## Figure S3

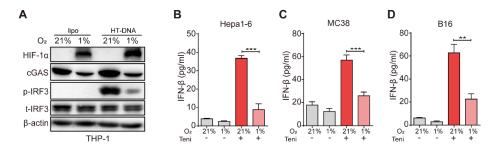


Figure S3. Hypoxia inhibited cGAS/STING-mediated IFN-I signaling in multiple tumor cell lines.

(A) THP-1 cells were cultured in normoxic (21%  $O_2$ ) or hypoxic (1%  $O_2$ ) environment for 18 h, then the cells were transfected with herring testes (HT)-DNA (5  $\mu$ g/ml), followed by continuous normoxic or hypoxic culture for another 6 h, the protein expression of cGAS and p-IRF3 were detected by immunoblotting,  $\beta$ -actin was used as a loading control. (B-D) Murine tumor cells were treated with teniposide at 20  $\mu$ M, followed by either normoxic or hypoxic culture for 24 h, then the supernatant IFN- $\beta$  of Hepa1-6 (B), MC38 (C), B16 (D) cells were measured by ELISA. Data in (A) are representative of 3 independent experiments. Data in (B, C, D) are shown as mean  $\pm$  SD of 3 independent experiments. \*:P < 0.05, \*\*:P < 0.01 and \*\*\*:P < 0.001.