

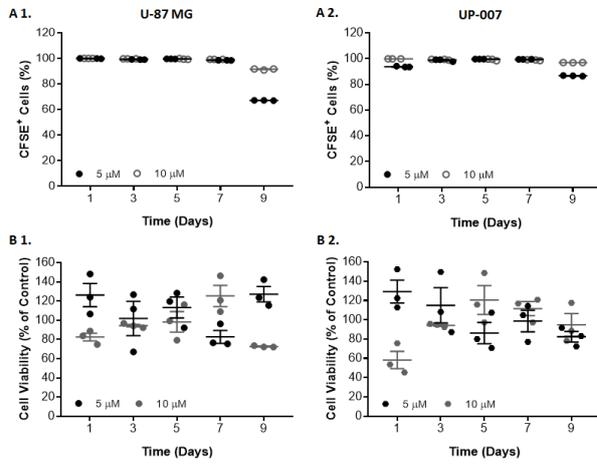
Supplementary Data.

Table S1. IC<sub>50</sub> values of TMZ, CLM and VCR in three GBM cell lines and astrocytes.

IC <sub>50</sub> (μM)*	Temozolomide	Vincristine	Clomipramine
<b>GBM</b>			
U87-MG	>1000 μM	3.60 ± 0.62 μM	16.37 ± 1.53 μM
UP-007	>1000 μM	6.92 ± 1.77 μM	6.48 ± 2.20 μM
<b>Astrocytes</b>			
UP-010	>1000 μM	0.14 ± 0.48 μM	2.40 ± 0.43 μM

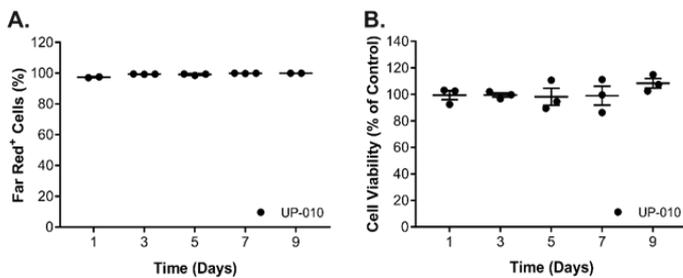
\*IC<sub>50</sub> value are mean ± standard error of means calculated from 3 independent experiments.

Figure S1.



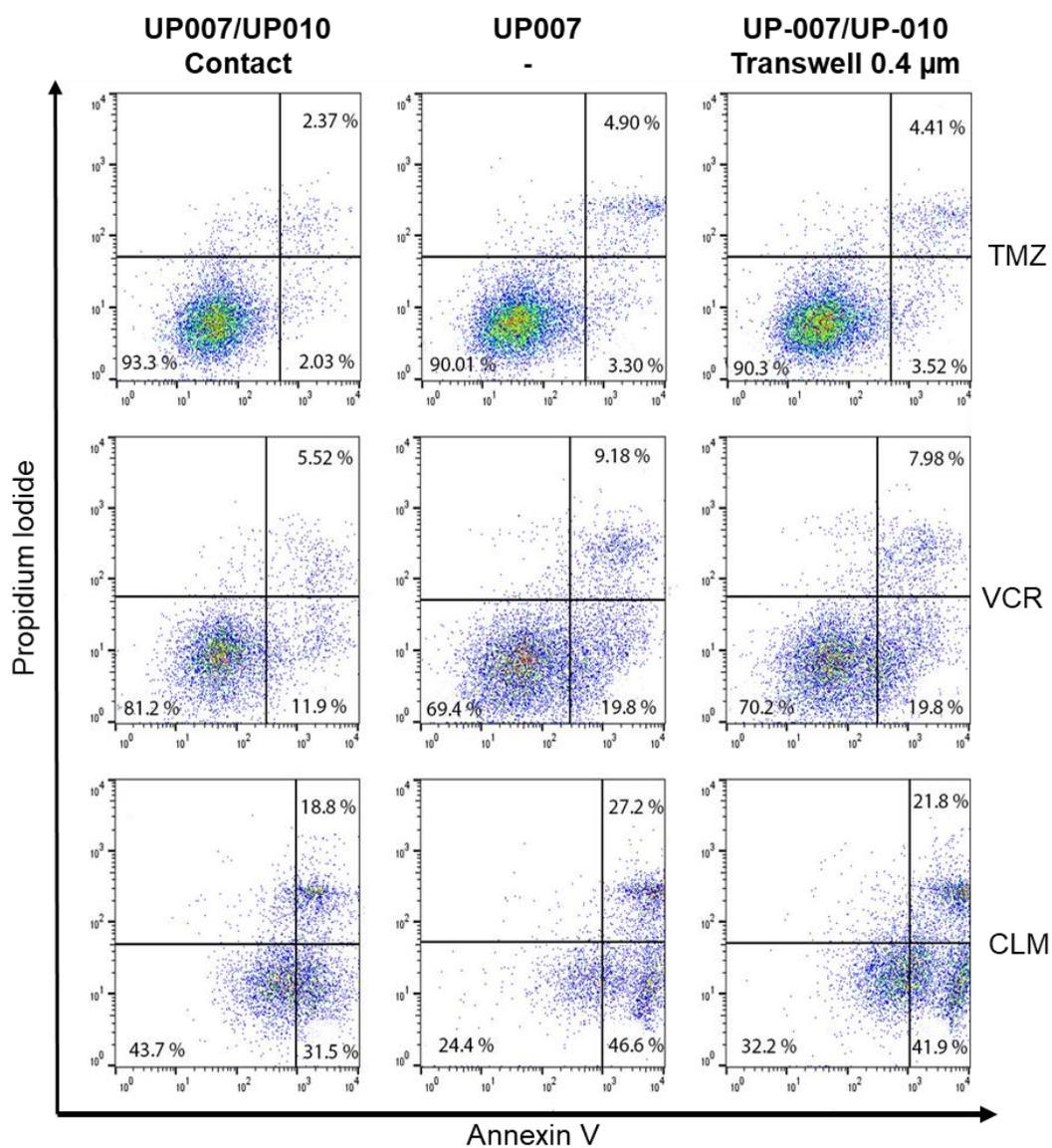
**Fluorescent labelling and viability of GBM cells marked with Cell Trace.** A1-A2) Percentage of CFSE-positive U-87 MG (A1) and UP-007 (A2) over 9 days of culture. Percentage of cells is given compared to an unstained control. B1-B2) Viability of U-87 MG (B1), UP-007 (B2) cells stained with Cell Trace CFSE (5-10 μM) compared to unstained control cells. Mean ± SEM (N=3).

Figure S2.



**Fluorescent labelling and viability of astrocytes (UP-010) marked with Cell Trace Far Red.** A) Percentage of Far Red-positive UP-010 cells along 9 days of culture. Percentage of cells is given compared to an unstained control. B) Viability of Far Red-positive UP-010 compared to unstained control cells. Mean ± SEM (N=3).

Figure S3.



**Astrocytes protect glioblastoma cells from apoptosis by direct contact.** To avoid cell-to-cell contact astrocytes cells (UP-010) were seeded on the top of transwell chamber (pore size 0.4 μm) while the CFSE glioblastoma cells (UP-007) were seeded on the bottom of 24 well plate. Representative dot plot showed TMZ (400 μM), VCR (2 μM) or CLM (20 μM) induced apoptosis of UP-007 in different culture system.