Use of the Crustacean Artemia franciscana for Alternative Biotests

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Abstract

Use of the crustacean Artemia franciscana for alternative biotests of the second generation was studied, and possible experimental design and applications of such tests outlined. In addition to the classical use in ecotoxicology, the test can be used in pharmacology as well, or to monitor the effects of ionizing radiation in co-exposure with some chemical compounds. The synergistic effect of co-exposure of PCB (DELOR 103), cadmium chloride and potassium dichromate with beta 89Sr irradiation was shown. We also demonstrated the anti-oxidative and pro-oxidative effects of the ascorbic acid in dependence on its concentration. Use of the pharmaco-toxicological screening in search for the novel inhibitors of cyclin-dependent kinases was demonstrated as well, showing that Artemia franciscana may be used as a suitable biosensor instead of the expensive tests on higher vertebrates.

Artemia salina, synergy effects, irradiation, pharmacotoxicology

An important difference between experiments on live animals and on their tissues or cells in culture lies in the complex response of a living organism to any agent in contrast to only partial response of their tissues or cells in culture to the same agent. Experiments on tissues or cells in culture are not studies on simplified organisms but only experiments on living components of organisms as a whole (Pazourek 1992). The environment including the aquatic ecosystems is often contaminated with various exogenous agents at low concentrations (Beňová et al. 2007; Nováková et al. 2007). Although the response of many test organisms to various agents is well-known, our knowledge of the toxicity of mixtures of such agents is rather limited. Exogenous agents at low concentrations may interact with various physical factors and/or pharmaceuticals. The effects of such co-exposure may not be revealed for some time, and there are good reasons to assume that they might be of rather chronic or subacute character (Dvořák and Beňová 2002; Beňová et al. 2006). In order to monitor the effects of such co-exposures and of mixtures of residues of various exogenous agents on the organism it is possible to extend the viability of test organisms under standard conditions (Dvořák et al. 2005).

The tests for monitoring co-exposures of low concentrations of exogenous agents should meet the following requirements: sufficient sensitivity to exogenous agents in short- or long-term tests, high homogeneity of test organisms and their low sensitivity to variable experimental conditions of tests, high reproducibility of experiments, they should represent as closed a system as possible (with minimum interactions with other environmental factors), high availability of the test organisms, and simple verification and statistical analysis of the results. But the above criteria can hardly be fulfilled by the conventional tests on vertebrates. Moreover, there are serious public demands to minimize the distress and suffering of experimental animals (Russell and Burch 1992). The European Convention for the Protection of Vertebrate Animals demands minimization of the number of experiments on laboratory animals. The Three R’s concept of Reduction, Refinement
and Replacement defined according to the Council of Europe has been accepted (Council of Europe 1976). These demands highly accentuate the urgency of development of the alternative biotests.

Subacute biotests of the second generation on *Artemia franciscana* (formerly *A. salina*) in various experimental designs fully match the above demands.

The aim of the study was to prove the usage of this biotest for different purposes: the co-exposure of some chemical agents combined with the exposure to ionizing radiation, the study of the anti-oxidative and pro-oxidative effects of the ascorbic acid (Vitamin C) in dependence on its concentration, and the pharmaco-toxicological screening for new cyclin-dependent kinase inhibitors (CDKI). For example, “Olomoucin” and “Roscovitin” used in this study belong to this very promising group of cytostatics with negligible side effects (Sklenář et al. 2006).

**Materials and Methods**

**General procedures**

The diapausing eggs of *A. franciscana* were purchased from the SANDERS Comp. (Utah, USA) under the name of “Maxima brine shrimp eggs”. The hatching was performed over a period of 24 h in continuously aerated salt water at 25 °C. The salt water contained the following chemicals of the analytical grade [g·L⁻¹]: 23.9 NaCl; 10.83 MgCl₂·6H₂O; 2.25 CaCl₂·6H₂O; 0.68 KCl; 9.06 Na₂SO₄·10 H₂O; 0.2 NaHCO₃; 0.04 SrCl₂·6H₂O; 0.099 KBr, and 0.027 H₃BO₃ (Dvořák 1995).

Each test included at least 50 specimens per each concentration of the agent tested. In a standard design 5 disposable Petri dishes (60 mm in diameter) were used. Ten specimens were plated into the dish in a total volume of 10 ml including the tested agents. The living nauplii were counted at 24-h intervals. The dead nauplii were not removed (the results remained unaffected by dead nauplii providing that the death rate in the control group had not exceeded 10%).

**Monitoring of co-exposures of low volume activity of ⁸⁹Sr and low cadmium, chromium and Delor 103 concentrations**

Delor 103 (4.5 ng·L⁻¹) was selected to represent PCBs. PCBs produce colloidal solutions. The initial Delor 103 colloidal solution in salt water was obtained by three-day agitation. Excessive undissolved Delor 103 was then removed. One ml of solution containing 45 ng Delor 103 in 1 litre of sea water was added to each dish. The concentration was determined by capillary gas column chromatography analysis, where 41 peaks were identified mainly as derivatives from di- to pentachlorides. The derivatives were identified by the Kovats index. The source of ionizing radiation was a beta emitter ⁸⁹Sr in a solution of 30 kBq·L⁻¹ volume activity. Strontium - isotope ⁸⁹Sr was supplied by the Amersham Comp. as strontium chloride dissolved in water. The manufacturer guarantees the specific ⁸⁹Sr activity of 1.85 - 7.4 GBq·g⁻¹ and purity < 0.5% ⁸⁴Sr < 0.1% ⁹⁰Sr, with pH within the 5.0 - 9.0 range. The resulting activity in dishes was obtained by recalculating the original activity to the reference date. The solution was then diluted with salt water to the activity ten times higher than that required in dishes.

Potassium dichromate K₂Cr₂O₇ of analytical grade was diluted in salt water. One ml of the solution at a concentration 10 times higher than that in the dish was added to each dish 0.05 g·L⁻¹.

In all experiments of this study, cadmium chloride dihydrate CdCl₂·2.5 H₂O (0.02 g·L⁻¹) of analytical grade (Lachema Brno) was used. Salt water was used in both the initial solution and subsequent dilutions. One ml of solution at a concentration 10 times higher than that used in a dish was added to each dish.

A standard design of the biotest was used (see General procedures).

**Anti-oxidative and pro-oxidative effects of the ascorbic acid**

The oxidizing agent was hydrogen peroxide at 0.4 g·L⁻¹ concentration. The anti-oxidative substance was the ascorbic acid at 0.3 or 0.1 g·L⁻¹ concentrations. The test had a standard layout described in General procedures.

**Comparison of the toxicities of cyclin-dependent kinase inhibitors (Olomoucin and Roskovitin) and the toxicity of risk elements**

Because of poor solubility of the agents, the biotest was performed in salt water at a total salt concentration of 0.9%. Solubility of Roskovitin in water was very poor. Consequently this agent was dissolved in a mixture of dimethyl sulphoxide (DMSO, 10 g·L⁻¹) and Tween 80 (TW, 5 g·L⁻¹). The toxicity of the solvents at the above concentrations was tested.

**Statistical methods**

Groups of 50 specimens were used in tests of each of the concentrations of the agents studied. The only indicator monitored was the lethality of experimental specimens. All values that failed to meet the requirements of Dean-Dixon test (Q test) at the significance level of α = 0.05 were excluded (compare Dvořák 1999).

The calculated and test values were plotted in 3D diagrams that expressed the dependence of *Artemia salina* death rate on the concentration of the tested agents and on time.
Differences between different concentrations, types of toxic agents and their combinations were tested using the method described by Wayland and Hayes (1991). To identify significant relationships between different sets of data in a specific experiment or between different experiments, test of independence in contingency tables was used (Anděl 1985).

Results
Monitoring of co-exposures of low volume activity of $^{89}$Sr and low cadmium, chromium and Delor 103 concentrations
None of the agents (Delor 103 (4.5 ng·l$^{-1}$, lethality 8%), strontium 89 (30 kBq·l$^{-1}$, lethality 10%), cadmium chloride (0.02 g·l$^{-1}$, lethality 12%) caused any significant toxic effect except for potassium dichromate (0.05 g·l$^{-1}$) whose lethality was 27%. However, the co-exposure of all the agents lead to the high lethality of 81% (Fig. 1).

![Fig. 1. Lethality of *A. franciscana* during co-exposure of $^{89}$strontium 30 kBq·l$^{-1}$, PCB - Delor 103 4.5 ng·l$^{-1}$, cadmium chloride (CdCl$_2$·2.5 H$_2$O) 0.02 g·l$^{-1}$, and potassium dichromate 0.05 g·l$^{-1}$.

Antioxidative and pro-oxidative effects of various concentrations of the ascorbic acid
The effects of co-exposure of hydrogen peroxide 0.4 g·l$^{-1}$ and the ascorbic acid are shown in Fig. 2. Contrary to the effects of these agents used separately, the chart shows a significant ($\alpha = 0.05$) pro-oxidative effect of the ascorbic acid at the concentration of 0.3 g·l$^{-1}$ used in combination with hydrogen peroxide at the concentration of 0.4 g·l$^{-1}$. On the other hand, co-exposure of hydrogen peroxide and the ascorbic acid at concentrations 0.4 g·l$^{-1}$ and 0.1 g·l$^{-1}$ respectively, revealed the anti-oxidative action of the ascorbic acid by a decrease in lethality ($\alpha = 0.05$). The quenching effect of the ascorbic acid at 0.1 g·l$^{-1}$ persisted throughout the experiment (24-120 h). It became most distinctive after a 96-h exposure when lethality had dropped by 34% compared to the effects of hydroxide peroxide alone.

Comparison of the toxicities of cyclin-dependent kinase inhibitors (Olomoucin and Roskovicin) and the toxicity of risk elements
Fig. 3 shows a comparison between Olomoucin and Roskovicin toxicities and the toxicities of risk elements, namely chromium, zinc and cadmium. All agents were used at the concentration of 0.1 g·l$^{-1}$. The test lasted 120 h at 0.9% salinity. Olomoucin dissolved in dimethyl sulphoxide (DMSO) showed higher toxicity than Olomoucin dissolved in a mixture of DMSO and Tween (TW). But only the results for 96 h exposure proved to be significant. Chromium in the form of potassium dichromate proved to be the most toxic risk element. Its toxicity was significantly higher than that of Olomoucin or Roscovitin (both dissolved in a mixture of DMSO+TW) in
all exposures. Cadmium had significantly higher toxicity than Olomoucin (at exposures higher than 72 h) or Roskovitin (at exposures higher than 96 h). Zinc had significantly higher toxicity than Olomoucin dissolved in DMSO+TW (at exposures higher than 96 h) or Roskovitin (at exposures higher than 120 h).

Discussion

The genus *Artemia* is a member of the order *Anostraca* representing a group of crustaceans of the class *Branchipoda*. The *Artemia* populations have been found in about 500 salt lakes and salt works in the temperate, subtropical and tropical climate zones. *Artemia* prefer the salinity of 47 g·l\(^{-1}\) but they can survive at salinity of up to 250 g·l\(^{-1}\) as well (Ruppert et al. 2003).
Test validation criteria depend on the lethality of control groups, which should not exceed 10% in a 96 h test and 20% at exposures exceeding 120 h (Dvořák et al. 2009). Since diapausing eggs of *A. franciscana* are commercially canned for aquaristic purposes, we can very easily and cheaply obtain more or less homogenous populations consisting of millions of specimens (Dvořák et al. 2009).

As $^{89}$Sr is a beta radiation emitter, we cannot evaluate the real dose received by specimens during an experiment. Consequently, the effects of strontium in this study are given in volume activity units of kBq·l$^{-1}$.

Hexavalent chromium damages DNA, decreases its synthesis and enhances oxidative processes in cells. Similar effects are produced by ionizing radiation. The effects of strontium 89 or Delor 103 were not significantly different from those of the control group. On the other hand, co-exposure of those agents together with potassium dichromate led to a significantly different value than that of the control group. We assume that the effects of both agents, i.e. Delor 103 and strontium 89 were due to the damage of membrane integrity resulting in higher penetration of both potassium dichromate and cadmium chloride. Cadmium, on the other hand, induces production of metallothioneins (Kovářová et al. 2009), i.e. substances that exhibit radioprotective effects (Beňová et al. 2006). A biotest using *A. franciscana* to monitor the cumulative effects of Cd, Cu, Pb and Zn in sediments has been published (Fichet et al. 1998).

Our experiments have demonstrated the possibilities of use of the presented biotest as a sensor for the co-exposure of a number of agents at various concentrations, as well as of ionizing irradiation.

The search for scavengers of reactive forms of oxygen (free radicals) is one of the priority fields of modern pharmacology. They include a number of agents and their effects depend largely on their concentrations. In our experiment, co-exposure of hydrogen peroxide and ascorbic acid at concentrations of 0.4 g·l$^{-1}$ and 0.1 g·l$^{-1}$, respectively, reduced the toxicity to a greater extent than hydrogen peroxide at 0.4 g·l$^{-1}$ per se. This fact confirms the generally well-known anti-oxidative efficiency of the ascorbate (Young et al. 1992). Lethality, however, was lower than that of the control group. Consequently, one can assume that ascorbic acid is able to partially eliminate the effects of hydrogen peroxide, at least in this experimental design. The increase of concentration of the ascorbic acid had led to higher elimination of the effects of hydrogen peroxide. On the other hand, lethality was increased already at the concentration of 0.3 g·l$^{-1}$. Results of our test are in agreement with the clinically well-known studies showing that ascorbic acid exposures higher than 1000 mg per day have pro-oxidative effects whereas lower concentrations act as scavengers of the reactive forms of oxygen. *A. franciscana* therefore proved its utilization as a convenient biosensor for pharmacological studies of relations between exposure and its biological effects. The experimental design described in these experiments has recently been used in studies with some other prospective agents including plant extracts but the results have not been published yet.

One of the most promising groups of antitumour chemotherapeutics is the cyclin-dependent kinase inhibitors (CDKI) (Benson et al. 2005). The first synthetic purine inhibitor of CDKI was Olomoucin (Hajduch et al. 1999; Kryštof et al. 2002). The development of new pharmaceuticals has inevitably involved studies of their toxicity. This fact offers exceptional opportunity for the second generation biotest that may replace some experiments on vertebrates.

Due to the low solubility of CDKI in salt water, we decreased the concentrations of salt water to 0.9% in experiments with those agents. This caused a reduction in the ionic strength of the solution and an increase in their solubility. However, in such experiments it was impossible to extend the tests to ten days by the addition of glucose (Dvořák et al. 2005). Dimethyl sulfoxide (DMSO) proved as a very useful solvent in these experiments. It has
excellent dissolving capacity for the agents tested and low toxicity for *A. franciscana*. The concentration of DMSO used for the dissolution of purine CDKI was 15 g·l⁻¹. This approach is not suitable in all applications (e.g. in case of Roskovitin). Other co-solvents may include nonionic tensides, such as Polysorbat 80 (TWEEN 80), and poloxamers (Pluronic F-68). To solve hardly soluble agents, combinations of DMSO (10 g·l⁻¹) and TWEEN 80 or Pluronik at concentrations of 5 g·l⁻¹ are mostly used (Sklenář et al. 2006). When DMSO was used in combination with another co-solvent, only 10 g·l⁻¹ concentration was applied.

The toxicity of Olomoucin dissolved in DMSO + TWEEN was significantly lower than that of the combination Olomoucin + DMSO without TWEEN. This effect was probably caused by the prolonged life span of *A. franciscana* in salt water containing TWEEN (Fig. 3).

Our biotests with *A. franciscana* have demonstrated considerable possibilities for the use of these organisms as biosensors. This experimental system was successfully proved in experiments with more than 10,000 specimens. It is obvious that the conventional tests on vertebrates cannot compete with these tests in terms of quantities. These tests offer new possibilities for toxicity testing of various environmental agents and their combinations at low concentrations. The biotests can be used in pharma-toxicological studies of newly synthesized CDKI as well. Of course, such alternative biotests cannot fully replace the conventional tests on laboratory vertebrates, but they have the potential to markedly reduce the number and extent of such experiments.

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