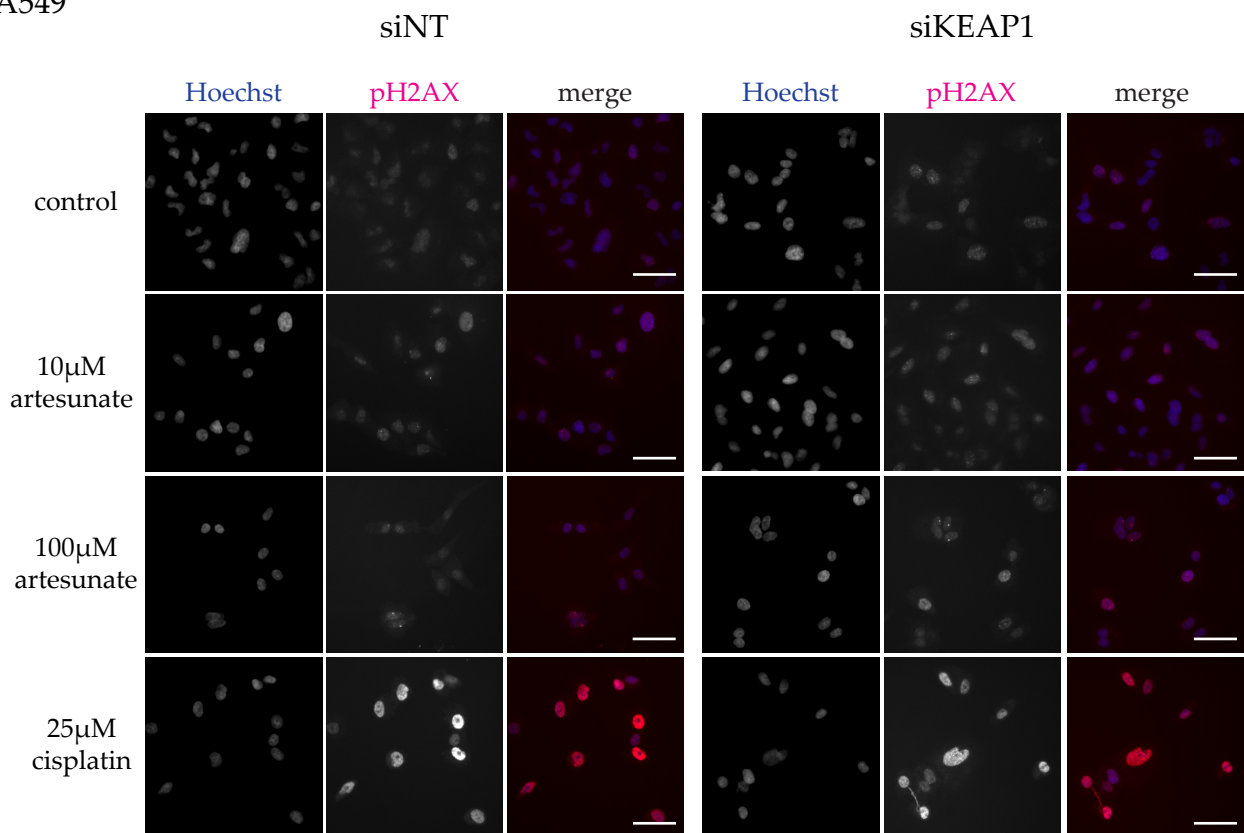
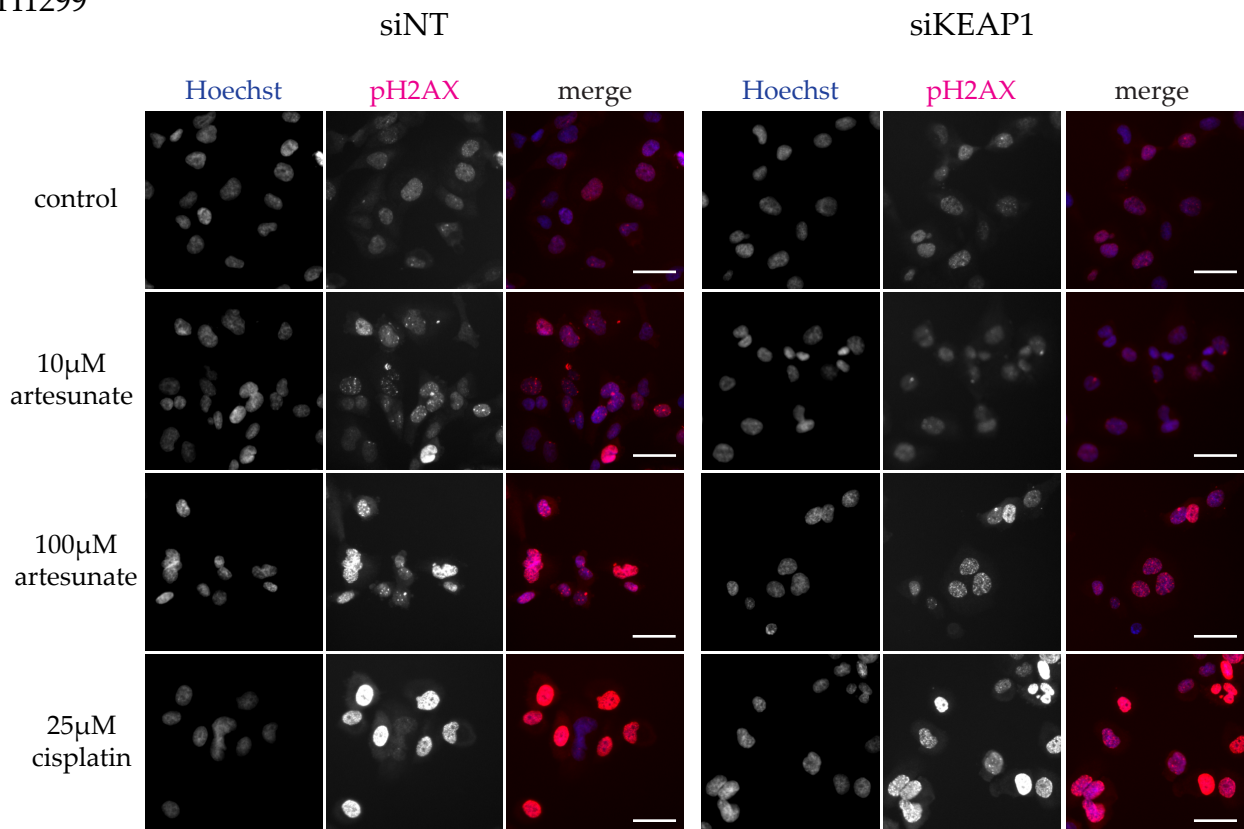


(a) A549

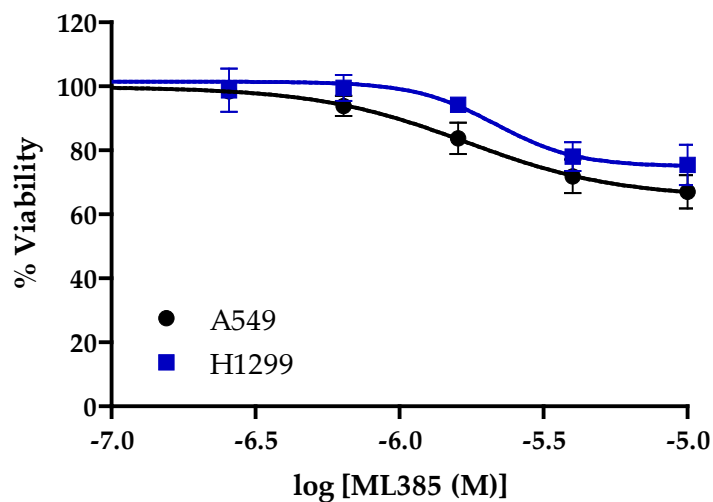


(b) H1299

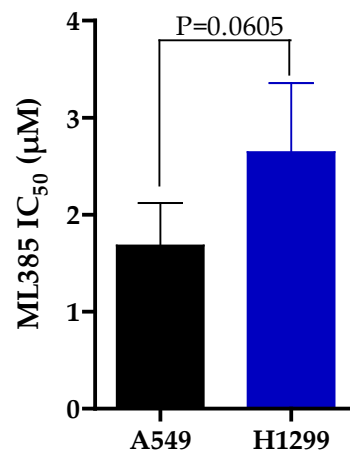


Supplemental Figure 1: Representative images of pH2AX staining in A549 (a) and H1299 cells (b) that had been transfected with non-targeting (siNT) or KEAP1 (siKEAP1) specific siRNA following 24 hour treatment with 0.1% DMSO (control), 10 μ M artesunate, 100 μ M artesunate, or 25 μ M cisplatin (bar = 50 μ m).

(a)

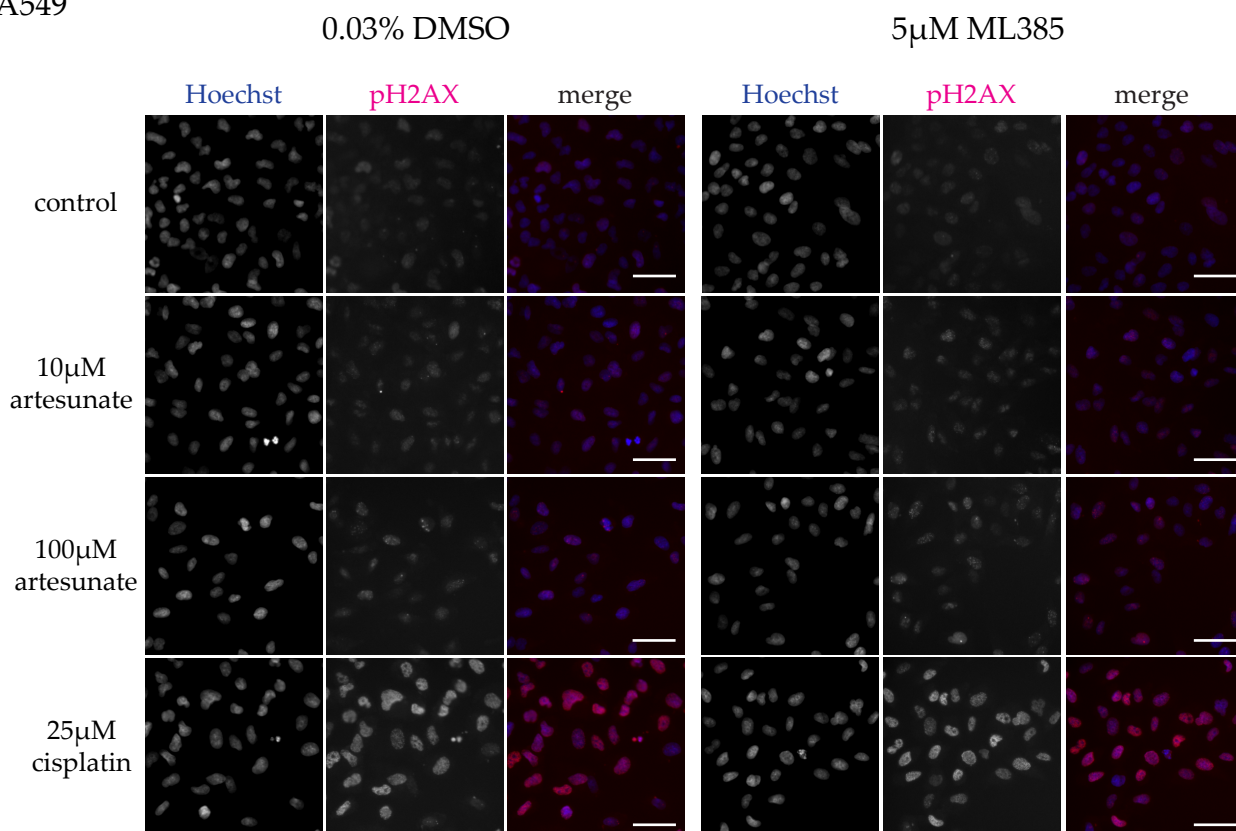


(b)

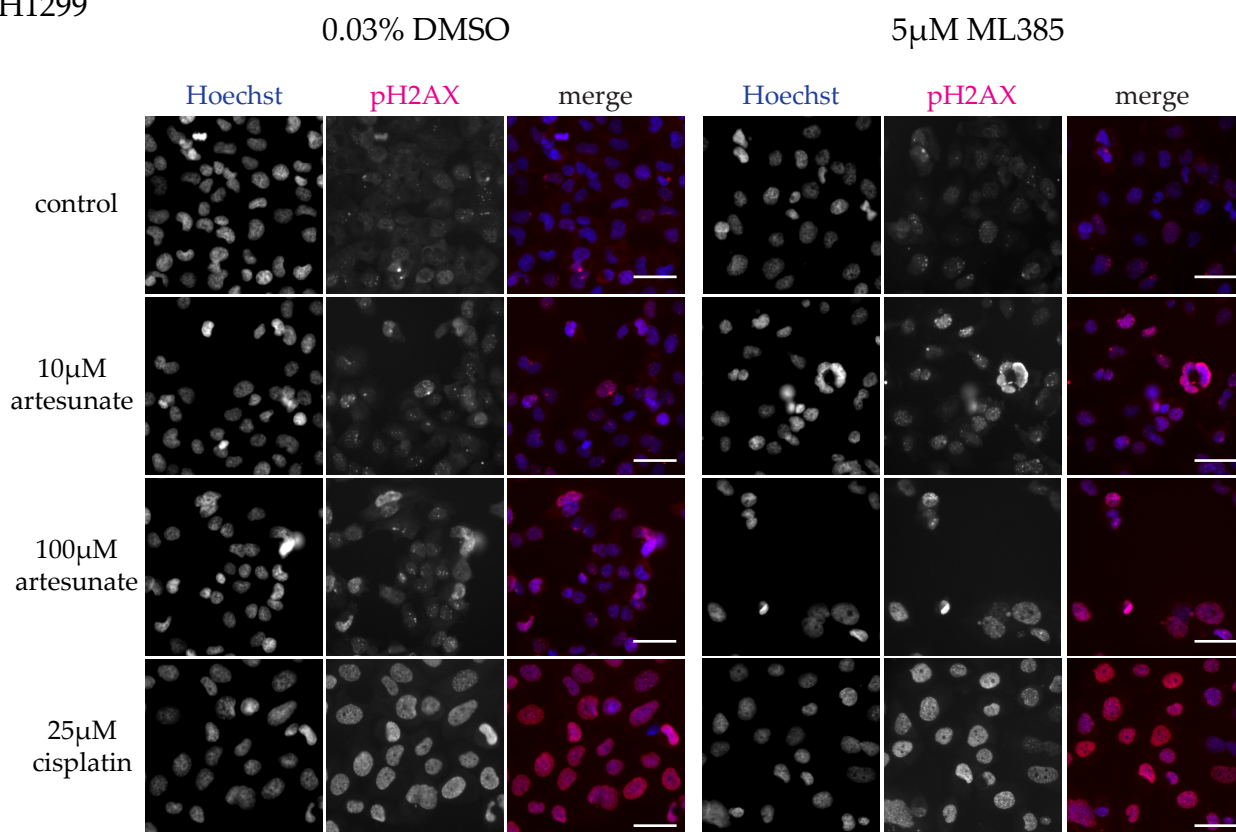


Supplemental Figure 2: ML385 (NRF2 inhibitor) has minimal effect of viability of A549 and H1299 NSCLC cell lines as a single agent (a) Cells were treated with a serially diluted concentrations of ML385 for 96hr. Each cell lines was normalized to cells treated with 0.1% DMSO as a control and is graphed as the mean \pm SD, $n=4$. (b) The mean IC₅₀ of ML385 in each cell line is graphed \pm SD. P-value was calculated using a two-tailed t-test comparing the mean IC₅₀ for each cell line tested.

(a) A549

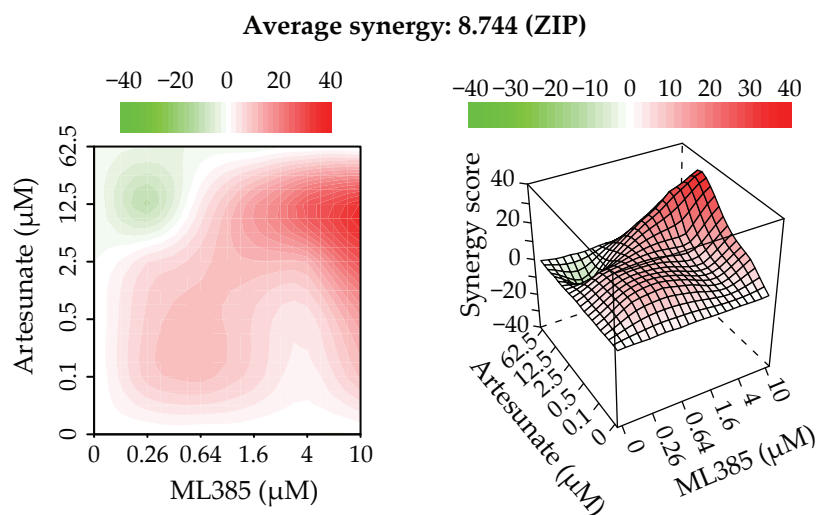


(b) H1299

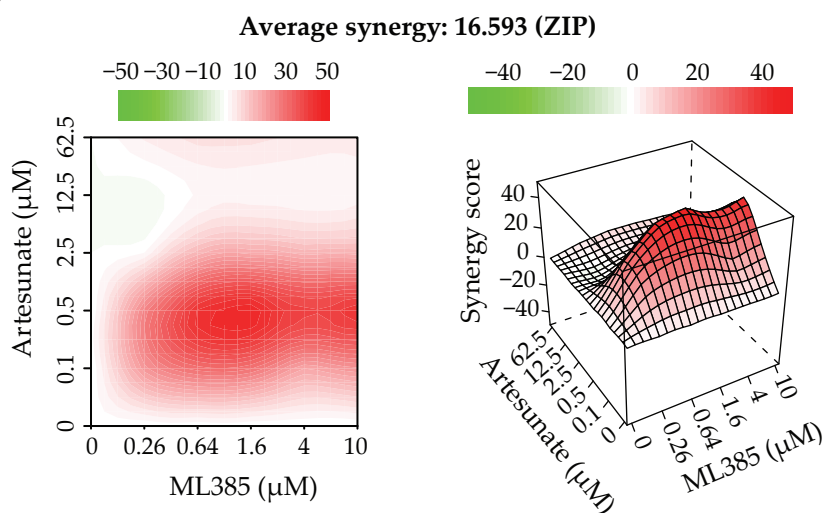


Supplemental Figure 3: Representative images of pH2AX staining in A549 (a) and H1299 cells (b) that had been treated with 0.03% DMSO or 5 μ M ML385 for 24 hour prior to treatment with 0.1% DMSO (control), 10 μ M artesunate, 100 μ M artesunate, or 25 μ M cisplatin for 24 hours (bar = 50 μ m).

(a)



(b)



Supplemental Figure 4: ZIP model demonstrating synergy between artesunate and ML385

Graphic representation of the ZIP model of synergy scoring as calculated by using a 6x6 dose-response matrix in A549 (a) and H1299 (b) cells. Red color indicates synergy, while green indicates antagonism between the drug combinations tested.