 055000 061000 061000 093947	031546 033100 033100 035900 052211 052211 055000 061000 094900 021300 NA	033100 033100 033100 033200 052211 052211 055000 061000 061000 083947 094900 021300 NA
GMTdate	COMMUNA	yyminidd	940528		940528		940528		940528		940529		940529			940529	940529	940529	940529	940529	940529 940529 940529	940529 940529 940529	940529 940529 940529	940529 940529 940529 940529	940529 940529 940529 940529	940529 940529 940529 940529	940529 940529 940529 940529	940529 940529 940529 940529 940529
Instrument Cast# STATION#			CI9407A.017		CI9407A.017		CI9407A.017		CI9407A.017		CI9407A.018		C19407A.018			CI9407A.019	Cl9407A.019	CI9407A.019 CI9407A.019	CI9407A.019 CI9407A.019	CI9407A.019 CI9407A.019 CI9407A.020	CI9407A.019 CI9407A.019 CI9407A.020	CI9407A.019 CI9407A.019 CI9407A.020	CI9407A.019 CI9407A.019 CI9407A.020 CI9407A.020	CI9407A.019 CI9407A.020 CI9407A.020 CI9407A.020	CI9407A.019 CI9407A.020 CI9407A.020 CI9407A.020	CI9407A.019 CI9407A.019 CI9407A.020 CI9407A.021 CI9407A.021	CI9407A.019 CI9407A.020 CI9407A.020 CI9407A.021 CI9407A.021	CI9407A.019 CI9407A.020 CI9407A.020 CI9407A.021 CI9407A.021 CI9407A.021
Cast#	×		910		012		8		802		210		ЭЗЭ			918	918	018	014	019 018	019	018 019 015	019 018	019 018 019 000	019 019 018	002 002	019 019 019 000 000 000	018 019 019 019 000 000 000 000 000 000 000
Instrument	臺		СТБ		Moc		ZPN		VPR		CTD		MOC			CTD	СТО	CTD	СТР	CTD MOC	MOC	MOC CTD MOC	MOC MOC	MOC CTD MOC ZPN	CTD MOC CTD ZPN	CTD MOC CTD ZPN ZPN	CTD MOC MOC ZPN ZPN DFT	CTD MOC CTD ZPN ZPN CTD
EVENT*	689		CI14894.009		CI14894.010		CI14894.011		CI14894.012		Cl149.94.001		CI14994.002	1			C114994.003											

EVENT*	Instrument Cast# STATION#	Cast#		GWTdate	GMTdate GMTdre Loccate Loctine Latitude Longitude Watte Institut	1 ocdate	Loctime	Lattede	Longitude	WatCle	cis()sty		Region	Commen
wooddyy,ees		××	*WCCCCI.888	YYNINIDD	IDD. FIFININSS: YYMMOD: HIMINISS DOMINEF DOMINEF	WININDO	SSMAHH	DOMM FF		pppp caca		NNN	iddd UUU.	⊥1 2003
				yymmdd	infinities:	ymmdd hhmmss		ddmm:#	domini:	place	OCC			
C114994.010	¥0C	910	CI9407A.021	940529	105100	940529	065100	4149.67	6612.78	9081	0000	Miller/Durbin	Stratffed	Clear
					111200		071200	4148.75	6612.93		9008		station	Calm
CI14994.011	ZPP	8	C19407a.021	940529	120000	940529	000080	4146.57	6610.31	0084	0000	Durbin	Stratffed	Clear
					131500		091500	4144.76	6609.70		0057		station	Calm
C114994.012	CTD	021	CI9407A.021	940529	134645	940529	094645	4144.12	6611.72	8800	0000	Gifford/Durbin	Stratified	Collect
					135800		008560	4143.73	6611.89		0020		station	water
CI14994.013	CIO	022	CI9407A.021	940529	143659	940529	103659	4143.58	6612.03	0087	0000	Gifford/Durbin	Stratified	Callect
					144900		104430	4143.01	661244		3900		station	water
CI14994.014	DPP	8	CI9407A.021	940529	151000	940529	111000	4143.48	6614.15	9800	0015	Durbin	Strattled	Callect
					161500		121500	4143.55	6614.87				station	water
C114994.015	CTO	023	CI9407A.021	940529	165056	940529	125056	4144.76	6617.20	9800	0000	Gifford/Durbln	Stratffed	
	TAP	88			170500		130500	4145.07	6619.96		0072	Berman	station	
C114994.016	MOC M	017	C19407A.021	940529	172100	679076	132100	4145.22	6617.34	6800	0000	Miller/Durbin	Strattfled	Clear
				-	173700		133700	4145.38	6617.92		0200		station	Sunny
Cl14994.017	ZPN	903	CI9407A.021	940529	180100	940529	140100	4145.98	6616.73	0800	0000	Runge	Strattfled	Live tow
					182500		142500	NA	NA		0020	Gifford	station	
CI14994.018	ZPN	004	CI9407A.021	940529	183000	940529	143000	4145.98	6616.73	0800	0000	Durbin	Stratified	Live tow
					184200		144200	4146.85	6616.49		0020		station	
CI14994.019	댐	002	CI9407A.021	940529	194600	940529	154600	4146.73	6615.54	0084	0000	Ourbin	Stratified	Retrieve
					NA		NA	NA	NA		NA		station	for repair
C114994.020	DFT	889	CI9407A.021	940529	195300	940529	155300	4147.37	6614.92	0084	0000	Durbin	Stratified	Deploy
					NA		NA	NA	NA		NA		station	drifter
CI14994.021	VPR	88	CI9407A.021	940529	203200	940529	161600	4147.16	6614.58	0084	NA	Davis	Stratffed	
	·	,			NA		193600	4144.54	6607.24		NA		station	
CI14994.022	СТD	024	CI9407A.021	940529	235000	940529	200000	4143.40	6607.05	9093	0000	Gifford/Durbln	Stratified	
	TAP	600		940530	245900		205900	4141.28	6606.07		0020	Вегтал	station	

The same	Strange Land	NOTIATE CASTA STATION	GMTdate	tte CNTfirte	Locdate	Loctime	Lattride Longitude WatUe naUsp	orgitude	P P P P P P P P P P P P P P P P P P P	: 1:		I DIRECT	1
	1) Curways		CICHNEY	1 SSM	DOMNER!	DIMMER I	0000	•••	777	HER PPF CCC	3
wdddy, egs	×	WCCCCI:988	T (MINITELL	T (Na) A (F. H.	The management of the state of	Thrift 94	# must	OCICIO Pippa # while	ppo	aga			
		_	Manimod 040530		940529	213900	4140.42	6607.03	9084		Davis	Stratffed	
CI15094.001 VPR	3	CI840/A.021	340000	040800			4138.61	6609.31		NA		station	
	100	C10407A 034	040530	043000	940530		4139.30	6611.31	0087	-1	Gifford/Durbin	Stratified	
CH 5094,002 CHD	S	CI340/A.021	Parata Parata	044500			4139.56	6611.99		0200			
	\dagger	C104074 021	940530	045800	940530	002800	4139.72	6612.64	991	-+	Miller/Durbin	교	Casi Casi
CI15094.003 MUC	0 0	CISTO COLOR		051300		011300	4139.69	6613.36	2600	-†		_	rigu
-+-	8	C19407A 021	940530	000090	940530	020000	4139.44	6607.71	885	\dashv	Durbin	Stratified	
CI15084.004 2rr	3	1		064500		024500	4139.52	6607.41		8057		station	
	+	C10407A 001	040530	070300	940530		4141.02	6613.59	0082	88	Gifford/Durbin	Stratffled	
C115094.005 CID	8 5	-	2	075200		_	4141.91	6612.41		820	Вегтап	station	
_	+	+	040530	000000	940530	040000	414217	6612.16	0082	NA	Davis	Strattled	
C115094.006 VPH	3	CISAUTAOR	3	110500		071500	4140.01	6606.93		NA NA		station	
	+	CIO407A 094	940530	115300	940530	075300	4139.19	6606.23	8600	800	Gifford/Durbln	Stratffed	
C115094.007 CID		\top		125800		082800	4137.34	6604.44	0032	0020	Berman	station	
_	1	-	040530	134000	940530	091000	4136.51	6604.63	9092	8	Miller/Runge	Strattfled	Live tow
CI15094.008 ZPN	8	CI940/A.UZI	200	20000		092200		6604.28		0020		station	
_	-	十	CCECAC	1	940630	093700	4136.09	6605.20	000	NA	Davis	Stratified	
C115094,009 VPR	8	C1949/A.UZ1	25042			121600	4135.95	6608.56		NA		station	
	1	-	0,1	-	040530	10300		09.8099	600	8	Glfford/Durbln	Stratified	
C115094.010 CTD	-	S CI9407A.021	200	23/20	200	133400			88	0050	Веппал	station	
TAP		\dashv		-	083080	- -	╄	_	600	8	Gifford/Durbln	Stratified	Collect
C115094.011 CTD	88	CI9407A.021	940530	-	2000	-1-	+	6609.35		0200		station	water
	+	-+		1/3400	940530	135300	+-	99.8099	808	0000	Miller/Durbin	Stratified	
C115094.012 MOC	019	GIS40/A.UZI	3	╬	╁╴	141800		6611.24		9093		station	
+	1	十	1	-	940530	+	-	6610.45	868	0000	Durbin	Stratffed	
CI15094.013 ZPP	8	3 CI940/A.UZI	200	4	200	+		6610.28		0057		station	-
_		一十	040530	-	940530	┷		6610.71	9094	0000	Miller/Durbin	Stratfled	
C115094.014 MOC	ည္က	0 CI940/A.021	\$	-}-	T	1	01 74 97 40	6611 00		8080		station	

CI9407 EVENT LOG

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Comment	CC III		Live tow														Clear	Calm	Clear	Calm	Clear	Calm			Live tow	Fog	Uve taw		Deploy	Orffter
Region	dad UEB		Stratified	station	Stratified	station	Stratified	station	Stratffed	station	Strattfied	station	Stratified	station	Stratified	station	Stratffed	station	Strattfled	station	Stratified	station	Stratffed	station	Mbed	station	Mixed	station	Mixed	station
	るされ		Runge		Davis		Giford/Durbin	Вегтап	Glfford/Durbin		Miller/Durbin		Durbin		Davis		Glffard/Durbln		Miller/Durbin		Durbin		Gifford/Durbin		Runge		Runge		Durbin	
insDep P	gaad	gogo	NA	NA	NA N	NA	0000	0020	0000	9200	0000	0083	0000	0057	NA	NA	0000	0900	0000	0082	0000	0057	0000	0200	0000	0030	0000	0040	0000	¥
WatDep InsDep		дорр.	9094		3600		0094	0083	0033		3600		3600		3600		3600		9600		9600		3600		0047		0047		0047	
Longitude	DOMNIFFF DODD:	ddmin #	6610.56	6610.48	6610.78	6608.05	6607.43	6605.24	6605.56	NA	6605.30	6605.84	6605.89	6607.20	6607.75	₩	6610.62	6610.13	6609.95	6609.83	6609.57	6608.45	6608.50	6607.99	6608.35	6807.89	6715.52	6715.50	6715.52	A A
Lattude	DOMMER	III WWDD	4137.97	4137.86	4137.53	4137.74	4137.45	4136.53	4136.71	NA	4136.33	4136.12	4135.83	3912.10	4135.40	NA	4135.40	4135.43	4134.72	4134.29	4134.66	4134.29	4134.25	4133.43	4133.76	4132.26	4119.66	4119.78	4119.66	¥
Loctine	HIMMISS	ssiuiutju	161600	162700	163900	193700	195545	210700	205812	210627	211600	212900	214000	223000	230000	043700	064100	062600	064700	020000	071500	081000	082743	083700	092000	09260	140000	141000	141000	ΑĀ
Locdate	HHMMSS YMMDD: HHMMSS DDMM.FF	ymmdd bbmmys	940530		940530		940530		940530		940530		940530		940531		940531		940531		940531		940531		940531		940531		940531	
GMTtme	HHWIMSS	hhrmmss	201600	202700	203900	223900	235545	010700	005812	010627	011600	012900	014000	023000	030000	083700	100944	102600	104700	110000	111500	121000	122743	123700	132000	132600	180000	181000	181000	Α̈́
GMTdate	YYNNINDD:		940530		940530		940530	940531	940531		940531		940531		940531		940531		940531		940531		940531		940531		940531		940531	
Instrum Cast# STATION#	XXX weecel sss		CI9407A.021		CI9407A.021		CI9407A.021		CI9407A.021		CI9407A.021		C19407A.021		CI9407A.021		CI9407A.021		C19407A.021		CI9407A.021		CI9407A.021		CI9407A.022		CI9407A 022		CI9407A.022	
Cast#	XXX		900		200		000	013	831		021		8		88		032		825		902		889		200		800		904	
list on			ZPN		VPR		СТО	TAP	CTD		MOC		ZPP		VPR		CTD		MOC		ddZ		CTD		ZPN		NdZ		DFT	
EVENT#	wooddy, ees		CI15094.015		CI15094.016		Cl15094.017		C115194.001		CI15194.002		CI15194.003		CI15194.004		CI15194.005		CI15194.006		CI15194.007		C115194.008		CI15194.009		CI15194.010		CH5194.011	

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			CMTrista	CINTUMB Locdate	ocdate 1	Localme Latitude Conglitude Praticipal	antide Lt	· · · · · · · · · · · · · · · · · · ·	Tarrior III	1		TIL CUC BOOK	41000
EVENT#	E Cent	+		WANTED GOOD CODING THE DOWN FIRE DOOR CODE CODE CODE	4 CUNIVEY	THANKS I	DIMMER	DANFFF	COC	dag		10.1	7
woddw.ese IIII	::::	XXX WCCCL SSS	31	CHAIN ALL	· · · · · · · · · · · · · · · · · · ·	# E-E-E-E-E-E-E-E-E-E-E-E-E-E-E-E-E-E-E-	# # # #		dodo ppp	2020			
			yymmdd	· · · ·		o seimini	7124 32		947	-	Gifford/Durbln N	Mixed	Fog
C115194.012 CTD	034	Cl9407A.022	940531	1	200		-	 	1	0040	S	station	
			→.	_+	2000	-	+	6709 82	0047	0015 E	Durbin	Mixed	Collect
C115194,013 DPP	P 002	CI9407A.022	940531		940531	-+-		6709.76	十一			station	water
			_	231000		-1-	4121 01	6707.68	9055	0000	Durbin	Mixed	Pump
CI15294.001 ZPP	900	CI9407A.022	940601	01000	1000	-	+-	6707.17		0047		station	fallure
			_	014500	040594	+-	╁	6707.18	9500		Miller/Durbin	Mixed	
CI15294.002 MOC	023	C19407A.022	940601	3000	100	-	4119.51	6707.36		0045		Station	
 1	-	-+	_	02020	040531	+	4118.88	6707.68	9900	0000	Gifford/Durbln	Mixed	
CI15294.003 CTD	0.055	C19407A.022	200	25555		+-	4118.61	6707.74		0040		station	
				023800	040604	-	4116.70	6710.51	9200	0040	Gifford, Durbin,	Mixed	Live tow
CI15294.004 ZPN	600 N	CI9407A.022	940601	24,000	1000	+	4117.22	6710.81		0000	Runge	station	
-			_	┽	2000	2000	+-	NA NA	9055	NA	Davis	Mixed	HEA.
C115294 005 VPR	80 80	CI9407A.022	2 940601	- †	200	3		ΝΔ		NA		station	frlad
_				-+	040604	V V	670	6710.51	90055	0040	Gifford, Durbin, Mixed	Mixed	Live tow
CI15294.006 ZF	ZPN 010	C19407A.022	2 940601	-		NA NA	NA	AN AN		0000	Runge	station	
				=+	_	200	4116 70	6710.51	9025	00040	Giffard, Durbin	Mixed	Live tow
CI15294.007 ZF	ZPN 011	Cl9407A.022	2 940601		2004			6710.81		9 8 8	Runge	Station	-
			-+	+	-	2000		671292	9048	0000	Gifford/Durbln	Mixed	Live tow
CI15294.008 C	CTD 036	S C19407A.022	2 940601	+	2004	265600		6711.56	9046	9835	Вептал	station	
	TAP 014		-	-+-		2000		6710.03			Miller	Mixed	
C115294.009 ZI	ZPN 012	2 CI9407A.022	2 940601	-	200	00000	_	6709.77				station	
				_	-	-1-		E71258	8054	8	Gifford/Durbin	Mbked	_
C115294.010 C	CTD 037	7 CI9407A.022	340601	_	940601		+-	+-	╁	9035	Вегтал	station	
-	TAP 015		_	-		+		┼	9054	000 000	Gifford/Durblin	Mixed	
C115294.011 C	CTD 038	8 CI9407A.022	22 940601	一十	2000	+-	+-	AN		35		Station	
	 			-		+	┪	+-	0053	0000	Durbln	Mixad	
CH 5294 012 Z	ZPP 007	7 CI9407A.022	22 940601	1/2005	200	+-	╫	↓_	╀╌	0047		station	_

Confrient	CCC TIT				Live tow		Live tow		Live tow		Collect	water															Repair	drifter
Hegion	ğ		Mixed	station	Mixed L	Station	Mixed	station	Mixed	station	Mixed	station v	Mixed	station	Mixed	station	Mixed	Station	Mbed	station	Mixed	station	Mbked	station	Mbced	station	Mixed	Statton
	NNN.		Miller/Durbin N	S	Runge	8	Runge	S	Runge N	8	Durbin	8	Gifford/Durbin	Berman	Giffard/Durbin N	93	Gifford/Drubin 1	Berman 8	Gifford/Durbin		Durbin	•	Miller/Durbin	•	Gifford/Durbln	Berman	Durbin	
insDep Pl	ddd: 'Y	godo	0000	0045	0000 H	0045	9000 H	0045	9000 H	0045	0015 D	NA	9 0000	0035 B	9000 G	9035	0000	0040 B	0000	0040	D000	0047	0000	0049	0000	0040 E	0000	¥ Y
WatDep	clade	pappa	0054		9000		9900		9999		0055		0049		0049		9055		0900		2000		1900		0054	9900	0054	
GMTtime Loctdate Loctime Latitude Longitude WatDep HisDep Pl	HHIMMSS YYMIMDO: FRIMMSS DOWINER DOWN FFF DODDO		6714.89	6715.38	6715.08	NA	6715.08	NA	6715.08	6716,32	6715.79	NA	6714.06	6712.32	6712.56	NA	6708.50	6707.27	6707.30	NA	6202.09	6709.53	6707.85	6708.33	6714.79	6714.08	6713.88	NA
Lathtide	COMME	yymindd thirmss ddmn.fff admn.fff	4116.47	4116.42	4118.08	NA	4118.08	NA	4118.08	4118.26	4118.96	NA	4120.93	4121.75	4121.68	NA	4118.37	4115.76	4116.14	NA	4118.12	4117.78	4112.97	4112.57	4116.16	4116.28	4116.38	NA
Loctime	HHIMINISS	HINTOTISS	142500	143400	151500	NA	153000	NA	154500	154500	155600	160400	181000	190400	185470	190500	220600	230100	225300	230000	230000	0002000	000200	003500	060500	020600	073000	NA
Locdate	COMMU	ymmdd	940601		940601		940601		940601		940601		940601		940601		940601		940601		940601	940602	940602		940602		940602	
GMTtime	SSIVIMIH	hhrimsa	182500	183400	191500	NA	193000	NA	194500	194500	195600	200400	221000	230400	225700	230500	020600	030100	025300	030000	030000	040500	042000	043500	100500	110600	113000	NA
GMTdate	COMMINA	ymmdd	940601		940601		940601		940601	-	940601		940601		940601		940602		940602		940602		940602		940602		940602	
Instituti Cast# STATION#	weecel sss		CI9407A.022		CI9407A.022		CI9407A.022		CI9407A.022		C19407A.022		CI9407A.022		CI9407A.022		CI9407A.022		CI9407A.022		CI9407A.022		CI9407A.022		C19407A.022		CI9407A.022	,
Cast#	ě		924		912		013		914		8		68	910	g		24	917	Q42		88		925		943	918	865	
Institut	隻		MOC		ZPN		ZPN		ZPN		ddO		CTO	TAP	CTD	,	CTD	TAP	CTD		ZPP		00 ₩		CTO	TAP	H	
EVENT#	weddy, eee		Cl15294.013		CI15294.014		CI15294.015		C115294.016		CI15294.017		C115294.018		CI15294.019		CI15394.001		CI15394.002		C115394.003		CI15394.004		CI15394.005		CI15394.006	

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Comment	111 232		Redeploy	drifter	Live tow												Live	tow												
Parion	Ď.		Mixed	station	Mixed	station	Mixed	station	Mixed	station	Mixed	station	Mixed	Station	Mixed	statton	Mixed	station	Hydro	section	Hydro	section	Hydro	section	Hydro	section	Hydro	section	Hydro	section
	223		Durbin		Miller		Giffard/Durbin	Вегтал	Gifford/Durbln		Durbin		Miller/Durbin		Gifford/Durbin		Runge		Gifford/Durbin		Miller/Durbin		Gifford/Durbin		Gifford/Durbin		Gifford/Durbin		Gifford/Durbln	
TisDen F	dddd	coco	0000	NA	NA N	NA	0000	00400	0000	0040	0000	0037	0000	0045	0000	0040	NA	NA	0000	3600	0000	6600	0000	0000	800	0040	0000	0040	0000	0070
WatDen	0000	popp	0054		0020		0057		0057		2900		9999		9500		NA	NA	0047		0043		0048		0057		8900		0800	
GMTtime Locotate Locotine Latitode Lonoitude Mathematical	HEIMMASS YYMMOD FIRHMIKSS DOMIKEFF DOMIKEFF DODDO:	ddmm.ff	6713.73	NA	671279	6712.75	6712.94	6714.29	6713.86	NA	6714.81	6715.85	6715.88	6716.33	6717.76	6717.77	6718.54	6718.87	6724.78	6724.78	6724.28	6724.14	6720.81	6720.17	6717.01	6715.38	6711.13	6710.64	6706.60	6706.29
Latitude	DDMMFF	ddmimiff	4116.29	NA	4116.63	4116.33	411238	4110.67	4111.01	NA	4112.08	4111.62	4111.64	4111.35	4112.87	4112.98	4114.34	4114.39	4120.38	4120.98	4120.79	4120.67	4115,49	4114.69	4110.26	4108.25	4102.13	4101.87	4055.90	223100 4055.59
Loctine	THMINSS	hitimiss	085000	NA	093300	094300	115845	130300	124900	130100	132500	140500	141300	142200	160100	160800	171600	172800	181849	183400	184800	185300	194613	195200	205259	204400	212916	214000	222400	223100
Locdate	DOMINUL	ymmeld			940602		940602		940602		940602		940602		940602		940602		940602	,	940602		940602		940602		940602		940602	
GMTfrie	THIMMSS	ការការកាន់ន	125000	NA	133300	134300	155845	170300	164900	170100	172500	180500	181300	182200	200100	200800	211600	212800	221849	223400	224800	225300	234413	235200	002259	244400	012916	014000	022400	023100
GMTdate	COMMUN	winitidd	940602		940602		940602		940602		940602		940602		940602		940602		940602		940602		940602		940603		940603		940603	
Instrum Cast# STATION#			CI9407A.022		Cl9407A.022		CI9407A.022		CI9407A.022		CI9407A.022		CI9407A.022		CI9407A.022		CI19407A.02		CI9407A.023		CI9407A.023		CI9407A.024		CI9407A.025		CI9407A.026		CI9407A.027	
Cast#	×		900		915		044	019	045		600		026		046		016		047		027		048		049		020		051	
Instrum	喜		DFT		ZPN		СТD	TAP	СТБ		ZPP		MOC		CTD		ZPN		CTD		MOC		СТО		СТО		СТD		СТБ	
EVENT#	wdddyy, ees		CI15394.007		CI15394.008		CI15394.009		CI15394.010		CI15394.011		CI15394.012		CI15394.013		CI15494.014		CI15394.015		C115394.016		C115394.017	***	CI15494.001		CI15494.002		CI15494.003	

CHUIS		nc.	P	JN	. 1					Γ¢	age	<i>5</i>	\ 1-	IU												
Commant CCC TTP	Clear	raugh	Clear	rough															Fly-by		S	bottles				
Region Comment Hash RPP CCC TTP	Hydro	section	Hydro	section	Hydro	section	Hydro	section	Hydro	section	Hydro	section	Hydra	section	Hydro	section	Hydra	section	Airbome	LIDAR	Hydro	section	Hydro	section	арқн	section
	Miller/Durbin		Gifford/Durbln	·	Gifford/Durbln		Gifford/Durbin		Miller/Durbin		Glfford?Durbln		Giffard/Durbin		Glfford/Durbin		Miller/Durbin		Primmerman		Gifford/Durbln		Gifford/Durbin		Miller/Durbin	
inspep P detet: DDDD:	- 1		8	0800	0000	0115	0000	0470	0000	0298	0000	9050	0000	0510	0000	0800	0000	1600	NA	NA	0000	0000	0000	9900	0000	8983
WatDep DDDD: dddd	1800		6800		0125		0480		0612	0430	1060		1082		0100		0101		NA		0085		0000		0020	
GMT-line Lacdete Lactrine Lattrude Longhude Wadbep InsDep Pt HHMMSS YYMMDD HHMMSS DDMMFF DDMMFFF DDDD: dddd: "NINNIN" NINNINS NYMMGD NHMMSS DDMMFFF DDMMFFF DDDD: dddd: DDDD NINNINS DDMMFFF DDMMFFFF DDDD DDDD NINNINS DDMM DDDD DDD	6706.22	6706.15	6701.96	6701.82	6657.57	6657.73	662239	6652.32	6652.38	6653.20	6639.00	6639.20	6625.00	6625.18	6631.24	6631.29	6631.50	6631.92	6735.50	NA	6637.44	6637.45	6643.77	6643.81	6643.81	6644.62
Locclete Locatine Lettinde Longitude YYMMIDD: H-HIMMSS DDMIN,FF DD	4055.33	4054.49	4049.60	4049.28	4043.29	4043.01	4036.93	4036.16	4036.14	4036.53	4046.07	4046.00	4056.08	4056.12	4101.04	4100.76	4100.60	4100.24	4117.60	NA	4106.04	4105.74	4111.18	4111.89	4110.76	4110.17
Loctime: Fit-IMMSS: nhmmss	223600	225400	234200	235300	004500	005900	015200	023100	023500	031600	044500	051400	062931	071000	081230	092900	083800	085800	094023	135900	101320	102200	112424	113200	113700	120000
Locdete YYWWDD wmmdd	940602		940602		940603		940603		940603		940603		940603		940603				940603		940603		940603			
GMTIME HHMMSS thmmss	023600	025400	034200	035300	044500	045900	055200	063100	063500	071600	084500	091400	102931	111000	121230	132900	123800	125800	134023	175900	141320	142200	152424	153200	153700	160000
GMTdate YYMMDD: yymmdd	940603		940603		940603		940603		940603		940603		940603		940603		940603		940603		940603		940603		940603	
instrum Cast# STATION#	CI9407A.027		CI9407A.028		CI9407A.029		CI9407A.030		CI9407A.030		Cl9407A.031		CI9407A.032		CI9407A.033		C19407A.033		C19407A.036		CI9407A.034		CI9407A.035		CI9407A.035	
	88		22		923		954		620		922		920		057		88		8		858		620		8	
instrum	ည ္		CTD		СТО		CTD		MOC		CTD		СТО		CTD		MOC		FLY		CTD		O TD		MOC	
EVENT# wdddyy aes	CI15494.004		CI15494.005		CI15494.006		CI15494.007		Cl15494.008		C115494.009		CI15494.010		C115494.011		CI15494.012		CI15494.013		C115494.014		CI15494.015		C115494.016	

mmern	E			No bottles									Orifter	repair	ρ	>	Deploy	drifter			ē	*	Collect	water				
Ö	<u>ម</u> ត			운		-			_		-			Te.	E A	₽		늄		_	₽ T	草		*	71		77	
Region Comment			Hydro	section	Hydro	section	Hydro	section	Hydro	section	Hydro	section	Stratified	station	Stratffled	station	Stratified	station	Stratified	station	Stratifled	station	Stratified	station	Stratffled	station	Stratifled	station
	羹		Glfford/Durbin		Glfford/Durbin		Gifford/Durbin		Gifford/Durbin		Miller/Durbin		Durbin		Durbin		Durbin		Gifford/Durbin		Durbin		Ourbin		Miller		Gifford/Durbln	
lrsDep	papa	CCCC	0000	2000	0000	0020	0000	0040	0000	0035	0000	0050	0000	NA	0000	0000	0000	NA	0000	1200	0000	9890	800	0015	NA	NA	0000	000
declism	codo		0200		0065		6300		0051		0020		0087		0087			8800	0088		0088		0088		NA		0091	
appunduo	JONN FF	1dmm.fft	6650.07	6650.30	6656.58		6702.98	6703.52	6708.43	6709.20	6709.76	6710.12	6609.30	NA	6609.01	60.6099	6615.01	NA	6614.95	6614.67	6614.62	6614.14	6614.21	6614.11	6618.71	6618.65	6616,10	6615.79
atitude	DIMMEE	dmmt	4116.28	4116.02	4121.39		4126.45	4126.53	4130.96	4131.67	4131.96	4132.23	4120.12	NA	4119.94	4119.83	4134.84	NA	4134.59	4134.12	4134.16	4133.30	4133.45	4132.92	4132.81	4132.72	4129.77	4129.32
actine .	LIMMSS	himmss ddmmitt ddmmitt: dddd	125913 4	130600	140115 4	141200	150952 4	151900 4	155604 4	162100	162000	163000	200000	NA	222300	223600	030500	NA	004300	002800	010000	012700	014700	024200	081400	082400	120401	122100
ocdate	T COMMY	yymmidd h	940603		940603		940603		940603		940603		940603	1	940603		940604		940604		940604		940604		940604		940604	
te (GMTfinne Locatore Lactime Latitude Langfaude WatiDea Institute Pt	FINNSS: Y	1 hhmmss y	165913	170600	118011	181200	190952	191900	195604	202100	202000	203000	0002000	NA	022300	023600	043500	AN	044300	045800	020000	052700	054700	064200	121400	122400	160401	162100
GIVITCHATE C	TYNNINDE (HEMINSS TYNNINDE) HEIMINSS ODINMER ODINMER ODDIN	yymmdd .	940603	-	940603		940603		940603		940603	-	940604		940604		940604		940604		940604		940604		940604		940604	
Instume Cast# STATION#	XXX WYCCCCI SSS		CI9407A.036		CI9407A.037	-	CI9407A.038		CI9407A.039		CI9407A.039		CI9407A.040		CI9407A.040		CI9407A.041		CI9407A.041		CI9407A.041		CI9407A.041		CI9407A.041		CI9407A.041	
Cast#	X		8		981		790		88		332		8		915		200		8		910		8		017		965	
Instructe	丰		CTD		CTD		CTD		CTD		MOC		DFT		ZPN		DFT		CTD		ZPN		ОРР		ZPN		GF3	
EVENT*	999	1::::	C115494.017		CI15494.018		CI15494.019		CI15494.020		C115494.021		C 15594.001		CI15594.002		CI15594.003		CH5594.004		C115594.005		CI15594.006		CI15594.007		CI15594.008	

site		9035		6721.19	4115.46	054800		094800					
Mixed	Glfford/Durbin	0000	0048	6720.53	4115.27	053468	940607	093468	940607	CI9407B.042	88	CTD	CH5894.003
site		ΑĀ		NA	NA	NA		NA					
Mixad	Durbin	8	84 1	6720.19	4115.07	050300	940607	000000	940607	CI9407B.042	940	DFT	CI15894.002
site		NA		NA	NA	NA		NA A					
Mbæd	Durbin	8	841	6744.72	4104.87	030000	940607	000020	940607	CI9407B.042	8	ᇤ	CI15894.001
site		NA		NA	NA	NA		NA					
Stratified	Durbin	8	888	6626.15	4122.04	060700	940605	100700	940605	CI9407A.041	88	DET	CI15694.005
site		060 060		6626.34	4122.06	063700		103700					
Stratffed	Glffard/Durbln	8	9093	6626.19	4122.04	063100	940605	103100	940605	CI9407A.041	290	CTD	CI15694.004
site		0087		6620.53	4123.72	015600		009930					
Stratifled	Miller/Durbin	000 000	99	6620.87	4123.76	013000	940605	053000	940605	CI9407A.041	88	MOC	C115694.003
site		0057		6620.86	4123.96	012000		052000					
Stratified	Durbin	0000	0600	6620.54	4125.09	000000	940605	043000	940605	CI9407A.041	911	ddZ	CI15694.002
slta		0070		6620.50	4125.47	001300		041300					
Stratifled	Gifford/Durbin	8	6600	6628,49	4125.34	000400	940605	040400	940605	CI9407A.041	990	CTD	C115694.001
site		NA		6622.46	4127.8	082400		242900					
Stratified	Miller	NA	NA	6622.47	4127.81	081400	940604	241900	940604	C19407A.041	020	NdZ	CH5594,014
site	Durbin	0080		6619.58	4126.87	155100		195100					
Stratffed	Runge	0000	0092	6619.75	4126.89	154500	940604	194500	940604	CI9407A.041	919	ZPN	CI15594.013
site	Durbin	0080		6619.75	4126.93	154500		194500					
Strattfled	Runge	0000	0091	6618.75	4126.89	151500	940604	191500	940604	CI9407A.041	910	ZPN	Cl15594.012
slte				6618.65	4126.93	151000		191000					
Stratified	Durbin	0015	0600	6618.49	4127.01	144500	940604	184500	940604	CI9407A.041	905	OPP	CI15594.011
stte		0080		6617.59	4126.73	143000		183000					
Stratified	Miller/Durbin	0000	1600	6616.82	4127.34	141100	940604	181100	940604	CI9407A.041	833	MOC	CI15594.010
site		2000		6616.69	4127.51	140500		180500					
Stratified	Durbin	0000	0091	6616.38	4128.45	130000	940604	170000	940604	C19407A.041	99	ZPP	CI15594.009
		OCCO						hhmmss	wmmdd:				
THE PPP	Z	dddd	DDDD:	DDIMMEE	DOWN FF	HINIMSS	COMMA	SSIVINI#1	COMMA	\$\$\$ (DDDDAA	×	喜	•
:::			watDep.	Langitude	Latitude	Localme	Locdate	GMTtime			Cast#	Instrume	EVENT*
	Stratified Stratified Stratified Stratified Collect site site site site site site site sit	MNAY: Stratified site in a	Durbin Durbin Burge Durbin Burge Durbin Miller/Durbin Miller/Durbin Miller/Durbin Durbin Durbin Durbin Durbin Ourbin Durbin Durbin Durbin Ourbin Durbin Durbin Durbin Durbin Durbin Durbin Durbin Durbin	Durbin Durbin Burge Durbin Burge Durbin Miller/Durbin Miller/Durbin Miller/Durbin Durbin Durbin Durbin Durbin Ourbin Durbin Durbin Durbin Ourbin Durbin Durbin Durbin Durbin Durbin Durbin Durbin Durbin	Durbin Durbin Burge Durbin Burge Durbin Miller/Durbin Miller/Durbin Miller/Durbin Durbin Durbin Durbin Durbin Ourbin Durbin Durbin Durbin Ourbin Durbin Durbin Durbin Durbin Durbin Durbin Durbin Durbin	Durbin Durbin Burbin Burbin Burbin Burbin Burbin Burbin Burbin Burbin Durbin Durbin	Durbin Durbin Burge Durbin Burge Durbin Miller/Durbin Miller/Durbin Miller/Durbin Durbin Durbin Durbin Durbin Ourbin Durbin Durbin Durbin Ourbin Durbin Durbin Durbin Durbin Durbin Durbin Durbin Durbin	Durbin Durbin Burge Durbin Burge Durbin Miller/Durbin Miller/Durbin Miller/Durbin Durbin Durbin Durbin Durbin Ourbin Durbin Durbin Durbin Ourbin Durbin Durbin Durbin Durbin Durbin Durbin Durbin Durbin	Durbin Durbin Burge Durbin Burge Durbin Miller/Durbin Miller/Durbin Miller/Durbin Durbin Durbin Durbin Durbin Ourbin Durbin Durbin Durbin Ourbin Durbin Durbin Durbin Durbin Durbin Durbin Durbin Durbin	GWITTERING Checkeles Locatories Latitudes Chiptignation Media (1998) FF WYMANDIDIS (H-MMMSS) Chiptignation Chiptignation	CANTIGNER CANTIGNER <t< td=""><td>CANTIGNER CANTIGNER <t< td=""><td> No. No.</td></t<></td></t<>	CANTIGNER CANTIGNER <t< td=""><td> No. No.</td></t<>	No. No.

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ā	ZZZZ.		Davis		Durbin		Gifford/Durbin		Durbin		Miller/Durbin		Davis		Miller		Gifford/Durbln		Durbin		Miller/Durbin		Runge		Runge		Davis	
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CI16194.001	CTD	075	Cl9507B.042	940610	02020	940609	220500	4107.97	6731.43	90955	0000	Gifford/Durbin	Mbed	
					021800		221800	4108.39	6731.42		0040		site	
CI16194.002	ZPP	015	Cl9407B.042	940610	030000	940609	230000	4108.75	6732.08	9055	0000	Durbin	Mixed	
					030400		234000	4109.06	6731.32		0047		slta	
C116194.003	MOC	989	Cl9407B.042	940610	040300	940610	000000	4109.25	6731.35	0051	0000	Miller/Durbin	Mixad	
					041700		001700	001700 4109.23	6731.43		0047		sitte	
CI16194.004	VPR	013	Cl9407B.042	940610	051444	940610	011444	4109.18	6730.19	0900	0040	Davis	Mixed	
					081765		041755	4100.45	6723.13		0000		site	
CI16194.005	CTD	9/0	CI9407B.042	940610	091900	940610	051900	4105.89	6729.70	9900	0040	Gifford/Durbin	Mixed	Last mixed
					093200		053200	4105.61	6729.74		0042		ste	site cast
CI16194.006	H	011	CI9407B.042	940610	094700	9406610	054700	4105.54	6730.13	9500	0000	Durbin	Mixed	Retrieve
à					NA		NA	NA	NA		NA		site	drifter
CI16194.007	H	012	CI9407B.043	9406610	115000	940610	075000	4102.01	6716.13	0057	0000		Startfled	Redeploy
					NA		NA	NA	NA		NA		site	drifter
CI16194.008	ZPN	031	Cl9407B.043	940610	110000	940610	000020	410215	6711.69	8500	0000	Durbin	Stratified	Live
					112000		072000	4101.86	6711.52		0055		site	tow
C116194.009	CTD	220	CI9407B.043	940610	120200	940610	080200	4101.89	6711.33	6900	0000	Gifford/Durbin	Startifled	
					121300		081300	4101.90	6711.72		0051		site	
CI16194.010	ΩРР	200	CI9407B.043	940610	132000	940610	092000	4102.23	6713.77	8900	0015	Durbin	Stratifled	Collect
					142000		102000	4103.39	6715.66		NA		site	water
CI16194.011	FLΥ	600	CI9407B.043	940610	133600	940610	093600	4211	6654.8	NA	NA	Primmerman	Startifled	Fly-by
					165000		125000	NA	NA		NA		site	
C116494.012	СТБ	078	Cl9407B.043	940610	150400	940610	110400	4103.21	6715.47	0065	0000	Gifford/Durbin	Stratified	
					151800		111800	4103.56	6715.45		0020		slte	
CI16194.013	ZPP	910	CI9407B.043	940610	152000	940610	112000	4104.17	6715.01	0064	0000	Durbin	Startifled	
					161500		121500	4104.28	6715.21		0057		site	

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Contiment	COC TIT				Live	tow	- Ke	to*	Live	tow	<u></u>	tow tow							Sunny, flat,	саш	Сотритег	problem	Repeat	cast				
Region	янные ссстт		Stratified	site	Stratified	site	Stratified	site	Strattfled	site	Stratified	site	Stratifled	stte	Stratified	site	Stratifled	site	Stratified	stra	Strattfled	site	Strattfled	site	Stratfled	site	Stratifled	site
H. W.	NEW.		Miller/Durbin		Durbin, Gifford	Runge	Durbln, Gifford	Runge	Durbin, Gifford	Runge	Durbin, Gifford	Runge	Glfford/Durbin		Durbin		Miller/Durbin		Davis		Gifford/Durbln		Gifford/Durbin		Durbín		Miller/Durbin	
dacisul		GCCO	88	0054	8	0900	8	0040	8	0055	8	0045	8	0020	88	0057	0000	6300	NA	NA	000	9055	800	0065	88	0057	0000	883
WatDap	COCO	cicia	0064		8		0062		0065		0065		0070		900		6900		1900		9200		0075		0073		1200	
ephylouo	DOMM FF	अंतामान्य वी	6715.50	6715.23	6715.17	6715.30	6715.30	NA	6715.30	NA	6715.30	6715,45	6715.15	6715.50	6715.71	6715.52	6715.43	6715.39	6715.39	6715.15	6720.11	6720.36	6720.41	6720.49	6720.64	6720.69	6720.55	6720.05
GMTtime: Locate Locatine Lattiude Longitude WatDap	HEIMINGS YMMINDD: HEIMINGS DOWN FE DOWN FF DOOD: dddd:	yymined: hhiminss ddiminiff ddiminiff	4104.35	4104.51	4104.28	4104.52	4104.52	NA	4104.52	NA	4104.52	4104.66	4101.31	4101.48	4101.63	4102.02	410200	4101.43	4101.84	4056.35	4057.50	4057.95	4057.97	4058.17	4058.41	4059.38	4059.33	4059.19
contine	THE INVISES:	hhmmss	122000	123000	130000	131000	131500	NA	133000	¥ Y	134500	133400	220200	221700	230000	234500	235100	001500	005020	071119	110000	111800	112400	113400	120000	125500	130100	131200
Locdate	COMMY	wmmdd	940610		940610		940610		940610		940610		940610		940610		940610		940611		940611		940611		940611		940611	
GMTtime	H-MASS.	hhmmss	162000	163000	170000	171000	171500	NA	173000	NA	174500	173400	020200	021700	030000	034500	035100	041500	005020	071110	150000	151800	152400	153400	160000	165500	170100	171200
GWTdate	1.75	1	940610		940610		940610		940610		940610		940611		940611		940611		940611		940611		940611		940611		940611	
Institute Cast# STATION#			CI9407B.043		CI9407B.043		Cl9407B.043		CI9407B.043		CI9407B.043		CI9407B.043		Cl9407B.043		CI9407B.043		CI9407B.043		CI9407B.043		CI9407B.043		CI9407B.043		CI9407B.043	
Cast#	Š		83		32		88		8		88		079		017		040		914		080		084		018		å	
Instruction	喜		MOC		ZPN		ZPN		ZPN		ZPN		CTD		Zpp		MOC		VPR		CTD		СТО	-	Zpp		MOC	_
#LVUV.U	é		CH6194.014	1	CH6194.015		CH6194.016		CH6194.017		C116194.018		C116294.001		C116294 002		CI16294 003		CI16294 004		CI19294.005		CH6294 006		C116294 007		Cl16294.008	

41170	Price and kerific	#teg()	The High Land 14 STATION #	GMTdate	GMTume (Locdate (Loctaine Lattinde Longleide) WatDeo (InsDep P	Locdate	Localne	atttude	application	WatDen	LISDAD		Region	Comment
•.1.•	2000	3		41 E	COLD WAYNES YOUR HENNINGS DOWN FF DOWN FF DODGE	CONTRACT.	- WANGE	DUMMEE	PINM FF	COCO	::::	之	THE PPP CCC TIT	FEC 11
widdly age		X	XXX VYCCCCLSSS	X YMMC/LA	CCIVIMILITY	ייייייייייייייייייייייייייייייייייייייי	The Manager	· ***	1	1	1.			
				yymindd	hhmmss	: Symmeta	hhmmss	hhmmss: ddmm.ff ddmm.ff dddd	odmini ita	::-	0000			
C116294 009	ZPN	989	CI9407B.043	940611	172500	940611	132500	4059.54	6720.43	9071	800	Runga	Stratffed	Lye
					NA		NA	NA	NA		980		site	tow
C11 F294 010	ZPN	037	CI9407B.043	940611	171500	940611	133000	4059.54	6720.43	0071	88	Runge	Strattfied	Live
01024.010		3			175100		135100	4059.80	6719.40		0020		stte	tow
C14 E204 044	/pp	ج ا	C19407B 043	940611	180900	940611	140900	4059.58	6719.14	2900	NA	Davis	Stratifled	
010834:011		3			214258		174258	4044.84	6712.07		NA		site	
C14 R294 012	NdZ	88	CI9407B.043	949611	230000	940611	190000	4056.50	6716.51	6200	8	Durbin	Stratffed	Live
1000000	:	3			232000		192000	4056.41	6716.48		00700		site	wat
C148304 001	NdZ	050	CI9407B.043	940612	009000	940611	200500	4056.16	6717.05	6/00	0000	Durbin	Strattfled	- Ke
100.10	:				001600		201500	4056.07	6717.30		0075		site	tow
C146304 002	NdZ	040	CI9407B.043	940612	9000	940611	210000	4055.82	6717.63	6200	000	Glfford	Stratffed	∏VB
Too Hoose To	i				011300		211300	4055.72	6717.60		0070		site	tow
C148304 003	QĮ.	80	C19407B 043	940612	020200	940611	220200	4056.08	6718.35	6200	0000	Gifford/Durbln	Stratified	No bottles
SON TRANSITO					022000		222000	4056.40	6718.72		0900		stte	fred
C14 8304 004	CTD	283	C19407B 043	9406612	022600	940611	222600	4056.24	6718.82	8200	0000	Gifford/Durbln	Stratified	Callect
10010		3					224100	4056.35	6718.82		0900		sitte	water
C116304 005	700	650	C19407B.043	940612	030000	940611	230000	4056.32	6718.93	8200	0000	Durbin	Stratffed	
200,100	_				035500		235500	4056.43	6719.61		290		stte	
CH6394 006	MOC	042	CI9407B.043	940612	035900	940611	235900	4056.48	6719.57	0078	8 8	Miller/Durbin	Stratified	
		-		-	041800	940612	001800	4055.90	6719.99		8		stte	-
CH 6394 607	VPR	910	CI9407B.043	940612	050722	940612	010800	4056.35	6719.67	9200	800	Davis	Stratfiled	
100	7				071900		031800	4103.06	6724.08		0900		sfte	
C116394 008	CTD	984	CI9407B.043	940612	150500	940612	110500	4054.66	6720.87	9200	8	Gifford/Durbln	Strattfled	
				-	151500		111500	4054.91	6720.91		88		site	
C116304 009	700	020	C19407B.043	940612	160000	940612	120000	4055.36	6720.79	6200	8	Durbin	Stratffed	
80.5	7				164500		124500	4055.31	6720.41		0057		sfte	

EVENT#	Instrutte	Cast#	Instrume Cast# STATION#	GMTdate	9 CM/Time Locdate Loctime Latitude Longhude WatDep InsDep Pl	Locdate	Locitine	Latitude	Longlatide	WatDep	InsDep		Region Comment	Comment
weddy, eee		xx	Wedeck sss	YYMINDD	A HEMMISS YYMMOD; HEMMISS DOWNER DOWNER DODG:	COMMINA	THIMMES	DOMW.FF	DDIMM FF	0000	gaq	ングを	THE PP CCC TIT	CCC IIII
				wmmdd	hhmmss	ymmdd	hhmmss	ddmm.ff	ddmm.ff. ddmm.ff.	adda	qoqo			
CI16394.010	MOC	043	CI9407B.043	940612	165000	940612	125000	4056.27	6720.66	8200	0000	Milter/Durbin	Stratified	
					170800		130800	4056.01	6719.92		0075		stte	
CI16394.011	ZPN	041	CI9407B.043	940612	172000	940612	132000	4055.99	6719.71	8200	NA	Runge	Stratifled	Live
					174500		134500	4056.19	6719.00		NA		site	tow
CI16394.012	VPR	017	CI9407B.043	940612	181030	940612	141030	4055.77	6718.51	0073	0000	Davis	Stratified	Sunny, flat
					193800		153800	4055.44	6717.35		0063		site	calm
Cl16494.001	стр	085	CI9407B.043	940613	020000	940612	220000	405221	6717.43	0085	0000	Gifford/Durbin	Stratified	
					021600		221600	4052.40	6717.50		3900		stra	
CI16494.002	ddZ	021	CI9407B.043	940613	025500	940612	225500	4052.59	6718.19	0084	0000	Durbin	Stratified	
					034000		234000	4053,11	6718.38		0065		site	
CI16494.003	MOC	044	CI9407B.043	940613	032000	940613	235000	4052.99	6718.46	0084	0000	Miller/Durbin	Stratified	
					041000		241000	4053.86	6718.15		0800		stte	
CI16494.004	VPR	018	CI9407B.043	940613	020800	940613	010800	4053,26	6718.63	0084	NA	Davis	Stratifled	
					008690		023800	4054.38	6718.12		NA		site	
CI16494.005	СТD	980	CI9407B.043	940613	070200	940613	030200	4054.60	6717.95	0081	0000	Gifford/Durbln	Stratified	
					071700		031700	4054.72	6717.76		0061		stte	
CI16494.006	DFT	013	CI9407B.043	940613	080300	940613	040300	4054.18	6716.65	0084	NA	Durbin	Stratffed	Recover
					NA		NA	NA	NA		NA		site	drifter
Cl16494.007	СТБ	087	CI19407B.04	940613	110400	940613	070400	4120.94	6725.06	0042	0000	Gifford/Durbin	Hydro	
					111100		071100	4120.80	6725.03		0031		section	
Cl16494.008	MOC	045	CI9407B.044	940613	112000	940613	072000	4120.63	6725.04	0044	0000	Miller/Durbin	Hydro	
					113000		000670	4153.86	6718.15		9035		section	
Cl16494.009	СТБ	088	CI9407B.045	940613	122000	940613	082000	4114.72	6720.48	0048	0000	Gifford/Durbin	Hydro	
					122900	,	082900	4114.67	6720.72		9035		section	
Cl16494.010	СТD	680	C19407B.046	940613	132000	940613	092000	4108.47	6715.85	9900	0000	Gifford/Durbin	Hydro	Severe
					133300		093300	4108.57	6716.15		0042		section	spiking

Contrient	CCC TIT		Repeat	cast				·																	Llva	tow		
Region	янн РРР ссс ттр		Hydro	section	Hydra	section	Hydro	section	Hydro	section	Hydro	section	Hydro	section	Hydro	section	Hydro	section	Hydro	section	Hydro	section	Transect		Georges	Basin	Georges	Basin
1	NAME:		Gifford/Durbin		Gifford/Durbin		Miller/Durbin		Gifford/Durbin		Gifford/Durbin		Glford/Durbin		Miller/Durbin		Glfford/Durbin		Gifford/Durbln		Glfford/Durbin		Davis		Hunge		Giffard/Durbin	
dacishi		adda	0000	0040	0000	0020	0000	0063	0000	0081	0000	9200	0000	0120	0000	0135	0000	0470	0000	0485	0000	0480	0000	0126	NA	NA	0330	
WattDep	adda		8900		0068		8900		0062		0600		0131		0143		0491		0783		1187		1783		0344		0344	
Langhude	DOMW,FF.	ddininitif	6716.24	6716.37	6711.42	6711.60	6711.56	6711.57	6706.66	6706.91	6702.12	6702.04	6657.12	6656.56	6657.06	6655.77	6652,95	6649.02	6638.86	6638.18	6624.76	6624.05	6623.39	6716.81	6710.09	6710.1	6710.13	6710.75
attructe	HHIMINICIC	dmm ff	4108.62	4108.69	4102.22	4102.38	4102.28	4102.31	4055.97	4055.99	4049.59	4049.67	4043.30	4043.15	4043.29	4042.55	4037.04	4037.76	4046.03	4046.00	4055.91	4055.12	4054.62	4238.35	4222.77	4222.56	4222.95	4222.75
octine i	SSMINI	i) ssuiuji	093900 4108.62	094600	105200	110000	110700	111800	122300	123300	133300	134400	144100	145400	145500	152800	162500	171800	182300	185400	201100	204300	210349	006000	005300	010900	013500	021100
ocdate L	HI GOMINA	Minned dold dominitia dominitia	940613		940613		940613		940613		940613		940613		940613	_	940613		940613		940613		940613	940615	940615		940615	
GMIdate GMItime Locdate Locathe Lathude Langhude WatDep InsDep	DD. HITHIMISS YYMMOD: HITHIMISS DOMM.FF ODMIN.FF ODDING HOUDDS:	(ssiuiujų p		134600	145200	150000	150700	151800	162300	163300	173300	174400	184100	185400	185500	192800	202500	211800	222300	225400	001100	004300	010349	040900	045300	020300	053600	061100
SMT date	COMMA	yyminid	940613		940613		940613		940613		940613		940613		940613		940613		940613		940614		940614	940615	940615		940615	
Institute Cast# STATION#	XXX: Vvcccclsss		CI9407B.046		CI9407B.047		CI9407B.047		CI9407B.048		CI9407B.049		CI9407B.050		C19407B.050		CI9407B.051		CI9407B.052		CI9407B.053		CI9407B.053		Cl9407B.054		CI9407B.054	
Cast#	×		8		9		046		260		88		094		740		360		960		097		919		<u>ş</u>		86	
Instrume	臺		CTD		CTD		MOC		CTD		CTD		CTD		Moc		CTD		CTD		CTD		VPR		ZPN		СТБ	
EVENT*	wdddyy ees		Cl16494.011		CH6494.012		C116494.013		C116494.014		CI16494.015		C116494.016		C116494.017		C116494.018		C116494.019		C116594.001	,	CI16594.002		Cl16694.001		CI16694.002	

Comment	CCC TIT		Diapause	collection			Dlapause	collection	Calanus	dnos
Recton	HAR PPP CCC TIT		Georges	Basin	Wilkinson	Basin	Wildnson Diapause	Basin	Wilkinson	Basin
			0000 Miller/Durbin		Gifford/Durbin Wilkinson		Miller/Durbin		0000 Miller/Durbin	
riscieri P	dd¢d;	COCO	0000	0322	0000	0190	0000	0139	0000	0035
WatDep	COCO		0338	6960	0220		0215		0218	
Longmide	DDMM.FF	ddmmi.tft	6710.71	4221.4 6712.92	6913.09	6913.21	6913.22	6911.92	691233	6906.72
Latinde	DDWM:FF	ddmmtt	022600 4222.49 6710.71	4221.4	134800 4153.33 6913.09	141000 4153.75 6913.21	940615 141500 4153.66 6913.22 0215	150700 4155.68 6911.92	154500 4153.07 6912.33	180100 4151.93 6906.72
Localme	L#-IMMSS:	hhmmss	022600	000000	134800	141000	141500	150700	154500	180100
Locdate	CICININIA	ymindd	940615		940615		940615		940615	
Givinitine	SSWMHH.	f himmiss gymmdd himmiss ddinnuff ddinnuff ddddi	062600 940615	073000	174800	181000	181500	190700	194500 940615	220100
GMTdate	YYNNDD HHMMSS YYNNDD HHMMSS DDMM:FF DDMM:FF DDDD: ddbd; I'NNNN	windd	5		2		5		5	
EVENT# Instrume Cast# STATION# GMTdate Coodate Locdate Latitude Latitude WatDep Inschep P	XXX vvccccl.sss		CI16694.003 MOC 048 CI9407B.054 94061		099 CI9407B.055 94061		CI16694.005 MOC 049 CI9407B.055 94061	42	CI16694.006 MOC 050 CI9407B.055 94061	
Cast#	ă		048		660		88		920	
Instrume	=		Moc		CTD		Moc		MOC	
EVENT#	weddly, asa		CI16694.003		C116694.004 CTD		CI16694.005		C116694.006	

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