Nitrate isotope investigations reveal future impacts of climate change on nitrogen inputs and cycling in Arctic fjords: Kongsfjorden and Rijpfjorden (Svalbard)

Marta Santos-Garcia¹, Raja S. Ganeshram¹, Robyn E. Tuerena^{1, 2}, Margot C.F. Debyser¹, Katrine Husum³, Philipp Assmy³ and Haakon Hop³

¹School of Geosciences, University of Edinburgh, Edinburgh, EH9 3FE, United Kingdom ²Scottish Association for Marine Science, Dunstaffnage, PA37 1QA, United Kingdom ³Norwegian Polar Institute, Fram Centre, Tromsø, 9296, Norway

10 Correspondence to: Marta Santos-Garcia (M.SantosGarcia@ed.ac.uk)

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Abstract. Ongoing climate change in the Arctic has caused tidewater glaciers to retreat while increasing the discharge of freshwater and terrestrial material into fjords. This can affect both nutrient inputs and cycling within the fjord systems. In particular, tidewater glaciers and the presence of associated sub-glacial meltwater plumes can have a large impact on fjord circulation and biogeochemistry. In this study, we assess the influence of tidewater glaciers on nitrogen inputs and cycling in

- 15 two fjords in Svalbard during the summer using stable isotopic analyses of dissolved nitrate (δ^{15} N and δ^{18} O) in combination with nutrients and hydrographic data. Kongsfjorden receives inputs from tidewater glaciers, whereas Rijpfjorden mainly receives surface inputs from land-terminating glaciers. Results showed that both fjords are enriched in nutrients from terrestrial inputs. Nutrient ratios indicate excess Si and P relative to N. In both fjords, terrestrial nitrate from snowpack and glacier melting are identified as the dominant sources based on high δ^{18} O-NO₃⁻ and low δ^{15} N-NO₃⁻ of dissolved nitrate. In
- 20 Kongsfjorden, mixed-layer nitrate is completely consumed within the fjord system which we attribute to vigorous circulation at the glacial front influenced by the subglacial plume and longer residence time in the fjord. This is in contrast with Rijpfjorden where nutrients are only partially consumed perhaps due to surface river discharge and light limitation. In Kongsfjorden, we estimate terrestrial and marine N contributions to the nitrate pool from nitrogen isotopic values (δ^{15} N-NO₃⁻) and this suggests that nearly half the nitrate in the subglacial plume (50 ± 3 %) and the water column (44 ± 3 %) originates from terrestrial
- 25 sources. We show that terrestrial N contributes significantly to the regenerated N pool (63-88 %) within this fjord suggesting its importance in sustaining productivity here. Given this importance of terrestrial nutrient sources within the fjords, increase in these inputs due to climate change can enhance the fjord nutrient inventory, productivity and nutrient export offshore. Specifically, increasing Atlantification and warmer Atlantic Water will encourage tidewater glacier retreat and in turn increase surface discharge. In fjords akin to Rijpfjorden this is expected to foster more light limitation and less dynamic circulation,
- 30 ultimately aiding the export of nutrients offshore contributing to coastal productivity. Climate change scenario postulated for fjords such as Kongsfjorden include more terrestrial N-fuelled productivity and N cycling within the fjord, less vigorous circulation due to the retreat of tidewater glaciers and the expansion of oxygen depleted deep waters isolated by the sill.

Short Summary. Terrestrial sources of nitrate are important contributors to the nutrient pool in the fjords of Kongsfjorden and Rijpfjorden in Svalbard during the summer and they sustain most of the fjord primary productivity. Ongoing tidewater glacier retreat is postulated to favour light limitation and less dynamic circulation in fjords. This is suggested to encourage the export of nutrients to the middle and outer part of the fjord system, which may enhance primary production within and in offshore areas.

1 Introduction

- 40 In Arctic marine ecosystems, temperature anomalies of +2°C have led to increased discharge of freshwater (Beszczynska-Möller et al., 2012) and fluxes of carbon and nutrients across the land-ocean interface with profound implications for coastal ecosystems and biogeochemical cycling (McGovern et al., 2020). Nitrate (NO₃⁻) is the predominant form of fixed nitrogen (N) used by organisms in the ocean and is normally deemed as the limiting-nutrient in the Arctic Ocean (Yamamoto-Kawai et al., 2006), whereas phosphate and silica are also essential for algal growth in nearshore environments (Egge, 1998). In fjordic and
- 45 coastal settings in the Arctic, terrestrial inputs of N could be important (Kumar et al., 2018) and these inputs are changing with polar warming through a myriad of factors such as increased river discharge, permafrost thaw, soil N cycling and vegetation changes (Holmes et al., 2012).

The assessment of the impact of N fluxes, cycling and fate in a fjordic setting due to climate change is complicated by seasonally varying circulation patterns. Fjordic circulation responds to winter cooling and sea ice formation as well as summer

- 50 freshwater discharge. In turn, these processes also influence the nitrogen fluxes, cycling and ultimate fate. Circulation in fjords is restricted owing to its narrow shape and presence of a sill (Svendsen et al., 2002; Dürr et al., 2011). The ineffective tidal mixing, alongside freshwater influxes, contributes to the development of a very sharp halocline and stratified water column during the summer months (Geyer and Ralston, 2011; Monteban et al., 2020). Deep convective mixing occurs in the autumn due to cooling of surface water (thermal convection) and in the winter due to sea ice formation and brine release (haline
- 55 convection) (Cottier et al., 2007). Remnants of winter cooled waters can persist into the summer and are characterised by low temperatures and a wide salinity range reflecting their variable origin (De Rovere et al., 2022). In spring, temperature rises and ice break-up begins, leading to a reduction in salinity and the re-establishment of the strong summer pycnocline (Cottier et al., 2010). This pattern of fjordic circulation is subject to climate change on a pan-Arctic scale through increased freshwater inputs, glacial retreat as well as by warm marine waters entering the fjords.
- 60 In Svalbard, freshwater discharge is expected to increase by 200% before 2100 (RCP4.5 scenario, Adakudlu et al., 2019). Increased freshwater discharge, glacial retreat, snow and permafrost melt and reduced sea-ice cover are ongoing climate change trends in many parts of the Arctic (Arctic Climate Impact Assessment (ACIA), 2005). Increased freshwater inputs are expected to alter convective mixing in fjords possibly isolating deep waters below the sill. Enhanced freshwater inputs and stratification can impact both terrestrial N supply and cycling processes. On a pan-Arctic scale, increased river discharge is accompanied by enhanced N inputs (Holmes et al., 2012; McGovern et al., 2020). Therefore, it is hypothesised that (i) increasing freshwater
- by enhanced N inputs (Holmes et al., 2012; McGovern et al., 2020). Therefore, it is hypothesised that (1) increasing freshwater inputs will alter N supply and cycling leading to changes in the nutrient status and availability within the fjord and (ii) this will alter the exchange of nutrients between the fjord and offshore.

Arctic fjords are experiencing widespread glacial retreat (Kohler et al., 2007; Østby et al., 2017). Loss of tidewater glaciers have a significant impact on fjord circulation as they contribute to additional freshwater flux from glacial melt (Cowan, 1992; Ingvaldsen et al., 2001). At marine terminating glacier fronts, subglacial discharge enters the fjord at depth- specifically at the glacial grounding line during the summer (Cowton et al., 2015; Carroll et al., 2015, 2016). As a result of its lower density, the submarine glacial melt forms an upwelling plume enhancing vertical mixing by the entrainment of fjord waters (Everett et al., 2018; Halbach et al., 2019). The plumes are strongest at the glacier fronts, and decrease with increasing distance from land

75 due to dilution and mixing (Darlington, 2015; Hopwood et al., 2020). Moreover, upwelling plumes can entrain and elevate remnant winter cooled waters from the bottom of fjords, leading to the presence of a distinct, cold and relatively saline water mass throughout the water column close to the tidewater glacier front (Torsvik et al., 2019). Recent studies have shown that meltwater plumes not only play a prominent role in vertical mixing at glacier fronts, but also enhance sub-surface lateral mixing (Torsvik et al., 2019). Importantly, plumes also promote the transfer of heat from the ocean to the glacier front of large 80 Greenland fjords, thereby drawing in warmer waters from offshore (Straneo et al., 2010; Cowton et al., 2015). However, in the smaller fjord systems in Svalbard, this invigorated lateral circulation is mainly confined within the fjords (Torsvik et al., 2019).

Ongoing "Atlantification" has caused the increased prevalence of warm Atlantic Waters displacing cold Arctic coastal waters off Svalbard (Polyakov et al., 2017). This phenomenon has been linked to increases in sea water temperatures, shortened sea ice-covered period, altered freshwater inputs and enhanced light penetration (David and Krishnan, 2017; Hop et al., 2019; Hop and Wiencke., 2019). Although it is evident that Svalbard fjords are subject to both marine and terrestrial climate change, the net impact of these changes on N sources and cycling processes is currently unclear. The purpose of this study is to close this gap by conducting a detailed study of N sources and cycling using novel isotopic tools in two Svalbard fjords, allowing (i)

90 improved predictions of changes in the inventory of nutrients within sub-Arctic and Arctic fjords and, (ii) an assessment of the future changes in exchange of nutrients between the fjords and offshore areas. This is the first study to present a quantitative account of terrestrial contribution to the nitrate pool and primary productivity in Arctic fjords.

2 Methodology

95 2.1 Sampling sites

The sub-Arctic fjord of Kongsfjorden (79.0°N, 11.7°E) and Arctic Rijpfjorden (80.0°N, 22.3°E), located on the islands of Spitsbergen and Nordaustlandet, Svalbard, respectively (**Fig. 1**), are ideal locations to document changes in N dynamics affected by freshwater influxes and the retreat of local glaciers (**Fig. A1**). Important hydrographic features around Svalbard include the Fram Strait and the Barents Sea (**Fig. 1a**). The Fram Strait is one of the main gateways into and out of the Arctic

- 100 Ocean (Ilicak et al., 2016). Kongsfjorden samples were collected during the Norwegian Polar Institute Monitoring Cruises on RV *Lance* between 29 July and 2 August 2017 (NP2017) and from 13 and 15 July 2018 (NP2018). These samples were from four areas, namely near glacier fronts (Kb5-7, only sampled during NP2018 cruise), fjord (Kb0-3), continental shelf (V12) and continental slope (V10 and V6) (Fig. 1b).
- 105 Rijpfjorden was sampled between 3 and 5 August 2017 (NP2017) (**Fig. 1c-d**), where sample locations are divided into inner fjord (R1,2), outer fjord (R3), continental shelf (R4-5) and continental slope (R6-7B) stations. The inner fjord and the outer fjord basins are separated by a shallow sill at around 40 m depth.

2.2 Nutrient and isotopic analysis

- 110 Water samples were collected from Niskin bottles mounted on a rosette equipped with a CTD system recording conductivity, temperature and pressure. In addition, other parameters measured were salinity (PSU), chlorophyll *a* fluorescence (mg m⁻³) and oxygen (mL L⁻¹). Dissolved inorganic nutrient concentrations (nitrate, nitrite, silicate and phosphate) were analysed by the Norwegian Polar Institute (NP2017, NP2018) at the Institute of Marine Research. The samples were collected in 20 mL scintillation vials, fixed with 0.2 mL chloroform and stored refrigerated until sample analysis. Nitrite, nitrate, phosphate and
- 115 silicate were measured spectrophotometrically at 540, 540, 810 and 810 nm, respectively, on a modified Scalar autoanalyzer. The measurement uncertainty for nitrite was 0.06 mmol L⁻¹ and 10% or less for nitrate, phosphate and silicate. Water samples for isotopic analysis were prefiltered and frozen immediately after collection.

The denitrifier method for dual nitrogen and oxygen isotopes was used for the isotopic analysis of dissolved nitrate (Sigman et al., 2001, McIlvin and Casciotti, 2011). The denitrifier method takes advantage of the denitrifying bacteria *Pseudomonas*

aureofaciens with limited N-reductase activity which transform NO₃ to N₂O (Sigman et al., 2001; Casciotti et al., 2002). Nitrous oxide was extracted from 20mL-vial headspace by a Combi PAL auto-sampler and transported by a continuous flow of helium gas through a GasBench II coupled with a Delta V Advantage.

- Two standards, USGS 34 and IAEA N3, were used as reference for isotope ratio mass spectrometry (IRMS) analysis. δ^{15} N_{AIR} values of these standards were 1.8±0.2 (USGS 34) and 4.7±0.2 (IAEA N3), and δ^{18} Ovsmow values were -27.9±0.6 (USGS 125 34) and 25.6 \pm 0.6 (IAEA N3). Each standard was individually prepared at 2, 5, 15 and 30 μ mol concentrations. To overcome the discrepancies in δ^{18} O, solutions were prepared in milli-Q H₂O and low nutrient seawater respectively (McIlvin and Casciotti, 2011). Also, internal standards, from North Atlantic Deep Water, were used in order to represent average Atlantic Water δ^{15} N signature. This standard was run with each batch to check for inter-run comparability. IRMS analysis was carried
- 130 out at the University of Edinburgh using ISODAT 2.5 software. Isotopic measurements were determined relative to a reference peak.

Measurements of δ^{15} N-NO₃ and δ^{18} O-NO₃ were corrected to AIR and VSMOW, respectively, with the use of the correction scheme in Weigand et al. (2016) and following Tuerena et al. (2021a, 2021b) and the reference standards.

Averaged data reproducibility (1 σ) across all runs was 0.3 and 0.5 for δ^{15} N and δ^{18} O respectively, determined using internal standards with isotopic values of 4.8 \pm 0.2 % (δ^{15} N_{AIR}) and 1.8 \pm 0.6 % (δ^{18} O_{VSMOW}). These deviations can be approximated 135 to normal variability (0.2 and 0.6‰ for δ^{15} N and δ^{18} O, respectively).

Nutrient and isotopic data from JR17005 and FS2018 cruise were previously reported in Tuerena et al. (2021a) and Debyser et al. (2022) respectively. All nutrient and isotopic data are combined with hydrographical data obtained from a conductivitytemperature-depth rosette and processed using Matlab_R2020a software to document N dynamics.

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2.3 Data analysis, processing and visualisation

Cross-section figures of Kongsfjorden were designed using a global Topo15.1 bathymetry dataset with a spatial resolution of 4 km (Smith and Sandwell, 1997). Bathymetric cross-section figures of Rijpfjorden are based on data from the Norwegian Mapping Authority Hydrographic Service and IBCAO database version 4.1.

The degree of stratification ($\Delta \rho$, kg m⁻³) was calculated as the difference in potential density between 10 and 40 m depth. In addition, apparent oxygen utilisation (AOU, μ mol kg⁻¹) was computed using the equation AOU= $[O_2]_{saturated} - [O_2]_{seawater}$. Meanwhile, the oxygen saturation (%) was given by $[O_2]_{seawater}/[O_2]_{saturated} \times 100$.

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The semi-conservative tracers N* and Si* were calculated from inorganic nutrient concentrations where N* = NO_x - 16 × PO₄ + 2.9 (Gruber and Sarmiento, 1997) and Si* = SiOH₄ - NO_x (Sarmiento et al., 2004) are indicative of deviations from Redfield stoichiometry and 1:1 proportions, respectively. Negative N* values reflect N deficit and excess phosphate while positive N* values reflect excess nitrate relative to phosphate.

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Regression analyses were computed to ascertain the significance of observed linear trends (p-value ≤ 0.05). All statistical analyses were included in Appendix B (Table B1).

3 Results

160 3.1 Environmental setting and water mass characterisation

Based on the temperature (θ in °C) and salinity (S in PSU), three different water masses were identified in Kongsfjorden (13-15 July 2018), Rijpfjorden (3-5 August 2017) and their adjoining continental shelves and slopes following Pérez-Hernández et al. (2017). On the western side of Svalbard, warm and salty Atlantic water (AW; $\theta > 1$, S > 34.9) is carried northward along the shelf edge by the West Spitsbergen Current (WSC; **Fig. 1**) and enters the Arctic Ocean, supplying the continental slope at

- 165 Kongsfjorden (Figs. 2, A2) and other fjords along the way (Fig. 3; Cottier et al., 2005; Cokelet et al., 2008; Renner et al., 2018; Skogseth et al., 2020). These warm waters also fill intermediate depths across the whole Arctic Basin (known as Arctic Intermediate Water, AIW; $-1 \le \theta \le 1$, S > 34.9; Figs. 2, 3, A2) and represent an important supply pathway for nutrients (Beszczynska-Möller et al., 2012; Randelhoff et al., 2018). Occasionally, AW protrudes onto the shelf area, and thus there is an interplay between the intrusion of AW at depth and meltwater at the surface (Onarheim et al., 2014). This was the case at
- 170 Kongsfjorden, where Atlantic water dominance extended onto the continental shelf (via a trough called "Kongsfjordrenna"), as shown by salinity and temperature profiles above 40 m-depth, with salinity 34.9 PSU and temperature 5.6°C (**Fig. A3**). This intrusion was also associated with a chlorophyll *a* maximum of 4-6 mg m⁻³ (**Fig. A3**). This is in agreement with recent studies reporting AW intrusions into Kongsfjorden (6.5 mg m⁻³ chlorophyll maximum, Payne and Roesler, 2019).
- 175 Similarly, in the northern coast of Svalbard, the Atlantic influence remains substantial as AW travels eastward towards the Nansen Basin (Cokelet et al., 2008; Renner et al., 2018). However, fjords on the northern coast- including Rijpfjorden- have a wider continental shelf and are under a weaker AW influence compared to those on the western coast with a narrower continental shelf (Hop et al., 2019). For instance, the mouth of Rijpfjorden is 60 km away from the shelf, whereas that of Kongsfjorden is 45 km from the shelf (Howe et al., 2010). Rijpfjorden does experience occasional inflow of Atlantic-origin
- 180 water during summer to late autumn (Wallace et al., 2010; Hop et al., 2019). Although warmer water extends between 25 and 100 m depth at the continental shelf off Rijpfjorden (**Fig. 3a**), the temperature was not warm enough to be characteristic of AW ($\theta < 1$, **Fig. 3a**), nor the fluorescence peak (**Fig. 3c**) values were as high as those seen in the AW intrusion in Kongsfjorden (Payne and Roesler, 2019). Instead, the shelf at Rijpfjorden was dominated by Polar Surface Waters (PSW) (**Fig. A2**), which are a mixture of AW, river runoff, precipitation and ice melt (Rudels, 1989).

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Within the fjords, two variants of Polar Surface Water (PSW) were recognised, namely inshore PSW (iPSW) and warm PSW (PSWw) (**Fig. A2**). In detail, iPSW is a mixture of ice melt and AW (iPSW, $\theta \ge 2$, $S \le 34.9$) while PSWw ($0 \le \theta \le 2$, $S \le 34.9$) is a mixture between PSW and warm AW (Rudels et al., 2005; Cokelet et al., 2008). Kongsfjorden was dominated by iPSW, whereas Rijpfjorden was governed by PSW, and in both fjords the presence of PSWw was limited.

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Kongsfjorden and Rijpfjorden not only experience different degrees of AW influence and variations in PSW, but also have other contrasting physical characteristics which are relevant to N cycling. These characteristics include fjord size and geometry as well as the presence of tidewater glaciers, all of which in turn affect circulation and residence times. Kongsfjorden is significantly deeper (350 m) than Rijpfjorden (c.a. 200 m), although their mean depths are similar (100 m) and so are their

- 195 sizes (Kongsfjorden is 26 km long, 6-14 km wide; Rijpfjorden is 40 km long, 7-12 km wide). The average water residence time for Kongsfjorden is 172 hours and for Rijpfjorden this is expected to be much less (Torsvik et al., 2019; Yang et al., 2022). While Kongsfjorden experiences contributions of subglacial freshwater from five tidewater glaciers including Kongsbreen and Kronebreen (Fig. A1; How et al., 2017) and direct runoff from the Bayelva river, Rijpfjorden only receives surface freshwater input from relatively small glacially-fed rivers with short-lived summer flow as inferred from satellite
- 200 imagery (Wang et al., 2013). One important element of Kongsfjorden circulation is the subsurface discharge of meltwater from

subglacial plumes (Darlington, 2015). The meltwater plumes play a prominent role in vertical and lateral exchanges, and consequently act as a nutrient pump - as documented by numerous studies (Darlington, 2015; How et al., 2017; Schild et al., 2017; Halbach et al., 2019; Torsvik et al., 2019). Notably, the magnitude of the effect of subglacial discharge and the glacier front plume in Kongsfjorden varies spatially along the glacial front owing to topographical differences along the Kongsbreen

- 205 and Kronebreen glacier fronts. In particular, the Kongsbreen front is deeper and narrower than that at Kronebreen, and this constrains the lateral movement of water, thus this plume has a clearer vertical structure (Torsvik et al., 2019). In contrast, the Kronebreen front shows a less pronounced effect of the subglacial discharge and glacier front plume as lateral movement of water is unconstrained allowing mixing with adjacent fjord waters. Nonetheless, a topographic barrier below 35 m restricts lateral mixing below this depth along the Kronebreen transect (Torsvik et al., 2019). In Rijpfjorden, marine terminating glaciers
- 210 are absent and freshwater inputs from land occur at the surface. This surface input, the smaller size and limited marine influence due to the broader shelf leads to contrasting conditions between Rijpfjorden and Kongsfjorden.

The degree of stratification (Δρ) in the upper 40m of the water column increases landwards in both fjords (**Table A1**; **Figs. 2b**, **3b**) reflecting freshwater discharge. Notably, stratification was stronger in Rijpfjorden, particularly in the most offshore stations, compared to Kongsfjorden (**Table A1**). While in Kongsfjorden the freshwater layer was thicker in the proximity of the tidewater glacier fronts (**Fig. 2b**), in Rijpfjorden, freshwater was confined to a thin surface layer that extends further offshore (**Fig. 3b**; **Table A1**). It is also worth noting that in Kongsfjorden, AW seem to be drawn in at the continental shelf over the same depth range as the freshwater layer. This feature is attributed to glacier-induced lateral circulation set up by subglacial plume in fjords with tidewater glaciers (Svendsen et al., 2002; Cottier et al., 2005; Straneo et al., 2010; Cowton et al., 2015. The hear table 2010)

220 al., 2015; Tverberg et al., 2019).

Additionally, the cross-sections identified a fourth water mass, Winter Cooled Water (WCW, Cottier et al., 2005), in the deeper part of the fjord basins (c.a. >300 m at Kongsfjorden and c.a. >100 m at Rijpfjorden) characterised by low potential temperatures ($\theta < 1.1^{\circ}$ C at Kongsfjorden, Fig. 2a; $\theta < -1^{\circ}$ C at Rijpfjorden, Fig. 3a), high apparent oxygen utilisation (AOU)

- of > 40 µmol kg⁻¹ (Figs. 2d, 3d) and oxygen saturation of < 90% (Figs. 2e, 3e). Isolation and retention of such winter waters is attributed to reduced vertical exchange in the deep fjord basins, mainly due to restricted circulation owing to the presence of a sill at 200 m outside Kongsfjorden (in "Kongsfjordrenna") (Fig. 2) and at 25 m depth in Rijpfjorden (Cottier et al., 2010; Fig. 3). The high AOU characteristics of these water masses is indicative of their long residence time and isolation from the atmosphere (Svendsen et al., 2002) and as such these water masses are sensitive to hypoxia impacted by small changes in productivity, nutrient cycling and isolation time.</p>
 - 3.2 Nutrient concentrations and isotopic ratios

Depth profiles of temperature, salinity, chlorophyll *a* fluorescence, nutrient concentrations, N*, Si* and isotopic ratios in Kongsfjorden and Rijpfjorden are illustrated in Fig. 4a-j and Fig. 5a-j, respectively. In addition, cross-sections of nutrients,
the semi-conservative tracers N* and Si* and isotopes (δ¹⁵N-NO₃⁻ and δ¹⁸O-NO₃⁻) are also shown in Fig. 6a-g and Fig. 7a-g for Kongsfjorden and Rijpfjorden, respectively. In Kongsfjorden, temperature throughout the water column at the glacier front (Kb5-7, at Kronebreen and Kongsbreen transects) was strikingly lower than in the fjord and further offshore (Fig. 4a), indicating a distinct water mass formed at the front that cannot be traced offshore. It is suggested that these cold and saline waters are remnants of winter cooled waters resulting from heat loss to the atmosphere and contact with the glacier front

240 (Torsvik et al., 2019; De Rovere et al., 2022). Only at Kronebreen front (Fig. A1, station Kb5), temperatures increased to values similar to fjord temperatures at 20-35 m depth (Fig. 4a). Here lateral movement is less constrained than in Kongsbreen front and, thus, more mixing is possible (Torsvik et al., 2019). In contrast, in Rijpfjorden, temperature profiles did not show a

distinct water mass at the fjord end (Fig. 5a). In addition, the halocline in Kongsfjorden was spread over a larger depth (~50 m-depth) and was less sharp than in Rijpfjorden (Figs. 4b, 5b), as supported by salinity cross sections (Figs. 2b, 3b). This is

245 a characteristic feature of fjords with tidewater glaciers where plume set-up disperses salinity due to mixing and entrainment (Torsvik et al., 2019). In summary, Kongsfjorden exhibits the features of dynamic circulation associated with the subglacial plume set-up, which is absent in Rijpfjorden.

Nutrient and N* concentrations in Kongsfjorden were low or below the limit of detection (<1 μmol L⁻¹ nitrate, <0.2 μmol L⁻¹
phosphate, < -12.7 μmol L⁻¹ N*) throughout the top 50 m in mid- to outer- fjord (Kb0-3) and continental shelf (V12) stations (Figs. 4d,f,g, 6a,c,d) due to uptake by phytoplankton as demonstrated by elevated chlorophyll concentrations (Fig. 4c). Such low nitrate and phosphate concentrations at these fjord stations were located within and above the halocline, whereas those at the continental shelf were related to the warmer AW intrusion (Fig. 4a,b). In Rijpfjorden, although lowest nutrient and N* concentrations were found above the halocline (~50 m-depth), in general they were higher with 1 μmol L⁻¹ nitrate, >0.2 μmol
L⁻¹ phosphate and > -12.5 μmol L⁻¹ N* both in the fjord and at the shelf stations (Figs. 5d,f,g, 7a,c,d). These slightly higher nutrient concentrations in the stratified layers of Rijpfjorden may indicate incomplete nutrient utilisation at the surface as

terrestrial inputs are unlikely to be much higher compared to Kongsfjorden.

In both fjords, coinciding with the base of the halocline, i.e., below 50 m depth, nitrate, silicate, phosphate and N*
concentrations increased in all stations (Figs. 4d-g, 5d-g) while the isotopic ratios and Si* decreased (Figs. 4h-j, 5h-j). Decrease in nutrient concentration towards the surface accompanied by increases in δ¹⁵N-NO₃⁻ and δ¹⁸O-NO₃⁻ (Figs. 4i,j, 5i,j) are trends associated with uptake by phytoplankton, while the increase in N* with depth are associated with nutrient regeneration. Higher Si* (> 0) towards the surface implies the presence of nutrient inputs carrying excess Si with respect to N (Figs. 4h, 6e). In contrast, nutrient concentrations were higher in the glacier front at Kongsfjorden (Kb5-7) than in other fjord sampling sites. These high nutrient values were associated with low salinity (Fig. 4b), showed no defined depth-dependent trend (Fig. 4d,f) and Si concentrations were generally high enough to overcome Si-limitation while N-deficiency persisted (Si*>0 and N*<0; Figs. 4g,h, 6d,e). Moreover, δ¹⁵N-NO₃⁻ was low and clustered at around 4.3 ± 0.1‰ (Fig. 4i) indicating a source dependence rather than uptake.

Conversely, nutrient and N* concentrations reached 3.2 μ mol L⁻¹ nitrate, 0.3 μ mol L⁻¹ phosphate and -10.4 μ mol L⁻¹ N* at 25mdepth in the continental slope at Kongsfjorden (**Figs. 4d,f,g, 6a,c,d**), decreasing towards the surface. There, elevated δ^{15} N-NO₃⁻ and δ^{18} O-NO₃⁻ with increasing proximity to water surface (**Figs. 4i,j, 6f,g**) are explained by uptake by phytoplankton. The same interpretations also apply to the continental slope of Rijpfjorden (**Figs. 5d-j, 7a-g**).

In Rijpfjorden, δ^{15} N-NO₃⁻ and δ^{18} O-NO₃⁻ values taken at the deepest sampling depth of 100 m showed enrichments in isotopic values from the shelf and slope stations into the fjord. Namely the δ^{15} N-NO₃⁻ at fjord stations was 5.8 ± 0.2 ‰, which was

higher than the shelf and slope signature of 5.4 ± 0.4 ‰ (Fig. 5i). Meanwhile, δ^{18} O-NO₃⁻ at fjord stations (2.7 ± 0.5 ‰) also showed an enrichment with respect to shelf and slope stations (2.2 ± 0.7 ‰) (Fig. 5j). Noticeably, δ^{15} N-NO₃⁻ and δ^{18} O-NO₃⁻ signatures at shelf and slope stations fall within the range of AW signatures (δ^{18} O_{AW}-NO₃⁻ = 2.8 ± 0.3 ‰; δ^{15} N_{AW}-NO₃⁻ = 5.1 ± 0.1 ‰; Tuerena et al., 2021b).

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4 Discussion

4.1 Nutrient limitation and terrestrial nutrient inputs

In Kongsfjorden, higher nutrient concentrations at the glacier front stations suggests nutrient inputs from glacial discharge plumes. Concentrations of 1-4 μ mol L⁻¹ nitrate, 0.2-0.4 μ mol L⁻¹ phosphate and 1.5-2.5 μ mol L⁻¹ silicate found within the

- 285 halocline in stations near glacier fronts can only be explained by this terrestrial supply as the halocline presence clearly restricts nutrient mixing from depths into the upper 50m -as was shown in nutrient profiles in other fjord stations (Figs. 4d-f, 5d-f). Indeed, glacial meltwater has been found to be enriched in these nutrients, with concentrations broadly ranging from 0.24-1.60 µmol L⁻¹ nitrate, 0- 0.19 µmol L⁻¹ phosphate and 1.72- 3.47 µmol L⁻¹ silicate (Halbach et al., 2019). In Rijpfjorden, where nutrient concentrations were generally higher than in Kongsfjorden, there are no tidewater glaciers, but this does not rule out
- 290 the presence of terrestrial inputs from surface meltwater flow through streams. In fact, terrestrial inputs are potentially responsible for the enrichment in silicate (>1μmol L⁻¹) and phosphate (>0.2 μmol L⁻¹) in inner Rijpfjorden. Marine inputs cannot explain higher concentrations in Rijpfjorden as AW intrusion is too weak to extend to the continental shelf.

On the continental shelf outside Kongsfjorden, a strong thermocline at ~40 m pointed to a strong marine influence through the intrusion of AW between 20-50 m (34.9 PSU, 5.6°C) into the fjord. This intrusion was associated with a chlorophyll *a* maximum of 4-6 mg m⁻³ and nutrient-depletion due to phytoplankton growth, thus suggesting a potential marine contribution to the nutrient pool of Kongsfjorden. Vertical supply of nutrients from underlying modified PSW (34.7 PSU, 3°C) was likely hindered by the strong thermocline. Likewise, outside of the sill, AW was well-mixed and dominates throughout the whole water column, hosting a phytoplankton bloom with a fluorescence peak (14 mg m⁻³) at 20 m depth.

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Marine nutrient contribution to productivity in both fjords is hindered by the strong halocline that develops in summer and as a result mixing is restricted to winter overturning. In addition, AW is relatively nutrient poor as nutrient depletion occurred before entering the fjord (Cottier et al., 2010). Therefore, terrestrial nutrient inputs which occur throughout the summer with higher nutrient concentrations than marine inputs can be more important in these fjords during the summer (Hopwood et al., 2020).

Terrestrial inputs, evident from glacier front profiles, can be distinguished using nutrient stoichiometry by plotting nitrate concentrations against phosphate (**Fig. 8a**) and against silicate concentrations (**Fig. 8b**). N and P concentrations in Kongsfjorden, Rijpfjorden and eastern Fram Strait are shown in **Fig. 8a** along with lines representing Redfield ratio (N:P, 1:15

- and 1:16). Linear trends with slope similar to Redfield ratios indicate phytoplankton uptake and regeneration stoichiometry. Thus, most of the fjord samples show the influence of nutrient uptake and recycling. In **Fig. 8a**, $\Delta N/\Delta P$ at Kongsfjorden and Rijpfjorden was 16.8 (R² 0.96, *p*-value ≤ 0.05) as slope conforms broadly to Redfield ratio. However, the x-intercept (0.14 µmol PO4³⁻ L⁻¹) suggests surplus supply of phosphate from riverine input into the fjords and the inner shelf area (**Fig. 8a**; McGovern et al., 2020). It has been reported that discharge from the Bayelva river into Kongsfjorden contributes 60 µmol
- 315 PO4³⁻ L⁻¹, equivalent to 5.4 tons per year using a total discharge of 29 x 10⁶ m³ recorded in 2012 (Zhu et al., 2016), whereas Rijpfjorden receives P-input by glacially fed rivers (Wang et al., 2013). In general, the P intercept is consistent with the suggestion that nutrients supplied from Arctic soils are enriched in phosphate relative to nitrate due to the loss of N via denitrification owing to waterlogging during the summer months (Hayashi et al., 2018).
- 320 In contrast, marine inputs cannot explain the P intercept recorded in Kongsfjorden and Rijpfjorden as AW in eastern Fram Strait shows an intercept of ~ 0 (Tuerena et al., 2021a; Fig. 8a). Thus, the nutrient stoichiometry of Kongsfjorden and Rijpfjorden are influenced by terrestrial nutrient inputs. Broadly, despite this terrestrial input, the nutrient data plotted in Fig. 8a indicates rapid N depletion in the upper water column of both fjords. This can be explained by stratification and excess terrestrial P inputs relative to the Redfield ratio.

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Inshore and fjord samples that show excess P also show higher Si:N ratios due to terrestrial Si supply both in Kongsfjorden and Rijpfjorden (**Fig. 8a**; Dugdale and Wilkerson, 2001). When nitrate *vs.* silicate concentrations are plotted using salinity as

a third variable, most stations follow a linear trend with a N:Si ratio of 2.5 (**Fig. 8b**). Outliers are associated with low salinity (<33.5 PSU) and have a lower N:Si ratio of 1.3 indicative of Si enrichment (relative to NO_3^{-}) from terrestrial discharge (**Fig.**

- 8b). This is supported by Si* profiles, where values are higher within the halocline (Figs. 4h, 6e). These outliers correspond to innermost stations at both fjords and are represented as high Si:N ratios (> 1) in Fig. 8a. Furthermore, in Kongsfjorden, data also suggest that terrestrial silicate is supplied through the glacial discharge plume (Fig. 8b), most likely from glacier meltwater enriched in silicate from weathering of siliceous rocks (Dugdale and Wilkerson, 2001; Halbach et al., 2019). Meanwhile, in Rijpfjorden, silicate is supplied through glacially fed rivers rather than directly by tidewater glaciers (Wang et al., 2013). In
- 335 contrast, AW is a poor source of silica, with silicate limitation (Krisch et al., 2020) and phosphate deficiency evident in surface waters (Tuerena et al., 2021a) and make only limited contribution to the halocline in both fjords. Therefore, it is suggested that Kongsfjorden and Rijpfjorden have the potential to act as a source of silicate and to a lesser extent phosphate to AW offshore. This aspect is discussed below taking into consideration the fjord residence times.
- 340 Both Kongsfjorden and Rijpfjorden receive terrestrial nutrients enriched in Si and P relative to N, via glacial plumes and riverine input, which are then consumed by phytoplankton uptake within the fjord (**Fig. 8**). Here we evaluate the potential impact of this on siliceous diatoms (Egge, 1998), by comparing Si:N uptake ratios within *vs.* outside of the fjord (**Fig. 9**).

In both fjords, data from surface waters (<100 m) indicate an Si:N uptake ratio of 0.3 (R²= 0.74, p ≤0.05), and in the eastern
Fram Strait of 0.4 (R²= 0.84, p ≤0.05) (Fig. 9). This suggests that the diatom production as a proportion of the whole phytoplankton community is higher in the eastern Fram Strait (c.a. 40% of phytoplankton are diatoms) compared to fjord stations (ca. 30%) given that diatom Si:N uptake ratio is ~ 1 (Brzezinski, 1985). This implies that terrestrial Si inputs have no significant impact on phytoplankton composition. This is probably because terrestrial Si inputs are insufficient to overcome the Si limitation as all fjordic samples fall below 5 μmolSi L⁻¹, which is generally thought to be the threshold for kinetic Si
limitation (Krause et al., 2018; Fig. 9). Si limitation in the fjord can be explained by the "silicate pump", whereby more Si is rapidly lost to the deep- through sinking of biogenic silica or Si(OH)₄-rich faecal pellets- compared to N, and Si resupply is

- further prevented by stratification (Dugdale et al., 1995). Nevertheless, it should be noted that diatom Si:N uptake ratio may be greater than 1 in the eastern Fram Strait owing to transient Fe limitation reported here (Krisch et al. 2020). Thus, the overall higher Si:N ratio in the eastern Fram Strait relative to Kongsfjorden may also result from this process.
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Several lines of evidence support higher diatom abundance outside of the fjord as diatoms generally dominate the phytoplankton assemblage in Arctic spring blooms (Hodal et al., 2012; Krause et al., 2019). Uptake ratio reported herein (ca. 0.4) in eastern Fram Strait suggests phytoplankton succession and a shift from diatom dominance in the spring bloom to nanoand picoplankton communities in summer months (Strom et al., 2006). Such phytoplankton succession is triggered by the

- 360 depletion of nutrients from surface waters following the spring bloom and has been previously reported outside of Kongsfjorden (Rokkan Iversen and Seuthe, 2011; Kulk et al., 2018). Previous studies have documented that small nano- and picophytoplankton dominate the phytoplankton community throughout the summer in Kongsfjorden (Piquet et al., 2014; Kulk et al., 2018), although phytoplankton abundances can be patchy and succession can vary from year to year due to sea ice duration and hydrological conditions. There are reports of diatoms being more abundant in cold years and small flagellates
- 365 during warmer years (Piwosz et al., 2015). The results presented here therefore reflect an integration over the summer growing season sampled. However, in general, small phytoplankton cells thrive in fresher and oligotrophic surface waters due to their large surface-area-to-volume ratio which provides effective acquisition of nutrient solutes and photons, and hydrodynamic resistance to sinking (Li et al., 2009).

Thus, our results document terrestrial nutrient inputs within the fjords, P inputs occur in excess of the Redfield ratio (N:P=16)

370 and the excess terrestrial Si inputs might not necessarily by itself favour greater diatom production later in the season due to progressive nutrient limitation within the fjords.

4.2 Identifying isotopic signatures of N source versus cycling in Svalbard fjords

- Dual isotope signatures of δ¹⁵N-NO₃⁻ and δ¹⁸O-NO₃⁻ as well as nitrate concentrations are used to delineate the relative importance of various terrestrial sources. δ¹⁵N-NO₃⁻ and δ¹⁸O-NO₃⁻ were used to trace nitrogen cycling processes as they
 exhibit characteristic isotopic fractionation trends (Sigman and Fripiat, 2019) and offer insight into the relative importance of potential nitrate sources (Sigman and Casciotti, 2001) (see Appendix C). In Figure 10, δ¹⁵N-NO₃⁻ is plotted against the log of nitrate concentration with salinity as a third variable. If biological uptake dominated changes in nitrate concentration and composition, data would show a linear relationship, as expected from Rayleigh fractionation, and the slope of the line would represent the fractionation factor (ε; Altabet and Francois, 2001). Fractionation during uptake would lead to ε within the range
 3-10 (Wankel et al., 2009). Indeed, ε of 4.9‰ was recorded in the Fram Strait (Tuerena et al., 2021b) and for comparison the trend line from this Fram Srait study is shown as a dotted line in Figure 10. In Kongsfjorden and Rijpfjorden, most fjordic samples do not show the linear trend conforming to Rayleigh fractionation. Deviation from the linear relationship increases landwards within the fjord, where innermost stations show a slope near 0 (Fig. 10). This suggests that the isotopic signatures of nitrate are not governed by assimilation processes alone and that mixing between marine and terrestrial nitrate sources
- 385 influence isotopic signals significantly. Nutrient dilution effect from freshwater input can be ruled out as salinity alone could not explain the isotope effect in **Figure 10** (see **Fig. A4** for justification). Additionally, the terrestrial source is significantly lower in δ^{15} N as the trends indicate lighter signatures with proximity to land in both fjords (**Fig. 10**). Importantly, the lowest δ^{15} N signatures with no slope are found in the Kongsfjorden glacial front samples where terrestrial inputs of P and Si are identified (**Fig. 8**). The remnants of winter cooled waters at Kronebreen and Kongsbreen (Halbach et al., 2019; Hopwood et
- al., 2020) have distinctly lighter isotopic values (δ^{15} N-NO₃⁻ of 3.9-4.3 ‰) relative to the rest of fjordic samples indicating a larger contribution from terrestrial sources rather than marine sources (AW δ^{15} N-NO₃⁻ of 5.1 ± 0.1 ‰, Tuerena et al., 2021b) causing significant deviations from uptake dominated fractionation trends. Meanwhile, the epsilon value of remaining fjordic samples (ε = 1.8‰) can be approximated to that in salinity stratified PSW and in northern Barents Sea (ε = 2‰; Tuerena et al., 2021b).
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A plot of δ^{15} N-NO₃⁻ *vs.* δ^{18} O-NO₃⁻ with depth as the third variable is shown in **Fig. 11a**. Data from Kongsfjorden and Rijpfjorden show three clear trends. Firstly, data from Kongsfjorden -with the exception of glacier front stations- and data from the continental slope outside of Rijpfjorden showed a positive correlation (R² 0.98, *p*-value \leq 0.05) with a slope close to 1, whereby isotopic signatures are heavier at shallower depth (**Fig. 11a**). This is indicative of nitrate assimilation by phytoplankton which fractionates both isotopes equally, that is δ^{15} N-NO₃⁻: δ^{18} O-NO₃⁻ ratio approaching 1 (DiFiore et al., 2009; Sigman et al., 2009). Although it is inferred that biological uptake mainly determines the isotopic trends in these samples, the slight deviation from 1 (c.a. 0.8) may also result from simultaneous uptake and recycling. During recycling, δ^{18} O of nitrate gets reset closer to water values with an added isotopic fractionation effect of 1.1‰ (Buchwald et al., 2012; Sigman et al., 2009). In these fjordic settings, δ^{18} O _{H2O} is estimated to be c.a. 0.3‰ according to the equation δ^{18} O _{H2O} = 0.54S-18.42 derived for Kongsfjorden (Tiwari et al., 2018) using the average salinity (S) of 34.8 ± 0.006 PSU in the winter cooled waters. For comparison purposes, another δ^{15} N-NO₃ *vs.* δ^{18} O-NO₃ plot is presented (**Fig. 11b**) using only stations from the continental

shelf and slope of Kongsfjorden as well as from eastern Fram Strait. Data show a positive correlation ($R^2 0.77$, *p*-value ≤ 0.05) with a slope of 0.8, with δ^{18} O-NO₃ values clustering at 2.0 $\pm 0.6\%$ and at 4.9 $\pm 0.2\%$ for δ^{15} N-NO₃⁻; these values are indicative of the initial isotopic signatures of the marine endmember. A comparison of **Fig. 11a** and **Fig. 11b**, reveals that the slope of

410 the line from Kongsfjorden samples is similar to the slope of offshore and Fram Strait samples (~0.8). In summary, biological uptake is the most likely determinant of the isotopic trends in these samples.

However, as expected, the glacier front stations in Kongsfjorden deviated significantly from the 1:1 line, with δ^{15} N-NO₃⁻ values clustering around 4.3 ± 0.1‰, meanwhile, δ^{18} O-NO₃⁻ ranged from 2.3 to 6.5 ‰ (**Fig. 11a**). This deviation was caused

- 415 by relatively light δ^{15} N-NO₃⁻ and enriched δ^{18} O-NO₃⁻ signatures, which were derived from terrestrial sources, in comparison to other fjordic stations. In addition to the deviation from 1:1 relationship, isotopic values of these samples did not show any linear trend. This indicates source dominance rather than uptake. These samples were previously identified within the glacial discharge plume (S <33.5PSU) with high silicate values deviating from the Si:N line (**Fig. 8b**). These samples incorporate terrestrial nitrate source signatures which are rapidly lost away from the glacial plume through biological uptake.
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In **Fig. 11a**, fjord and continental shelf samples at Rijpfjorden also deviated from the 1:1 line – falling on a slope of 0.6 (R² 0.81, *p*-value \leq 0.05). Interestingly, the extension of the line of best fit of Rijpfjorden data crosses the cluster of data points comprising glacier front Kongsfjorden stations (i.e., the terrestrial endmember signature). However, the samples themselves do not fall within the cluster and instead show enriched isotopic signatures. This is indicative of the input of terrestrial nitrate in Rijpfjorden with relatively light δ^{15} N-NO₃⁻ and enriched δ^{18} O-NO₃⁻ signatures similar to the glacial plume samples of

Kongsfjorden but subsequently modified by uptake and recycling as described below.

Unlike Kongsfjorden, nitrate is not completely depleted in Rijpfjorden indicating partial nitrate utilisation which leads to heavy residual nitrate signatures where δ^{18} O-NO₃⁻ reach values as high as 14‰ and δ^{15} N-NO₃⁻ up to 10‰ (**Fig. 11a**). The lower gradient (0.6 instead of 0.8) is indicative of partial uptake and regeneration. The partial uptake leads to heavy signatures of

- δ^{15} N-NO₃⁻ and δ^{18} O-NO₃⁻ but simultaneous recycling forces δ^{18} O towards lower values close to δ^{18} O H2O. This δ^{18} O H2O can be assumed to be the same in both fjords (0.3‰). Thus, isotopic composition reveals the importance of the terrestrial nitrate source in both fjords as well as differences between the two fjords in the way nitrate is consumed and recycled.
- In Kongsfjorden, nitrate uptake is complete with near-zero nitrate values above the halocline with the exception of those in proximity to the subglacial plume at the glacial front. When nitrate is present it reflects the additional terrestrial source and its isotopic signature. In contrast, nitrate is not fully consumed in near surface waters of Rijpfjorden leading to significant isotopic fractionation. There, the isotopic trends in **Figure 11a** -where the fractionation line passes through the glacial plume samples representing terrestrial endmember in this setting- suggest significant terrestrial nitrate contribution in Rijpfjorden which is subsequently masked by mixing with partially utilised nitrate. This partly explains why the terrestrial source signatures with
- low δ^{15} N-NO₃⁻ signature, evident in Kongsfjorden, are not as obvious in Rijpfjorden (**Fig. 11a**). The difference in nitrate utilisation suggested above in these two fjords can arise from the nature of circulation in these fjords which is discussed in detail below in the context of the role of subglacial plume in nutrient dynamics.

445 **4.3** The role of the subglacial plume in nitrate use and cycling

The difference in nitrate utilisation suggested above in these two fjords can arise from the nature of circulation in these fjords and the difference in light limitation resulting from surface and subsurface discharge. In Rijpfjorden, the fjord stations show strong freshwater stratification, and the fresher water is confined to a shallow surface layer resulting from surface discharge

450 from rivers (Figs. 3b, 5b). In contrast, at the glacial front of Kongsfjorden meltwater influences a larger depth in the water column with less fresh surface waters (Figs. 2b, 4b, A2). This reflects the presence of the subglacial plume in Kongsfjorden where the plume rises from the grounding line of tidewater glaciers at 60 m depth (Darlington, 2015) increasing mixing

between meltwater and the entrained saltwater over a larger depth range near the surface (Everett et al., 2018). This is in agreement with a previous study which documented the plume to extend to 40 m depth in this fjord (Torsvik et al., 2019). It is

455 worth noting that the meltwater plume in Kongsfjorden is mainly subglacial freshwater as opposed to glacial surface melt (How et al., 2017).

Importantly, the magnitude of the effect of the subglacial discharge and glacier front plume in Kongsfjorden varies spatially along the glacial front owing to topographical differences along Kongsbreen (Kb6-7) and Kronebreen (Kb5) transects. Along both fronts, a distinct cool and saline water mass was identified as remnant winter cooled waters formed due to heat loss to the atmosphere and contact with the glacier front (De Rovere et al., 2022). Studies by Torsvik et al. (2019) attributed the presence of this water mass in Kongsfjorden to its entrainment and elevation by the glacial plume dynamics. However, along Kronebreen front these winter cooled waters undergo mixing with fjord water as shown by temperature above 3.5 °C and salinity above 33.5 PSU at 20-35 m depth (Fig. 4a,b) and the presence of the subglacial discharge at glacier front plume is less distinct. At
Kronebreen front, mixing associated with the plume is topographically unconstrained -as opposed to in Kongsbreen – as it is wider thus allowing lateral water movement (Torsvik et al., 2019). The lack of mixing below 35m-depth is explained by a topographic barrier that restricts the exchange of water masses as suggested by particle analysis (Torsvik et al., 2019).

Whether the release of glacially eroded, entrained sediments (Elverhøi et al., 1983; Trusel et al., 2010; Kehrl et al., 2011)
occurs through surface or subsurface discharge can impact light limitation. Rijpfjorden experienced direct surface discharge release through glacially fed rivers (Wang et al., 2013). This surface release of sediments is expected to encourage stronger light limitation at the surface than subsurface discharge, as suspended sediments remain near the surface due to strong stratification (Cowton et al., 2015; Carroll et al., 2015, 2016). Near-zero fluorescence values in the inner fjord of Rijpfjorden support that productivity could be hindered by light limitation. In contrast, in Kongsfjorden, turbidity associated with sediment discharge is subject to dynamic mixing associated with subglacial plumes (Cowton et al., 2015; Carroll et al., 2015, 2016)

- 475 discharge is subject to dynamic mixing associated with subglacial plumes (Cowton et al., 2015; Carroll et al., 2015, 2016) which invigorate vertical mixing and lateral exchange of water. In such a dynamic setting, with weak stratification, the turbidity caused by glacially eroded sediments is relatively more dispersed, increasing light penetration away from the plumes which facilitates plankton blooms (Calleja et al., 2017). Such tidewater glacier-related circulation dynamics have been documented in Kongsfjorden (Darlington, 2015; How et al, 2017; Schild et al., 2017; Halbach et al., 2019). The longer residence times and
- 480 the dynamic circulation associated with tidewater glaciers in Kongsfjorden lead to complete utilisation of nutrients at the surface and recycling at depth within the fjord (i.e., increased regenerated nutrient storage), inhibiting the export of nutrients offshore. In Rijpfjorden, light limitation retards nutrient utilisation (Torsvik et al., 2019) leading to only partial use as suggested by the isotope data. Moreover, the differences could also result from delayed nutrient utilisation in Rijpfjorden compared to Kongsfjorden due to its more extensive ice cover. Other additional factors could be the shallower depth of Rijpfjorden and
- 485 thus shorter residence time which facilitates shelf exchange before nutrients are fully utilised (Straneo and Cenedese, 2015; Morlighem et al., 2017). This increases the opportunity for exporting the heavy, partially-utilised nitrate from Rijpfjorden into the shelf where it can be subsequently utilised and impact productivity.

As a final note, the scenario described for Rijpfjorden is representative of what is happening in Greenland and other Arctic fjords that are currently experiencing tidewater glacier retreat onto land (Nuth et al., 2013; Meire et al., 2017; Kanna et al., 2018). The implication is that, with increasing Atlantification and warmer AW in the future, AW intrusion in Arctic fjords will strengthen and thus encourage tidewater glacier retreat. This will favour surface discharge of terrestrial nutrients, less dynamic circulation, leading to light limitation and retarded use of nutrients and ultimately, aiding the export of nutrients offshore. The significance of this stems from the fact that the Greenland ice sheet and its associated tidewater glaciers have been suggested

495 to be an important source of nutrients to the wider Arctic Basin and thus exert an important control on overall Arctic net

primary production (Hawkings et al., 2015), with terrestrial nutrient inputs currently estimated to support 28-51% of net primary production basinwide (Nuth et al., 2013; Terhaar et al., 2021). It is worth noting some differences between Svalbard and Greenland settings. Some Greenland fjords have very deep tidewater glaciers and receive smaller terrestrial N inputs due to less soil, thus marine sources of nutrients could be relatively more important (Cape et al., 2019).

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4.4 Fingerprinting terrestrial N sources signatures and contributions to N inventory

In a fjordic setting, various sources of terrestrial nitrogen are discharged and mixed with marine sources, subjected to varying degrees of uptake and recycling with the fjords and the residual exported to the shelf. Previous studies show that terrestrial N inputs exhibit high seasonality, often peaking in the summer months (McGovern et al., 2020), and have δ¹⁵N-NO₃⁻ values typically ranging from 0 to 5‰ (Holmes et al., 2012; Sigman and Fripiat, 2019). Snowpack melting and release of nitrate from atmospheric sources also has significant implications for the isotopic signatures. The snowpack acts as a sink for ¹⁵N-depleted atmospheric reactive nitrogen (δ¹⁵N-NO₃⁻: ~-6.5‰) (Heaton et al., 2004; Björkman et al., 2014; Vega et al., 2015). Some studies highlight role of nitrification and nitrified ammonium, from a variety of sources to be an additional contributor of nitrate from melting snowpack (Hodson et al., 2005; Wynn et al., 2007; Halbach et al., 2019). Other terrestrial inputs, such as tidewater glacier melting may contain N from bird faeces (δ¹⁵N-NO₃⁻, ~8‰; Skrzypek at al., 2015) and N from microbial degradation of organic matter in the glacier ice (D'Angelo et al., 2018), which can contribute to moderately N-rich meltwater runoff (Shi et al., 2018). Although N₂ fixation in diazotrophic bacteria has been reported in the Arctic regions (Blais et al., 2012; Zehr and Capone, 2020), including the glacial environment of Svalbard (Telling et al., 2011), the magnitude of this source remains poorly constrained. Low isotope values of δ¹⁵N-NO₃⁻ have been attributed to terrestrial N, fixation in Svalbard

515 (~-0.5‰) (Skrzypek at al., 2015).

The isotopic studies indicate significant terrestrial N inputs in both fjord systems considered. In Kongsfjorden the glacier front samples identified with terrestrial sources of nitrate have isotopic signatures that do not show evidence for uptake and isotopic fractionation. These samples have δ^{15} N-NO₃⁻ values clustering around 4.3± 0.1‰, meanwhile, δ^{18} O-NO₃⁻ ranges from 2.3 to

- 520 6.5 ‰. Although the isotopic signature of the terrestrial N is masked by mixing and partial use of nitrate in Rijpfjorden, the isotope fractionation trendline passes through the glacial front samples of Kongsfjorden, indicating similar terrestrial source signatures in both fjords (**Fig. 11a**). This provides the opportunity (1) to identify the origin of terrestrial sources and (2) to delineate terrestrial and marine component of N in the upwelling plume at the glacial front.
- 525 High δ¹⁸O-NO₃⁻ values (60-86‰; Heaton et al., 2004) have been reported within atmospherically-deposited nitrate related to ozone-depleted air in Svalbard's snowpack (Björkman et al., 2014). Meanwhile low δ¹⁵N-NO₃⁻ in snowfall has been attributed to long range atmospheric transport deposition over the Arctic Ocean (δ¹⁵N_{snowfall} of -4‰; Heaton et al., 2004; Vega et al., 2015). In general, nitrate from this source is expected to have lighter δ¹⁵N and heavier δ¹⁸O relative to marine values (δ¹⁸O-NO₃⁻ of 2.0 ± 0.6‰ and δ¹⁵N-NO₃⁻ of 4.9 ± 0.2 ‰; Fig. 11b; Appendix C). In this regard it is important to note that during
- 530 warm summers as much as 50% of the annual snowpack accumulation in Svalbard may melt (Pohjola et al., 2002). Therefore it is expected that the dominant terrestrial source could be melting of seasonal snowpacks and glacier ice because it acts as a sink for atmospheric reactive nitrogen with high δ^{18} O-NO₃⁻ and low δ^{15} N-NO₃⁻ signatures as previously mentioned (Björkman et al., 2014; Vega et al., 2015). The predominance of this source is consistent with isotopic composition (lighter δ^{15} N and heavier δ^{18} O) of nitrate in Kongsfjorden glacial front and from the intersection of the line of best fit at Rijpfjorden through this
- 535 data (Fig. 11a).

Additional contributor of nitrate from melting snowpack is nitrified ammonium (Hodson et al., 2005; Wynn et al., 2007; Halbach et al., 2019). However, this ammonium-sourced nitrate is lighter ($\delta^{15}N$ =-4.4 to -3.5 ‰, $\delta^{18}O$ = -0.3‰; Wynn et al., 2007). Other terrestrial inputs such as bird guano would enrich the $\delta^{15}N$ -NO₃⁻ signature in the fjord (c.a. 8 ‰; Bokhorst et al., 2007; Szpak et al., 2012; Skrzypek et al., 2015). We regard these as minor sources as they cannot produce the combination of

540 light $\delta^{15}N$ (4.3± 0.1 ‰) and heavy $\delta^{18}O$ values (2.3- 6.5 ‰) that we record at the glacial front stations. In particular the heavy $\delta^{18}O$ values (2.3- 6.5 ‰) recorded in the glacial front can only come from atmospheric deposited nitrate in ice-packs subsequently transported through meltwater discharge.

Overall, the δ¹⁵N-NO₃⁻ value of 4.3± 0.1 ‰ (Fig. 11a) could be considered as nitrate sourced from the upwelling plume. The influence of biological uptake in the glacier front discharge plume is expected to be low as high suspended matter hinders light penetration and phytoplankton production in the plume regions (Kumar et al., 2018). This is supported by very low chlorophyll contents (<0.8 mg m⁻³, this work), and high mineral matter load in GF/F filters supports this contention. The fact that the δ¹⁵N-NO₃⁻ values strongly cluster around 4.3 ‰ without any relationship to nitrate concentration as would be expected from uptake also support this (Fig. 10). However, the plume dynamics involve some degree of mixing with marine sources due to entrainment of underlying marine waters. This mixing is evaluated below.

- In Kongsfjorden, published terrestrial endmember value for δ^{15} N-NO₃⁻ is 3.5 ‰ (Kumar et al., 2018). This number is derived from dual C and N isotope mixing model based on Kongsfjorden sediments. Thus the 4.3 ‰ value evident for δ^{15} N-NO₃⁻ at the fjord front can be regarded as an admixture of terrestrial N (c.a., 3.5 ‰; Kumar et al., 2018) and marine source (c.a., 5.1 ± 0.1 ‰; Tuerena et al., 2021b) in equal proportions ($\frac{4.3-3.5}{(4.3-3.5)+((5.1\pm0.1)-4.3)} \times 100 = 50 \pm 3$ %). Note that the marine endmember of 5.1 ± 0.1 ‰ (Tuerena et al., 2021b) is used in the mixing calculation instead of that estimated from continental shelf and
- slope samples ($4.9 \pm 0.2 \%$; Fig. 11b) as it is a better representation of pure AW signal, while the latter is likely altered by advected terrestrial nutrients from the glacier front. The glacial front samples containing marine N in equal proportion to terrestrial N is consistent with entrainment and mixing that occurs during upwelling of the subglacial plume that draws nitrate from remnant winter cooled waters through lateral mixing as suggested by recent studies at Kronebreen and Kongsbreen fronts (Halbach et al., 2019; Hopwood et al., 2020).
- 560 Additionally, the contribution of terrestrial *vs* marine sources to the overall nutrient inventory of Kongsfjorden can be estimated. Using the average δ^{15} N value calculated from fjord basin samples below 100 m of $4.4 \pm 0.3 \ \text{\%}-\delta^{15}$ N-NO₃⁻ (**Fig. 4i**) and avoiding isotopic fractionation during phytoplankton assimilation which is evident in shallower samples. This value provides an estimate of marine contribution of $(\frac{4.4-3.5}{((4.4\pm0.3)-3.5)+((5.1\pm0.1)-(4.4\pm0.3))} \times 100 =) 56 \pm 3 \ \text{\%}$. Thus marine contribution gains some importance and terrestrial inputs remain significant (44 ± 3 %).
- Errors in these estimates stem from applying annually integrated terrestrial endmember estimates from sediments to water column snapshot of marine and terrestrial mixing documented during this study. For example, terrestrial inputs from early-season melt will give signatures closer to snowmelt, while late season incorporates heavier values due to denitrification in waterlogged soils and pockets of local anoxia in glaciers and stronger washout of guano (Wynn et al., 2007; Hayashi et al., 2018). These processes can also have large spatial variability within the ice pack (Ansari et al., 2013). Nevertheless, this
- 570 assessment indicates that terrestrial sources of N make an important contribution to the dissolved pool of nitrate in the glacial front as well as to the water column N inventory of the fjord.

In summary, we conclude that (i) snow and ice melt are the major sources of terrestrial nitrate to these fjords, (ii) at the glacial front, the nitrate is sourced from equal admixture of terrestrial and marine sources and (iii) terrestrial contribution remains significant to the whole fjord nutrient inventory even when considering deep waters. The prevalence of terrestrial N

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contribution at depth can be explained by N regeneration from sinking organic matter. Winter convection can also mix

terrestrial N to the deep water but this source is likely to be small. In the next section, nutrient regeneration is quantified relative to winter convection which will allow understanding the relative contribution of terrestrial *vs* marine nutrients to overall fjord productivity.

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4.5 Terrestrial N contribution to primary productivity in Kongsfjorden during the summer

Terrestrial inputs are an important contributor to the nutrient pool in the fjords, but what is its relative contribution to productivity in the fjord? To estimate this we use the WCW in Kongsfjorden, which occupies the deep basin between the land and the sill, and has the longest residence time in Kongsfjorden as illustrated by high AOU and low oxygen saturation (**Fig. 2d,e**; Svendsen et al., 2002). Given low WCW renewal rates, this water mass should have received prolonged supply of regenerated nitrogen via settling particulate organic nitrogen with limited physical exchange with the surface and the isotopic signatures are not affected by biological uptake since water mass formation. We can therefore use WCW to quantify the terrestrial contribution to productivity in the fjord as seen in the integrated regenerated nitrate pool and the preformed nitrate pool inherited by the water mass during formation through winter convection using dual isotopes of N and O.

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This methodology exploits the difference in isotopic fractionation engendered during remineralisation and nitrification processes. The principle enabling this calculation is that, while the $\delta^{15}N$ of remineralised NO₃⁻ records isotopic signatures of reactive N assimilated at the sea surface, δ^{18} O of remineralised NO₃ is reset close to that of ambient water in which regeneration occurs (see Appendix C). Using δ^{18} Owcw (1.9 ± 0.2 ‰) measured in this study, ambient water δ^{18} O_{H20}=0.3‰ estimated in Kongsfjorden (Tiwari et al., 2018) and $\delta^{18}O_{AW}=2.8 \pm 0.3\%$ previously measured for Atlantic waters outside Kongsfjorden 595 (Tuerena et al., 2021b) we estimate 65 ± 15 % of the nitrate in the WCW is regenerated (see Appendix D for more details). Furthermore the estimated N isotopic composition of this remineralised nitrate (δ^{15} Nreg: 3.7-4.1 ‰, Appendix D) demonstrates a major contribution of terrestrial sources (63-88%) to fjord primary productivity and recycling as opposed to marine sources (12-37%) (summarised in Table 1). The higher proportion of terrestrial N in the regenerated N pool demonstrates its 600 importance to primary production in the fjord and reflects the fact that terrestrial sources are supplied directly above the halocline throughout the summer growing season, whereas uptake of marine nutrients is hindered by the strong halocline that develops during the summer. This should be considered as a very broad estimate given the uncertainties associated with $\delta^{15} N_{reg}$ estimates. These include (i) limited sampling points in WCW (n=3),(ii) empirical data used as an approximation of $\delta^{18}O_{AW}$ and $\delta^{18}O_{H2O}$ and (iii) the percentage of regenerated nitrate (65 ± 15%). Nevetheless, these estimates demonstrate the 605 importance of terrestrial N inputs in supporting productivity (63-88%) while advection of marine nutrients is an important contributor (56 ± 3%) to the overall N inventory of Kongsfjorden (Hegseth et al., 2019). The former is consistent with

- contributor (56 \pm 3%) to the overall N inventory of Kongsfjorden (Hegseth et al., 2019). The former is consistent with McGovern et al. (2020) showing the important influence of terrestrial nutrients on fjord productivity in Svalbard. The latter conclusion is supported by Hegseth et al. (2019).
- 610 One important aspect of this conclusion is that terrestrial N sources account for 63-88% of fjord productivity and contribute in similar proportion to the regenerated pool of N in the deep isolated water masses of the fjord. Terrestrial inputs are expected to increase with climate warming and with the relocation of N held in soils and permafrosts (Vonk et al., 2015). Such enhanced terrestrial N inputs will have a direct effect of increasing surface productivity, the pool of regenerated/recycled N and the fjord N inventory. Such enhanced N cycling in extreme conditions would result in oxygen depletion of the the isolated water masses
- 615 such as the WCW. An added effect is the retreat of tidewater glaciers and absence of vigorous lateral circulation associated with subglacial plumes, which can further inhibit mixing, increase isolation of deep water masses and cause an expansion of oxygen depleted waters.

4.6 Future changes

Our results and analysis suggest that terrestrial N inputs contribute to nearly one half of the summer nitrate inventory in these fjords and support a large proportion of the productivity in summer due to its input to the surface stratified layer. Secondly, the presence or absence of subglacial plume can influence the relative use of nitrate within the fjord and its exchange with the open ocean. Both of these aspects are subject to future change.

It is expected that future warming will increase the source and magnitude of terrestrial inputs with key implications to the nutrient inventory, fjord primary productivity and nutrient export offshore. Permafrost melting is likely to gain importance, mobilising nutrients and in turn increasing terrestrial input into the marine environment (Vonk et al, 2015). This study shows that terrestrial N contributions account for nearly half of the N inventory and much of the productivity in Kongsfjorden and therefore, future increase in terrestrial N supply is likely to increase productivity as well as the storage and cycling of N within Arctic fjords.

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This comparative study between two fjords illustates the importance of tidewater glaciers in nitrate use and cycling within the fjords. The widespread retreat of tidewater glaciers in the Arctic and Greenland (Østby et al., 2017; Slater et al., 2019) has important implications on N exchange between fjords and the open ocean. It is expected that tidewater glaciers will retreat in Kongsfjorden to become land-terminating glaciers with strengthening Atlantification and warmer AW (Torsvik et al., 2019).

635 The surface meltwater flow can decrease primary producitvity via light and nutrient limitation as documented in the case of Rijpfjorden. Ultimately, given that N is the limiting nutrient in the Arctic Ocean and marine settings in general, this would aid the export of N offshore and contribute to the overall net primary production in the Arctic Ocean (Terhaar et al., 2021).

The broad differences discussed here are likely to change with Arctic settings, for instance in Greenland fjords where terrestrial N supply is limited, due to the lack of N storage in soils, marine N supply is more important (Cape et al., 2019). Here subglacial plumes invigorate marine N supply through upwelling from depth in the fjord and this fuels much of the productivity (e.g., Meire et al., 2017). The retreat of glaciers can reduce this marine N supply from upwelling thus inhibiting productivity. In the case of Svalbard fjords, terrestrial N supply is significant and is likely to gain further prominence in the future due to remobilisation of soil and permafrost N. In addition, we show that roughly half the N stored in deep fjord waters is also of terrestrial origin recycled within the fjord. Thus climate change is likely to increase productivity and N cycling and storage within the fjord. However, on longer time scales, if the distance between the fjord and the retreating glaciers becomes large enough, there will be more time for denitrification during transport, which may deplete the nutrient pool before it is discharged into the fjord. Once glaciers fully melt, nutrient inputs will be mainly associated with seasonal snow melt with opportunities for enhanced N cycling and removal in soils potentially reducing nutrient inputs.

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Regarding the export of nutrients offshore, climate change can have contrasting results depending on the size of the fjords and its circulation. Increased terrestrial N inputs and freshwater discharge are expected to increase export rather than storage in Rijpfjorden, as unused nutrients are flushed offshore given its shorter residence time. Thus, fjords with fast exchange rates like Rijpfjorden have more potential to alter pan-Arctic productivity as nutrients are exported offshore. In fjords like Kongsfjorden,

655 additional N inputs in the future can increase terrestrial N recycled within the fjord as biological uptake here is more complete. In Kongsfjorden, recycled nutrients are stored in isolated waters such as the WCW and this pool will increase due to the larger nutrient supply. In addition, the retreat of tidewater glaciers has the effect of reducing mixing in the fjord and potential for increasing retention of winter waters (Torsvik et al., 2019). The combination of greater productivity fuelled by terrestrial N, larger pool of N cycling within the fjord and reduced circulation in the absence of tidewater glaciers can lead to reduced oxygen 660 levels and possibly hypoxia in isolated deep waters of these fjords. The postulated contrasting responses of Kongsfjorden and Rijpfjorden to climate change are summarised in **Table 2**.

Conclusion

Kongsfjorden and Rijpfjorden are highly influenced by terrestrial nutrient inputs of N, P and Si. Particularly, terrestrial nitrate

- sources are key contributors to the NO₃⁻ pool in the fjords and are roughly equal in proportion to marine contributions from AW. These terrestrial inputs carry excess silicate and phosphate relative to nitrate with respect to marine sources. Nitrate from snowpack and glacier melting are identified as dominant sources of terrestrial NO₃⁻ in both fjords based on high δ^{18} O-NO₃⁻ and low δ^{15} N-NO₃⁻. In Kongsfjorden, stratification contributes to both nutrient limitations. Biological uptake of these nutrients indicates that nitrate and possibly silicate limits primary production despite terrestrial input. N and Si limitation is partly caused
- 670 by excess terrestrial P and in the case of Si also through effective removal through the "silicate pump". In Rijpfjorden, isotopic signatures indicate that nitrate is not fully utilised which may reflect light limitation and shallow stratification associated with the surface runoff. The contrast in nitrate use in these two fjords is attributed to the nature of meltwater inputs which is subglacial in Kongsfjorden and surface in Rijpfjorden and the resulting differences in circulation and stratification. As a result Rijpfjorden contributes terrestrial NO₃⁻ to the coastal sea whereas in Kongsfjorden terrestrial NO₃⁻ is mainly retained and
- 675 recycled within the fjord.

Given the significance of terrestrial nitrate sources in Svalbard fjords, it is postulated that continuing Arctic warming and enhanced meltwater discharge and terrestrial NO₃⁻ inputs will impact fjordic primary productivity as well as the NO₃⁻ exchange offshore. The larger size and vigorous circulation associated with surface glacial plumes currently leads to complete use of nutrients in surface waters of Kongsfjorden. Here, increase in terrestrial inputs and the retreat of tidewater glaciers due to climate change is postulated to increase productivity and nutrient cycling within the fjord possibly leading to more oxygen depletion in isolated deep waters which may expand given the less vigorous circulation in the absence the subglacial plume. In contrast, such changes in smaller fjords will lead to conditions akin to Rijpfjorden where surface inputs lead to strong stratification and light limitation which limits NO₃⁻ use and favours the export offshore of unused nutrients. This condition 685 could become more common with ongoing glacial retreat in many small Arctic fjords. The implication is that future increases

- 685 could become more common with ongoing glacial retreat in many small Arctic fjords. The implication is that future increases in terrestrial nutrient inputs in smaller fjords such as Rijpfjorden has greater potential to impact primary productivity offshore by exporting the unused nutrients.
- 690 Author contribution. MSG and RSG wrote the paper. MSG measured nitrate isotopes for NP2017 and NP2018. RET and MCFD measured nitrate isotopes for JR17005 and FS2018, respectively. All authors contributed to the final version of the paper.

Competing interests. The authors declare that they have no conflict of interest.

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700 captain and crew of the RV Lance.

Table 1. Summary of isotopic signatures (‰), endmember values (‰) and terrestrial and marine contributions (%)705calculated in Kongsfjorden.

REPORTED δ^{15} N SIGNATURES	‰	Terrestrial contribution (%) Marine contribution (%)
Nitrate in upwelling plume (at glacier front)	4.3 ± 0.1	50 ± 3	50 ± 3
Nitrate in fjord basin (>100m depth)	4.4 ± 0.3	44 ± 3	56 ± 3
Regenerated nitrate (65 \pm 15 %) in Winter Cooled Waters (integrated contribution)	3.7 - 4.1	63-88	12-37
Preformed nitrate (35 \pm 15 %) in fjord basin - Ranges betweeen 4.3 \pm 0.1 % (most constant)	stly terrestri	ial) and 5.1 ± 0.1 ‰ (fully mar	rine)

ENDMEMBER VALUES	%0
Terrestrial δ^{15} N endmember. From <i>Kumar et al. (2018)</i>	3.5
Marine δ^{15} N endmember. From <i>Tuerena et al. (2021b)</i>	5.1 ± 0.1
WCW δ^{15} N value	4.2 ± 0.2

		Kongsfjorden	Rijpfjorden
		Duc-Alcuc Join (12.0 14, 11.1 L)	(7 C.7.7 1)010 (00.0 1), 77.7 7
	Physical characteristics Fjord size Maximum fjord depth Mean fjord depth	~26 km long, ~6-14 km wide Deeper (~350 m) ~100 m	~40 km long, ~7-12 km wide Shallower (~200 m) ~100 m
	Continental shelf length	Narrower (~45 km) (\therefore stronger AW influence)	Wider ($\sim 60 \text{ km}$) (:: weaker A W influence)
suompuos tuss	Hydrology Glacial influence Riverine influence Dominant type of freshwater discharge Strength of fjordic circulation Estimated water residence time	Tidewater glaciers feed into fjord Bayelva river Subsurface discharge Vigorous circulation (driven by subglacial plume) ~172 hours (Yang et al., 2022)	Land-terminating glaciers feed into fjord Small, short-lived rivers Surface discharge Weaker vertical and lateral mixing Shorter (< 172 hours)
Pre	Nutrient cycling Stratification	Weaker - freshwater spread over a larger depth and is less fresh	Stronger -freshwater is confined to a fresher, thin layer extending further
	Light conditions Nitrate utilisation	Weaker turbidity throughout fjord (except at glacier fronts) Complete	ottshore Stronger turbidity throughout fjord Partial
	N storage and cycling Nitrate export offshore	More N recycling within fjord→ Higher regenerated N storage Lower	Less N recycling within fjord→ Lower regenerated N storage Higher
	Climate change effects Increased intrusion of warmer AW and higher air temperatures	\rightarrow Encouraged tidewater glacier retreat	→ Increased surface discharge
enoitoil	Increased permafrost/snowpack/glacier melting, increased riverine discharge and increased remobilisation of soils	\rightarrow Increased terrestrial nitrate input to fjords (<i>note: in longer time</i> .	scales, this might be reversed by increased denitrification rates)
Future prec	Climate change consequences Stratification Fjord N inventory Fjordic circulation Light conditions	(while tidewater glaciers are still present) Increased Increased (fuelled primarily by terrestrial inputs) Reduced Increased turbidity	Increased Increased (fuelled primarily by terrestrial inputs) Reduced Increased turbidity
	N storage and cycling Nitrate export offshore Ovvrom levels at isolated doon waters	Increased Decreased Hymoxia will notantially develors	Decreased Increased
	Net impact on primary productivity	Increased fordic productivity	Increased productivity in offshore coastal regions



21°E 22°E 23°E 24°E Longitude 1 IBCAO v.3 by Jakobsson et al. (2012). the eastern side of Fram Strait, illustrates sampling sites of the ow triangles; 2018, red dots) for δ^{15} N-NO₃⁻ and δ^{18} O-NO₃⁻ s well as on the shelf (V12) and continental slope (V10 and V6).

005 (blue triangles) cruises were also used for better coverage of offshore conditions. Shipboard measurements from cruise JR17005 were taken from the RSS *James Clark Ross* as part of the UK Changing Arctic Oceans programme in May-June 2018. Cruise FS2018 took place on the R.V *Kronprins Haakon* as part of the Fram Strait Arctic Outflow Observatory in August-September 2018. Colour coding corresponds to depth in metres.

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In (c) and (d), the bathymetric map of Rijpfjorden, Svalbard, illustrates sampling sites of the Norwegian Polar Institute monitoring cruises (NP2017, yellow triangles) for δ^{15} N-NO^{3⁻} and δ^{18} O-NO^{3⁻} determination. In (c) all sample locations are shown, namely in the inner fjord (R1-2), outer fjord (R3) as well as on the shelf (R4-R5) and continental slope (R6-R7B). In (d) a zoom-in map of inner and outer fjord region is shown. Colour coding corresponds to depth in metres.



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(NSd) 33.5 (

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AOU (µmol kg⁻¹ 10

Figure 2. Cross-sections across Kongsfjorden of (a) potential temperature (°C); (b) salinity (PSU); (c) fluorescence (mg m⁻³), (d) apparent oxygen utilisation (AOU, µmol kg⁻¹) and (e) oxygen saturation (%). Data from CTD casts from NP2018 cruise. Grey vertical lines indicate the individual CTD casts where water samples were collected for isotopic analysis, with sampling depths indicated by the black dots and station number indicated above each line. Distance is shown as cumulative distance between successive stations. WCW; Winter Cooled Waters.



Figure 3. Cross-sections across Rijpfjorden of (a) potential temperature (°C); (b) salinity (PSU); (c) fluorescence (mg m⁻³),
(d) apparent oxygen utilisation (AOU, μmol kg⁻¹), (e) oxygen saturation (%). Data from CTD casts from NP2017 cruise. Grey vertical lines indicate the individual CTD casts where water samples were collected for isotopic analysis, with sampling depths indicated by the black dots and station number indicated above each line. Distance is shown as cumulative distance between successive stations. 'If', inner fjord; 'Of', outer fjord.



Figure 4. Depth profiles of (a) Temperature (°C), (b) salinity (PSU), (c) fluorescence (mg L⁻¹), (d) nitrate (µmol L⁻¹), (e) silicate (µmol L⁻¹), (f) phosphate (µmol L⁻¹), (g) N*(µmol L⁻¹), (h) Si* (µmol L⁻¹), (i) δ¹⁵N-NO₃⁻ (‰) and (j) δ¹⁸O-NO₃⁻ (‰) from Kongsfjorden using NP2018 (circles and solid lines) and NP2017 (triangles) cruise samples. Colour denotes locations in Kongsfjorden at glacier fronts, fjord, continental shelf and slope. Note: Standard deviations of the isotopic values reported in (i) and (j) are 0.33 and 0.47 ‰ for δ¹⁵N and δ¹⁸O, respectively.



Figure 5. Depth profiles of (**a**) Temperature (°C), (**b**) salinity (PSU), (**c**) fluorescence (mg m⁻³), (**d**) nitrate (μmol L⁻¹), (**e**) silicate (μmol L⁻¹), (**f**) phosphate (μmol L⁻¹), (**g**) N*(μmol L⁻¹), (**h**) Si* (μmol L⁻¹), (**i**) δ^{15} N-NO₃⁻ (‰) and (**j**) δ^{18} O-NO₃⁻ (‰) from Rijpfjorden using NP2017 cruise samples. Colour denotes locations in Rijpfjorden (at inner fjord, outer fjord, continental shelf and slope). Note: Standard deviations of the isotopic values reported in (**i**) and (**j**) are 0.33 and 0.47 ‰ for δ^{15} N and δ^{18} O, respectively.



Figure 6. Cross-sections across Kongsfjorden of (**a**) nitrate (μmol L⁻¹), (**b**) silicate (μmol L⁻¹), (**c**) phosphate (μmol L⁻¹), (**d**) N* (μmol L⁻¹), (**e**) Si* (μmol L⁻¹), (**f**) δ^{15} N-NO₃⁻ (‰) and (**g**) δ^{18} O-NO₃⁻ (‰). Nutrient cross sections use data from NP2018 while isotope cross sections use NP2017 and NP2018 data. Distance is shown as cumulative distance between successive stations. Note: Standard deviations of the isotopic values reported in (**f**) and (**g**) are 0.33 and 0.47 ‰ for δ^{15} N and δ^{18} O, respectively.



805 Figure 7. Cross-sections across Rijpfjorden of (a) nitrate (μ mol L⁻¹), (b) silicate (μ mol L⁻¹), (c) phosphate (μ mol L⁻¹), (d) N*(μ mol L⁻¹), (e) Si* (μ mol L⁻¹), (f) δ^{15} N-NO₃⁻ (‰) and (g) δ^{18} O-NO₃⁻ (‰) using NP2017 data. Distance is shown as cumulative distance between successive stations. Note: Standard deviations of the isotopic values reported in (f) and (g) are 0.33 and 0.47 ‰ for δ^{15} N and δ^{18} O, respectively.



Figure 8. Correlation of nutrient concentrations throughout the whole water column of Kongsfjorden and Rijpfjorden. (a) nitrate (μ mol L⁻¹) *vs.* phosphate (μ mol L⁻¹) in relation to Redfield ratio (dashed lines for 1:15 and 1:16). In (a) colour coding for circles corresponds to Si:N ratio. Circles correspond to data measured in Kongsfjorden during NP2018 cruise. Triangles

- 815 correspond to data measured in Kongsfjorden during NP2017 cruise. Squares correspond to data measured in Rijpfjorden during NP2017 cruise. Dark grey squares correspond to data from eastern Fram Strait (JR17005 and FS2018 cruises). (b) nitrate (µmol L⁻¹) vs. silicate (µmol L⁻¹) in Kongsfjorden (NP2018, circles; NP2017, triangles) and Rijpfjorden (NP2017, squares). Black regression line includes all points with salinity over 33.5 PSU. Light brown regression line includes all points with salinity below 33.5 PSU. In (b) colour coding corresponds to salinity. All *p*-values ≤ 0.05. Note: Regression results are
 820 in Table P1
- 820 in **Table B1**.



825 Figure 9. Correlation of silicate (µmol L⁻¹) vs. nitrate (µmol L⁻¹) in upper water column (<100 m) within the fjord (Kongsfjorden; NP2018 - dark red circles and NP2017- dark red triangles, Rijpfjorden; dark red squares) and outside of the fjord (eastern Fram Strait; black circles). Dark red dashed line is the regression line for Kongsfjorden (NP2018,2017) and Rijpfjorden (NP2017) surface data. Black dashed line is the regression line for eastern Fram Strait (JR17005 and FS2018) surface data. All *p*- values ≤0.05. Note: Regression results are in Table B1.



Figure 10. δ^{15} N-NO₃ vs. ln (nitrate concentration) in the top 600 m in Kongsfjorden (NP2017, triangles; NP2018, circles) and 835 Rijpfjorden (NP2017, squares) with salinity as a third variable. Shapes represent locations, namely at glacier fronts (circles), fjord (triangles), continental shelf (stars) and slope (squares). The gradients of the regression lines are representative of fractionation factor (ε) for samples at glacier front (light green dotted line, $\varepsilon = 0.2\%$), fjord (dark green dotted line, $\varepsilon = 1.8$ ‰) and continental slope (magenta dotted line, $\varepsilon = 2.8$ ‰). Regression line for continental shelf samples was excluded due to insufficient data points. Black dotted line represents biological assimilation in Fram Strait where ε is equal to 4.9% as reported 840 in Tuerena et al., (2021b). Equation of the line (y = -4.9x + 17.35) was estimated knowing that the gradient was equal to 4.9 and assuming initial δ^{15} N-NO₃⁻ of 5.1‰ and initial nitrate concentration of 11.8µM as reported in Tuerena et al (2021b). Note:

Standard deviations of the isotopic values reported are 0.33 and 0.47 % for δ^{15} N and δ^{18} O, respectively.

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squares). Black dashed line is the regression line taken data from all offshore data (NP2018, NP2017, JR17005 and FS2018). All *p*-values ≤ 0.05 . Note: Rijpfjorden samples under AW influence include shelf stations (i.e., R4, R5) below 25 m depth, and slope stations (i.e., R6, R6B and R7B) below 100 m.

860 Note: Regression results are in **Table B1**. Standard deviations of the isotopic values reported in **A** and **B** are 0.33 and 0.47 ‰ for δ^{15} N and δ^{18} O, respectively.

Appendix A- Supplementary Tables and Figures

Table A1. Spatial distribution of the degree of stratification ($\Delta \rho$, kg m⁻³) in Kongsfjorden and Rijpfjorden with an indication of location (at glacier fronts, fjord, continental shelf and slope) and stations.

Kongsfjorden	Fjord (at glacier front)		Fjord		Continental S	helf	Conti	nental slo	pe
(MOSJ18)	KB5 KB6 KB7	KB3	KB2 KB	1 KBO	V12		V10	V6	
$\Delta \rho \ (\frac{kg}{m3})$	2.7 ± 0.3		1.7 ± 0.5		0.1		0	.1 ± 0.1	
Rijpfjorden			Fjord		Continental S	helf	Conti	nental slo	pe
(MOSJ17)		R1	Fjord R2	R3	Continental S R4 R5	h elf R6	Conti R6B	nental slo R7	ре R7B



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Figure A1. Seabird colonies, tidewater glaciers and river in Kongsfjorden, with an indication of the innermost NP2018 stations. Basemap Satellite image from the Norwegian Polar Institute (<u>https://geodata.npolar.no/</u>). Seabird colony data from Strøm et al (2008) and glacier names from Meslard et al. (2018).



Figure A2. T-S diagrams for Kongsfjorden and Rijpfjorden study areas (a) Temperature-salinity plot for all CTD casts from NP2018 cruise taken in Kongsfjorden. Colours are indicative of different regions along Kongsfjorden, namely, light green illustrates the fjordic region at the glacier front (i.e., closest to land); dark green are the remaining stations in the fjord; dark blue is the shelf and magenta is the continental slope and offshore. Dashed lines represent density (kg m⁻³). (b) Temperature-salinity plot for all CTD casts from NP2017 cruise taken in Rijpfjorden, Svalbard. Colours are indicative of different regions



along Rijpfjorden, namely, light green illustrates the inner fjord region; dark green are the outer fjord stations; dark blue is the shelf and magenta is the continental slope and offshore.
Polygons indicate water types. AW, Atlantic Water; AIW, Arctic Intermediate Water; PSW, Polar Surface Water; iPSW, inshore Polar Surface Water; PSWw, warm Polar Surface Water. Marginal plots show density distribution of 1m binned salinity (x-axis) and temperature (y-axis) values. Notice how at lower densities (shallower depths) polar surface waters are

885 salinity (x-axis) and temperature (y-axis) values. Notice how at lower densities (shallower depths) polar surface waters are dominant, whilst AW are more predominant at greater densities (greater depths). Water mass classification was based on that by Pérez-Hernández et al. (2017).







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Figure A4. Nitrate concentrations *vs* salinity in the top 600m in Kongsfjorden (NP2017, triangles; NP2018, circles) and Rijpfjorden (NP2017, squares). Colours represent locations, namely at glacier fronts (light green) and fjord (dark green). Note how changes in nitrate concentrations are independent of salinity. Specifically, in (A) changes in salinity are not accompanied by a change in nitrate concentrations and vice versa in (B).

Appendix B- Statistical analyses

Table B1. Regression analyses (i.e., number of observations and *p*-values) for key linear models used in this study with an indication of the figure they correlate to.

	N° of observations	p-value
Fig.8(a), Redfield ratio	127	1.82E-91
Fig.8(b), black line	57	5.61E-21
Fig.8(b), light brown line	22	3.07E-06
Fig.9, red line	30	9.57E-10
Fig.9, black line	29	3.43E-12
Fig.11(a), black line	25	4.95E-21
Fig.11(a), purple line	8	2.44E-03
Fig.11(b)	109	1.27E-35

Appendix C- Using stable isotope tools to determine N fluxes and cycling

Biologically-mediated N transformation processes result in kinetic stable isotope fractionation often with unique fractionation factors which can be exploited to delineate these processes (Sigman and Fripiat, 2019). For instance, photosynthetic uptake preferentially incorporates the lighter isotope ¹⁴N as opposed to the heavier ¹⁵N, leading to kinetic fractionation (Ryabenko, 2013). The ¹⁵N/¹⁴N ratio resulting from that fractionation is presented as $\delta^{15}N$ (**Eqn.1**).

 $\delta^{15}N(\%) = (({}^{15}N/{}^{14}N) \text{ sample } / ({}^{15}N/{}^{14}N) \text{ standard } -1) \times 1000 \text{ Eqn.1}$

where $({}^{15}N/{}^{14}N)_{standard}$ is the reference standard, measured in atmospheric N₂.

Isotopic fractionation caused by a given biological process is known as the kinetic effect, ε. According to Eqn.2, ε is defined
by the difference in rates with which the two N isotopes are converted into product in such a manner that each N cycle process also has a characteristic isotope effect (Sigman and Casciotti, 2001).

 ϵ (‰) = (¹⁴k /¹⁵k - 1) × 1000 Eqn.2

where ¹⁴k and ¹⁵k are the rate coefficients of the reaction for ¹⁴N- and ¹⁵N -containing reactant, respectively.

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This ε value can be used to distinguish two types of fractionation during assimilation. These are, firstly, Rayleigh fractionation corresponding to a closed system, and secondly, steady state fractionation, which occurs in open systems where there is resupply of nutrients (Sigman and Casciotti, 2001).

930 In addition, δ^{18} O values of N derivatives offer additional important constraints on natural processes (Kendall, 1998), where the ¹⁸O/¹⁶O ratio of nitrate is measured relative to the standard Vienna Standard Mean Ocean Water (VSMOW).

Dual isotope signatures of δ^{15} N and δ^{18} O can provide insight of nitrate inputs, such as the percentage contribution of terrestrial *vs.* marine sources as well as various N cycling processes (Sigman and Casciotti, 2001; Casciotti et al., 2002). Effect of various

- 935 N cycling processes on N and O isotopes isotopic signatures nitrate and its concentrations are summarised in **Figure C1**. The balance between N₂ fixation and denitrification renders mean δ^{15} N and δ^{18} O values of 5‰ and 2‰ respectively in the deep ocean, where δ^{15} N ranges between ~1 and 20‰ (Brandes and Devol, 2002; Sigman et al., 2009a; Tuerena et al., 2015). Values higher than 5‰ (δ^{15} N) and 2‰ (δ^{18} O) in near surface waters results from fractionation during assimilation by phytoplankton at the ocean surface which enriches the residual pools of both δ^{15} N-NO₃⁻ and δ^{18} O-NO₃⁻ (**Fig. C1**) (Altabet and Francois, 2001;
- Sigman and Casciotti, 2001). In comparison with marine nitrate, terrestrial sources of nitrate often have low values of δ^{15} N-NO₃⁻ and high values of δ^{18} O-NO₃⁻ subject to variable N inputs from N-fixation and atmospheric deposition on land (Heaton et al., 2004) and are further modified by soil processes such as denitrification and ammonia volatilisation to varying degrees (**Fig. C1a**; Carpenter et al., 1997; Kendall, 1998).
- Isotopic analyses of δ^{15} N and δ^{18} O of nitrate can also provide quantitative estimates of the regenerated nitrate pool as opposed to preformed nitrate (Sigman and Fripiat, 2019). The principle behind this is that regeneration moves the δ^{18} O-NO₃⁻ signature towards that of δ^{18} O (H₂O) while conserving the δ^{15} N-NO₃⁻ signature (Granger et al., 2018) (**Fig. C1b**). Variations in δ^{15} N-NO₃⁻: δ^{18} O-NO₃⁻ provide information about the δ^{15} N-NO₃⁻ from remineralisation (regenerated nitrate pool) and thus, also of the N exported and recycled from the euphotic zone (Sigman et al., 2005; Casciotti and Buchwald, 2012; Smart et al., 2015).
- Processes that remove nitrate such as assimilation and denitrification fractionate (δ^{15} N-NO₃⁻: δ^{18} O-NO₃⁻) in 1:1 ratio (**Fig. C1c**; DiFiore et al., 2009; Sigman et al., 2009b). The isotopic effects are best expressed in the residual nitrate pool when the processes are incomplete leaving the residual pool enriched as uptake preferentially utilises the lighter isotopes (Sigman and Casciotti, 2001). On complete biological utilisation of nitrate, the mass balance dictates that the products of the reaction (biomass) reflect the isotopic signatures of the initial nitrate source. This allows for the assessment of relative nitrate utilisation
- 955 by various uptake processes.



Figure C1. The effect of different marine N cycle processes on (a) δ^{15} N-NO₃⁻ and [NO₃⁻], (b) δ^{18} O-NO₃⁻ and [NO₃⁻] and (c) δ^{15} N-NO₃⁻ and δ^{18} O-NO₃⁻. The values of δ^{15} N-NO₃⁻ and δ^{18} O-NO₃⁻ designated by the black dot in (a), (b) and (c) represent global oceanic values, namely 5‰ and 2‰, respectively.

Dashed arrows denote processes that add or remove fixed N from the ocean, while solid arrows denote a component of the internal cycling of oceanic fixed N.

In (a), the orange shading represents variation of δ^{15} N-NO₃⁻ caused by the δ^{15} N of the organic N being remineralised.

In \mathbb{O} , the blue dashed line represents the 1:1 ratio at which N-assimilation and denitrification occur. Moreover, the orange shading represents variation in the δ^{15} N: δ^{18} O ratio of the organic N being remineralised.

Compiled from Sigman and Fripiat (2019), Ryabenko (2013), Dähnke and Thamdrup (2013) and Sigman and Casciotti (2001).

Appendix D- Calculations for the quantitative measure of the contribution of terrestrial inputs vs. marine inputs to the fjord nitrate pool

Step 1- Calculating the percentage of regenerated nitrate in the Winter Cooled Water (WCW)

960

1) Using equation in Tiwari et al. (2018) (Eqn.1).

 $\delta^{18}O(H_2O)=0.54S-18.42=0.3$ ‰ Eqn.1 where S in WCW= 34.8 ± 0.006 PSU (NP2018)

965

Previous field and modelling studies have used a nitrifying δ^{18} O value of **1.1% plus** δ^{18} O_{H20} (Buchwald et al., 2012; Sigman et al., 2009b; Eqn.2). Thus,

 $\delta^{18}O_{reg}(H_2O)=1.1\% + 0.3\% = 1.4\%$ in WCW Eqn.2

975

2) Using equation in Tuerena et al. (2021)

Since regeneration is calculated in the most saline and dense marine waters of the fjords the δ^{18} O values of Atlantic Waters can be used (Granger et al., 2018; Tuerena et al., 2021). Thus, the following equation from Tuerena et al. (2021) (Eqn.3) was used to calculate the proportion of regenerated nitrate (NO₃⁻reg/NO₃⁻tot), and resulted in a value of 50-80% regeneration (65% average).

$$\frac{NO3_{reg}}{NO3_{tot}} = \frac{(\delta^{18}O_{meas} - \delta^{18}O_{AW})}{(\delta^{18}O_{reg} - \delta^{18}O_{AW})}$$
 Eqn.3

980

where $\delta^{18}O_{meas} = \delta^{18}O-NO_3^-$ measured in WCW= 1.9 ± 0.2% (data at >300 m, n=3) and $\delta^{18}O_{AW} = 2.8 \pm 0.3\%$ (Tuerena et al., 2021). δ^{18} Oreg was 1.4‰, calculated using Eqns.1-2 and NO_{3reg}/NO_{3tot}= proportion of regenerated nitrate.

 $= 65 \pm 15 \%$

985

Step 2 - Calculating δ^{15} N_{reg}

The 65% regeneration estimate was then used to determine the δ^{15} N of remineralised NO₃⁻ (*x*, Eqn.4). The δ^{15} N of preformed NO₃ is given as a range between the conservative estimate of 4.3 ± 0.1 ‰ (terrestrial estimate in this study) and 5.1 ± 0.1 ‰ 990 (marine endmember; Tuerena et al., 2021) which assumes all preformed nitrate is of marine origin.

$$\delta^{15} N_{WCW} = \frac{65}{100} x + \left(\frac{35}{100} \times \delta^{15} N_{\text{preformed}}\right)$$
 Eqn. 4

where x is δ^{15} N of remineralised NO₃⁻, δ^{15} N_{WCW} = δ^{15} N-NO₃⁻ measured in WCW= 4.2 ± 0.2 ‰ (data at >300 m, n=3), δ^{15} N_{preformed} ranges between 4.3 ± 0.1 and 5.1 ± 0.1 ‰ (Tuerena et al., 2021), $\frac{65}{100}$ is the proportion of regenerated nitrate and $\frac{35}{100}$ is the proportion of preformed nitrate (derived from Eqn. 5). 995

(1) Conservative estimate (i.e., most of the preformed (2) Assuming that all of the preformed nitrate is marine

nitrate is terrestrial)	Regenerated Preformed
Regenerated Preformed	$\delta^{15} N_{WCW} = \frac{65}{100} x + (\frac{35}{100} \times 5.1 \pm 0.1 \%)$
$\delta^{15} \mathbf{N}_{\text{WCW}} = \frac{65}{100} \chi + \left(\frac{35}{100} \times 4.3 \pm 0.1 \%\right)$	$4.2 \pm 0.2 \% = \frac{65}{100} x + \left(\frac{35}{100} \times 5.1 \pm 0.1 \%\right)$
$4.2 \pm 0.2 \% = \frac{65}{100} x + \left(\frac{35}{100} \times 4.3 \pm 0.1 \%\right)$	$x=3.7\pm0.4$ ‰
$x=4.1\pm0.4$ ‰	

The resulting δ^{15} N values of 3.7- 4.1 ‰ of remineralised NO₃⁻ demonstrates the major contribution of terrestrial sources (63-88%) to fjord primary productivity as opposed to marine sources (12-37%) (summarised in Table 1). This is explained by the fact that terrestrial sources occur in constant supply, whereas uptake of marine nutrients is hindered by the strong halocline that develops during the summer (see section 4.5).

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