

Supporting Information

for

Highly stable selenadiazole derivatives induce bladder cancer cell
apoptosis and inhibit cell migration and invasion through activation of
ROS-mediated signaling pathways

Results and Discussion

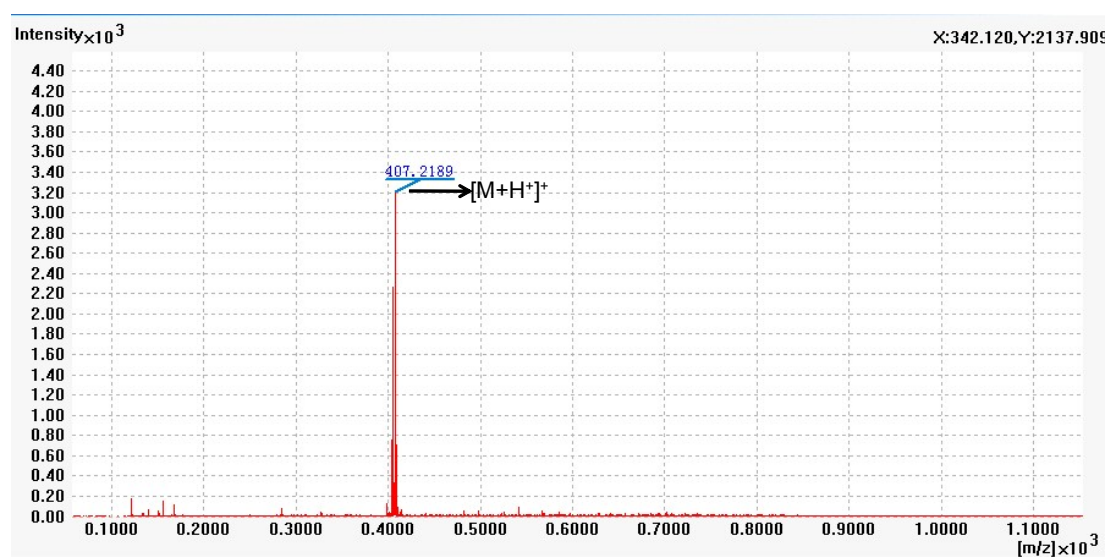


Figure S1. The ESI-MS spectrum of **1b**.

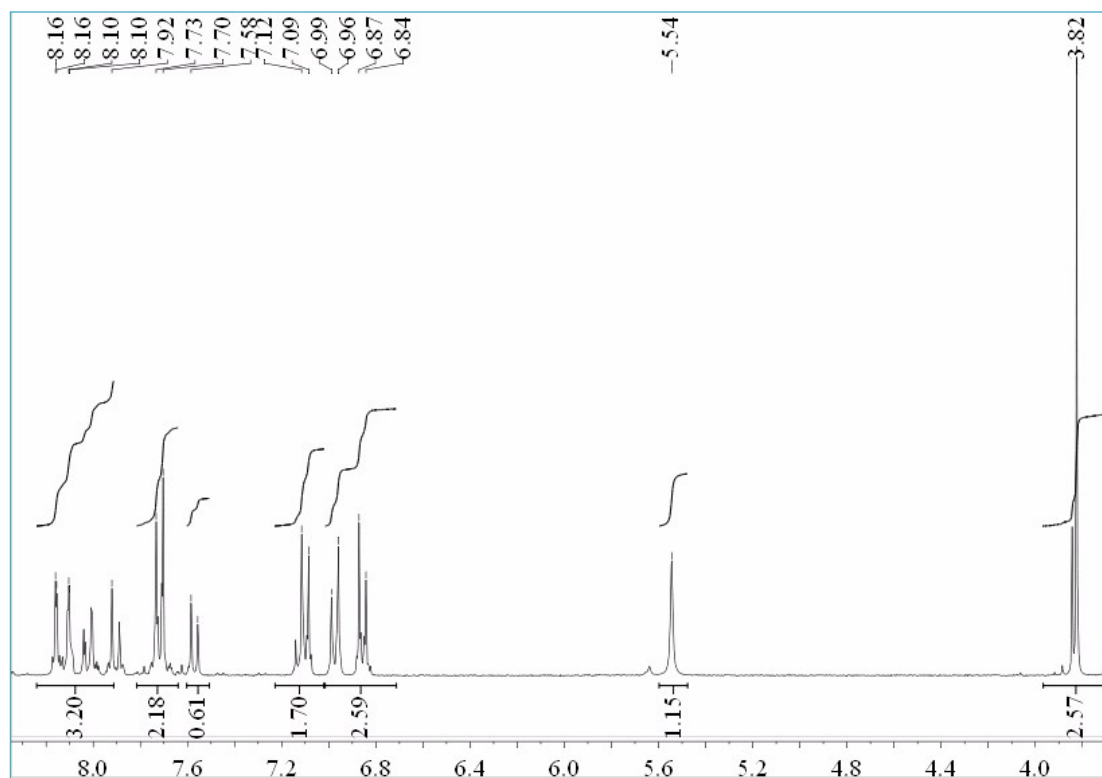


Figure S2. The ^1H NMR spectra of **1b**.

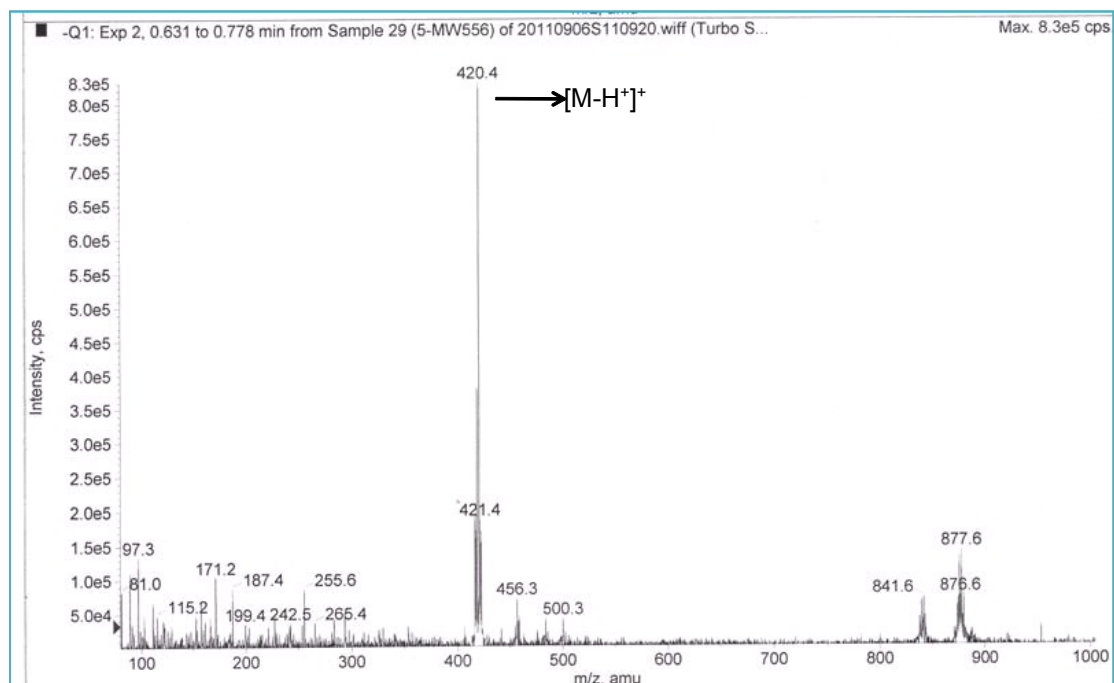


Figure S3. The ESI-MS spectrum of **1d**.

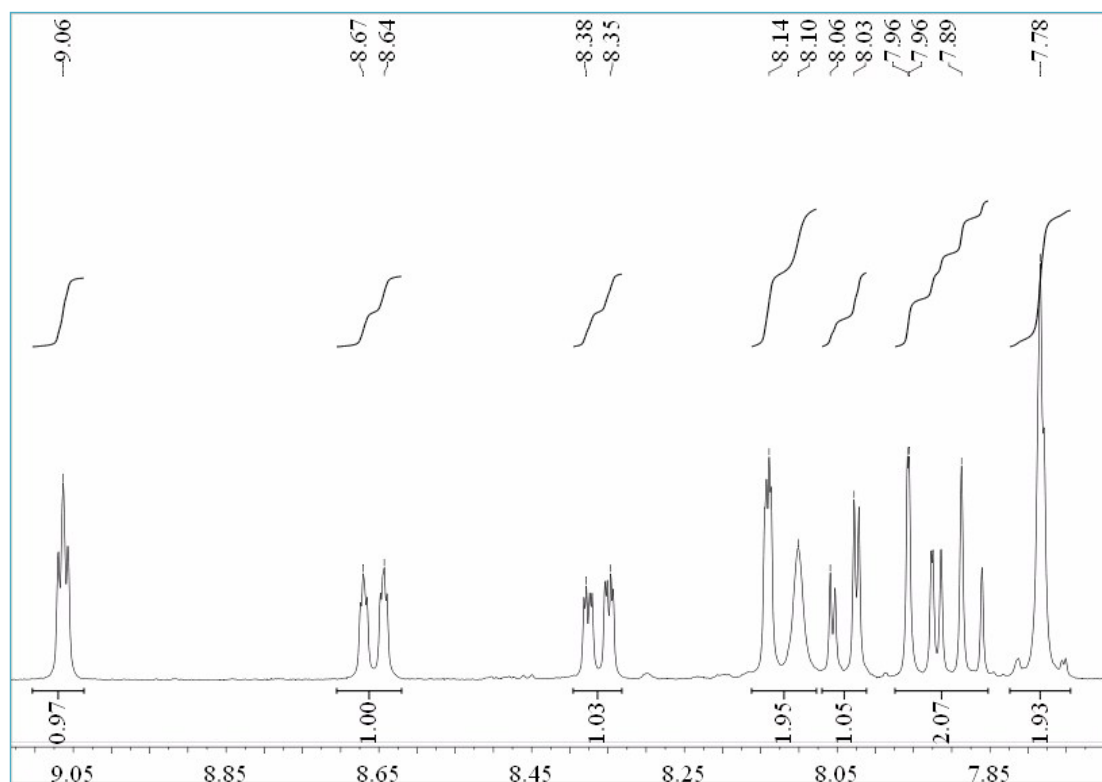


Figure S4. The ^1H NMR spectra of **1d**.

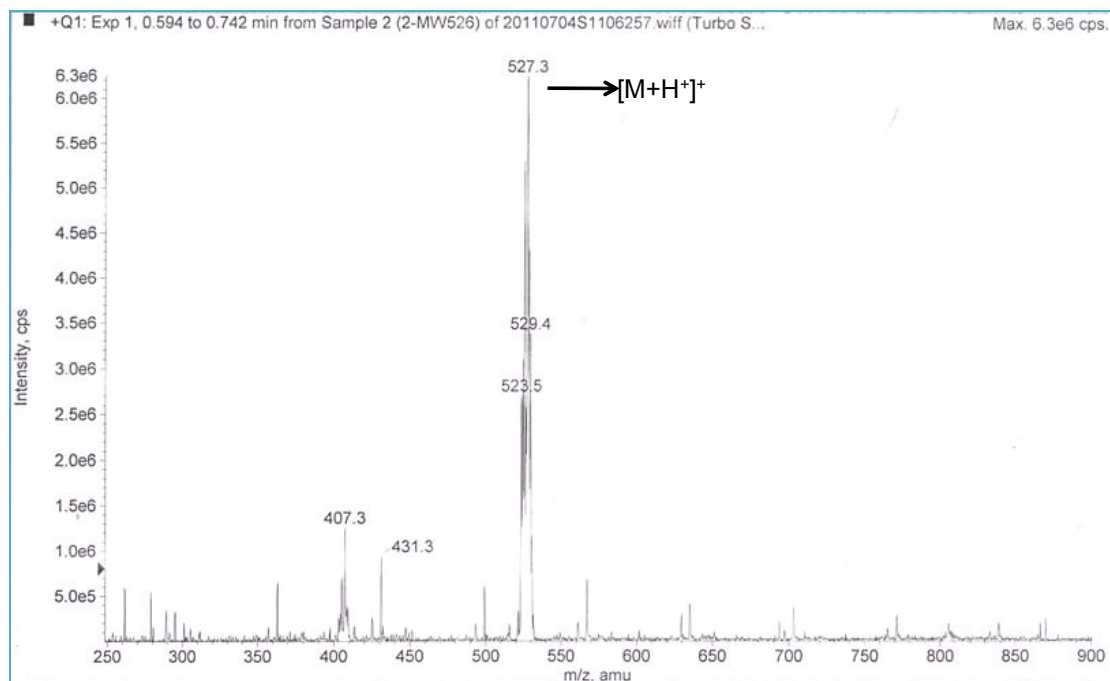


Figure S5. The ESI-MS spectrum of **2b**.

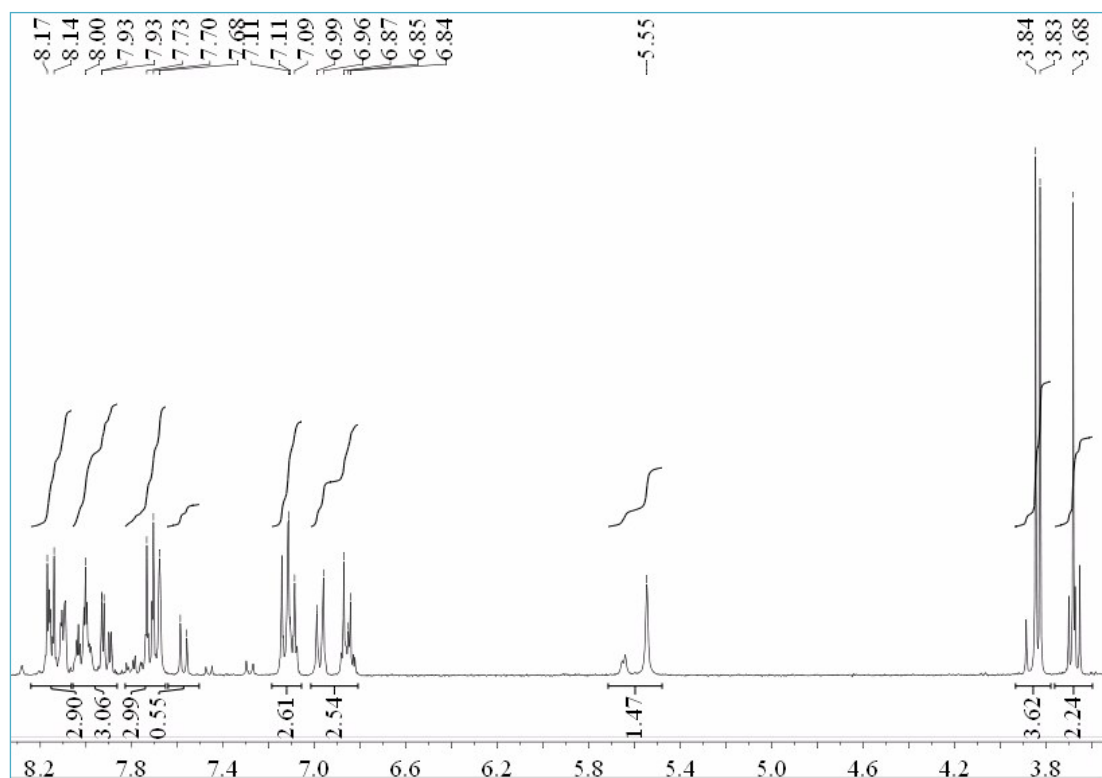


Figure S6. The ^1H NMR spectra of **2b**.

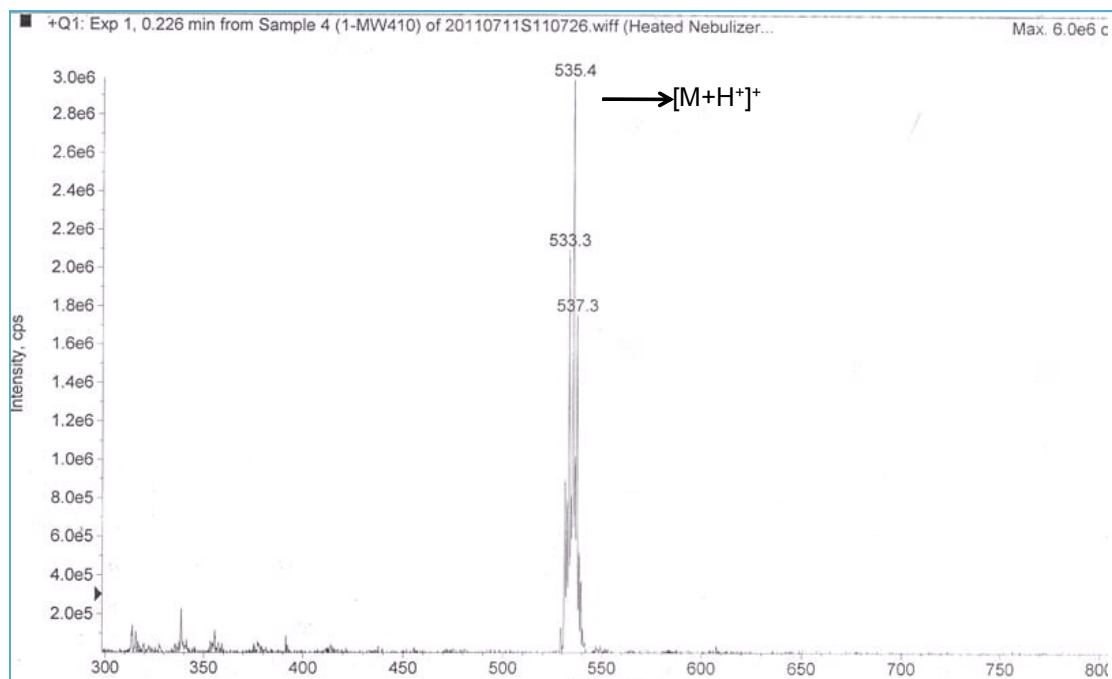


Figure S7. The ESI-MS spectrum of **2c**.

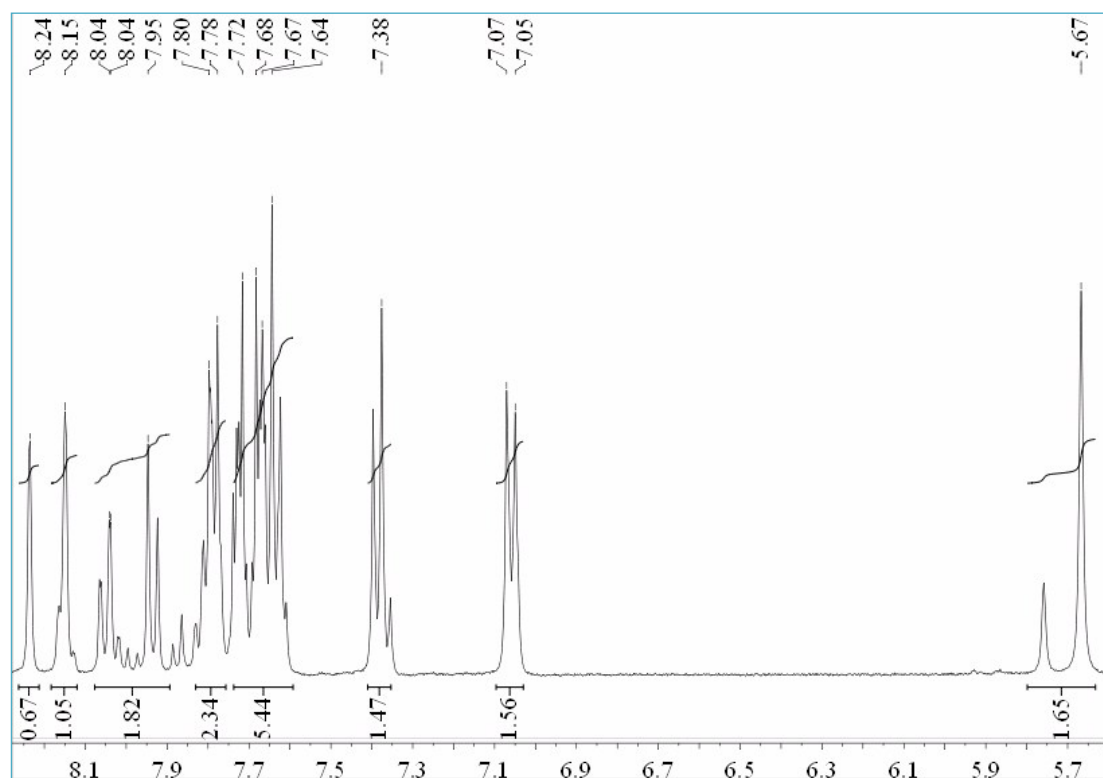


Figure S8. The ^1H NMR spectra of **2c**.

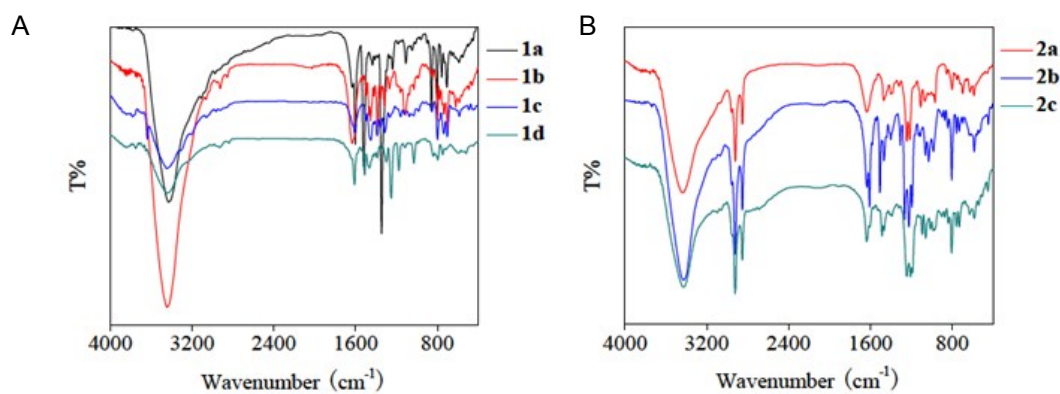


Figure S9. Fourier transform infrared spectroscopy (FTIR) of SeDs.

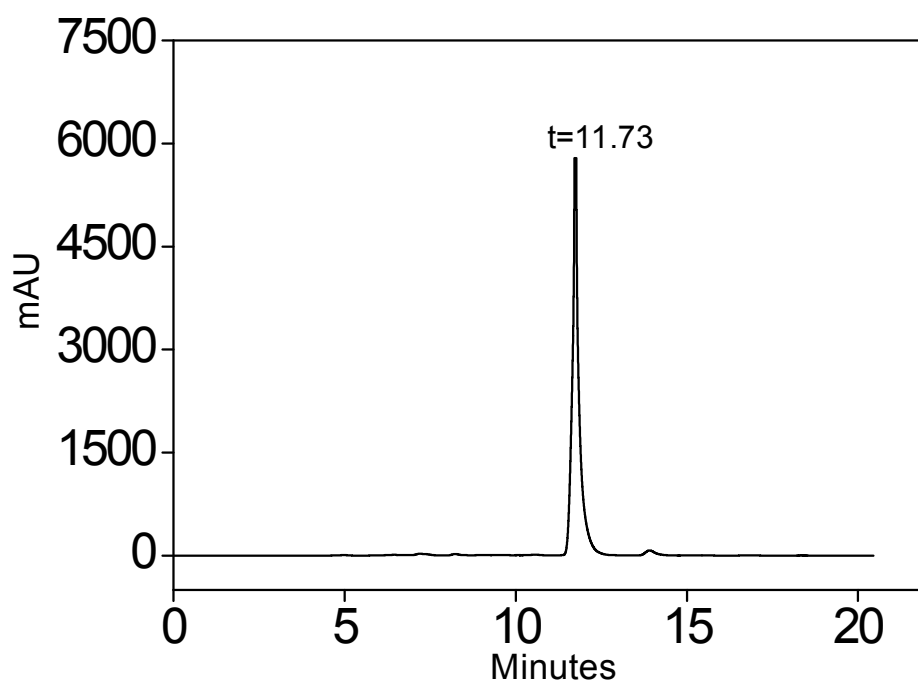


Figure S10. HPLC chromatogram of **1b**.

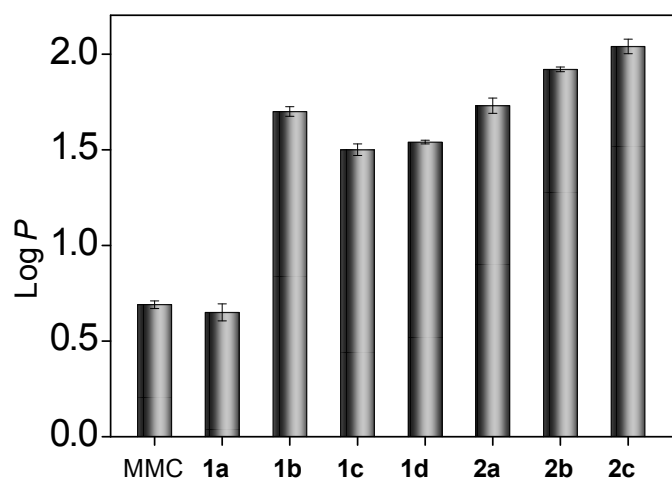


Figure S11. The lipophilicity ($\log P$) of **1a-2c** and MMC.

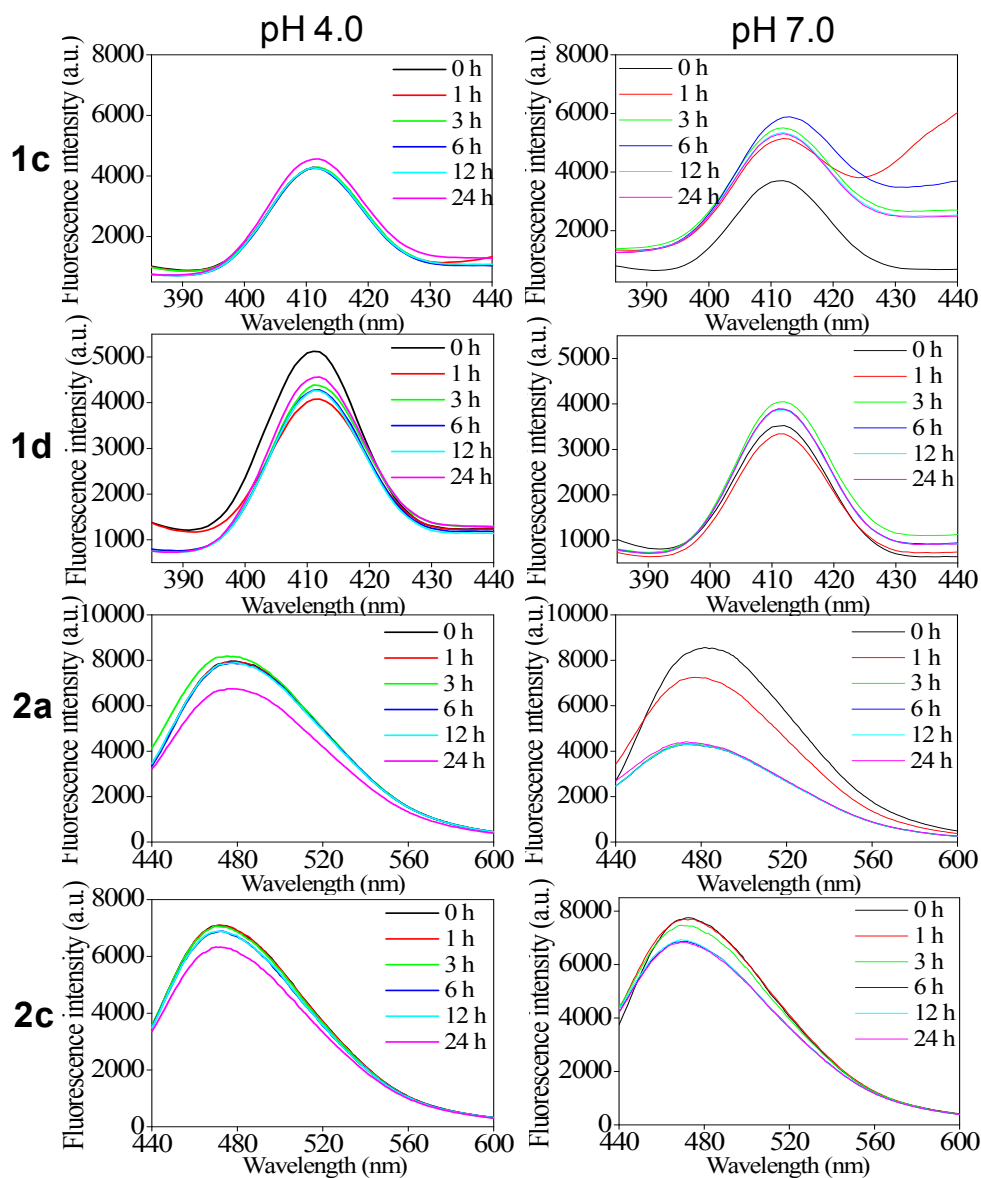


Figure S12. Fluorescence spectra of **1c**, **1d**, **2a** and **2c** (5 μ M) incubated in PBS at pH 4.0 and 7.0 for various periods of time.

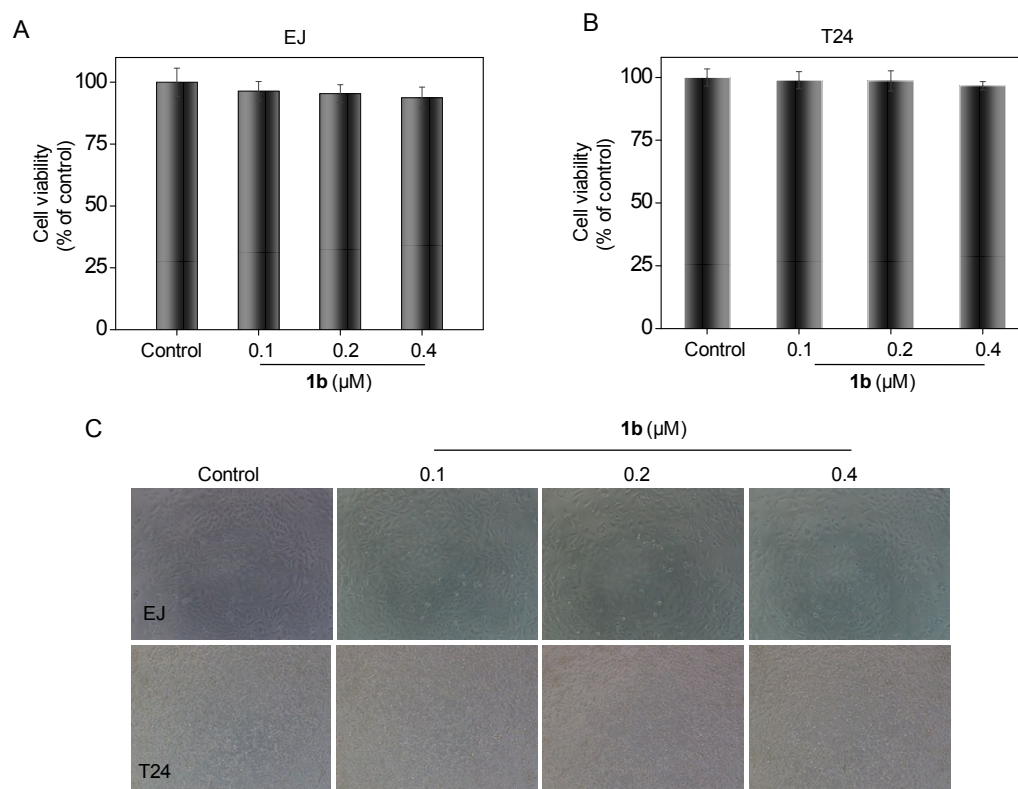


Figure S13. (A) (B) Cell viability of bladder cancer cells (EJ, T24) treated with low concentrations of **1b**. (C) Light microscopy images of exposed to low concentrations of **1b**.

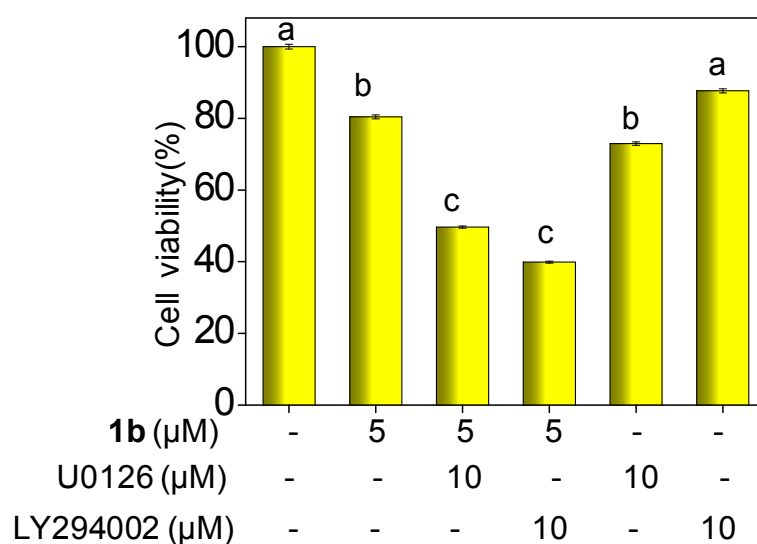


Figure S14. Effects of LY294002 and U0126 on **1b**-induced inhibition on the growth of EJ cells. Cells were pretreated with 10 μM LY294002 or U0126 for 2 h and co-treated with **1b** for another 24 h. Bars with different characters (a–c) are statistically different at the $P < 0.05$ level.