# Egg hatching rate of the cyclopoid copepod Oithona similis in arctic and temperate waters

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Running head: Egg hatching rate of Oithona



An equation is presented to facilitate estimation of the production of the cosmopolitan cyclopoid copepod Oithona similis The egg hatching rate was studied from Arctic, subarctic and temperate waters covering a temperature interval from -1 to 20.5 °C. Within this temperature range the hatching rate (HR) increased from 0.03 to 0.42 d<sup>-1</sup>. Results from all experiments were fitted to a function HR (% d<sup>-1</sup>) = 4.2176+1.7545\*T (r2=0.98; P<0.0001). When combined with site-specific information on temperature, egg:female ratios and the carbon content of females and eggs, secondary production of this ubiquitous species can be readily

### Introduction

Egg production of free spawning copepod species have routinely been used to estimate copepod production, assuming that adult female copepods do not grow, but rather allocate the ingested carbon into the production of eggs. The weight specific egg production rate (SEP) of the females is often assumed to equa the growth rate of the younger stages and the production can therefore be easily estimated from the SEP and the standing stock (but see Hopcroft & Roff, 1998, Sabatini & Kiørboe 1994). While the SEP of free spawning copepods is generally estimated over a single 24 hour interval, this method can not be directly applied to egg carrying species that typically produce clutches of eggs less constantly and then carry the same clutch for several days. These include all cyclopoids, poecilostomatoids and harpacticoids plus the important calanoid genera Pseudocalanus, Euchaeta, Clausocalanus that combined constitute a significant fraction of marine copepods.

Of the egg carrying marine copepods, the small cyclopoid Oithona similis is a cosmopolitan species with a wide geographical distribution, from the poles to equator (Nistida 1985, Mazzocchi et al. 1995). Where investigated, *Oithona* has been shown to be one of the most abundant marine copepod genera (Turner, 1982: Paffenhöffer 1993, Calbet & Agusti 1999), Unfortunately the recommended use of nets with a mesh size of 200 µm for sampling of copepods (UNESCO 1968) still bias our knowledge about the quantitative importance of many small copepod species such as *Oithona*. Resent investigations using nets with smaller mesh size (e.g. 45 to 64 µm) or water bottles have documented that *Oithona* contributes significantly to the standing stock of copepods in many marine ecosystems (Paffenhöffer 1993, Gonzales & Smetacek 1994, Nielsen & Sabatini 1996, Hopcroft et al.. 1998). Knowledge about its production and potential grazing impact is therefore of key importance to the understanding of the productivity and dynamics of the Sea

The population specific egg production rate (SEP, d-1) of egg carrying copepods can be accurately estimated by the egg-ratio method (Edmondson, 1971). This method requires knowledge of the egg/female ratio of the population (i.e. including females not carrying eggs), the egg hatching rate (HR, d<sup>-1</sup>) at *in situ* temperature, and the carbon content of the egg and female:

## SEP = (Egg /female) \* HR \* (egg C/female C)

Sabatini & Kiørboe (1994) have previously estimated the relationship between carbon content and the size of both eggs and females for *Oithona similis*. The aim of this paper is to establish the quantitative relationship between temperature and egg hatching rate for *Oithona similis* to provide a simple method of estimating the production of this abundant copepod without routine experiments.



Table 1 Range in surface salinity, temperature, chlorophyll a, female cephalothorax length + SE and sidered Numbers in a

-55	Northwater	Disko Bay	Greenland Sea	Gulf of Alaska	North Sea 14	
Temperature,	-1.551.51	5-7	7.3-0.4	5-15		
in situ*	-0.4	2	-1.2	5-8	7	
Experimental temperature	-1, 1,4	0.2, 4.5, 7.5	-0.8, 2.3, 4.2	5,10,16, 18.5, 20.5	12	
Salinity (PSU)	30.3-30.4	32.6-33.6	32.7-35.0	29.0-32.4	34-35	
Chlorophyll a (µg chl a l-1)	4.1-5.0	1-3	0.3-1.0	0.73-2.0	0.1-0.5	
Female length	477±5	441 ±9	473±3	454±6	532±26	
(µm)	(83)	(229)	(106)	(28)	(440)	
Egg diameter	67.2±1.5	58.3±0.4	63.1±0.4	64.5±1.1	56.9±0.9	
(um)	(35)	(165)	(439)	(36)	(600)	

<sup>\*</sup> If a thermocline was present, second line indicates the temperature below the thermocline.

### Results

Across the different systems, the environment spanned a broad range (Table 1). The salinity at the different sites was the same (29.0-35 psu), while the temperature obviously increased from the arctic to the temperate regions. Chlorophyll varied an order of magnitude between locations, but in no systematic pattern with respect to water temperature. Egg size was comparable at all locations.

The eggs in the sacs developed relatively synchronously until hatching. On several occasions we observed that nauplii escaped from the egg sacs within minutes of having hatched. In other cases hatching appeared to occur over as several hours, with nauplii frequently remaining attached to the female for some time by remnants of the opened egg sac. In general the hatching success was high (> 95%). During the experiments no female mortality was observed, although during the Disko Bay and Gulf of Alaska cruises where some of the females were lost from the wells due to rough seas.

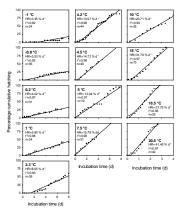
The egg hatching time was inversely related to the water temperature, decreasing from 25.7 d<sup>-1</sup> to 2.8 d<sup>-1</sup> across the temperature range tested (-1.0 to 20.5 °C) (Figure 2). Several equations were fit to the data, many of which provided good overall statistical fit (Table 2). However, for those models with 3 fitted parameters at least one parameter was not significant. The linear models (Figure 3) gave the consistently better fit compared to the exponential models, however, the Bělehrádeks model with exponent fixed at 2.05 (McLaren et al. 1969) proved as satisfactory as the linear model and gave comparable fit. We advocate the linear models between hatching rate (HR, % day1) or hatching time (HT, in days) and temperature (T, °C) which are mathematically simpler:

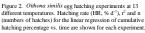
HR = 4.2176+1.7545\*T, r2 = 0.98, P < 0.0001, n=16

HT = (0.0464+0.0145\*T)<sup>-1</sup>, r<sup>2</sup> = 0.97, P <0.0001, n=16

Table 2. Statistical summary of different models examined for HR and HT vs. temperature. For the

		a	(±S.E).	b	(±S.E.)	c	(±S.E.)	r
Hatching rate	a+b*T	4.2176	(0.6721)	1.75451	(0.0665)			0.98
vs. temperature (T)	a*e <sup>(b*T)</sup>	7.968	(0.7663)	0.0845	(0.0060)			0.94
	c+a*e(6*T)	67.229	(43.211)	0.0214	(0.0114)	-62.351	(46.6143)	0.98
	a*(T+c) <sup>b</sup>	0.7027	(0.4981)	1.2655	(0.1954)	4.6412	(1.9087)	0.98
	a*(T+c)2.05	0.0327	(0.0029)			12.793	(1.1072)	0.98
Hatching time	(a+b*T)"	0.0464	(0.0014)	0.0145	(0.0012)			0.97
vs. temperature (T)	a*e(-b*T)	23.610	(0.9293)	0.2088	(0.0194)			0.95
	c+a*e(-b*T)	19.942	(0.5414)	0.3107	(0.0277)	3.2955	(0.5414)	0.98
	a*(T+c)'b	344.12	(444.29)	1.5759	(0.4258)	5.6103	(1.9311)	0.98
	a*(T+c)-2.05	1504.5	(202.13)			7.6998	(0.4970)	0.98





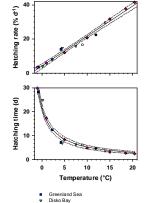


Figure 3. Oithona similis egg hatching rate a) and hatching time

Halifax (Faton 1971)

### Discussion

Of the egg carrying marine copepods, the cyclopoid Oithona similis exists over a wider range of temperatures and salinity than most other marine copepods from temperate brackish coastal water to subtropical oligotrophic oceans (Nistida 1985, Mazzocchi et al. 1995). In cold areas like the arctic and temperate regions, Oithona is often the most important winter copepod genus present, and reproduces year round in surface waters (Kiørboe & Nielsen 1994, Uye & Sano 1995)

One potential shortcoming of this method is that it presumes the animals incubated are randomly distributed throughout their egg-carrying cycle. If egg-laying (and hatching) follow a strongly diel cycle (Hopcroft & Roff 1996, Ambler et al.. 1999), then there will be a bias introduced, creating a step-like pattern in the percentage hatching. At cold temperatures, when hatching time is long, this causes relatively little error in the fina estimation of hatching rate. If the method is applied in the tropics, it would appear necessary (and be logistically feasible) to observe both the production and hatching of clutches to estimate the hatching time (e.g. Hopcroft & Roff 1996).

Previous investigations of Oithona species batching or development time cover a higher or smaller temperature range than this study e.g. Oithona davisa – 10 to 30 °C (Uye & Sano 1995, 1998) and Oithona similis - 4.5 to 14 °C (Eaton 1971). Eaton noted that that her value at 4.5 °C might be suspect, as we have confirmed, limiting her reliable data to only 9 & 14 °C. Thus, our hatching rate measurements at colder and extended temperatures, make the equations applicable for a much larger geographical range.

To our knowledge, this note is the first attempt to establish a general equation for estimation of hatching rates of this very important copepod covering the full range of temperatures from arctic to temperate waters. The applied multi-well technique is low cost, space efficient, and allows rapid handling of many replicates yielding an easy establishment of temperature-dependent hatching rate relationships for sac spawners. This facilitates routine estimation of productivity. More importantly, for preserved finer-mesh samples that contain both females and their detached egg sacs, our equations provide a critical step that allows for prediction of secondary production of this abundant but often ignored component of the copepod community.

Ambler, J.W., F.D. Ferrari, J.A. Fornshell & E.J. Buskey. 1999. Diel cycles of molting, mating, egg sac production and hatching in the swarm forming cyclopoid copepod Dioithona oculata. Plankton Biol. Ecol. 46: 120-127.

Calbet A, Agusti S. (1999) Longitudinal changes of copepod egg production rates in Atlantic waters: temperature and food availability as the main driving factors. Mar. Ecol. Prog. Ser. 181: 155-162.

Eaton J M (1971) Studies on the feeding and reproductive biology of the marine cyclopoid copepod Oithona similis, Claus. PhD. Thesis.

Edmondson, W.T. 1971, Reproductive rates determined directly from egg ratio, pp. 165-169, In: W.T. Edmondson & G.G. Winberg (ed.) A manual on methods for assessment of secondary production in fresh waters, Blackwell Scientific, Oxford

Gonzales HE, Smetacek V.(1994) The possible role of the cyclopoid copepod Oithona in retarding vertical flux of zooplankton faecal material. Mar. Ecol. Prog. Ser. 113: 233-246

Hopcroft R R, Roff JC (1996). Zooplankton growth rates: diel egg production in the copepods Oithona, Euterpina and Corycaeus from tropical waters. J. Plankton Res. 18: 789-803.

Hopcroft RR Roff JC (1998). Zooplankton growth rates: the influence of female size and resources on egg production of tropical marine cooecods. Mar. Biol. 132: 79-86.

Hopcroft RR. Roff JC, Lombard D (1998). Production of tropical copepods in the nearshore waters off Kingston, Jamaica: the importance of small species. Mar. Biol. 130: 593-604.

Kiørboe T, Nielsen TG (1994). Regulation of zo-oplank-ton biomass and produc-tion in a temperate, coastal ecosy-stem. I Copepods.

Mazzocchi MG, Zagami G, lanora I, Gugliemo L, Crescenti N, Hure J (1995). Atlas of Marine Zooplankton Strait of Magellan, Copepods. In: L. Guglielmo and A. Inora (eds.) Springer-Verlag Berlin Heidelberg New York

McLaren IA, Walker DA, Corkett CJ (1968). Effects of salinity on mortality and development rate of eggs of the copepod

nus minutus. Can. J. Zool. 46: 1267-1269.

McLaren IA., Corkett CJ. Zillioux EJ (1969). Temperature adaptations of copepod eggs from the Arctic to the tropics. Biol. Bull. 137:

Nielsen TG, Sabatini M (1996): The role of the copepod Oithona spp. in North Sea plankton communi-ties. Mar. Ecol. Prog. Ser.139:79-

Nistida S (1985), Taxonomy and distribution of the family Oithonidae (Copepoda, Cyclopoida) in the pacific and Indian Ocean, Bull

Paffenhöffer G-A (1993) On the ecology of marine cyclopoid copepods (Crustacea, Copepoda). J. Plank. Res. 15: 37-55 Sabatini M, Kierboe T (1994). Egg production, growth and development of the cyclopoid copepod Oithona similis. J. Plank. Res. 16:

Turner JT (1982). The annual cycle of zooplankton in a Long Island estuary. Estuaries 5: 261-274

UNESCO (1968), Monographs on oceanographic methodology, No 2, Zooplankton sampling, UNESCO, Paris,

Uye S-I, Sano K (1995) Seasonal reproductive biology of the small cyclopoid copepod Oithona davisae in a temperate eutrophic inlet. Mar. Ecol. Prog. Ser. 118: 121-128

Uye S-I, Sano K (1998) Seasonal variation in biomass, growth rate and production rate of the small cycloppoid copepod Oithona davisae in a temperate eutrophic inlet. Mar. Ecol. Prog Ser. 163: 37-44

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