

# **Session IX**

**PROPAGATION AND *IN VITRO* MANIPULATION**

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## MICROPROPAGATION OF SELECTED ADULT PLANTS OF *ARBUTUS UNEDO* L. (STRAWBERRY TREE)

Filomena GOMES<sup>1\*</sup>, Maria L. LOPES<sup>2</sup>, Jorge AGRELA<sup>1</sup>, Jorge M. CANHOTO<sup>2</sup>

<sup>1</sup>CERNAS, Dep. Florestal, Escola Superior Agrária de Coimbra, Bencanta, 3040 – 316 Coimbra, Portugal, <sup>2</sup>Instituto do Ambiente e Vida, Departamento de Botânica, Faculdade de Ciências e Tecnologia, Universidade de Coimbra, 3001-455 Coimbra, Portugal  
[fgomes@esac.pt](mailto:fgomes@esac.pt)

*Arbutus unedo* L. is a species characteristic of Mediterranean climates, widely represented in southern Europe. The genus *Arbutus* (*Ericaceae*) includes about 20 species from which *Arbutus unedo*, known as strawberry tree, is the most interesting from an economic point of view. Several aspects contribute to the economic interest of this plant including 1) fruit production (fresh or processed to make jellies); 2) production of a spirit called “Medronheira”, that represents the main income for strawberry tree producers and (3) the more recent utilization of shoots in the floral industry. Moreover, the plant is highly resistant to forestry fires, as a result of its low resin content and its ability to produce new shoots after burning. From an ecological perspective it should also be mentioned that *A. unedo* has the capacity to grow in poor as well as in water deficient soils making the tree an ideal species to recover degraded lands and to prevent forestry fires.

Adult plants of strawberry tree, growing in different regions of Portugal were selected for its potential for fruit production. Branches (30 – 40 cm length) of these trees were collected in the field and maintained in the greenhouse until epicormic shoots start to develop. Following sterilisation, shoot tips (< 2 mm) and nodal segments (10-20 mm) were then used to establish the *in vitro* cultures.

Best results (38.7 ± 9.8 %, survival rate) were obtained when shoot tips (< 2 mm) were used. Optimum shoot proliferation was achieved on a basal De Fossard medium (FS; De Fossard *et al.*, 1974), containing Murashige and Skoog (1962) micro-nutrients, FS organics, sucrose, and 9 µM benzyladenine (BA). Rooting of the formed shoots occurred following auxin treatments. The highest root (93.3 %) rates were achieved when shoots were inoculated in root induction medium, Knop (Gautheret, 1959), with 24.7 µM 3-indolebutyric acid (IBA, during 6 days) or dipped on 9.8 x10<sup>3</sup> µM IBA (for 15 sec), and followed by its subculture (5 weeks) on the same medium without growth regulators and containing charcoal (1.5 %). Rooted plantlets were transferred to pots and 84.7 ± 4.6 % of them acclimatized.