The Northeast Greenland shelf as a potential late-summer CO₂ source to the atmosphere

Esdoorn Willcox¹, Marcos Lemes¹, Thomas Juul-Pedersen³, Mikael Kristian Sejr⁴, Johnna Michelle Holding⁴, and Søren Rysgaard²

¹Centre for Earth and Observation Science, University of Manitoba, Manitoba, Canada

²Arctic Research Centre, Aarhus University, Aarhus, Denmark

³Pinngortitaleriffik, Greenland Institute of Natural Resources, Kivioq 2, PO Box 570, 3900 Nuuk, Greenland

⁴Institut for ecoscience, Aarhus University, Aarhus, Denmark

Abstract. The Northeast Greenland shelf is a region currently considered to be an annual net sink of carbon dioxide (CO_2) from the atmosphere. Water from the Northeast Greenland shelf is advected to the formation regions of North Atlantic Deep Water and therefore any carbon uptake may be stored for ocean thermohaline circulation timescales. We present the most extensive study of carbon chemistry on the Northeast Greenland shelf to date made possible by opportunistic sampling due to

- 5 a sudden decrease in sea ice concentration in late August and September 2017. These are the first full-depth measurements of total alkalinity and dissolved inorganic carbon at latitudes between 75 and 79 °N with additional data collected in the region of the Northeast Water Polynya and outside Young Sund. We find that surface mixed layer concentrations are variable and for many stations higher than the interpolated atmospheric concentration for the region during the sampling period. Below the surface mixed layer, CO₂ concentrations increase linearly with decreasing apparent oxygen utilisation. The mixed layer
- 10 deepens during the study period which is associated with apparent changes in CO_2 uptake. The Northeast Greenland shelf is a hydrologically complex region with many processes influencing the carbonate system at smaller scales than our sampling density. The scatter in the dataset are more than mere outliers and their lack of relationship to any measured variable indicates a strong influence of currently undescribed process(es) or variable(s) at the sampled scales. These data were collected during a time of radically low sea ice concentrations for the region and may be an indication of future conditions. Since they indicate the
- potential of the region to act as a seasonal source of CO_2 to the atmosphere this may modify our current estimate of the region as a strong annual net sink relatively protected from the immediate influence of atmospheric warming and climate change.

1 Introduction

The Arctic Ocean and adjecent continental shelves are changing rapidly under the influence of climate change (????). The Northeast Greenland shelf is an Arctic outflow shelf (??) and one of the two gateways (the other the Canadian Arctic Archipelago)

20 through which water from the Arctic Ocean is transported southward into the North Atlantic Ocean (?). Together with the alongslope East Greenland Current (EGC), the shelf acts as a gateway through which water from the Arctic Ocean can be advected to the Greenland Sea, the Irminger Sea, and the Labrador Sea, regions that are crucial to the Atlantic Meridional Overturning Circulation through the formation of intermediate and deep water masses (?). This means that any carbon stored in the region may be retained in the global oceans on the timescales of the thermohaline circulation (??). The consensus is that the North-

- 25 east Greenland shelf has been a net annual carbon sink like other Arctic shelf regions, though this appears to be changing in response to changing conditions. The initial determination of the region as a sink was made through interpolation studies (?), Self Organising Maps (SOM, ?), and temporally and/or spatially limited observations using various methods, most focused on the Northeast Water Polynya or the near-coastal regions and fjords (???). Recent studies have indicated the potential for the region to become corrosive in terms of aragonite saturaton (?) and highlight the difference between the carbon system on
- 30 the eastern side of Greenland versus the west in terms of the relationship between carbon chemistry and depth (??). The latter is far more pronounced in western than eastern shelf areas and may be related to differences in their respective hydrography. Higher benthic production nearer to the shelf edge may be indicative of stronger primary productivity in this area, though the shelf is considered to be oligotrophic and the previously strong benthic-pelaic benthic-pelagic coupling in the region may be weakening (?). Due to high sea ice cover during all seasons the shelf is challenging to access, making it difficult to consistently

35 measure all the parameters required to determine the conditions and processes influencing dissolved CO_2 concentrations.

40

The northern North Atlantic and the Greenland Sea are more accessible and studies in these regions receiving water from the Northeast Greenland shelf and EGC (e.g. ?, ?) show that waters sourced from the Arctic remain undersaturated in dissolved CO_2 while Atlantic waters can act as a weak seasonal source. Water from the North Atlantic that might be entrained into the EGC also tends to be undersaturated (??), but has rapidly increasing concentrations, particularly in Autumn below latitudes of 78 °N when concentrations are at or near that of the atmosphere.

The uptake of CO_2 gas from the atmosphere in the northern North Atlantic (>50 °N) is partially driven by the cooling of warm water at the surface during northward transport which increases gas solubility, including CO_2 . High stratification and primary productivity in summer combined with deep convective mixing in winter enable the exposure of a more water to the atmosphere which further facilitates uptake. The Arctic Ocean carbon system is less well understood due to low (spatial and

- 45 temporal) sampling densities though the Eurasian Basin uptake of anthropogenic CO_2 is thought to be increasing (?). There are also additional processes at play in northern latitudes that influence CO_2 gas exchange, affecting both the solubility and biological pumps in the region such as the sea ice related processes of brine expulsion and sea ice melting, and the input of 10-11% of global meteoric river water (?). Each of these is characterised by their seasonality (??). The Atlantic Water being transported north into the Arctic (and into the EGC as part of the return Atlantic Current) is much warmer than it was in
- 50 previous decades (?). This increase in heat has been associated with changes in the Arctic halocline which shields sea ice from melting from below (?) and is likely to also stimulate heating in the EGC and potentially the Northeast Greenland shelf where the warm surface water from the return Atlantic Current (RAC) comes in direct contact with sea ice advected from the Arctic Ocean. Since the RAC is a surface current this energy is directly available for the melting of ice (icebergs, melange, and sea ice). While the melting of melange and icebergs merely reduces the temperature and freshen the surface water, changing the
- 55 gas solubility, melting sea ice can release ikaite ($CaCO_3 \cdot 6H_2O$) which facilitates additional CO_2 gas dissolution (?). Reduced sea ice cover is thought to facilitate primary productivity through enhanced availability of light, providing nutrients are also available (?) and rates of net primary productivity are thought to be increasing (?) even in the face of increasing stratification and the associated nutrient limitation (?). The Arctic Ocean surface waters are nutrient limited (?) and the regions of extreme

nitrate limitation are expanding, though primarily in the Western Arctic (??). As a result of surface water nutrient limitation

- 60 primary producers are generally found under sea ice (?), in the sea ice marginal zone, particular where there is upwelling (?), or as a 'deep chlorophyll maximum' (DCM) below the nitrogen depleted surface layer (? and references therein). Since the DCM is not directly in contact with the atmosphere the uptake by primary producers is not directly associated with drawdown from the atmosphere unless the strong stratification is broken and has a chance to equilibrate prior to sea ice freeze up. This equilibration needs to occur before the produced organic carbon is remineralised and before sea ice cover is extensive enough
- to form a barrier between ocean and atmosphere. This dominance of DCM may be a recent development. During the 1990s, primary productivity on the northern Northeast Greenland shelf was found near the surface in the Northeast Water Polynya and the required nutrients were associated with water from beneath the landfast and glacial ice (?). This led to the development of the 'seasonal rectification hypothesis' which describes strong uptake of atmospheric CO_2 during the sea ice melt season by primary producers followed by a season of inhibited autumn CO_2 release to the atmosphere by the development of an
- 70 extensive sea ice cover (?). Since then, the open water fraction in the region has changed dramatically as has the temperature of the Arctic river influenced Polar Water layer (?). In the summer of 2017 the Northeast Greenland shelf experienced a sudden drop in sea ice cover starting in August 2017 initiating a previously unseen decline in Arctic Ocean sea ice export which persisted throughout 2018 (?). These ice-free conditions allowed unprecedented access to previously unstudied parts of the Northeast Greenland shelf (Figure 1 a). The observations for this study were made opportunistically in these suddenly ice-free
- 75 waters and may offer some insight into the response of the CO_2 system on the Northeast Greenland shelf to an increasingly warm and ice-free Arctic.

2 Materials & methods

2.1 Cruise & hydrographical setting

Data for this study was collected during two cruises (DANA2017 and NEGREEN2017). The hydrography of the Northeast Greenland shelf during these cruises was described in our previous paper (?). To summarize briefly, several water types were found to be superimposed on much of the shelf albeit in different ways in different geographical areas. The hydrography is dominated by freshwater from various Arctic Ocean sources with different total alkalinity (Figure 1a) down to the depths of the Eurasian Basin Atlantic Water (EBAW) and Return Atlantic Water (RAW) which have similar practical salinities of respectively 34.8 and 35. The freshwater is primarily sourced from the Russian Shelf (particularly the Laptev Sea) where vast amounts of riverine freshwater are introduced changing the salinity and surface water geochemical properties.

- 85 amounts of riverine freshwater are introduced changing the salinity and surface water geochemical properties. This water is further geochemically modified in the Siberian shelf seas prior to cross-Arctic transport as a result of shallow bathymetry combined with high winds and extensive polynyas adding a measurable denitrification signal (???) and changing the isotopic fractionation (?). Finally the surface water masses are advected off of the Siberian continental shelves and entrained into the Transpolar Drift (TPD). Once entrained into the TPD the annual sea ice freeze-melt cycle will continue to freshen the
- surface layer by the export of brine and dilution with meltwater. This process diverts the slope of the surface water from that between Atlantic Water and meteoric freshwater toward the sea ice melt end-member in both TA-S and δ^{18} O-S diagrams. A

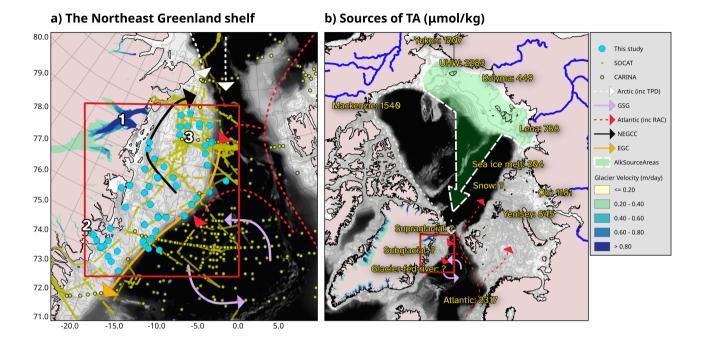


Figure 1. (a) Overview of carbon system chemistry and CO₂ fugacity (fCO₂) samples on the Northeast Greenland shelf. Arrows indicate known major currents. White indicates the advection of Arctic and Atlantie sourced water, including Return Atlantie Water (RAW), Polar Surface Water (PSW), and Eurasian Basin or Arctic Atlantic Water (EBAW/AAW). Red arrows indicate Atlantic sourced water, including Return Atlantic Water (RAW) which is transported by the Return Atlantic Current (RAC) from the east side of Fram Strait towards Northeast Greenland. Black is the Northeast Greenland Counter current (NEGCC) which transports water west then northward in a counterclockwise direction directly past the coast, purple is the Greenland Gyre, and orange is the East Greenland Current (EGC) which roughly follows the continental slope. SOCAT surface water fCO₂ measurement coordinates from ?, CARINA full depth carbon chemistry stations from ?. Numbers 1, 2, and 3 refer to the Northeast Greenland Ice Stream (culminating in 79N glacier or Nioghalvfjerdsbrae and Zachariae Isstrom), Young Sund, and the Northeast Water Polynya region respectively. (b) Known sources of total alkalinity to the Arctic Ocean highlighting the source regions of the Transpolar drift (green area with dashed white outline) of the Transpolar drift (TPD) and the location of the study area (red rectangle). Sources to the Arctic Ocean include Arctic rivers with variable catchment geology, sea ice and snow melt, and the Pacific Water coming in through the Bering Strait. River TA values from ?, Pacific from ?, and Atlantic from ?. Sea ice TA is from own measurements during these cruises ?. Locally, there is an unknown contribution of both sub- and supraglacial sources as well as glacier-fed rivers. Bathymetry was sourced from IBCAO (?), sea ice extent from OSTIA (?), and ice velocity from QGreenland v2 (?)

comparison between the Laptev Sea and Northeast Greenland shelf in terms of apparent oxygen utilisation (AOU) against the nutrients phosphate and silicate, and nitrate to phosphate ratio, confirm the strong link between the Laptev Sea and the Northeast Greenland Shelf via the TPD. The surface water, located above the maximum Brunt-Väisälä frequency squared

- 95 (N²), and above the remnant of the winter mixed layer, is almost entirely depleted in nitrogen (median NO₃⁻ = 0 μ mol/kg). Directly below this is a remnant of the winter mixed layer which exists at freezing temperatures and a practical salinity of ~ 31.4 psu ($\sigma_T \sim 25$). This is fresher than this inflection point was in previous decades (???). This layer contains, and apparently traps, the oxygen maximum indicating that it is not actively ventilated during the time when sampling occurred. From the salinity at the inflection point (and the oxygen maximum), there is a cold halocline layer which follows the freezing line up to
- 100 a salinity of 34.0, the Lower Halocline Water. At this point we find another inflection away from the freezing line with a sharp temperature increase in temperature to EBAW at 4 °C (S = 34.8) and AW at 6 °C (S = 35). The saltiest and warmest Atlantic Water found is likely sourced from the West-Spitsbergen Current. This water can be found at the surface just off the continental shelf and can make incursions onto the continental shelf, particularly further south. Surface conditions in terms of sea ice cover and surface temperature during the three weeks of the two cruises were variable (Figure 2) with warm surface temperatures and
- 105 patchy sea ice dominating the first part, after which the sea ice fraction increased, particularly toward the north and north-west part of the shelf.

2.2 Sample Analysis

Descriptions for the analysis of the A full description of the hydrography based on the Conductivity, Temperature, and Depth (CTD) instrument data, nutrients, and total alkalinity are included in the methods section of **?**. To analyse DIC, seawater samples were transferred from the CTD Rosette to gas-tight vials (12 mL Exetainer, Labco High Wycombe,UK), poisoned with 12 μ L solution of saturated HgCl₂, and stored in the dark at room temperature until analysis. DIC was measured on a DIC analyzer (Apollo SciTech, Newark, DE, USA) by acidification of a 0.75 mL subsample with 1 mL 10% H₃PO₄ (Sigma-Aldrich, Saint-Louis, MO, USA), and quantification of the released CO₂ with a nondispersive infrared CO₂ analyzer (LI-COR, LI-7000, Lincoln, NE, USA). Results were then converted from μ mol L⁻¹ to μ mol kg⁻¹ based on sample density, which was estimated from salinity and temperature. An accuracy of $\pm 2 \ \mu$ mol kg⁻¹ was determined for DIC from routine analysis of

certified reference material (A.G. Dickson, Scripps Institution of Oceanography, San Diego, CA, USA). CTD measurements of temperature and salinity were combined with the TA and DIC bottle data to calculate the pCO_2 using

the program CO2SYS (?) with the dissociation constants k1 and k2 of ? refitted by ? and the hydrogen sulfite dissociation constant from ?.

120 2.3 Mixed layer depth determination

We estimate the depth of the mixed layer by determining that of the pycnocline through estimating the pycnocline from the determination of the maximum Brunt-Väisälä frequency (N^2) (?) for all stations with bottles taken shallower than 120 m depth. Our previous study (?) indicated this would be a good proxy since the pycnocline acted as a barrier, trapping dissolved oxygen

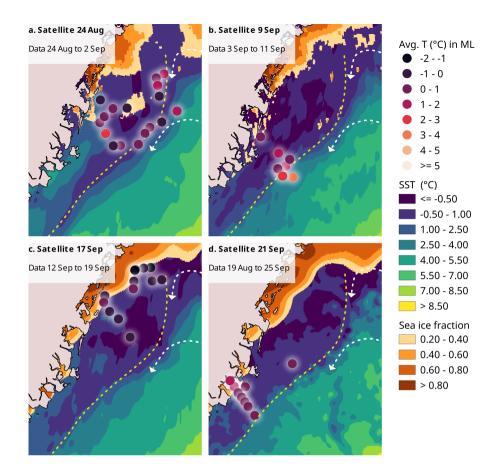


Figure 2. Surface conditions (sea ice fraction and sea surface temperature) on the shelf and average mixed layer depth temperature per station subdivided into four sampling periods. ESA sea surface temperature and sea ice fraction were obtained from ? (?). Average station mixed layer depth temperatures are the average temperature for all sampled depths above the maximum Brunt-Väisälä frequency squared (N^2)

below it indicating that this water was not ventilated during the period of our study. The maximum N^2 was calculated for each 125 CTD cast individually and depths varied between 1 and 30 m, with shallower depth closer to the coast and further north.

2.4 Normalisation of carbon chemistry bottle data

The bottle data were normalised by the application to the data of a fitted polynomial. The polynomial captures the effects of both the sea ice melt and <u>meteopric meteoric</u> freshwater dilution. A full justification, including a comparison with more traditional normalisation techniques, is provided in the <u>Supplement section 3 of the Supplement associated</u> with this manuscript.

Because the mean is heavily influenced by the extreme outliers in these data, parametric methods are not representative. Nonparametric methods relying on the median are more representative. The modified Z-score is one such method, it relies on the Mean Absolute Deviation (MAD). Data are marked as outliers when the modified Z-score is larger than a value D. Our choice of D (1.5) is discussed in section 4 of the Supplement.

135 3 Results and discussion

Based on previous studies, the region is expected to act as a sink for atmospheric CO_2 . Periods of high drawdown are specifically thought to occur when the light returns in spring allowing for autotrophic production during phytoplankton blooms including under ice blooms (??), and during upwelling events in the marginal ice zone (?). The release of CO_2 during the dark season, when no photosynthesis can occur and the region becomes (net) heterotrophic, is inhibited by extensive sea ice cover

- 140 (?). This ice-covered period can be associated with CO_2 supersaturation (?). Autumn is a transition period between a summer highly stratified environment where light is available and is dominated by sea ice and meteoric freshwater flux, and a winter environment that is dark, unproductive, and influenced by sea ice growth and brine rejection. In the northern North Atlantic, autumn is associated with the breakdown of stratification near the surface due to higher wind speeds and storms. This pattern is repeated in the Greenland Sea where average wind speeds tend to increase during the period of this study (days of year 240
- 145 256, ?), and August and September are associated with increasing concentrations of dissolved CO_2 after a seasonal low in July (?).

The fall of 2017 had exceptionally low sea ice cover for the region (?), allowing unpecedented access to undersampled regions of the shelf. The parameters which usually explain most of the variability in carbon dioxide fugacity (fCO_2) in the ocean surface are temperature (T), salinity (S), total alkalinity (TA), and dissolved inorganic carbon (DIC). Gas solubility is

- 150 expected to increase with decreasing temperature, change with salinity as a result of variable dissociation constants through their dependence on ion activities. With increasing TA, the fCO_2 is expected to decrease since these are the ions associated with increasing the ocean buffer capacity (?), and DIC is taken up by autotrophs during primary production and converted to organic matter. The data collected on the Northeast Greenland shelf in fall of 2017 do not clearly show the patterns expected (Figure 3). The data are scattered and outliers do not follow a discernable pattern with respect to salinity or temperature. Outliers occur
- 155 during both cruises, in measurements from both labs, toward high and low TA and DIC concentrations, and at different depths. There is no clear correlation between the outliers and any variable measured. We therefore have to surmise that at this time we are missing a (set of) variable(s) and/or process(es) with which to describe the extreme values in these data, and we do not have sufficient justification to remove any of the outliers from the dataset. We cannot discard any data without a good reason to flag it as an outlier, and with this amount of variability in the dataset using linear correlations loses some efficacy. Mean values are
- 160 not representative of the data therefore any attempt at statistical analysis necessarily relies on non-parametric techniques such as the modified Z-score.

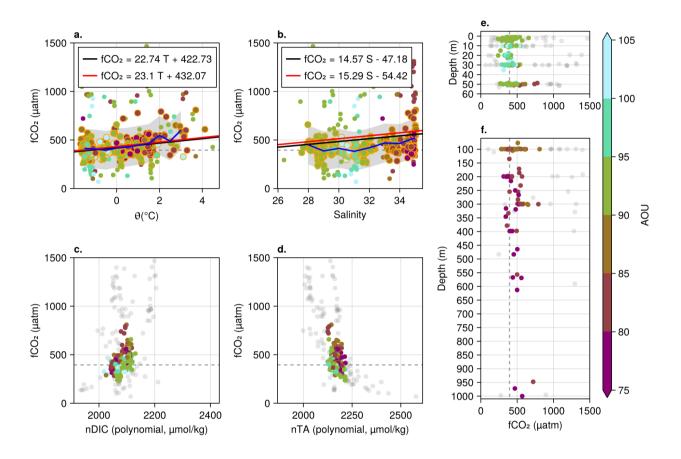


Figure 3. Carbon dioxide fugacity fCO_2 plotted as a function of potential temperature (a), practical salinity (b) normalised DIC (c), normalised TA (c), and with depth for shallower (e) and deeper waters (f). The orange lines in a,b are the best fit line for the median $\pm 200 \ fCO_2$ for each step in controlling variable. The median for steps in salinity is shown as the blue line where values included in the median calculation (± 200) are bounded by the grey region. Data with orange stroke in a,b and in colour in c,d,e,f are values with a modified Z-score of within D ± 1.5 . The red line in (a,b) is the modified Z-score data best fit. All data are coloured according to the Apparent Oxygen Utilization (AOU) calculated as the difference between the TEOS-10 calculated oxygen solubility and the observed oxygen concentrations.

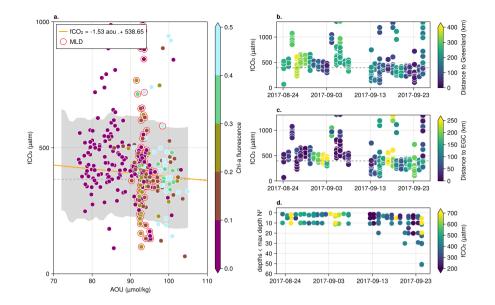


Figure 4. (a) the change in fCO_2 with changes in apparent oxygen utilisation (AOU). (b,c) fCO_2 changes with date, coloured respectively by distance of station to Greenland and the EGC (d) Data in the mixed layer depth (MLD) for each station by date

Using median values of fCO₂ for steps of each controlling variable (T, S, etc.) rather than the mean and picking values for fCO₂ between which the correlation is to be made or using an extreme modified Z-score outlier flag (> D=1.5), a linear relationship can be established for temperature (Figure 3a). The same method fails for salinity (Figure 3b) because the median fCO₂ follows a slightly polynomial shape which means the line is an overestimate compared to the values calculated from CO2SYS. The median and inter quartile range (iqr) for the mixed layer depth based on the N² are 410.49 and 147.58 μatm, which is above the projected atmospheric value for the region of 395 μatm based on SeaFlux (?) though this is for the entire time period which may not be representative (Figure 4). If we divide further by time, the period before 10 September has a median 477.66 with iqr 201.96 μatm and after this date the median goes down to 367.89 with iqr 110.66 μatm. This indicates
170 a change in conditions, either between the sampling period or the sampling locations where the region turns from a source to a sink. For samples taken near the surface, the apparent oxygen utilisation is under 95% indicating either its use in biochemical processes, or the active ventilation of or mixing with waters with even lower dissolved oxygen concentrations. In case of the former, this may also be responsible for the some of the higher concentrations in surface layer fCO₂ though we have no

175 One of the reasons that the relationships between fCO_2 and temperature and salinity respectively is unpredictable and highly variable is that water types with different histories found on the Northeast Greenland shelf can have similar end-member values for certain parameters. For example, meteoric freshwater from the longer fjords has had time to heat up before being advected onto the shelf, with air temperatures in summer as high as 10 to 12 °C (?). Atlantic Water (AW) from the return current also has temperatures of over 4 °C and a salinity of 35. Eurasian Basin Atlantic Water (EBAW) i.e. Arctic Atlantic Water that has

additional evidence to show active remineralisation.

- 180 circumnavigated the Eurasian Basin and lost heat is cold but has a salinity of 34.8. This is very close to AW salinity (34.8/35 = 0.99). The proportionality of the difference in TA between AW is similar to that in salinity, e.g. 0.98 for a EBAW TA of 2274 (?) and an AW TA of 2317 (?) but the 1% difference may be indicative of additional processes rendering TA non-conservative in this layer, diluted by a large volume. The region is also known for its small diameter (5 10 km) but deep penetrating eddies which offshelf can reach down to over 1000 m (???). With our sampling density which has distances between stations that are frequently over 30 km such features could create heterogeneous results for neighbouring station locations.
 - At depths below the surface mixed layer, as defined by the Brunt-Väisälä frequency squared (N²), the AOU and fCO₂ are inversely correlated (Figure 3e). This ranges from the remnant of the winter mixed layer which is supersaturated with respect to dissolved oxygen, AOU > 100 %, and the median fCO₂ is lower than atmospheric values at 383.39 with an iqr of 130.40 μ atm, to depths where AOU < 80 % and the fCO₂ has a median of 453.32 with an iqr of 119.61 μ atm. The maximum
- AOU corresponds with higher Chloropyll *a* fluorescence in the remnant winter mixed layer which indicates that the dissolved bioactive gas concentrations in this layer are at least partially driven by the presence of a Deep Chlorophyll Maximum (DCM) (Figure 4a). While surface conditions were variable during the sampling period in terms of sea ice fraction and temperature (Figure 2), the region off-shelf, to the east of the EGC, is generally associated with warmer temperatures and higher salinity, while waters across the shelf itself have colder surface temperatures. Sea ice is most persistent in the north. The first part of the
- 195 sampling period had warmer surface temperatures on the shelf itself, especially in the south along the coast . This is associated with higher , particularly at higher distances from the Greenland coast and smaller distances to the slope (EGC). (Figure 2a). In addition, this period (before 13th September) had more stations further from land (Figure 4b) and closer to the EGC (Figure 4c), and generally exhibit higher fCO₂.

During the sampling period, the surface temperature cools and the sea ice in the north becomes more consolidated (Figure

200 <u>2b,c,d</u>). The fCO₂ during the later period are much lower and trend below atmospheric saturation, potentially indicating a seasonal shift (Figure 4b,c). The increasing Mixed Layer Depth (MLD) near the end of the study (Figure 4d) could support this though this could also be attributable to another process such as the presence of a front. A comparison of mixed layer values ignoring temporal variability is included in section 6 of the Supplement.

The algorithms established by ? to determine TA and DIC for the North Atlantic (surface layer) fit our data well for TA, albeit with a lot of considerable scatter (Figure 5 a). This is not entirely surprising since the dataset used for the algorithm was in part obtained from measurements of the northern part of the Northeast Greenland shelf (?). To determine the best fit for DIC, they removed values for nearshore waters proximal to riverine meteoric freshwater sources from the dataset due to those measurements being lower than the algorithmically predicted values. Our measurements are also lower than the values predicted using their algorithm even though they are not directly near a meteoric freshwater source (Figure 5 b). Similar linear

210 regressions were fitted by ? and ?. The former do not provide an accurate reflection of our data and the second have a similar slope but a lower intercept. The ? equation is therefore the best predictor for TA on the shelf.

The difference between TA and DIC drives much of the fCO_2 variability calculated using CO2SYS, and increases (on average) between the first and second parts of the cruise (Figure 6a,b) as a result of a reduction in DIC. The average TA in

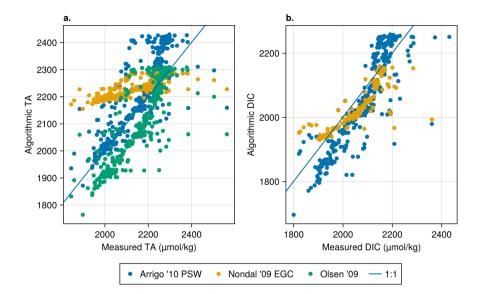


Figure 5. Measured concentrations of TA (a) and DIC (b) compared to values predicted using the algorithms from ?, ?, and ?. Since NO_3^- concentrations were only available for the last two weeks of the cruise, these are the only data shown for the ? fit in (b)

the mixed layer remains the same throughout the study period. The reduction in mixed layer DIC relative to mixed layer TA is most pronounced at the lower latitudes in the southernmost transect near Young Sound (Figure 6 c).

As previously described in ?, the depth-depedence of carbonate chemistry on the Northeast Greenland shelf is non-linear. Whether the surface mixed layer will act as a sink or a source of CO_2 with respect to the atmosphere seems to vary though it is clear that the region is not as strong a sink as previously expected and may be a net source. Increases in freshwater, both meteoric as well as sea ice melt, are associated with more corrosive surface waters near the coast in the region (?) but this can be compensated for by high productivity stimulated by nutrient input from local ice melt (??). If this is the case this may be

- 220 be compensated for by high productivity stimulated by nutrient input from local ice melt (??). If this is the case this may be another reason for the extreme variability of results we obtained. The mixed layer at the surface on the Northeast Greenland shelf that is advected in from the Arctic Ocean is already severely nitrogen depleted (?) which impacts opportunities for local primary producers to exist at the surface away from areas where local features such as eddies or actively melting sea ice might contribute nutrients to the surface water. Where sea ice melting, glacier melting, or potentially even iceberg fertilisation
- contribute nutrients to the surface, primary productivity can be quickly stimulated and the associated removal of DIC would allow for increased buffering by the TA and result in a lower fCO_2 in these areas. Sea ice and iceberg melt can be patchy and major continental meteoric freshwater contributions directly onto the shelf happen primarily at the termini of the 79N glacier and Zachariaea Isstrom, therefore the extent of surface primary productivity influencing the carbon system is likely limited during the sampling period. The rest of the shelf receives local freshwater input from long fjords where all the nutrients added in the surface have likely already been fully utilised before they reach the shelf (?). This means that primary productivity is
- necessarily limited to a deep chlorophyll maximum (DCM) below the nitracline. During years of more extensive sea ice cover

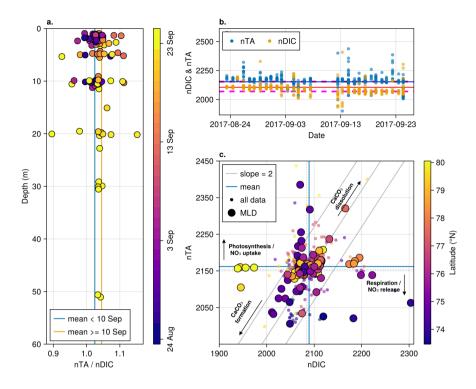


Figure 6. (a) Mixed layer nTA/nDIC with depth for the surface mixed layer. (b) nTA and nDIC respectively by measurement date. Blue line is TA and orange line DIC best fit between 24 Aug - 10 Sep. Dashed lines are for dates after 10 Sep. (c) nTA against nDIC

primary production may occur closer to the surface and stimulate more direct uptake of CO_2 from the atmosphere. Higher benthic productivity has been observed closer to the EGC which, with strong benthic-pelagic coupling in the region, indicates higher productivity at the surface near the slope (?) and an associated higher uptake of CO_2 . In the absence of sea ice melt it is possible that this is stimulated by along-shelf upwelling or by EGC-associated eddies which are particularly prominent in areas where the density of the warm Atlantic and the cool Polar Water are the same (?).

4 Summary

240

235

We present the first full depth carbon system observations of the area of the Northeast Greenland shelf between 75 and 79 °N, with additional measurements outside of Young Sund and in the region of the Northeast Water Polynya. Our total alkalinity (TA) measurements correspond well to the predictive algorithm created by **?**, whereas the dissolved inorganic carbon (DIC) measurements are lower than predicted by these authors. We find that the shelf does not act as a consistent sink as expected per the calculated fugacity of carbon dioxide (fCO₂) from samples of TA and DIC. Using non-parametric methods due to the large number of outliers in the dataset, we find that the surface of the region can act as either a sink or source of CO₂ with respect to the atmosphere. The highest uptake is associated with a maximum in apparent oxygen utilisation (AOU) and chlorophyll within

- the remnant of the winter mixed layer where there is both light and nutrient availability. This water is not actively ventilated and therefore cannot contribute directly to atmospheric carbon exchange. The middle of the study period saw an apparent breakdown in stratification based on an increase in mixed layer depth as determined by the maximum Brunt-Väisälä frequency squared (N²). This was associated with a reduction in surface layer fCO₂ to median values below the expected atmospheric concentration, apparently due to a corresponding reduction in DIC. The many outliers in fCO₂, particularly in the surface
- 250 mixed layer, are not clearly associated with any known process or measured variable. It is likely that the shelf is characterised by influences at smaller scales than the sampling density of this study. August and September 2017 were extraordinary in terms of low sea ice cover, which was the reason that opportunistic sampling of this previously unsampled area could take place. Our results may therefore not represent a baseline for the region when ice covered but rather may act as an example of the response of the region to future increases in oceanic and atmospheric heat and reductions in sea ice.

255 Data availability. The data for this study can be found at 10.1594/PANGAEA.968792 and 10.1594/PANGAEA.968793

Author contributions. The fieldwork component including taking samples from the CTD was performed by Thomas Juul-Pedersen, Johnna Michelle Holding, and Søren Rysgaard Marcos Lemes and Mikael Sejr performed the geochemical laboratory measurements for TA and DIC in their institutes respectively. Subsequent data analysis, writing of code, and initial drafting of the manuscript was performed by the primary author. Extensive feedback on first and second drafts of the manuscript was obtained from all co-authors.

260 Competing interests. The authors declare no competing interests.

Acknowledgements. We would like to thank our manuscript reviewers for providing detailed and relevant feedback and making a positive contribution to this manuscript. The cruises were funded through the Greenland Ecosystem Monitoring Program (Leg 1), the Danish Centre for Marine Research, the Natural Sciences and Engineering Research Council of Canada (NSERC), and the Independent Research Fund Denmark (G-Ice Project) (Grant 7014-00113B/FNU) (Leg 2). This study received financial support from the Aage V Jensens Foundation,
the Arctic Research Centre, Aarhus University, and Independent Research Fund Denmark (GreenShelf project (Grant 0135-00165B/FNU) to MSS). The captain and crew of RV DANA are acknowledged for excellent assistance during our field cruise to NE Greenland. Egon Frandsen is acknowledged for support on logistics and operations.