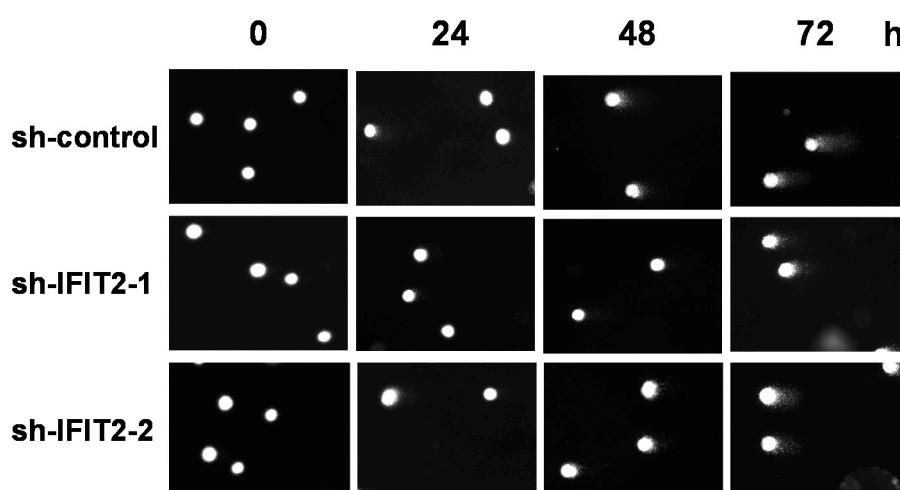


# Supplementary Materials: SAHA Overcomes 5-FU Resistance in IFIT2-Depleted Oral Squamous Cell Carcinoma Cells

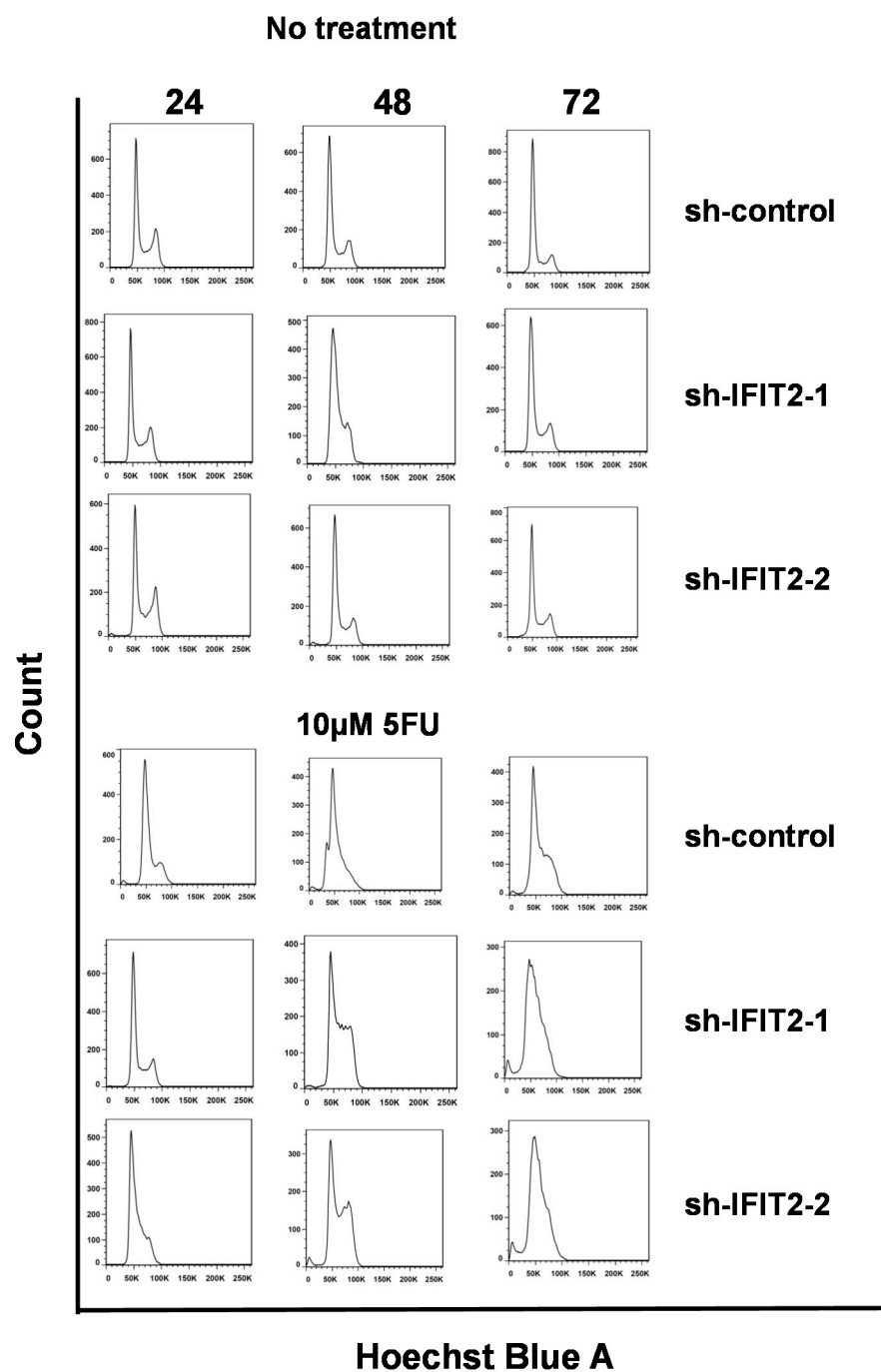
Prabha Regmi, Kuo-Chu Lai, Chung-Ji Liu and Te-Chang Lee

**Table S1.** Primer sequences for quantitative real-time PCR analysis.

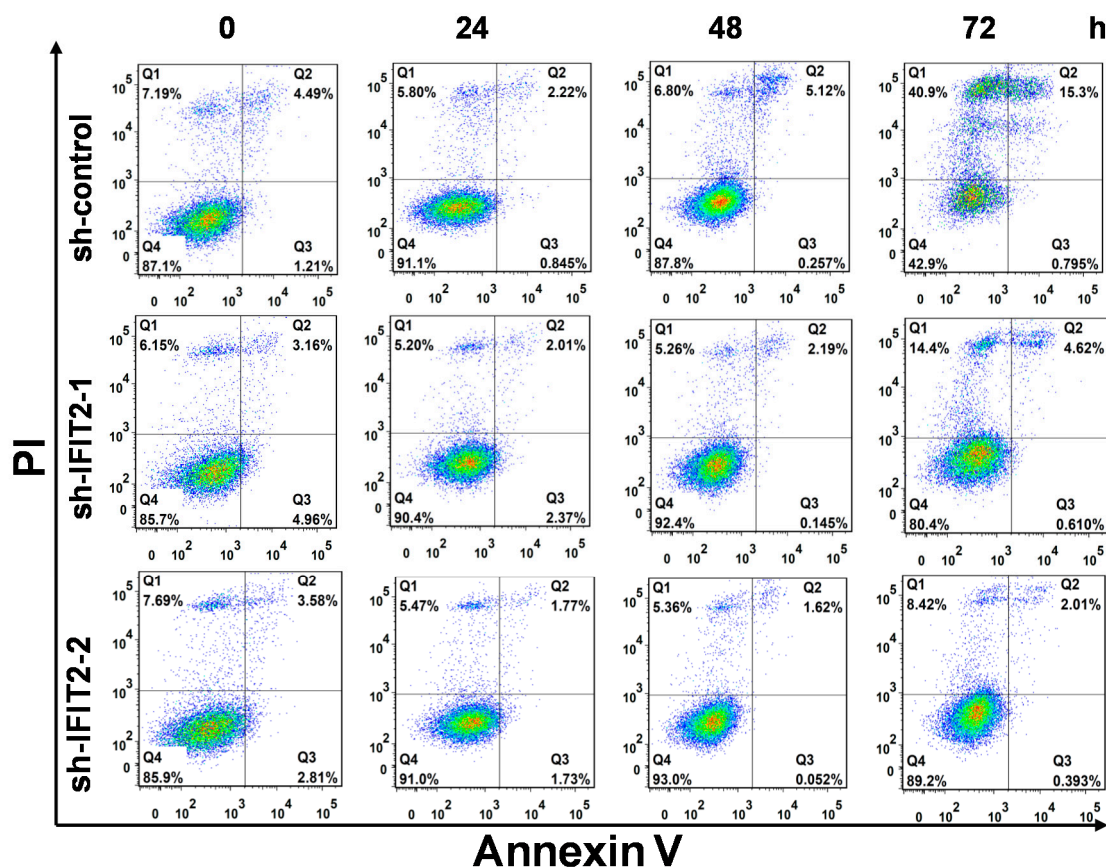
Gene	Forward Primer	Reverse Primer
TS	5'-GGAAGGGTGTTTTGGAGGAGTT-3'	5'-AGATTTTCACTCCCTTGGAAGACA-3'
TP	5'-GCTGGAGTCTATTCTGGAT-3'	5'-ACTGAGAATGGAGGCTGTGATG-3'
OPRT	5'-GCGTCTTCTGAGTCAGGTTG-3'	5'-GCATCTGCTAGCTGCAACAG-3'
TK1	5'-GTTTTTCCCTGACATCGTGGA-3'	5'-CGAGCCTCTTGGTATAGGCG-3'
GAPDH	5'-GGAGTCCCTGCCACACTCA-3'	5'-GCCCCTCCCCTCTTCAAG-3'



**Figure S1.** Representative images of Comet assay. Growing sh-control, sh-IFIT2-1 and sh-IFIT2-2 cells were treated with 10mM 5FU for 12, 24, 48 and 72 h. At the end of treatment, the cells were harvested and subjected to a comet assay as described in the Materials and Methods.



**Figure S2.** Representative histograms of cell-cycle analysis. Growing sh-control, sh-IFIT2-1 and sh-IFIT2-2 cells were treated with 10 µM 5-FU for 24, 48 and 72 h. and the cell-cycle analysis was performed using the Hoechst dye as described in methods.



**Figure S3.** Representative image of annexin V staining for apoptosis analysis using flow cytometry. Cells were treated with 5-FU for 24, 48 and 72 h, stained with annexin V-APC and PI, and subjected to flow cytometric analysis. Q1, Q2, Q3, Q4 represent necrosis, late apoptosis, early apoptosis and live cells respectively.

Figure 1

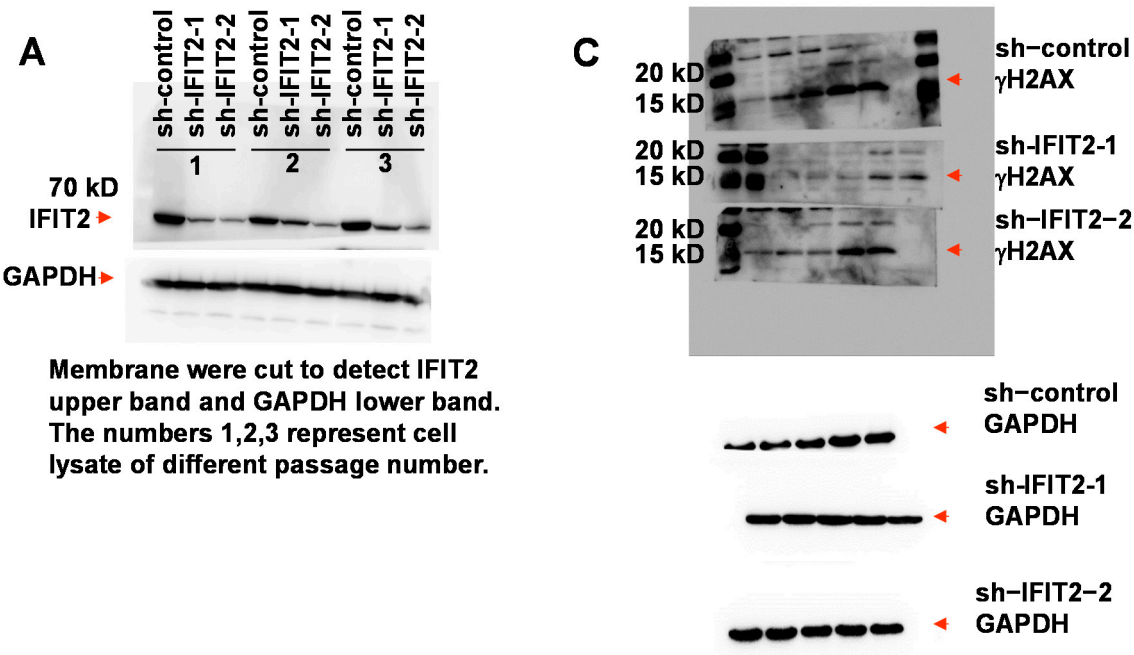


Figure 2B

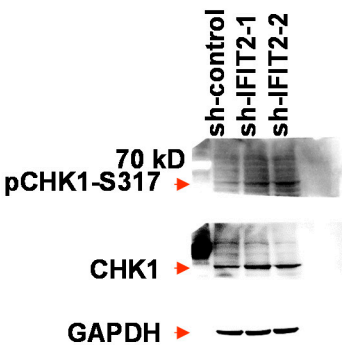
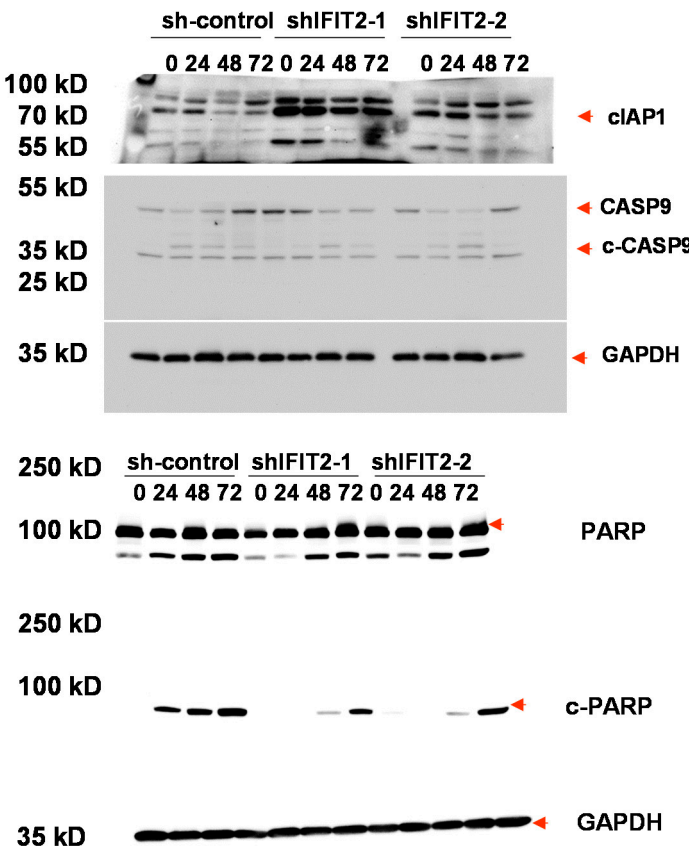
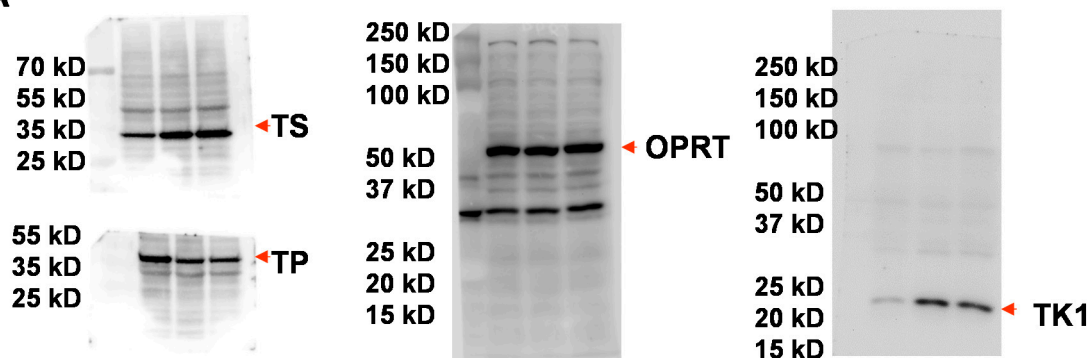
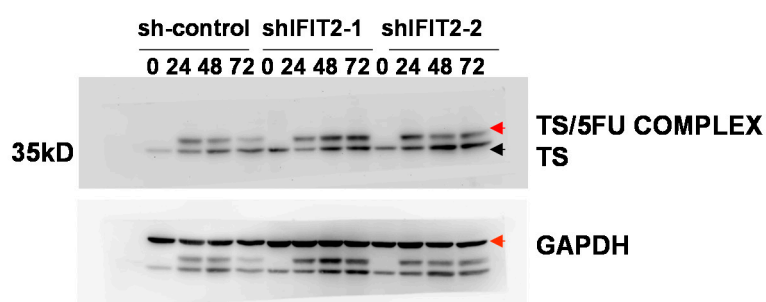
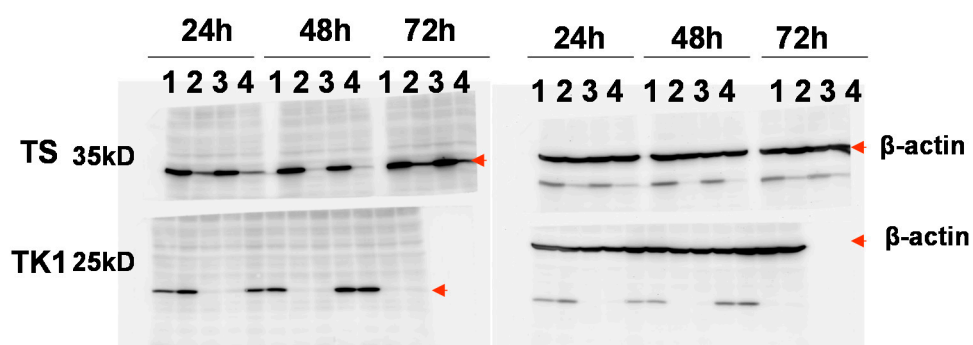


Figure 3C



**Figure 4****A****B****Figure 5A**

Label 1,2,3,4 represent different siRNA used as follows:

1. CONTROL siRNA

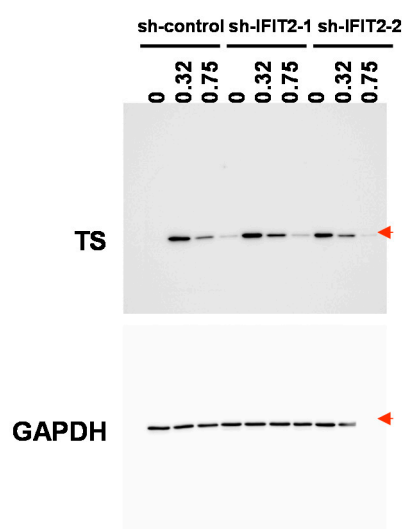
2. TS siRNA

3. TK1 siRNA

4. TS+TK1 siRNA

24h panel represents transfection efficiency after 24h

## Figure 7A



**Figure S4.** Original images of Western blots.

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