Introduction

In recent years, the anti-tumour potential of cannabinoids has highlighted the importance of this system in the generation of new anti-cancer therapies (Freimuth et al., 2010; Patsos et al., 2005). The aim of the present study was to investigate the potential anti-tumour activity of a cannabinoid extract rich in cannabidivarin on breast tumour cells.

Method

MCF-7 cells (American Type Culture Collection) were grown and maintained in RPMI 1640 medium supplemented with 10% fetal bovine serum at 37°C, 5% CO₂. The cells were plated in 96-well culture plates at a density of 1x10⁴ cells/well and allowed to adhere at 37°C for 24 hours. The following day, various doses of extract in the absence and presence of AM251, SR144528 and capsazepine, were added to the cells and further incubated for 4 days. Then the supernatant was removed and MTT (3-(4,5-dimethylthiazol-2-yl)-2, 5-diphenyltetrazolium bromide) was added for 4 hours. The ability of cells to form formazan crystals by active mitochondrial respiration was determined by using a Microplate reader after dissolving the crystals in DMSO. Cytotoxicity was expressed as a relative percentage of the absorbance measured at 540 nm in the control and extract-treated cells. Data were presented as the mean ± s.e.mean and analysed using ANOVA followed by Dunnet’s t-test; n=4.

Results

The extract induced dose-dependent cytotoxic effects on MCF-7 cells with an IC50 of 0.067 mg/ml. Pre-treatment with AM251, SR144528 and Capsazepine, CB1, CB2 and TRPV1 receptor antagonists, respectively, did not reverse the cytotoxicity afforded by the extract. Interestingly, the cytotoxicity was potentiated by the application of AM251 with an IC50 of 0.017± 0.01 mg/ml. Single application of antagonists alone or vehicle did not affect the survival rate of the MCF7 cells.

Conclusion

The data confirms that the cannabinoid system is involved in the apoptosis of MCF-7 tumour cells. Further experiments are required to investigate the receptor type/subtypes involvement and the mechanism of cell death.

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References
