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FEATURE ARTICLE

Feeding-related controls on microbial nitrogen cycling associated with the Arctic marine copepod *Calanus hyperboreus*

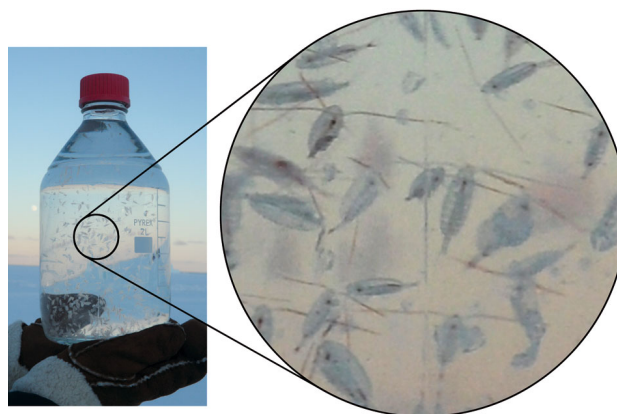
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ABSTRACT: The copepod *Calanus hyperboreus*, a key species of Arctic marine ecosystems, has a partially anoxic gut that is suspected to host anaerobic microbial activities. So far, however, only dead specimens have been studied in which gut processes are quickly abolished by microbial carcass degradation. Here, live specimens were exposed to different feeding regimes and ambient oxygen levels to study the controls on copepod-associated microbial nitrogen cycling in ¹⁵N-enrichment experiments. Algae-fed copepods exhibited consistently high rates of denitrification and low or variable rates of dissimilatory nitrate reduction to ammonium (or nitrite), nitrification, and nitrous oxide release. Thus, live *C. hyperboreus* contribute to pelagic fixed-nitrogen loss through denitrification that is otherwise absent from oxygenated Arctic surface waters. The feeding-related controls on copepod-associated anaerobic nitrogen cycling are supported by significantly reduced rates in starved copepods. In addition, freshly released fecal pellets showed similar patterns of nitrogen cycling as fed copepods. Unlike in previous studies on copepod carcasses, low ambient oxygen levels did not stimulate anaerobic nitrogen cycling in live specimens, supporting that the *C. hyperboreus* gut is anoxic, irrespective of ambient oxygen levels. Fecal pellets and carcasses retained denitrification activity for 4 d of simulated sinking. We estimate that during the time of the Arctic spring bloom, pelagic denitrification associated with live specimens, fecal pellets, and carcasses of *C. hyperboreus* is equivalent to ~12% of the benthic fixed-nitrogen loss. This estimate increases to ~28%, provided that the abundant sibling species *C. glacialis* and *C. finmarchicus* exhibit denitrification activities in proportion to their smaller body size.



A bottle full of copepods dominated by *Calanus hyperboreus* specimens (collected in Disko Bay, West Greenland) in which microbial denitrification activity was found.

Photo: Peter Stief

KEY WORDS: Copepods · Arctic · Denitrification · Nitrification · Nitrous oxide · Oxygen · Feeding · Fecal pellets · Carcasses · ¹⁵N stable-isotope labeling

INTRODUCTION

The zooplankton community in Arctic marine ecosystems is dominated by the co-occurring calanoid copepod species *Calanus hyperboreus*, *C. glacialis*, and *C. finmarchicus* (Hirche 1991, Madsen et al. 2001). The timing of the Arctic phytoplankton spring bloom dictates the life cycle of these herbivores, which show peak abundances in surface waters

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around the time of the bloom (Conover 1988, Swalethorp et al. 2011). Aside from holding a central position in the Arctic food chain (Falk-Petersen et al. 2009), calanoid and other copepods play an important role in pelagic biogeochemistry as colonization sites for diverse microorganisms (Møller et al. 2007, Tang et al. 2010, Turner 2015). Major biogeochemical implications of the elevated microbial densities and activities on copepods include mineralization and respiration processes that may result in microscale O_2 depletion inside the gut (Tang et al. 2011), decaying carcasses (Glud et al. 2015), and fecal pellets (Alldredge & Cohen 1987), despite high O_2 levels in the surrounding water. Notably, the potentially anoxic interior of copepods and fecal pellets might host anaerobic microbial activities that are not present in the oxygenated water column.

Recently, intense anaerobic nitrogen cycling (N-cycling) was demonstrated in sinking phytodetritus aggregates that had an anoxic center, especially when exposed to low-oxygen conditions (Klawonn et al. 2015, Kamp et al. 2016, Stief et al. 2016). Similarly, sinking carcasses of diverse copepod species from different aquatic ecosystems displayed significant anaerobic N-cycling (Glud et al. 2015, Stief et al. 2017, 2018). The rates of carcass-associated anaerobic N-cycling increased with decreasing O_2 concentrations in the surrounding water, suggesting that the spatial extent of internal anoxia in copepod carcasses is mainly controlled by ambient O_2 levels. In *C. finmarchicus* carcasses, for instance, denitrification activity increased exponentially with decreasing ambient O_2 levels (Glud et al. 2015). The denitrification pathway reduces NO_3^- to N_2 gas, which leads to fixed-nitrogen loss from aquatic ecosystems because of the relatively low bioavailability of N_2 . In small marine and freshwater copepod carcasses, dissimilatory nitrate reduction to ammonium (DNRA) and dissimilatory nitrate reduction to nitrite (DNRN) appear to be important anaerobic N-cycle pathways (Stief et al. 2017, 2018). These 2 pathways do not directly contribute to fixed-nitrogen loss. In strictly anoxic settings, however, carcass-associated DNRA and DNRN may supply free-living anammox bacteria with NH_4^+ and NO_2^- , respectively, which are then converted to N_2 (Thamdrup 2012). Copepod carcasses may thus contribute directly and indirectly to pelagic fixed-nitrogen loss, depending on the microbial populations colonizing them and the ambient O_2 levels. Fixed-nitrogen loss coupled to pelagic microsites is commonly overlooked by incubation-based rate measurements due to pre-filtration of water samples or small incubation volumes that exclude the presence of copepods,

fecal pellets, and phytodetritus aggregates. Thus, by taking into account the fixed-nitrogen loss coupled to copepods and other pelagic microsites, the models of oceanic nitrogen dynamics might be considerably improved, especially for low-oxygen environments (Bianchi et al. 2018).

For calanoid copepods in subarctic waters, a relative carcass abundance of ~25% has been found (Yamaguchi et al. 2002). Daase et al. (2014) report highly variable relative carcass abundances ranging from 9–14% in an Arctic fjord to as high as 94% in a basin of the Arctic Ocean at 200–3000 m water depth. Thus, the total contribution of copepods to pelagic N-cycling is in any case higher than estimated from rate measurements with carcasses only. It is not known, however, whether live copepods display the same N-cycling patterns and rates as dead copepods. Anaerobic N-cycling is expected to occur either inside the anoxic gut of live copepods (Tang et al. 2011), or inside the disintegrating body of dead copepods (Glud et al. 2015, Stief et al. 2017, 2018). In either case, internal anoxia may establish due to high O_2 consumption rates of microorganisms colonizing the copepods, whereas labile organic carbon would be either food derived and continuously supplied by the copepod's feeding activity, or body derived and thus finite.

In this study, the hypothesis is tested that anaerobic N-cycling associated with the large marine copepod *C. hyperboreus* is subject to feeding-related controls. To this end, live copepods were exposed to different feeding regimes and the rates of anaerobic N-cycling were measured in ^{15}N -incubations. In addition, live copepods were exposed to low-oxygen conditions to test whether anaerobic N-cycling in the gut or on the exoskeleton can be further stimulated or induced, respectively. Patterns of N-cycling were compared between algae-fed copepods and freshly released fecal pellets. The persistence of anaerobic N-cycling associated with fecal pellets and copepod carcasses was followed for a simulated sinking period of 4 d. The results of these experiments were used for a first assessment of the *in situ* contribution by live specimens, fecal pellets, and carcasses of *C. hyperboreus* to fixed-nitrogen loss in Arctic marine ecosystems.

MATERIALS AND METHODS

Collection of study organisms

Calanus hyperboreus was sampled in Disko Bay near the southern tip of Disko Island off West Green-

land (69° 14' 15" N, 53° 31' 15" W) in April 2014, April 2015, and February 2016. Zooplankton was collected by vertical hauls (150–0 m) with a WP-2 zooplankton net (200 µm mesh size) with a large non-filtering cod end through a hole in the sea ice. Seawater was collected with a 30 l Niskin bottle at 150 m water depth. During transport to the laboratory, zooplankton and seawater were kept at *in situ* temperature in a cooling box. Live females of *C. hyperboreus* were isolated in an ice-cooled tray within ~6 h of collection. In a thermostated room (2.5 ± 0.5°C), the copepods were maintained at a density of 3 individuals (ind.) l⁻¹ in 10 l buckets filled with aerated *in situ* seawater and fed once per day with the cryptophyte *Rhodomonas salina* (2014) or the diatom *Skeletonema marinoi* (2015) until used for experiments at the Arctic Station (Qeqertarsuaq, Disko Island, Greenland; University of Copenhagen). In 2016, live copepods were transported to the University of Southern Denmark (Odense, Denmark) where they were used for experiments. These copepods were maintained at a density of 3 ind. l⁻¹ in 1 l bottles filled with aerated artificial seawater (ASW; Red Sea Aquatics) stored in a fridge (4.0 ± 0.5°C) and fed every second day with *S. marinoi*. Before each feeding event, one-third of the seawater in the maintenance buckets and bottles was discarded and replaced with fresh seawater and dead specimens were removed. The 2 algae species, *S. marinoi* (CCMP 1332, NCMA) and *R. salina*, had been cultured in F/2 medium prepared in 0.2-µm-filtered, autoclaved seawater with and without silicate addition, respectively (Guillard & Ryther 1962).

The maintenance density of *C. hyperboreus* of 3 ind. l⁻¹ was high compared with *in situ* densities of up to 0.09 ind. l⁻¹ as observed in Disko Bay during the spring bloom (Swailethorp et al. 2011), but similar to very high calanoid copepod densities encountered in single net hauls (Madsen et al. 2001). The animals appeared to behave and feed normally as most specimens had completely filled guts when retrieved for experiments. Respiration and excretion rates were within the previously reported ranges (Gilfillan et al. 1986, Auel et al. 2003) and were not significantly different between specimens maintained in large and small maintenance containers (respiration: $t = 0.053$, $df = 29$, $p = 0.958$; excretion: $t = -0.017$, $df = 20$, $p = 0.987$).

Overview and preparation of experiments

¹⁵N-stable-isotope enrichment experiments in natural seawater or in ASW were made with live and dead specimens and fecal pellets of *C. hyperboreus* (if applicable, fed with *R. salina* or *S. marinoi*) depending on availability of animals, algae, and seawater during 3 measurement campaigns in 2014, 2015, and 2016 (Table 1).

- The effect of filled versus empty gut (i.e. gut filling degree) of live specimens on copepod-associated N-cycling was addressed in 2 experimental series. In 2014, the algae-fed group of copepods was fed with *R. salina* at a saturating cell density of ~50 000 cells ml⁻¹ at least 24 h before and also during ¹⁵NO₃⁻-incubations in non-filtered seawater,

Table 1. Overview of experiments on microbial nitrogen cycling associated with *Calanus hyperboreus* (fed with *Rhodomonas salina* or *Skeletonema marinoi*) and its fecal pellets and carcasses. Wet wt: wet weight of single specimens; data are means ± SE of 3–12 batches of 1–3 specimens. ASW: artificial seawater. Oxix: 85–95% air saturation (AS); hypoxic: 10–25% AS; near-anoxic: 0–4% AS

Year	Seawater	¹⁵ N-tracer	Experimental treatments	Wet wt (mg)	n	Algae species	
2014	Disko Bay (150 m)	¹⁵ NO ₃ ⁻ (10 µM)	Live copepods	Algae-fed, oxix	15.2 ± 0.3	(20)	<i>R. salina</i>
				Algae-starved, oxix	16.3 ± 0.5	(20)	–
				Algae-fed, hypoxic	15.1 ± 1.0	(14)	<i>R. salina</i>
			Dead copepods	Fresh	11.8 ± 1.4	(12)	<i>R. salina</i>
				Aged	10.4 ± 1.5	(12)	<i>R. salina</i>
			Fecal pellets	Fresh	No data		<i>R. salina</i>
				Aged	No data		<i>R. salina</i>
2015	Disko Bay (150 m)	¹⁵ NO ₂ ⁻ (5 µM)	Live copepods	Algae-fed, oxix	13.8 ± 1.0	(15)	<i>S. marinoi</i>
				Algae-starved, oxix	11.9 ± 0.7	(15)	–
		0.2 µm filtered	¹⁵ NH ₄ ⁺ (5 µM)	Bacteria-starved, oxix	13.0 ± 0.4	(15)	–
2016	ASW	¹⁵ NO ₂ ⁻ (5 µM)	Live copepods	Algae-fed, oxix	13.4 ± 1.1	(30)	<i>S. marinoi</i>
				Algae-fed, near anoxic			<i>S. marinoi</i>

while the algae-starved control group was kept in non-filtered seawater without algae addition. In 2015, the algae-fed group of copepods was fed with *S. marinoi* at a saturating cell density of $\sim 15\,000$ cells ml^{-1} at least 24 h before and also during $^{15}\text{NO}_2^-$ -incubations in non-filtered seawater, while the algae-starved and the bacteria-starved control groups were kept in non-filtered seawater without algae addition and in 0.2- μm -filtered seawater with all bacteria removed, respectively. The particulate organic carbon (POC) levels in the 2 algae-fed treatments in the 2014 and 2015 measurement campaigns were estimated at 1.125 and 1.15 $\mu\text{g C ml}^{-1}$, respectively (Olenina et al. 2006). The POC levels in the algae-starved and bacteria-starved control groups were estimated at 0.03 and 0 $\mu\text{g C ml}^{-1}$, respectively (Fukuda et al. 1998). The gut filling degree of the transparent copepods was assessed visually and is reported as the percentage gut length that was filled with food particles.

- The effect of the ambient oxygen level on N-cycling associated with live copepods was studied in 2 experimental series. In 2014, algae-fed specimens were exposed to 85–95 % (oxic) versus 10–25 % air saturation (hypoxic) during $^{15}\text{NO}_3^-$ -incubations in non-filtered seawater. In 2016, algae-fed specimens were exposed to 85–95 % (oxic) versus 0–4 % air saturation (near anoxic) during $^{15}\text{NO}_2^-$ -incubations in 0.2- μm -filtered ASW.
- N-cycling associated with fresh versus aged fecal pellets was studied in 2014 by feeding *C. hyperboreus* kept in a 10 l bucket with *R. salina* for 24 h and then transferring the algae-fed specimens to a new bucket with non-filtered seawater. Fecal pellets that were released during the following 12 h and 96 h, respectively, were collected by sieving onto a 125 μm mesh. Afterwards, fecal pellets were incubated in $^{15}\text{NO}_3^-$ -enriched, non-filtered seawater under oxic conditions.
- N-cycling associated with fresh versus aged carcasses (i.e. dead copepods) was studied in 2014 after killing algae-fed specimens in deoxygenated seawater for 1 h and then keeping them suspended in aerated, non-filtered seawater on a plankton wheel for 24 h and 96 h, respectively. Afterwards, the carcasses were incubated in $^{15}\text{NO}_3^-$ -enriched, non-filtered seawater under oxic conditions. Note that only about one-third of the specimens died during the 1 h exposure to anoxia.

Live and dead copepods were incubated in batches of 1–3, while fecal pellets were incubated in batches of 60–120 in 25 ml glass bottles (see next section). Control experiments without copepods and fecal pel-

lets were run in parallel in non-filtered (if applicable, containing *R. salina* or *S. marinoi*) or 0.2- μm -filtered seawater.

Rate measurements

The live and dead copepods and fecal pellets that had been prepared as described in the previous section were used in 24 h incubation experiments for quantification of their O_2 and N-turnover rates. The 25 ml glass bottles that were used as incubation vials had oxygen-sensitive optodes (SensorSpot, PyroScience) fixed on the inside, which allowed contactless O_2 measurements (Firesting, PyroScience). The bottles were filled with seawater that was specifically adjusted for each incubation experiment regarding filtration, ^{15}N -tracer amendment, algae addition, and ambient O_2 level (Table 1). Filtration of seawater for removal of microorganisms was done with Monta Mil® nylon membrane filters (0.2 μm ; Frisenette) and using acid-washed glassware. ^{15}N -tracers (Sigma-Aldrich) were added at final concentrations of 10 μM ($^{15}\text{NO}_3^-$) and 5 μM ($^{15}\text{NO}_2^-$, $^{15}\text{NH}_4^+$). Background NO_3^- concentrations in the *in situ* and ASW were 13.2 μM and 1.3 μM , respectively. Background NO_2^- and NH_4^+ concentrations were generally < 0.2 μM . The addition of $^{15}\text{NO}_2^-$ and $^{15}\text{NO}_3^-$ increased the *in situ* concentration of oxidized dissolved inorganic nitrogen by 38 % and 76 %, respectively, which may have stimulated the activity of anaerobic N-cycle pathways to some degree. The added dose of $^{15}\text{NO}_2^-$ was not expected to have toxic effects on copepods and bacteria, since the lethal concentration that kills 50 % of the exposed population (LC_{50}) is at least 1–2 and 3–4 orders of magnitude higher, respectively, than used here (Camargo & Alonso 2006, Lotti et al. 2012). The addition of $^{15}\text{NH}_4^+$ was high compared with *in situ* NH_4^+ concentrations, which may have stimulated nitrification activity substantially, but the nitrification rates measured here were still very low.

The algae-fed treatments received aliquots of the respective algae culture to arrive at cell densities of *R. salina* ($\sim 50\,000$ cells ml^{-1}) and *S. marinoi* ($\sim 15\,000$ cells ml^{-1}). Before addition, algae cultures were washed to remove excess nutrients. To this end, the cultures were centrifuged ($1000 \times g$, 20 min), re-suspended in seawater, and subsampled for microscopic cell counts. After algae addition, the seawater was split into 2 portions, one to be degassed for 20 min with O_2 and the other one with He or Ar (depending on availability). Degassing served to (1) lower the natural N_2 background for improving

denitrification measurements at low rates and (2) allow the adjustment of the targeted ambient O_2 level by mixing the appropriate volumes of O_2 - and He- or Ar-flushed seawater. This procedure worked for most of the experiments, with the exception of the denitrification measurements in 2015 because the used Ar gas had a higher $^{30}N_2$: $^{28}N_2$ ratio than air.

The 25 ml glass bottles were filled with the prepared seawater, copepods, or fecal pellets and sealed with butyl rubber stoppers without entrapping gas bubbles. The bottles were mounted on a plankton wheel to keep the copepods, fecal pellets, or carcasses in suspension and were then incubated in darkness at $2.5 \pm 0.5^\circ C$ for up to 24 h. This incubation temperature is very similar to the *in situ* temperature that *C. hyperboreus* is exposed to in Disko Bay at a water depth of 150 m during spring–early summer (Swalethorp et al. 2011). At time intervals of 1.5–3.0 h for the first 6–12 h and again at the end of the incubations (i.e. after 24 h), O_2 measurements were made and water samples for N-analyses were withdrawn through the stopper as described in detail in Stief et al. (2016). Ambient O_2 concentrations were maintained at stable levels by re-injecting seawater adjusted to an O_2 concentration that would compensate the concentration decrease observed since the last sampling time point (Fig. S1 in the Supplement at www.int-res.com/articles/suppl/m602p001_supp.pdf). The water sample withdrawn was used for analyses of N_2 , N_2O , NO_3^- , NO_2^- , and NH_4^+ .

Nitrogen analyses

Nitrate plus nitrite concentrations were measured on a NO_x analyzer (CLD 60, Eco Physics) using the VCl_3 reduction assay (Braman & Hendrix 1989). Nitrite concentrations were measured with the Griess reaction and using a microplate reader (Multiskan GO Photometer, Thermo Fisher Scientific) after García-Robledo et al. (2014). Nitrate concentrations were then obtained by subtraction. Ammonium was analyzed with the salicylate method (Bower & Holm-Hansen 1980). Isotopically labeled dinitrogen ($^{15}N-N_2$) was analyzed in the headspace of 3 ml exetainers (Labco) on a gas chromatography-isotopic ratio mass spectrometer (Thermo Delta V Plus, Thermo Fisher Scientific) (Dalsgaard et al. 2012) with the excess above natural abundance calculated according to Nielsen (1992). Nitrous oxide concentrations were measured in the headspace of the same exetainers on a gas chromatograph (GC 7890, Agilent Technologies) and corrected for the prior headspace sampling

for $^{15}N-N_2$ analysis and the proportion of N_2O dissolved in the seawater. ^{15}N -labeled NO_3^- , NO_2^- , and NH_4^+ were analyzed with the cadmium/sulfamic acid, sulfamic acid, and hypobromite assay, respectively (Warembourg 1993, McIlvin & Altabet 2005, Füssel et al. 2012).

Calculations and statistics

Rates of oxygen and nitrogen turnover in the 25 ml bottles were calculated from linear concentration changes, which in many cases were limited to the first 4–5 sampling time points (i.e. until 8–12 h after the start of the incubations). The dilution effect on solutes and gases caused by the sampling process was accounted for in the calculations (Stief et al. 2016). Oxygen and nitrogen turnover directly associated with the copepods and fecal pellets were approximated by subtracting turnover rates (if any) measured in control incubations with seawater only.

The total denitrification rate ($P^{Total}N_2$) was calculated from $^{15}NO_3^-$ and $^{15}NO_2^-$ incubations using the principles of random isotope pairing in the absence of anammox (Nielsen 1992) as $P^{Total}N_2 = (P^{29}N_2)^2 / (4 \times P^{30}N_2) + P^{29}N_2 + P^{30}N_2$. The concurrent production of $^{15}N-N_2$, $^{15}NO_2^-$, and $^{15}NH_4^+$ indicated that aside from denitrification, DNRN and DNRA were also occurring (i.e. rather than $^{15}NO_3^-$ assimilation followed by $^{15}NO_2^-$ and $^{15}NH_4^+$ recycling due to the decay of microbial biomass, which would have occurred with a time delay). The net DNRN rate was calculated from the $^{15}NO_2^-$ production rate divided by the initial $^{15}NO_3^-$ -labeling fraction in $^{15}NO_3^-$ incubations. The net DNRA rate was calculated from the $^{15}NH_4^+$ production rate divided by the initial $^{15}NO_3^-$ - or $^{15}NO_2^-$ -labeling fraction in $^{15}NO_3^-$ and $^{15}NO_2^-$ incubations, respectively. The nitrification rate was calculated in 2 ways: (1) for $^{15}NH_4^+$ incubations, the $^{15}NO_3^- + ^{15}NO_2^-$ production rate was divided by the initial $^{15}NH_4^+$ -labeling fraction to arrive at the total nitrification rate; and (2) for $^{15}NO_3^-$ incubations, the $^{14}NO_3^-$ production rate was calculated as the temporal change in the difference between the total NO_3^- concentration and the $^{15}NO_3^-$ concentration to arrive at the rate of NO_2^- oxidation, the second step of nitrification. The net NH_4^+ release rate was calculated from the increase in total NH_4^+ concentrations. No data are available for (1) DNRN in 2015 and 2016 due to lack of $^{15}NO_3^-$ incubations and (2) denitrification in 2015 due to the excessively high $^{30}N_2$: $^{28}N_2$ ratio of the used Ar gas (see previous section).

Significant differences in mean rates that had been measured in independent treatments were determined by Student's *t*-tests (2 factor levels) and 1-way ANOVA (3 factor levels) with $\alpha = 0.05$. Homogeneity of variances was tested with Levene's test and, depending on the outcome, Tukey's and Games-Howell's tests for homogeneous and non-homogeneous variances, respectively, were used as posterior tests. Significant differences in mean rates that had been measured at 3 consecutive time points in the same experiment were determined by paired-sample *t*-tests with a Bonferroni-corrected $\alpha = 0.05/3$. One-sample *t*-tests with $\alpha = 0.05$ were used to check whether rates were significantly different from zero. All statistical analyses were performed with SPSS Version 11.0.

RESULTS

Nitrogen-cycle pathways

In incubations with live, algae-fed *Calanus hyperboreus* from the campaign in 2014, the rates of O_2 uptake and net NH_4^+ release (as combined metabolic activities of the animal and its associated microorganisms) by far exceeded the rates of microbially mediated N-cycling (Fig. 1). The copepod-associated microbial N-cycling itself was clearly dominated by denitrification with activities ~20-fold higher than nitrification, DNRA, DNRN, and net N_2O release. During the campaigns in 2015 and 2016, much higher DNRA rates were measured, which in 1 case rivaled denitrification rates (see below).

The patterns of O_2 and N-turnover associated with freshly released (≤ 12 h) fecal pellets resembled the patterns observed for algae-fed copepods, except for the complete absence of net NH_4^+ release and nitrification activity (i.e. the rates equaled zero after subtraction of the minute rates measured in the respective controls). Based on reported length ranges of the gut and fecal pellets of female adults of 5.5–7.0 mm and 400–700 μm , respectively (Urban-Rich 1999, Juul-Pedersen et al. 2006, Olli et al. 2007, Tang et al. 2011), we estimate that a completely filled gut corresponds to 10 fecal pellets. Nitrogen turnover rates of 10 freshly released (and then incubated under oxic conditions) fecal pellets make up as much as 40%, 72%, and 28% of copepod-associated denitrification, DNRA, and DNRN activity, respectively (Fig. 1). In contrast, fecal-pellet-associated microbial N-turnover rates make up only 7% of the copepod-associated net N_2O release and 0% of net NH_4^+ release and nitrification.

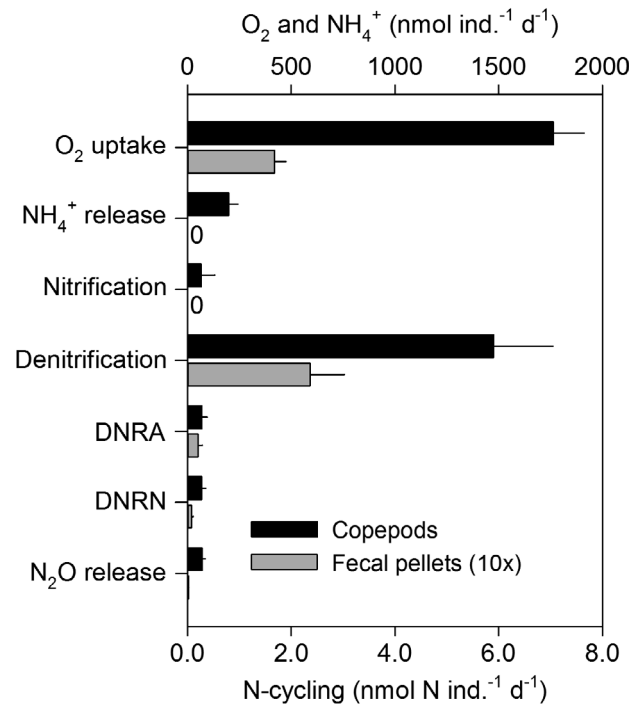


Fig. 1. Oxygen and nitrogen turnover associated with *Calanus hyperboreus* and its fecal pellets. Algae-fed copepods and freshly released fecal pellets were incubated in $^{15}NO_3^-$ -enriched seawater under oxic conditions, in darkness, and at 2.5°C for up to 24 h. Rates are expressed per 1 copepod specimen or per 10 fecal pellets. Means \pm SE of $n = 12$ (copepods) and $n = 5$ (fecal pellets) incubations are shown. Note different scales on the 2 x-axes. DNRA/DNRN: dissimilatory nitrate reduction to ammonium/nitrite, respectively

Since only total N_2O concentrations were measured by gas chromatography, the microbial N_2O source could not be identified from any ^{15}N -labeling. Thus, the net N_2O release by live copepods was plotted against nitrification and denitrification activity in search for covariation (Fig. 2A,B). However, no significant correlation was found between net N_2O release and either nitrification or denitrification activity in 3 different live copepod treatments (Table S1 in the Supplement). Instead, highly significant correlations between net N_2O and NH_4^+ release were found for algae-fed and algae-starved copepods under oxic conditions (Fig. 2C, Table S1).

Gut filling degree

The starvation of copepods for algae before and during the ^{15}N -incubations reduced the gut filling degree from $81.3 \pm 6.3\%$ (mean \pm standard error) to $5.0 \pm 1.7\%$ (campaign in 2014) and from $56.3 \pm 14.4\%$

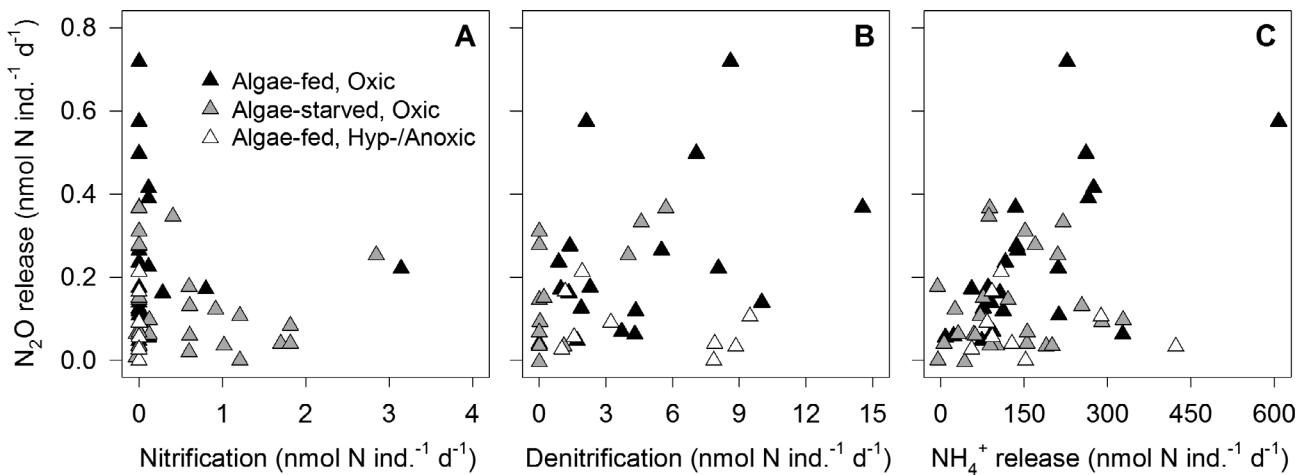


Fig. 2. Nitrous oxide release from *Calanus hyperboreus*. Correlation between net N_2O release and (A) nitrification activity, (B) denitrification activity, and (C) net NH_4^+ release associated with live copepods. Incubation conditions as specified in 'Materials and methods' and Table 1. The only significant correlations were found between net N_2O release and (C) net NH_4^+ release by algae-fed and algae-starved specimens under oxidic conditions (Table S1 in the Supplement)

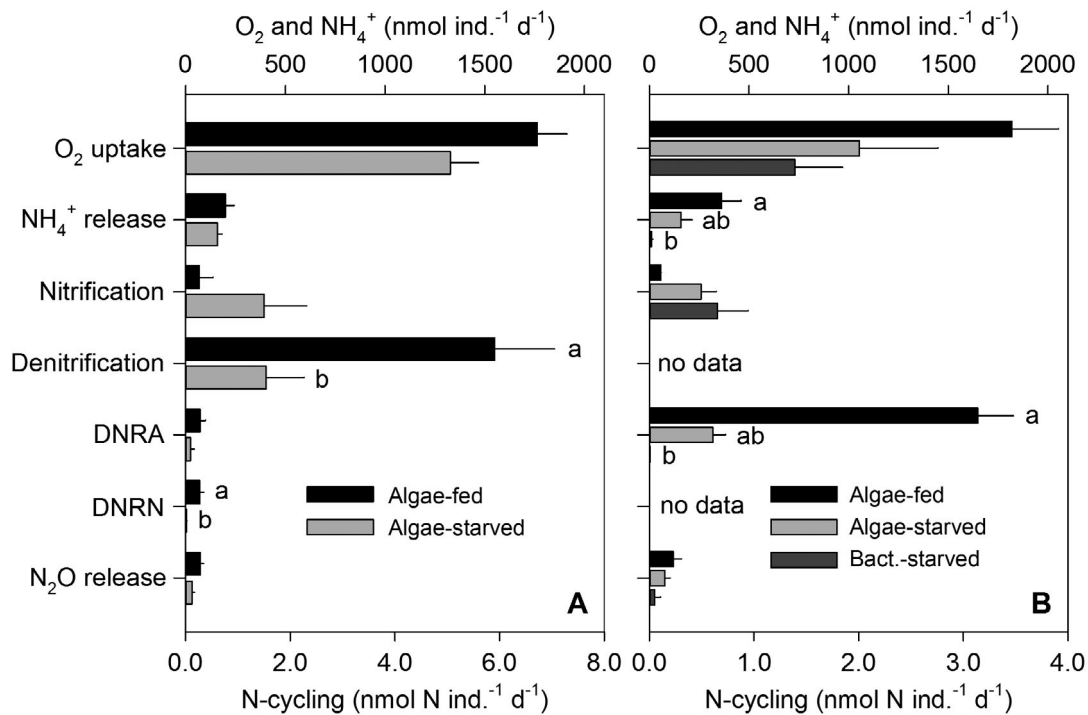


Fig. 3. Effect of gut filling degree. Algae-fed, algae-starved, and bacteria-starved *Calanus hyperboreus* were incubated in (A) $^{15}\text{NO}_3^-$ -enriched or (B) $^{15}\text{NO}_2^-$ -enriched seawater. Incubation conditions as specified in 'Materials and methods' and Table 1. Means \pm SE of $n = 5-12$ (algae fed), $n = 5-10$ (algae starved), and $n = 5$ (bacteria starved) incubations are shown. Note different scales on the 2 x-axes. Lower-case letters indicate statistically significant differences between algae-fed, algae-starved, and bacteria-starved copepods. DNRA/DNRN: dissimilatory nitrate reduction to ammonium/nitrite, respectively

to $5.0 \pm 3.3\%$ (campaign in 2015). The additional starvation of copepods for seawater bacteria and particles larger than $0.2 \mu\text{m}$ reduced the gut filling degree further to $3.3 \pm 2.0\%$ (campaign in 2015). The reduced gut filling degree led to significantly reduced

rates of denitrification, DNRA, DNRN, and net NH_4^+ release (Fig. 3; Table S2 in the Supplement), though not necessarily in both measurement campaigns (i.e. DNRA and NH_4^+ release). Rates of O_2 uptake, nitrification, and net N_2O release were not significantly

changed by differences in the gut filling degree (Fig. 3, Table S2).

Ambient oxygen level

Oxic (85–95 % air saturation), hypoxic (10–25 % air saturation), and near-anoxic (0–4 % air saturation) conditions were tested for their effect on copepod-associated, anaerobic N-cycling that may be stimulated by the expansion of internal anoxia under low-oxygen conditions (Fig. 4). Note that in the near-anoxic treatment, fully anoxic conditions were achieved at the last sampling time point (Fig. S1C). In any case, no significant effect of low-oxygen conditions on denitrification, DNRA, and DNRN was observed (Fig. 4A,B; Table S3 in the Supplement), which supports that the gut of *C. hyperboreus* is anoxic even at high ambient O_2 levels. In contrast, low-oxygen conditions led to significantly lower rates of copepod-associated net N_2O release (Fig. 4, Table S3) and nitrification rates that equaled zero (Fig. 4). Taken together, these observations suggest that copepod-associated denitrification is not a source of N_2O , but nitrification potentially is. The mortality of *C. hyperboreus* was ~70% at the end

of the hypoxic treatment and 100% after 3 h of the near-anoxic treatment.

Temporal trends

Nitrogen cycling associated with copepods and fecal pellets was followed for 4 d after death and defecation, respectively (Fig. 5; Table S4 in the Supplement). One day after killing, N-cycling rates were not significantly different between carcasses and live copepods, indicating that the relative partitioning between different N-cycling pathways also remained the same (Fig. 5A, Table S4). Furthermore, N-cycling rates were not significantly different between 1-d-old and 4-d-old carcasses (Fig. 5A, Table S4). Noteworthy, the 4-d-old carcasses still remained net sinks of NO_x^- (Fig. S2A in the Supplement). The wet weight of the 4-d-old carcasses was about one-third lower than that of live copepods (Table 1). The only significant change experienced by aging fecal pellets was a strong increase of nitrification activity (Fig. 5B, Table S4), which turned the fecal pellets from a net sink into a net source of NO_x^- (Fig. S2B), despite the stable denitrification activity associated with fecal pellets (Fig. 5B, Table S4).

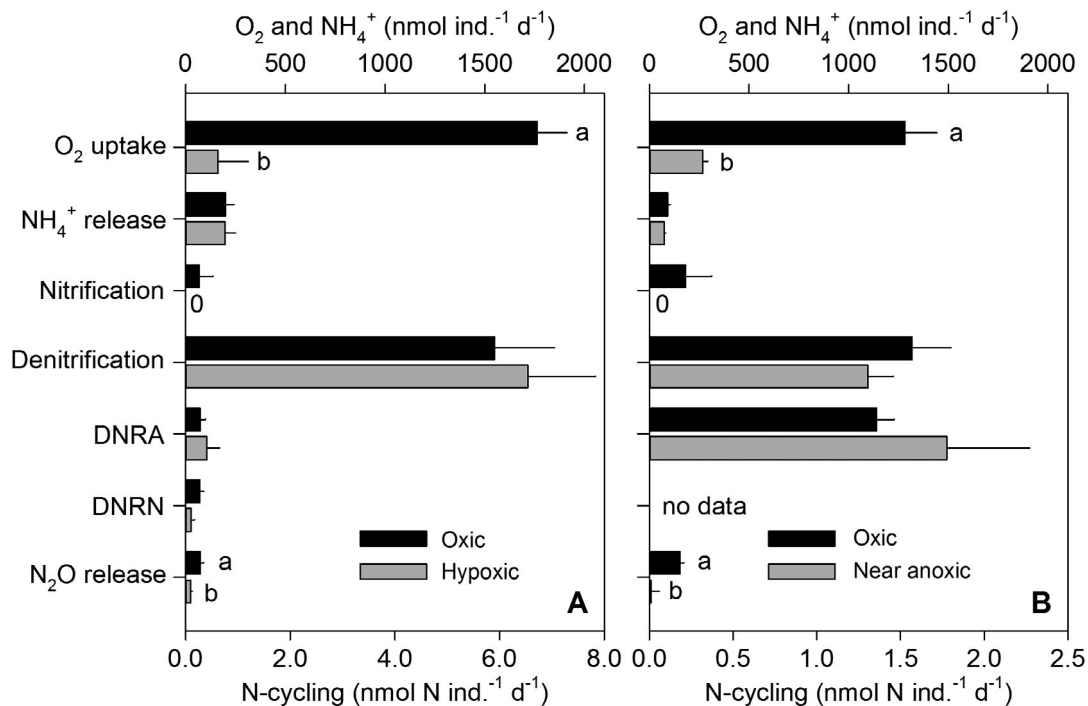


Fig. 4. Effect of ambient oxygen level. Algae-fed copepods *Calanus hyperboreus* were incubated in (A) $^{15}NO_3^-$ -enriched or (B) $^{15}NO_2^-$ -enriched seawater under oxic vs. hypoxic or near-anoxic conditions. Incubation conditions as specified in 'Materials and methods' and Table 1. Means \pm SE of $n = 5$ –12 (oxic), $n = 6$ (hypoxic), and $n = 5$ (near anoxic) incubations are shown. Note different scales on the 2 x-axes. Lower-case letters indicate statistically significant differences between oxic, hypoxic, and near-anoxic conditions. DNRA/DNRN: dissimilatory nitrate reduction to ammonium/nitrite, respectively

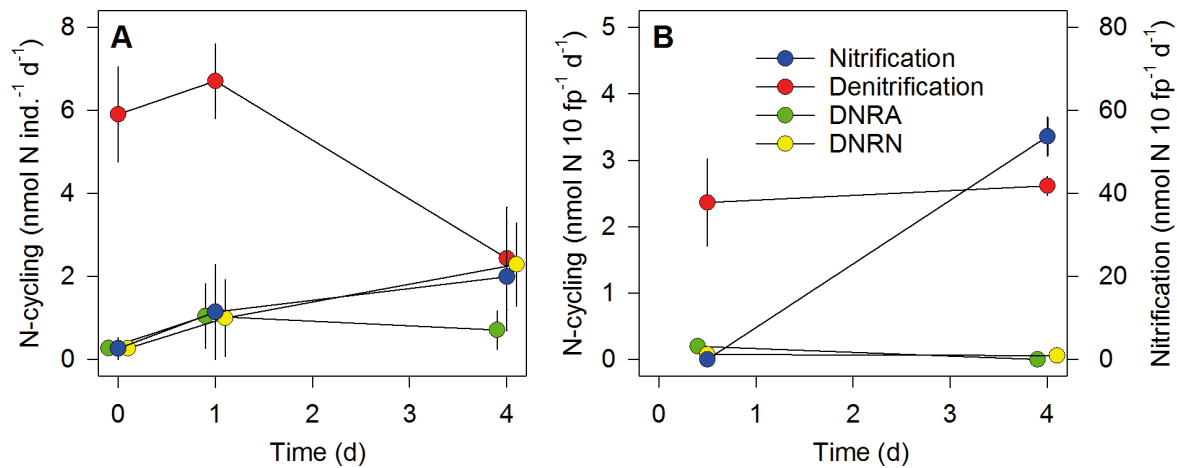


Fig. 5. Effect of time. The rates of different N-cycle pathways associated with (A) *Calanus hyperboreus* and (B) its fecal pellets were measured for up to 4 d after death and defecation, respectively, under incubation conditions as specified in 'Materials and methods' and Table 1. Means \pm SE of $n = 12$ (live copepods), $n = 6$ (dead copepods), and $n = 3-5$ (fecal pellets) incubations are shown. Note that in (B), rates are expressed per 10 fecal pellets (fp) and nitrification activity is shown on a dedicated right-hand y-axis. DNRA/DNRN: dissimilatory nitrate reduction to ammonium/nitrite, respectively

Contribution by calanoid copepods to fixed-nitrogen loss in Arctic waters

The pelagic N_2 production by live copepods (through denitrification) and the benthic N_2 production in sediments (through denitrification and anamox) were compared on an areal basis (Table S5 in the Supplement). During the Arctic spring bloom, the depth-integrated *in situ* abundance of adult *C. hyperboreus* at a 250 m deep station in Disko Bay reaches maximum levels of ~ 2250 ind. m^{-2} (Swalethorp et al. 2011). For our extrapolation, 75% of the *C. hyperboreus* specimens were assumed to be alive and 25% dead (Yamaguchi et al. 2002). Denitrification associated with the live specimens amounts to $10 \mu\text{mol N m}^{-2} \text{d}^{-1}$ (Table S5), which is equivalent to 3.5% of the benthic N_2 production (Rysgaard et al. 2004). However, the *C. hyperboreus* contribution increases to as much as 11.6%, if denitrification activities of dead specimens and fecal pellets are included in the estimate (Table S5).

The other 2 calanoid copepod species in the Arctic, *C. glacialis* and *C. finmarchicus*, are even more abundant than *C. hyperboreus* (Hirche 1991, Madsen et al. 2001, Swalethorp et al. 2011), but they have a much smaller individual biomass (Hirche 1991, Madsen et al. 2001). Since denitrification activity has recently been found in carcasses of the small *C. finmarchicus* (Glud et al. 2015), it can be assumed that the medium-sized *C. glacialis* also exhibits denitrification activity. It can be further assumed that the gut volume decreases proportionately to biomass and

therefore, the denitrification rates measured for *C. hyperboreus* were downscaled linearly to the much smaller individual biomass of *C. glacialis* and *C. finmarchicus* (Table S5). The total integrated denitrification activity associated with the 3 calanoid copepod species and their fecal pellets and carcasses amounts then to $82 \mu\text{mol N m}^{-2} \text{d}^{-1}$ (Table S5), which is equivalent to as much as 28.3% of the benthic N_2 production (Rysgaard et al. 2004).

DISCUSSION

Feeding-related controls on copepod-associated microbial nitrogen cycling

Live *Calanus hyperboreus* specimens hosted both aerobic and anaerobic N-cycle activities among which denitrification was quantitatively the most important and consistent. The other 2 dissimilatory nitrate reduction pathways, DNRA and DNRN, showed lower rates and varied more strongly between measurement campaigns. Nitrification activity was generally low in live copepods, but gained in importance in aging fecal pellets and carcasses. Several lines of evidence indicate that this pattern of copepod-associated microbial N-cycling is directly and indirectly controlled by the animal's feeding activity and thus the gut filling degree: (1) anaerobic N-cycling was drastically reduced in animals with empty guts, (2) ambient O_2 levels did not influence anaerobic N-cycling in algae-fed specimens (i.e.

their filled guts were anoxic anyway), and (3) freshly released fecal pellets exhibited similar N-cycling as algae-fed copepods.

A copepod gut filled with food is the predestined site for anaerobic N-cycling because it is at least partially anoxic and also rich in labile organic carbon from digestive processes (Tang 2005, Tang et al. 2011). The gut microbial community itself is involved in creating these microenvironmental conditions. First, the aerobic activity of heterotrophic microorganisms in the food promotes gut anoxia (Tang et al. 2011, Heisterkamp et al. 2016), which triggers a shift to anaerobic metabolism in many microorganisms ingested by aquatic invertebrates (Stief et al. 2009, Heisterkamp et al. 2010). Second, the microbial community of the copepod gut contributes to the solubilization of organic food particles through exoenzyme activities (Lawrence et al. 1993). In the light of these microbe–microenvironment interactions, it is not surprising that the rates of anaerobic N-cycling in live copepods were significantly reduced by starving the animals, which probably rendered the gut richer in oxygen and poorer in organic carbon and microbial biomass. Similar results were found for the aquatic larvae of the midge *Chironomus plumosus* in which denitrification and N_2O production were exclusively mediated by microorganisms in the gut contents, but not in the gut wall (Stief et al. 2009).

Net N_2O release was not significantly different in algae-fed, algae-starved, or bacteria-starved specimens of *C. hyperboreus*. However, a very strong and uniform control on both net N_2O release and nitrification activity was exerted by the ambient O_2 level, suggesting that N_2O was mainly formed through aerobic nitrification. The positive correlation between net N_2O and NH_4^+ release further suggests that nitrification was stimulated by NH_4^+ accumulating in the incubation bottles. The individual-specific rates of net N_2O release by algae-fed specimens of *C. hyperboreus* translate into weight-specific rates of 14–18 pmol N mg wet wt⁻¹ d⁻¹ that fall within the range of net N_2O release due to gut denitrification reported for benthic invertebrates (without a shell or exoskeleton) from freshwater and marine habitats of 13–44 and 0.5–23 pmol N mg wet wt⁻¹ d⁻¹, respectively (Stief et al. 2009, Heisterkamp et al. 2010). In addition, benthic aquatic invertebrates with a shell or exoskeleton host microbial N_2O production by nitrification in external biofilms (Heisterkamp et al. 2010, 2013, Svenningsen et al. 2012), which also seems likely for live and dead specimens of *C. hyperboreus*.

Freshly released fecal pellets showed a similar pattern of anaerobic N-cycling as algae-fed copepods,

with high denitrification and low DNRA and DNRN rates, despite the incubation under oxic conditions. The denitrification activity of 10 fecal pellets was still significantly lower than that of 1 copepod with a filled gut, possibly because (1) only the very center of the released fecal pellets remained anoxic or (2) denitrification in the fecal pellet proceeded under micro-aerobic conditions, but at a significantly lower rate. Oxygen microsensor measurements have previously been made in large crustacean fecal pellets ($3.4 \times 10^9 \mu\text{m}^3$ in volume; Alldredge & Cohen 1987) and in the boundary layer surrounding the much smaller fecal pellets of *Temora longicornis* ($2.2\text{--}10.0 \times 10^5 \mu\text{m}^3$ in volume; Ploug et al. 2008). Internal anoxia was evident in the large fecal pellets, but could not be inferred for the small fecal pellets. The fecal pellets of *C. hyperboreus* ($3.1\text{--}10.7 \times 10^6 \mu\text{m}^3$ in volume; Urban-Rich 1999, Juul-Pedersen et al. 2006) take an intermediate position, which makes it difficult to estimate whether they were oxic or anoxic in the center. If the freshly released fecal pellets were indeed not anoxic in the center, then denitrification activity may have also persisted under hypoxic conditions (Lloyd 1993). Such ‘aerobic denitrification’ has been a long-debated N-cycle pathway, but meanwhile studies are accumulating that demonstrate this process convincingly (e.g. Schreiber et al. 2010, Marchant et al. 2017). Aerobic denitrification is facilitated by the constitutive expression of denitrification genes independent of the ambient O_2 level and is thus advantageous in environments with fluctuating O_2 levels, as may be the case for the guts of aquatic invertebrates (Tang et al. 2011, Heisterkamp et al. 2016).

In carcasses, the feeding-related controls on copepod-associated N-cycling must fade away because food uptake and processing stop and the copepod body disintegrates. This does not mean, however, that N-cycling declines or takes on a different pattern immediately. In fact, within the first 4 d after death, N-cycling remained remarkably stable and did not experience significant transformations (see below). Only recently, anaerobic N-cycling has been observed in copepod carcasses of different species and geographic origin (Glud et al. 2015, Stief et al. 2017, 2018). The intensity of carcass-associated N-cycling is mainly controlled by the ambient O_2 level through variation in the spatial extent of internal anoxia, which is conceptually different from the anoxic gut scenario described above and resembles more the situation in phytodetritus aggregates that sink through different O_2 regimes (Klawonn et al. 2015, Stief et al. 2016).

Contribution by calanoid copepods to pelagic fixed-nitrogen loss

The relatively high denitrification activity associated with *C. hyperboreus* corresponds to fixed-nitrogen loss through N_2 production. Arctic surface waters are generally well-oxygenated, which will definitely inhibit pelagic fixed-nitrogen loss mediated by free-living microorganisms (Kalvelage et al. 2011, Dalsgaard et al. 2014). Thus, copepods (similar to sinking phytodetritus aggregates) are unique microsites of fixed-nitrogen loss that is otherwise not represented in the water column of Arctic marine ecosystems. However, fixed-nitrogen loss does occur in Arctic marine sediments that are anoxic a few millimetres or centimetres below the surface (Rysgaard et al. 2004). We estimate that during spring–early summer, the total denitrification activity associated with all calanoid copepods and their fecal pellets and carcasses is equivalent to ~28% of the benthic N_2 production through denitrification and anammox. Thus, it appears that the copepod-associated fixed-nitrogen loss reaches a dimension that should be taken into account for budgets and models of marine N-cycling in Arctic marine ecosystems.

Our estimate of pelagic denitrification associated with all calanoid copepods can be regarded as conservative because (1) single abundance peaks of adult copepods were ignored in favor of more commonly observed abundance values (Swalethorp et al. 2011), (2) juvenile stages were not considered because their small body size may (or may not) prohibit internal anoxia and thus denitrification activity, (3) *in situ* abundances of fecal pellets in Disko Bay by far exceed the abundances calculated from individual-specific fecal pellet production rates (Juul-Pedersen et al. 2006, Swalethorp et al. 2011), and (4) calanoid fecal pellets have similar dimensions, despite the large differences in body size (Juul-Pedersen et al. 2006). However, our estimate is only valid for spring–early summer (i.e. for approximately one-quarter of the year) when the abundance of calanoid copepods in the water column is high. Taking this short abundance maximum into account, the annual rate of pelagic denitrification associated with all calanoid copepods will amount to $7.5 \mu\text{mol m}^{-2} \text{yr}^{-1}$, which is equivalent to 7.1% of the annual rate of benthic N_2 production (Rysgaard et al. 2004) and thus considerably lower than the 28.3% estimated for spring–early summer only. Such estimates of the copepod contribution to fixed-nitrogen loss clearly need to be improved by the annual monitoring of copepod abundances and N-cycling rates (ideally in simultaneous

studies at the same location), which is a major undertaking in the Arctic.

Contribution by settled fecal pellets and carcasses to benthic nitrogen cycling

Once released into the water, fecal pellets and the associated microorganisms are exposed to a macro-environment that is rich in O_2 and poor in labile organic carbon and thus almost the inverse of the gut microenvironment. Anaerobic N-cycling remained remarkably stable under these new conditions, but nitrification activity increased strongly with time. From the O_2 consumption rate of the *C. hyperboreus* fecal pellets and the typical C:N ratio of copepod fecal pellets of 7–9 (Morales 1987), it is estimated that nitrification activity can be sustained by NH_4^+ produced through organic matter degradation inside the fecal pellets (Table S6 in the Supplement). Importantly, the strong increase in nitrification activity turned the fecal pellets from net NO_x^- sinks into net NO_x^- sources. At an estimated sinking velocity of the *C. hyperboreus* fecal pellets of $\sim 50 \text{ m d}^{-1}$ (Urban-Rich 1999, Wassmann et al. 1999), the fecal pellets would sink $\sim 200 \text{ m}$ during the experimental aging period of 4 d. Fecal pellets may thus reach the seafloor in many coastal marine areas and still contribute to both benthic nitrification and denitrification.

The *C. hyperboreus* carcasses remained microsites of denitrification and net NO_x^- consumption throughout the entire aging period of 4 d. Denitrification rates showed first signs of a decrease (but not statistically significant) in 4-d-old carcasses. The decline of this anaerobic, heterotrophic activity is likely explained by the loss in organic matter content and the potential reduction of internal anoxia. Sinking velocities of small copepod carcasses are $\sim 50\text{--}100 \text{ m d}^{-1}$ (Elliott et al. 2010, Kirillin et al. 2012, Dubovskaya et al. 2015) and thus carcasses settling on the seafloor at shallow sites may persist as microsites of denitrification as is known from diatom aggregates (Marzocchi et al. 2018).

CONCLUSION

Our results suggest that N-budgets in Arctic marine ecosystems need to include the newly discovered pelagic fixed-nitrogen loss that is associated with live calanoid copepods and their remains in addition to the already known benthic fixed-nitrogen loss that takes place in anoxic sediment layers. Such pelagic

fixed-nitrogen loss enabled by anoxic microsites has also been discovered for phytodetritus aggregates and copepod carcasses (Glud et al. 2015, Klawonn et al. 2015, Stief et al. 2016), in polar, temperate, and tropical latitudes (Glud et al. 2015, Klawonn et al. 2015, Stief et al. 2016, 2017, 2018, this study), in low-oxygen environments (Stief et al. 2017), and in freshwater ecosystems (Stief et al. 2018). Thus pelagic fixed-nitrogen loss enabled by anoxic microsites is emerging as a widespread phenomenon in various aquatic ecosystems and is consequently also adopted in current models of oceanic N-dynamics (Bianchi et al. 2018).

It can be anticipated that other globally abundant groups of zooplankton (e.g. krill, amphipods, salps, and jellyfish) will also be identified as pelagic hotspots of fixed-nitrogen loss. Anaerobic N-cycling in these hotspots is reinforced by low ambient O₂ levels that continue to spread in the open ocean, but also in eutrophic coastal regions (Diaz & Rosenberg 2008, Schmidtke et al. 2017). Interestingly, evidence is increasing that copepods and other zooplankton are abundant in and around oxygen minimum zones (e.g. Teuber et al. 2013, Hirche et al. 2014), which may allow higher rates of zooplankton-associated fixed-nitrogen loss due to the higher temperatures and lower ambient O₂ levels compared with Arctic marine ecosystems. Similarly, substantial zooplankton-associated fixed-nitrogen loss may prevail in coastal areas of temperate zones with longer-lasting periods of high productivity than in the Arctic.

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