# INFLUENCE OF LOW AND HIGH TEMPERATURE ON DIURNAL PERIODICITY OF TETRASPORE SHEDDING IN SOME RED ALGAE

# ABSTRACT

Influence of low and high temperature on diurnal periodicity in tetrespore output from Gelidium pusilium, Petrocladia heteroplatos and Gelidiopsis variabilis was investigated. In general, peak shedding of spores was delayed than the normal periodicity in G. pustllum and G. variabilts at low (0 and 20°C) as well as high (40°C) temperature. Temperature plays a major role in altering the peak liberation of tetraspores in a day in these three red algae.

Information on the spore producing capacity of commercially important seaweeds is necessary to cultivate them by reproductive propagation method. Studies were made by several workers on spore shedding of different marine algae Only very fragmentary information is available

Subbarangajah, 1981; Umamaheswara Rao and Kaliaperumal, 1983; Subbarangajah, 1985; Kaliaperumal and Umamaheswara Rao. 1987). The effect of temperature on the liberation of tetraspores in the agar yielding seaweeds Gelidium pusillum (Stackhouse) Lat regarding the influence of environmental factors Jolis, Pterocladia heteroplatos (Boergesen) on spore liberation and its diurnal rhythm Umamaheswara Rao and Kaliaperumal and (Katada, 1955; Umamaheswara Rao and Gelidiopsis variabilis (Graville) Schmitz had

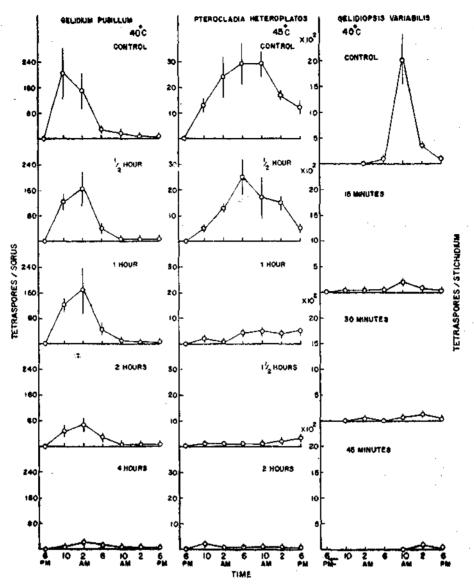


Fig. 1. Changes in diurnal periodicity of tetraspore output from fronds of G. pusillum and P. heteroplatos treated for short periods at 0°C and G. variabilis at 0 and 20°C (Vertical lines show standard errer to the mean values).

been studied (Umamaheswara Rao and Kaliaperumal, 1983; Kaliaperumal and Umamaheswara Rao, 1987). The influence of low and high temperatures on diurnal periodicity of tetraspore output from these three red algae growing at Visakhapatnam Coast are presented in this note. The author expresses his thanks to the Head of the Department of Botany, Andhra University, Waltair for the laboratory facilities. His thanks also are due to Prof. M. Umamaheswara Rao, Dept. of Botany, Andhra University, Waltair for suggesting this problem, encouragement and going through

the manuscript. He is grateful to Dr. V. A. Parastem n for financial assistance and also to CSIR, New Delhi for the award of Senior Research Fellowship.

### Materials and methods

Plants of G. pusillum, P. heteroplatos and G. variabilis were collected in March-May and September-October 1978 from the intertidal region of Visakhapatnam Coast during spring tides in the afternoon. As described by Umamaheswara Rao and Kaliaperumal (1983) fertilte terasporic thalli were selected and used for spore liberation experiments. The experiments were conducted in the laboratory by keeping the petri dishes with materials in a temperature controlled dark incubator and deep freezer. Since there was no sporulation from tetrasporic fronds of G. pusillum and P. heteroplatos and stichidia of G. variabilis treated for 24 hr at low and high temperature, experiments were conducted by treating the plants at these temperature for short periods. The un reated sets kept for 24 hr at room temperature in a dark chamber were treated as controls. G. pusillum was treated for 1, 2, 3 and 4 hr at 0°C and 1, 1, 2 and 4 hr at 40°C, P. heteroplatos was treated for 4, 8, 12 and 16 hr at 0°C and for 1, 1, 11 and 2 hr at 45°C. G. variabilis was treated for 1, 1, 11 and 2 hr at 0°C; 4, 8, 12 and 16 hr at 20°C and 15, 30 and 45 minutes at 40°C.

After the temperature treatment all experimental sets were maintained at room temperature in the dark chamber for remaining period of the day. The experiments were commenced from 1800 hr and the spores liberated into the petri dishes at 4 hr intervals were counted following the method of Umancheswara Rao and Kaliaperumal (1983). Mean values of 10 experiments conducted for each temperature treatment using G. pusillum and P. heteroplatos and 5 experiments with G. variabilis are plotted in Figs. I and 2 and the data are presented as tetraspores/sorus and tetraspores/stichidium.

Results

Data collected on shedding of tetraspores from tetrasporic fronds of G. pusillum and P. heteroplatos tres.ed for short periods at 0°C and 20°C are given in Fig. 1. Data obtained in control experiments are all plotted for comparison. In the experiments conducted at 0°C with G. pusillum during September-October, peak output was observed between 6 PM and 10 PM in control and thalli treated for 1 hr. The peak shedding was delayed by 4 hr i.e. from 10 PM to 2 AM in fronds treated for 2 he and 3 hr. The spore output markedly decreased at different periods of the day in fronds treated for 4 hr with maximum outputbetween 2 AM and 6 AM. In P. heteroplato more number of spores were found between 2 AM and 6 AM in control and thalli treated for 4 hr at 0°C. Four hour delay in the maximum shedding was observed in fronds treated for 8 hr. The spore output was very less in 12 hr and 16 hr treatments and maximum liberation was found from 2 PM to 6 PM.

In experiments conducted during September. October with G. variabilis at 0°C, the diurnal periodicity for 1 hr and 1 hr treatment were similar to those of controls with peak libertion between 6 AM and 10 AM. At 11 hr and 2 hr treatments, the peak output of spores was delayed by 4 hr from 10 AM to 2 PM. In experiments conducted at 20°C during the month of April, there were no changes in the diurnal periodicity between control and sitchidia treated for 4 hr, with peak shedding of spores from 6 AM to 10 AM. But in sitchidia treated for 8 hr, peak spore output was delayed for 4 hr from 10 AM to 2 PM. At 12 hr and 16 hr treatmen. 8 hr delay in the peak liberation was seen with maximum sporulation between 2 PM and 6 PM.

Figure 2 shows the results obtained on diurnal periodicity in tetraspore liberation from G. pusillum and G. variabilis treated at 40°C and P. heteroplatos treated at 45°C for short

periods. The data collected from the control experiments are also plotted. In the experiments conducted with G. pusillum, during March-May, at 40°C peak spore output was observed between 6 PM and 10 PM in control and between 10 PM and 2 AM with 4 hr delay

in ½ hr to 4 hr treatments. In P. heteroplatos maximum sporulation was found from 2 AM to 10 AM in control experiments. Peak output was seen between 2 AM and 6 AM in fronds treated for 2 hr at 45°C. The spore output values were very low and irregular in fronds

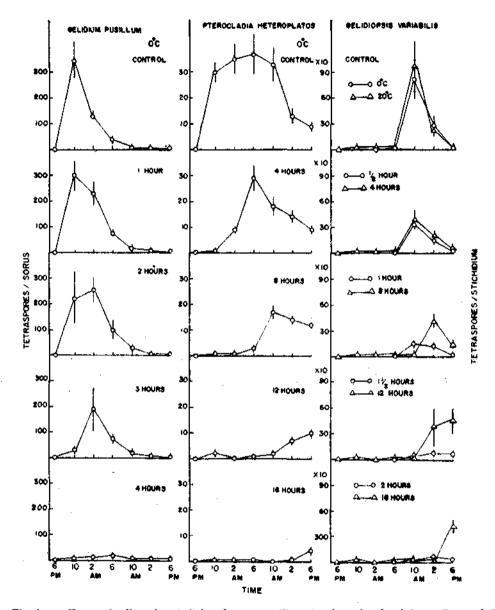


Fig. 2. Changes in diurnal periodicity of tetraspore liberation from fronds of G, pusillum and G, variabilis treated for short periods at 40°C and P, heteroplatos at 45°C (Vertical lines show standard error to the mean values).

treated for 1, 1 and 2 hr. Experiments conducted with G. variabilis at 40°C during September-October, peak spore output was observed between 6 AM and 10 AM in control and in 15 minutes treatment. The spores liberated at 15 minutes treatment were only 10% of the control experiments. The number of spores liberated was also less in 30 and 40 minutes treatments with 4 hr delay in peak liberation from 10 AM to 2 PM.

#### Discussion

Regular diurnal periodicity in tetraspore output with peak liberation during night time in G. pusillum and during day time in G. variabilis was observed. The pattern of the diurnal curves changed seasonally in these two red algae with 4 hr delay in peak shedding of spores during winter months from December to February/March. But in P. heteriplates, there was no regular trend in the diurnal periodicity of tetraspore release and there was no seasonal variation in the pattern of diurnal curves (Umamaheswara Rao and Kaliaperumal, 1987).

The low and high temperature altered the peak shedding of spores in G. pusillum, P. heteroplatos and G. variabilis (Figs. 1 and 2) while in Iridophycus cornucoptae water temperature had no effect on the diurnal periodicity

of spore liberation (Fukuhara, 1957). Peak liberation of tetraspores was delayed in G. pusillum and G. variabilis when sea water temperature was below 30°C. In fronds of Gracilaria corticata and Hypnea valentiae treated for short periods at 10°C, peak shedding of tetraspores was delayed for 4 to 8 hr (Subbarangaiah, 1985). Similarly in the present study also peak shedding of tetraspores was delayed in G. pusillum and G. variabilis treated for short periods at low temperatures of 0°C and 20°C.

In fronds of G. corticata and H. valentiae treated for short periods at 40°C, peak output of tetraspores was observed 4 hr earlier than the normal periodicity (Subbarangaiah, 1985). But in the present investigation, peak shedding of tetraspores was delayed in G. pusillum and G. variabilis treated for short periods at high temperature of 40°C. More detailed studies are needed to understand this variation observed in the daily periodicity of spore liberation at high temperatures in these algae. In Gelidium amansti the time of peak shedding of spores varied depending on the seasonal changes in sea water temperature (Katada et al., 1953; Katada, 1955). The present study agrees with the findings on G. amansii and it also confirms the relationship between diurnal rhythm in spore release and sea water temperature.

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