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COMPARING THE CYTOTOXIC POTENTIAL OF WITHANIA SOMNIFERA WATER AND METHANOL EXTRACTS

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Abstract

The plant *Withania somnifera* (Linn.) (Solanaceae) is a well-known herbal medicine used in many parts of the world. It has anti-inflammatory, antioxidant, and antitumor as well as neural protective properties. It seems as if the two most active withanolide components, namely withaferin A and withanolide D, found in methanol (MeOH) extracts, are responsible for the anti-inflammatory and antioxidant properties of the plant. The current research evaluated and compared the cytotoxic potential of water and methanol extracts of *W. somnifera* using a combined crystal violet MTT and Neutral Red assay. MRC-5 cells, a human embryonic lung-derived diploid fibroblast cell line, were the cells of choice. We found that the three lowest concentrations (0.007, 0.042, 0.250 µg/ml) of the plant material extracted in double distilled H₂O and MeOH do not differ significantly in any of the assays. We therefore suggest that low concentrations of MeOH extracts (up to 0.250 µg/ml plant material) do not cause cell damage to the MRC-5 cells, however, higher levels should be avoided as cell viability and cell numbers are negatively influenced.

Key words: *Withania somnifera*, anti-inflammatory; water extract; methanol extract

Introduction

Withania somnifera (Linn.) is a member of the plant family Solanaceae and is also known as Ashwagandha, Indian Ginseng or Winter Cherry. It is an important component of the Ayurvedic pharmacopoeia of India, where it has been used for hundreds of years (Henderson and Anderson, 1966; Ahmad et al., 2005; Glaser, 1988). It has a wide distribution in Africa, Southern Europe and Asia, and is considered to be indigenous to South Africa (Henderson and Anderson, 1966). It also grows in the drier tropical regions and is also found from Liberia to Nigeria and the Southeast Islands (Iwu, 1993).

The plant is chemically very complex and more than 80 compounds have been isolated (van Wyk *et al.*, 2000). The withanolides (steroids with an ergostane skeleton), particularly Withaferin A, 3- β -hydroxy-2,3-dihydrowithanolide F as well as Withanolide D are of particular medicinal interest (Chu *et al.*, 1988; van Wyk *et al.*, 2000). Research has shown that *W. somnifera* has anti-inflammatory, antioxidant anti-tumour and immunomodulatory properties (Anbalagan and Sadique, 1981; Begum and Sadique, 1988; Somasundaram *et al.*, 1983; Kulkarni *et al.*, 1991; Al-Hindawi *et al.*, 1992; Begum and Sadique, 1987; Bhattacharya *et al.*, 1997; Parihar and Hemnani, 2003; Parihar *et al.*, 2004).

Some of the important active compounds reported in the herb include the following: Withaferin A, sitoindosides VII-X, 5-dehydroxywithanolide-R, Withasomniferin-A, 1-oxo-5 β ,6 β -epoxy-witha-2-ene-27-ethoxy-olide,4-(1-hydroxy-2,2-methylcyclopropanone) -2,3-dihydrowithaferin A, 2,3-dihydrowithaferin A, 24,25-dihydro-27-desoxywithaferin A, physagulin D (1-6)- β -D-glucopyranosyl-(1-4)- β -D-glucopyranoside, 27-O- β -D-glucopyranosylphysagulin D, physagulin D, Withanoside I-VII, 27-O- β -D-glucopyranosylviscosalactone B, 4,16-dihydroxy- β ,6 β -epoxyphysagulin D, Viscosalactone B and Diacetylwithaferin A (Uma Devi, 1996; Ganzera *et al.*, 2003; Jayaprakasam *et al.*, 2003.; Kaur *et al.*, 2003; Matsuda *et al.*, 2001).

The two most active Withanolide components, Withaferin A and Withanolide D are found in methanol extracts (Kuboyama *et al.*, 2002; Ganzera *et al.*, 2003; Choudhary *et al.*, 2004). Also, the main antioxidant activity of *W. somnifera* was found in the methanol fraction (Parihar *et al.*, 2004). Although water extracts are also known to have great medicinal potential, the methanol extracts seem to extract the active components the best. The question, however, arises whether the methanol extracts of the herb may be cytotoxic to cells in culture. Therefore, because of the potential use of the herb as an anti-inflammatory product, the current research aimed to evaluate and compare the cytotoxic potential of water and methanol extracts of *W. somnifera* using a combined Crystal Violet assay, the MTT assay and the Neutral Red assay. The cells of choice were MRC-5 cells, which is a human embryonic lung-derived diploid fibroblast cell line. Statistical analyses were employed to determine whether there is a correlation between the two extract methods, as well as the concentrations used.

Experimental Procedures

Herb material

Root material (WS1PRETORIA) was collected in the Pretoria area (South Africa) and identified by botanists from the Department of Botany, University of Pretoria, South Africa. A herbarium specimen was prepared and compared to an authentic specimen in the HGJW Sweikerdt herbarium at the University of Pretoria.

Preparation of extracts

Leaves, stems and roots were fragmented and dried at 45°C for 48 h and MeOH and water extracts were made. Plant material (140 mg) was added to 1,400 ml double distilled (dd) H₂O of water, yielding a final concentration of 100mg plant material per ml. 2.1039 g of MeOH extracted plant material was added to 4.2 ml MeOH, yielding a final concentration of 0.5 g of plant material per ml MeOH. This was diluted 1:4 with dd H₂O, yielding a final concentration of 100 mg plant material in 1 ml. Five concentrations of extracts of both ddH₂O and MeOH were added to the cells in culture. The concentrations were: 0.007, 0.042, 0.250, 1.151, and 9.090 μ g/ml.

MRC-5 Permanent cell line

MRC-5 cells were obtained from Highveld Biological Company, Johannesburg, South Africa. The cells were plated at a cell concentration of 2×10^4 cells per ml in 24 flat well plates and were kept for 24 h at 37°C and 5% CO₂ before conducting the experiments. Solution of 100mg/ml of each of the plant

extracts was prepared either in dd H₂O or MeOH and exposed to MRC-5 cells for 48 h. The cytotoxicity of MeOH alone was also determined.

Combined Crystal Violet, MTT and Neutral Red assay

Cell number, cell viability and lysosomal membrane integrity, were determined using a combined Crystal Violet, MTT and Neutral Red (CV/MTT/NR) assay. To each well 100µl 0.15% NR prepared in DPBS (A 10x DPBS stock solution was prepared by dissolving 2g/l KCl, 2g KH₂PO₄, 80g/l NaCl and Na₂HPO₄ in ddH₂O that was diluted 1:9 with ddH₂O prior to use) was added and then incubated for 90 min at 37°C in a CO₂ incubator. A further 50µl volume of a 0.1 mg/ml 3-(4,5- Dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) solution prepared in DPBS was added to each well and the cell culture plates were maintained for a further 60 min at 37°C. The medium was then carefully removed and the plates were blotted dry. The cells were fixed for 10 min with 200µl of a 1% acetic acid and 1% formaldehyde solution in ddH₂O. The fixative was removed and the NR was solubilized with 200µl of a 1% acetic acid and 50% ethanol solution prepared in ddH₂O. The dissolved NR was transferred into a 96-flat well plate. The MTT formazan crystals were then dissolved with 200µl DMSO by shaking of the plates for 20 min before being transferred into a 96-flat well plate. Each well was washed once with DPBS and was dried in a microwave oven for a few mins at 60% power. The cells attached to the bottom of the plate were stained by adding a volume of 300µl of a 0.1% (weight/ volume (w/v) CV solution prepared in 200mM of formic acid pH 3.5, each well for 60 min. After the microplate was washed with ddH₂O and dried, the bound dye was dissolved in 10% acetic acid prepared in ddH₂O. The solution was transferred to a 96-well and the absorbance of all bioassays was determined spectrophotometrically at 570 nm using the Microplate Reader (ELx800).

Statistical analysis

The experiment was conducted in a factorial design with main effects the extracts at two levels (plant material extracted in ddH₂O and also in MeOH) and *concentration* at five levels (0.007, 0.042, 0.250, 1.151, and 9.090 µg/ml). For each of the three assays the outcome variable was analysed in an appropriate analysis of variance (ANOVA) with two fixed effects, namely extract and concentration as well as the interaction. Pair-wise comparisons were done using the LSD method of Fisher. Testing was done at the 0.05 level of significance.

Results and Discussion

Combined Crystal Violet, MTT and Neutral Red assay

The CV assay measures cell number in the presence of the 5 concentrations of the two extracts. All analyses were performed on the dataset and bar graphs of the results compiled (Figure 1 – 3). From the ANOVA the two extracts differ significantly ($p = 0.109$) but not the concentrations ($p = 0.5676$). However, a significant interaction between extracts and concentrations exists ($p < 0.0001$). This interaction is displayed in Figure 1. Note that the first 3 concentrations of the two extracts react similarly but behave very differently over the last two concentrations, causing the significant difference between extracts ($p = 0.0109$) at these high concentrations, the ddH₂O extract increased cell numbers, while the MeOH extract caused a decrease.

Figure 2 displays the results of the MTT assay, showing cell viability. ANOVA showed that the two extracts differ significantly ($p = 0.0005$) and that the concentrations also differ significantly ($p < 0.0001$). In particular, following pair wise comparisons, the first three concentrations do not differ significantly but differ from the last two concentrations which also differ from each other. There is also a significant interaction between extract and concentration ($p = 0.0110$) which is clear from Figure 2, where

except for the fourth concentration the extracts behave similarly. However, what can be concluded from the graph is that cell viability decreases when using the two highest concentrations of the MeOH extract and the highest concentration of the ddH₂O extract.

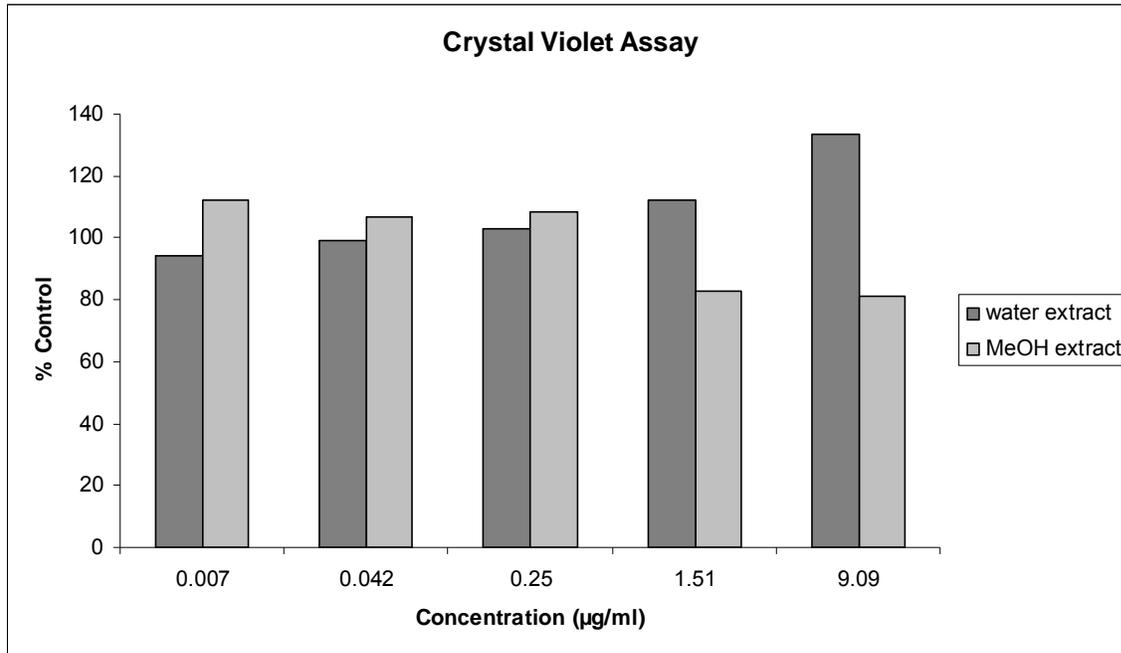


Figure 1: Effects of 5 concentrations of *W. somnifera*, extracted in ddH₂O and MeOH and prepared in cell culture medium on MRC-5 cell number after 48 h exposure: results from Crystal Violet Assay.

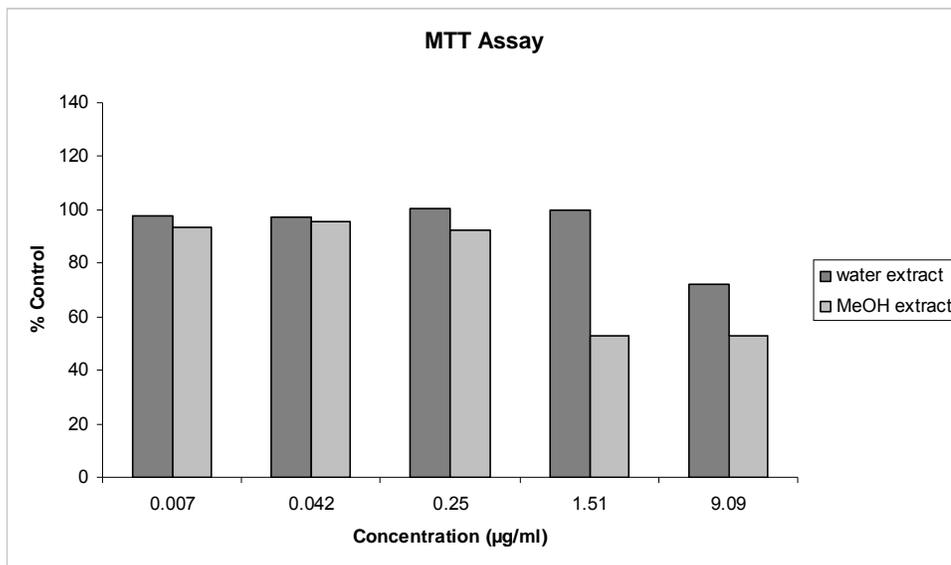


Figure 2: Effects of 5 concentrations of *W. somnifera*, extracted in ddH₂O and MeOH and prepared in cell culture medium on MRC-5 cell viability after 48 h exposure: results from MTT Assay.

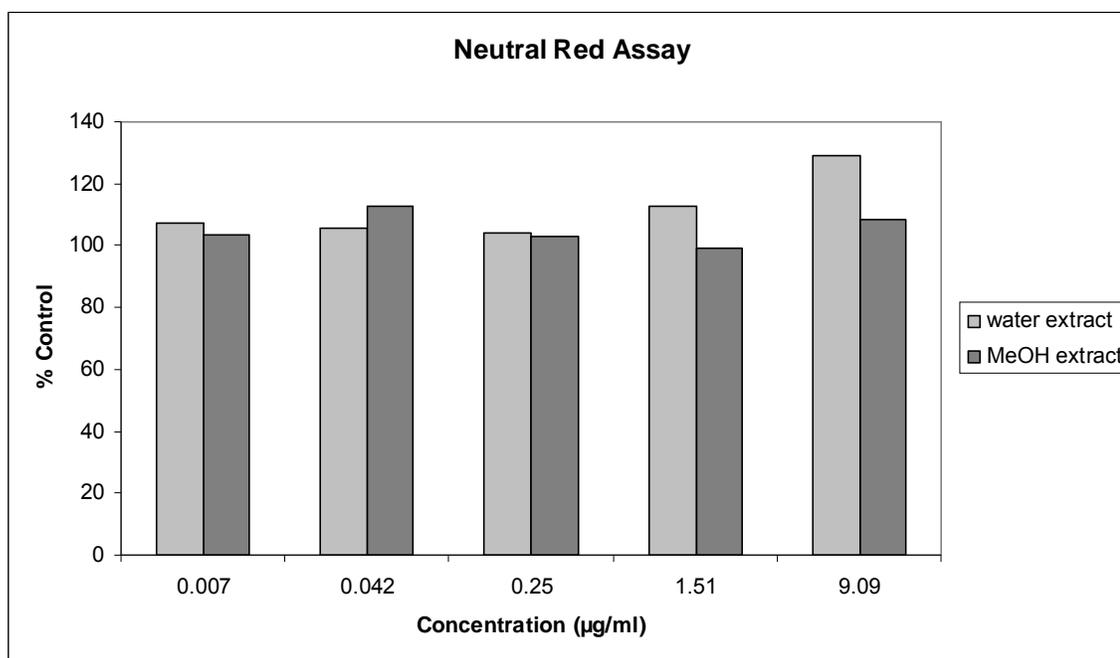


Figure 3: Effects of 5 concentrations of *W. somnifera*, extracted in ddH₂O and MeOH and prepared in cell culture medium on MRC-5 lysosomal integrity after 48 h exposure: results from Neutral Red Assay.

The final assay, namely the NR assay, which determines lysosomal membrane integrity, is presented in Figure 3. ANOVA showed that respectively the extracts and the concentrations are significantly different ($p = 0.0304$ and $p = 0.0170$). In particular, LSD pair-wise comparisons show that the first four concentrations do not differ significantly; however, these four concentrations differ from the highest concentration. The significant interaction between extract and concentration is also clear in Figure 3, where the two extracts behave similarly over the first three concentrations and quite differently over the last two concentrations.

We therefore conclude that the three lowest concentrations (0.007, 0.042, and 0.250 µg/ml) of the plant material extracted in ddH₂O and MeOH seems not to be significantly different in any of the assays. At these three concentrations cell numbers, cell viability and lysosomal membrane activity were preserved.

Thus low concentrations of MeOH extracts (up to 0.250 µg/ml plant material) do not cause cell damage to the human embryonic lung-derived diploid fibroblast cell line, however, higher levels should be avoided as particularly cell viability and cell numbers are negatively influenced. The two active components Withaferin A and Withanolide D are most successfully extracted using MeOH (Kuboyama et al., 2002; Ganzera et al., 2003; Choudhary et al., 2004). Research therefore investigating the anti-inflammatory or antioxidant activity of *W. somnifera* can use MeOH extracts, but care should be taken not to use concentrations that are too high.

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