DNA-NUCLEAR MATRIX INTERACTIONS AND IONIZING RADIATION SENSITIVITY

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Ionizing radiation produces primarily deletion mutations in exposed mammalian cells (Breimer, 1988; DeMarini *et al.*, 1989; Sankaranarayanan, 1991). The kinetics of gene and chromosome mutation induction and the nature of the genetic changes suggest that these mutations develop from the recombination of two DNA double-strand breaks. Recombination requires DNA double-strand breaks to be in close spatial and temporal proximity. As chromosome structure will determine the proximity of DNA sequences, it should influence the types of recombinations that can occur. In addition, recent studies suggest that chromosome structure, and in particular, DNA supercoiling, will also influence recombination frequency.

Inherent Radiation Sensitivity and DNA Supercoil Unwinding

The DNA in chromosomes is organized in many different levels (Nelson *et al.*, 1986). At one level of organization, the DNA exists as a supercoiled loop attached to a protein structure, the DNA-nuclear matrix. When breaks are induced in this supercoiled loop, the loop relaxes and unwinds. Analyses of many different rodent and human cell lines (including Chinese hamster V79 cells [Olive, 1992; Gordon *et al.*, 1990], mouse L5178y cells [Kapiszewska *et al.*, 1989], human tumor cell lines [Schwartz and Vaughan, 1989], cell lines from patients with ataxia telangiectasia [Taylor *et al.*, 1991], and the radiosensitive Chinese hamster ovary [CHO] cell line xrs-5 [Schwartz *et al.*, 1990, in press]) have shown an association between inherent radiation sensitivity and the ability of these loops to relax and unwind. The evidence for this association comes from DNA neutral filter elution, alkaline unwinding, and nucleoid-based assays of DNA damage (reviewed in Olive, 1992).

The neutral filter elution assay (Bradley and Kohn, 1979) measures the amount of DNA which elutes from cells lysed on a filter. It is used to estimate DNA double-strand break frequency. The assumption is that the rate at which DNA elutes depends on the

size of DNA fragments and therefore on DNA break frequency. After low dose exposures, radioresistant cell lines show reduced rates and extents of DNA elution as compared to radiosensitive cells (Schwartz *et al.*, 1991; Olive, 1992), suggesting that radioresistant cells are more resistant to DNA double-strand break induction than sensitive cells. However, no difference in break induction is seen after high dose exposures, and other independent measures of DNA double-strand break induction usually fail to detect differences between radioresistant and radiosensitive cell lines in DNA break induction (Olive, 1992). Wlodek and Olive (1990) have suggested that the reduced rate of DNA elution seen after low dose exposures in radioresistant cell lines is due to some residual protein structure which limits DNA unwinding and therefore DNA elution under the conditions used in the assay. This model is supported by the observation that differences between resistant and sensitive cells in elution rates can be seen even in unirradiated samples (Schwartz *et al.*, 1991).

Evidence for constraints to DNA unwinding in radioresistant cells is also seen with the alkaline unwinding assay. In the alkaline unwinding assay (Ahnstrom, 1973), the amount of single-stranded DNA that forms when cells are lysed and denatured under alkaline condition is quantified. Single-stranded regions of the genome form when the DNA unwinds at sites of single-strand breaks. The relative amount of single-stranded DNA should therefore be directly related to the number of DNA single-strand breaks. Following radiation exposure, less single-stranded DNA is found in radioresistant cells than in radiosensitive cells (Olive *et al.*, 1986; Olive, 1992; Schwartz *et al.*, 1992) which would suggest fewer induced breaks. However, as mentioned above, other independent assays of DNA breaks find that sensitive and resistant cells do not differ in radiation-induced break frequencies (Olive, 1992). Therefore the reduced amounts of single-

stranded DNA in radioresistant cells must reflect some constraint to DNA unwinding in these cells. This constraint is also seen in unirradiated cells (Schwartz *et al.*, 1992).

Perhaps the best evidence for an association between radiation sensitivity and DNA supercoil unwinding comes from studies with nucleoids. Nucleoids are nuclei isolated such that most of the proteins have dissociated from the cell and what remains are intact supercoiled DNA loops attached to the nuclear matrix (Cook and Brazell, 1975; Cook *et al.*, 1976). DNA strand breaks induce relaxation and unwinding of the supercoils, which are seen as increases in nucleoid size. This can be detected by sedimentation assay (Cook and Brazell, 1975), microscopic analysis (Roti Roti and Wright, 1987), or flow cytometric analysis (Milner *et al.*, 1987). Smaller increases in nucleoid size are seen in radioresistant cell lines following ionizing radiation exposure than in radiosensitive cell lines (Gordon *et al.*, 1990; Kapiszewska *et al.*, 1989; Milner *et al.*, 1987; Vaughan *et al.*, 1992). Because radiosensitive and resistant cell lines are equally sensitive to break induction, the smaller increase in nucleoid size must reflect some constraint to unwinding in resistant cells. The structure responsible for the attachment of DNA to the nuclear matrix, the matrix attachment region (MAR), may be the structure controlling the extent of DNA unwinding, as differences in the nature of the MAR in resistant and sensitive cells have been reported (Schwartz and Vaughan, 1989; Vaughan *et al.*, 1991).

DNA-Nuclear Matrix Attachment and DNA/Chromosome Break Repair

Limiting DNA unwinding may facilitate the repair of DNA strand breaks by keeping broken ends in closer proximity for a more rapid and accurate rejoining. The radioresistant cell lines mentioned above all rejoin DNA and chromosome breaks more rapidly than sensitive cells (Schwartz *et al.*, 1988; Schwartz and Vaughan, 1989). This has been demonstrated with a variety of biochemical- and cytogenetic-based assays. The effect is only on the kinetics of break rejoining, as there is no evidence for a reduced

capacity to rejoin breaks in most of these radiosensitive cells. The rate of rejoining also appears closely linked to the frequency of radiation-induced intra- and interchromosome exchanges. Radioresistant cell lines show smaller frequencies of interstitial chromosome deletions, dicentrics, and rings following radiation exposure. Thus, the constraint to DNA unwinding is associated with faster rates of DNA/chromosome break rejoining and reduced frequencies of intra- and interchromosomal exchanges in radioresistant cells.

The repair of DNA double-strand breaks is a complex process involving both exonuclease and DNA polymerase activities. One early event that follows break induction is the modification of proteins near the break site, which opens up the chromatin structure and initiates unwinding of DNA supercoiled loops. This unwinding may extend to adjacent loops (Mullenders *et al.*, 1983). An analogous situation is found in transcribing regions of the genome where RNA polymerases drive DNA unwinding.

Recently, it has been shown that the MAR in actively transcribing regions of the genome may exist in a base-unpaired or cruciform-like structure (Kohwi-Shigematsu and Kohwi, 1990; Leonard and Patient, 1991; Bode *et al.*, 1992). This structure tends to be more resistant to endonuclease digestion and has a greater affinity for the nuclear matrix. It has been suggested that this structure insulates transcribing regions of the genome from adjacent nontranscribing regions by preventing the unwinding of adjacent loops. Their open loop structure may also provide sites for transcription factors and polymerases to bind, and the torsional energy stored at these sites may be useful for driving gene transcription.

This open loop MAR structure could also act to insulate a radiation-damaged loop from the rest of the genome, preventing the unwinding of adjacent loops and helping to maintain DNA ends in closer proximity. The closer proximity might lead to faster break rejoining and fewer intra- and interchromosomal exchanges. These open loop MAR

configurations might also provide a recognition site at which repair proteins collect, which would facilitate repair by placing the required proteins close to the damaged site, and the torsional energy stored at this site could drive some of the enzymatic repair reactions as well.

This model suggests that differences in radiation sensitivity should be associated with alterations in gene expression. In fact, in the cell systems mentioned above, radiation sensitivity is just one aspect of a complex phenotype that distinguishes resistant and sensitive cell lines. Such a model also suggests that actively transcribed regions of the genome should be more resistant to radiation-induced change than nontranscribed regions. And, as the MARs are dynamic structures showing both tissue-specific and transformation-related alterations (Pienta and Coffey, 1992; Dickinson *et al.*, 1992), one would predict that tissue-specific and transformation-related alterations in radiation sensitivity will reflect MAR alterations.

Conclusion

The association between inherent ionizing radiation sensitivity and DNA supercoil unwinding in mammalian cells suggests that the DNA-nuclear matrix attachment region (MAR) plays an important role in radiation response. In radioresistant cells, the MAR structure may exist in a more stable, open configuration, limiting DNA unwinding following strand break induction and maintaining DNA ends in close proximity for more rapid and accurate rejoining. In addition, the open configuration at these matrix attachment sites may serve to facilitate rapid DNA processing of breaks by providing (1) sites for repair proteins to collect and (2) energy to drive enzymatic reactions.

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