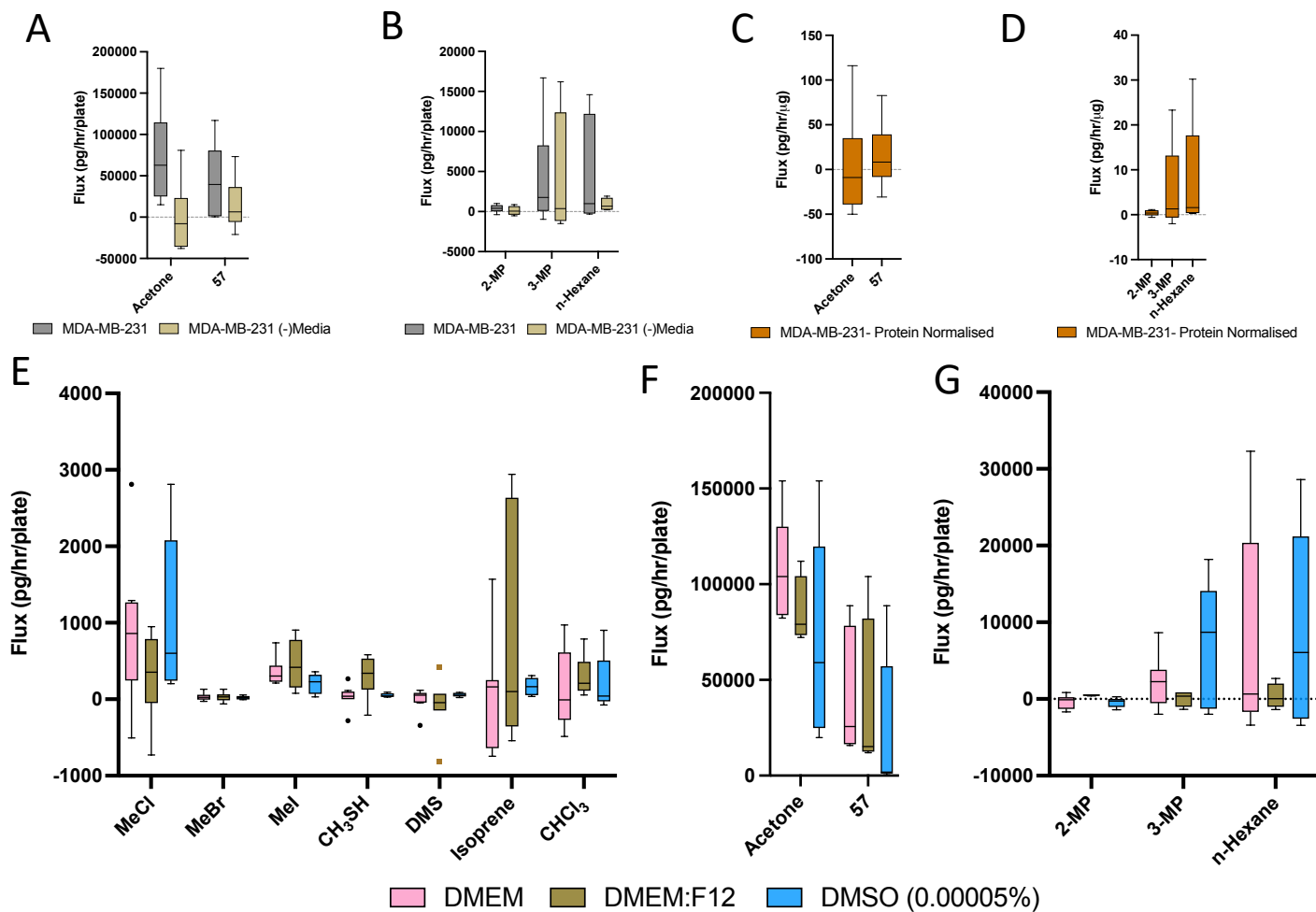
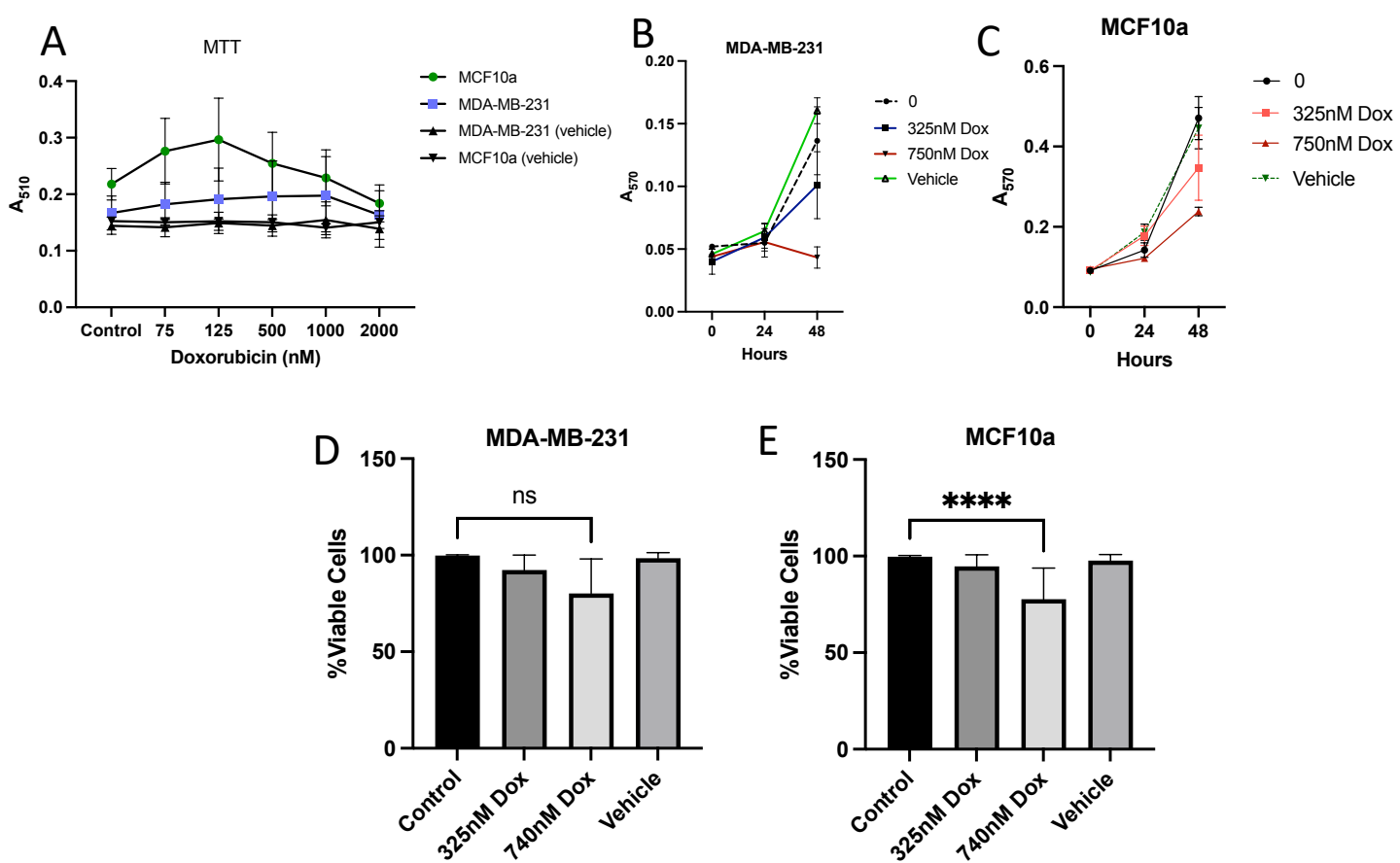




Supplementary figure 1. Cellular volatiles and media backgrounds. Volatiles from cellular headspace vs cellular headspace with media control deducted (**A, B**). Media subtracted and protein normalised VOC flux for MCF10a (n = 9); MCF7 (n = 4); MDA-MB-231 cells (n = 6) (**C,D**). Volatiles released from media alone. DMEM (n = 6), DMEM:F12 (n = 4), DMSO addition (n = 6) (**E, F, G**). CHCl₃ = Chloroform, DMS = Dimethyl sulfide, MeBr = Methyl bromide, MeCl = Methyl Chloride, MeI = methyl iodide, MeSH = Methanoethiol Boxplot whiskers show median ± Tukey distribution. ANOVA followed by Tukey or Bonferroni post hoc test was performed



Supplementary figure 2. Doxorubicin treatment of MDA-MB-231 and MCF10a. **(A)** MTT assay of varying concentrations of doxorubicin for both MDA-MB-231 and MCF10a. Relative levels of vehicle (DMSO) are provided at each stage (mean $\pm$ SEM; $n = 3$). **(B, C)** Sulforhodamine B assay over time for with doxorubicin treatment (DOX) and vehicle (DMSO, 0.00008%) for MDA-MB-231 and MCF10a cells (mean $\pm$ SEM; $n = 3$). **(D, E)** Trypan blue exclusion assays following 24hr doxorubicin treatment and vehicle (DMSO, 0.00008%) for MDA-MB-231 and MCF10a. ANOVA followed by Bonferroni post hoc test was performed; **** $p < 0.0001$.