

East-West Distribution of Nutrients and Dissolved Inorganic Carbon in the Northern North Pacific in Autumn

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Abstract: We measured nutrient and dissolved inorganic carbon (DIC) concentrations from west to east across the northern North Pacific Ocean from station K2 to Ocean Station Papa during September and October 2005. At 2000 m depth, silicate levels increased and dissolved oxygen levels decreased from west to east. This trend was attributed to seawater in the layer flowing from west to east in the northern North Pacific. Distributions of normalized (salinity = 33) nutrients and DIC showed the general trend of “west high, east low” (WHEL) in the surface layer. Normalized nitrate and DIC concentrations above 30 m depth ranged from 7 to 13 $\mu\text{mol kg}^{-1}$ and from 2027 to 2045 $\mu\text{mol kg}^{-1}$, respectively. Seasonal drawdowns of nutrients and DIC from winter to autumn showed the WHEL trend, similar to previous studies. The ratio of silicate drawdown to nitrate + nitrite drawdown also showed the WHEL trend, which might result from longitudinal differences in the proportion of diatoms in the phytoplankton community in the northern North Pacific.

Keywords: Nutrient, dissolved inorganic carbon, northern North Pacific.

1. INTRODUCTION

Studies of the carbon cycle have been conducted at time-series stations in the northeastern and northwestern North Pacific Ocean to estimate the strength of the biological pump. Ocean observations have been conducted at Ocean Station Papa (OSP; 50°N, 145°W) in the eastern subarctic Pacific since 1956 [1, 2]. Carbon and nutrient concentrations have been measured at station KNOT (44°N, 155°E) in the western subarctic Pacific since 1992 [3], and intensive time-series observations were conducted at station KNOT between 1998 and 2000 [4]. Since 2001, we have been conducting time-series observations at station K2 (47°N, 160°E) in the northwestern North Pacific [5-8]. As a result of these various studies, it became clear that seasonal net community production from winter to autumn in the Western Subarctic Gyre (37-61 gC m^{-2} [8-10]) was higher than that in the Alaska Gyre (26-35 gC m^{-2} [2]). Spring blooms, consisting mainly of diatoms, are known to occur only in the western part of the subarctic Pacific [11, 12]. Diatoms in this area, because of their large size and resultant high settling velocity, are suspected of playing a key role in transporting particulate organic carbon to the deep ocean [13-16].

In addition, sea surface measurements of CO_2 and nutrients from commercial vessels have been made between the eastern and western North Pacific since 1995 [17, 18]. These results also indicate that the strength of the biological pump

in the western part of the northern North Pacific is higher than that in the eastern part. There have been many previous studies of the biological pump at each time-series station in the northern North Pacific, and the observations of sea surface CO_2 and nutrient levels were previously routinely conducted from east to west across the northern North Pacific. However, simultaneous hydrographical observations of CO_2 and nutrients from east to west across the northern North Pacific were never conducted during autumn when the drawdown of nutrients in the upper layer (approximately 100 m) is at an annual maximum [1, 8].

In September and October 2005 we conducted biogeochemical observations across the northern North Pacific from station K2 to OSP [19, 20]. In this paper, we present our vertical profiles of dissolved inorganic carbon (DIC) and nutrient concentrations and compare the strength of the biological pump from east to west across the northern North Pacific.

2. SAMPLING AND ANALYSIS

2.1. Sampling Locations

Observations were conducted during the cruise of R/V *Mirai* in September and October 2005 in the northern North Pacific (Fig. 1). Samples were collected at stations K1 (51°N, 165°E), K2 (47°N, 160°E), EW1 (47°40N, 169°16E), EW2 (47°N, 175°E), EW3 (46°N, 180°E), EW4 (46°N, 175°W), EW7 (49°30N, 175°W), and OSP (50°N, 145°W). Stations K1 and K2 are located in the northern and central parts of the Western Subarctic Gyre, respectively. OSP is located in the Alaska Gyre.

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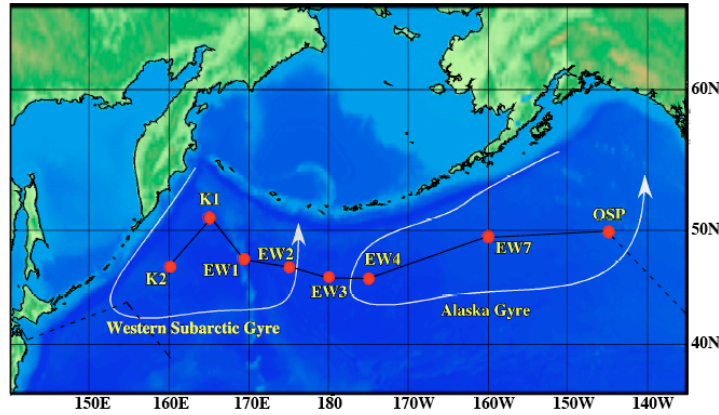


Fig. (1). Location of sampling stations and main ocean currents in the northern North Pacific Ocean. Gray lines with arrows indicate the subarctic gyres in the eastern and western regions [1].

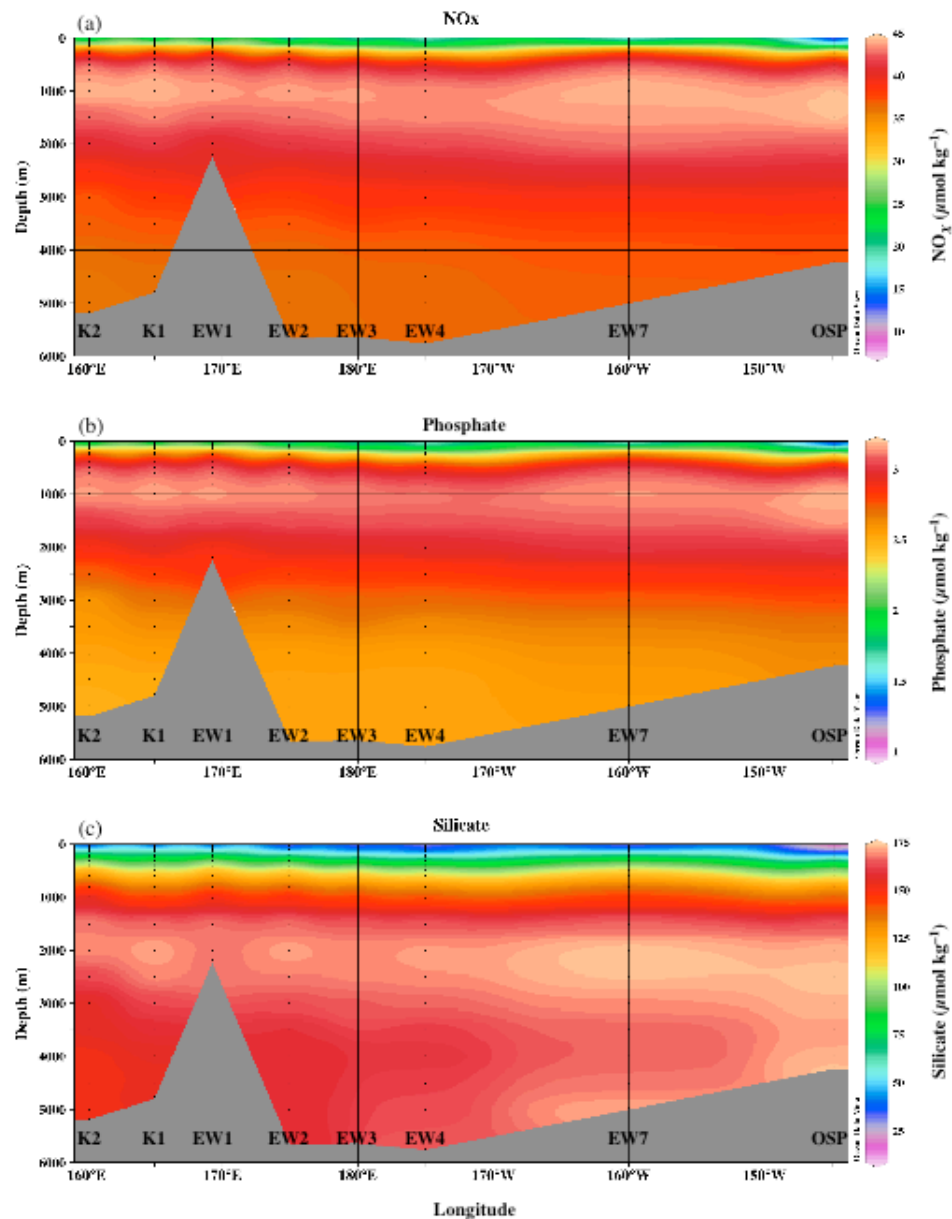


Fig. (2). Distributions of nitrate + nitrite (NO_x) (a), phosphate (b), and silicate (c) concentrations from surface to bottom in the northern North Pacific in September and October 2005. To illustrate east-west trends, data are arranged from west to east ignoring differences in latitude. Dots and gray areas indicate sampling depths and the sea floor, respectively.

2.2. Water Sampling and Sample Analysis

Seawater samples were collected from the surface to the bottom layer by Niskin bottle samplers attached to a CTD (SBE 911plus, Sea-Bird Electronics Inc.).

Nutrient concentrations were determined by using a continuous flow analyzer (TRAACS 800 system, Bran+Luebbe). The precision (1σ) of nutrient analysis was less than 0.3%. DIC was measured by coulometer (carbon dioxide coulometer Model 51012, UIC Inc.). The coulometer was calibrated using Dickson's certified reference material [21]. The precision (1σ) of DIC measurements was $\pm 1.0 \mu\text{mol kg}^{-1}$. Total alkalinity (TALK) was determined by potentiometric titration [22]. The precision (1σ) of TALK measurements was $\pm 1.3 \mu\text{mol kg}^{-1}$. Dissolved oxygen (DO) was analyzed by potentiometric titration [23]. The precision (1σ) of DO measurements was $\pm 0.05 \mu\text{mol kg}^{-1}$. Apparent oxygen utilization (AOU) was calculated as the difference between observed DO concentration and equilibrium

concentration with the atmosphere at the measured temperature and salinity [24].

3. RESULTS AND DISCUSSION

3.1. Distributions of Nutrient, DIC, TALK, and DO from Surface to Bottom

Station EW1 is located above the Emperor Seamount Chain. The bottom depth at station EW1 is approximately 2200 m, whereas the bottom depth at the other stations range from 4200 to 5500 m (Fig. 2). All physical and chemical data from all stations are included in the Supplementary Material.

Nitrate plus nitrite (NO_x) and phosphate showed maximum concentrations around 1000 m depth across the northern North Pacific (Fig. 2a and b). We attributed the NO_x and phosphate maxima to decomposition of sinking particulate organic matter (POM). However, silicate

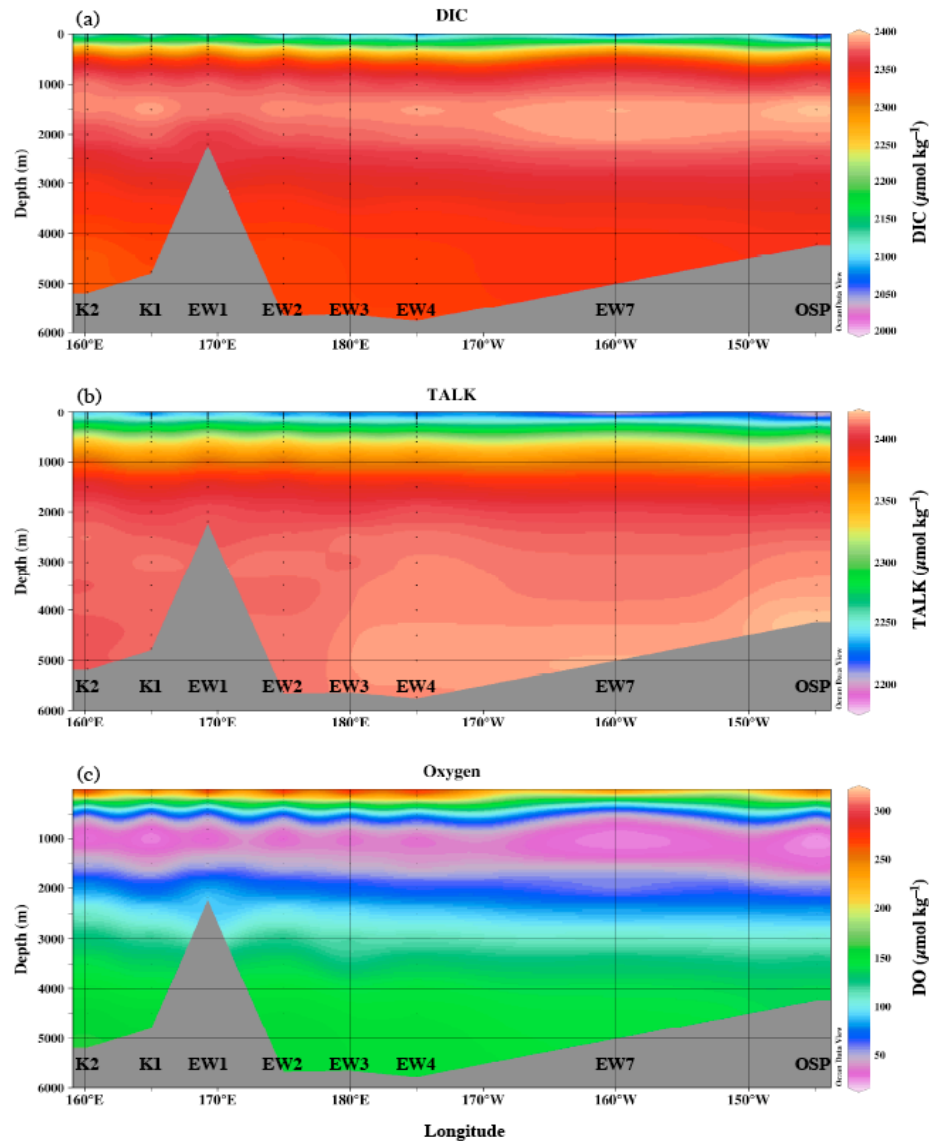


Fig. (3). Distribution of DIC (a), TALK (b), and DO (c) concentration from surface to bottom in the northern North Pacific in September and October 2005. To illustrate east-west trends, data are arranged from west to east ignoring the differences in latitude. Dots and gray areas indicate sampling depths and the sea floor, respectively.

concentrations showed maxima from 2000 to 2500 m depth across the northern North Pacific and also near the bottom at stations EW7 and OSP (Fig. 2c). The silicate maximum in the intermediate layers was attributed to dissolution of sinking opal. We attributed the difference between the depth of the silicate maximum and that of the other nutrients to the different rates of decomposition for opal and other POM. The silicate maximum near the bottom at stations EW7 and OSP could be caused by dissolution of the opals from the sediment.

Maximum DIC concentrations were found at approximately 1500 m depth across the northern North Pacific (Fig. 3a). The DIC maximum is attributed to decomposition of sinking POM, similar to NO_x and phosphate. TALK reached the maximum levels near the bottom at stations EW7 and OSP (Fig. 3b). These TALK maxima might be caused by the dissolution of carbonates from sediment. Minimum DO concentrations were measured at approximately 1000 m depth (Fig. 3c). We attribute the DO minimum to the consumption of DO accompanying decomposition of the sinking POM. Therefore, the DO minimum layer coincided with the depth of maximum NO_x and phosphate concentrations (Figs. 2a, 2b, and 3c).

At 2000 m depth, silicate levels increased and DO levels decreased from west to east (Figs. 2c and 3c). There is significant negative correlation between silicate and DO at 2000 m ($r = 0.89$, $P < 0.003$). We attribute this trend to the

fact that seawater at this depth in the eastern region is older than that in the western region because seawater in this layer flows from west to east in the northern North Pacific [25].

3.2. Distributions of Physical Parameters in the Surface Layer (0-200 m)

Potential temperature above 200 m depth tended to increase slightly from west to east (Fig. 4a). The potential temperature above 30 m depth was 10.3-11.8 °C. There was clear temperature minimum layer around 100 m depth in the western part of the study area, and this layer became less distinct from west to east. At OSP, the temperature minimum layer was unclear and potential temperature declined with increasing depth.

Salinity above 50 m depth decreased slightly from west (salinity = 32.9) to east (32.4), and was lowest in the study area at station EW7 (Fig. 4b). At 150 m, salinity at station EW7 (33.64) was higher than at the other stations (<33.48).

The distribution of potential density above 30 m depth showed to a trend of higher values in the west and lower in the east ("west high, east low" [WHEL]; Fig. 4c). The surface mixed layer depth, defined by a criterion of a 0.125 kg m^{-3} potential density difference from that of the surface [26], tended to increase from west (about 39 m) to east (50 m) in the study area (Fig. 4c). At 100 m depth, the potential density at OSP (25.70) was substantially lower than those at the other stations (26.14-26.58).

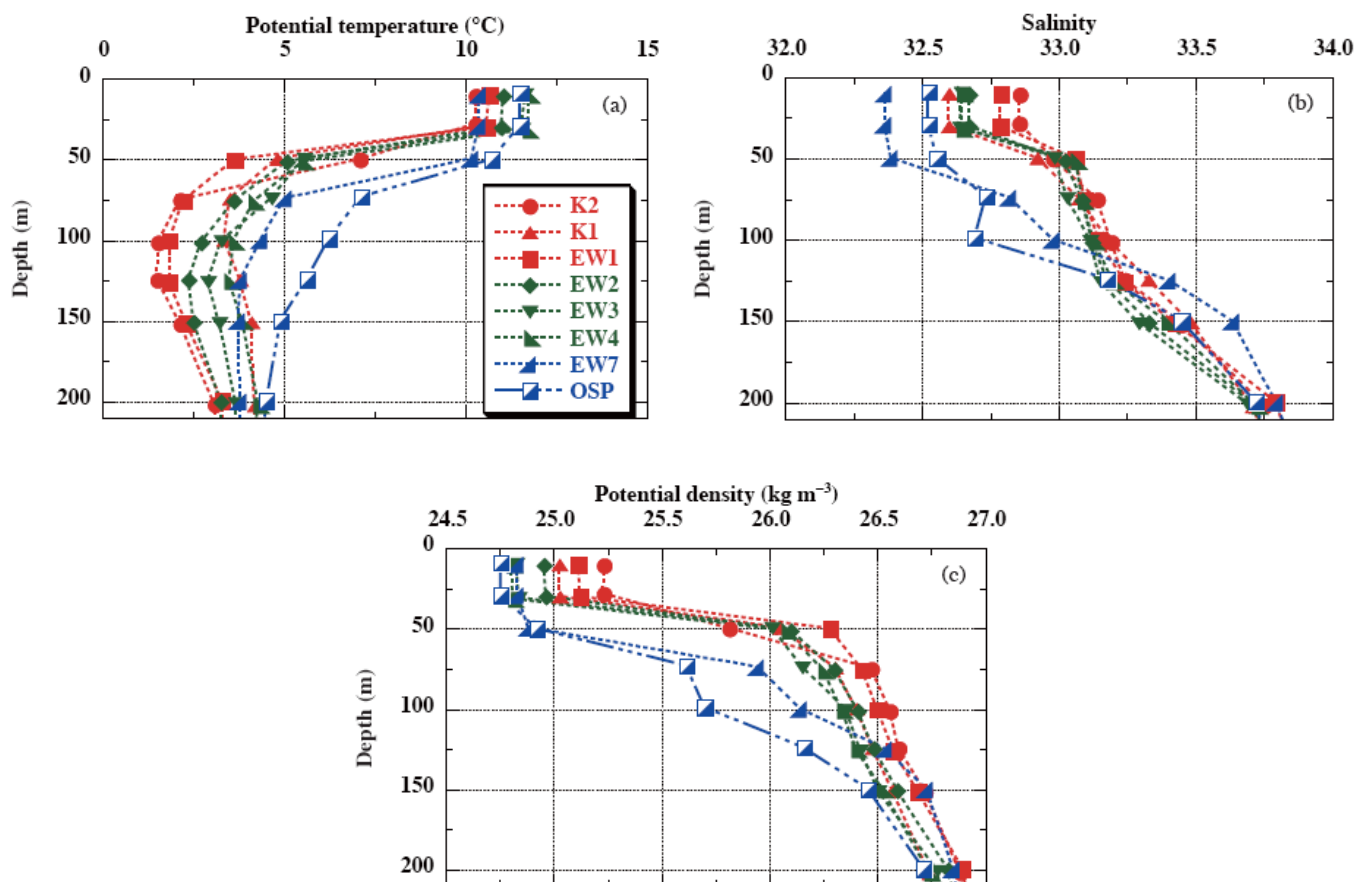


Fig. (4). Vertical distribution of potential temperature (a), salinity (b), and potential density (c) in the surface layer (10-200 m) at all stations, September-October 2005.

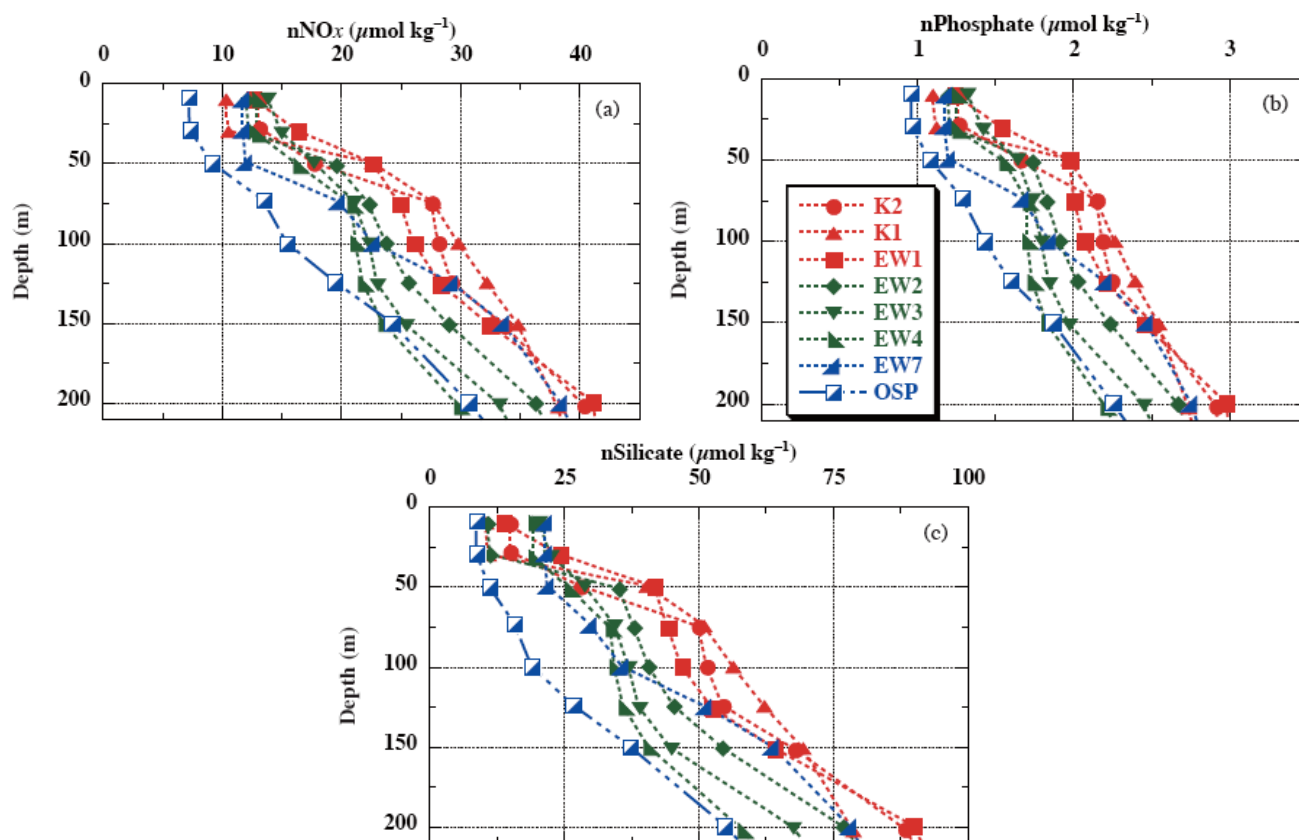


Fig. (5). Vertical distribution of normalized (to salinity of 33) NO_x ($n\text{NO}_x$) (a), phosphate ($n\text{Phosphate}$) (b), and silicate ($n\text{Silicate}$) (c) in the surface layer (10-200 m) at all stations, September-October 2005.

3.3. Distribution of Nutrient, DIC, TALK, DO, and AOU in the Surface Layer

Nutrients (NO_x , phosphate, and silicate), DIC, and TALK were normalized to a constant salinity of 33 to remove the effects of evaporation and precipitation. The distribution of normalized NO_x ($n\text{NO}_x$) showed the general trend of WHEL above 200 m depth (Fig. 5a). $n\text{NO}_x$ above 30 m depth ranged from 7 to 13 $\mu\text{mol kg}^{-1}$ (Fig. 5a). At 150 m depth, $n\text{NO}_x$ concentrations at stations EW3, EW4, and OSP (approximately 24 $\mu\text{mol kg}^{-1}$) were lower than those at the other stations (approximately 33 $\mu\text{mol kg}^{-1}$). The other nutrients showed similar trends (Figs. 5b and c).

Distribution of normalized DIC ($n\text{DIC}$) generally followed the WHEL trend above 200 m depth (Fig. 6a), similar to nutrients. $n\text{DIC}$ above 30 m depth ranged from 2027 to 2045 $\mu\text{mol kg}^{-1}$ (Fig. 6a). Normalized TALK ($n\text{TALK}$) showed vertically constant profiles at all stations (Fig. 6b).

Above 100 m depth, DO concentrations showed little variation with depth or between stations (Fig. 7a). Below 100 m depth, DO decreased with increasing depth and the differences in DO concentrations between stations were larger than those above 100 m depth. Values for AOU above 50 m depth were usually negative (*i.e.*, water was

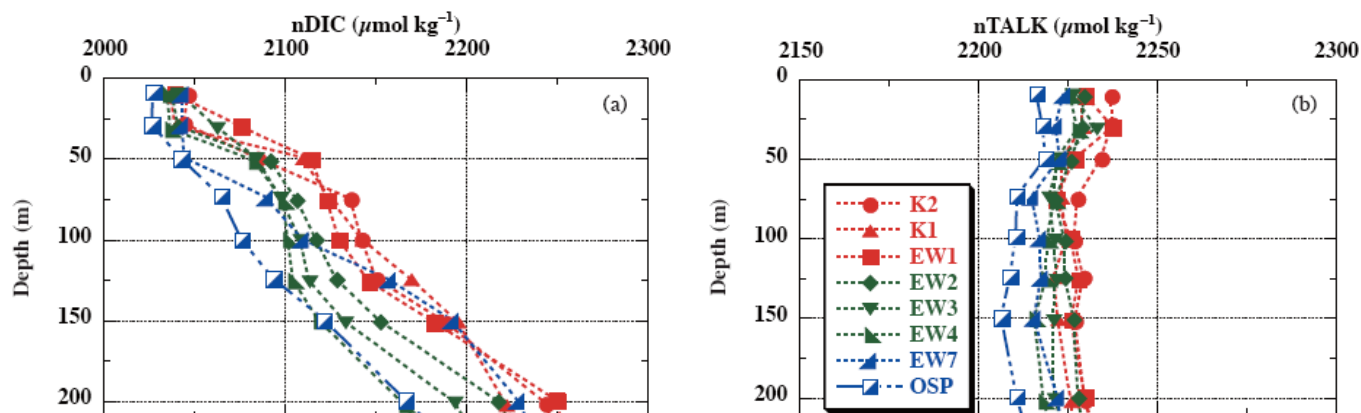


Fig. (6). Vertical distribution of normalized (to salinity of 33) DIC ($n\text{DIC}$) (a) and TALK ($n\text{TALK}$) (b) in the surface layer (10-200 m) at all stations, September-October 2005.

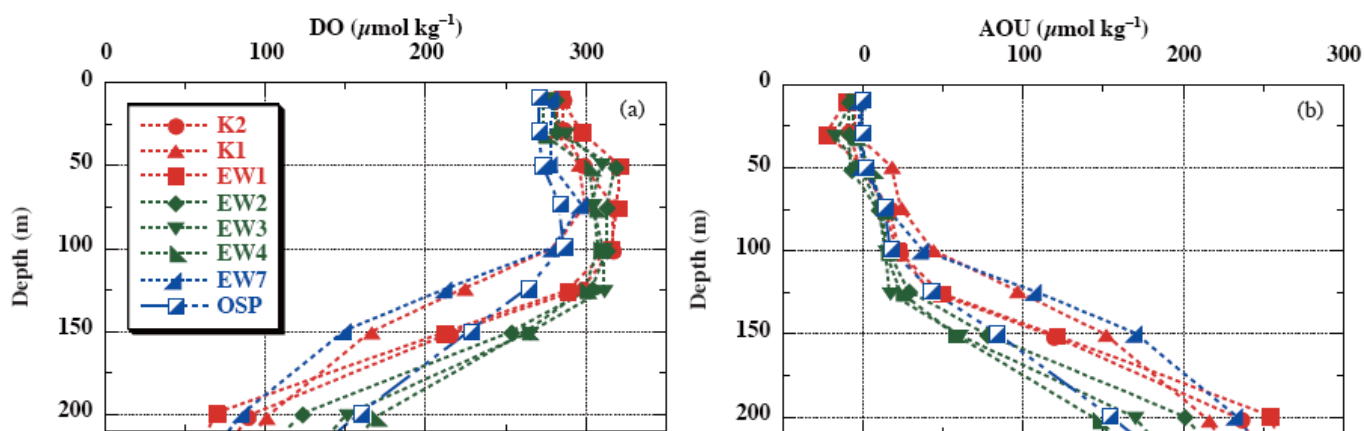


Fig. (7). Vertical distribution of dissolved oxygen (DO) (a) and apparent oxygen utilization (AOU) (b) in the surface layer (10-200 m) at all stations, September-October 2005.

supersaturated with respect to atmospheric oxygen; Fig. 7b). It is possible that these negative AOU values resulted from photosynthesis by phytoplankton. AOU increased with depth. At most stations, the rates of AOU increase with depth were different above and below 100 m depth. The consumption of oxygen during the remineralization of POM was apparent below 100 m depth.

3.4. East-West Distributions of Seasonal Drawdown of DIC and Nutrients

From liner regression analysis of nDIC, normalized nutrient concentrations, and AOU in the depth range of 100-200 m at all stations, we estimated the average regeneration ratio of C : N : Si : P : O₂ at $(116 \pm 4) : (16.2 \pm 0.2) : (44.7 \pm 0.8) : 1 : (-172 \pm 17)$ in the northern North Pacific. The concentrations of nDIC or normalized nutrients in the surface mixed layer in winter (A_w) were estimated by subtracting the regenerated fraction of each component from its concentration at 100 m (A_{100}) in autumn as follows [27]:

$$A_w = A_{100} - R_{i/O_2} \Delta AOU_{100}, \quad (1)$$

where R_{i/O_2} is the regeneration ratio for each component (i) and ΔAOU_{100} is the difference between AOU at 100 m depth in winter and the observed AOU. AOU at 100 m in winter was defined as $15 \mu\text{mol kg}^{-1}$ which is the average value in early April in the Western Subarctic Gyre and at OSP [8, 28].

The seasonal drawdown of DIC or nutrient concentrations in the surface mixed layer (ΔA) was calculated from the following equation:

$$\Delta A = \int_0^{100} (A_w - A_z) dz \quad (2)$$

where A_z represents the nDIC or normalized nutrient concentration in each layer above 100 m depth. ΔA represents the column-integrated drawdown of DIC or nutrients due to biological activity and/or air-sea CO₂ exchange in the surface mixed layer from winter to autumn. Seasonal drawdowns were integrated from sea surface to 100 m depth. A depth of 100 m was chosen for five reasons: 1) it was close to the temperature minimum layer at most stations (Fig. 4a), 2) DO above 100 m depth was almost constant with depth and at all stations (Fig. 7a), 3) it was the depth reached by the mixed layer in winter [1, 8], 4) it was the

depth at which the concentrations of DIC and nutrients were close to those at the sea surface in late winter [8, 29], and 5) decreases in DIC and nutrients from spring to autumn occurred only above 100 m [8, 29]. Furthermore, column-integrated drawdowns estimated from measurements taken during this season (September and October) were considered comparable to the annual drawdowns [1, 8].

The trend in seasonal drawdown of nDIC and normalized nutrients (ΔC , ΔN , ΔP , and ΔSi) above 100 m depth from winter to autumn was generally WHEL (Figs. 8a-d). This trend is comparable with the results of previous studies [9, 17]. However, seasonal drawdowns at station EW1 were smaller than those at the neighboring stations. It is possible that the small seasonal drawdowns at station EW1 resulted from the slightly higher nDIC and normalized nutrients levels at 30-50 m depth at this station compared with those at the other stations (Figs. 5 and 6a). At station K2 and OSP, seasonal net community production from winter to autumn in the surface layer as estimated from ΔN and the Redfield ratio (C : N = 106 : 16 [30]) was 57 and 36 gC m⁻², respectively. These results are comparable to previous values [2, 8-10].

Ratios of the seasonal drawdown of normalized silicate to that of nNO_x ($\Delta Si/\Delta N$) showed the WHEL trend (Fig. 8e). Diatoms have high Si/N ratios (1.1-3.0) [31-33], and the distribution of diatoms in the northern North Pacific follows the WHEL trend [29, 34]. In addition, dissolved iron concentrations in the surface layer in the northwestern North Pacific were nearly twice those in the northeastern North Pacific [35, 36]. Dissolved cobalt and nickel—which, like iron, are micronutrients [37]—were also found at higher concentrations in the western part of the northern North Pacific compared to the eastern part [38, 39]. Consequently, we suspected that the horizontal variations in trace metal concentrations influenced the horizontal variations in phytoplankton species composition and, further, in the strengths of the biological pump in the northern North Pacific Ocean.

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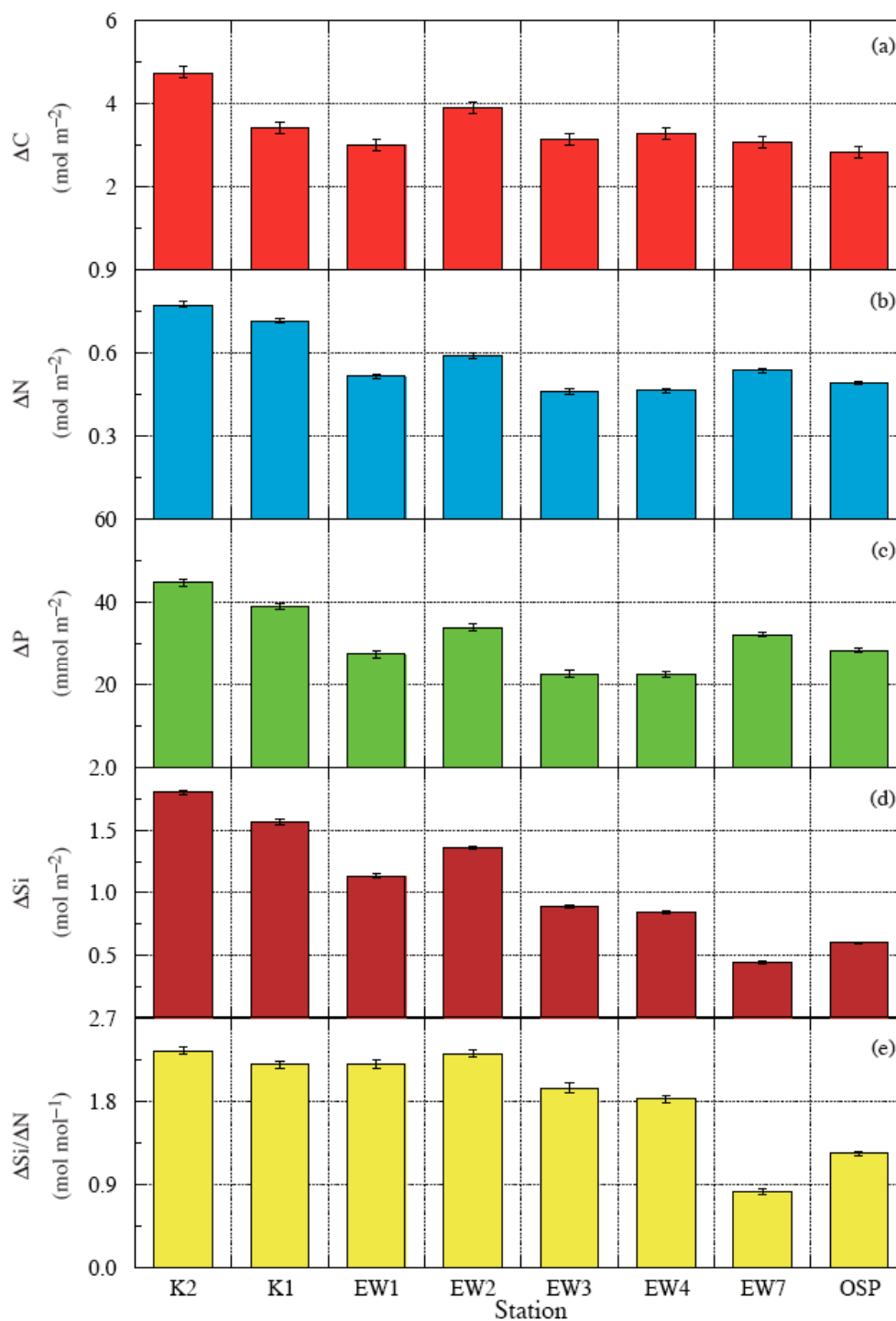


Fig. (8). East-west distributions of seasonal drawdown of DIC and nutrients (ΔC , ΔN , ΔP , and ΔSi) (a-d) and $\Delta Si/\Delta N$ ratio (e) above 100 m depth from winter to autumn in the northern North Pacific, September-October 2005. The error bars indicate the precision (1σ) of each measurement.

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SUPPLEMENTARY MATERIAL

Dataset of physical and chemical parameters at stations K2, K1, EW1, EW2, EW3, EW4, EW7, and OSP in September and October 2005. The data have not been corrected for systematic errors among stations.

REFERENCES

- [1] Harrison PJ, Boyd PW, Varela DE, Takeda S, Shiomoto A, Odate T. Comparison of factors controlling phytoplankton productivity in the NE and NW subarctic Pacific gyres. *Prog Oceanogr* 1999; 43: 205-34.
- [2] Wong CS, Waser NAD, Whitney FA, Johnson WK, Page JS. Time-series study of the biogeochemistry of the North East subarctic Pacific: reconciliation of the C_{org}/N remineralization and uptake ratios with the Redfield ratios. *Deep-Sea Res II* 2002; 49: 5717-38.

- [3] Wakita M, Watanabe S, Watanabe YW, Ono T, Tsurushima N, Tsunogai S. Temporal change of dissolved inorganic carbon in the subsurface water at station KNOT (44°N, 155°E) in the western North Pacific subpolar region. *J Oceanogr* 2005; 61: 129-39.
- [4] Saino T, Bychkov A, Chen CA, Harrison PJ. The Joint Global Ocean Flux Study in the North Pacific. *Deep-Sea Res II* 2002; 49: 5297-301.
- [5] Honda MC, Kawakami H, Sasaoka K, Watanabe S, Dickey T. Quick transport of primary produced organic carbon to the ocean interior. *Geophys Res Lett* 2006; 33: 10.1029/2006GL026466.
- [6] Honda MC, Watanabe S. Utility of an automatic water sampler to observe seasonal variability in nutrients and DIC in the northwestern North Pacific. *J Oceanogr* 2007; 63: 349-62.
- [7] Kawakami H, Honda MC. Time-series observation of POC fluxes estimated from ²³⁴Th in the northwestern North Pacific. *Deep-Sea Res I* 2007; 54: 1070-90.
- [8] Kawakami H, Honda MC, Wakita M, Watanabe S. Time-series observation of dissolved inorganic carbon and nutrients in the northwestern North Pacific. *J Oceanogr* 2007; 63: 967-82.
- [9] Tsurushima N, Nojiri Y, Imai K, Watanabe S. Seasonal variations of carbon dioxide system and nutrients in the surface mixed layer at station KNOT (44°N, 155°E) in the subarctic western North Pacific. *Deep-Sea Res II* 2002; 49: 5377-394.
- [10] Honda MC. Biological pump in northwestern North Pacific. *J Oceanogr* 2003; 59: 671-84.
- [11] Saito H, Tsuda A, Kasai H. Nutrient and plankton dynamics in the Oyashio region of the western subarctic Pacific Ocean. *Deep-Sea Res II* 2002; 49: 5463-486.
- [12] Yamaguchi A, Watanabe Y, Ishida H, *et al.* Structure and size distribution of plankton communities down to the greater depths in the western North Pacific Ocean. *Deep-Sea Res II* 2002; 49: 5513-30.
- [13] Tsunogai S, Noriki S. Particulate fluxes of carbonate and organic carbon in the ocean. Is the marine biological activity working as a sink of the atmospheric carbon? *Tellus* 1991; 43B: 256-66.
- [14] Kemp AES, Pike J, Pearce RB, Lange CB. The "Fall dump" - a new perspective on the role of a "shade flora" in the annual cycle of diatom production and export flux. *Deep-Sea Res II* 2000; 47: 2129-154.
- [15] Smetacek VS. The giant diatom dump. *Nature* 2000; 406: 574-75.
- [16] Honda MC, Imai K, Nojiri Y, Hoshi F, Sugawara T, Kusakabe M. The biological pump in the northwestern North Pacific based on fluxes and major components of particulate matter obtained by sediment-trap experiment (1997-2000). *Deep-Sea Res II* 2002; 49: 5595-625.
- [17] Wong CS, Waser NAD, Nojiri Y, Whitney FA, Page JS, Zeng J. Seasonal cycles of nutrients and dissolved inorganic carbon at high and mid latitudes in the North Pacific Ocean during the *Skaugran* cruises: determination of new production and nutrient uptake ratios. *Deep-Sea Res II* 2002; 49: 5317-338.
- [18] Zeng J, Nojiri Y, Murphy PP, Wong CS, Fujinuma Y. A comparison of $\Delta p\text{CO}_2$ distributions in the northern North Pacific using results from a commercial vessel in 1995-1999. *Deep-Sea Res II* 2002; 49: 5303-315.
- [19] Fujiki T, Matsumoto K, Honda MC, Kawakami H, Watanabe S. Phytoplankton composition in the subarctic North Pacific during autumn 2005. *J Plank Res* 2009; 31: 179-91.
- [20] Kawakami H, Honda MC, Matsumoto K, Fujiki T, Watanabe S. East-west distribution of POC fluxes estimated from ²³⁴Th in the northern North Pacific in autumn. *J Oceanogr*; accepted.
- [21] Dickson AG, Goyet C. Handbook of methods for the analysis of the various parameters of the carbon dioxide system in seawater, Version 2. 1994.
- [22] Dickson AG, Afghan JD, Anderson GC, Eds. Sea water based reference materials for CO₂ analysis: 2: A method for the certification of total alkalinity, unpublished, 2002.
- [23] Dickson AG, Ed. Determination of dissolved oxygen in sea water by Winkler titration: WHP Operations and Methods. WHP Office Report Woods Hole, Massachusetts, USA 1994; 1-14.
- [24] Weiss RF. The solubility of nitrogen, oxygen and argon in water and seawater. *Deep-Sea Res* 1970; 17: 721-35.
- [25] Reid JL. On the total geostrophic circulation of the Pacific ocean: flow patterns, tracers, and transports. *Prog Oceanogr* 1997; 39: 263-352.
- [26] de Boyer Montégut C, Madec G, Fischer AS, Lazar A, Iudicone D. Mixed layer depth over the global ocean: An examination of profile data and a profile-based climatology. *J Geophys Res* 2004; 109: C12003. doi:10.1029/2004JC002378.
- [27] Midorikawa T, Umeda T, Hiraishi N, *et al.* Estimation of seasonal net community production and air-sea CO₂ flux based on the carbon budget above the temperature minimum layer in the western subarctic North Pacific. *Deep-Sea Res I* 2002; 49: 339-62.
- [28] Steiner N, Vagle S, Denman K, McNeil C. Oxygen and nitrogen cycling in the northeast Pacific - simulations and observations at Station Papa in 2003/2004. *J Mar Res* 2007; 65: 441-69.
- [29] Varela DE, Harrison PJ. Seasonal variability in nitrogenous nutrition of phytoplankton assemblages in the northeastern subarctic Pacific Ocean. *Deep-Sea Res II* 1999; 46: 2505-538.
- [30] Redfield AC, Ketchum BH, Richards FA. In: Hill MH, Ed. The Sea: The influence of organisms on the composition of seawater. Wiley-Interscience, New York 1963; 26-77.
- [31] Brzezinski M. The Si/C/N ratio of marine diatoms: interspecific variability and the effect of some environmental variables. *J Phycol* 1985; 21: 347-57.
- [32] Hutchins DA, Bruland KW. Iron-limited diatom growth and Si/N uptake ratios in a coastal upwelling regime. *Nature* 1998; 393: 561-4.
- [33] Takeda S. Influence of iron availability on nutrient consumption ratio of diatoms in oceanic waters. *Nature* 1998; 393: 774-7.
- [34] Mochizuki M, Shiga N, Saito M, Imai K, Nojiri Y. Seasonal changes in nutrients, chlorophyll *a* and the phytoplankton assemblage of the western subarctic gyre in the Pacific Ocean. *Deep-Sea Res II* 2002; 49: 5421-39.
- [35] Martin JH, Gordon RM, Fitzwater SE, Broenkow WW. VERTEX: phytoplankton/iron studies in the Gulf of Alaska. *Deep-Sea Res* 1989; 36: 649-80.
- [36] Nakabayashi S, Kusakabe M, Kuma K, Kudo I. Vertical distributions of iron (III) hydroxide solubility and dissolved iron in the northwestern North Pacific Ocean. *Geophys Res Lett* 2001; 28: 4611-14.
- [37] Bruland KW, Donat JR, Hutchins DA. Interactive influences of bioactive trace metals on biological production in oceanic waters. *Limnol Oceanogr* 1991; 36: 1555-77.
- [38] Fujishima Y, Ueda K, Maruo M, *et al.* Distribution of trace bioelements in the subarctic North Pacific and the Bering Sea (the R/V Hakuho Maru cruise KH-97-2). *J Oceanogr* 2001; 57: 261-73.
- [39] Ezoe M, Ishita T, Kinugasa M, Lai X, Norisuye K, Sohrin Y. Distributions of dissolved and acid-dissolved bioactive trace metals in the North Pacific Ocean. *Geochem J* 2004; 38: 535-50.

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