Supplementary Figure S4



Supplementary Figure S4. Loss of ATR induces DNA damage in APOBEC3A-expressing cells. A. Levels of γH2AX in A3A^{WT} or A3A^{E72A}-expressing cells transfected with ATR siRNA were analyzed by Western blot. **B.** Inducible A3A^{WT}-expressing cells were left uninduced or induced with DOX, treated with ATRi (3 µM) for 8 h, and immunostained for γH2AX and PCNA. Representative images are shown. **C.** Quantification of γH2AX intensity in 4,000 cells shown in B. Cells were colored according to the intensity of γH2AX staining. The same cells and quantifications were also used in Figure 1E. **D.** Quantification of γH2AX and TUNEL intensities in 5,000 inducible A3A^{WT}-expressing cells induced with DOX and treated with ATRi (3 µM) for 8 h. TUNEL positive cells were shown in red. **E-F.** U2OS-derived cells were induced to express A3A^{WT} or left uninduced. Cell survival was analyzed after 5 days of continuous Chk1i treatment (E), or after 1 day of Chk1i treatment followed by a 3-day release (F). Error bar: S.D. (n=3)