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# ANTIFUNGAL ACTIVITY OF *Plinia jaboticaba* (Vell.) Berg. LEAVES EXTRACT ON *Candida* sp. CULTURES

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## **INTRODUCTION**

The species Plinia jaboticaba (Vell.) Berg, popularly known as "jabuticaba-sabará" is an endemic plant of Southeast of Brazil, belonging to the Myrtaceae family. The literature describes its application in popular medicine on treatment of angina, dysentery, erysipelas and asthma, though the few studies on the species concentrate analyzing themselves in the properties. The aim of this study was to evaluate their antifungal effect on Candida cultures and their monocytes SD. cytotoxicity.

### **MATERIAL AND METHODS**

To evaluate the action of P.jaboticaba leaves extract on Candida sp., it was used the doses of 50 mg/mL, 25 mg/mL and 12.5 mg/mL. Fungal growth was assessed by viable fungi recovery from co-cultures for 24 hours, after plating in Potato Dextrose agar. It was considerate the average of co-culture to the results. The cell viability of human monocytes against the different doses of the extract was performed by MTT reduction test, which finalized to verify cytotoxicity. experiment was submitted to the Human Ethics Committee under approval number 53361116.3.0000.5370/2016.

### **RESULTS**

After evaluating the duplicate cocultures, following results were obtained: in the cultures of 12.5mg/ml there was no antifungal action, as well as in control fungal culture. However, in the 25 and 50mg/mL doses, there was a significant decrease in fungal growth. The average fungal recovery at a dose of 25mg/mL was

696 CFU and for 50mg/mL it was 188 CFU. There was no cytotoxicity in all concentrations when compared to the control group, a positive sign for further studies of therapeutic application of the plant specimen.

### **CONCLUSIONS**

According to the results obtained in the present work, it is concluded that the P.jaboticaba extract of leaves antifungal effect when used at doses of 25 to 50mg/mL. It is important to remember that the analysis performed preliminary to confirmatory experiments and that it is necessary to carry out cocultures containing phagocytic cells to verify the potentiation of the antifungal action of this extract.

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