

# Organic compounds isolated from Juniperus brevifolia bark

Amélia Vieira,<sup>a</sup> Marcele Martins,<sup>a</sup> Ana M. L. Seca,<sup>a,b</sup> Diana C. G. A. Pinto,<sup>a</sup> Artur M. S. Silva<sup>a</sup>





<sup>a</sup>Department of Chemistry & QOPNA, University of Aveiro, 3810-193 Aveiro, Portugal; <sup>b</sup>DCTD, University of Azores, 9501-801 Ponta Delgada, Portugal

### INTRODUCTION

In continuation of our study on the chemical characterization of endemic plants of the Azores archipelago, we have examined Juniperus brevifolia. Our interest in this plant was stimulated by the wide range of biological activities reported for other species of this genus and/or of their constituents.<sup>1</sup>

Previous work showed that leaves extracts, for instance dichloromethane extract and chloroform-soluble fraction of the acetone extract, showed high activity against HeLa and Hep-2 tumour cell lines.<sup>2</sup> Chemical investigation of these extracts afforded more than thirty compounds, mainly abietane and pimarane derivatives, eight of them were new natural compounds and other exhibit interesting antitumor activity.3



Recent investigation showed that the bark acetone extract has antioxidant activity similar to that of quercetin; and also showed activity against *Bacillus* cereus, B. subtilis and Micrococus luteus. Furthermore the bark acetone extract showed higher anti-AChE activity. These findings prompted us to study bark extracts.

### RESULTS AND DISCUSSION

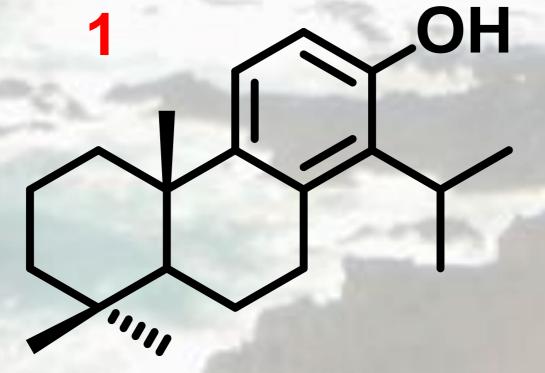
From the hexane and acetone extracts of *Juniperus brevifolia* bark were isolated several compounds (1-7) by preparative chromatographic techniques.

 $\delta_{\rm C}$  26.7

<sub>C</sub> 133.4

The structural unequivocal identification of these compounds (1-7) were obtained by detailed spectroscopic methods.

**(1)**, previously **Totarol** identified by GC-MS in bark of hexane extract Juniperus brevifolia, is a potent antibacterial and fungicide diterpene.<sup>5</sup>



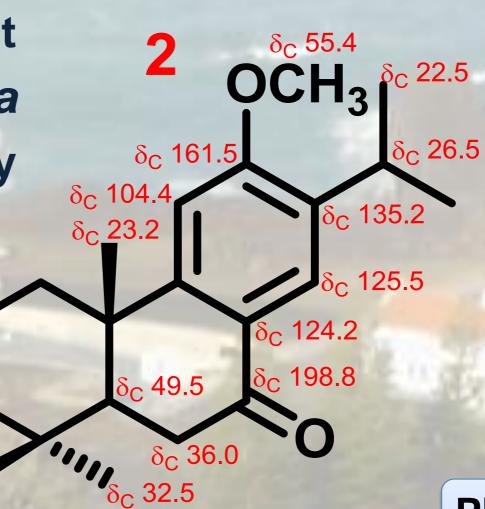
 $\delta_{\rm C}$  37.

In conclusion, from the bark of *Juniperus brevifolia*, were isolated 7 compounds, mainly diterpenes.

Compound 2 is identified for the first time in *Juniperus brevifolia* while compound 3 is reported here for the

PO

12-Methyl-sugiol ether (2), was first isolated from Juniperus formosana and exhibits antibacterial activity against MRSA.6

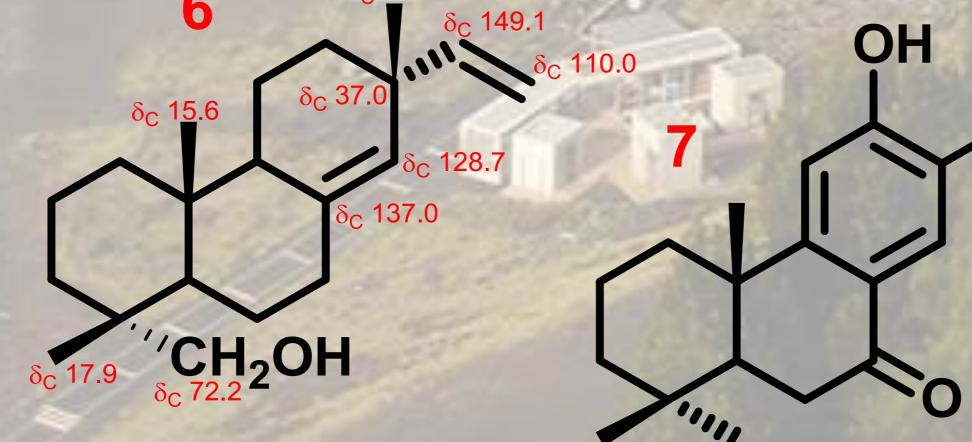


Cupresol (3), is a norditerpene with a suggested lactone ring, for androgen treatment of receptorassociated diseases.7

## MATERIAL AND METHODS Plant collected in Pedreira, S. Miguel

2001 (Azores) in June (AZB-Ana Seca-1)

Sitosterol (5), sandaracopimarinol (6) and sugiol (7) are compounds already isolated from dichloromethane extract from leaves of Juniperus brevifolia.3a Compound 6 is cytotoxic against Hela and MCF-7 tumor cell lines and antibacterial.3b



triacylglicerol of saturated fatty acid (4), was also isolated from bark of Juniperus brevifolia.

Bark (517 g) was air-dried, powered and extracted with hexane, CH<sub>2</sub>Cl<sub>2</sub>, acetone and EtOH using a soxhlet apparatus (24 h each)

**Hexane extract** (8.3 g)

Acetone extract (25.9 g)

Silica CC gel (gradient mixtures hexane and ethyl acetate)

Liquid-liquid partion with hexane, CH<sub>2</sub>Cl<sub>2</sub> and CHCl<sub>3</sub>.

Silica gel TLC (different polarity eluent mixtures)

Compounds

CH<sub>2</sub>Cl<sub>2</sub> fraction

Silica gel CC and TLC (different polarity eluent mixtures)

Compounds 1,

first time in Juniperus genus. The presence of diterpenes, well known for their bioactivities, can partially explain the durability of the J. brevifolia wood when protected by the bark and showed the potential of this species as source of drugs.

### REFERENCES

1. Seca A. M. L.; Silva A. M. S. The chemical composition of the *Juniperus* genus (1970–2004). In: Govil JN, Singh VK, editors. Recent progress in medicinal plants, vol 16-phytomedicines. Houston: Studium Press LLC; 2006. p. 401–522.

2. Moujir L. M.; Seca A. M. L.; Silva A. M. S.; Barreto M. C. Planta Med. 2008, 74, 751-753.

3. a) Seca A. M. L.; Silva A. M. S.; Bazzocchi I. L.; Jimenez A. I. Phytochemistry, 2008, 69, 498-505; b) Moujir L. M.; Seca A. M. L.; Araujo L.; Silva A. M. S.; Barreto M. C. Fitoterapia 2011, 82, 225-229.

4. Oliveira N.; Medeiros S.; Barreto M. C.; Seca A. M. L.; Rosa J. S. XXIII ENSPQ, Aveiro, Junho, 2013, 214, P2.50.

5. Seca, A. M. L.; Silva, A. M. S. Nat. Prod . Res., 2008, 22, 975-983.

6. Kuo, Y.-H.; Yu, M.T. Phytochemistry, 1996, 42, 779-781.

7. Hsiao, P.-W.; Lin, F.-M.; Wang, S.-Y. Patent USA no, 2010/0022632 A1, **2010**.









### **ACKNOWLEDGEMENTS**

Thanks are due to Fundação para a Ciência e a Tecnologia (FCT, Portugal), the European Union, QREN, FEDER, COMPETE, for funding the Organic Chemistry Research Unit (QOPNA) (project PEst-C/QUI/UI0062/2013) and the Portuguese National NMR Network (RNRMN). M. Martins is grateful to CNPq, Conselho Nacional de Desenvolvimento Científico e Tecnológico – Brasil, for the scholarship granted.