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Impacts of Intraguild Predation on Arctic Copepod Communities

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Communities of large copepods form an essential hub of matter and energy fluxes in Arctic marine food webs. Intraguild predation on eggs and early larval stages occurs among the different species of those communities and it has been hypothesized to impact its structure and function. In order to better understand the interactions between dominant copepod species in the Arctic, we conducted laboratory experiments that quantified intraguild predation between the conspicuous and omnivorous *Metridia longa* and the dominant *Calanus hyperboreus*. We recorded individual egg ingestion rates for several conditions of temperature, egg concentration, and alternative food presence. In each of these experiments, at least some females ingested eggs but individual ingestion rates were highly variable. The global mean ingestion rate of *M. longa* on *C. hyperboreus* eggs was 5.8 eggs ind⁻¹ d⁻¹, or an estimated 37% of *M. longa* daily metabolic need. Among the different factors tested and the various individual traits considered (prosome length, condition index), only the egg concentration had a significant and positive effect on ingestion rates. We further explored the potential ecological impacts of intraguild predation in a simple 1D numerical model of *C. hyperboreus* eggs vertical distribution in the Amundsen Gulf. Our modeling results showed an asymmetric relationship in that *M. longa* has little potential impact on the recruitment of *C. hyperboreus* (<3% egg standing stock removed by IGP at most) whereas the eggs intercepted by the former can account for a significant portion of its metabolic requirement during winter (up to a third).

Keywords: intraguild predation, copepods, *Metridia longa*, *Calanus hyperboreus*, Arctic regions, numerical modeling

INTRODUCTION

Arctic and subarctic marine food webs are characterized by the presence of large calanoid copepods that channel primary production toward secondary consumers. Copepods have developed life cycle strategies that allow them to thrive in these highly seasonal environments. In the Arctic Ocean, several species of *Calanus* dominate the mesozooplankton biomass (Head et al., 2003; Hopcroft et al., 2010). During the short productive period in spring and summer, copepods feed on ice algae (when available in ice-covered regions) and phytoplankton and concentrate this energy into lipids, mostly stored as wax ester in hypertrophied oil sacs (Lee et al., 2006). In winter, they survive thanks to those lipid reserves that fuel their reduced metabolism during an extended period of dormancy

(the diapause). All year long, copepods are an essential food source for many predators such as bowhead whale, little auk, and above all Arctic cod, a cornerstone component of the Arctic food web (Fort et al., 2010; Falardeau et al., 2013; Pomerleau et al., 2014). Thus, communities of large copepods form a critical hub of matter and energy fluxes in Arctic and subarctic marine food webs.

In the Arctic, copepod community biomass is mainly composed of the large species *Calanus hyperboreus* and *Calanus glacialis* (adult female median prosome length of 6.7 and 4.1 mm, respectively), the medium-sized *Metridia longa* (2.8 mm) and the small *Pseudocalanus* spp. (1.1 mm) (Darnis et al., 2012). Following the current dynamics, the boreal species *C. finmarchicus* is also regularly found in marginal Arctic seas, especially in the eastern Greenland Sea and Barents Sea (Conover and Huntley, 1991). Moreover, *C. finmarchicus*' biogeographic distribution in the surface layer is projected to move even farther northward in response to surface circulation and temperature forcing induced by climate change (Reygondeau and Beaugrand, 2011). The structure and functions of copepod communities are critical from an energetic point of view for marine ecosystems and depend on the actual assemblage of species. From one species to the other, the oil sac size is different and therefore the energetic content differs. *C. hyperboreus* and *C. glacialis* are bigger and contain more lipids than the boreal *C. finmarchicus*. Consequently, many small visual predators such as fish larvae and juveniles, would reap a larger energetic reward for a similar harvesting effort, making these Arctic copepod species the preys of choice. Hence, changes in the assemblage of copepods communities could impact marine predators' recruitment, resulting in a form of bottom-up control (Mills et al., 2013; Greene and Pershing, 2014). For example, along the West coast of the Spitsbergen island in the Svalbard archipelago, the increase of warm Atlantic water masses that bring along abundant *C. finmarchicus* may have a negative impact on the reproductive success of little auks by reducing the relative abundance of its preferred prey, *C. glacialis* (Kwasniewski et al., 2010). In order to understand and predict impacts of environmental changes on Arctic marine ecosystems, it is necessary to better understand the mechanisms responsible for the specific assemblages of copepod communities.

In addition to physical forcing, relationships occurring within copepod communities can influence their structures and functions. Species that follow each other and co-occur in a community not only share food and space resources, but also develop complex interactions between them. Intraguild predation (IGP) has been proposed as an ecological strategy that could structure copepod communities (Irigoiien and Harris, 2006; Plourde et al., 2009; Darnis, 2013; Melle et al., 2014). This particular type of predation occurs between members of a group of species that share the same food resources (Polis et al., 1989). This phenomenon is widespread across a variety of marine and terrestrial ecosystems and through all taxonomic and trophic levels (Polis et al., 1989; Holt and Polis, 1997; Arim and Marquet, 2004). This complex interaction is particularly interesting since it results in an immediate energy gain for the

predator as well as a long term reduction of its competition (Hiltunen et al., 2013).

In copepod populations, survival rate to adulthood is strongly influenced by egg and nauplii mortality (Davis, 1984; Plourde et al., 2009). Egg mortality is particularly high in broadcast spawning species that release their eggs in the water column (Ohman et al., 2004), such as *Calanus* spp and *M. longa*. Although predominantly considered to be herbivorous, most calanoid copepod species have a flexible diet and can be omnivorous or even cannibalistic (Landry, 1981; Ohman and Hirche, 2001; Bonnet, 2004; Basedow and Tande, 2006). These species usually generate a filtration current to obtain small prey, essentially phytoplankton and micro-zooplankton, or cruise through the water and attack when a prey is detected (Kiørboe, 2013). Thus, they can consume eggs and young nauplii stages with limited mobility.

Cannibalism on eggs and nauplii can control the phenology of *C. finmarchicus* recruitment (Ohman and Hirche, 2001). It is therefore likely that IGP can impact recruitment as well as influencing the temporal succession of dominant species (Irigoiien and Harris, 2006). In environments as contrasted as the Beaufort Sea and the St-Lawrence estuary, reduced *C. hyperboreus* recruitment and abundance co-occurred with an increase in *M. longa* abundance (Plourde et al., 2002; Darnis, 2013). During the time of peak egg production by *C. hyperboreus*, the gut of individual *M. longa* is often observed to be orange, a particularity linked to the probable ingestion of lipid-rich *C. hyperboreus* eggs (Conover and Huntley, 1991). Hence it has been hypothesized that *M. longa* individuals that remain active at intermediate depths throughout winter (no diapause) could intercept and ingest buoyant *C. hyperboreus* eggs that are spawned in deep waters (Plourde et al., 2003; Darnis, 2013). Given that *M. longa* population dynamics resources are scarce in winter, lipid-rich eggs could represent an important energy source for such an omnivorous (opportunistic) species. Later in early spring, numerous *C. hyperboreus* and *C. glacialis* that emerge from diapause and ascend toward the surface ahead of the phytoplankton bloom could also be feeding on eggs and young nauplii stages. Consequently, IGP could impact the recruitment of the true Arctic *C. hyperboreus*.

Surprisingly, only few studies have been conducted on predation within calanoid copepod communities (Landry, 1981; Huntley and Escritor, 1992; Bonnet, 2004; Basedow and Tande, 2006; Vestheim et al., 2013). The majority of the studies targeted cannibalism and while none was focused on IGP between the dominant Arctic species *M. longa* and *C. hyperboreus*, Huntley and Escritor (1992) reported ingestion rates of the vicariant species *M. gerlachei* on eggs of the dominant Antarctic *Calanoides acutus* in the Austral Ocean. In all experiments, ingestion rates on eggs (and nauplii) vary according to their concentration. The influence of alternative food source on ingestion produced ambiguous experimental results. On one hand, experiments with female *C. pacificus* suggest a switch between herbivorous and carnivorous behavior that depends on the relative abundances of phytoplankton and their own nauplii (Landry, 1981). On the other hand, experiments with *C. finmarchicus* have shown that ingestion rates on its own

nauplii are independent of the ambient algae concentration (Basedow and Tande, 2006).

Copepod community models have been developed with a focus on development, growth and competition for food resources (Record et al., 2012, 2013). However, community level processes involving interspecific interactions such as IGP have not yet been implemented into these models. In order to provide a better understanding of IGP and to provide a better parameterization for models of copepod communities, we conducted laboratory experiments that quantified the ingestion of copepod eggs by some of the dominant Arctic copepod species, *M. longa* and *C. hyperboreus*. We further conducted a simple numerical experiment in order to assess the potential ecological implications of our findings in Arctic marine ecosystems. Copepods were sampled in the Lower St-Lawrence estuary (LSLE) and feeding experiments on eggs were conducted under different conditions of temperature, egg concentration, and alternative food availability.

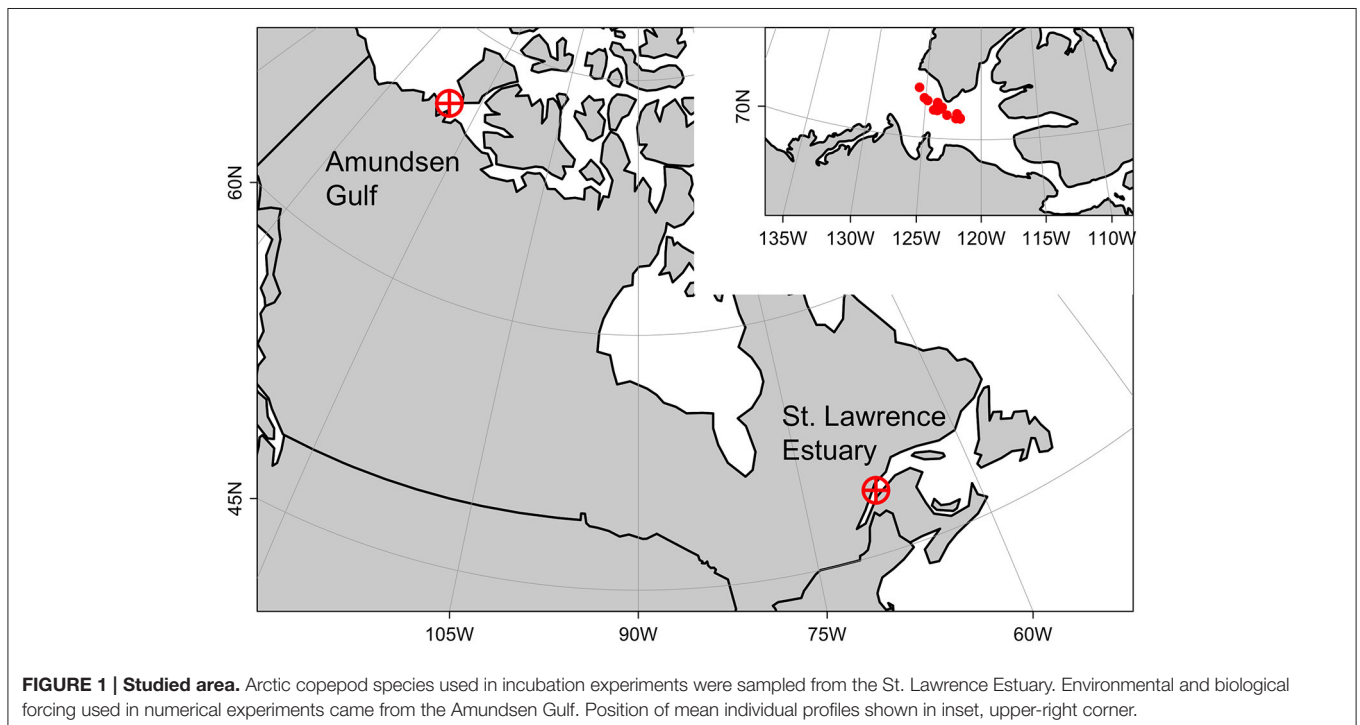
METHODS

Area of Study and Sampling

The LSL is the southernmost sea directly influenced by Arctic water masses and that presents Arctic features in the North Atlantic (**Figure 1**). Arctic water masses enter in the Gulf of St-Lawrence (GSL) by the Strait of Belle Isle in the North, transporting Arctic copepods species such as *C. hyperboreus*, *C. glacialis*, and *M. longa*. Warmer North Atlantic water masses enter the system by Cabot Strait in the South and carry along boreal species such as *C. finmarchicus*. Both Arctic and temperate species thrive in the GSL system where cold and warm water

masses are segregated vertically between the thin seasonal surface layer, the cold intermediate layer renewed locally each winter (core temperatures can be negative) and the deep Atlantic layer (between 4 and 5°C). Colder water masses dominate the eastern and northern parts of the Gulf and the deep lower Estuary where upstream tidal pumping brings cold and nutrient-rich waters to the surface, whereas warm surface waters (up to 20°C) develop throughout the summer over the shallow southern half of the Gulf (Saucier et al., 2003; Le Fouest et al., 2005). These particular water masses and species mix make the LSL an exceptional experimental model to study IGP in the context of the rapid environmental changes that the Arctic is currently facing. Zooplankton was sampled in the LSL (48°40'N, 68°35'W) in October 2014 with a vertical plankton net (158 µm mesh size, 1 m diameter) at 125 or 200 m from the surface at a towing speed of 0.5 m s⁻¹. The catch was diluted into 4 L jars filled with filtered seawater and maintained close to ambient temperature at 5 to 6°C in coolers during transport to the laboratory (less than an hour).

In a subsequent numerical experiment, we applied our observations to a truly Arctic environment, the Amundsen Gulf. The Amundsen Gulf is located in the Canadian territory of Nunavut at about 71°N and bordered by Banks Island and Victoria Island. It connects south-eastern Beaufort Sea to the Canadian archipelago (**Figure 1**). The Amundsen Gulf water masses are generally cold (around 0°C) and are mainly discriminated by their different salinities: the Polar-Mixed Layer from the surface to c.a. 50 m ($S \sim 31.6$), the Pacific Halocline below until 200 m ($32.4 < S < 33.1$) and the slightly warmer Atlantic Layer below 200 m ($S \sim 34$) (Carmack and Macdonald, 2002). The Gulf covers about 60,000 km² and the maximum



depth of this large channel is 630 m. In winter, it is entirely ice-covered except for sporadic polynyas and flaw leads and the spring ice-breakup is highly variable (Galley et al., 2008).

Live Animal Sorting

Owing to the harsh conditions at sea during winter over the LSLE, copepods were sampled at the end of October 2014 prior to the beginning of *C. hyperboreus* reproduction and the formation of dense sea ice. Plourde et al. (2003) have previously shown, for a similar experimental setup, that capture and handling of *C. hyperboreus* females triggered gonad maturation, leading to egg production a few weeks after collection and approximately a month earlier than expected according to the *in situ* timing. A visual inspection of appendages and behavior under a binocular microscope allowed us to select adult female *M. longa* and *C. hyperboreus* in good condition from the live samples, within 48 h of the catch. Animals were kept in the dark between 3 and 5°C in groups of 25 to 50 in 1 L beakers equipped with egg separators (mesh size = 333 μm) and filled with filtered seawater. Female *M. longa* were fed with solutions of concentrated diatoms *Thalassiosira weissflogii* (Instant Algae® TW 1200), whereas female *C. hyperboreus* were not fed since egg production is entirely fuelled by internal lipids during the dormant part of their life cycle. The beakers were inspected daily for egg production for about a month until female *C. hyperboreus* spawned enough eggs to start the experiments.

Predation Experiments

Within 48 h of the start of the experiment, individuals used as predators were photographed laterally using a PixeLINK camera of 5 Mb (PL-E425CU) mounted on a LEICA MZ6 stereoscope. In order to minimize stress, individuals were kept in cooled seawater until the very moment that the picture was taken and gently manipulated with handling needles. Prosome length (distance between the tip of the cephalosome and the tip of the last thoracic segment), prosome area, oil sac length, and area were measured with the software ImageJ v. 1.49. The condition index was estimated as the oil sac area divided by the prosome area of the individual in order to give an indication of its lipid content. Predators' carbon content was estimated from species and stage-specific seasonal relationships between individual carbon mass and prosome length (Forest et al., 2010). The photographed animals were then placed individually in 45 mL Petri dishes filled with filtered seawater and equipped with egg separator (mesh size = 333 μm) for acclimation at the experimental temperatures.

Experiments were carried in an atmosphere-controlled chamber in November 2014 with female *M. longa* fed with *C. hyperboreus* eggs. Predators were placed individually in a bottle (1.35 or 2.37 L) filled with filtered seawater containing a precise number of eggs spawned within the previous 48 h. Bottles were turned upside down once per minute on a rotating wheel in order to maintain the eggs in suspension. Manipulations were conducted under a dimmed red light and the experiments were carried in the dark. Incubation time was kept relatively short (less than 24 h) in order to avoid a complete consumption of the eggs and hence a failure to accurately estimate ingestion rates. It also varied according to incubation conditions, with longer

incubation times for lower egg concentration (lower encounter probability between eggs and the predator). After 4 to 22 h, the contents of the bottles were filtered with a 73 μm sieve and the remaining eggs were counted. A minimum of two controls without predator for each treatment was set up in order to check the accuracy of the egg recovery method. The status of each individual was verified at the end of the experiment. Ingestion rates were discarded for the few dead individuals, and the sluggish or unhealthy-looking ones as well.

We tested the effect of temperature on *M. longa* ingestion of *C. hyperboreus* eggs. Incubations were carried at 1, 4, and 8°C to reflect the potential range of *in situ* water temperature encountered by this species between the Arctic and subarctic regions. Unfortunately, we could not test for negative water temperatures. In order to characterize functional responses, we tested the effect of *C. hyperboreus* egg concentration on *M. longa* ingestion rate. We were not aware of actual data about *in situ* *C. hyperboreus* egg concentrations in the water column, but we estimated it to be low (Huntley and Escritor, 1992). Hence we chose concentrations of 5, 10, 20, and 30 egg L⁻¹. Finally, we checked whether there was an influence of alternative food availability, by adding an additional food source in the form of concentrated diatoms *T. weissflogii* (Instant Algae® TW 1200) in half of our replicates. Algae were offered at about 50 μg C L⁻¹ according to cell concentration determined with a Hausser Bright-Line Hemacytometer (counting chamber) and carbon to volume relationships for diatoms (Menden-Deuer and Lessard, 2000).

Estimation of Ingestion Rates

The instantaneous feeding rate on eggs g (h⁻¹) was derived from an exponential equation (Båmstedt et al., 2000):

$$g = \frac{\ln\left(\frac{E_f}{E_0}\right)}{t} \quad (1)$$

where E_0 and E_f are respectively the egg concentration at beginning and the end of experiment (egg L⁻¹) and t is incubation time (h). The number of *C. hyperboreus* eggs obviously did not increase during the experiments, and experiment duration was not long enough for hatching to occur (no nauplii was ever found in any of the control or experiment bottles).

We deduced the clearance rate F (L ind⁻¹ h⁻¹), which corresponds to the volume of water processed assuming 100% capture efficiency and a homogeneous food concentration, from both g and the volume of the incubation bottle V (L):

$$F = g \times V \quad (2)$$

Finally, we obtained the ingestion rate I (egg ind⁻¹ h⁻¹) with:

$$I = F \times [E] \quad (3)$$

where $[E]$ is the average egg concentration as given by:

$$[E] = \left(E_0 \times \frac{1 - e^{(-g \times t)}}{g \times t}\right) \quad (4)$$

Daily ingestion rates were assumed to be 24 times the hourly rates since *M. longa* is known to be active and swimming almost continuously (Hirche, 1987). Egg ingestion rates were then converted in carbon units using an egg carbon content of 0.84 ($\mu\text{g C egg}^{-1}$; Plourde et al., 2003). The proportion of daily metabolic needs (%) satisfied by egg ingestion was estimated from the ratio of carbon ingestion rate $I_C = 0.84 * I$ ($\mu\text{g C ind}^{-1} \text{d}^{-1}$) and a mean and constant respiration rate ($\mu\text{g C ind}^{-1} \text{d}^{-1}$) measured by Seuthe et al. (2006).

Data Analysis

The number of ingested eggs followed a Poisson distribution (an asymmetric right-skewed distribution of discrete values). Hence, in order to minimize estimating errors, a generalized linear model (GLM) for Poisson distribution was used to predict the number of eggs ingested (EI) (the raw data) according to temperature (T), egg concentration (E), presence of additional food source (AC), prosome length (PL), and condition index (CI) of the predator. Bottle volume (V) and duration of experiment (D) were taken into account by using them in an offset term. Several models were tested (with and without interactions) and we computed Akaike's information criterion corrected for overdispersion (QAIC) as a decision-support metric. The GLMs formulae were of the form (here is the one with all the independent variables but no interactions):

$$\log\left(\frac{EI}{D \times V}\right) = \beta_0 + (\beta_T \times T) + (\beta_E \times E) + (\beta_{AC} \times AC) + (\beta_{PL} \times PL) + (\beta_{CI} \times CI) \quad (5)$$

where β_i are the coefficient estimates for each variable. The number of eggs ingested per unit of time and volume can easily be obtained from this model results.

Model of Egg Vertical Distribution

In order to assess the implications of our findings in the context of Arctic marine ecosystems, we developed a simple water column (1D) model of the vertical distribution of *C. hyperboreus* eggs in the Amundsen Gulf. The model computed the time evolution of egg concentration at a given depth according to advection and diffusion, gains by egg production and losses by development and predation (Figure 4). The rate of change of egg concentration followed the classical advection-diffusion-reaction formulation (Soetaert and Herman, 2009):

$$\frac{\partial E}{\partial t} = -w \frac{\partial E}{\partial z} + K \frac{\partial^2 E}{\partial z^2} + P_z - I * M + 1/H * E \quad (6)$$

where the first right-hand side term represents the effect of egg vertical velocity, the second the effect of diffusion and the others several biological reaction terms. More specifically, E was the egg concentration (egg m^{-3}), t the time (h), w the egg velocity (m h^{-1}) defined positive downward, z the depth (m), K the vertical eddy diffusivity coefficient ($\text{m}^2 \text{h}^{-1}$), P_z the depth-dependent egg production rate ($\text{egg m}^{-3} \text{h}^{-1}$), H the egg hatching time (h), M the females *M. longa* concentration (ind m^{-3}) and I the ingestion rate by other copepods (egg ind^{-1}

h^{-1}). The egg velocity w was given by Stokes' law (Visser and Jónasdóttir, 1999):

$$w = 3600 \times \frac{g \times d^2 \times (\rho_{\text{egg}} - \rho_{\text{water}})}{18 \times \mu} \quad (7)$$

where g is the gravitational constant (9.81 m s^{-2}), d is the egg diameter (m), ρ_{egg} is the egg density (g m^{-3}), ρ_{water} is the water density (g m^{-3}) and μ is the dynamic viscosity of the seawater ($1.85 \text{ g m}^{-1} \text{ s}^{-1}$), here taken as a constant (Table 1).

The source term for eggs (P_z) came from the average daily production of *C. hyperboreus* population ($30,000 \text{ eggs m}^{-2}$) observed between February and April 2008 in the Amundsen Gulf (Darnis, 2013). *C. hyperboreus* females released more than 90% of their eggs during this 3 month-period, while remaining at depths between 200 and 300 m (Darnis, 2013). As a result, we computed the vertical profile of egg production rate P_z according to a normal distribution whose mean was centered at 250 m, its standard deviation 15 m and its integral equal to $30,000 \text{ eggs m}^{-2}$ (99.9% of the eggs were released between 200 and 300 m).

The IGP rate I (h^{-1}) exerted on *C. hyperboreus* eggs was simply the product of *M. longa* females' abundance (ind m^{-3}) and the individual filtration rate ($\text{m}^3 \text{ ind}^{-1} \text{ h}^{-1}$) found in our grazing experiments.

Egg hatching time H (h) followed an empirical Bělehrádek's function obtained from observed hatching times of *C. hyperboreus* eggs (Jung-Madsen et al., 2013):

$$H = a \times (T - \alpha)^{-b} \quad (8)$$

where T was water temperature ($^{\circ}\text{C}$), a ($\text{d } ^{\circ}\text{C}^{-1}$), α ($^{\circ}\text{C}$) and b constants.

A simple ordinary differential equation framework was not optimal for the modeling of developing eggs because of the "numerical diffusivity" caused by the hatching rate (Gentleman et al., 2008). Simply put, with a development (hatching) rate, the progression through development stages is treated as a continuous process within the population, instead of a discrete event highly synchronized among many individuals. This can

TABLE 1 | Model parameters and references.

Parameter	Unit	Value	References
H , egg hatching time			JM2013
a	$\text{d } ^{\circ}\text{C}^{-1}$	1196	JM2013
α	$^{\circ}\text{C}$	12.7	JM2013
b	-	-2.05	JM2013
d , egg diameter	m	1.92×10^{-4}	JM2013
P_z , egg production	$\text{egg m}^{-2} \text{ d}^{-1}$	30,000	D2013
ρ_{egg} , egg density	g m^{-3}	0.6×10^3 (min)	JM2013
		19.4×10^3 (mean)	JM2013
		26.8×10^3 (max)	JM2013
μ , dynamic viscosity	$\text{g m}^{-1} \text{ s}^{-1}$	1.9	VJ1999

VJ1999, Visser and Jónasdóttir (1999); D2013, Darnis (2013); JM2013, Jung-Madsen et al. (2013)

lead to unrealistic and spurious results, such as a small fraction of the simulated egg population that has already hatched after the first time step! In order to prevent this, we used the simple approach of spreading the egg development over 20 numerical stages of equal length (Gentleman et al., 2008) and we further integrated the development throughout these stages with a flux limiting numerical scheme (Record and Pershing, 2008).

Simulations

The 1D water column model was split into 5 m vertical layers between 0 and 300 m, and the time step of integration was 12 h. For the model forcing, we used physical and biological datasets from the Circumpolar Flaw Lead System Study (CFL; Barber et al., 2011). We only selected profiles from stations that were at least 300 m deep and located within the Amundsen Gulf. As a result, the physical forcing came from mean vertical profiles of eddy diffusivity coefficient (K), temperature (T), and water density (ρ_{water}) obtained at 20 different stations. These stations were sampled in November and December 2007 under dense sea-ice cover with a vertical microstructure profiler (VMP500, Rockland Scientific International). For two long-term stations the number of casts used in the average profile were respectively 24 and 25, whereas at least 5 profiles were used to build the 18 other mean profiles. Missing values in averaged profiles (e.g., near the surface or below 250 m) were dealt with according to Equation (1) from Bourgault et al. (2011). Further details about this dataset can be found in their study.

We also used 18 vertical profiles of *M. longa* female abundance obtained with a Hydrobios® multinet sampler. Details of the sampling procedure can be found in Darnis and Fortier (2014). We selected the stations sampled between February and April 2008, during the peak of *C. hyperboreus* reproduction. The layer thickness for the vertical sampling ranged from 10 to 144 m, with a median of 20 m.

In order to test the sensitivity of the model to both the physical properties of the water column and the vertical distribution of the predators (*M. longa*) we ran 360 simulations, one for each possible combination of the physical and biological forcing fields. In addition, we ran this ensemble of simulations for three different scenarios of egg density in order to verify the impacts of different egg velocities: the mean (scenario D_0), minimum (D_{min}) and maximum (D_{max}) egg densities observed by Jung-Madsen et al. (2013). Simulations ran for 15 days in order to reach a quasi-steady state where the maximum local rate of change in egg concentration $\delta E/\delta t$ was less than 10^{-6} .

RESULTS

Egg Ingestion Rates

Predation of *C. hyperboreus* eggs by female *M. longa* occurred in each of the incubation experiments. However, in each experiment there was high individual variability and several individual incubations showed no egg ingestion. After discarding incubations within which dead or unhealthy individuals were found at the end of the experiment, for each treatment approximately 8 individual replicates out of the initial 10 were used for further analyses. The frequency distribution of ingestion

rates was positively skewed, i.e., the median value was lower than the mean. The global mean ingestion rate of *C. hyperboreus* eggs by *M. longa* was $5.8 \text{ eggs ind}^{-1} \text{ d}^{-1}$ ($SE = 0.57$, $n = 141$) and the median was $3.7 \text{ eggs ind}^{-1} \text{ d}^{-1}$ (Figure 2). In terms of carbon, the mean ingestion rate was $4.9 \mu\text{g C ind}^{-1} \text{ d}^{-1}$ and the median $3.1 \mu\text{g C ind}^{-1} \text{ d}^{-1}$ ($SE = 0.48$, $n = 141$; Figure 2). *C. hyperboreus* eggs constituted an energy-rich food source ($0.84 \mu\text{g C egg}^{-1}$; (Plourde et al., 2003) and the average daily ration of *M. longa* females feeding on *C. hyperboreus* eggs was 37% of their estimated metabolic needs based on respiration rates ($SE = 4$, $n = 141$). Individual variability resulted in a contrasted pattern where about a quarter of *M. longa* did not ingest any eggs, while an equivalent proportion filled more than 50% of their daily energetic requirements through egg ingestion. Some individuals actually largely exceeded their daily metabolic needs (i.e., over 100%, Figure 2), even when eggs were offered at low concentrations.

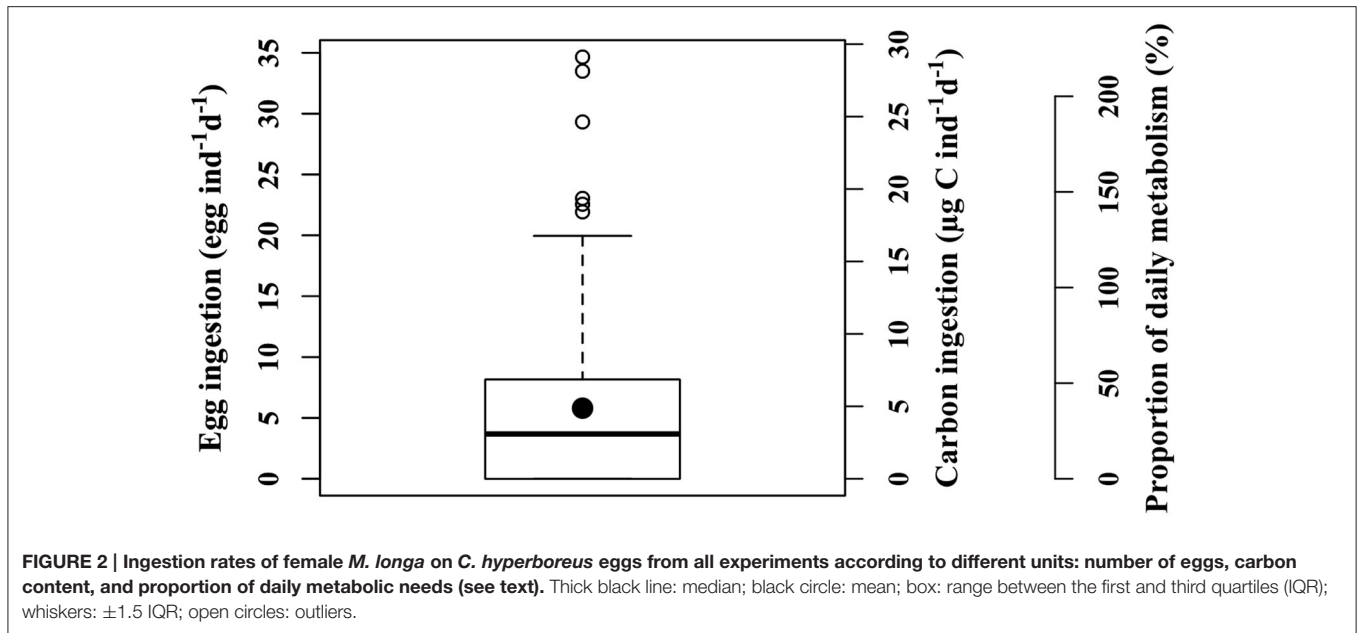
Influence of Incubation Conditions

According to the GLM analysis, presence of additional food source, prosome length, or condition index of the individuals had no discernable effect on egg ingestion (Table 2). The only predictor that improved the model and that significantly affected the number of eggs ingested by *M. longa* was egg concentration (Figure 3). In the full model with no interactions, temperature seemed to have a significant negative effect on ingestion rates (Table 2). However, in the model that only kept egg concentration and temperature as independent variables, the influence of temperature and the interaction term between them did not remain significant. Moreover, the size effect of the temperature and interaction term coefficients was small compared to the impact of egg concentration and the QAIC values were very close between the models. Hence, we decided to use the simplest model with only egg concentration as predictor of egg ingestion (Figure 3). We did not observe any feeding saturation for the range of egg concentrations offered.

Simulated Egg Vertical Distribution

The velocity of *C. hyperboreus* eggs estimated with Equation (7) ranged between -9 and -4.1 m d^{-1} for the mean egg density scenario (D_0) over the 20 physical forcing profiles. Negative velocities meant that eggs were positively buoyant, from the bottom of the water column up to the surface. Egg velocity was dependent on the water density profile and it decreased slowly with decreasing depth (Figure 4). This pattern was conserved among the 20 physical forcing profiles whose overall variability was low. Egg velocity was strongly influenced by egg density itself. *C. hyperboreus* egg density is highly variable, both between individual females and within clutches of the same female (Jung-Madsen et al., 2013). When we used the minimal egg density (scenario D_{min}) the velocity tripled to range between -26.2 and -21.3 m d^{-1} , whereas for maximal egg density (scenario D_{max}) the associated velocity was not negative throughout the water column. The egg velocity ranged between -2.3 and 2.6 m d^{-1} with a converging depth of neutral buoyancy around 100 m.

Egg velocity was critical for egg vertical distribution. In the ensemble of simulations for scenario D_0 , egg vertical



distributions showed higher concentrations between 150 and 275 m and peaked around 225 m to reach about 3000 eggs m^{-3} (or 3 eggs L^{-1}) once the simulation reached its steady state (Figure 4). For scenario D_{max} , the denser eggs ascend only slightly in the water column before hatching. They remained concentrated between 200 and 300 m with a maximum egg concentration a little less than 4000 eggs m^{-3} near 250 m (Figure 4). Contrarily, in the D_{min} scenario, eggs moved rapidly upward and some even managed to reach the first 5 m of the water column to attain a concentration of a little less than 300 eggs m^{-3} (Figure 4). Eggs were spread over the entire column and concentration peaked above 160 m at about 1200 eggs m^{-3} .

Impact of *M. longa* Predation on *C. hyperboreus* Eggs

For each egg density scenario tested, the proportion of egg biomass eaten by *M. longa* was more sensitive to the profiles of *M. longa* abundance than to the physical forcing (see Figure 5 for scenario D_0). Over the ensemble of 360 simulations of the D_0 scenario, the percentage of *C. hyperboreus* egg standing stock ingested by *M. longa* ranged between 0.1 and 1.1% with most of the values being in the low end (Figure 6). The percentage of egg standing stock ingested by *M. longa* varied between 0.2 and 3.2% for scenario D_{min} and between 0 and 0.8% for maximal egg density (scenario D_{max} ; Figure 6).

This modest impact of *M. longa* IGP is hardly noticeable between the egg concentration profiles simulated with and without egg predation, even for the combination of physical conditions and *M. longa* profiles that lead to the maximum difference (Figure 7). The corresponding daily egg ingestion of the whole population of *M. longa* ranged between a little more than 1 to almost 20 eggs $m^{-3} d^{-1}$ (Figure 7). It is noticeable, though, that such egg ingestion values could allow *M. longa* individuals to satisfy almost 10% of their metabolic

TABLE 2 | Results from the generalized linear models (GLM) fitted to predict the number of *C. hyperboreus* eggs ingested by female *M. longa* according to egg concentration (*E*), temperature (*T*), alternative food source (*AC*), condition index (*CI*), prosome length (*PL*), and interactions between the terms.

GLM	Predictor	β	SE	P-value	QAIC
<i>E, T, AC, CI, PL</i>	<i>Intercept</i>	-1.870	3.340	ns	215
	<i>E</i>	0.068	0.012	***	
	<i>T</i>	-0.093	0.040	*	
	<i>AC</i>	0.0007	0.004	ns	
	<i>CI</i>	-0.927	1.118	ns	
	<i>PL</i>	-0.300	1.165	ns	
<i>E, T, E×T</i>	<i>Intercept</i>	-3.365	0.379	***	208
	<i>E</i>	0.111	0.024	***	
	<i>T</i>	0.031	0.078	ns	
	<i>E×T</i>	-0.008	0.004	ns	
<i>E</i>	<i>Intercept</i>	-2.944	0.199	***	215
	<i>E</i>	0.053	0.009	***	

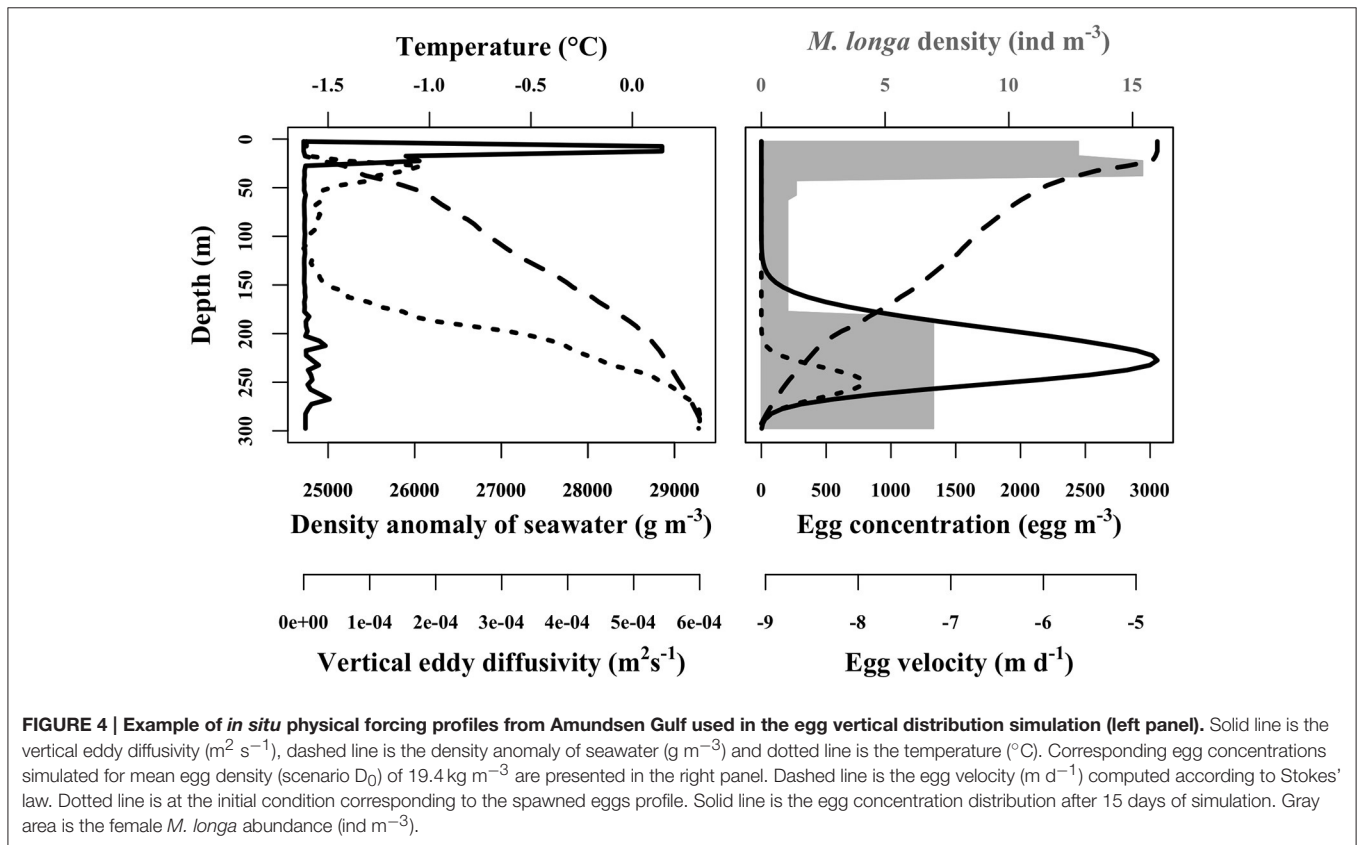
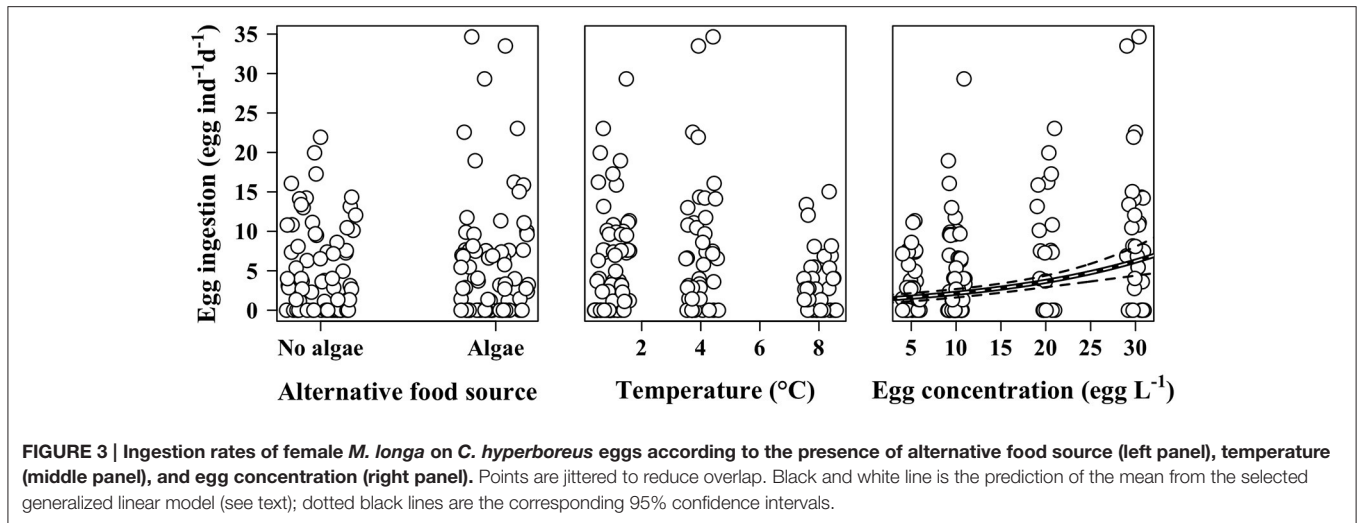
Estimated coefficients *b*, standard error SE, P-values (*** $P < 0.001$; ** $P < 0.01$; * $P < 0.1$; ns not significant) and Akaike's information criterion corrected for overdispersion QAIC. Model entry in bold indicates the one selected for further numerical experiments.

needs according to respiration rates from Seuthe et al. (2006), or up to 37% if we consider the lower respiration rates reported by Hirche (1987).

DISCUSSION

Individual Variability

Our results show high individual variability of egg ingestion rates, with a quarter of all the individuals not ingesting any eggs and about the same proportion satisfying more than half of



their daily metabolic needs from egg grazing. This asymmetric and widely spread distribution has a geometric coefficient of variation of 149%. This pattern is consistent across the range of incubation conditions we tested, and this level of variability is common in any experimental setting measuring individual biological features (size, structural or storage weight, swimming behavior, etc.) and physiological rates (respiration, ingestion, growth, etc.) (e.g., Basedow and Tande, 2006). We do not

have clear explanations for the level of variability we observed, but it is a useful observation to report (see Supplementary Material for a spreadsheet of individual observations). Individual variability has long been recognized as a key property of plankton ecology since population dynamics and trophic interactions (that are of primary interest for marine ecologists) are emerging properties of individual characteristics and behaviors (Båmstedt, 1988). Modern experimental and *in situ* observation methods

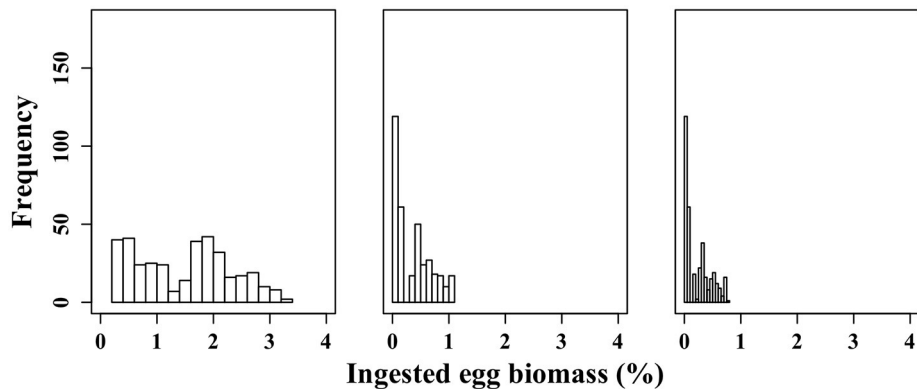


FIGURE 6 | Frequency distribution of the proportion of *C. hyperboreus* eggs standing stock ingested by female *M. longa* according to the minimal (0.6 kg m^{-3}) (scenario D_{\min}) (left panel), mean (19.4 kg m^{-3}) (scenario D_0) (middle panel) and maximal *C. hyperboreus* egg density (26.8 kg m^{-3}) (scenario D_{\max}) (right panel).

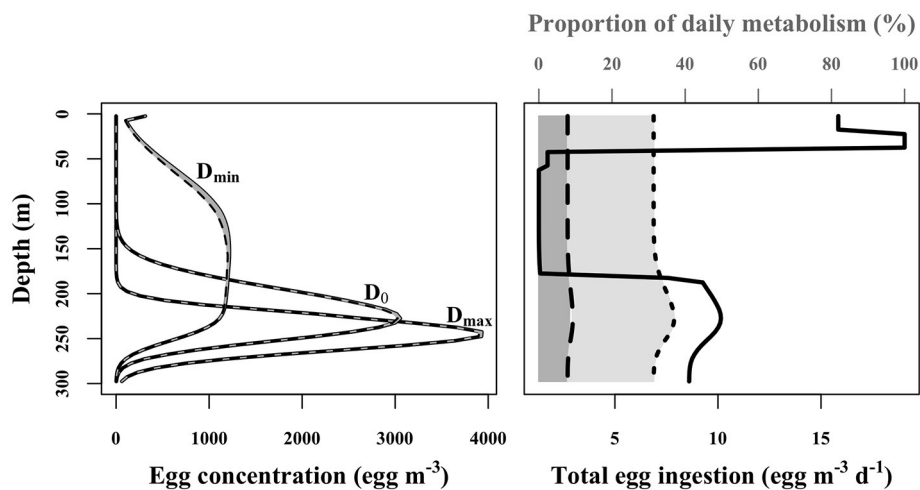


FIGURE 7 | Simulated egg concentration profiles for three different scenarios (scenario D_{\min} , D_0 , and D_{\max} ; left panel). Solid line is without female *M. longa* intraguild predation, dashed line is with intraguild predation and the gray area is the difference between both. On the right panel, solid line is the *C. hyperboreus* egg biomass ingested by the population of female *M. longa* ($\text{egg m}^{-3} \text{ d}^{-1}$) according to the simulated egg concentration with scenario D_0 and *M. longa* abundance profile presented in **Figure 4**. Proportion of individual daily metabolic needs satisfied by egg ingestion (%) according to respiration rates from Seuthe et al. (2006; Dashed line) and Hirche (1987; Dotted line).

C. hyperboreus eggs. As already mentioned, adults and advanced copepodite stages of the large *C. glacialis* emerge from diapause and initiate ascent to surface layers prior the spring bloom to feed on ice algae (Daase et al., 2013). Therefore, they could also benefit from the energy rich *C. hyperboreus* eggs (and potentially the nauplii) and add to the predation pressure. Thus, we likely have underestimated IGP pressure on *C. hyperboreus* and its impact on its recruitment in several ways. However, it may remain very dependent on the seasonal timing of the feeding activity of these potential predators, as well as their vertical position in the water column as already demonstrated.

It seems that there is actually an ecological trade-off for *C. hyperboreus* females to lay low (lipid-rich) or high-density (lipid-poor) eggs. On one hand the obvious metabolic advantage

for lipid-rich eggs is that the offspring can rely on abundant reserves to develop. The downside however could be that low-density eggs reach faster and “en masse” the layer where *M. longa* and other species are abundant, hence increasing their mortality risk. On the other hand, dense eggs ascend very slowly and actually hatch before having reached these dangerous depths. The wide range in egg density that has been observed among eggs from different *C. hyperboreus* females but also within the same egg clutch (Jung-Madsen et al., 2013) could actually be part of a strategy that mitigates predation risk by spreading the eggs across a range of ascent speeds. However, the delicate balance of both opposing effects on offspring fitness could only be assessed in a more detailed modeling study supported by finely resolved and

concurrent observation of the zooplankton community vertical distribution.

C. hyperboreus Importance in the Arctic System

Our results could offer an interesting contrast with the earlier study of Huntley and Escritor (1992) on a couple of homologous species from Antarctica, *M. gerlachei* and *C. acutus*. They showed from incubations of a group of individuals that *M. gerlachei* could reach daily rations ranging between 4 to 11% of its body weight when offered 1000 egg L⁻¹. The authors further estimated that this concentration was about three orders of magnitude higher than what it could be *in situ* (1 egg L⁻¹) and thus concluded that this type of predation is likely insignificant. However, our experimental results showed that *M. longa* individuals could meet up to 75% of their daily metabolic requirements on eggs of *C. hyperboreus* offered at a concentration of 5 L⁻¹, based on respiration rates observed in Amundsen Gulf (Seuthe et al., 2006). We also simulated that the mean *in situ* ration of eggs for *M. longa* females should vary between 8 and 37% of their metabolic needs, depending on the respiration rate. The lower boundary is based on Seuthe et al. (2006), whereas the upper boundary is based on lower respiration rates (Hirche, 1987). Neither value is sufficient to ensure complete metabolic maintenance, but during wintertime it could be combined to some level of lipid stores to cope with the otherwise scarce food background available to these copepods. Lipid-rich *C. hyperboreus* eggs (and potentially nauplii) are a reliable and valuable food source that at least some *M. longa* individuals seem prone to take advantage of during several months of the year.

Our study stresses the importance of *C. hyperboreus* as a linchpin of Arctic marine trophic network. This key species has adapted to the extreme environment by evolving its ability to store considerable amounts of lipids. *C. hyperboreus* efficiently concentrate the energy from the short-lived primary production bloom and supports the entire marine trophic network. Higher trophic levels rely heavily on its large copepodite and adult development stages, while zooplankton species from a similar trophic level (from the same guild) also benefit from the smaller packaging of the bounty within its eggs. Moreover, since *C. hyperboreus* reproduction occurs for several months during a period of the polar year when primary production has shut down, it likely provides a precious and unparalleled resource for several species of planktonic predators that remain active in the dead of winter. Hence, IGP should not only be considered as an extra mortality source that could affect recruitment of the species that is preyed upon, but also as a crucial survival strategy that could shape the life cycle strategies of some opportunistic species.

IGP needs to be studied further as we move toward an integrated approach of marine ecology that recognizes the influence of both individual variability and community-level interactions. Other implications of IGP than those evoked in our study could be important. In the North Atlantic for example, the survival of *C. finmarchicus* early stages follows different

seasonal patterns in areas where it is the dominant *Calanus* species than in areas where its larger congeners, *C. glacialis* and *C. hyperboreus*, co-occur. In such areas where all three species live together, the *C. finmarchicus* recruitment peak can occur several weeks after the spring bloom, much later than it usually does (Melle et al., 2014). In early spring, *C. hyperboreus* and *C. glacialis*, that are already active when *C. finmarchicus* initiates its reproduction fuelled by the phytoplankton bloom, could ingest *C. finmarchicus* eggs and affect its recruitment. Interestingly, IGP could contribute to the resistance of marginal Arctic marine ecosystems to the northward advance of the boreal *C. finmarchicus* under the pressure of climate change.

AUTHOR CONTRIBUTIONS

KD designed and ran the laboratory experiments, participated to the *in situ* sampling, did the statistical analysis, ran the modeling experiment, and wrote most of the paper. FM contributed to the design of the laboratory experiment, contributed to the data analysis, designed, and executed most of the modeling experiment and contributed to the writing. SP contributed to the design of the laboratory experiment, provided access to the laboratory facility, and contributed to the writing. PJ contributed to the design and the execution of the laboratory experiment, and ran the *in situ* sampling. FC contributed to the design of the numerical experiment, provided data from Amundsen Gulf, and contributed to the writing.

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SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <http://journal.frontiersin.org/article/10.3389/fmars.2016.00185>

The R code and forcing fields for the model can be found online at: https://github.com/NEOLab-Git/Chyp_Egg_1D.git

The detailed table of individual observations is provided.

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Conflict of Interest Statement: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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