Extreme Resistance of Bdelloid Rotifers to Ionizing Radiation

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Accessibility
Extreme resistance of bdelloid rotifers to ionizing radiation

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Contributed by Matthew Meselson, January 30, 2008 (sent for review November 9, 2007)

Rotifers of class Bdelloidea are common invertebrate animals with highly unusual characteristics, including apparently obligate asexuality, the ability to resume reproduction after desiccation at any life stage, and a paucity of transposable genetic elements of types not prone to horizontal transmission. We find that bdelloids are also extraordinarily resistant to ionizing radiation (IR). Reproduction of the bdelloids Adineta vaga and Philodina roseola is much more resistant to IR than that of Euchlanis dilatata, a rotifer belonging to the desiccation-intolerant and facultatively sexual class Monogononta, and all other animals for which we have found relevant data. By analogy with the desiccation- and radiation-resistant bacterium Deinococcus radiodurans, we suggest that the extraordinary radiation resistance of bdelloid rotifers is a consequence of their evolutionary adaptation to survive episodes of desiccation encountered in their characteristic habitats and that the damage incurred in such episodes includes DNA breakage that is repaired upon rehydration. Such breakage and repair may have maintained bdelloid chromosomes as colinear pairs and kept the load of transposable genetic elements low and may also have contributed to the success of bdelloid rotifers in avoiding the early extinction suffered by most asexuals.

Results

Reproduction. The reproduction of Adineta vaga and Philodina roseola was much more resistant to IR than that of Euchlanis dilatata. The relative parental fecundity of E. dilatata was reduced 10-fold by a dose of ~200 Gy, whereas a corresponding reduction in bdelloid fecundity required a dose five times greater. A similar difference between the monogonont and the bdelloids was seen in relative parental fertility. The reproductive effects of IR extended to the daughters of irradiated parents, with dose-dependent reductions in fecundity and fertility almost as great as seen for the parents. Even at the highest doses tested, irradiated rotifers remained active for the 1–2 weeks during which they were observed.

Table 1 presents data on parental and F1 reproduction of rotifers exposed to doses up to 280 Gy for the monogonont and 1,120 Gy for the bdelloids. For each dose group of 96 animals, including nonirradiated controls, Table 1 gives the number of animals that deposited eggs, the number that produced at least one daughter, the proportion of animals giving daughters normalized to the corresponding proportion for nonirradiated rotifers (relative parental fertility), and, for E. dilatata and P. roseola, the number of hatched eggs per daughter normalized to the number produced per daughter by daughters of nonirradiated rotifers (relative F1 fecundity). For E. dilatata, the number of hatched eggs was counted directly, whereas for the bdelloids it was taken as the number of progeny, as described in Materials and Methods.

The proportion of parental rotifers not depositing eggs averaged ~5% for bdelloids and 12% for E. dilatata, independent of dose. Such animals were apparently infertile or had reached the end of their reproductive lifespan, ~5 days for E. dilatata and 2 weeks for the bdelloids under our conditions. The total number of eggs deposited per individual averaged 12 and 19 for A. vaga and P. roseola, respectively, whereas the average for E. dilatata was ~3 in the first two experiments and 6 in the third, a variation possibly reflecting differences in the age of the parents.

Relative parental and F1 fecundity and fertility are plotted as a function of dose in Fig. 1. The two bdelloid species are similar in their dose-response relations, with an indication that reproduction of P. roseola may be a little more radiosensitive than that of A. vaga. Parental fecundity, which measures the total number of daughters produced by irradiated animals, is only moderately more radiosensitive than parental fertility, which requires only the production of at least one daughter. Bdelloid parental and F1 fertility and fecundity display an extended shoulder in their

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dependence on dose, whereas only a slight shoulder is found for monogononts. Beyond their respective shoulders, reproduction falls off rapidly, with no indication of a resistant fraction. The reproduction of daughters of irradiated rotifers is also radiosensitive, with a dose–response relation similar to that of their parents but shifted to somewhat higher doses. Again, *P. roseola* appears to be a little more radiosensitive than *A. vaga*.

**DNA Breakage.** Fig. 2 shows the gel electrophoretic distribution of DNA found after exposing *A. vaga* to doses in the range of 0 to 1,120 Gy. The average molecular sizes at 500, 840, and 1,120 Gy are 350, 250, and 190 kbp, respectively, with an average of 0.005 double-strand breaks (DSBs) Gy⁻¹. Some of the DSBs we observed may have been measured for many arthropod species at various life stages, mainly for purposes of pest control. In a tabulation of such sterilizing doses for male arthropods, which are generally more radiosensitive than females, including 285 species representing 61 families of insects and other arthropods, irradiated mainly as pupae or nymphs and generally achieving at least 90% parental and F₁ sterility, the average dose is ~110 Gy (ref. 15 and www-ididas.iaea.org/ididas/). For the most radio-resistant species in the tabulation, the lepidopteran Spilosoma (*Diacrisia*) obliqua, a dose of 200 Gy to pupal or adult males, the most resistant sex, resulted in >99% F₁ sterilization. Among other invertebrates, sensitivity to IR has been measured for X-irradiation of Caenorhabditis elegans eggs in buffer, for which ~30 Gy reduced the number developing into adults by 90%.

**Discussion**

Reproduction of Irradiated Rotifers. We find that reproduction of the bdelloid rotifers *A. vaga* and *P. roseola* is much more resistant to ¹³⁷Cs γ-irradiation than that of the monogonont rotifer *E. dilatata*. A dose of ~200 Gy reduced the number of daughters produced by the monogonont by ~90%, whereas a corresponding reduction in bdelloid fecundity required a dose five times higher. The principal difference between the bdelloid and monogonont dose–response relations lies in the extended shoulder over which bdelloid fertility and fecundity remain high before falling steeply, whereas only a much less extended shoulder is found for the monogonont. The sterilizing effect of IR extends to the daughters of irradiated parents, with dose–response relations shifted to somewhat higher doses. Taking the size of the *A. vaga* genome as ~180 Mbp (R. Gregory, personal communication), the average molecular size of 350 kbp observed after a dose of 560 Gy, causing a reduction in fecundity of only 20%, corresponds to ~500 DSBs per genome.

The reproduction of *A. vaga* and *P. roseola* is more resistant to IR than that of any other metazoan for which we have found relevant data. Such resistance to IR is likely to be characteristic of the Bdelloidea generally, as *A. vaga* and *P. roseola* represent families that diverged millions of years ago (14) and because of the probable association, discussed below, of bdelloid radiation resistance with anhydrobiosis. Doses of IR or high-energy electrons required to prevent reproduction have been measured for many arthropod species at various life stages, mainly for purposes of pest control. In a tabulation of such sterilizing doses for male arthropods, which are generally more radiosensitive than females, including 285 species representing 61 families of insects and other arthropods, irradiated mainly as pupae or nymphs and generally achieving at least 90% parental and F₁ sterility, the average dose is ~110 Gy (ref. 15 and www-ididas.iaea.org/ididas/). For the most radio-resistant species in the tabulation, the lepidopteran Spilosoma (*Diacrisia*) obliqua, a dose of 200 Gy to pupal or adult males, the most resistant sex, resulted in >99% F₁ sterilization. Among other invertebrates, sensitivity to IR has been measured for X-irradiation of Caenorhabditis elegans eggs in buffer, for which ~30 Gy reduced the number developing into adults by 90%.

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**Table 1. Reproductive performance of bdelloid and monogont rotifers exposed to IR**

<table>
<thead>
<tr>
<th>Species</th>
<th>Wells with eggs, n</th>
<th>Wells with F₁, n</th>
<th>Total eggs, n</th>
<th>Total eggs hatched, n</th>
<th>Relative parental fertility</th>
<th>Relative parental fecundity</th>
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<tr>
<td><em>E. dilatata</em></td>
<td></td>
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<tr>
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<td>249</td>
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<td>278</td>
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<td>86</td>
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</table>

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and the snail Zonitoides arboreus, which produced only sterile eggs after exposure to 70 Gy (16, 17). Among the very few animal species known to be capable, like bdelloid rotifers, of eggs after exposure to 70 Gy (16, 17). Among the very few animal species known to be capable, like bdelloid rotifers, of reproduction. Shown are A. vaga (triangles), P. roseola (squares), and E. dilatata (circles). Relative fecundity (filled symbols), relative fertility (empty symbols), and DSBs per Mbp (empty diamonds) are indicated. Left vertical axis shows relative fecundity or fertility; right vertical axis shows DSBs per Mbp; horizontal axis shows dose in Gy.

Fig. 1. Dose dependence of bdelloid and monogonont reproduction. Data on reproduction are from Table 1. (A) Parental reproduction. (B) F₁ reproduction. Shown are A. vaga (triangles), P. roseola (squares), and E. dilatata (circles). Relative fecundity (filled symbols), relative fertility (empty symbols), and DSBs per Mbp (empty diamonds) are indicated. Left vertical axis shows relative fecundity or fertility; right vertical axis shows DSBs per Mbp; horizontal axis shows dose in Gy.

As we find for bdelloid and monogonont rotifers, the effect of IR on other animals generally extends beyond the irradiated parents to their progeny. F₁ sterility induced by parental irradiation, widely made use of in insect control programs, is associated with chromosome aberrations, mutations, and possibly deleterious epigenetic effects expressed as impaired reproduction in generations beyond that which is irradiated.

The observation that rotifers rendered sterile by IR nevertheless remain active is consistent with the fact that somatic cell division is already complete at the time of hatching (20–23) and with the general observation that metabolic processes are less radiosensitive than cell division. Egg production, however, involves two maturation divisions starting from primary oocytes present in the hatched animal (24, 25) yet is also essentially unaffected even at the highest doses tested, suggesting either that egg production can proceed without maturation divisions or that the sterilizing effect of IR is manifested only later in development.

**Basis of Radiation Resistance.** The extraordinary resistance of bdelloid reproduction to IR cannot be attributed to an unusually small genome size, as the genome sizes of A. vaga and P. roseola are comparable to or larger than those of C. elegans, Drosophila melanogaster and many other animals that display no comparable radiation resistance (15, 21, 26, 27). Nor can bdelloid radiation resistance be explained by genetic redundancy or a multiplicity of templates for homologous repair. Bdelloid primary oocytes are in G₁ (21, 27) and, as described in ref. 28, bdelloids appear to be degenerate tetraploids, with only two copies of most genes.

The possibility that bdelloid radiation resistance reflects a novel chromosome alignment that keeps homologous regions together, facilitating access of broken ends to homologous templates, has been proposed for D. radiodurans where it appears unlikely for a number of reasons (29). Moreover, neither chromosome alignment nor the novel variant of synthesis-dependent strand annealing proposed to facilitate DSB repair at high levels of DNA breakage (30), provide an explanation for the extended shoulder in bdelloid and D. radiodurans dose–response relations that distinguishes them from radiosensitive organisms.

It might be thought that the measured efficiency of IR breakage, 0.005 DSB per Gy per Mbp, refers only to somatic DNA and that bdelloid germ line DNA is somehow protected. We think this unlikely as no such resistance is seen in D. radiodurans or in other radiation-resistant prokaryotes, where the number of breaks made per Mbp per Gy, ~0.004 (31, 32), is not significantly different from that for radiosensitive bacteria, fungi, and animals. More likely, the ability of bdelloids to remain fertile after extensive DNA breakage and other damage inflicted by IR derives from attributes of the systems that repair such damage or those that protect the repair systems, or both.

The finding that IR killing of D. radiodurans is paralleled by oxidative damage to its proteins has led to the proposal that the extraordinary radio resistance of D. radiodurans results from unusually effective protection of its proteins against toxic products of IR (33), a conclusion consistent with the ability of extracts of D. radiodurans to protect E. coli from radiation killing (34) and with the much higher scavenging ability for reactive oxygen species in extracts of D. radiodurans than in similarly prepared extracts of E. coli (35). Although protein damage in highly radiosensitive species may cause lethality even before significant DNA damage occurs (36, 37), this is clearly not the case for bdelloid rotifers, where hundreds of DSBs are made by a dose of IR that causes but little reduction in fecundity.
The picture that emerges from the above considerations is that a major defense against radiation damage in *D. radiodurans* and bdelloid rotifers, accounting for the distinctive shoulders in their dose–response relations, is an enhanced capacity for scavenging the reactive molecular species generated by IR and that the proteins and other cellular components thereby protected include those essential for the repair of broken DNA. This, however, leaves the question of why DNA, in both *D. radiodurans* and bdelloid rotifers, is not similarly protected but instead accumulates DSBs in direct proportion to dose and to the same extent per Gy per Mbp as in radio-sensitive organisms (Fig. 1). A possible answer lies in evidence that IR-induced DSBs result from closely spaced scission events on opposite chains of the DNA duplex that are caused by clusters of hydroxyl radicals or other reactive species generated in close association with DNA that are relatively immune or inaccessible to elimination by scavengers (38, 39).

**IR and Desiccation.** Naturally occurring bacterial isolates that are highly resistant to IR are also resistant to desiccation, and mutations that diminish radiation resistance in *D. radiodurans* also reduce its resistance to desiccation. These observations and the low levels of IR in the natural habitats of *D. radiodurans* and other radioresistant bacteria indicate that their radiation resistance is a consequence of evolutionary adaptation to survive desiccation (40). Similarly, the extraordinary radiation resistance of bdelloid rotifers is almost certainly an adaptation to survive in their characteristic ephemeral aquatic habitats.

The association of resistance to IR with anhydrobiosis and the observation that desiccation, like exposure to IR, is accompanied by the production of reactive oxygen species (41) and protein oxidation in diverse biological systems and that proteins of desiccation-resistant bacteria are protected against such damage (33, 42) indicates that at least part of the damage caused by desiccation is the same as that caused by radiation. In *D. radiodurans* this includes DNA breakage, which increases with desiccation time and can reach a high level before there is appreciable killing (40). Although DNA breakage has not been investigated in desiccated bdelloid rotifers, their ability to resume reproduction as a function of desiccation time also exhibits an extended shoulder (2), and it seems likely that, as in *D. radiodurans*, desiccation is accompanied by extensive DNA breakage, implying that the evolution of bdelloid genomes has been accompanied by unusually frequent and extensive DNA breakage and repair.

**Implications for Bdelloid Genome Structure and Evolution.** Accurate repair of a DSB requires the presence of a homologous template. Although bdelloid primary oocytes are in G1 and therefore lack sister chromatids, the requirement for homologous templates would be satisfied if bdelloid primary chromosomes were present as colinear pairs. In fact, this appears to be the case, as described in ref. 28, showing that bdelloids are probably degenerate tetraploids. Homology of colinear chromosomes sufficient for efficient repair could be maintained by selection against clones in which such homology becomes inadequate to support DSB repair by template-dependent repair processes. Local relaxation of the requirement for homology might occur, however, if subor neo-functionalization becomes established at particular sites and where the decreased efficiency of repair is more than offset by selection against clones in which such sites have been homogenized.

There are several ways in which the anhydrobiotic lifestyle of bdelloid rotifers may have contributed to their success in avoiding the early extinction that is the usual fate of asexuals. Desiccation can be of benefit to surviving bdelloids by facilitating their dispersal and freeing them from desiccation-sensitive competitors, parasites, pathogens, and predators. In addition to such ecological factors, bdelloid anhydrobiosis may have genetic benefits. In degenerate tetraploids, homogenization associated with the repair of desiccation-damaged DNA may significantly accelerate the appearance of clones homozygous for recessive beneficial mutations (43).

Finally, frequent DSB repair may act to keep the deleterious load of transposable elements low and account for the remarkable paucity in bdelloid genomes of transposable elements (TEs) of types that are not prone to horizontal transfer, namely long interspersed element LINE-like and other nonviral retrotransposons (44–46). TEs could be deleted or truncated by single-strand annealing mediated by microhomologies and, if not homozygous, by removal of nonhomologous 3′ ends in the course of synthesis-dependent strand annealing (47). Synergistic selection against TEs could occur, for example, if in the course of DSB repair an extended invading strand dissociates after copying a donor TE. Reinsertion and continued replication may then occur either at a eutopic site, leading to faithful repair, or at an ectopic TE, giving a lethal or semilethal nonreciprocal translocation (48).

As the probability of a DSB being near a TE and the probability of ectopic reinsertion are both proportional to the density of homologous TEs, such template switching would cause synergistic selection against them. Nonreciprocal translocations, probably mediated by such template switching, accompany the formation of substantial palindromic IR-induced DNA. Insertion of Ty retrotransposons in diploid *S. cerevisiae* (J.L. Argueso, J. Westmorland, P.A. Mieczkowski, M. Gawell, T.D. Peters, and M.A. Resnick, unpublished data). DNA damage and repair associated with anhydrobiosis may therefore have enabled bdelloids to become free of vertically transmitted TEs and limit their load of horizontally transmitted elements, avoiding the unchecked and ultimately lethal increase of deleterious TEs that may otherwise occur in asexuals (49, 50).

**Materials and Methods**

**Rotifer Culture.** The bdelloid rotifers *A. vaga* and *P. roseola* were provided by Claudia Ricci (University of Milan, Milan, Italy) and purchased from Carolina Biological Supply, respectively. Our cultures of these species were established from single eggs and have been maintained continuously in spring water on a diet of *E. coli* for 11 (*A. vaga*) and 19 (*P. roseola*) years. The monogonont rotifer *E. dilatata* was provided by Elizabeth Walsh (University of Texas, El Paso, TX). Rotifers were cultured in MBL medium made up in spring water (51) and fed *E. coli* grown in tryptone broth (bdelloids) or *Chlamydomonas reinhardtii* (*E. dilatata*). *C. reinhardtii* was provided by Joel Rosenbaum (Yale University, New Haven, CT) and cultured with aeration in MBL medium with continuous illumination in a flask 20 cm from a 40-W cool-white (4100 K) spiral fluorescent lamp. Bacteria and algae were centrifuged and resuspended in distilled water twice, refrigerated, and used within 10 days. Rotifers were cultured in 150 × 25-mm plastic Petri dishes at room temperature, transferring ~1,000 animals by pipette to a dish containing fresh medium every week or so. *P. roseola* and *E. dilatata* are free-swimming and may easily be collected by pipette. *A. vaga* generally crawls leech-like on surfaces from which it is not readily dislodged but also gathers in swarms from which animals may easily be collected by pipette.

**Irradiation and Scoring Reproductive Performance.** Approximately 1,000 rotifers were transferred by pipette to a 5-cm plastic Petri dish containing 5 ml of half-strength MBL medium and kept 6–10 h at room temperature to clear the animals of food organisms. As rapid cooling was found to be lethal to *E. dilatata*, the dish was wrapped in a cloth towel, placed in a small cardboard box, and kept for ~6 h in the cold room before being taken, on ice, to the irradiation facility, a procedure followed for both monogononts and bdelloids. After addition of 0.5 ml of freshly prepared 2 mM L-cysteine to prevent the possible accumulation of toxic radiation products in the medium, the dish was placed on a block of ice on a slowly rotating platform beneath a 12,000-Gy/h Cs source delivering a dose of 140 Gy/h at the position of the reproductive compartment. After 10 min, the reproductive compartment was removed. A small fraction (52), young monogononts, distinguished by their small size, were generally selected for irradiation.

For determination of reproductive performance, the dish was removed approximately 20 min after irradiation. The number of eggs hatched, the number of *E. dilatata* per dish, and the number of *P. roseola* that were motile served as reproductive indices. The eggs hatched into adults, the adults matured, and the numbers of eggs were scored in two Petri dishes, 10 adults in each, for a total of 20 adults. This was repeated 5 times for *E. dilatata* and 3 times for *P. roseola*. Several unique index values are reported for the natural isolates and the strain from each species. The corresponding index values are shown when appropriate. Each index is an average of the number of animals or eggs scored.
from the irradiator at specific intervals, and individual animals were transferred by pipette to each well of a 96-well microtiter dish containing 0.15 ml of MBL medium per well and a small amount of food. The dish was kept on ice during sampling, making it relatively easy to dislodge A. vaga by pipetting. Relative parental fertility was scored as the proportion of irradiated animals producing at least one active daughter, normalized to the corresponding proportion for nonirradiated animals. Each day or two until egg deposition ceased, –1 week for E. dilatata and 2 weeks for the bdelloids, any daughters present (distinguished from the parent by their smaller size) were removed to avoid including the eggs of daughters in counts of eggs deposited by their mothers. A daughter from each fertile parental well was placed in a well of another microtiter dish containing food and 0.15 ml of MBL medium, and every few days any granddaughters present were removed. Relative F1 fertility was scored as the proportion of daughters producing at least one grand-daughter, normalized to the proportion produced by daughters of nonirradiated parents. This procedure explained below, bdelloid daughters were counted as they were removed from parental wells, as were bdelloid granddaughters removed from daughter wells.

Hatched and nonhatched eggs of E. dilatata and nonhatched bdelloid eggs were easily counted, but the transparency of hatched bdelloid eggs prevented their reliable identification. As inactive progeny were only rarely encountered, the number of blanched bdelloids egglays was therefore taken to be the same as the number of bdelloid progeny. At each dose, relative parental and F1 fecundity was taken as the total number of hatched eggs (monogononts) or progeny (bdelloids), normalized to the corresponding number from the nonirradiated animals. Three such experiments were performed with E. dilatata and one each with A. vaga and P. roseola.

**Pulsed-Field Gel Electrophoresis.** For examination of DNA breakage ~1,000 starved, prechilled rotifers were irradiated as described above for 0, 1/6, 2, 4, 6, or 8 h, harvested by centrifugation in the cold at ~10,000 g for 5 min, resuspended in 0.2 ml of cold 50 mM EDTA and 10 mM Tris (pH 8.0), and centrifuged again. After removing 0.16 ml of supernatant, tubes were placed in a 42°C water bath and supplemented with 40 μl of freshly melted 1% low melting point agarose (LMPA; NuSieve GTG) in 2× Tris-EDTA (200 mM EDTA, 100 mM Tris, pH 8.0) at 42°C, mixed well by pipetting, transferred to a plug mold, and left to solidify for 30 min in the cold room. Each plug and an additional plug containing S. cerevisiae chromosomes (New England BioLabs MN03455) was individually placed in 0.2 ml of digestion buffer (1× Tris-EDTA supplemented with 1% sarcosyl and 1 mg/ml freshly dissolved proteinase K) and kept for 1 h in the cold room and 18 h at 55°C. Plugs were then rinsed with 0.5× 40 mM Tris acetate, 1 mM EDTA, pH 8.5 (TAE), kept with gentle rocking for 3 h in 1 ml of 0.5× TAE at room temperature and embedded in a 5-mm deep slab of 0.7% LMPA in 0.5× TAE. Electrophoresis was done in a BioRad CHEF-DR III electrophoretic system at 14°C and 5 V/cm, with a switch angle of 20° and switch times of 50 to 250 s for 18 h.

After electrophoresis, the gel was placed in SYBR Gold (Invitrogen) freshly diluted 1:1000 in water, gently rocked overnight, and scanned with a BioRad Molecular Image FX with Quantity One quantitation software (SYBR Gold settings: 488-mw excitation, 530-mw band pass). Under the conditions used, signal intensity is a linear function of DNA concentration in the gel (53). After baseline subtraction, each photometric scan was divided into 33 equal intervals along the direction of migration and the number average molecular size of DNA was calculated as \[\text{size} = \frac{\text{area} \times \text{bandpass}}{\text{intensity}}\] by summation over intervals, taking molecular sizes from a plot of the migration distance of the yeast markers against their sizes.

**ACKNOWLEDGMENTS.** Karine VanDoninck participated in the early phase of this work. This work was supported by the Eukaryotic Genetics Program of the National Science Foundation.


