

## **Antiproliferative and antibacterial activity of some *para* substituted Benzylideneacetophenones and establishing their structure activity relationship (SAR)**

### **Among**

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### **Abstract**

We herein reports *in-vitro* antiproliferative and antibacterial activity of *para* substituted benzylideneacetophenones and established their structure activity relationship (SAR) to optimize *para* position as a *biologically-oriented-synthetic* target for design of small molecule based future anticancer/antibacterial agents. Among synthesized compounds, **1c** exhibits excellent antiproliferative activity against human osteosarcoma cell line (MG-63) compared to **1b** and **1a** suggesting dimethylamino (-N(CH<sub>3</sub>)<sub>2</sub>) functionality as a better *para*-substituted analogue for *in-future* anticancer agents. Similarly antibacterial screening of aforesaid compounds against against different strains of Gram negative and Gram positive bacteria reveals methoxy (-OCH<sub>3</sub>) rather than dimethylamino (-N(CH<sub>3</sub>)<sub>2</sub>) as a better *para*-substituted functionality on ring B comparatively. Furthermore, we in this study concluded and justify our theory "*lipophilicity affects antibacterial activity*".

### **Keywords**

Benzylideneacetophenone, Antiproliferative, MTT assay, Antibacterial assay