SUPPORTING INFORMATION

The *Bis*-Electrophile Diepxybutane Cross-links DNA to Human Histones but Does Not Result in Enhanced Mutagenesis in Recombinant Systems

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Chem. Res. Toxicol. 22, 000-000 (2009)

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Figure S3. Cross-linking of purified histone H2b to single-stranded or double-stranded oligonucleotides by diepoxybutane.

Histone Protein	Uniref100 ID	% Coverage
Histone H2A.m	P04908	41
Histone H3/b	Q93081	13
Hisonte H4	Q6FGB8	17
Histone protein	Q5R2W0	27
Histone 1, H2aj	Q5JXQ5	14
Histone 4	Q4A487	22
Core histone macro-H2A.	1 075367	11
Histone protein	Q5R2W0	38
Histone H3/b	Q93081	20
Histone H4	Q6FGB8	50
Histone H2A.m	P04908	36

Table S1. Histone proteins identified in two independent screens for cross-link candidates.





Figure S2. Mutagenicity (S2A) and survivorship (S2B) in TRG8 cells (expressing AGT or histone H2b or containing pINIII vector treated) with 1,2-dibromoethane for 30 min at 37 °C before growing the cells on his^+ and his^- plates.



Figure S3. Cross-linking of purified histone H2b to single-stranded or double-stranded oligonucleotides by diepoxybutane. Gel shift assays were performed by incubating histone H2b $(1 \ \mu g)$ with ³²P-5'-endlabeled 16-mer single-stranded or double-stranded oligonucleotides in reactions containing various concentrations of diepoxybutane overnight at 37 °C. Samples were separated by SDS-polyacrylamide gel (15% w/v) electrophoresis and DNA-protein cross-links were detected via autoradiography and quantified using Quantity One software (BioRad).