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STUDIES ON THE EFFECTS OF SOME AGENTS ON FERTILIZED SEA URCHIN EGGS, AS A PART OF THE BASES FOR MARINE POLLUTION BIOASSAY I¹⁾

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Introduction

The author examined the inhibitory degree of the polluted sea water in the Inland Sea of Japan upon the fertilization and further development of the sea urchin eggs (Kobayashi, 1972), after the manual of marine pollution bioassay proposed by him in 1971 (Kobayashi, 1971).

To supplement the previous two works on the one hand and to offer some more solid base for future works on the other hand, a few series of control experiments were made on four factors for marine pollution, which have not yet been dealt with in previous papers; namely, tannic acid which can be the most significant pollutant derived from timber-reservoirs or lumber industry, the pH showing a side of acid or alkaline contamination, Kaolin which may be taken to show the effect of muddy water caused by reclamation or shed from denuded lands, and the thermal pollution which is met with generally around thermal power stations.

Material and Methods

Eggs of *Anthocardis crassispina* (A. Agassiz) were experimented with in each test water to check the rate of fertilization, first cleavage, gastrulation and some anomalies in the development. Eggs were obtained by the current KCl-method, being washed several times with fresh sea water, and used as soon as possible, within 1 hour at the latest. Sperms were obtained from testes within 1 hour after being taken out of the test. The standard sperm density for insemination was maintained at about 1 dry sperm: 1,000 sea water in volume. When it was necessary, the preliminary check of eggs was done in the control laboratory water to see if the fertilization membrane was elevated in 3 minutes after insemination in over 85% of eggs and if the well synchronized first cleavage occurred in over 80% of them. The eggs were inseminated in respective test water within 2-3 minutes as far as possible. Firstly the percent of eggs with the elevated fertilization membrane to the total eggs observed was read. The first cleavage occurred in most cases 45 to 60 minutes after the insemination at

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26°C or 28°C. Then, the rate and state of the first cleavage, namely proportions of undivided cells, normal two cells and multi-cells caused by polyspermy were checked at some adequate time. Two hundred eggs were fixed with 5% formaldehyde at a time for this examination. Lastly, the state of swimming embryos exclusive of those deposited on the bottom, namely proportions of permanent blastulae, normal gastrulae and abnormal exogastrulae, was checked 12 to 15 hours after the insemination. Two hundred embryos were fixed at a time for this check. The test was repeated over 3 times on different batches. The results representing typically are described. The experiments were done with tannic acid and kaolin on July 25, 1971 at about 26°C in water temperature and with pH and higher temperature on August 19, 1971 at 28°C in water temperature. The state of *Anthocidaris crassispina* eggs could not be exactly uniform throughout the period from July to August, but the effect of respective factors could never be influenced significantly by the difference in the state of eggs in this period, because the normal fertilization and development of these eggs were confirmed in advance by control experiments mentioned above.

Tannic acid and kaolin to be tested were in the preliminary experiments diluted each into a series of solution by successive dilution by ten times of the original solution (chemicals 1 : 10⁴ sea water). In these series was found first the gross concentration range of respective agents, in which some anomalies appeared on developing sea urchin eggs. Then further series of solutions by successive dilution by two times were made within the gross concentration range of respective agents to see their inhibitory effects in detail.

To see the effects of pH and higher water temperature a series of sea water of successive grades of these factors were prepared on the bases of their usual ranges in the control laboratory sea water. Table 2 show the results of experiments.

Results

1. Tannic acid and Kaolin (Table 1)

In Table 1, the concentrations of these agents are given in the column A by the dilution of the original solutions (Sea water for dilution/original solution, in volume), and the same is expressed in the column B by the absolute molecular concentration (exclusive of bound water). The maximal ineffectual and minimal effectual concentrations are given in ppm. At checking the fertilization membrane formation 3 minutes after the insemination, the formation of lower fertilization membrane was excluded. In some solutions, there were found some eggs in which no cleavage followed the formation of the fertilization membrane.

At higher concentrations of tannic acid, the fertilization never occurred, and all eggs remained as they were at the beginning of experiments. The lower the concentration was, the more the fertilization was recovered, till the following cleavages were regained to usual rates. The polyspermy did not appear. Permanent blastulae and some retardation in development occurred at higher concentrations, but not any exogastrulae.

In the original suspension of kaolin (1 : 10⁴; average size of kaolin particles about

Table 1. Effects of tannic acid and kaolin upon the fertilization and development of *Anthocardis crassispina* eggs. Date: July 25, 1971. Test water temperature: 26°C. Original solution, chemicals 1 : 10⁴ laboratory sea water. The column A of "concentrations" shows the dilution of the original solution by the volume in—original solution 1 : volume of sea water; B shows the same dilution in the absolute molecular concentration.

Chemicals	Concentrations		3 min. after insem.	60 min. after insemination			15 hrs. after insemination			Ultimate state of eggs
	A	B	Fertiliz. membra. form. %	1 cell state %	2 cell state %	Multi- cells: polysp. %	Perma- blast. %	Norm. gast. %	Exo- gast. %	
Control sea water			86	16	84	0	1	99	0	normal
Tannic acid $C_{76}H_{52}O_{46}$ min. eff. conc.: 6.3 ppm max. ineff. conc.: 3.1 ppm	4	16×10^{-6}	0	100						no fertilization development retarded development retarded normal
	8	8×10^{-6}	26	85	15	0	18	82	0	
	16	4×10^{-6}	53	49	51	0	1	99	0	
	32	2×10^{-6}	85	17	83	0	0	100	0	
Kaolin (acid wash.) $H_2Al_2Si_2O_8 \cdot H_2O$ min. eff. conc.: 500 ppm max. ineff. conc.: 250 ppm	0.1	4×10^{-3}	35	67	33	0	2	98	0	development retarded
	0.2	2×10^{-3}	48	55	45	0	1	99	0	development retarded
	0.4	10^{-3}	87	15	84	1	1	99	0	normal

1 μ in diameter), did not appear any inhibitions in the fertilization and subsequent development. Moreover, at concentrations higher than 1 : 10³ no observation was allowed to learn in what state the treated eggs were. As the turbidity was as high as 324 even at 1 : 10³. Then, the suspension of 1 : 10³ was diluted by successive dilution by two times. At the concentrations over 1 : 4 $\times$ 10³, rates of fertilization and subsequent cleavages were more or less reduced but the polyspermy did not appear. With the increase in concentration, later developmental stages were more or less retarded.

2. PH and higher temperature (Table 2)

The pH of the sea water was raised or lowered by addition of respectively caustic soda and hydrogen chloride, and checked by colorimetry. The pH of the control running sea water of the laboratory (7.8) was a little lower than that of the open sea water (8.2–8.3). With the decrease of pH, rates of fertilization and subsequent cleavages were reduced, and the treated eggs remained as they were at the beginning of experiments. At pH 7, the fertilization occurred on most eggs, but almost polyspermic, and the cleavage ceased at earlier states. In the range from 7.8 to 8.6, the fertilization, cleavages and further development appeared to be carried out normally. With the increase of pH over 8.6, rates of fertilization and subsequent cleavages decreased, and the cytolysis occurred at earlier stages.

The temperature experiments were made in two series of different temperature degrees; this was not intended but due to the thermostat used. With the raise of temperature, rates of fertilization and subsequent cleavages were reduced, while the formation of multi-cells by polyspermy was promoted and thus the cytolysis occurred

Table 2. Effects of pH and higher temperature upon the fertilization and development of *Anthocardis crassispina* eggs. Date: July 28 and August 19, 1971. Test water temperature: 28°C.

Agents	Grades	3 min. after insem.	45 min. after insemination			12 hrs. after insemination			Ultimate state of eggs
		Fertiliz. membra. form. %	1 cell state %	2 cell state %	Multi- cells: polysp. %	Perma. blast. %	Norm. gast. %	Exo- gast. %	
Control sea water	pH 7.8	98	3	97	0	1	99	0	normal
pH	5.8	7	100						almost no fertilization
	6.2	16	81	19	0				almost no fertilization
	6.6	82	15	79	6				remain before 2-4 cell-stage
	7.0	97	6	21	73				remain before 8 cell-stage
	7.4	98	3	85	12	19	80	1	almost normal
	7.8	98	3	96	1	3	97	0	normal
	8.2	99	1	98	1	1	99	0	normal
	8.6	99	2	97	1	3	97	0	normal
	9.0	97	3	95	2	43	57	0	development retarded
	9.4	83	20	79	1				cytolysis occurred before blastula
	9.8	52	69	31	0				cytolysis
	10.2	13	100						cytolysis
Control sea water	28°C	100	1	99	0	0	100	0	normal
High temperature 1.	30°C	97	4	93	3	11	88	1	almost normal
	32	92	26	36	38				cytolysis occurred before blastula
	34	84	39	7	54				cytolysis occurred before 4 cells
	36	10	92	2	6				cytolysis
Control sea water	28°C	99	2	98	0	0	100	0	normal
High temperature 2.	29°C	99	1	99	0	1	99	0	normal
	31	94	19	57	24	73	26	1	development retarded
	33	85	36	31	33				cytolysis occurred before 16 cell
	35	37	61	17	22				cytolysis

at earlier states. Another test indicated that the fertilization did not occur at higher temperature over 37°C.

Considerations

It is obvious that various pollutants affect differently at different developmental stages of sea urchin eggs at different concentrations (Kobayashi 1971).

The inhibitory effects of the turbid water along the northern coast of Tanabe

Bay, i.e. the water from Mori and Egawa Harbours, upon the fertilization and development of sea urchin eggs are seemingly attributable to tannic acid derived from the timber reservoir and deposited bark pieces in the sea areas. A similar phenomenon was suggested by Tabata (1956) who supposed that tannins was the main toxic components of extracts from timbers stored in the sea. The minimal concentration of tannic acid to inhibit the fertilization, cleavages and further development of sea urchin eggs was 6.6 ppm, this concentration coincides almost with that given in the paper of Okubo and Okubo (1962) reporting a bioassay for the sea water pollution using developing eggs and embryos of sea urchins and bivalves. Further, it must be noted that the toxicity of tannins is comparable to that of chrome and manganese and much stronger than that of waste oil.

The effects upon marine organisms of muddy water derived from various sources have not yet been studied fully, though the effects will differ according to the nature of substances, size of particles, and their density. Thus, the effect of kaolin, the main component of clayey soil, upon the sea urchin eggs was observed. The suspension of pure kaolin washed by acid and consisting of particles ca. $1\ \mu$ in diameter is neutral, non poisonous, in a state of hydrosol and adsorptive of some ions or large molecules in the sea water. At the concentrations over 250 ppm, the fertilization, cleavages and later development were more or less affected. Possibly the clayey muddy water, as well as the suspension of kaolin, may affect the respiration and surface charge etc. of eggs of marine animals or the condition of the sea water, or simply adhesion of particles on the egg or embryo surface may induce sinking of them down to the sea-floor that must be their grave.

The normal fertilization and cleavages are maintained in *Anthocardis crassispina* between pH 7.8 to 8.6, but the decrease of pH below 6.6 will reduce the rate in fertilization and subsequent cleavages. Polyspermy is induced at neutral pH. The increase of pH over 9.0 also reveals the reduction in rates of fertilization and cleavages, followed by cytolysis. It has been noted since many years ago that the sea water of lower or higher pH is unavailable to secure the normal fertilization and development of sea urchin eggs. It was observed with *Arbacia* by Smith and Clowes (1924) that the fertilization was blocked below pH 6.8 and above 10.0, normal cleavages (if fertilized in sea water) maintained at 5.8 to 8.2, but were retarded below 5.2 and above 9.4, the maximal polyspermy appeared near 7.2 (also by Clark 1936) and the greatest cytolysis occurred above 9.3. Throughout the results of the present experiments and those obtained by previous researchers, the normal fertilization and development of sea urchin eggs seem to be maintained in the range from pH 7.4 to 8.6. This range seemingly coincides fairly with the pH-range allowing the normal reproduction of marine fishes, etc.

With the raise of temperature, rates of fertilization and cleavages will be reduced, while polyspermy and cytolysis will be increased. There have been made many experiments with higher temperature to induce such anomalies in the fertilization and further development of sea urchin eggs as inhibition of cleavage (Hoadley and Brill 1937), polyspermy (O. and R. Hertwig 1887) and cytolysis (Dan 1936, etc.).

All these results seem to indicate that the temperature higher by over 3 degrees centigrade than the usual summer water temperature may affect considerably the reproduction of marine organisms.

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Summary

1. A few series of experiments were made to clear the effects of four factors for marine pollution, tannic acid, kaolin, pH and higher temperature, after the manual of bioassay using sea urchin eggs and embryos (Kobayashi, 1971).

2. The minimal concentration of tannic acid, the main toxic component of the extracts from timbers stored in the sea, to inhibit the fertilization, cleavage and later development of sea urchin eggs was 6.3 ppm.

3. Kaolin, the main component of clayey soil, at the concentrations over 250 ppm more or less affected the fertilization, cleavage and later development.

4. The normal fertilization and cleavages were maintained in *Anthocardis crassispina* between pH 7.8 to 8.6, the neutral pH remarkably induced polyspermy, the decrease of pH below 6.6 reduced rates of fertilization and cleavage, and the increase of pH above 9.0 also brought about the reduction in fertilization and cleavage rates followed by cytolysis.

5. The raise of temperature by more than 3 degrees Centigrade above the usual summer water temperature reduced rates of fertilization and cleavage, but promoted polyspermy and cytolysis.

REFERENCES

- Clark, J. M. 1936. An experimental study of polyspermy. Biol. Bull. 70: pp. 361-384.
- Dan, K. 1936. Electrokinetic studies of marine ova. III. The effect of dilution of sea water, and of sodium and calcium upon the surface potentials of *Arbacia* eggs. Physiol. Zool. 9: pp. 43-57.
- Hertwig, O. and R. 1887. Über den Befruchtungs- und Teilungs-vorgang des tierischen Eies unter dem Einfluss äusserer Agentien. Jena. Zeitsch. N.F. 13: pp. 120-241 and 477-510.
- Hoadley, L. and Brill, E. R. 1937. Temperature and the cleavage rate of *Arbacia* and *Chaetopterus*. Growth 1: pp. 234-244.
- Kobayashi, N. 1971. Fertilized sea urchin eggs as an indicatory material for marine pollution bioassay, preliminary experiments. Publ. Seto Mar. Biol. Lab. 18: pp. 379-406.
- Kobayashi, N., Nogami, H. and Doi, K. 1972. Marine pollution bioassay by using sea urchin eggs in the Inland Sea of Japan (the Seto-Naikai). Publ. Seto Mar. Biol. Lab. 19: pp. 359-381.
- Okubo, K. and Okubo, T. 1962. Study on the bioassay method for the evaluation of water pollution II. Use of the fertilized eggs of sea urchins and bivalves. Bull. Tokai Reg. Fish. Res. Lab. 32: pp. 131-140.
- Smith, H. W. and Clowes, G. H. A. 1924. The influence of hydrogen ion concentration on the fertilization process in *Arbacia*, *Asterias* and *Chaetopterus* eggs. Biol. Bull. 47: pp. 333-344.
- Tabata, K. 1965. Systematic studies on toxic components in industrial wastes with reference to the tolerance of aquatic lives-III. On acute toxic components in water-extract liquor from bark. Bull. Tokai Reg. Fish. Res. Lab. 42: pp. 17-21.