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**IN VITRO EVALUATION OF ANTICANCER AND ANTIBACTERIAL  
ACTIVITY OF INDIUM CURCUMIN AND  
INDIUM DIACETYLCURCUMIN**

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Curcumin, a yellow pigment isolated from the rhizomes of *Curcuma longa* (turmeric) has demonstrated wide range of pharmacological activities; including antioxidant, anticancer, anti-inflammatory, antibacterial, and antiviral activities. There are also some reports of therapeutic potential of curcumin for Alzheimer's disease, Parkinson disease, diabetes and hypercholesteremia [1]. Curcumin has low systemic bioavailability, because of its low aqueous solubility and poor stability [2]. Complexation of curcumin with metals improves its stability. Indium curcumin (In(cur)<sub>3</sub>) and indium diacetylcurcumin (In(DAC)<sub>3</sub>) was synthesized and characterized by Mohammadi et al. In this study, the cytotoxic effect of these complexes was assessed on bladder and stomach cancer cell lines by MTT (Methyl thiazolyl tetrazolium) assay. The result indicated that both complexes inhibit the cell lines viability to different extent. Exposure of bladder cell to In(cur)<sub>3</sub> and In(DAC)<sub>3</sub> induces almost 50%-60% cell death. The cytotoxic effect of the indium curcumin complexes on the stomach cell was concentration dependent. Exposure of the stomach cancer cell line to 20 μM concentration of In(cur)<sub>3</sub> and In(DAC)<sub>3</sub> causes to 61% and 36% cell death respectively indicating indium curcumin is more cytotoxic than indium diacetylcurcumin in MTT assay. Furthermore, antibacterial activity of the complexes against *Bacillus pumilus* and *Escherichia coli* was investigated by dilution test method. The result showed that both complexes inhibit *E. coli* growth in a dose-dependent manner, but indium curcumin is considerably more effective than the other indium complex. The result also indicated that none of the complexes have antibacterial effect against *B. pumilus*. On the whole, it can be concluded that the indium curcumin complexes have anticancer and antibacterial potential.

#### References

- [1] Shen, L.; Ji, H. F. *Trends in molecular medicine*. **2012**, *18*, 138–144.  
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