

## SHORT NOTE

## International Study on *Artemia*\*. XV. Effect of Light Intensity on Hatching Rate of *Artemia* Cysts from Different Geographical Origin

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**ABSTRACT:** *Artemia* cysts from 4 geographical strains were incubated under standard hatching conditions at 5 different light intensities. Hatching rates increased with increasing light intensities. The light-intensity threshold at which the maximal hatching rate was attained, varied from strain to strain. Differences in light sensitivity between the *Artemia* strains studied can, at least partly, be attributed to variation in chorion characteristics. The hypothesis is discussed that light might act as a diapause inhibitor in marine and freshwater branchiopods.

In addition to temperature, salinity, pH, and oxygen content of the incubation medium, light conditions influence the hatching efficiency of *Artemia* cysts (Sorgeloos, 1980). Increased hatching efficiencies in light, as compared to controls incubated in darkness, was first described for *Artemia* by Sorgeloos (1973). In the meantime this phenomenon has been confirmed for the same as well as for several other strains by Meade (1976), Royan (1976), Person-Le Ruyet and Salaun (1977) and Spektorova and Syomik (1979).

From a recent review on cyst hatching in *Artemia* (Sorgeloos, 1980), it appears that hatching rate is much more susceptible than hatching efficiency to fluctuations of temperature and salinity. As a consequence one may expect that varying light intensities also affect the hatching rate in *Artemia* cysts. Furthermore, since important differences have been reported for various

cyst-characteristics (Vanhaecke and Sorgeloos, 1980) the light effect may vary from one *Artemia* strain to another. We have studied the effect of varying light intensities on the hatching rate of *Artemia* cysts from 4 geographical strains: Buenos Aires (Argentina), Chaplin Lake (Canada), Great Salt Lake (Utah), and San Pablo Bay (California). From the latter strain both untreated and decapsulated cysts (treated according to methods described in Bruggeman et al., 1979, 1980) have been used for experimentation.

Cylindroconical hatching tubes made of glass were placed at different distances from a fluorescent light tube (Philips, 65 W) in order to expose the tubes to light intensities of 20, 100, 500, 1000 and 2000 lux, respectively. For each strain 5 replicate tubes were set up each with 250 mg cysts incubated in 100 ml natural seawater (35 ppt) at 25 °C. The cysts were kept in suspension by gentle air-bubbling. The hatching rate was followed by subsampling at intervals of 1 h : i. e. 5 subsamples of 250 µl each were taken with an automatic micropipet and the number of nauplii counted. This sampling procedure was continued until the maximum hatching efficiency (determined prior to experiments, following the method of Sorgeloos et al., 1978, at 1000 lux) was attained, or until no increases in hatching efficiency were noted in 3 consecutive series of samples. The mean values for the 5 subsamples, taken every hour, were expressed as percentage of the maximal hatching efficiency. For each light experiment non-linear regression lines were used to calculate non-linear regression lines, employing the method of orthogonal polynomials. Subjecting these regression lines to an analysis of variance revealed the best fit representing the hatching curve.

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The results for the 4 strains studied are graphically represented in Figure 1. As a general trend, hatching rates increase with increasing light intensities until, from a certain intensity onwards, they remain constant. Great variation is noted among the strains studied: (1) Great Salt Lake cysts, revealed minimal differences in hatching rate in different light intensities. (2) Buenos Aires and San Pablo Bay cysts exhibit, increased sensitivities for light; their light-intensity thresholds are situated between 500 and 1000, respectively, 100 and 500 lux. (3) Chaplin Lake cysts appear to be very sensitive to light; not only their hatching rate but also their hatching efficiency varies considerably as a function of experimental light intensities. (4) In San Pablo Bay cysts, chorion removal results in a drop of the light intensity threshold from 100–500 lux in untreated cysts (Fig. 1 C) to 20–100 lux in decapsulated cysts (Fig. 2).

According to Sorgeloos and Persoone (1975) *Artemia* cysts are susceptible to light triggering as soon as they have reached full hydration under aerobic conditions. Sorgeloos and Persoone also noted that a minimal dose of light energy is needed to trigger onset of metabolism in the encysted embryo. We extrapolate from our present findings that, as illumination intensity is lowered below the critical light intensity threshold for maximal hatching rate, progressively more accumulation time is

needed to reach the triggering dose. As a consequence hatching rate is affected.

Light absorption by the cyst chorion has been reported in *Triops* by Hempel-Zawitkowska (1970) and in *Artemia* by Iwasaki et al. (1980). Lowering of the light-intensity threshold in decapsulated versus untreated cysts exemplifies this phenomenon. Strain differences could eventually be attributed to differences in chorion characteristics, e.g. in Buenos Aires, San Pablo Bay and Great Salt Lake *Artemia* a positive correlation was found between chorion thickness (10.4, 7.6 and 4.7  $\mu\text{m}$ , respectively; Vanhaecke and Sorgeloos, 1980) and light-intensity threshold ranges for maximal hatching rate (500–1000, 100–500 and 0–20 lux, respectively). This correlation, however, cannot be extrapolated to all *Artemia* strains; Chaplin Lake cysts, for example, have a thin chorion (5.35  $\mu\text{m}$ ; Vanhaecke and Sorgeloos, 1980) but require 1000 to 2000 lux to assure a maximal hatching rate. Other parameters could interfere with the chorion-thickness criterium: e.g. differences in concentration of the chorion pigment haematine (Gilchrist and Green, 1960) – which is known to be responsible for the light-absorption characteristic of the chorion (Hempel-Zawitkowska, 1970) – or variations in the number of those cysts that do not need a light stimulus and can hatch in

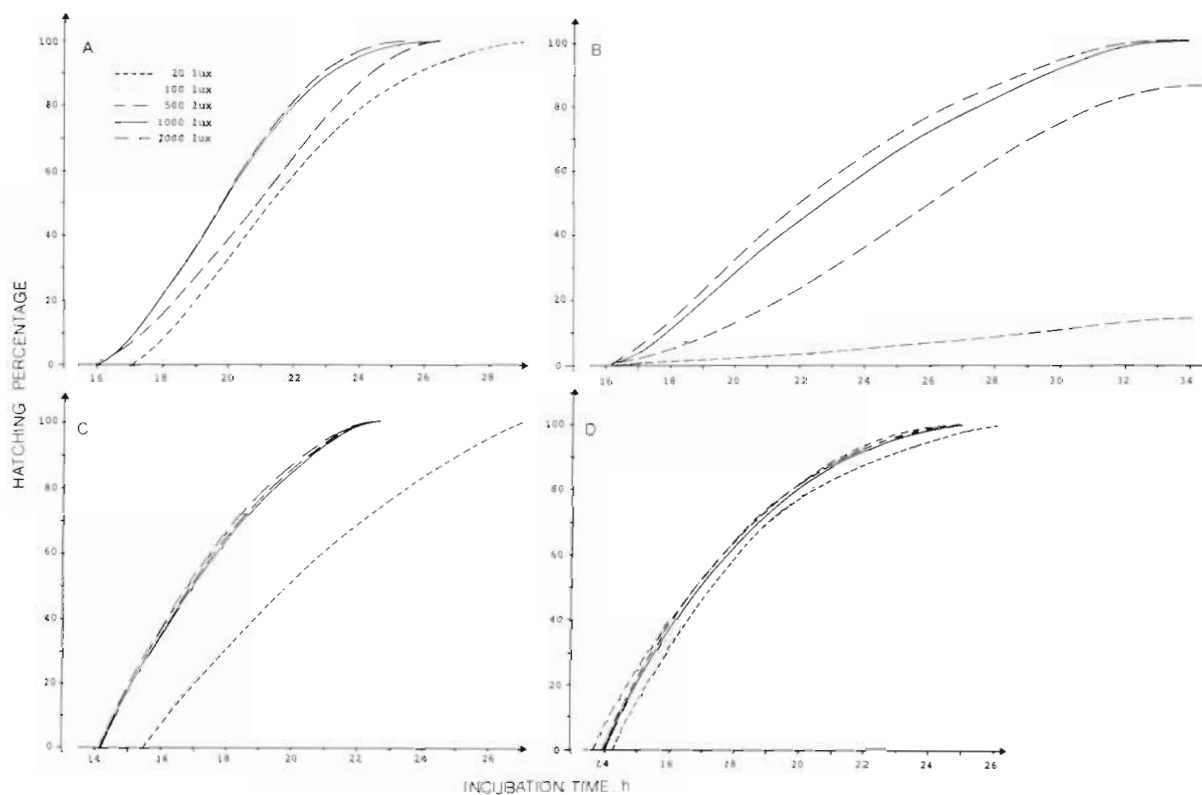


Fig. 1. *Artemia* sp. Effect of light intensity on hatching rate of cysts from (A) Buenos Aires, (B) Chaplin Lake, (C) San Pablo Bay, (D) Great Salt Lake

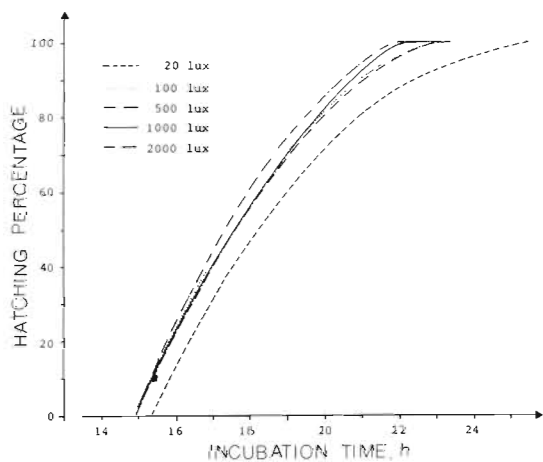


Fig. 2. *Artemia* sp. Effect of light intensity on hatching rate of decapsulated cysts from San Pablo Bay

darkness (Sorgeloos, 1973), possibly due to differences in cyst-processing and/or cyst-storage conditions (Sorgeloos et al., 1976). In view of important differences between strains reported for many other cyst characteristics (Bruggeman et al., 1980; Vanhaecke and Sorgeloos, 1980), strain-specific differences in the light sensor mechanism can also be expected. As long as this mechanism is not identified, all hypotheses on light triggering of the hatching metabolism in hydrated *Artemia* cysts must remain speculative.

Since the need for light stimulation in cyst hatching of the freshwater conchostracan *Eulimnadia antlei* can be removed by drying 'at least in some cases' (Belk, 1972), the light effect on cyst hatching in branchiopods could be a diapause inhibitor rather than a trigger of hatching metabolism. Indeed various kinds of 'environmental shocks' (Prophet, 1963) have been reported to inhibit diapause and to stimulate nauplii production in branchiopod cysts, e. g. osmotic shock in *Branchipus* (Mathias and Bouat, 1934); drying in *Artemia* (Dutrieu, 1960) and in *Triops* (Klekowski and Hempel-Zawitkowska, 1968); UV-radiation in *Triops* (Hempel-Zawitkowska, 1970); freezing in *Artemia* (Hempel-Zawitkowska, 1971a) and in *Triops* (Hempel-Zawitkowska, 1971b); magnetic fields (Dolgopol'skaya et al., 1969), ionic radiation (Metalli and Ballardini, 1972), exposure to organic solvents (Tazawa and Iwanami, 1974) and cosmic radiation (Planel et al., 1975) all in *Artemia*. Hatching of a small percentage of brine shrimp cysts in darkness (Sorgeloos, 1973) could then be explained by partial diapause inhibition in the dried cysts.

In conclusion, a better understanding of diapause inhibition in *Artemia* cysts could eventually lead to improved processing techniques for assuring maximal hatching rates in darkness. In the meantime, minimal light intensities of 1000 lux are advised for hatching

*Artemia* cysts from origins different from the ones referred to in this paper.

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