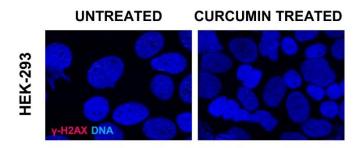
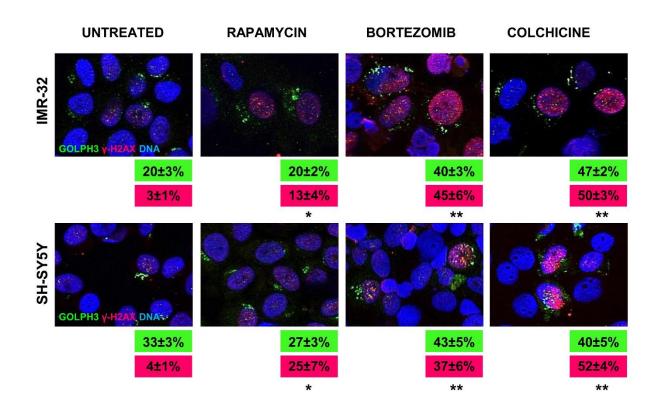
Supplementary Material, Ognibene M. et al.



Supplementary Figure 1. Immunofluorescence analysis of non tumoral control HEK-293 cells cultured with 20 μ M curcumin 48 hours using anti- γ H2AX (red). Cells were counterstained with DAPI to visualize nuclei (blue). Untreated cells were cultured with 0.1% DMSO. (Magnification 40x).



Supplementary Figure 2. Immunofluorescence analysis of IMR-32 and SH-SY5Y cells cultured respectively with 300 nM or 500 nM Rapamycin, with 2 nM or 3.5 nM Bortezomib and with 0.02 μ M or 0.05 μ M Colchicine for 48 hours using anti-GOLPH3 (green) and anti- γ H2AX (red) and compared to untreated cells. Cells were counterstained with DAPI to visualize nuclei (blue). (Magnification 40x). In green and red boxes are reported the percentages of GOLPH3 and γ H2AX positive cells respectively. Data are representative of three independent experiments ± S.D.