Impact of aragonite saturation state changes on migratory pteropods

Steeve Comeau1,2,* , Jean-Pierre Gattuso1,2, Anne-Marin Nisumaa1,2 and James Orr3

1 CNRS-INSU, Laboratoire d’Océanographie de Villefranche, BP 28, 06234 Villefranche-sur-Mer Cedex, France
2 Université Pierre et Marie Curie-Paris 6, Observatoire Océanologique de Villefranche, 06230 Villefranche-sur-Mer, France
3 LSCE/IPSL, Laboratoire des Sciences du Climat et de l’Environnement CEA-CNRS-UVSQ, F-91191 Gif-sur-Yvette Cedex, France

Thecosome pteropods play a key role in the food web of various marine ecosystems and they calcify, secreting the unstable CaCO3 mineral aragonite to form their shell material. Here, we have estimated the effect of ocean acidification on pteropod calcification by exploiting empirical relationships between their gross calcification rates (CaCO3 precipitation) and aragonite saturation state (Ωa) combined with model projections of future Ωa. These were corrected for modern model-data bias and taken over the depth range where pteropods are observed to migrate vertically. Results indicate large reductions in gross calcification at temperate and high latitudes. Over much of the Arctic, the pteropod Limacina helicina will become unable to precipitate CaCO3 by the end of the century under the IPCC SRES A2 scenario. These results emphasize concerns over the future of shelled pteropods, particularly L. helicina in high latitudes. Shell-less L. helicina are not known to have ever existed nor would we expect them to survive. Declines of pteropod populations could drive dramatic ecological changes in the various pelagic ecosystems in which they play a critical role.

Keywords: ocean acidification; zooplankton; Arctic

1. INTRODUCTION

Since pre-industrial time, large quantities of carbon dioxide (CO2) have been released into the atmosphere as a result of fossil-fuel burning and deforestation [1]. Atmospheric CO2 concentrations have increased roughly exponentially from about 280 ppm in the eighteenth century to about 390 ppm today. The oceans have buffered this increase by taking up the equivalent of about one-fourth of anthropogenic CO2 emissions [1–3]. As this extra CO2 dissolves in sea water, it forms additional carbonic acid and reduces ocean pH. Since pre-industrial time, surface ocean pH has fallen by about 0.1 unit [4], and models project that under the IPCC IS92a scenario, there will be a further decline of 0.3–0.4 units by the end of the century [4]. Yet, ocean acidification does not only affect the physical chemistry of the sea; they have an affect on some organisms and alter ecosystems. Previous studies demonstrate that some key calcifiers, such as coccolithophores [8], corals and coralline algae [9–11], molluscs [12] and echinoderms [13] exhibit lower calcification rates under the chemical conditions that are expected by the end of the century. However, some studies have found contradictory results (e.g. [14]).

Pteropods are among the calcifiers that are expected to be greatly affected by declining aragonite saturation state (Ωa) (e.g. [15]). These macro-zooplankton are holoplanktonic molluscs that are widely distributed throughout all oceans. Their diel vertical migrations can cover several hundred metres. They are a major dietary component for some zooplankton and higher predators, such as herring, salmon, whales and birds, and they have an active role in the food web of pelagic ecosystems, particularly in high latitudes [16]. Pteropods are also major contributors to fluxes of calcium carbonate [17] and organic matter [18] to the deep ocean. It is expected that pteropods will be highly sensitive to these future changes in sea water chemistry, because they maintain an external calcareous shell made of aragonite [6,19,20], a relatively soluble form of calcium

* Author and address for correspondence: Department of Biology, California State University, 18111 Nordhoff Street, Northridge, CA 91330-8303, USA (steeve.comeau@csun.edu).

carbonate [21]. Recent studies have demonstrated a strong relationship between the $\Omega_a$ and the rates of calcium carbonate precipitation in the Arctic pteropod species *Limacina helicina* [15,22] as well as the Mediterranean species *Creseis acicula* [23].

This study aims to link model projections for chemistry with laboratory data on pteropod calcification rates to estimate future changes in pteropod calcification. Thus, we combined model projections of future $\Omega_a$ with laboratory-derived empirical relationships relating gross calcification rates to $\Omega_a$ in order to estimate the *in situ* calcification rates of pteropods within the depth range of their vertical distribution.

### 2. MATERIAL AND METHODS

#### (a) Data acquisition

For our analysis, we chose sites having co-location of three types of data: (i) for the vertical distribution of pteropod species that have known responses of their calcification rate to the aragonite saturation state, (ii) for nearby carbonate chemistry, and (iii) for model projections of the aragonite saturation state projections. Data on the vertical migration of pteropods were obtained from six publications and one personal observation (table 1). These data provide information on vertical ranges of the high-latitude species *L. helicina* at five sites in the Arctic Ocean and two sites in the Southern Ocean (figure 1). Because the Arctic data from Kobayashi [24] covered a wide area, that dataset was divided into three different sites. Additionally, data on another species, *C. acicula* were available at two study sites (Sargasso Sea and Hawaii).

For simplicity, we consider that both species migrate vertically from the surface to 200 m, because vertical migrations at the different sites generally varied from 150 to 250 m for both *L. helicina* and *C. acicula*. For brevity, we have selected to provide details for two sites in the Arctic, one site in the Southern Ocean, and one site at temperate latitudes (sites 1–4, respectively). Particular attention was paid to the Arctic, the first region that is expected to become undersaturated with respect to aragonite [7]; it is also a region where pteropods play a major ecological role.

#### (b) Institut Pierre Simon Laplace model

Based on previous comparison efforts [6], differences between model projections over the next 100 years are relatively small after each model is corrected for its modern-day model-data bias. For simplicity, we used output from one Earth System Model (from Institut Pierre Simon Laplace, IPSL-CM4) forced under the IPCC SRES A2 scenario. Simulated fields of carbonate chemistry variables from the IPSL-CM4 model were used in this analysis. The climate system in this coupled model includes an atmospheric general circulation model (LMDZ4 [28]) that is coupled through the OASIS coupler to an ocean general circulation model (OPA8 [29]) that is itself coupled to a sea ice model (Louvain-la-Neuve ice model). Imbedded in this climate system are modules for the terrestrial biosphere (ORCHIDEE [30]) and the marine carbon cycle (PISCES [31]).
More about the IPSL-CM4 Earth System Model and its validation can be found in Cadule et al. [32,33].

(c) Model correction

Data from stations in the CARINA database (CARbon IN the Atlantic [34]) nearest to the study sites were used both to evaluate present-day model behaviour and improved model projections. Observed total alkalinity (\(A_T\)), dissolved inorganic carbon (\(C_T\)), pressure, salinity, temperature and phosphate and silicate concentrations, collected between 2000 and 2005 depending on the site, were used to determine the corresponding \(V_a\) with the \texttt{R} package seacarb [35]. The calculated saturation states were then compared with model results from the same year. The model-minus-data difference, i.e. the model bias, was then subtracted following the approach of Orr et al. [6] in order to improve future projections.

(d) In situ calcification

The relationship between gross calcification (\(G_{\text{gross}}\)) and \(\Omega_a\) for the Arctic pteropod \textit{Limacina helicina helicina} (\(G_{\text{gross}} = 0.57 \ln (\Omega) + 0.25\)) was taken from Comeau et al. [15]; the relationship for the temperate pteropod \textit{C. acicula} (\(G_{\text{gross}} = 0.39 \ln (\Omega) + 0.3\)) was taken from [23]. Known genetic differences between the Arctic pteropod \textit{L. helicina}, and the Antarctic pteropod \textit{Limacina helicina antarctica} [36], were not considered for this analysis. Rates measured on Mediterranean species \textit{C. acicula} were used in the two subtropical study sites, (Sargasso Sea and near Hawaii), where \textit{C. acicula} is also found. As described in Comeau et al. [15], temperature effect was mainly reflected by its influence on the \(V_a\). Relationships between CaCO$_3$ precipitation and \(V_a\) were not significantly different on pteropods incubated at 0°C and 4°C. Differences in temperature were then neglected. Annual-mean model output for the \(\Omega_a\) was used to compute the corresponding \(G_{\text{gross in situ}}\). Mean \(G_{\text{gross}}\) were calculated by assuming that pteropods migrate daily between the surface and 200 m. They are expressed in per cent relative to the estimated pre-industrial \(G_{\text{gross}}\).

3. RESULTS

The projections of the IPSL model for the \(\Omega_a\) are generally consistent with values derived from field measurements (figure 2, table 2 and electronic supplementary material,
The mean of the model minus data for \( V_a \) over the top 200 m along with its propagated error are 0.11 ± 0.10, 2.0 ± 0.35, 2.0 ± 0.26 and 0.38 ± 0.17 at sites 1–4, respectively. The bias-corrected model projects that at site 1 in the Arctic, the mean \( V_a \) within the migration-depth range of pteropods was 1.5 in 1860, 1.2 in 2005, but only 0.4 at the end of the century (figure 3). On the west coast of Spitsbergen, at site 2, the mean \( V_a \) within the same depth range was 2.0 in 1860, 1.8 in 2005 and only 1.1 at the end of the century (figure 3). On an annual average, in the Arctic, these near-surface waters become undersaturated with respect to aragonite at all study sites by the end of the century (figure 3 and electronic supplementary material, figure S2). The decline is more pronounced at study site 1 (83.5º N, 98.5º W), where undersaturation is expected to occur over the top 200 m by 2050 (figure 3), relative to waters around Spitsbergen (site 2; 79º N, 11º E), where undersaturation is projected to occur only by the end of the century (figure 3 and table 3). Near Spitsbergen though, at the end of the century, pteropods still maintain some positive \( G_{\text{gross}} \) relative to preindustrial rate because...
The projected shoaling of less-saturated waters (figure 3) is also directly measurable in the field. In the Iceland Sea, measurements [40] demonstrate that the ASH is currently shoaling at a rate of 4 m yr⁻¹. This shoaling and the general reduction of \( \Omega_a \) in near-surface waters will result in increased exposure of pteropods to less-saturated waters throughout their diurnal migrations. It is unknown if these changes in water chemistry could prompt pteropods to reduce their vertical migration length scale in order to reduce their exposure to less-saturated waters. Pteropods can be found as deep as 1200 m in some seasons [24], and thus may be already exposed to undersaturated waters during seasonal migration. At present, any such exposure must be limited in time. Although the exact reasons for these deep vertical migrations of pteropods and the associated physiological impacts are not known, zooplankton generally migrate diurnally or seasonally to reduce the predatory pressure [41]. Zooplankton typically feed near the surface, which is rich in food, but also in predators; they migrate to deeper layers to hide from their predators. For example, in the Kongsfjorden (Svalbard), juveniles of \( L. helicina \) overwinter at about 200 m and switch to feed on lower quality food [42,43]. If pteropods reduce their depth range in response to acidification or other anthropogenic pressures, that would also result in greater predatory pressure, notably on juveniles.

Our results highlight the critical situation that \( L. helicina \) will face in the coming decades in the Arctic Ocean. Moreover, these projections are conservative because they concern rate of \( G_{\text{gross}} \) rather than rates of net calcification, which is the balance between \( G_{\text{gross}} \) and dissolution of calcium carbonate. Calcification rates were indeed measured using the \(^{45}\text{Ca} \) incorporation technique [15], which estimates the active incorporation of \( \text{CaCO}_3 \) occurring mostly at the growing edge of the shell (aperture), but does not provide information about shell dissolution rates. In other words, even when pteropods precipitate \( \text{CaCO}_3 \) at the shell aperture, there may be simultaneous dissolution across the entire shell surface [23]. At some point, as \( G_{\text{gross}} \) declines and dissolution of \( \text{CaCO}_3 \) increases, rates of dissolution would become larger than rates of \( G_{\text{gross}} \) and net calcification would become negative.

In recent decades, as the \( \Omega_a \) has declined, effects from corresponding reductions in pteropod calcification rate on abundance have yet to be detected. The 1951–2008 time series of zooplankton in Californian coastal waters reveals no evidence for declines in the abundance of \( \text{Thecosomata} \) pteropods [44], despite a measurable reduction of \( \Omega_a \) in the same region [39]. Similar time series have not been investigated elsewhere.

To what extent might \( \text{Thecosomata} \) pteropods be able to survive without a shell? A laboratory study on larvae of the Mediterranean pteropod \( Cavolinia inflexa \) shows that larvae can be viable without a shell when they are reared in waters that are undersaturated with respect to aragonite [45]. Another study reports that larvae of the Arctic pteropods \( L. helicina \) are viable after 29 days of...
laboratory incubation in waters that are undersaturated with respect to aragonite ($\Omega_a = 0.86$), although their shells exhibit extensive degradation [43]. We hypothesize that pteropod shells defend against small predators, such as small decapods, small medusa or small fish larvae, thus preventing them from feeding on shielded pteropods. Pteropod shells also offer their hosts a means to regulate their buoyancy and thus their diurnal migrations [46]. Moreover, for pteropod species of the order Limacinidae, the shell helps maintain the structural integrity of their soft body parts [45], which could make life without a shell unviable. We hypothesize that as the ocean warms and L. helicina migrate poleward, confined by its affinity for waters cooler than 5°C [47], it will be trapped by the shoaling aragonite horizon pushing pteropods nearer to equator. Throughout the northern oceans, L. helicina could thus disappear entirely by the end of the century after becoming incapable of calcifying enough to offset dissolution and other factors of shell erosion.

The loss of Arctic pteropods could have major biogeochemical, ecological and economic impacts. The potential impacts of their disappearance on fish populations and other predators, such as zooplankton, seabirds or whales will depend on the capacity of these predators to modify their diet. Fish populations could be affected from the bottom-up control of zooplankton, as already demonstrated regarding cod recruitment [48,49]. Another example is the potential impact on juvenile pink salmon, whose diet can consist of up to 60 per cent of L. helicina [50]. Without L. helicina, how would the required permanent switch to other prey of different nutritional value and abundance impact juvenile pink salmon? In any case, reductions in pink salmon, the most abundant salmon in the Pacific and the mainstay of North Pacific fisheries, would have major economic repercussions. Evaluating how the disappearance of pteropod populations would affect both economics and ecosystems remain as critical future research challenges.

We have provided regional estimates of the changes in pteropod calcification rates during the industrial era until the end of this century by linking model projections for chemical changes with laboratory data correlating those changes to pteropod calcification rates. More than ever, there appears little future for high-latitude shielded pteropods, particularly the dominant species L. helicina, assuming that atmospheric CO$_2$ continues to increase during the coming decades.

This work is a contribution to the ‘European Project on Ocean Acidification’ (EPOCA) which received funding from the European Community’s Seventh Framework Programme (FP7/2007-2013) under grant agreement no. 211384. It is also a contribution to the MedSeA project. We thank P. Cadule and L. Bopp for allowing us to use results from their simulations with the IPSL-CM4 model. We also thank A. Olsen for providing the CARINA data and Gaby Gorsky for fruitful discussions.

REFERENCES


