

Figure S1. The distribution of CFSE fluorescence was analyzed by flow cytometry at 0, 24, 48 and 72 h. (a) and (b) were the fluorescence of KB and KBvin cells at different time points. (c), (e) and (g) demonstrated the fluorescence peak of KB cells after pinostrobin or tectochrysin treatment for 24 h, 48 h and 72 h, respectively. (d), (f) and (h) suggested the fluorescence peak of KBvin cells.

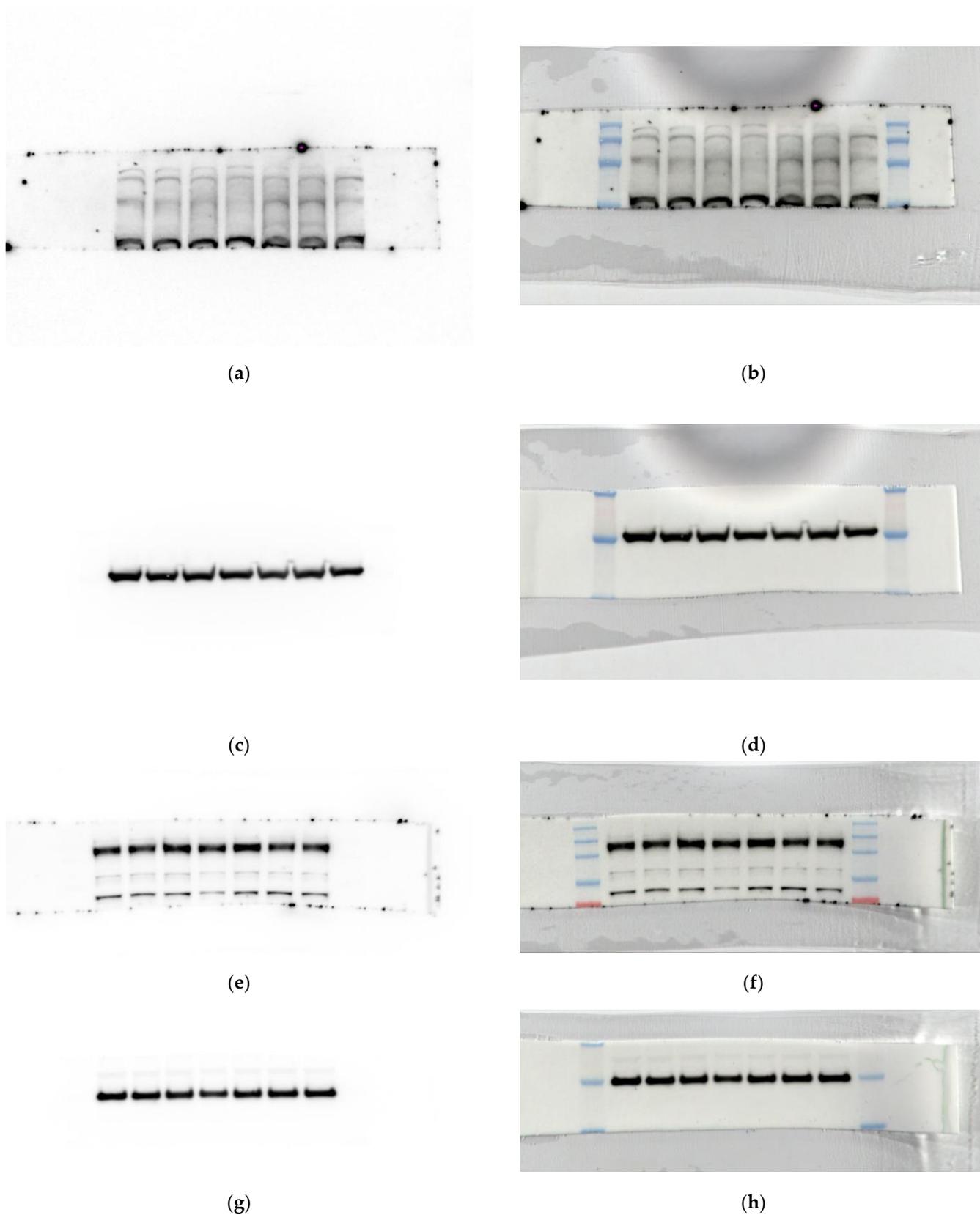


Figure S2. The raw data of Western blotting. (a), (c), (e) and (g) were the raw data of chemiluminescence. (b), (d), (f) and (h) demonstrated the data containing marker. (a–d) presented the bands of P-gp and beta-actin in KB cells, respectively. (e) ~ (h) indicated the bands of P-gp and beta-actin in KBvin cells.

Table S1. IC₅₀ values (72 h) of tariquidar in HeLa S3 and KBvin cell lines.

Treatment	HeLa S3 (drug sensitive)	KBvin (resistant)
	IC ₅₀ (μM) ± SE	IC ₅₀ (μM) ± SE
Tariquidar	16.09 ± 1.01	8.66 ± 0.27

SE, standard error.

Table S2. IC₅₀ values (72 h) of the combination of tariquidar with paclitaxel in HeLa S3 and KBvin cells.

Treatment	HeLa S3 (drug sensitive)		KBvin (resistant)	
	IC ₅₀ (nM) ± SE	RF	IC ₅₀ (nM) ± SE	RF
Paclitaxel	10.36 ± 0.30		1178.73 ± 28.45	
+ 1 μM Tariquidar	1.06 ± 0.08	9.78	0.39 ± 0.17	3022.39
+ 2.5 μM Tariquidar	1.19 ± 0.25	8.71	0.24 ± 0.03	4911.38

RF, reversal fold; SE, standard error. RF = IC₅₀ values of paclitaxel divided by those of the combination of paclitaxel with tariquidar.