Element analysis of the eutardigrades *Richtersius coronifer* and *Milnesium cf. asiaticum* using particle induced X-ray emission (PIXE)

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**ABSTRACT**

Semi-terrestrial tardigrades are well-known for their tolerance to a variety of environmental extremes, including desiccation, freezing, and radiation. Despite several attempts to reveal the genetic and molecular mechanisms behind the resilience of tardigrades, it is still unknown how these animals are able to maintain the integrity of their cellular components under severe stress. Quantitative or qualitative changes in molecular compounds (e.g., carbohydrates, proteins) are expected, and have been the main line of research towards understanding the tolerance of tardigrades. In radiation tolerant bacteria, a tolerance mechanism based on manganese has been proposed. We evaluate this hypothesis in tardigrades and provide the first data on element composition in desiccated and non-desiccated specimens of two eutardigrade species, *Richtersius coronifer* and *Milnesium cf. asiaticum*. A focused 2 MeV proton microbeam was utilised to determine the elemental content, distributions and concentrations, using the ion beam analytical technique particle induced X-ray emission (PIXE). The presence of six elements – phosphorus, sulphur, chlorine, potassium, calcium and iron – were confirmed in all tardigrade specimens, at levels up to a few mg g⁻¹. However, manganese was found in less than 10% of the analysed specimens, and in low amounts, thus our study provides no evidence for the manganese hypothesis. We also show that the distributions and/or concentrations of some elements differ between the two species as well as between the dehydrated and hydrated state. In particular, very low levels of iron were found in dehydrated *M. cf. asiaticum*. Our analysis shows that the PIXE technique is a useful tool for investigating questions on the distribution of elements both in dehydrated and hydrated tardigrades.

Key words: tardigrades, *Richtersius coronifer*, *Milnesium cf. asiaticum*, PIXE, radiation tolerance.

**INTRODUCTION**

Tardigrades inhabiting semi-terrestrial habitats are well-known for their tolerance to extreme environmental conditions such as desiccation, freezing, and radiation (Wright et al., 1992; Jönsson, 2003; Møbjerg et al., 2011). These abilities are shared with some species from a few other animal groups such as arthropods, nematodes and rotifers, and represent adaptations (or by-products of adaptations) to survive under very dry or cold conditions. For other organisms, exposure to these environmental stresses induces lethal damage to cells, and tardigrades must therefore possess a way to either protect their cells from damage, or efficiently repair damage that arises. Most of the early work on mechanisms of desiccation tolerance was directed towards studies on membrane stabilization molecules (Crowe et al., 1992; Crowe, 2002). More recent research has had a general molecular approach on detecting differences between animals in the desiccated (anhydrobiotic) and the hydrated states, in order to reveal the molecular and physiological mechanisms behind the tolerance (Mali et al., 2010; Reuner et al., 2010; Rizzo et al., 2010; Welnicz et al., 2011). Although these efforts have certainly improved our knowledge on the tolerance to environmental extremes in tardigrades, we are still far from understanding the systems of protection and actual components involved. Documented tolerance to ionizing and UV radiation not only in desiccated tardigrades but also in active hydrated animals (May et al., 1964; Jönsson et al., 2005; Horikawa et al., 2006; Altiero et al., 2011) have raised the question if DNA repair mechanisms may play a role in their tolerance to radiation (Jönsson et al., 2005; Jönsson, 2007). Few studies on DNA damage and repair in tardigrades have appeared, but Neumann et al. (2009) and Rebecchi et al. (2009) reported evidence of DNA degradation after exposure to periods of anhydrobiosis, with an increase in damage with time spent in the dry state. Evidence that DNA repair mechanisms are an important part of the tolerance to desiccation and radiation in anhydrobiotic animals also comes from studies on insect larvae (Gusev et al., 2010) and rotifers (Gladysh and Meselson, 2008).

In a study on *Deinococcus radiodurans* (Brooks and Murray, 1981), the most radiation tolerant organism on Earth, Daly et al. (2004) reported much higher intracellular concentration of manganese in radiation tolerant strains compared to sensitive strains. The authors suggested that radiation tolerance may rely on mechanisms...
that prevent or inactivate the damaging reactive oxygen species (ROS) that arise when radiation hits the water molecules, and that manganese performs such a role to protect the cell from ROS after irradiation. Later studies showed that also another radiation tolerant bacterium (Deinococcus geothermalis) has high manganese/iron (Mn/Fe) accumulation (Daly et al., 2007), and that not only the levels but also the distributions of Mn and Fe in these two bacteria are essentially the same (Makarova et al., 2004). In these analyses, and also in a study of elemental composition of the gram-negative bacterium Pseudomonas fluorescens (Flügge) Migula, 1895 (Kemner et al., 2004), elements were determined using X-ray fluorescence analysis (XRF).

From these patterns of Mn and Mn/Fe in bacteria, the hypothesis was suggested that survival following irradiation by ionizing radiation is governed by the amount of oxidative protein damage rather than by the amount of DNA damage (Daly, 2009). According to this hypothesis, high intracellular Mn:Fe ratios enable redox cycling, such that scavenging of O$_2^–$ (superoxide) is facilitated without producing HO (hydroxyl radicals) in the process. This combination of more manganese and less iron would result in less protein oxidation overall. Both D. radiodurans and D. geothermalis have Mn:Fe ratios only slightly below 1 and survive radiation doses up to 12 kGy, as opposed to e.g. the bacterium Shewanella oneidensis, which has a Mn:Fe ratio of only 0.0005 and survive radiation doses of only 0.07 kGy (Daly, 2009). Thus, the key to extreme radiation tolerance could lie in simple molecular components such as manganese that protect proteins involved in repair of DNA damage, rather than in the evolution of new specific proteins.

Since tardigrades belong to the most radiation tolerant metazoa, not very far behind the bacteria in tolerated doses, it is of interest to evaluate the manganese-hypothesis of Daly (2009) in these animals. Very little is however known about the composition of chemical elements in tardigrades, or whether the element composition changes between the different states of hydration. Also in other radiation tolerant metazoa, we have found no studies on manganese levels. The purpose of this paper is therefore to evaluate if manganese-levels or distribution differ between tardigrades in the anhydrobiotic state and those in a hydrated state. The study also provides the first information on a number of other elements. In addition, we wanted to investigate possible effects of different preparation methods (freezing) of hydrated samples, as well as differences between animals with a full and an empty stomach. While the above studies in prokaryotes used X-ray fluorescence to analyse elements, we have employed the ion beam analytical technique of Particle Induced X-ray Emission, PIXE. To our knowledge, this is the first analysis of tardigrades using PIXE analysis.

**METHODS**

**Specimen collection**

We used the eutardigrade species Richtersius coronifer Richters, 1903 and Milnesium cf. asiaticum in our experiment, the former of which is one of the most investigated tardigrade species with respect to tolerance to environmental stress. The Milnesium population has preliminarily been identified as Milnesium asiaticum Tumanov, 2006, using the diagnostic key by Michalczuk et al. (2012a, 2012b), but until statistically satisfying morphometric data are available we prefer to classify it as M. cf. asiaticum. Both species were obtained from moss collected on carbonite rock fences at Öland, South-Eastern Sweden, 1-3 weeks before preparation of the experimental samples, and stored dry at room temperature. The moss was rehydrated and tardigrades extracted overnight in Baermann funnels with tap water (4-4.5 °dH). Extracted tardigrades were kept in refrigerator (4-5°C) for 12-48 h until prepared for the experiment.

**Sample preparation**

Three types of samples with regard to treatments were used: desiccated animals, hydrated quick-frozen animals, and hydrated slow-frozen animals. The frozen samples represented active animals, and the two categories were used in order to evaluate if the rate of freezing affected the results. All samples were placed and analysed in groups of five (but individually separated) on Kimfol™ foil stretched over an acrylic glass frame before treatment (desiccation or freezing). Excess water was removed before treatment. The desiccated animal samples were prepared by placing them in a desiccator at 94.5% relative humidity and 20°C for 2-3 days over a saturated potassium nitrate (KNO$_3$) salt solution. Relative humidity was monitored by a hair hygrometer (Lambrecht GmbH, Göttingen, Germany; accuracy 2.5% RH), inside the desiccator. In order to see if animals with food content differed in their element content from animals without food, two sets of samples with desiccated tardigrades were prepared. Hydrated slow-frozen samples were prepared by placing them in a deep freezer at -20°C for approximately 3 h, then transferred (without thawing) to an aluminium block pre-cooled in liquid nitrogen and slowly evacuated overnight (approximately 24 h) in a vacuum chamber. Hydrated quick-frozen samples were frozen in liquid nitrogen (-196°C, for half a minute), then kept at -20°C for approximately 3 h, before drying in a vacuum chamber as described above. Once the treatment process (desiccation or freezing) was finished, another layer of Kimfol was attached to the frame over the animals, to form a Kimfol sandwich.

After treatment and sealing of the Kimfol sandwich, all sandwiches were carbon coated, in order to minimise
charging-up of the sample during the PIXE analysis. All in all, a total of 55 tardigrades were analysed, divided into 7 experimental groups (based on species and state), with a varying number of specimens in each group (Tab. 1).

More specimens were in fact prepared – however, despite the careful carbon coating of the Kimfol sandwiches, some prepared samples still suffered from charging-up, resulting in detachment from the backing of the specimen when the proton beam approached. In particular the hydrated, quick-frozen samples suffered from detachment from the backing, leading to uneven sample sizes in different groups.

**Elemental analysis**

Particle induced X-ray emission (PIXE) is an ion beam analytical method, developed in the 1970s at Lund University (Johansson et al., 1970). The method employs MeV (megaelectronvolt) energy ions, produced in an accelerator, which are used to bombard the sample to be analysed. As the energetic ions hit the atoms of the sample, the ions knock out electrons in the sample atoms and create vacancies in inner electron shells. As these vacancies are filled by electrons from outer shells, the excess energy is emitted as characteristic X-rays, unique to the elements present in the sample. Elements heavier than sodium can be detected using this method (Johansson and Campbell, 1988), but in this study the focus was on heavier elements, *e.g.* Mn, at the expense of the lighter ones, *i.e.* in practice the lower limit was aluminum. By scanning the ion beam across the sample element maps showing the distribution of the elements in the sample can be acquired. The technique is qualitative as well as quantitative, provided that the sample matrix and the beam charge deposited is well-known. Quantitative results can also be obtained by analyzing known standards. Quantitative PIXE analysis is based on the ratio of the measured amount of X-rays to the number of irradiating protons giving rise to the corresponding characteristic X-ray peak. Sensitivity factors like cross sections for interactions, yields, absorption *etc.* are well-known and available in the software used for spectrum evaluation. For a thick sample, the quantitative result will be in units of mass concentration (mg g⁻¹ or µg g⁻¹) in the volume covered by the irradiating beam. The entire experimental set-

### Tab. 1. Concentrations (dry weight), including standard error of the mean, of six different elements – phosphorus, sulphur, chlorine, potassium, calcium and iron – in each of the seven experimental groups of tardigrades.

|                  | P (mg g⁻¹) | S (mg g⁻¹) | Cl (mg g⁻¹) | K (mg g⁻¹) | Ca (mg g⁻¹) | Fe (mg g⁻¹) |
|------------------|------------|------------|-------------|------------|-------------|-------------|
| *R. coronifer*   |            |            |             |            |             |             |
| Dehydrated       |            |            |             |            |             |             |
| Full stomach     | 2.0±0.8    | 3.4±1.3    | 0.1±0.1     | 1.4±0.4    | 1.9±0.5     | 0.1±0.1     |
| *Dehydrated*     |            |            |             |            |             |             |
| Empty stomach    | 2.6±1.2    | 4.0±1.6    | 0.3±0.2     | 1.6±0.3    | 2.7±1.0     | 0.2±0.1     |
| *Hydrated*       | 1.9±0.3    | 2.0±0.3    | 0.6±0.2     | 1.2±0.2    | 2.0±0.4     | 0.3±0.05    |
| *Hydrated*       | 2.8±0.6    | 3.6±0.8    | 0.4±0.2     | 1.5±0.3    | 2.1±0.9     | 0.5±0.2     |
| *Hydrated*       | 1.6±0.4    | 2.1±0.3    | 0.2±0.1     | 0.7±0.4    | 1.9±1.0     | 0.04±0.02   |
| *Hydrated*       | 2.4±0.7    | 2.1±0.5    | 0.6±0.2     | 1.2±0.4    | 4.7±1.1     | 0.2±0.1     |
| *Hydrated*       | 1.6±0.7    | 1.8±0.7    | 0.3±0.1     | 0.9±0.4    | 2.6±1.2     | 0.2±0.1     |
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up is calibrated on a regular basis by analysing commercially available element standards that cover the periodic table. The procedure, used for a similar set-up, is found in Shariff et al. (2002).

Particle induced X-ray emission analysis was carried out on dehydrated as well as hydrated, frozen tardigrades of *R. coronifer* and *M. cf. asiaticum*. The primary scope of this study was qualitative rather than quantitative analysis, as the main focus was determination of any redistribution of elements between the hydrated and the dehydrated state, as well as differences between species. Also, the stomach content of some *R. coronifer* samples could be analysed, by comparing samples with a full stomach to samples with an empty stomach. In addition to the primarily qualitative investigation of differences between hydrated and dehydrated specimens of the same species, possible species-related differences between the two states in *R. coronifer* and *M. cf. asiaticum* were also investigated using quantitative PIXE analysis.

The PIXE analysis was carried out using 2 MeV protons and the resulting data was analysed using the GeoPIXE software (GeoPIXE II software; http://nmp.csiro.au/GeoPIXE.html). A focused beam of approximately 15 µm size was scanned over the sample, providing two-dimensional maps showing element distributions across the sample. Samples were analysed using a proton current of about 250 pA and the analysis time per sample was typically around 40 minutes. The detector used was an 8 element Canberra HPGe detector (Shariff et al., 2004), equipped with a 25 µm thick Mylar absorber. To achieve data for quantitative analysis, beam current and analysis time was monitored and beam charge was measured in a Faraday cup before the sample. In addition, a Ti standard (Micromatter™, Vancouver, Canada) was also analysed, to be able to normalize against this standard in GeoPIXE. The sample matrix, needed for the X-ray yield calculation and absorption correction, was chosen as chitin, C₈H₁₃O₅N, with a density of 1 g cm⁻³. From the elemental map, a region of interest is chosen, which completely covers the animal. The data is merged into a spectrum which represents the whole animal. When quantifying a thin sample the results represent mass per unit area, and for a thick sample the results represent ppm or µg g⁻¹. For additional information, see Johansson and Campbell (1988).

For the results reported here, the values represent averages of elemental concentrations taken from the whole tardigrade and referring to dry weight. As seen in Figs. 1-4, there are strong local variations of elemental concentrations; the signals shown here were normalised for each element and given as relative values.

![Fig. 1. Particle induced X-ray emission images showing the distribution of phosphorus, sulphur, chlorine, potassium, calcium and iron in desiccated *R. coronifer*, with an empty stomach.](image_url)
Statistical analysis

Differences in the element concentration between i) the two species, ii) hydrated and dehydrated specimens of the same species, iii) quick- and slow-frozen hydrated specimens of the same species, and finally iv) dehydrated *R. coronifer* with full and empty stomach were tested using Analysis of Variance, one-way ANOVA (SYSTAT 12, SYSTAT Software Inc.), with P>0.05 considered non-significant and P<0.05 as significant. The Shapiro-Wilk test was used to test the data for normality distribution.

RESULTS

Qualitative elemental analysis of tardigrades

Figs. 1-4 show some representative elemental maps of a dehydrated and a hydrated, quick-frozen tardigrade of each of the two analysed species. All maps are 384 µm×384 µm in size. Significant levels of phosphorus, sulphur, potassium and calcium are present in both states and in both species. The levels of iron and chlorine are low, but the elements are found in all samples, irrespective of species and state. In contrast, the distribution of chlorine is different between the two states. In the hydrated animals, chlorine is to a higher extent found along the outer edge of the body, whereas in the dehydrated animals, chlorine is more evenly distributed in the body. Iron also appears to be differently distributed between the states, as there appears to be cavities in the dehydrated tardigrades, where this element is not found. A similar cavity pattern is seen in the calcium maps and to a certain extent in the phosphorus maps. Moreover it can be seen that, at least in the *M. cf. asiaticum* specimens, the elements calcium and phosphorus overlap nicely, as do sulphur and potassium and chlorine as well (Figs. 3 and 4).

The presence of manganese in these tardigrade specimens could only be confirmed in five out of 55 samples (both *R. coronifer* and *M. cf. asiaticum* were represented) and in all five cases, the levels were very low – 0.01-0.02 mg g⁻¹. In the rest of the samples, the manganese level was below the detection limit (0.01 µg g⁻¹), which means that the presence of lower levels of manganese in tardigrades cannot be excluded. Thus, no elemental maps of manganese are presented in this paper. Calcium is known to be a component of the buccal-pharyngeal structures of tardigrades, and particularly of the stylet (Bird and McClure, 1997; Guidetti et al., 2012). This was also confirmed in our analyses, where the stylet was the most clearly visible structure of the buccal-pharyngeal apparatus. Also in line with the study by Guidetti et al. (2012), the presence of calcium was much more prominent in the stylets of *R. coronifer* compared to *M. cf. asiaticum* with strong signals (as in Fig. 2) in about 55% of the *R. coro-
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nifer specimens. Only about 25% of the Milnesium specimens showed clear signs of the stylets, and then mainly in the piercing part of the stylet, as also shown by Guidetti et al. (2012). Why the stylet was not clearly visible in calcium maps of all R. coronifer specimens is unclear, but in some cases the animal may have been in the simplex stage connected with moulting, during which the stylet and buccal tube is absent and the animal is not feeding. The presence of simplex stages was not checked in the used specimens, but R. coronifer in the empty stomach category had a considerably lower percentage of visible stylets (23%) than the average, which indicates that many of those specimens may have been in simplex stage. In a few specimens of both R. coronifer and Milnesium calcium was also indicated in assumed areas of the claws.

Quantitative elemental analysis of tardigrades

Tab. 1 gives the mean concentration (in mg g⁻¹) and standard error of the mean for the six elements phosphorous, sulphur, chlorine, potassium, calcium and iron. One should keep in mind that these values describe the average content across the whole animal, not taking into account the fact that the concentration varies in the body of the tardigrade, as can be seen from the figures in the previous section. Roughly half (25) of the samples were found to contain very low, but measureable levels of copper (mean value=0.08 mg g⁻¹) and all but nine of the samples contained zinc (mean value=0.2 mg g⁻¹) as well. Comparing all seven experimental groups, we found significant differences among groups for sulfur, chlorine, potassium, calcium, iron and phosphorus (S, Cl, K, Ca, Fe and P, respectively) (P=0.000 in all cases except for element P, where P=0.022). Significant and marginally significant results from pairwise comparisons are shown in Tab. 2.

Slow- versus quick-frozen hydrated samples

Comparing slow- and quick-frozen R. coronifer, we found significant differences for elements S, Cl and Fe, with higher levels of S and Fe in the slow-frozen group, and higher levels of Cl in the quick-frozen group. In M. cf. asiaticum an opposite but non-significant pattern was observed for P and S, while levels of Cl and Ca were significantly higher in the quick-frozen group. These results indicate that the method of freezing influences the measurement of some elements. Whether these differences represent adaptive physiological adjustments taking place in the body of the slow-frozen tardigrades, or is due to some physical reason related to the freezing rate is unclear, but in our comparisons between hydrated and dehydrated tardigrades we have analysed slow-frozen and quick-frozen tardigrades separately.

Fig. 3. Particle induced X-ray emission images showing the distribution of phosphorus, sulphur, chlorine, potassium, calcium and iron in desiccated M. cf. asiaticum.
Dehydrated versus frozen hydrated samples

Comparing dehydrated versus quick-frozen *M. cf. asiaticum* we found significant differences for Cl, Ca and Fe. In all cases, lower levels were found in the dry samples. For dehydrated versus slow-frozen *M. cf. asiaticum*, only Fe showed a significant difference between groups, with five times higher levels in the frozen samples.

In *R. coronifer*, significant differences between dehydrated (similar outcome independent on whether samples had stomach content or not) and quick-frozen samples were found for Cl and S, with dehydrated samples having higher levels of S and lower levels of Cl. Comparing dehydrated versus slow-frozen *R. coronifer*, only one difference, common to both full and empty stomach, was found—a significantly lower level of Fe in the dehydrated samples. In addition, a significantly lower level of Cl in the dehydrated samples with full stomach was also found.

Full stomach versus empty stomach in *R. coronifer*

Dehydrated *R. coronifer* with and without stomach content showed significant differences in element levels only for Cl and Ca (Tab. 2) and in both cases element levels were higher in the animals with empty stomach.

Richtersius coronifer versus *M. cf. asiaticum*

Including all dehydrated samples, we found significant differences in elements S and K levels between *R. coronifer* and *M. cf. asiaticum*. Dehydrated *M. cf. asiaticum* had in both cases lower levels than *R. coronifer*. A more pronounced difference was found between *R. coronifer* with empty stomach and *M. cf. asiaticum*, with significant differences also in P, Cl and Fe—in all cases higher levels in *R. coronifer*. For quick-frozen samples, a significant difference between *R. coronifer* and *M. cf. asiaticum* was found for Ca only, where *M. cf. asiaticum* had on average more than twice the levels of *R. coronifer*. For slow-frozen samples, however, there were more significant differences between the species in levels of P, S, K and Fe, and in all cases *R. coronifer* had higher levels than *M. cf. asiaticum*.

DISCUSSION

Our analysis did not provide any support for the manganese hypothesis proposed by Daly et al. (2004, 2007), a hypothesis which suggests that manganese plays a role in tolerance to desiccation and radiation by preventing the action of damaging reactive oxygen species. The two tardigrade species investigated had very low levels of manganese.
manganese, and the element was only found in a minority of specimens. The levels of iron were considerably higher, resulting in a Mn:Fe ratio well below 0.1. Thus, from this investigation it seems clear that manganese, or the Mn:Fe ratio, is not involved in the radiation and desiccation tolerance of tardigrades, as suggested for bacteria.

No studies on tolerance have been done in *M. cf. asiaticum*, but another species (reported as *Milnesium tardigradum* Doyère 1840 but probably also *M. asiaticum* or *M. cf. asiaticum*) has been reported to be more tolerant than *R. coronifer* to ultraviolet radiation, freezing, and high temperature (Ramløv and Westh, 1992, 2001; Jönsson et al., 2008; Hengherr et al., 2009a, 2009b). Also differences in biochemistry connected with desiccation have been documented in these two species (Westh and Ramløv, 1991; Hengherr et al., 2008; Jönsson and Persson, 2010). One of the differences in element content between *M. cf. asiaticum* and *R. coronifer* in our study was a clearly lower level of iron in dehydrated *M. cf. asiaticum*. This level contrasted also against the hydrated specimens of the same species, indicating a possible involvement of this element in the biochemical adjustment to the dry state. Also in *R. coronifer*, dehydrated specimens had lower levels of iron than hydrated specimens. Further studies are obviously of interest to reveal why both tardigrade species, and *M. cf. asiaticum* in particular, show such low levels of iron in the dehydrated state. For several other elements, *M. cf. asiaticum* had lower values than *R. coronifer*. The reason for this is hard to explain, but our results may be used to generate ideas and hypotheses about differences in physiology between the two species. The same holds for the qualitative results, where some differences in distribution were found, e.g., in the distribution of iron, chlorine and calcium.

The reason for the differences found between slow- and quick-frozen samples is currently unclear, but one possibility is that our slow-freezing method allowed the animals to prepare their bodies biochemically for a dry state. If so, they should be more similar in their element levels to the dehydrated samples than to the quick-frozen samples.

### Tab. 2. Statistical significances (P values) from pair-wise post-hoc ANOVA comparisons of element amounts (see Tab. 1) among experimental groups, using Fisher’s least-significant-difference test. Element name is given as subscript. Marginally significant results are given in parentheses.

|                | R. cor. | R. cor. | R. cor. | R. cor. | M. cf. asiaticum | M. cf. asiaticum | M. cf. asiaticum |
|----------------|---------|---------|---------|---------|-----------------|-----------------|-----------------|
|                | Dehyd. FS | Dehyd. ES | QF | SF | Dehyd. | | SF |
| R. coronifer   |         |         |         |         |                 |                 |                 |
| Dehydrated     |         |         |         |         | (P<0.055)      | (P<0.055)      | (P<0.055)      |
| FS             | P<0.004 |         | P<0.018 | P<0.006 | (P<0.051)      | P<0.014        | P<0.040         |
|                | (P<0.065) | (P<0.006) | (P<0.055) | (P<0.051) | (P<0.051)      | P<0.014        | P<0.040         |
| R. coronifer   |         |         |         |         |                 |                 |                 |
| Dehydrated     |         |         |         |         | (P<0.076)      | (P<0.010)      | (P<0.003)      |
| ES             | P<0.004 | P<0.001 | P<0.001 | P<0.037 | (P<0.076)      | (P<0.010)      | (P<0.003)      |
|                | (P<0.004) | (P<0.001) | (P<0.001) | (P<0.037) | (P<0.076)      | (P<0.010)      | (P<0.003)      |
| R. coronifer   |         |         |         |         |                 |                 |                 |
| QF             |         |         |         |         |                 |                 |                 |
|                | (P<0.059) | (P<0.010) | (P<0.003) | (P<0.002) | (P<0.059)      | (P<0.010)      | (P<0.003)      |
| R. coronifer   |         |         |         |         |                 |                 |                 |
| SF             |         |         |         |         |                 |                 |                 |
|                | P<0.009 | P<0.008 | P<0.026 | P<0.000 | P<0.009         | P<0.024        | P<0.001         |
|                | (P<0.053) | (P<0.026) | (P<0.000) | (P<0.000) | (P<0.053)      | (P<0.026)      | (P<0.000)      |
| M. cf. asiaticum |         |         |         |         |                 |                 |                 |
| Dehydrated     |         |         |         |         |                 |                 |                 |
|                | (P<0.055) | (P<0.000) | (P<0.000) | (P<0.024) | (P<0.055)      | (P<0.000)      | (P<0.024)      |
| M. cf. asiaticum |         |         |         |         |                 |                 |                 |
| QF             |         |         |         |         |                 |                 |                 |
|                | (P<0.002) | (P<0.000) | (P<0.000) | (P<0.024) | (P<0.002)      | (P<0.000)      | (P<0.024)      |
| M. cf. asiaticum |         |         |         |         |                 |                 |                 |
| SF             |         |         |         |         |                 |                 |                 |

R. cor., Richtersius coronifer; dehyd., dehydrated; FS, full stomach; ES, empty stomach; QF, quick-frozen; SF, slow-frozen.
hydrated samples. This tends to be the case, since in both tardigrade species we found more significant differences in element levels between dehydrated and quick-frozen samples (M. cf. asiaticum: Cl, Ca, Fe; R. coronifer: Cl, S) than between dehydrated and slow-frozen samples (both species: Fe). Thus, when freezing of tardigrades is used to get specimens representing hydrated states, utilization of rapid freezing media such as liquid nitrogen is essential. However, it should be remembered that in analyses of the distribution of elements in the body, and perhaps also in quantitative analyses, rapid freezing may give rise to structural changes or damage induced by the freezing treatment that lead to a different distribution of elements compared to specimens treated in other ways.

A challenge in analysing whole tardigrades is that we cannot distinguish at what depth the X-rays originate. We know, from the data analysis software GeoPIXE (GeoPIXE II), that for the lightest elements in our analysis, e.g. phosphorus, we get 90% of the information about the element content (90% of the X-ray yield) in the first 40 µm of the sample, whereas for the heavier elements of our analysis, e.g. potassium and iron, we get 90% of the information in the first 80 µm of the sample. Thus, the thickness of the tardigrades and the random orientation of them on the backing may influence the results more in the case of the lightest elements in our analysis. The maximum thickness of dehydrated tardigrades has been found to exceed 150 µm (Nilsson et al., 2010), but frozen tardigrades are thinner. Thus, especially in the case of the lightest elements, we have in our present analysis not been able to probe the center of the samples. We plan to overcome this in future studies by analyzing sectioned tardigrades. Analyzing thin slices (of the same thickness) will give us the opportunity to obtain an improved quantification as well as resolution, which in turn would better correlate the element maps to structures in the tardigrade. Another option would be a micro-tomographical analysis. To get an indication if sample orientation may have effects on estimated element levels, one of our tardigrade samples was analysed twice, turning the sample back to front between the two analyses. The results for our six main elements (data not shown) did not indicate any obvious differences in the quantification due to orientation of the specimen, but as this comparison was based on one specimen only we cannot draw any firm conclusions.

CONCLUSIONS

Despite the methodological issues discussed above, our elemental analysis shows that PIXE is a suitable method for determining the element content in both hydrated and dehydrated tardigrades, with high spatial resolution. Even better results should be acquired if sectioning the tardigrades before analysis. Thus, this method can be used to evaluate specific hypotheses about redistribution of elements in connection with dehydration, and the role of different elements in tolerance to dehydration and radiation.

ACKNOWLEDGMENTS

We are grateful to J. Genberg for valuable input regarding the statistical analysis, and to referees for valuable comments. Financial support from The Gyllenstierna Krapperup’s Foundation (to E.J.C. Nilsson), and the Swedish Space Agency (to K.I. Jönsson) is gratefully acknowledged.

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