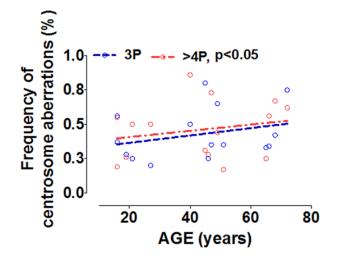
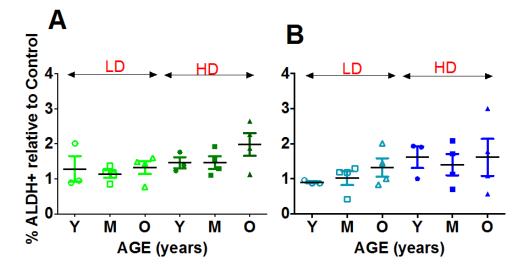
SUPPLEMENTARY MATERIAL



Supplementary Figure S1. Centrosome aberrations in control samples. The fraction of cells with 3 pericentrin foci (3P) and \geq 4 pericentrin foci (\geq 4) were plotted in unexposed cells for all strains as a function of age. Regression analysis was carried out to define the relationship between the frequency of cells with centrosome aberrations and age in unirradiated cells.



Supplementary Figure S2. Comparison of stem cell numbers in strains of various age groups. S/P cells were assessed in HMEC strains based on flow cytometry assessment of ALDH+ signal. The fraction of ALDH+ cells relative to the unexposed control was assessed. Strains were sub-grouped into young, middle and old represented by circles, rectangles and triangles, respectively. Filled symbols represent the high dose and open symbol represent the low dose. The frequency of S/P cells relative to control was compared between the different groups. Error bars represent SD within each age group. Data are based on two independent experiments for high and low dose. Data from both Cesium exposure (A) and Ti ion exposure (B) are presented. A slight age-dependent increase in S/P cell populations is noted with a low dose of Ti and a high dose of Cs.

Supplementary Table S1. Correlations between various endpoints.

Markers	CA	ALDH+	pATF2 (Ser ^{490/498})
CA		+ (p<0.05)	-
ALDH+	+ (p<0.05)		-
pATF2 (Ser ^{490/498})	-	-	
p38 (Thr ¹⁸⁰ /Tyr ¹⁸²)	+	-	+
p70 S6K (Thr ⁴¹²)	+	-	+
pAKT(Ser ⁴⁷³)	+	-	+
pCREB(Ser ¹³³)	+	-	+
pERK/MAPK(Thr ¹⁸⁵ /Tyr ¹⁸⁷)	+	-	+
pJNK (Thr ¹⁸³ /Tyr ¹⁸⁵)	-	-	+
pNFκβ (Ser ⁵³⁶)	-	-	+
pSTAT3(Ser ⁷²⁷)	-	-	+
pSTAT5 A/B (Tyr ^{694/699})	-	-	+
TGFRβII	+	+	+
pSMAD2 (Ser ⁴⁶⁵ /Ser ⁴⁶⁷)	+	+	+
pSMAD3 (Ser ⁴²³ /Ser ⁴²⁵)	-	-	+
SMAD4	+	+	+ (p<0.05)

Pearson's correlation analysis

Pearson's correlation was used to examine the association between centrosome aberration (CA) frequency, ALDH+ signal, persistent pATF2 foci, and the relationship of these parameters to the levels of various phospho-proteins in the stress activated signaling pathways.