

Genome-wide study predicts promoter-G4 DNA motifs regulate selective functions in bacteria: radioresistance of *D. radiodurans* involves G4 DNA-mediated regulation

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ABSTRACT

A remarkable number of guanine-rich sequences with potential to adopt non-canonical secondary structures called G-quadruplexes (or G4 DNA) are found within gene promoters. Despite growing interest, regulatory role of quadruplex DNA motifs in intrinsic cellular function remains poorly understood. Herein, we asked whether occurrence of potential G4 (PG4) DNA in promoters is associated with specific function(s) in bacteria. Using a normalized promoter-PG4-content (PG4_p) index we analysed >60 000 promoters in 19 well-annotated species for (a) function class(es) and (b) gene(s) with enriched PG4_p. Unexpectedly, PG4-associated functional classes were organism specific, suggesting that PG4 motifs may impart specific function to organisms. As a case study, we analysed radioresistance. Interestingly, unsupervised clustering using PG4_p of 21 genes, crucial for radioresistance, grouped three radioresistant microorganisms including *Deinococcus radiodurans*. Based on these predictions we tested and found that in presence of nanomolar amounts of the intracellular quadruplex-binding ligand N-methyl mesoporphyrin (NMM), radioresistance of *D. radiodurans* was attenuated by ~60%. In addition, important components of the RecF recombinational repair pathway *recA*, *recF*, *recO*, *recR* and *recQ* genes were found to harbour promoter-PG4 motifs and were also down-regulated

in presence of NMM. Together these results provide first evidence that radioresistance may involve G4 DNA-mediated regulation and support the rationale that promoter-PG4s influence selective functions.

INTRODUCTION

Guanine-rich sequences are known to adopt non-canonical secondary structure forms known as guanine quadruplex or G4 DNA motifs. These are four-stranded, Hoogsteen base-paired self-assembly of DNA strands in parallel/antiparallel orientation stabilized by charge co-ordination with monovalent cations (Figure 1) (1–4). Intramolecular quadruplex motifs result from folding of a single-nucleotide chain with the guanine tetrads linked by loops of varying sizes and can adopt multiple conformations (5). On the other hand, combinations of nucleotide chains that contribute towards tetrad formation give intermolecular motifs. Sequence with potential to form quadruplex motifs are present in various regions of the genome including telomeres (3,6) and promoters (7–12). In telomeres, quadruplex motifs have been implicated in mechanisms that reduce activity of the ribonucleoprotein telomerase (13) and also in telomere capping in *Saccharomyces cerevisiae* (14). In promoters work from our and other groups demonstrate quadruplex motifs as potential regulatory elements that influence gene expression (15–21). Furthermore, recent findings predict role of quadruplex motifs in chromatin packaging (12,22), recombination (23), CpG methylation (24) and genomic translocations in cancer tissues (25). Following the finding that potential G4 (PG4) motifs are enriched in promoters of *Escherichia coli* and several other bacteria (7,11), several

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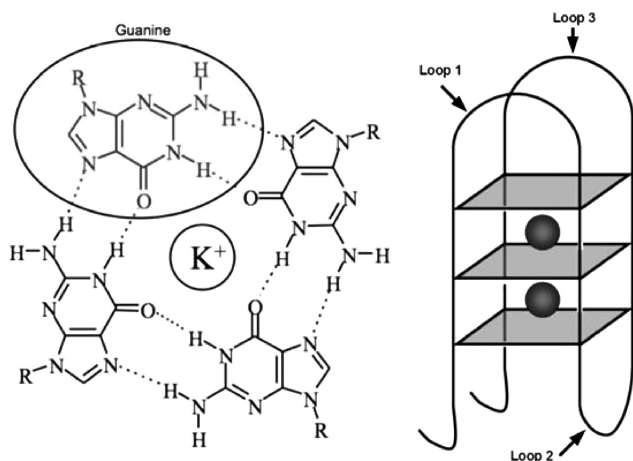


Figure 1. Structure of the guanine quadruplex motif. Hydrogen-bonded self-assembly of guanine bases stabilized by monovalent cations form tetrads (left) that make the core of the four-stranded structure. Intramolecular quadruplex motifs are made of guanine tetrads linked by three loops of variable nucleotide length comprised of any of the four bases A, T, G or C (right).

reports showed that promoters of many other species including human are not only replete with quadruplex-forming sequences (7–9,11,12) but that such motifs are also conserved across human, chimpanzee, mouse and rat promoters (10).

c-MYC was the first *in vitro* case where a G-quadruplex upstream of the P1 promoter was shown to affect transcription (15). This finding was further substantiated by other observations showing gene expression was influenced by G-quadruplexes within the core promoter of human *c-KIT* (26,27) and *k-RAS* (16) oncogenes. In addition, promoter-quadruplex motifs were reported for many genes, including *VEGF*, *PDGF*, *HIF1 α* , *BCL-2*, *RB*, *RET* (28,29), *HRAS* (30) and human telomerase hTERT (31,32). Furthermore, recently we found a non-canonical quadruplex motif, formed by two guanine repeats instead of three, to be functionally active in case of human *thymidine kinase 1* (33).

Further studies using chromatin immunoprecipitation (ChIP) experiments demonstrated that the non-metastatic factor NM23-H2 associates with the *c-MYC* promoter through a G-quadruplex motif providing relatively direct evidence in support of G-quadruplex-mediated transcription (17). In addition, interaction of recombinant hnRNP A1/Up1 with the *KRAS* promoter G-quadruplex (34); Myc-associated zinc finger protein (MAZ)/poly(ADP-ribose) polymerase 1 (PARP-1) binding to the G-quadruplex element in the murine *KRAS* promoter (35); and binding of nucleolin/hnRNP proteins to the G-quadruplex-forming sequences of the *VEGF* promoter (36) built more support for quadruplex-mediated transcription. Similarly, quadruplex motifs in the promoters of human sarcomeric mitochondrial creatine kinase, muscle creatine kinase and integrin α -7 of mouse were also shown to associate with the dimeric form of MyoD *in vitro* (37,38). In line with these reports, transcriptome

profiling in presence of intracellular G-quadruplex-binding ligands suggested a wide spread regulatory role of quadruplex motifs in transcription (18).

These studies gave credence to the possibility that quadruplex motifs, like many other regulatory elements, hold functional significance. However, unlike most established regulatory elements, association of quadruplex motifs with intrinsic cellular function(s) is poorly understood. But, given the complexity of eukaryote gene regulation, it is possible that regulatory role, if any, of quadruplex motifs can be better understood from the analysis of relatively less complex bacterial transcription. With this in mind, we sought to study possible links between gene function and the presence in gene promoters of motifs that potentially fold into quadruplex structures. This was done by analysing the relationship between functional classes of genes and quadruplex occurrence in their promoters, both at a genome-wide level and in individual genes (Figure 2a). Interestingly, this showed that promoter-quadruplex motifs occur in a fashion that is likely to impart specific functional attributes in species. We tested this prediction experimentally in *Deinococcus radiodurans* and *Deinococcus geothermalis* which withstand high levels of radiation. Findings suggest that quadruplex motifs present in promoters of key genes may play a critical role in response to radiation.

MATERIALS AND METHODS

Quadruplex detection

An algorithm written in Java was developed to identify sequence patterns with quadruplex-forming potential which was designed to find quadruplex and loop length combinations and count as well as perform sequence randomizations that were required for computing statistical significance of the results (see below). The algorithm is based on previous developed strategies (7) from our group and uses a tree structure. Briefly it assumes that:

- the stem size is constant in a single quadruplex and between 2 and 5;
- stems are only made of G (C on the complementary strand);
- the loop sizes are between 1 and 7; and
- one quadruplex is made of four stems and three loops.

For a given sequence, the algorithm checks every sliding window with size equal to

$$4 \times \text{max_stem_size} + 3 \times \text{max_loop_size}$$

which is the maximum size of a quadruplex. After the end of a given sequence is reached, all detected quadruplexes were returned with specific loop and stem size combinations. When quadruplexes with more than four stems were found, the first four stems were considered as a single quadruplex, the extra stem was considered for the subsequent sliding window.

Promoter-wise PG4 content—PG4_p

In order to compare PG4 motif content across genes and organisms we devised a method of attributing each promoter with a normalized value of PG4 motif content (PG4_p) based on PG4 motif density in a particular promoter which was controlled for both GC% and sequence content of the promoter. To compute significance of the presence of PG4 motifs we shuffled each promoter 100 times (while sequence content and GC% of individual promoters were maintained) and 100 simulated PG4_p values were used to find the simulated mean m and standard deviation σ . Assuming that the distribution was normal in random condition, PG4_p of a

particular promoter was considered statistically significant when it was at least two standard deviations above the simulated average PG4_p for that promoter

$$(\text{observed_value} > m + 2\sigma).$$

Z-score was computed for PG4_p and functional class enrichment (see below) using the classical formula:

$$(\text{obs} - \mu) / \sigma$$

where obs is the observed value of the variable (PG4_p or occurrence of genes for functional class analysis); μ is the expected means and σ is the standard deviation of the

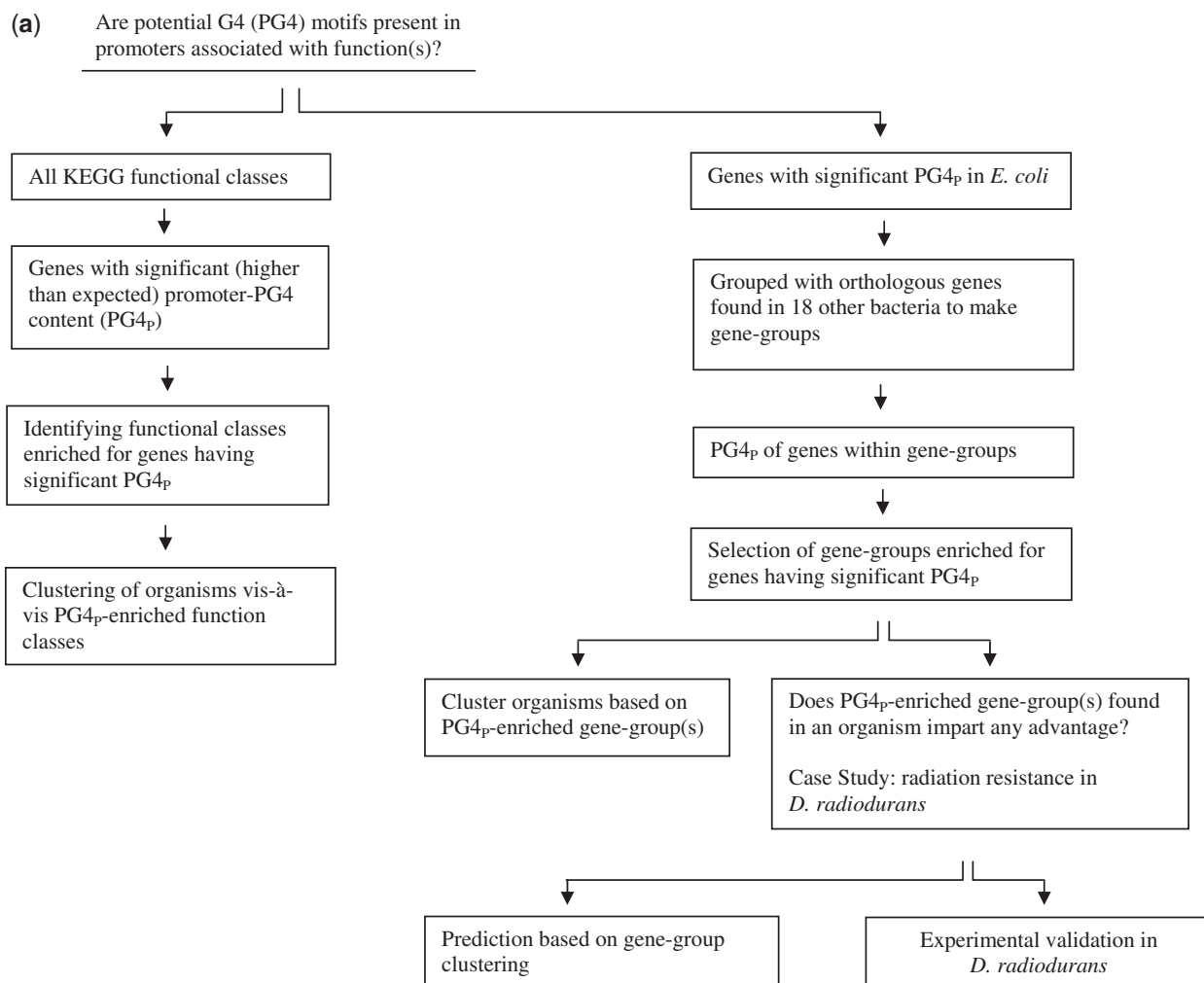


Figure 2. Promoter PG4 motifs define distinct functional classes in organisms. (a) Scheme for quadruplex-occurrence analysis in function classes (left panel) and individual genes (right panel). The index PG4_p was used for normalized content of potential quadruplex-forming sequence within individual promoters. Enrichment of promoters with significant PG4_p in a functional class was computed using the mean of randomly expected number drawn from 1000 simulations as described in Materials and Methods section. (b) Functional classes with higher than expected number of promoters having significant presence of PG4 motifs in *E. coli*; classes at least two standard deviation above expected are denoted with asterisk. (c) Heat map representation of cluster showing enrichment (z-score) of genes with significant PG4_p in functional classes across different organisms; blank squares indicate no gene with higher than expected PG4_p was found. (d) Heat map of clustering showing promoters of individual genes that have significant PG4_p across organisms. *Escherichia coli* was chosen as the reference organism and orthologues of the *E. coli* gene were used to construct gene-groups across 19 other organisms. Gene-groups where significant PG4_p ($z > 2.0$) was observed in at least three organisms (in addition to *E. coli*) are shown; Aca, *Acidobacterium capsulatum*; Afe, *Acidithiobacillus ferrooxidans* (ATCC 53993); Afo, *Acidimicrobium ferrooxidans*; Afr, *Acidithiobacillus ferrooxidans* (ATCC 23270); Art, *Arthrobacter sp.*; Bpf, *Bacillus pseudofirmus*; Bsu, *Bacillus subtilis*; Cai, *Catenulispora acidiphila*; Cbu, *Catenulispora burnetii*; Ddr, *Deinococcus deserti*; Dra, *Deinococcus radiodurans*; Eco, *Escherichia coli*; Gox, *Gluconobacter oxydans*; Hpy, *Helicobacter pylori*; Kra, *Kineococcus radiotolerans*; Msl, *Methylocella silvestris*; Nph, *Natronomonas pharaonis*; Rru, *Rhodospirillum rubrum*; Sul, *Sulfurihydrogenibium sp.*

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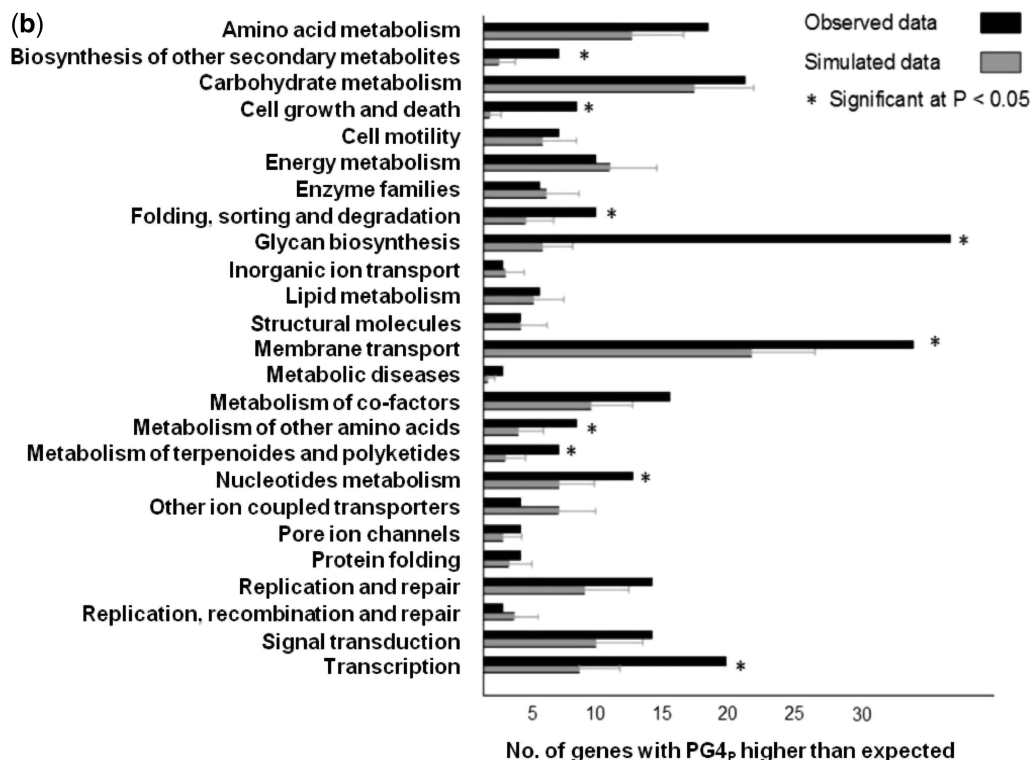


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variable were obtained from the simulations of PG_{4p} or functional classes enrichment.

Functional classes enrichment

For functional class annotations, 19 well-annotated bacterial genomes were obtained from KEGG (39). In order to avoid very general classifications, the second layer in KEGG annotation hierarchy was used for all analysis. For each gene the KEGG orthologue annotation was extracted and manually curated to avoid redundancy. In case of multiple annotations all possible functions were considered. To compute whether a particular function class was enriched for genes having significant PG_{4p} first the number of such genes in a particular class was found, which comprised the actual or observed set. Identical number of genes was randomly pulled from the same genome and 1000 such sets were prepared (randomly expected sets)—for each of the 1000 sets, number of genes with significant PG_{4p} were calculated; average number of genes across 1000 sets was used as the randomly expected number. A function class was considered to be enriched in genes with significant PG_{4p} when the actual or observed number was at least two standard deviations above the randomly expected average (that is z -score > 2).

Orthologous groups analysis

All *E. coli* genes with PG_{4p} higher than expected by at least two standard deviations from simulation (see above) were selected. Orthologues of these genes across the

remaining 18 genomes of our set were identified using KEGG orthologous cluster information. Fifty groups of orthologues were obtained in this way. For all these genes, z -score of PG_{4p} by comparison with simulation was computed. When an organism had more than one orthologue corresponding to the *E. coli* gene, only the gene with highest z -score was considered. In the same way orthologues of genes involved in radioresistance in *D. radiodurans* [taken from (40–42)] were selected using KEGG orthologous cluster information.

For cluster analysis we used ‘Cluster’ developed by Eisen *et al.* (43).

Growth conditions/treatment with quadruplex-binding ligands and gamma irradiation

To directly probe for the role of PG_{4p} in *D. radiodurans* IAM 12271 (MTCC 4465) and *D. geothermalis* (DSM 11300), porphyrin derivatives, N-methyl mesoporphyrin (NMM) or its un-methylated analogue mesoporphyrin IX dihydrochloride (MIX), and 5,10,15,20-tetrakis-(N-methyl-4-pyridyl)porphyrin (TMPyP4) or its positional isomer 5,10,15,20-tetra-(N-methyl-2-pyridyl)porphyrin (TMPyP2) were used. *D. radiodurans* and *D. geothermalis* were grown at 32°C in TGY (0.5% tryptone, 0.3% yeast extract, 0.1% glucose) broth containing respective concentrations of NMM or MIX (25 nM and 50 nM) and TMPyP4 or TMPyP2 (1.5 μM and 3 μM). Only cultures in exponential growth ($OD_{600\text{ nm}} = 0.2–0.5$) were evaluated for their ability to survive ionizing gamma radiation. Exponential phase bacterial cultures in TGY medium with or without NMM

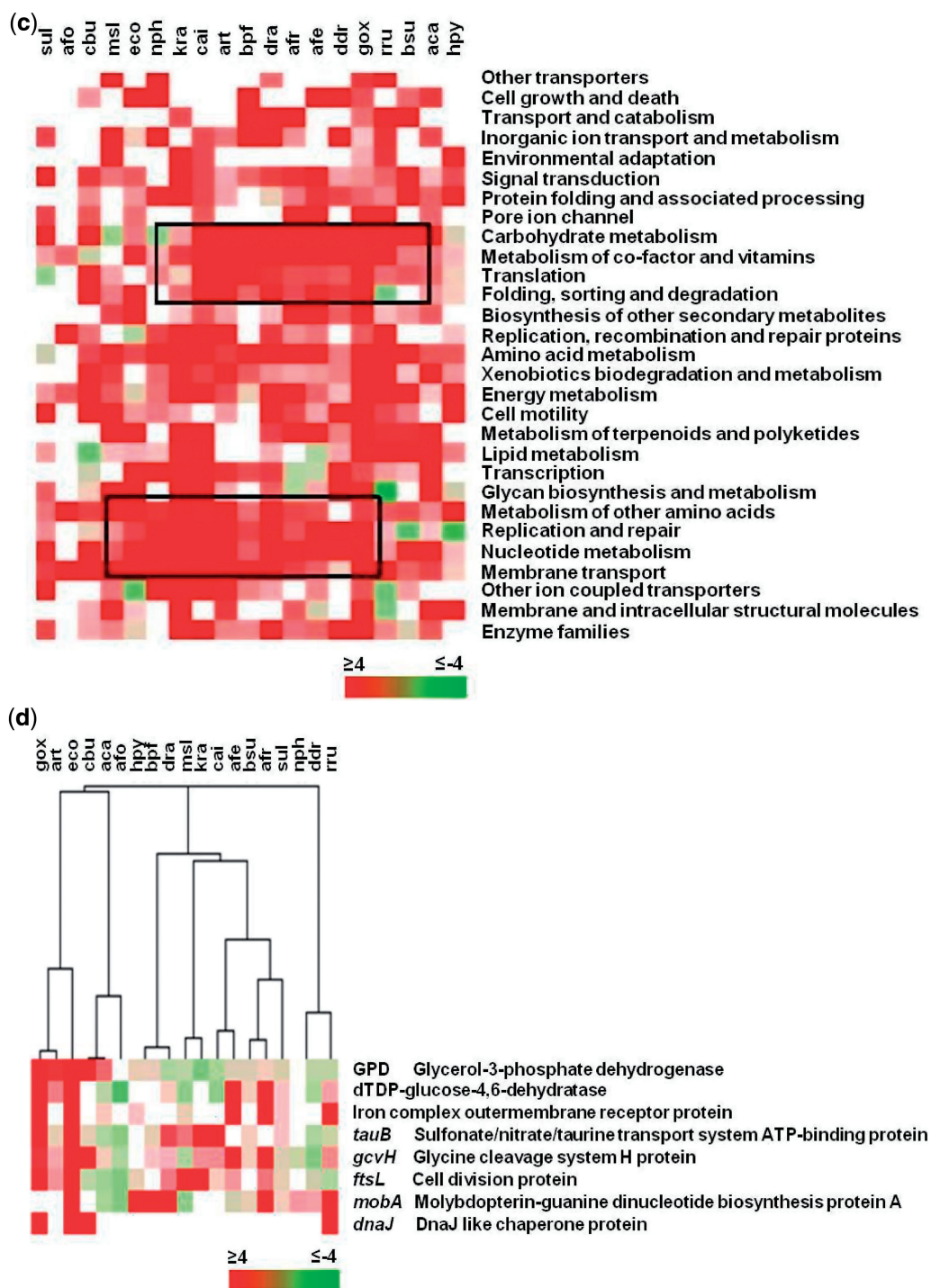


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(or MIX) and TMPyP4 (or TMPyP2) were exposed to gamma irradiation (see Supplementary Figure S1 for details of method used).

Radiation resistance was evaluated as described earlier (44). Briefly, mid log-phase culture ($OD_{600\text{nm}} = 0.3$) was divided into 40 ml aliquots, placed in 50 ml falcon tubes and were exposed to 5 kGy or 10 kGy of ^{60}Co γ -rays, at a dose rate of 2.57 kGy/h using ^{60}Co gamma chamber

(Gamma Cell 5000, BRIT, Mumbai, India) installed at Nuclear Research Laboratory, IARI, Delhi. All the irradiation experiments were done at 20°C. Another aliquot, kept outside the radiation source at 20°C, served as control. All the irradiated and unirradiated control samples were transferred into fresh TGY broth and incubated on a rotary shaker at 200 $\text{rev}\cdot\text{min}^{-1}$ at 32°C. For gene expression analysis RNA was isolated at 3 h

following irradiation based on earlier observations that found up-regulation/down-regulation of most genes after gamma irradiation was at 3 h post-irradiation recovery (45). Bacterial growth was observed by measuring turbidity at 600 nm of liquid cultures (TGY broth) and the viability of irradiated cells were evaluated after 24–90 h of post-irradiation recovery at 32°C as described earlier (46). All the assays were performed in triplicates.

Escherichia coli was grown at 37°C in Luria-Bertani (LB) broth containing 2 μM or 4 μM of NMM (or MIX) and TMPyP4 (or TMPyP2), which did not alter the growth of *E. coli*, and exposed to 1 kGy or 2 kGy of ⁶⁰Co γ-rays. The viability of *E. coli* in response to gamma irradiation was evaluated using the procedure described above, after 6–20 h of post-irradiation recovery at 37°C because of its lesser doubling time, as compared to *D. radiodurans*.

Gene expression analysis following irradiation

All samples were harvested in exponential phase by centrifugation at 10 000 rev.min⁻¹ for 10 min for RNA extraction. Total RNA was extracted from irradiated (at 3 h post-irradiation recovery) and unirradiated cultures using RNeasy RNA isolation kit (Qiagen) following manufacturer's protocol. Total RNA derived from each sample condition was treated with DNase I (Fermentas) and RNA quality and quantity were evaluated by determining UV absorbance at 260 nm and 280 nm. Two micrograms of each DNase I treated and purified RNA sample were reverse transcribed using the High-Capacity cDNA Reverse Transcription Kit (Applied Biosystems) as described in manufacturer's protocol. PCR primers were designed (Supplementary Table S1) to amplify each open reading frame based on fully sequenced *D. radiodurans* R1 and *E. coli* K-12 MG1655 genomes. Expression of *recA*, *recF*, *recO*, *recR* and *recQ* along with 16S rRNA transcript (endogenous control), expression of which was unaffected by ionizing radiation (47), was determined before and after irradiation. To analyse the relative intensity of the PCR bands, gel image of the PCR products was scanned and then analysed using AlphaEaseFC 4.0 software (Alpha Innotech, USA). All experiments were performed in triplicates.

RESULTS

To investigate the connection between promoter-PG4 motifs and function of genes, each promoter was assigned a normalized value of promoter-PG4 motif content, PG4_p, which was based on motif density (see Materials and Methods section). Significance of PG4_p was computed relative to 100 simulated PG4_p values obtained for respective promoters. For analysing function-PG4_p relationships we undertook two complementary approaches: (i) function class-specific, where PG4_p enrichment within orthologous function groups built from the KEGG database was analysed (Figure 2a, left panel) and (ii) gene-specific, where genes with significant PG4_p in a reference organism were used to query

across other organisms (Figure 2a, right panel). It was necessary that we studied only well-annotated organisms so that associations with function could be analysed with relative confidence, therefore 19 well-annotated organisms were considered.

Genes with PG4 motifs in promoters influence specific functions including carbohydrate metabolism

For each annotated function class in KEGG, we first determined the number of genes with significant PG4_p in a given organism. Next a rigorous method was used to ascertain whether the function class was enriched for PG4_p-genes by mere chance. Each class was randomly populated 1000 times to estimate the proportion of PG4_p-genes expected by chance; enrichment was considered significant when actual occurrence of PG4_p in any gene was at least two standard deviations above the random expectation (z -score > 2; see Materials and Methods section). All 19 organisms were analysed in this way to check for enriched function classes; a representative example for *E. coli* is shown in Figure 2b.

Next, we checked for correlation between enriched function classes, determined using PG4_p, and organisms: 29 function classes, found to be enriched in at least one organism, were clustered based on z -scores for enrichment of genes with significant PG4_p (Figure 2c, see Materials and Methods section for z -score analysis). Furthermore, to avoid any bias from sparsely populated function classes, we excluded ones that had <5 genes in any of the 19 species. Two major clusters were evident. The first cluster comprising 'carbohydrate metabolism', 'metabolism of co-factor and vitamins', 'translation' and 'folding, sorting and degradation' was enriched in 11 species. The second group of classes—'metabolism of other amino acids', 'replication and repair', 'metabolism of nucleotides' and 'membrane transport' were enriched in 13 species. Furthermore, we noted four functional classes to be largely predominant across most of the 19 organisms—'carbohydrate metabolism', 'amino acid metabolism', 'membrane transport' and 'energy metabolism'. As 'carbohydrate metabolism' involves glucogenesis, this function is closely linked to 'energy metabolism', therefore, it was interesting to find that both the classes were represented in majority of the organisms analysed. Taken together, it was apparent that metabolism of several essential entities like, carbohydrate, vitamins and amino acids appear to be important in the context of PG4 motif-controlled regulation.

Genes with promoter-PG4 motifs are organism specific

As mentioned above, in the second approach for function analysis we focused on individual genes that had significant PG4_p (Figure 2a, right panel). *Escherichia coli* was used as the reference organism and genes that had significant PG4_p ($z > 2.0$) in *E. coli* were selected and corresponding orthologues in the 18 other organisms identified using KEGG classifications; 150 gene-groups were obtained in this way. In cases where an organism had more than one gene corresponding to the *E. coli* gene, the one with the highest z -score for PG4_p was considered.

In order to see if there were any gene-groups that had significant PG4 content across organisms we clustered all the 150 groups across 19 organisms (Supplementary Figure S3a). Interestingly, most gene-groups were found to have only few organisms with z -score above two suggesting that it was unlikely that there was any particular gene(s) that had significant PG4_p that could be considered across several species. To ascertain that this was not due to the choice of *E. coli* as the reference organism, we performed similar analyses using either *Catenulispora acidiphila* (cai), *Kineococcus radiotolerans* (kra) or *Gluconobacter oxydans* (gox) as reference organisms; again, in each case we found very few gene-groups that were significant across organisms (Supplementary Figure S3b–d).

Next, we focused on gene-groups made with *E. coli* as reference that had at least three organisms with z -score of 2 or more, in addition to *E. coli*. Eight out of 150 orthologue groups that remained after this selection were clustered (Figure 2d), which revealed that for all genes except 1, *G. oxydans* (gox) and *E. coli* (eco) had high PG4_p. Similarly, *Sulfurihydrogenibium sp.* and *Bacillus subtilis* had significantly high PG4_p for at least three genes (glucose hydratase, an iron outer membrane receptor and a protein required for glycine cleavage). Overall, we noted that many genes with high PG4_p did not share this feature with other organisms suggesting that PG4_p enrichment of genes was likely to be specific to a particular organism.

Genes imparting resistance to radiation have enriched promoter-PG4 motifs in radioresistant bacteria

The observations described above suggest possible role of promoter-PG4 motifs in imparting specific functions to organism(s). Therefore, we reasoned, if one selects genes specific for a function and uses PG4_p to discriminate, it should be possible to segregate organisms. On the other hand, if PG4_p is not important one should not see any difference in the segregation pattern.

As a test case we selected radiation resistance because it has been extensively studied and genes for replication, repair and recombination delineated in radioresistant as well as other species (40–42). Furthermore, we reasoned that response to radiation as a functional readout would be relatively distinct. Twenty-one genes directly involved in radiation resistance were considered and orthologues of the selected genes in the remaining 18 bacteria from our set were determined using KEGG, giving 21 gene-groups. Next, we used PG4_p- z score for clustering, as done earlier, using both organisms and gene-groups (Figure 3). In doing so, we noted with interest that three radioresistant bacteria (*dra*, *D. radiodurans*; *ddr*, *Deinococcus desertii* and *kra*, *K. radiotolerans*) clustered together. Interestingly, using this analysis we found both *recA* and *recQ*, demonstrated to be important in imparting radioresistance, as part of a cluster which also had other important DNA repair genes like *radA* and the ATP-dependent protease *clpX* (48). On the other hand, and perhaps more importantly, it was evident that most organisms that were not radioresistant did not have enriched

PG4_p in these genes. Taken together, it was noteworthy that unbiased clustering with respect to genes involved in radiation resistance could segregate radioresistance bacteria vis-à-vis function.

Radiation resistance is compromised in presence of quadruplex-binding ligands

Based on above findings, we reasoned if quadruplex motifs are important for radiation resistance a ligand that could interact with quadruplex motifs inside cells would adversely affect radiation sensitivity. This was tested using the ligand NMM which specifically binds G-quadruplex motifs intracellularly and has no detectable binding to other nucleic acid structures, including ssDNA, dsDNA, triplex DNA, Z-DNA, duplex RNA and DNA–RNA hybrids (49,50). MIX, an unmethylated analogue of NMM, which does not bind to G-quadruplex motifs was used as a negative control (14,51). In addition, we used a second intracellular G-quadruplex-binding ligand TMPyP4 and its positional isomer TMPyP2, which does not bind G-quadruplex motifs (18,52).

We selected *D. radiodurans* and *D. geothermalis*, the best studied species among the members of *Deinococcus* for further studies (40–42,53). Experiments were performed at two doses of gamma irradiation, 5 or 10 kGy in presence or absence of NMM (or MIX) and TMPyP4 (or TMPyP2), as described under Materials and Methods section. We found that relative survival of *D. radiodurans* and *D. geothermalis* decreased in a dose-dependent fashion in presence of NMM: while ~65–80% of the bacteria survived at 5 kGy, at 10 kGy survival dropped to around 40–45% in presence of 50 nM NMM (Figure 4a and b, left panel). Relative survival of *D. radiodurans* and *D. geothermalis* also decreased in presence of TMPyP4; ~45–55% of the bacteria survived at 5 kGy whereas at 10 kGy survival dropped to about 20–35% in presence of 3 μM TMPyP4 (Supplementary Figure S4a and b, left panels). On the other hand, relative survival (%) of *D. radiodurans* and *D. geothermalis* was minimally affected in presence of MIX (Figure 4a and b, right panel) or TMPyP2 (Supplementary Figure S4a and b, right panel) in response to 5–10 kGy irradiation. As a control organism for radioresistance we used *E. coli*, which is far more sensitive to ionizing radiation than *D. radiodurans* and *D. geothermalis* (54); survival of *E. coli* was relatively unaffected in presence of ligands NMM/MIX and TMPyP4/TMPyP2 in response to irradiation (Supplementary Figures S5 and S6).

Promoters of key genes that confer radiation resistance harbour quadruplex motifs and are repressed in presence of quadruplex-binding ligand

Next, we sought to find out if compromised radiation resistance of *D. radiodurans* and *D. geothermalis* in presence of NMM was due to altered expression of key genes. In earlier studies we found that *D. radiodurans* in absence of *recA*, an important component of the DNA double-strand repair pathway, exhibits extreme sensitivity to ionizing radiation (55–57). Furthermore, as expected, we noted that *recA* expression was up-regulated on irradiation of

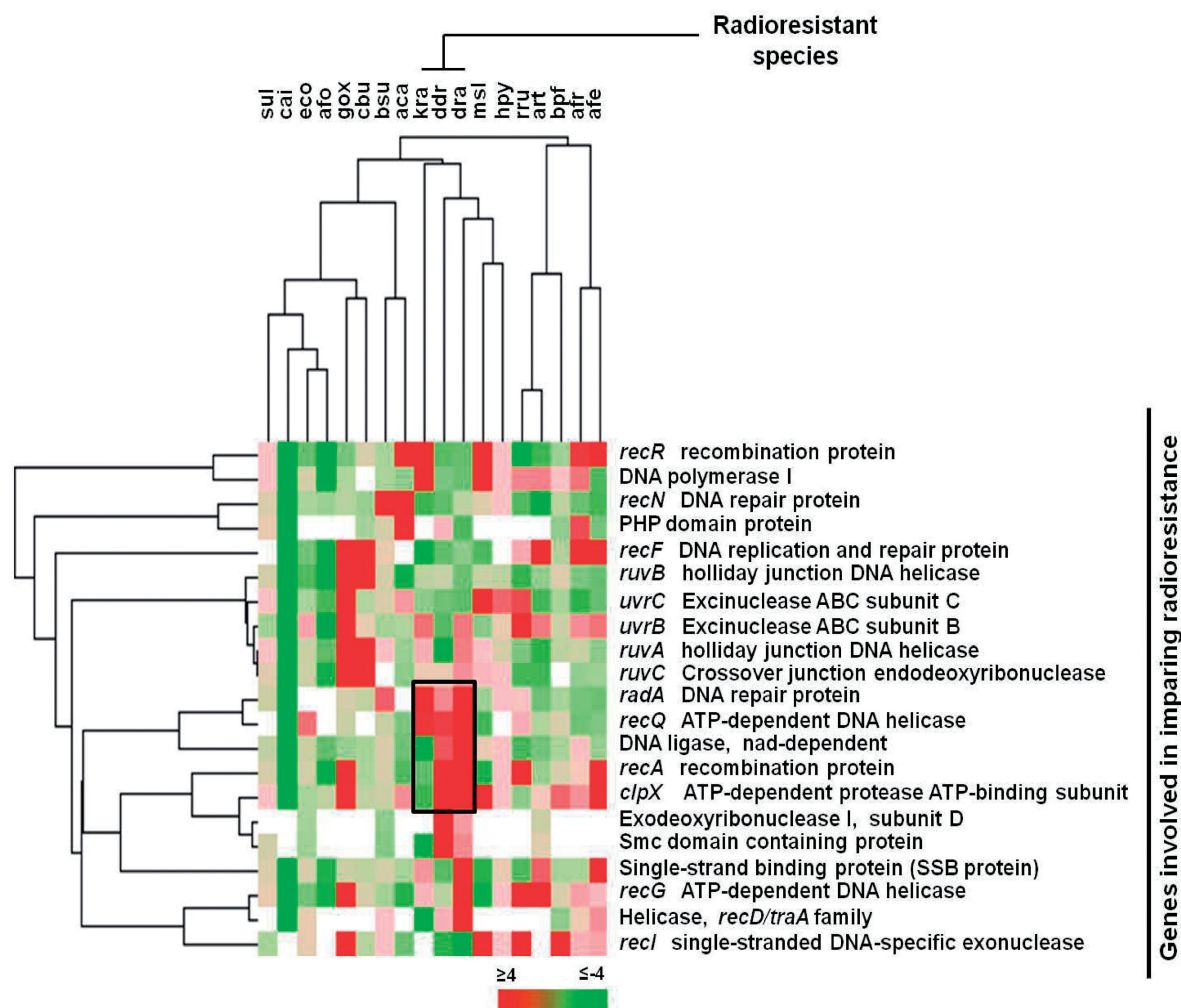


Figure 3. PG_{4p}-based analysis independently clusters radiation resistance species. Heat map representation of cluster diagram showing promoters of individual genes involved in imparting radiation resistance. Cluster was drawn for 18 organisms based on PG_{4p} of genes, with *E. coli* as the reference organism.

D. radiodurans at 5 kGy or 10 kGy in a manner that was dependent on the dosage of irradiation, supporting the role of *recA* in radiation resistance (Supplementary Figure S7). Based on this and our cluster analysis, which indicated significant PG_{4p} for *recA* (Figure 3), we first focused on *recA*. A closer analysis revealed presence of one/two distinct PG₄ motifs within 200 bases of the first gene start site in the *recA* operon in both *D. radiodurans* and *D. geothermalis*, respectively (Figure 4c and Supplementary Figure S8a). To directly test whether presence of NMM affects expression of *recA*, *D. radiodurans* was treated with either 25 or 50 nM of NMM. In presence of NMM, down-regulation of *recA* was clearly observed whereas expression of the 16S rRNA (endogenous control) gene remained unchanged (Figure 4d). Furthermore, we noted that inhibition of *recA* expression was dependent on the concentration of NMM; higher NMM levels resulted in relatively increased inhibition at both 5 kGy and 10 kGy irradiation. To further test whether the effect of NMM was due to G-quadruplex DNA binding, we treated *E. coli*-K12

MG1655, where *recA* is devoid of PG_{4p} motifs (Supplementary Figure S8b), with NMM. In contrast to *D. radiodurans*, we did not find any decrease in expression of *recA* in presence of 2 or 4 μ M NMM following irradiation (Supplementary Figure S9).

In case of *D. radiodurans* apart from *recA*, *recF*, *recO* and *recR* genes are also essential for radioresistance as demonstrated by greatly impaired growth in mutant strains devoid of these genes (42,58,59). Additionally, *recQ* mutants were also found to be radiation sensitive (60) though this has been contradicted by a recent study (58). Taking cue from our computational predictions, we independently analysed the promoters of *recF*, *recO*, *recR* and *recQ* operons and found multiple PG₄ motifs within 200 bp upstream of the first gene of operons (Figure 4e and Supplementary Figure S8a). Based on this we tested expression of all four genes in presence/absence of NMM following irradiation, though other than *recQ*, PG_{4p} of the *recF*, *recO* and *recR* promoters were not above the statistical threshold considered in our computational analysis. Interestingly, NMM significantly repressed

expression of all the four genes, *recF*, *recO*, *recR* and *recQ*, and not the endogenous control 16S rRNA gene, such that higher NMM levels resulted in relatively increased inhibition at both 5 kGy and 10 kGy irradiation (Figure 4f).

DISCUSSION

Herein independent lines of analyses revealed characteristics of promoter-G-quadruplex motifs that suggest role in specific functions. Not only did we find putative involvement in regulation of gene-classes related to selected functions, we also noted that the type of function was not analogous across organisms. On asking whether PG4 motif content of individual promoters (the PG4_P index)

is significantly associated with function, we first found that functional groups could be segregated based on PG4_P. Secondly, we noted with interest that this was specific for related group of species. In a complementary approach, using individual genes instead of functional classes, we again found that genes clustered in a fashion that was specific to organisms.

In order to experimentally test these predictions, we performed a case study using *D. radiodurans* and *D. geothermalis*, which have remarkable DNA repair systems that can withstand lethal doses of ionizing radiation (40,42,53). There are mainly two RecA-dependent recombinational DNA repair pathway in bacterial populations; the RecBCD and RecFOR pathways, which normally operates independently. The RecFOR pathway

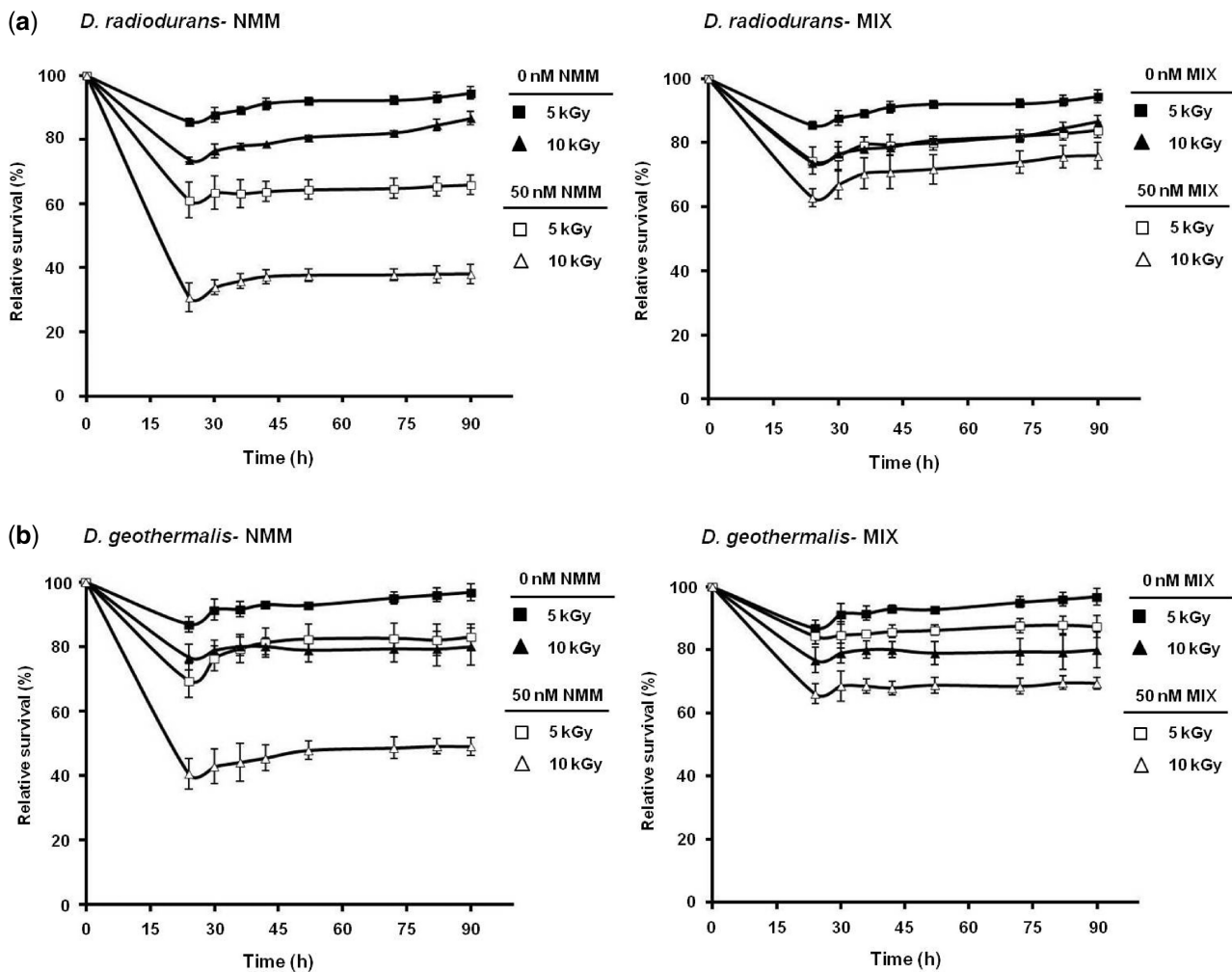


Figure 4. Quadruplex-binding ligands and radiation resistance. NMM attenuates growth of (a) *D. radiodurans* and (b) *D. geothermalis*, following exposure to gamma irradiation; *D. radiodurans* and *D. geothermalis* levels at 0 kGy were used to estimate relative survival following 24–90 h of post-irradiation recovery at 32°C in presence of 0 nM or 50 nM NMM (left panel) and MIX (right panel), respectively. Values are mean \pm standard error of three independent experiments. (c) Scheme showing PG4 motif positions within putative regulatory region of the *recA* operon. (d) Expression levels of *recA* and 16S rRNA genes in presence/absence of NMM in response to gamma irradiation (upper panel); quantification of relative intensities of the *recA* and 16S rRNA RT-PCR bands with respect to 0 nM NMM at respective radiation doses (lower panel). (e) Scheme showing PG4 motif in putative promoters of respective operons with *recF*, *recO*, *recR* and *recQ* genes. (f) Expression levels of *recF*, *recO*, *recR*, *recQ* and 16S rRNA genes in presence/absence of NMM in response to gamma irradiation (left panel); relative quantification of the *recF*, *recO*, *recR*, *recQ* and 16S rRNA RT-PCR results with respect to 0 nM NMM at respective radiation doses (right panel). All PCR assays were performed in triplicates and values are presented as mean \pm standard error; * denotes $P < 0.05$.

(continued)

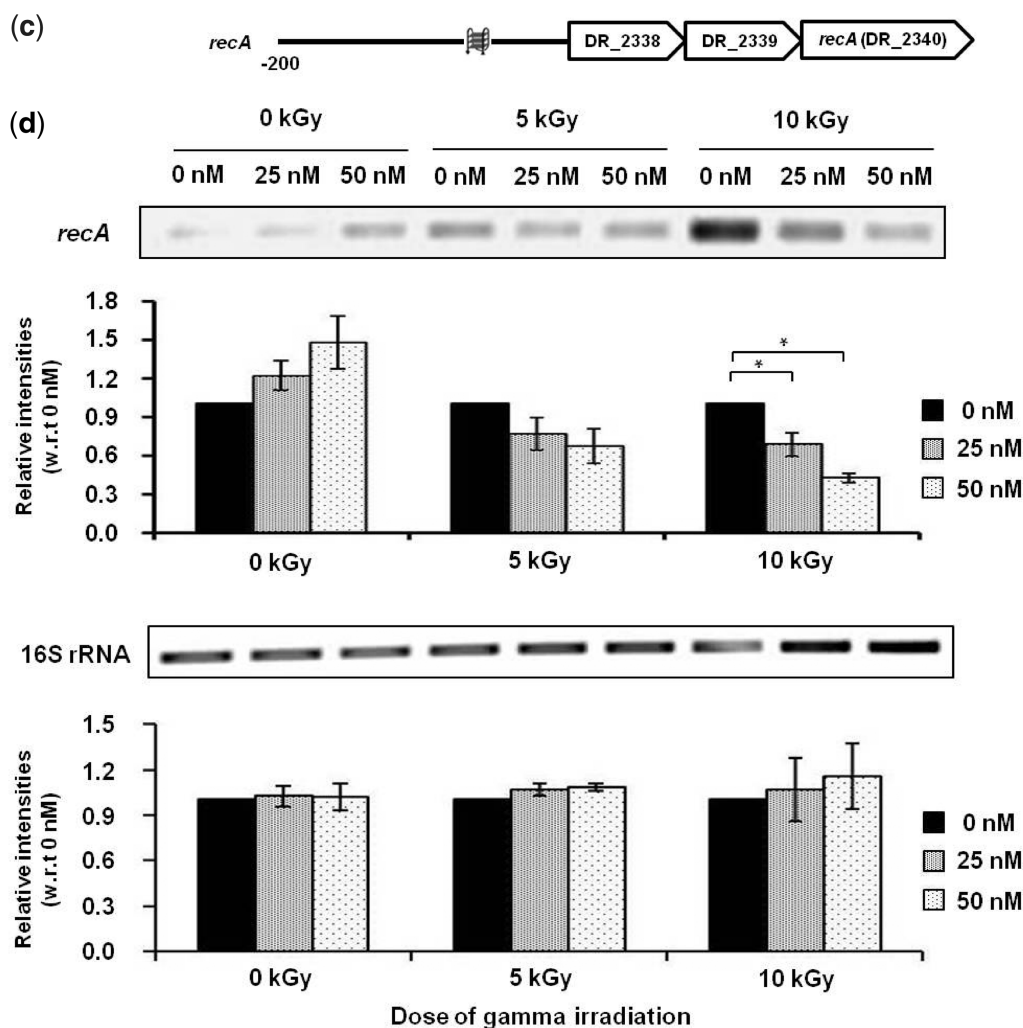


Figure 4. Continued.

comprising *recA*, *recF*, *recO*, *recR*, *recQ*, *recJ*, *recN*, *ruvA*, *ruvB* and *ruvC*, is used mainly during recombinational repair in *D. radiodurans*, as it lacks the RecBCD system like many other bacteria (41,46,59,61). Furthermore, *D. radiodurans* also does not encode homologs of the SbcB nuclease, which is an inhibitor of the RecF pathway (62). In support of our predictions we found that radiation resistance of *D. radiodurans* and *D. geothermalis* was attenuated by >50% in presence of 50 nM NMM, a ligand that is known to specifically bind G-quadruplex motifs inside cells. We also found that *recA*, *recF*, *recO*, *recR* and *recQ* genes have promoter-PG4 motifs and are repressed in a dose-dependent fashion in presence of NMM in *D. radiodurans*. Together, these findings support involvement of G-quadruplex-mediated regulatory mechanisms in radioresistance.

We observed relative increase in *recA*, *recF* and *recO* on NMM treatment in absence of irradiation. Though up-regulation of genes with promoter-quadruplex motifs has been reported in presence of NMM (22), this contrasts our observations related to NMM-mediated suppression of genes that confer radioresistance to *D. radiodurans*

following irradiation. On the other hand, in case of *E. coli*, which does not have any G-quadruplex motif in the *recA* proximal promoter, also we found enhanced expression of *recA* on NMM treatment in absence of irradiation (Supplementary Figure S9), though no change was detected following irradiation. Considered together with the specific intracellular G-quadruplex binding reported for NMM earlier (22,50), it is possible that the NMM-induced specific effects are observed on irradiation while the pre-irradiation changes are more general effects of the ligand treatment. It is also possible that the role of quadruplex motifs become more effective following irradiation, however further experiments will be required to test this speculation.

In an earlier study, we reported the genomic distribution of PG4 motifs in bacterial genomes. This provided first evidence for possible regulatory role of quadruplex motifs in any organism based on remarkable prevalence of PG4 motifs in bacterial promoters (7), which we subsequently found was a hallmark across more than 140 bacterial species (11). Furthermore, our genome-wide analyses also suggested that promoters with PG4 motifs

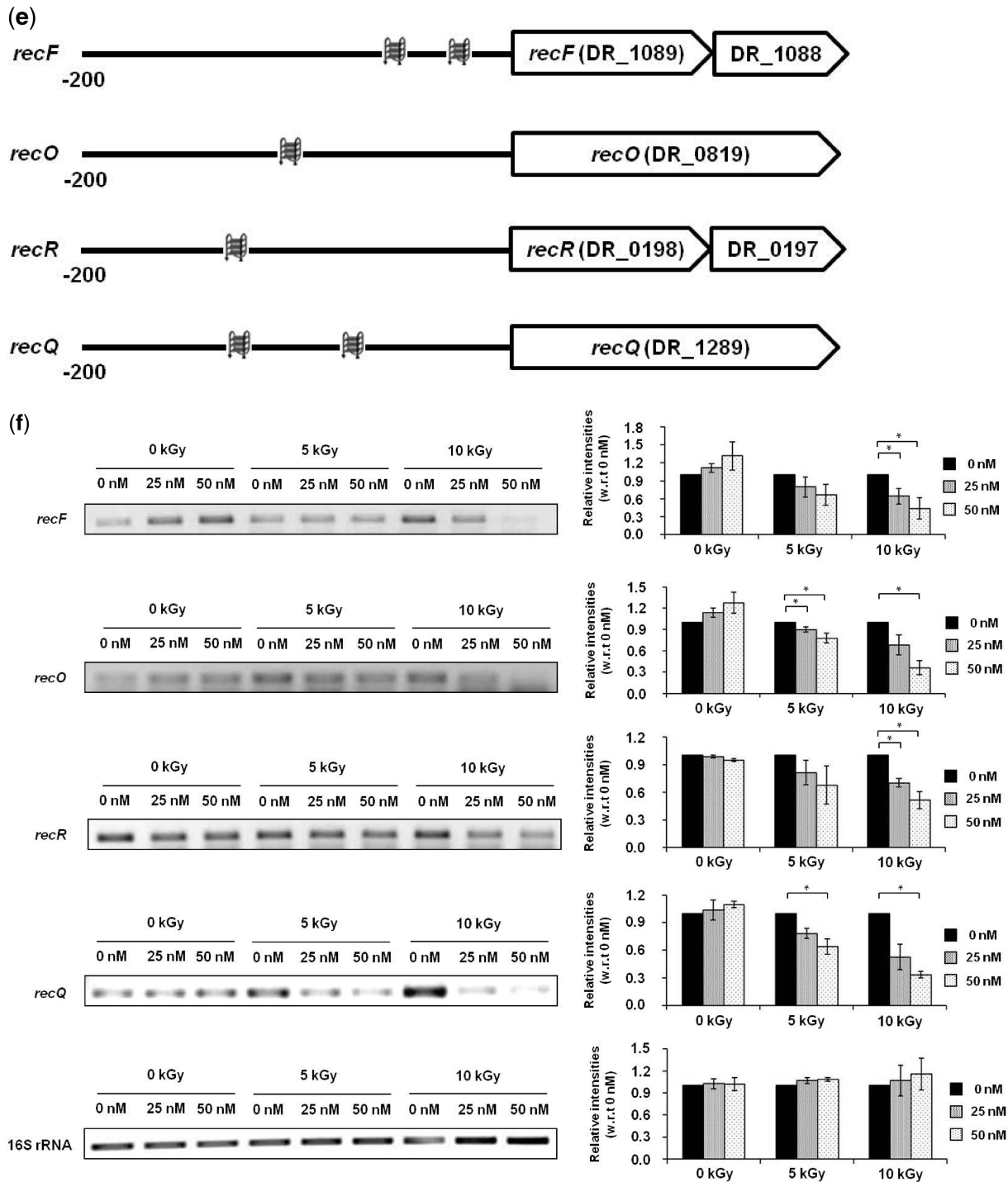


Figure 4. Continued.

in *E. coli* could be induced by supercoiling that leads to destabilization of the duplex DNA. In the current work, we have extended this initial finding to ask whether typical gene(s) pertaining to a particular functional class(es) have higher propensity for promoter PG4 content. This was done by focusing on relative estimation of PG4 motif content of promoters, both within and across several organisms, using the normalized index

PG4_p, which also allowed us to test the significance of PG4 motif occurrence in promoters across functional classes.

Despite arguments against the stability/formation of G-quadruplex motifs of stem size 2, we opted to consider them. This was primarily because in previous studies we have experimentally tested and found that such motifs randomly selected from *E. coli* readily adopt

G-quadruplex motifs in solution (7). Further support stems from work demonstrating that a two-tetrad non-canonical quadruplex motif regulates expression of human *thymidine kinase 1* (33) and recent work from Mergny group showing that G-quadruplex motifs with stem size of 2 remain stable *in vitro* (63). Moreover, one of the most studied quadruplex sequence, the thrombin aptamer constitutes of a stem size of 2 bases (64).

It is possible that radiation resistance involves PG4 motifs as a fortuitous connection due to high PG4_p content. By a similar analogy radioresistant species would be expected to cluster with *E. coli* when genes with high PG4_p were considered. This was not the case. Moreover, promoter-level analysis showed that PG4 motifs may have functional roles that were specific for at least a group of similar organisms. This also argues against a random association of PG4 motifs with radiation resistance genes. Furthermore, though unexpected, in addition to radioresistant species, we observed segregation of acidophiles when clustered for genes involved in radiation resistance. This further suggests that it is unlikely radioresistant species were found merely due to PG4-content. Finally when experimentally tested we found that indeed occurrence of G-quadruplex motif and its interaction with a specific G4-binding ligand impacts how *D. radiodurans* withstands radiation damage.

In summary, the notion of selective advantage prompted this study of promoter-wise PG4 motif content with the understanding that a finer analysis would provide distinct indication about possible PG4 motif-associated functions. This was first substantiated by cluster analysis of functional classes, where the distribution of promoters having significant PG4_p appeared to be non-random and associated to specific function(s). Secondly, a closer analysis of the cluster (Figure 2d) suggested an interesting pattern: groups of genes with high PG4_p appeared to be specific to related organisms. Keeping this in mind, we hypothesized that PG4 motif-function relationships may be interesting when considered with respect to functions that are specific to related organisms. Unbiased clustering of genes involved in radiation resistance segregated radioresistance organisms lending credence to this understanding. Taken together, based on these results, it is tempting to speculate that emergence of PG4 motifs as regulatory units not only influences function but also imparts directed advantage to counter environmental pressures. Further work addressing this possibility in different organisms and in relation to functional advantages acquired by the organism will be required to better understand this aspect of G-quadruplex function.

SUPPLEMENTARY DATA

Supplementary Data are available at NAR Online: Supplementary Table 1 and Supplementary Figures 1–9.

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