

In vitro study of the antifungal potential of **Apiaceae** hydroalcoholic extracts against *Candida* species

Natália Martins,^{1,2} Isabel C.F.R. Ferreira¹ Lillian Barros,¹ Mariana Henriques², Sónia Silva²

¹IBB - Institute for Biotechnology and Bioengineering, Centre of Biological Engineering, University of Minho, Braga, Portugal.

²Mountain Research Centre (CIMO), ESA, Polytechnic Institute of Bragança, Portugal.

The use of medicinal plants is an ancient practice, but recently there is an increasing interest towards the evaluation of their bioactive properties. Opportunistic fungal infections, linked with higher rates of fungal resistance to the current antifungal drugs, have deserved special relevance in the last decades. *Candida albicans* was identified as the main responsible agent for those infections, but other non-*Candida albicans* *Candida* (NCAC) species have been also found [1]. Thus, it is urgent to discover new alternatives against those pathogens with high resistance.

In the present work, the antifungal potential of hydroalcoholic extracts obtained from two **Apiaceae** plants (*Coriandrum sativum* L. and *Pimpinella anisum* L.), commonly used in folk medicine, were evaluated against a total of 19 *Candida* strains (from the species: *C. albicans*, *C. glabrata*, *C. parapsilosis* and *C. tropicalis*).

The fruits of *Coriandrum sativum* L. (coriander) and *Pimpinella anisum* L. (anise) showed similar antifungal potential considering the studied strains, being effective against three of the nineteen strains. However, regarding the tested *Candida* species, the extracts presented considerable variations. Whereas coriander was effective against *C. parapsilosis* (ATCC22019 and 513143) and *C. tropicalis* (ATCC750), anise was effective against *C. parapsilosis* (513143 and 491861) and *C. albicans* (558234). Furthermore, the inhibitory zones were different at 24 and 48h. Further studies are being carried out in order to characterize the mechanism of action and the compounds responsible for the bioactivity, but the use of these extracts seems to have potential in antifungal therapy.

Acknowledgments: The authors are grateful to FCT (Portugal) for financial support to CIMO (PEst-OE/AGUI0690/2011), N. Martins (SFRH/BD/87658/2012), L. Barros (*Compromisso para a Ciência 2008) and S. Silva (NORTE-07-0124-FEDER-000027; QREN) contracts.

References:

[1] Martins N, Ferreira ICFR, Barros L, Silva S, Henriques M. (2014). *Mycopathol.* 177, 223–240.