

77
II Meeting

of REQUIMTE

Monte da Caparica

. 28 and 29 .

September 2001

Book of Abstracts



r e q u i m t e

rede de química e tecnologia

DEVELOPMENT OF AN HPLC/DAD METHOD FOR DETERMINATION OF PHENOLIC PROFILE IN PORTUGUESE OLIVE FRUITS

Ana F. Vinha¹, Paula B. Andrade¹, Branca M. Silva¹, José A. Pereira², Patrícia Valentão¹, Rosa M. Seabra¹, M. Beatriz Oliveira³

CEQUP/ ¹Serviço de Farmacognosia and ³Serviço de Bromatologia, Faculdade de Farmácia, Universidade do Porto, R. Aníbal Cunha, 164, 4050-047 Porto. ²Escola Superior Agrária, Instituto Politécnico de Bragança, Quinta de Sta. Apolónia, Apartado 172, 5300 Bragança.

INTRODUCTION

Polyphenolic compounds influence the sensorial properties of olive fruits and virgin olive oils and are important markers for studying fruit characteristics of different cultivars and for controlling oil production processes (Romani *et al.*, 1999).

A few chromatographic methods have been used to study the phenolic compounds of olive fruit (Soler-Rivas *et al.*, 2000; Capasso *et al.*, 1992; Ficarra *et al.*, 1991; de Laurentis *et al.*, 1997; Baracco *et al.*, 1995). This communication reports the development of a new HPLC/DAD methodology to separate, identify and quantify the phenolic compounds from Portuguese olive fruit cultivars (*Cobrançosa*, *Madural* and *Verdeal*).

EXPERIMENTAL

Extraction of Phenolic Compounds from Olive Fruits. Each olive fruit sample (ca. 1.5 g) was thoroughly mixed with methanol until complete extraction of the phenolic compounds (negative reaction to NaOH 20%). The methanolic extract was filtered, concentrated to dryness under reduced pressure (40°C) and redissolved in acid water (pH 2 with HCl) (≈ 50 mL). The aqueous solution was then passed through an ISOLUTE C18 (NEC) column, previously conditioned with 60 mL of methanol and 140 mL of acid water (pH 2 with HCl). The loaded cartridge was washed with n-hexane and phenolic compounds were eluted with methanol. The methanolic extract was evaporated to dryness under reduced pressure (40°C), redissolved in methanol (4 mL) and 20 µL were analysed by HPLC.

HPLC Analysis of Phenolic Compounds. Separation of phenolics was achieved with an analytical HPLC unit (Gilson), using a Spherisorb ODS2 (25.0 x 0.46 cm; 5µm, particle size) column. The solvent system used was a gradient of water-formic acid (19:1) (A) and methanol (B): 0' - 5% B, 3' - 15% B, 13' - 25% B, 25' - 30% B, 35' - 35% B, 39' - 40% B, 42' - 45% B, 45' - 45% B, 50' - 47% B, 60' - 48% B, 64' - 50% B, 66' - 100% B. The solvent flow rate used was 0.9 mL/min. Detection was achieved with a Gilson diode array detector (DAD), and chromatograms were recorded at 280 and 320 nm. Phenolic compounds quantification was achieved by the absorbance recorded in the chromatograms relative to external standards.

RESULTS AND DISCUSSION

The results demonstrate that the *Cobrançosa* cultivar has the same qualitative composition as *Madural*, being characterised by the presence of nine identified compounds: hydroxytyrosol, 5-*O*-caffeoylquinic acid, verbascoside, luteolin 7-*O*-glucoside, oleuropein, rutin, apigenin 7-*O*-glucoside, quercetin 3-*O*-rhamnoside and luteolin (Figure 1). *Verdeal* cultivar exhibited a similar phenolic composition, but verbascoside was not present.

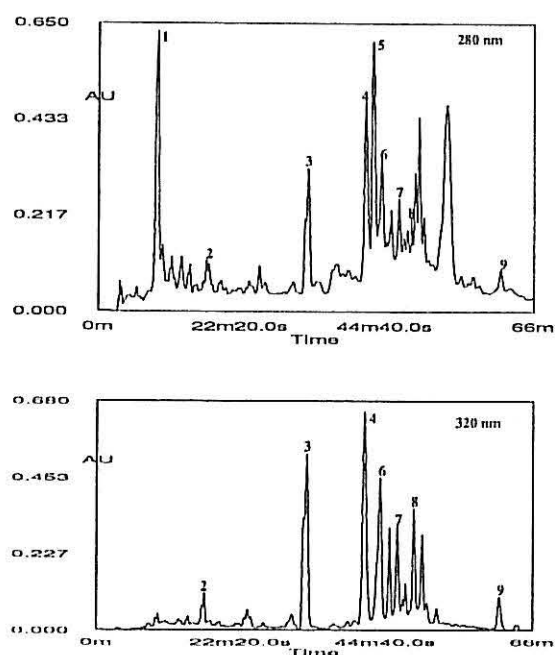


Figure 1. HPLC phenolic profile of a olive fruit sample. (1) hydroxytyrosol; (2) 5-*O*-caffeoylquinic; (3) verbascoside; (4) luteolin 7-*O*-glucoside; (5) oleuropein; (6) rutin; (7) apigenin 7-*O*-glucoside; (8) quercetin 3-*O*-rhamnoside; (9) luteolin.

ACKNOWLEDGMENTS

Branca M. Silva is grateful to Fundação para a Ciência e a Tecnologia for a grant (PRAXIS XXI/BD/21339/99).

REFERENCES

- A. Romani, N. Mulinacci, P. Pinelli, F.F. Vincieri, A. Cimato, *J. Agric. Food Chem.* **47** (1999), 964-967.
- C. Soler-Rivas, J.C. Espín, H. Wichers, *J. Sci. Food Agric.* **80** (2000), 1013-1023.
- Capasso, R., Evidente, A., Scognamiglio, F. *Phytochem Anal.* **3** (1992), 270-275.
- P. Ficarra, R. Ficarra, A. de Pasquale, M.T. Monforte, M.L. Calabro, *Farmaco* **46** (1991), 803-815.
- N. de Laurentis, G. Crescenzo, O.R. Lai, M.A. Milillo, *Pharm. Pharmacol. Lett.* **7** (1997), 27-30.
- A. Baracco, G. Bergin, E. Gnocco, M. Legorati, S. Sedocco, S. Catlinella, D. Fravretto, P. Traldi, *Rapid Commun. Mass Spectrom.* **9** (1995), 427-436.