

A Novel Compound from the Phenylsulfonylpiperazine Class: Evaluation of In Vitro Activity on Luminal Breast Cancer Cells

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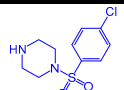
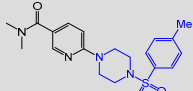
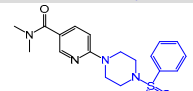
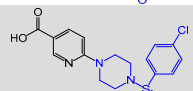
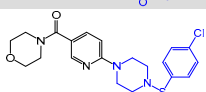
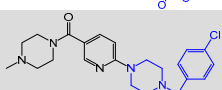
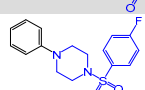
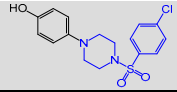
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Table S1: Compounds with half-maximal inhibitory concentration (IC₅₀) > 160 µM. Cells lines were treated for 48 h and viability was determined using the MTT assay.

Compound	Structure	IC ₅₀ (mM)			
		MCF-10A	MCF7	MD-AMB 231	MD-AMB 453
13		> 160	> 160	> 160	> 160
14		> 160	> 160	> 160	> 160
15		> 160	> 160	> 160	> 160
16		> 160	> 160	> 160	> 160
17		> 160	> 160	> 160	> 160
18		> 160	> 160	> 160	> 160
19		> 160	> 160	> 160	> 160
20		> 160	> 160	> 160	> 160

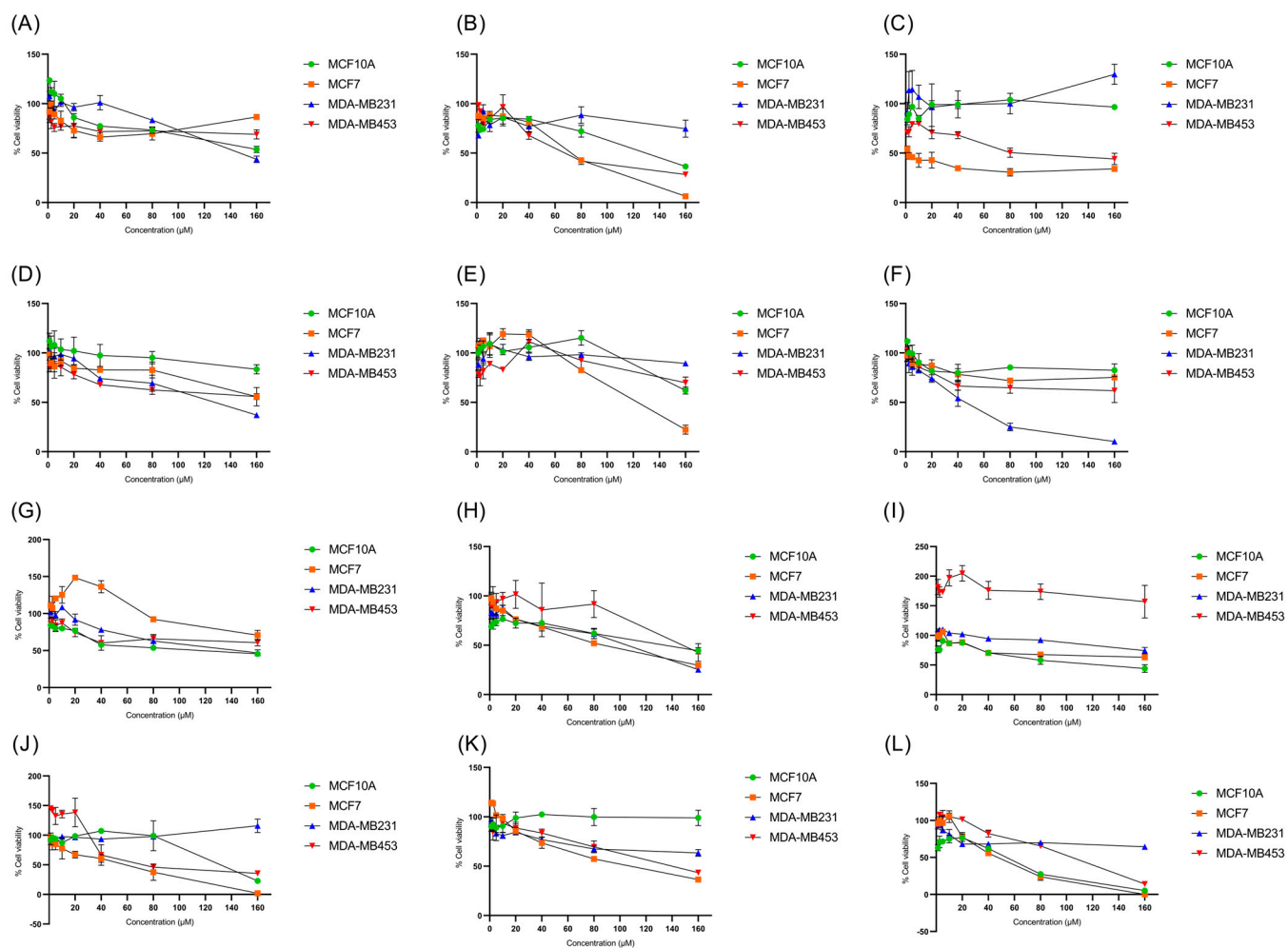


Figure S1: Graphical representation of compounds with half-maximal inhibitory concentration (IC_{50}) < 160 μ M in at least one of the tested cell lines, compounds **1 - 12** (A-L). Cell lines were treated for 48 h and viability was determined using the MTT assay.

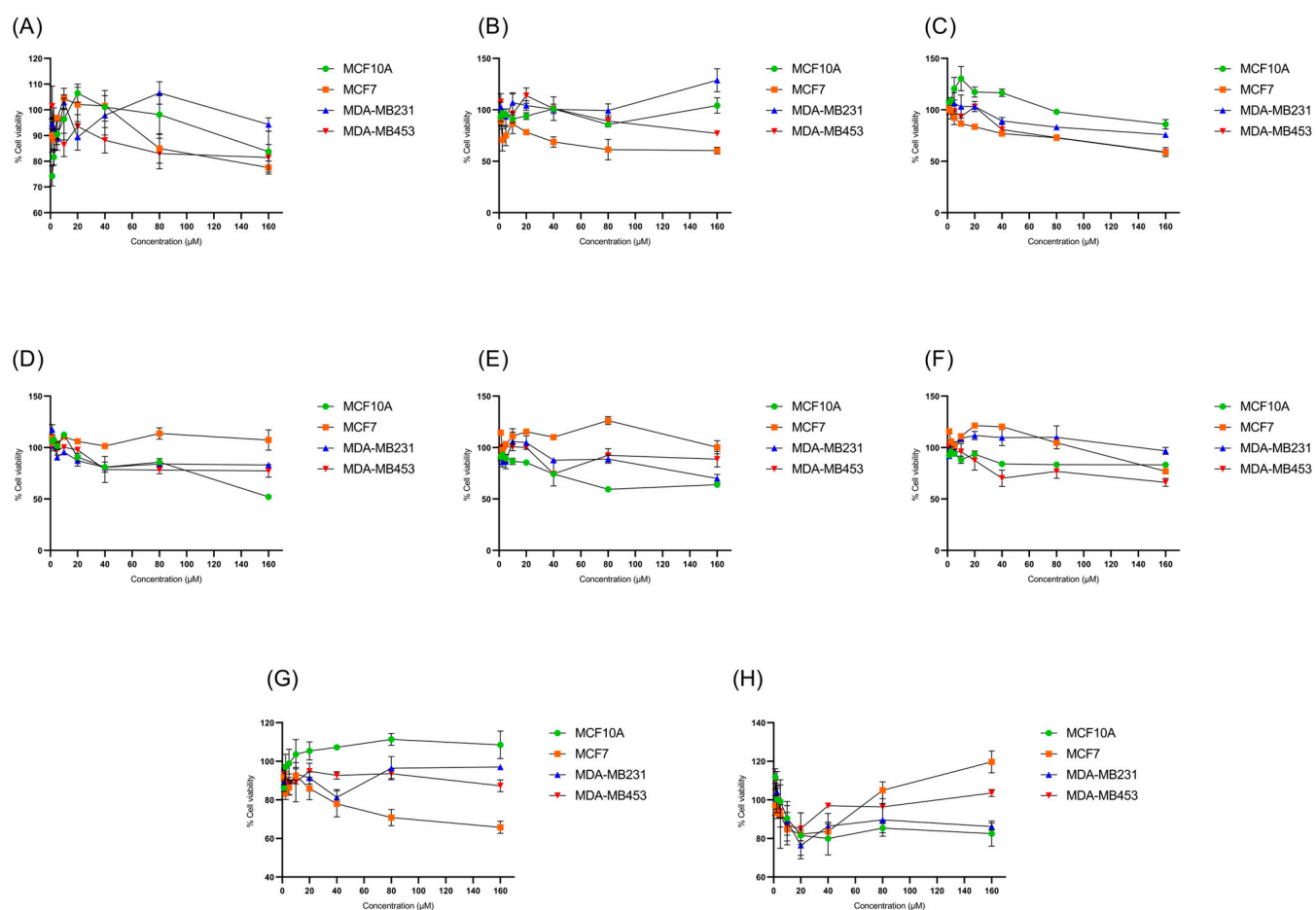


Figure S2. Graphical representation of compounds with half-maximal inhibitory concentration (IC_{50}) > 160 μM , compounds 13 - 20 (A-H). Cell lines were treated for 48 h and viability was determined using the MTT assay.

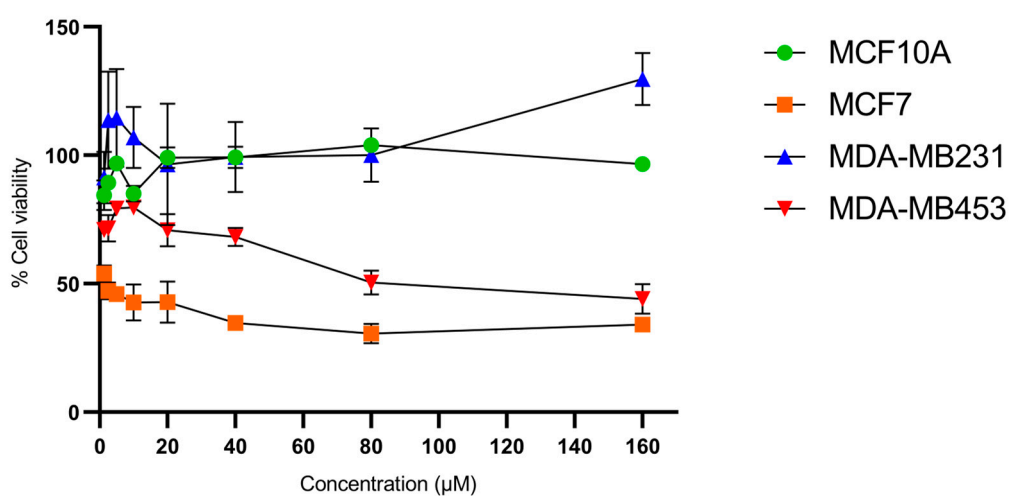


Figure S3: Graphical representation of the cytotoxic activity of compound 3 in the cell lines tested.