Resistance to metal contamination by historically-stressed populations of *Ceriodaphnia pulchella*: Environmental influence versus genetic determination

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Received 8 June 2004; received in revised form 25 January 2005; accepted 22 February 2005
Available online 19 April 2005

Abstract

Field populations of daphnids historically-stressed by metal contamination may show increased resistance to those contaminants. This study was undertaken aiming to confirm/infirm three main hypotheses: (1) field populations living in historically-impacted environments are more tolerant to metal stress than populations from reference sites; (2) resistance differences are genetically-determined, i.e., differences persist after controlling for environmental and maternal effects, by acclimating cloned lineages to similar conditions; and (3) resistance to stress in field populations living in historically-impacted environments is due to the disappearance of sensitive individuals rather than the appearance of highly resistant ones, i.e., the shift in the central tendency of resistance is linked to a decrease in the range of population resistance and not to an increased upper limit of the population resistance. Three populations of the cladoceran *Ceriodaphnia pulchella* Sars in Southern Portugal were sampled; one of which has been historically-stressed by acid mine drainage (AMD) from an abandoned cupric-pyrite mine and two from reference sites within the same watershed. To assess if resistance differences were genetically-determined, the three populations were acclimated for at least five generations under the same controlled conditions. Assays with AMD contaminated water samples were performed with both non-acclimated and acclimated individuals from all studied populations. Reproduction results in sub-lethal assays revealed significant differences between the reference and stressed populations. Significant differences in resistance to lethal levels of toxicity were observed for both non-acclimated and acclimated populations, individuals from population I being more resistant than those from reference populations. The existence of genetically-determined sensitivity differences was attested by the presence of significant differences in resistance to lethal levels of toxicity in acclimated individuals from reference and stressed populations. Results from cumulative mortality assays revealed that sensitive individuals were most probably present in the original population, but no conclusion could be draw about the presence of extreme resistant individuals in the historically-stressed population. Finally, it was shown that responses among populations converged from high to low levels of contamination.

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0045-6535/S - see front matter © 2005 Elsevier Ltd. All rights reserved.
doi:10.1016/j.chemosphere.2005.02.072
1. Introduction

The toxicity of heavy metals to aquatic organisms is well documented in terms of both lethal (Mance, 1987; Wiederholm and Dave, 1989; Gerhardt, 1995; Jak et al., 1996; Villaescusa et al., 1997) and sub-lethal effects (Mance, 1987; Hickey, 1989; Gerhardt, 1995; Miliou et al., 1998). Heavy metals can stress organisms by different mechanisms affecting the cell membrane (e.g., ion transport mechanisms), the DNA (e.g., inhibiting the transcription process), by causing oxidative damage (Stohs and Bagchi, 1995), and by causing damage to the nervous system and to the function of enzymes (Dedegle et al., 1994; Gerhardt, 1995). Several metal detoxification mechanisms are found in individuals, for example, the production of metallothioneins, which captures and retain metal ions (Hamer, 1986; Roesijadi, 1992) and the accumulation and retention (e.g., by mineralisation) of heavy metals in specific regions and organs (Borgmann et al., 1993; Cope et al., 1994; Lucan-Bouché et al., 1999; Munger et al., 1999). The diversity in these physiological responses allows a population living in an impacted environment, i.e., under a strong selection pressure, to acquire resistance to metals (Brown, 1978; Klers and Weis, 1987; Bodar et al., 1990; Posthuma et al., 1992; Stuhlbacher and Maltby, 1992; Donker et al., 1993; Shirley and Sibly, 1999). Tolerance to heavy metals can be acquired either by environmentally-induced physiological alterations, including those due to stress-induced gene expression (Stuhlbacher and Maltby, 1992; Stuhlbacher et al., 1993; Maltby and Crane, 1994; Lam, 1996, 1999) or by quantitative and/or qualitative changes in gene frequencies (Klers and Weis, 1987; Gill et al., 1989; Posthuma et al., 1992; Lam, 1999). Some complications may arise when trying to differentiate these hypotheses. For example, techniques that control for environmental influence involve screening tolerance after acclimation to controlled conditions (Maltby et al., 1987; Lam, 1999). One of the major problems is to which extent maternal influence persist in the F1, F2 and, even later generations. This problem can be diminished by culturing animals under laboratory conditions and screen tolerance to stress in F1 or later generation progeny.

The acquisition of resistance to toxicants may lead to a decrease in the genetic diversity of the population (Van Straaalen and Timmermans, 2002). Actually, there is some support for the genetic erosion hypothesis, more specifically at the molecular basis. Krane et al. (1999) analysed RAP DNA polymerase chain reaction-generated DNA from eight different population of Orconectes rusticus and found that changes in genetic diversity was significantly correlated with the extent to which the population had been exposed to contaminants. Also, Murdoch and Hebert (1994) observed that diversity of mitochondrial DNA of brown bullhead fish was lower in populations from contaminated rivers relatively to populations from relative pristine sites.

To study the resistance of field populations under exposure to long-term metal stress, three field populations of the lake-dwelling cladocerans Ceriodaphnia pulchella Sars were selected to carry out this study. Daphnids are particularly suitable to investigate genetically-determined resistance to contamination since they exhibit facultative asexual reproduction by ameiotic parthenogenesis. Furthermore, data acquisition on cladocerans’ resistance is particularly relevant since these individuals are widely used, as standard test species (particularly, Daphnia magna and Ceriodaphnia dubia) in hazard assessment studies. Surprisingly, scarce information related to resistance acquisition by field population of Cladocera is available from published literature (revision made by Reznick and Ghalambor, 2001).

In this study, two experimental approaches were undertaken aiming to test three questions: (1) are field populations living in historically-impacted environments more tolerant to metal stress than populations from reference sites? (Experiment 1); (2) are these differences genetically-determined, i.e., do differences persist after controlling environmental and maternal effects, by acclimating individual lineages to similar conditions for several generations? (Experiment 2); (3) is the resistance to metal stress observed in field populations living in historically-impacted habitats due to the disappearance of sensitive individuals or to the appearance of highly resistant individuals, i.e., is the shift in the central tendency of resistance linked to a decrease in the population resistance range of responses rather than to an increased upper limit of the population resistance? (Experiment 2). To investigate metal resistance in this species, field populations were sampled in a well studied, isolated aquatic system impacted by acid mine drainage (AMD), where other industrial, agricultural or urban runoff sources are absent.

2. Materials and methods

2.1. Study site

An aquatic system impacted with AMD, from an abandoned cupric-pyrite mine (active from 1859 to
was selected for the development of this study (Fig. 1). The oxidation of the pyrite environment produces an effluent with high concentrations of heavy metals and very low pH (Kelly, 1988; Sengupta, 1993). The ongoing oxidation of abandoned mine tailings continuously leads to the production of a very acid effluent, highly contaminated with heavy metals (Fe, Al, Zn, Cu, Mn, Co, Ni, Cd, Pb, Cr, As; in decreasing order, Lopes et al., 1999a; Pereira et al., 2000) which is discharged without a treatment into the Chancer Reservoir located in the Guadiana River Basin (SE Portugal). The pH of this reservoir is close to neutrality since the low pH of the effluent is neutralised by dilution. For the water samples used in bioassays (V1–V3 and T1), data on 12 metals, measured by atomic absorption spectroscopy, from previously collected samples with almost matching conductivity values are presented in Table 1. Further metal analyses were considered unnecessary since no concentration-effect relationships were intended to be established, but solely the comparison of relative sensitivities between populations.

This aquatic system is highly suited for the study of adaptations occurring in natural populations exposed to historical chemical stress, for three reasons: (i) the source of contamination is isolated and identified (pH and, mainly, heavy metals); no agriculture or industrial activities are present and there is no urban runoff; (ii) reference sites free of metal mining contamination exist close to the contaminated area; (iii) extensive data on the biology, ecology and water chemistry of the system are available (Pereira et al., 1995; Ribeiro et al., 1995; Lopes et al., 1999a,b; Pereira et al., 1999, 2000; Castro et al., 2003, 2004; Moreira-Santos et al., 2004a,b).

2.2. Assay organisms and culture conditions

Two reference populations (R and RR) and one historically-stressed (I) population of Ceriodaphnia pulchella Sars were collected from two lakes (R—pH 7.5, 189 μS/cm, DO 8.7 mg/l; and RR—pH 7.61, 193 μS/cm, DO 9.3 mg/l) located upstream the contaminated area, and one site located near the confluence of the AMD effluent with the Chancer Reservoir (I—pH 7.0, 231 μS/cm, DO 9.4 mg/l), respectively (Fig. 1).

The procedure to sample field populations was as described by Lopes et al. (1999a). Individuals were brought to the laboratory, and, using a dissection microscope, 15 egg-bearing females from each population were transferred to culture beakers filled with 50-μm filtered site water. Randomly sampled juveniles born the next day from those field-collected females were kept in 50-μm filtered site water until used to perform toxicity assays, the remaining juveniles were reared in the laboratory and acclimated to controlled conditions (for at least five generations), which were as follows: 25 ± 1°C with a 14:10 h L:D cycle in American Society for Testing and Materials (ASTM) hardwater (ASTM, 2002), with added vitamins and the organic additive Marinure 25 (Baird et al., 1989); animals were fed daily with the green

| Table 1 |
|---|---|
| Conductivity | 332–280 | 202 |
| Previous conductivity | 312 | 271 |
| Previous alkalinity | 6.5 | 27.0 |
| Al | 11.3 | <0.1 |
| As | <0.005 | <0.005 |
| Cd | 0.0286 | 0.00082 |
| Cr | <0.015 | <0.015 |
| Cu | 1.7 | 0.027 |
| Fe | 1.64 | 0.129 |
| Mn | 3.07 | 0.081 |
| Ni | 68.0 | <0.03 |
| Pb | 4.60 | 0.640 |
| Hg | <0.0002 | <0.0002 |
| Co | 90.0 | <0.03 |
| Zn | 12.10 | 0.288 |
algae *Pseudokirchneriella subcapitata* (Koršhikov Hindak (formerly known as *Selenastrum capricornutum* Printz) \(3 \times 10^5\) cells/ml/d).

2.3. Toxicity assays

Lethal and sub-lethal toxicity assays were performed with neonates from the three *C. pulchella* populations under controlled conditions of temperature (25 ± 1 °C) and photoperiod (14:10 h L:D cycle). Assays were conducted with laboratory-acclimated neonates (6–24-h old), from the third and fourth broods, and with non-acclimated neonates (6–24-h old) from recently field-collected egg-bearing females (USEPA, 1991). In lethal assays no food was added, while in sub-lethal assays individuals were fed daily with *P. subcapitata* \(3 \times 10^5\) cells/ml/day, which corresponds to \(1.2 \times 10^7\) cells/individual, with water being renewed every other day.

Conductivity (WTW, LF92, Weilheim, Germany), pH (WTW, 537) and dissolved oxygen (DO) (WTW, OXI 92) were monitored at each sampling site and at the beginning and at the end of each assay.

Two experiments were carried out:

2.3.1. Experiment 1—Sensitivity of reference versus stressed populations

This experiment investigated differences in sensitivity to AMD contamination between the reference and the historically-stressed populations. Lethal (survival time—death within minutes to hours) and sub-lethal (reproduction) toxicity assays were carried out with non-acclimated neonates exposed to two Very Toxic (samples V1 and V2) and to one Moderately Toxic waters (sample T1). In survival time assays (neonates from the populations R and I were exposed to water samples V1 and V2), five individuals were introduced, one by one, in 42-ml glass vessels filled with 15 ml of water sample, with four replicates per treatment (20 individuals/treatment). Immobilisation (no movement after gentle prodding) was checked every 3 min from 15 to 30 min, every 5 min from 30 to 120 min, every 15 min from 2 to 6 h, every hour from 6 to 12 h (Lopes et al., 1999b, 2000; Ribeiro et al., 2000). In reproduction assays (neonates from the populations R, RR and I were exposed to the water sample T1), each individual was introduced into 42-ml glass vessels filled with 15 ml of water sample, with four replicates per treatment. Reproduction was assessed daily by counting released neonates. Reproduction assays ended when all daphnids released the fourth brood.

2.3.2. Experiment 2—Environmental versus genetic components of resistance

This experiment investigated if differences in resistance to AMD contamination between reference and historically-stressed populations were due to environmental-induced physiological alterations or to local adaptation, and/or if those differences were due to the disappearance of most sensitive genotypes. Lethal (cumulative mortality—death within hours to days—with the Very Toxic water V3) assays were conducted with acclimated neonates from the three populations. In cumulative mortality assays, five individuals were introduced simultaneously in 42-ml glass vessels filled with 30 ml of water sample, with four replicates per treatment. Immobilisation was checked every 24 h. An assay ended when all the daphnids died or after 240 h of exposure if at least one individual survived that long.

2.4. Data analysis

Survival time and reproduction data were analysed with ANOVAs followed by post-hoc comparisons with the Tukey HSD test, and with *t*-tests when comparing only two samples (Zar, 1996). Survivorship curves with censored data from cumulative mortality assays were compared with the Gehan–Wilcoxon \(\chi^2\)-test (Pyke and Thompson, 1986). All these statistics, were performed using the program Statistica for Windows 4.3 (StatSoft, Aurora, CO, USA). The ET50 (median lethal time), with the respective 95% CI and slopes, was computed using PriProbit 1.63 (Sakuma, 1998; http://bru.gmprc.ksu.edu/proj/priprobit/download.asp).

3. Results

Conductivity, pH and DO of water samples at the time of collection are presented in Table 2. No significant variations were observed in these parameters either between collection of the water and its use in the assays. Differences never exceeded 0.1 pH units, 3 μS/cm conductivity or 0.2 mg/l DO in any experiment.

3.1. Experiment 1—Sensitivity of reference versus stressed populations

In the first experiment, non-acclimated neonates, from recently field-collected egg-bearing females, were exposed to two Very Toxic (V1 and V2) and one Moderately Toxic (T1) water samples, aiming to detect

<table>
<thead>
<tr>
<th>Table 2</th>
<th>Values of pH, conductivity (Cond., μS/cm), and dissolved oxygen (DO, mg/l) measured in the water samples used in toxicity assays</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>V1</td>
</tr>
<tr>
<td>pH</td>
<td>3.5</td>
</tr>
<tr>
<td>Cond.</td>
<td>332</td>
</tr>
<tr>
<td>DO</td>
<td>9.7</td>
</tr>
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differences, at lethal and sub-lethal levels, in sensitivity between the reference and the historically-stressed local populations. Lethal assays revealed that the population I was significantly more resistant to AMD contamination than the population R, when exposed to water samples V1 and V2 (\(t\)-tests: \(t_{38} = 3.48, p < 0.001\) and \(t_{38} = 4.67, p < 10^{-5}\), respectively for V1 and V2) (Fig. 2). The ET50s (CI) for population I and R were 1.62 h (1.50–1.72) and 3.71 (3.42–4.02), when exposed to water sample V1, and, 1.31 h (1.24–1.38) and 1.97 h (1.83–2.12), when exposed to V2. The slope of the survival curves was smaller in population I than in population R, both when exposed to V1 or V2 water samples (\(b = 9.2, 5.29; b = 13.6, 5.38\), respectively).

When exposed to the water sample V1, individuals from population R started to die 0.30 h earlier than those from the population I. During this 0.30-h period population R presented already 15% mortality (Fig. 2). Furthermore, 30% of population I was still alive when the last individual from the population R died, and it took 1.10 h for those 30% to perish (Fig. 2). Similar results were obtained with the sample V2: 40% of population R died before any mortality in population I being observed, which started to occur only 1.17 h later (Fig. 2). Furthermore, by the time 100% of mortality was observed in population R, population I still presented 55% of survivals and it resisted for an extra 4.60-h period (Fig. 2).

When exposed to the water sample T1, individuals from the population I presented a significantly higher reproductive output than populations R and RR (1-way ANOVA: \(F_{2,33} = 4.68, p = 0.016\)) (Fig. 3).

3.2. Experiment 2—Environmental versus genetic components of resistance

In the second experiment, all three populations (R, RR, and I) were fully acclimated to controlled conditions and exposed to the water sample V3. This phase tended to assess if differences in sensitivity between populations were due to environmental-induced physiological alterations or to local adaptation.

Individuals from the reference populations exposed to water sample V3 showed significantly reduced survival over those from the historically-stressed population (Gehan–Wilcoxon test: \(\chi^2 = 5.90, p = 0.05\)) (Fig. 4). After 240 h of exposure mortalities of 100%, 83% and 67% were registered for the populations R, RR and I, respectively. Furthermore, after 24 h of exposure only 17% of mortality was observed in the population I, while 100% and 75% were already attained by the populations R and RR, respectively (Fig. 4).

4. Discussion

4.1. Sensitivity of reference versus stressed populations

The first question asked in this study was if there were differences in sensitivity to AMD between reference and historically-stressed populations. The results
obtained in the first experiment, by exposing non-acclimated individuals to two Very Toxic (V1 and V2) and one Moderately Toxic (T1) water samples, showed that individuals from population I presented a higher tolerance than those from uncontaminated site populations, both at lethal and sub-lethal levels of contamination (Figs. 2 and 3). These results were expected, since similar tolerance to heavy metals has been reported in literature for a wide range of organisms; bacteria (Bruins et al., 2000), protists (Devars et al., 1998), plants (Macnair, 1997; Monni et al., 2000), and animals (Klerks and Weis, 1987; Posthuma and van Straalen, 1993). Furthermore, several authors have shown that tolerance to heavy metals can be induced in laboratory after a short pre-exposure period to metals (Bodar et al., 1990; Münzinger, 1990; Stuhlbacher and Malby, 1992; Stuhlbacher et al., 1993). Reinecke et al. (1999) pre-exposed the terrestrial oligochaete Eisenia fetida for more than ten generations to sub-lethal concentrations of cadmium sulphate, and found that pre-exposed individuals were more tolerant to high cadmium concentrations than unexposed ones. This higher tolerance, acquired by the pre-exposed individuals, was due to environment-induced physiological alterations (e.g., through the increased synthesis of binding proteins).

Results from the cumulative mortality assays with populations R and I exposed to waters samples V1 (pH 3.5, 332 μS/cm) and V2 (pH 3.8, 293 μS/cm) indicated that (Fig. 2): (i) the most sensitive individuals (the first ones to die) were most probably only present in the reference population, since 15% of individuals died before any of the individuals from population I perished, and (ii) the most tolerant individuals (the latter ones to die) were most probably only present in the historically-stressed population, since, when the last individuals from population R died, 30% (V1) and 55% (V2) of individuals from the population I were still alive. Since these individuals were neonates from recently field-collected egg-bearing females, which had been maintained in local water, differences in the extremes of lethal tolerance could be exclusively due to previous exposure of females from the population I (Klerks and Weis, 1987; Lam, 1999).

The highest lethal tolerance to intense AMD contamination by the stressed population was associated to a higher sub-lethal tolerance to moderate AMD contamination. After exposing individuals from the three populations to the water sample T1 (Fig. 3), the population I showed a higher fitness (neonates/female) than the reference ones (R and RR).

4.2. Environmental versus genetic components of resistance

Besides environmental-induced physiological alterations (including those due to stress-induced gene expression), populations can evolve resistance by genetic adaptation (Klerks and Weis, 1987; Lam, 1999). This was the second goal of the present study: to assess whether the higher resistance shown by the historically-stressed population I was genetically-determined. The results obtained with acclimated individuals exposed to the water sample V3 (pH 6.0, 280 μS/cm) supported this hypothesis (Fig. 4): since significant differences (p = 0.05) between the historically-stressed and reference populations, were still evident after controlling for maternal and external environmental influences (Klerks and Weis, 1987; Lam, 1999). These results indicate that probably a directional selection occurred in population I, since a directional shift in survival curve took place towards an increase in survival in heavy metal contaminated environments. Such genetically-determined resistance acquisition was found for other organisms and toxicants (Klerks and Weis, 1987; Reznick and Ghulambor, 2001; Klerks, 2002). Posthuma et al. (1993) observed that Collombola populations living in heavy metal contaminated sites had an increased excretion efficiency of metals than reference populations and were more tolerant to this type of contamination. They suggested that this was related with a strong directional selection pressure, induced by cadmium, which led to the selection of traits linked with tolerance. The results obtained in this study do not allow concluding for sure if the genetically-determined higher resistance of the population I was due to the disappearance of sensitive individuals or to the appearance of highly resistant ones. Results obtained with non-acclimated individuals exposed to the water samples V1 and V2 showed that the most sensitive individuals were most probably absent in population I, since individuals in population R started to die much earlier. Furthermore, individuals from the population I died much later than the ones from population R, suggesting that most tolerant individuals only occurred in the historically-stressed population. When exposed to the water sample V3 the most sensitive individuals were present in all the populations, since after 24 h of exposure mortality occurred in all populations. On the contrary, the most tolerant individuals were only present in the populations RR and I, corresponding to the 17% and 23% of survivors, respectively (in population R all individuals died). This difference between results obtained with non-acclimated (samples V1 and V2) and acclimated individuals (V3) might indicate that the higher resistance of non-acclimated individuals from population I was due to both environmental and genetic components, while after acclimation to controlled conditions (the environment influence disappeared) individuals from population I become less resistant, and, thus the most sensitive among these individuals died at the same time as the ones from the populations R and RR. Furthermore, the frequency of the most sensitive individuals was much lower in the historically-
stressed population than in reference populations, since most of the individuals from I died much latter than most of those from the reference populations. No conclusion could be drawn about the presence of individuals with an extreme genetically-determined resistance in the historically-stressed population. In addition, it was reported in literature (Forbes and Depledge, 1996) that impacted populations exhibiting higher EC50 and lower slopes of survival curves indicates intermediate selection, since sensitive individuals are still present in the population. This is in agreement with the results obtained with acclimated organisms, thus involving genetic differences, which suggests the presence of sensitive individuals both in reference and impacted populations.

In conclusion, a strong genetically-determined increase in resistance in population I was demonstrated in the present study, and was probably due to genetic erosion: the loss of genetic variability through the disappearance of sensitive individuals present in the original founder population.

Although being beyond this study aims, results here obtained also suggested that resistance differences between the reference and the stressed populations increased with increasing toxicity, as indicated by individuals exposed to the contaminated samples V1 and V2 ($p < 0.001$) and to T1 ($p = 0.016$). Furthermore, a smaller difference was found in the assay with acclimated individuals exposed to the water sample V3 (null hypothesis rejected at $p = 0.05$) than the difference found with non-acclimated individuals exposed to samples V1 and V2 ($p < 0.001$). Therefore, at lethal levels of toxicity, results here obtained suggest a convergence in resistance responses between the reference and the stressed populations from high to low levels of contamination: V1 and V2 to V3. Furthermore, a convergence is also present between lethal and sub-lethal levels of toxicity (V2 and V1 to T1, and V3 to T1). Although it is not common to find works where responses to different lethal levels of toxicity are compared, some authors compared responses at different toxicity levels and found a convergence in responses from high to low levels of contamination. Barata et al. (2000) compared toxicity in resistant and sensitive clones of *D. magna* exposed to lethal and sub-lethal levels of cadmium, and found that genetic variance in resistance to cadmium converge from lethal to sub-lethal responses and, also, from high to low levels of sub-lethal effects. Furthermore, Leppänen et al. (1998) compared survival and feeding activity (egestion rates) between natural populations of the midge *Chironomus riparius* Meigen originated from reference and heavy metal impacted sites, and found that larvae from contaminated sites exhibited a higher lethal resistance to contaminated sediments than larvae from reference sites, though differences were not found at sub-lethal levels of contamination.

Acknowledgement

This work was partially funded by Fundação para a Ciência e a Tecnologia (Portugal)—PRAXIS XXI (the PIN project, contract POCTI/CED/34891/99).

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