

## **Acalypha wilkesiana ethyl acetate extract enhances the in vitro cytotoxic effects of $\alpha$ -tocopherol in human brain and lung cancer cells**

### ABSTRACT

Multi-combinatorial approaches are considered nowadays to enhance the effectiveness of cancer treatment. In this study,  $\alpha$ -tocopherol was tested in combination with the ethyl acetate extract from *Acalypha wilkesiana* for cytotoxicity activity against U87MG and A549 cell lines. The GI50 values for  $\alpha$ -tocopherol against U87MG and A549 cells were  $0.923 \pm 0.411 \mu\text{g/ml}$  and  $5.290 \pm 1.952 \mu\text{g/ml}$  respectively in cell viability tests; when *A. wilkesiana* extract was added in adjunct with the treatment of  $\alpha$ -tocopherol in minimum inhibitory concentration (MIC), the GI50 values of  $\alpha$ -tocopherol improved significantly ( $p < 0.05$ ) to  $< 0.43 \mu\text{g/ml}$  ( $1 \mu\text{M}$ ) for both cell lines tested. Histological staining signified that both  $\alpha$ -tocopherol and *A. wilkesiana* extract treated cancer cell lines exhibited apoptotic morphological characteristics. Single cell gel electrophoresis (SCGE) comet assays revealed that  $\alpha$ -tocopherol caused only double strand DNA breaks; whereas *A. wilkesiana* extract caused both single strand and double strand DNA breaks in U87MG and A549 cells. It is proposed that  $\alpha$ -tocopherol and *A. wilkesiana* extract might trigger apoptosis in both U87MG and A549 cells through different apoptotic pathways that might complement each other to enhance their antiproliferative efficacy against the cancer cells.

**Keyword:** *Acalypha wilkesiana*; Tocopherol; Apoptosis; DNA damage