

A ferulic acid derivative FXS-3 inhibits proliferation and metastasis of human lung cancer A549 cells via positive JNK signaling pathway and negative ERK/p38, AKT/mTOR and MEK/ERK signaling pathways

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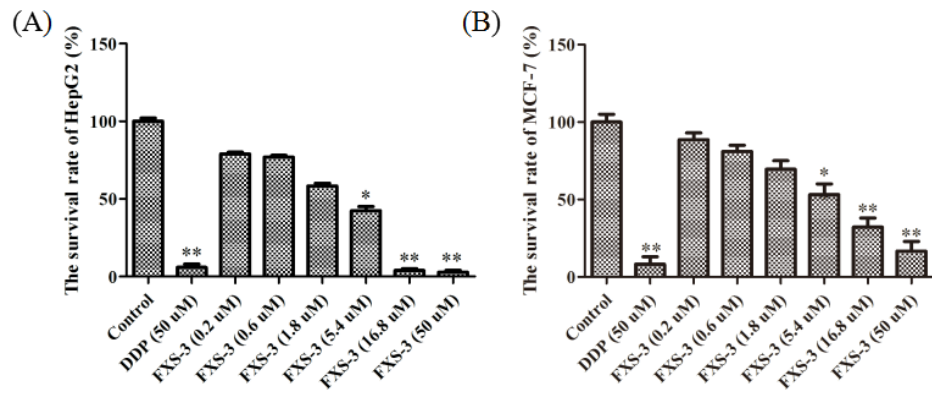


Fig. S1. FXS-3 inhibits proliferation of other cancer cells. (A) MTT assay showed that the FXS-3-treated group had a lower proliferation rate than the control group did in the liver cancer cells ($n = 3$). (B) MTT assay showed that the FXS-3 group had a lower proliferation rate than the control group did in breast cancer cells ($n = 3$). Statistical significance relative to the control group is indicated: *, $P < 0.05$; **, $P < 0.01$.

Table S1. Effects of FXS-3 on body weight of A549 xenograft in nude mice ($\bar{x} \pm s$, $n = 8$)

Group	Body weight (g)										
	1d	3d	5d	7d	9d	11d	13d	15d	17d	19d	21d
Control	20.6±0.6	21.4±0.5	22.2±0.4	22.7±0.4	23.3±0.4	23.3±0.8	23.7±0.5	24.2±0.7	24.3±0.6	24.4±0.2	24.5±0.4
Paclitaxel (8 mg/kg)	20.9±1.3	20.8±1.3	20.7±0.9**	20.2±0.7**	19.4±0.6**	19.0±0.6**	18.4±0.7**	18.0±0.6**	17.8±0.5**	17.8±0.4**	17.7±0.5**
FXS-3 (25 mg/kg)	20.1±1.2	21.0±0.9	21.8±0.6	22.5±0.9	23.0±0.9	23.5±0.7	23.9±0.6	24.2±0.5	24.4±0.5	24.3±0.5	24.4±0.4
FXS-3 (50 mg/kg)	20.7±0.7	21.5±0.5	22.0±0.5	22.5±0.6	22.9±0.6	23.3±0.6	23.6±0.5	23.9±0.5	24.2±0.6	24.4±0.5	24.5±0.4
FXS-3 (100 mg/kg)	20.6±1.5	21.1±1.2	21.5±1.1	21.7±1.1*	21.9±0.8**	22.1±1.3*	22.4±1.1*	22.5±1.0*	22.5±1.2*	22.6±1.2*	22.7±1.1*

* $P<0.05$ and ** $P<0.01$, *versus* the control group.