

Valorisation of chestnut shells as a prominent sustainable ingredient for cosmetics

Pinto Diana a, Cádiz-Gurrea María de la Luz a, Garcia Juliana b, Saavedra Maria José b, Dall'Acqua Stefano c, d, Costa Paulo e, Delerue-Matos Cristina a, Rodrigues Francisca a,*

a REQUIMTE/LAQV, Polytechnic of Porto, School of Engineering, Porto, Portugal, diana.pinto@graq.isep.ipp.pt (D.P.);

b CITAB – Centre for the Research and Technology of Agro-Environmental and Biological Sciences, University of Trás-os-Montes and Alto Douro, Vila Real, Portugal;

c DAFNAE, Department of Agronomy, Food, Natural Resources, Animals and Environment, University of Padova, Legnaro, Italy;

d DSF, Department of Pharmaceutical and Pharmacological Sciences, University of Padova, Padova, Italy;

e REQUIMTE/UCIBIO, MedTech-Laboratory of Pharmaceutical Technology, Department of Drug Sciences, Faculty of Pharmacy, University of Porto, Porto, Portugal.

*francisca.rodrigues@graq.isep.ipp.pt; franciscapintolisboa@gmail.com.

The demand for novel active ingredients for cosmetic purposes, mainly recovered from natural sources, has raised in the last decade owing to society's awareness of sustainability issues and the importance of skin appearance. Food by-products arise as promising alternatives to obtain eco-friendly cosmetic ingredients with high added value [1]. Chestnut shells, an abundant and undervalued agro-industrial waste generated during chestnut (*Castanea sativa*) processing, are particularly rich in bioactive compounds, such as polyphenols and vitamin E, endowed with pro-healthy skin effects [1-3].

The purpose of this study was to recover bioactive compounds from chestnut shells using clean and proficient technology, namely ultrasound-assisted extraction (UAE). The phenolic composition of the extract prepared at 70 °C for 40 min was evaluated by LC-MS, while the antioxidant/antiradical and antimicrobial activities, as well as elastase and hyaluronidase inhibitory effects, were assessed by in-vitro assays. The safety on skin cell lines (HaCaT and HT29-MTX) was assessed by an MTT assay (0.1 – 1000 µg/mL).

The chestnut shells extract revealed to be rich in ellagic acid, caffeic acid derivative, and epigallocatechin. These compounds were probably responsible for the high antioxidant, antiradical, and antimicrobial effects observed. The outcomes disclosed pronounced antibacterial effects against *Staphylococcus aureus*, *Klebsiella pneumoniae*, and *Escherichia coli* strains. The elastase inhibition was 74.8% at 0.4 mg/mL, while the IC₅₀ from the hyaluronidase assay was 760 µg/mL. The safety of the extract was proved on HaCaT and HFF-1 cells up to 1000 µg/mL (viability >70%). Overall, these findings pointed out that the chestnut shells extract is an appealing source of bioactive compounds with promissory in-vivo skin effects, such as anti-wrinkles. Noteworthy, further studies should be performed to assess the ex vivo permeation of the bioactive compounds through Franz diffusion cells coupled to human skin as well as the assessment of the extract safety in human volunteers by in vivo patch test.

This research was funded by FCT/MCTES through national funds (UIDB/50006/2020) and project PTDC/ASP-AGR/29277/2017 - *Castanea sativa* shells as a new source of active ingredients for Functional Food and Cosmetic applications: a sustainable approach, financially supported by national funds by FCT/MCTES and co-supported by Fundo Europeu de Desenvolvimento Regional (FEDER) throughout COMPETE 2020 - Programa Operacional Competitividade e Internacionalização (POCI-01-0145-FEDER-029277). This work was also supported by Project UIDB/04033/2020 (CITAB-Center for the Research and Technology of

Agro-Environmental and Biological Sciences). Diana Pinto (SFRH/BD/144534/2019) is grateful for the Ph.D. grant financed by FCT/MCTES and POPH-QREN and supported by funds from European Union (EU) and Fundo Social Europeu (FSE) through Programa Operacional Regional Norte. Francisca Rodrigues is thankful for her contract (CEECIND/01886/2020) financed by FCT/MCTES—CEEC Individual 2020 Program Contract.

[1] D. Pinto, M.d.l.L. Cádiz-Gurrea, A. Vallverdú-Queralt, C. Delerue-Matos, F. Rodrigues, *Food Res. Int.*, 144 (2021) 110364.

[2] F. Lameirão, D. Pinto, E.F. Vieira, A.F. Peixoto, C. Freire, S. Sut, S. Dall'Acqua, P. Costa, C. Delerue-Matos, F. Rodrigues, *Antioxidants*, 9 (2020) 267.

[3] D. Pinto, M.d.l.L. Cádiz-Gurrea, J. Garcia, M.J. Saavedra, V. Freitas, P. Costa, B. Sarmiento, C. Delerue-Matos, F. Rodrigues, *Sustain. Mater. Technol.*, 29 (2021) e00309.