The Isolation and Some Properties of Radiation-sensitive Mutants of *Micrococcus radiodurans*

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**SUMMARY**

Treatment of the radiation-resistant bacterium *Micrococcus radiodurans* with ultraviolet (u.v.) radiation and *N*-methyl-*N'*-nitro-*N*-nitrosoguanidine resulted in the isolation of two mutants highly sensitive to u.v. radiation. They were also sensitive to ionizing radiation and to the action of *N*-methyl-*N'*-nitro-*N*-nitrosoguanidine. The concentrations of sulphydryl groups in bacteria of the wild type and the mutants were not significantly different. Although the mutants were more sensitive to mitomycin C than the wild type the resistance of the latter was low. It is suggested that the DNA repair mechanism in the wild type operates very efficiently for the removal of single strand damage but not for that which involves cross-linking.

**INTRODUCTION**

*Micrococcus radiodurans*, a pigmented non-sporing bacterium, is characterized by its extreme resistance to ionizing radiation (Anderson, Nordan, Cain, Parrish & Duggan, 1956) and to ultraviolet (u.v.) radiation (Duggan, Anderson, Elliker & Cain, 1959). The bacterial DNA, considered to be the radiation-sensitive target in other, more sensitive, bacteria, is not unusual in *M. radiodurans* either in its base composition or its quantity per cell (Moseley & Schein, 1964) and it is not exceptionally resistant to u.v. radiation damage as measured by the amount of the lethal photoproduct, thymine-dimer, formed on irradiation (Setlow & Duggan, 1964). There is now evidence for the existence in *M. radiodurans* of an efficient enzymic mechanism for the repair of u.v. damage, which excises thymine-dimers from its DNA with such efficiency that the eventual death of the bacteria appears to be due to other causes such as damage to deoxycytidine and protein (Setlow & Boling, 1965; Boling & Setlow, 1966). The repair mechanism is of the dark-repair type, similar to that found in u.v.-resistant strains of *Escherichia coli* (Setlow & Carrier, 1964; Boyce & Howard-Flanders, 1964a) but apparently operating with a much greater efficiency, since *M. radiodurans* can survive much higher doses of u.v. radiation. For example, the dose of u.v. radiation required to inactivate 90% of the organisms in a culture of the u.v.-resistant strain of *E. coli K 12 AB 1157* is about 1000 ergs/mm² (Boyce & Howard-Flanders, 1964a) while 15,000 ergs/mm² is needed to achieve the same effect with *M. radiodurans*. There is also evidence that the resistance of *M. radiodurans* to ionizing radiation, which unlike u.v. radiation does not form thymine-dimers in DNA, is due to an enzymic repair mechanism for DNA damage (Moseley & Laser, 1965a; Dean, Feldschreiber & Lett, 1966). This mechanism

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appears to be essentially the same as that which operates for the repair of u.v. damage
(Moseley & Laser, 1965 b). The comparison of events which occur in u.v.-sensitive
and u.v.-resistant strains of *E. coli* after u.v. radiation has greatly contributed to an
understanding of the repair processes occurring in the resistant strains. Attempts have
therefore been made to isolate radiation-sensitive mutants of *M. radiodurans* in
order to establish by comparison the mechanism which confers on the wild type the
unique property of being the vegetative bacterium which is most resistant to radiation.

**METHODS**

**Organisms.** The strain of *Micrococcus radiodurans* used was originally isolated by
Anderson et al. (1956) and has been propagated in this laboratory for about 9 years.
The bacteria grow in tetrads and contain a carotenoid pigment which gives colonies a
salmon-pink appearance. The pigment can be extracted from dried bacteria with warm
methanol and has an absorption peak at 475 m\(\mu\).

**Media.** TGYA broth for growth contained Bacto-Tryptone (Difco), 5 g.; glucose,
1 g.; yeast extract (Difco), 3 g.; aspartic acid, 2 g.; distilled water, 1 l., adjusted to
pH 7.2 with NaOH. TGYA agar for colony counts and replica plating was made by
solidifying this medium with 15 g. Bacto agar/litre.

A buffer solution was used for suspending organisms during irradiation. It contained:
\(\text{KH}_2\text{PO}_4, 13.6 \text{ g.} ; \text{Na}_2\text{SO}_4, 2 \text{ g.}; \text{MgSO}_4\cdot7\text{H}_2\text{O}, 0.2 \text{ g.}; \text{Ca(NO}_3)_2\cdot4\text{H}_2\text{O}, 0.01 \text{ g.}; \text{FeSO}_4\cdot7\text{H}_2\text{O}, 0.5 \text{ mg.};\)
distilled water to 1 l., adjusted to pH 7.2 with KOH. The same
buffer was used for washing the bacteria and for dilution of suspensions for colony
counts.

**Preparation of organisms for irradiation.** Colonies from an agar plate were inoculated
into 12 ml. TGYA broth in \(\lambda\)-tubes and the tubes shaken at 30\(^{\circ}\) for 18 hr. The or-
ganisms were centrifuged down, washed and resuspended in buffer solution at a con-
centration of about \(10^8\) colony forming units/ml. Care was taken to break up clumps
of organisms by homogenising the suspensions immediately before irradiation by
using an MSE homogeniser (Cat. No. 7700).

**U.v. irradiation** was done by using a Hanovia germicidal lamp (Model 12). 5 ml.
samples of washed bacterial suspension were irradiated in Petri dishes (9 cm. diam.)
at a distance of 40 cm. from the lamp, the dose rate being 22.5 ergs/mm.\(^2/\sec.\)
The suspension was agitated during irradiation by means of a magnetic stirrer to
prevent sedimentation of the bacteria and to maintain uniformity of the absorbed dose.

**\(\gamma\)-irradiation** was carried out in a \(\gamma\)-source at a dose rate of 16.7 Krad/min.
Three ml. volumes of bacterial suspension were irradiated, oxygen being bubbled
during the irradiation.

**Isolation of mutants sensitive to u.v. radiation.** Optimum conditions for the isolation
of bacterial mutants usually result in the diminution of the colony count to about
\(0.1-1.0\%\) of the initial count by treatment with the mutagen. However, \(N\)-methyl-\(N'\)-
nitro-\(N\)-nitrosoguanidine (NG) has been found to give high mutation rates in *Escherichia
coli* at concentrations which caused only a \(50\%\) loss of viability (Adelberg, Mandel &
Chen, 1965). In contrast, very much higher concentrations of NG (500 \(\mu g./ml.)\) did not
produce any loss of viability in *Micrococcus radiodurans* even after incubation for 3 hr.
In view of the additive effect of ionising and u.v. damage (Moseley & Laser, 1965 b) it
was decided to combine u.v. radiation with application of the mutagen. This was done
Radiation sensitive \textit{M. radiodurans}

by irradiating the bacteria with a non-lethal dose of u.v. radiation before exposing them to the mutagen. This combined treatment produced the desired decrease in viability with optimal chances of producing sensitive mutants.

Four ml. of a suspension (about $10^8$ colony forming units/ml.) irradiated with 10,000 ergs/mm.$^2$ of u.v. radiation were added to 5 ml. TGYA broth and the suspension shaken at $30^\circ$ for 30 min. One ml. NG solution (5 mg./ml.) was added to a concentration of 500 $\mu$g./ml. After 45 min. further incubation the colony count was decreased to 0.1-1.0% of the original count. Samples (0.1 ml.) were diluted 100-fold in TGYA broth, to dilute out the mutagen, and shaken at $30^\circ$ to allow one or two divisions to take place. Agar plates were then spread with dilutions of culture to give about 80-100 colonies/plate and incubated at $37^\circ$ for 2 days. Two replicate copies of each plate were made by the felt-pad technique (Lederberg & Lederberg, 1952). One copy was irradiated under the u.v. lamp for 20 min. (a dose determined by sterilising small colonies of u.v.-sensitive \textit{Salmonella typhimurium} CLT 22). After incubation for 24 hr. at $37^\circ$ the non-irradiated and irradiated plates were compared for growth. Colonies on the non-irradiated plates which were absent from the irradiated plates were isolated and screened for u.v. sensitivity.

\textit{Resistance of Micrococcus radiodurans to the lethal action of NG and to mitomycin C.} A sample (2-5 ml.) of an 18-hr culture of \textit{M. radiodurans} was added to 7.5 ml. TGYA broth and shaken at $30^\circ$ for 100 min. (about 1 generation time). NG or mitomycin C was added to 100 $\mu$g./ml. and 20 $\mu$g./ml., respectively, and 0.1 ml. samples removed at suitable time intervals, diluted in buffer, and colony counts made on TGYA agar plates.

\textit{Determination of sulphydryl-group concentration in Micrococcus radiodurans.} The technique was based on that of Hamm & Hofmann (1965). To 1 ml. of a thick suspension of organisms (equiv. 50-70 mg. dry wt), 35 ml. of 8 M-urea and a slight excess of AgNO$_3$ (3.5 $\mu$ mole) were added. After stirring at room temperature for 1 hr to allow complete reaction between the SH-groups and AgNO$_3$, 4.5 $\mu$ mole of glutathione was added to give a slight excess of SH-groups. The excess of glutathione (as SH) was then titrated amperometrically with $10^{-3}$ M-AgNO$_3$.

\textbf{RESULTS}

\textit{Isolation of mutants}

Two mutants of wild-type \textit{Micrococcus radiodurans} which were sensitive to u.v. radiation were isolated. The two mutants are referred to as \textit{M. radiodurans} uv 17 and \textit{M. radiodurans} uv 38. It was necessary before studying the new strains to show that they were mutants of \textit{M. radiodurans} and not contaminants. The colonies of all three strains, wild type, uv 17 and uv 38 were similar in size and appearance, and all contained the carotenoid pigment which shows an absorption peak at 475 m.$\mu$. The concentration of pigment/unit dry wt organism was not significantly different in the three strains. Morphologically the organisms were identical and had the same mode of division which led to the formation of tetrads. Comparison of turbidity measurements with colony counts over the growth cycle gave similar plots, indicating that the dimensions of the organisms were similar.

Growth rates in TGYA broth were measured by using turbidity and colony count methods. In log-phase growth at $30^\circ$ with adequate aeration the generation time for
the wild type was 90 min.; UV 17 had the same generation time, but that of UV 38 was 150 min., i.e. about 1.7 times longer. During the study of mutant UV 17 another mutant arose spontaneously which was non-pigmented but which had all the other properties of UV 17, e.g. identical cell morphology, growth rate, and radiation sensitivity. This strain has been called *M. radiodurans* UV 17 w.

**Radiation resistance**

The dose response curves of the two mutants to u.v. radiation, compared with that of the wild type, are plotted in Fig. 1 which shows an enormous increase in sensitivity. The dose response curves to γ-radiation (Fig. 2) shows that even though the mutants were isolated on the basis of their u.v. sensitivity they were also sensitive to ionizing radiation. All the survival curves were sigmoidal. The shapes of the dose response curves have been defined by their intercept number, which is the dose obtained by extrapolating the exponential part of the curve to unit survival (see Fig. 1). A summary of the results is given in Table 1.

Although the mutants were much more sensitive than the wild type, the increase in sensitivity to both types of radiation was not the same. A direct comparison of the survival curves is complicated by the fact that, with one exception, the increase in the exponential slope was not proportional to the reduction in the shoulder. Thus the reduction in the length of shoulder of strain UV 38 was much greater (28-fold for u.v. and five-fold for γ-radiation) than the increase in exponential slope, viz. five-fold for u.v.-radiation and two-fold for γ-radiation. Nevertheless, the data show that for
Radiation sensitive *M. radiodurans*

Table 1. Analysis of the dose-response curves shown in Figs. 1 and 2

The D_{10} value is the dose required to decrease the colony count of a culture by 90% on the exponential part of the survival curve. The intercept value, in terms of dose, is obtained by extrapolating the exponential part of the curve to unit survival. The ratios are derived from the D_{10} and intercept values of the various strains divided by those of the wild type.

<table>
<thead>
<tr>
<th>u.v. radiation dose response curves</th>
<th>γ-radiation dose response curves</th>
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</thead>
<tbody>
<tr>
<td>Strain</td>
<td>D_{10} (ergs/mm^2)</td>
</tr>
<tr>
<td>Wild type</td>
<td>3,400</td>
</tr>
<tr>
<td>UV 17</td>
<td>340</td>
</tr>
<tr>
<td>UV 38</td>
<td>680</td>
</tr>
</tbody>
</table>

Strain UV 38 the increase in sensitivity to u.v. radiation was greater than that for γ-radiation. The same is true for strain UV 17, the reduction in the shoulder being 28-fold for u.v. radiation and 7-fold for γ-radiation, while the exponential slope increased by factors of 10 and 7 respectively.

The lethal effect of N-methyl-N'-nitro-N-nitrosoguanidine (NG)

Wild type *Micrococcus radiodurans* was completely resistant to the effect of 100 µg. NG/ml., showing no loss of viability after 80 min. of incubation. The radiation-sensitive mutants UV 17 and UV 38 were sensitive to the lethal action of NG, less than 10^{-8} of their populations surviving after 10 and 15 min. of incubation, respectively (Fig. 3). From this it follows that the increase in sensitivity of the mutants to NG was greater than that towards radiation. It was noted that practically all the colonies derived from wild-type organisms which had been incubated in the presence of NG 100 µg./ml. for 80 min. showed abnormalities, e.g. sectoring caused by partial loss of pigment, roughness, growth-rate variations, leading to the formation of very irregular edges to the colonies. On the other hand, colonies derived from surviving organisms of strains UV 17 and UV 38 were normal as compared with colonies from untreated organisms.

The lethal effect of mitomycin C

Figure 4 shows the lethal effect of mitomycin C at a concentration of 20 µg./ml. on *Micrococcus radiodurans* and its mutants UV 17 and UV 38. The radiation-sensitive mutants were more sensitive to the action of mitomycin C than the wild type but the disparity was less than that for the lethal effect of u.v. radiation.

Sulphhydryl-group content

Three samples of each bacterial strain were analysed for sulphhydryl group content. Values obtained were 21, 24, and 27 µmole SH/g. dry wt of wild type, strain UV 17 and strain UV 38, respectively. The dry weight of a colony-forming unit of *Micrococcus radiodurans* obtained previously was 2.5 × 10^{-12} g. (Moseley & Laser, 1965a) which gives values of 0.5, 0.6, and 0.7 × 10^{-16} mole SH/colony-forming unit for the three strains.
DISCUSSION

The fact that Micrococcus radiodurans is the most resistant vegetative bacterium, so far investigated, to u.v. radiation as well as ionizing radiation has often been overlooked, and explanations have been sought for its resistance to ionizing radiation which have no relevance to its resistance to u.v. radiation. Some explanations have postulated that an intracellular carotenoid pigment (Kilburn, Bellamy & Terni, 1958) or a sulphydryl compound (Bruce, 1964; Bruce & Malchman, 1965) act as protectors. The isolation of non-pigmented mutants of M. radiodurans which have the same resistance as the wild type of ionizing radiation indicated that the pigment is not responsible for such resistance (Moseley & Laser, 1965a) but it could not be excluded that pigment precursors were present which acted as energy-transfer substances. This possibility has now been excluded by the isolation of pigmented sensitive mutants.

The value of $0.5-0.6 \times 10^{-18}$ mole SH/colony-forming unit is in reasonable agreement with that obtained by Bruce & Malchman (1965) of $0.8-2.0 \times 10^{-18}$ mole/organism based on the binding of $p$-hydroxymercuribenzoate. However, this concentration of sulphydryl groups is present in the radiation-sensitive organisms as well as in the radiation-resistant organisms. The mere presence of sulphydryl groups is not an indication of the presence of an intracellular protective compound able to confer such high resistance as is shown by the wild-type Micrococcus radiodurans. However, some of the residual resistance to ionizing radiation in the sensitive strains may yet be due to the presence of sulphydryl groups and thus account for the fact that they
are slightly less sensitive to ionizing radiation than to u.v. radiation. This is a very small part of the total resistance of the wild-type *M. radiodurans* to ionizing radiation, the major part of which is due to the presence of a repair mechanism.

The dark repair system which operates in u.v. resistant strains of *Escherichia coli* does not only recognize damage to DNA of the thymine-dimer type. The excision of defective bases and the incorporation of new ones follows treatment of organisms with the bifunctional alkylating agent nitrogen mustard (Hanawalt & Haynes, 1965) and with NG (Hanawalt, 1966). Thus the precise base defect appears to be less important than some associated secondary structural alteration in the phosphodiester backbone of the DNA. This situation is also true for *Micrococcus radiodurans*. Its remarkable capacity to survive extremely high doses of ionizing radiation and of u.v. radiation is complemented by its resistance to the decay of radioactive phosphorus (\(^{32}\)P) incorporated in its DNA and to the action of NG, which probably acts as a monofunctional alkylating agent. *Micrococcus radiodurans* organisms remain viable even after 50,000 \(^{32}\)P disintegrations/nucleus, as compared with values of 10–50 for *E. coli* (M. Swann, personal communication). Since *M. radiodurans* has such great resistance to radiation and to compounds which cause DNA defects, it is surprising that it is sensitive to mitomycin C which cross-links DNA *in vivo* (Iyer & Szybalski, 1963) and which functions partially as a bifunctional alkylating agent (Iyer & Szybalski, 1964). Although wild-type *M. radiodurans* is more resistant to mitomycin C than are the radiation-sensitive mutants, the order of resistance is quite low, being the same as that shown by u.v.-resistant strains of *E. coli* (Boyce & Howard-Flanders, 1964b). This suggests that *M. radiodurans* has a repair system of extremely high efficiency for damage within one or both strands of the DNA, but which does not operate for damage involving cross-linkage. The higher resistance of wild-type *M. radiodurans* as compared with that of the sensitive mutants need not be interpreted in terms of removal of cross-links since only 1 out of 5 to 10 mitomycin C molecules participates in the formation of cross-links while the others react with one DNA strand only (Iyer & Szybalski, 1964; Weissbach & Lisio, 1965). This suggests that the differential sensitivity of the three strains of *M. radiodurans* to mitomycin C reflects the inability of the radiation-sensitive strains to repair monofunctional alkylating damage which the wild type repairs very efficiently (hence its high resistance to NG) but is inactivated by cross-link damage. Preliminary biochemical studies indicate that the radiation-sensitive mutants and wild-type *M. radiodurans* are able to excise u.v. radiation-induced thymine dimers from their DNA and are able to incorporate fresh bases. Their sensitivity appears to be due to the inability to control the excision process (Moseley, 1967).

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References


