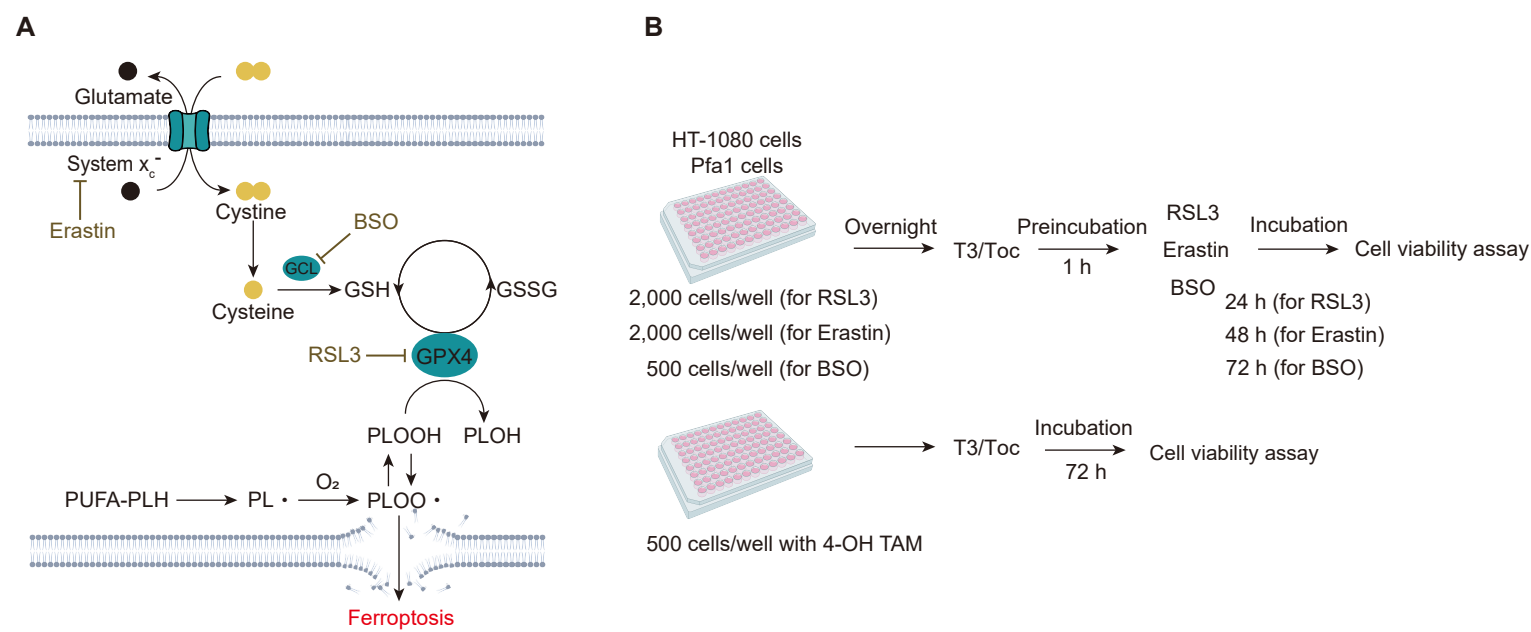


Supplementary Figure 1



Supplementary Figure 1. Schematic illustration of ferroptosis-inducing mechanisms and experimental workflow.

(A) Ferroptosis-inducing mechanisms. Erastin inhibits the cystine/glutamate antiporter (system X_c^-), reducing intracellular cystine and depleting glutathione (GSH). BSO inhibits glutamate–cysteine ligase (GCL), the rate-limiting enzyme in GSH synthesis. RSL3 directly inhibits GPX4.

(B) Schematic illustration of experimental workflow.

Supplementary Figure 2

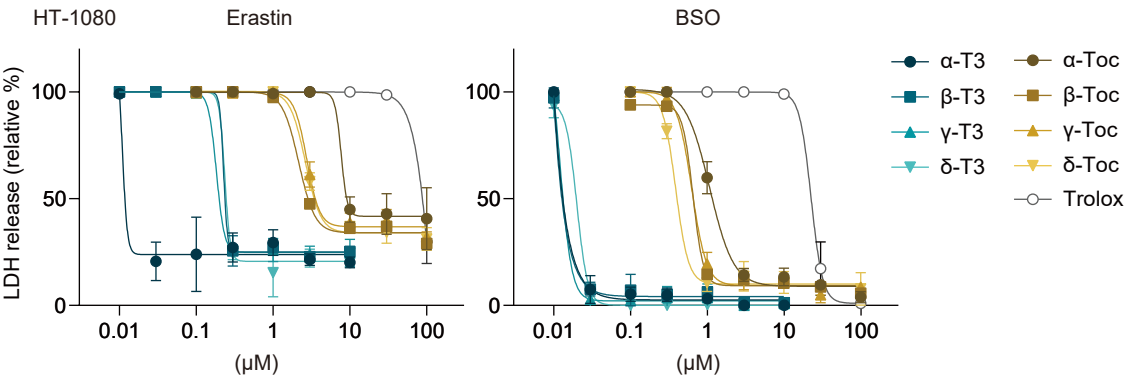
A

RSL3									
	α -T3	β -T3	γ -T3	δ -T3	α -Toc	β -Toc	γ -Toc	δ -Toc	Trolox
HT-1080	0.03	0.03	0.05	0.15	45.5	6.96	6.92	12.1	7.49
Pfa1	0.12	0.12	0.13	0.25	43.6	8.60	10.0	8.67	24.9

Erastin									
	α -T3	β -T3	γ -T3	δ -T3	α -Toc	β -Toc	γ -Toc	δ -Toc	Trolox
HT-1080	0.45	0.51	0.51	0.52	4.67	3.27	3.43	3.41	59.4
Pfa1	0.20	0.43	0.31	0.35	3.73	2.21	2.80	2.52	36.3

BSO									
	α -T3	β -T3	γ -T3	δ -T3	α -Toc	β -Toc	γ -Toc	δ -Toc	Trolox
HT-1080	0.11	0.18	0.15	0.30	0.83	0.46	0.46	0.33	10.0
Pfa1	0.18	0.41	0.22	0.38	3.74	1.72	1.78	0.69	30.6

B

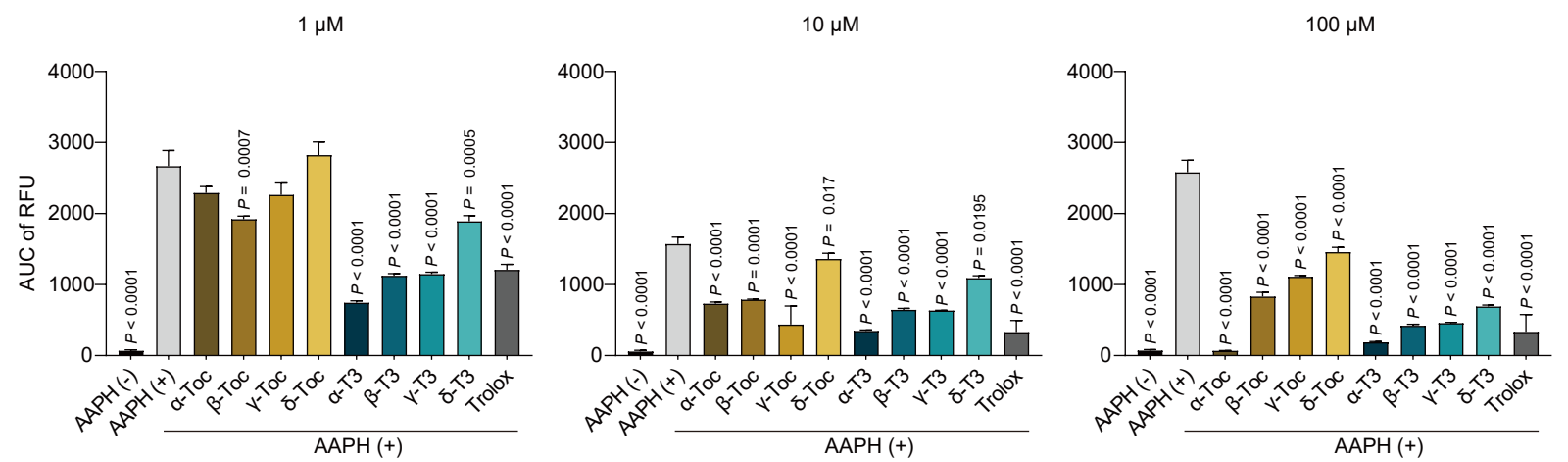


Supplementary Figure 2. EC₅₀ values of vitamin E analogs under ferroptosis-inducing conditions and LDH assay.

(A) EC₅₀ (μ M) values of each vitamin E analog for preventing ferroptosis induced by RSL3, erastin, and BSO based on cell viability assay.

(B) LDH assays were performed using erastin (1 μ M, 48 h) or BSO (10 μ M, 48 h) in the presence of tocotrienols (0–10 μ M), tocopherols (0–100 μ M), or Trolox (0–100 μ M) for 48 h. Data are presented as mean $\pm$ SD (n = 3).

Supplementary Figure 3



Supplementary Figure 3. Quantification of the area under the curve from FENIX assay data.

Area under the curve (AUC) values derived from FENIX assay fluorescence curves were shown. One-way ANOVA using AAPH (+) as the control group, followed by Dunnett’s post hoc test. Values of $P < 0.05$ are indicated.