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## MICROPROPAGATION OF DIFFERENT AROMATIC PLANTS

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### Introduction

Aromatic plants have been used for centuries as species, natural flavor, raw material for essential-oil industry and other purposes. Some plants are endowed with aroma characteristics and this is where the definition aromatic comes from. Micropropagation of aromatic plants has advantage over conventional propagation because of high multiplication rate, but it depends on the performance of the starting material, media composition, growth regulators and environmental factors.



### Results and discussion

Different explants of Eruca sativa L., cultured on MS supplied with certain concentrations and combinations of growth regulators, proliferated in shoots, leaf rosettes, roots and callus. The apical meristem and hypocotils of *Coriandrum sativum* L. gave shoots, leaf rosettes and callus when cultured on MS with 1 mg/L Kin.

The explants for Rosmarinus sp. did not show reaction on the cultivation media, while seeds of selected genotype of Origanum vulgare L. and Metha piperita L. did not germinate.

#### Conclusion

Micropropagation is an alternative method to traditional propagation and it offers improvements over traditional vegetative propagation because of faster rate of multiplication. In this research, Eruca sativa L. and Coriandrum sativum L. are species with the highest potential for in vitro micropropagation when cultivated on selected media with addition of different combinations of growth regulators.

### Materials and methods

Selected genotypes of salad rocket, coriander, rosemary, oregano and peppermint were tested for their micropropagation potential on different media supplied with different concentrations of growth regulators. Apical buds and meristem, cotyledons and hypocotyls were used as starting material and/or explants.

The starting material/explants were object of certain sterilization protocol and cultured on different media supplied with different concentrations of growth regulators. After cultivation, the explants were placed in growth chamber with controlled conditions for future development and observation.







no germination

Table 1. Species of medical plants micropropagated in in vitro conditions on different media

Species	Explants/ Starting material	Medium + growth regulators (mg·l <sup>-1</sup> )	Results
	apical meristem	MS + 1 mg/L BAP	roots
			leaf rosettes
		MS + 1 mg/L BAP + 0,5 mg/L IAA	shoots
			roots
			callus
	hypocotyls		leaf rosettes
		MS + 1mg/L BAP	shoots
			callus
Coriandrum sativum L.	apical meristem		shoots
		MS + 1mg/L Kin	leaf rosettes
			callus
	hypocotyls		
		MS + 1mg/L Kin	shoots
Rosmarinus sp.	apical buds	$MS + 0.1 \text{ mg/L IAA} + 5 \text{ mg/L GA}_3$	,
	apical meristem	$MS + 0.1 \text{ mg/L IAA} + 5 \text{ mg/L GA}_3$	,
	hypocotyls	$MS + 0.1 \text{ mg/L IAA} + 5 \text{ mg/L GA}_3$	,
	cotyledons	$MS + 0.1 \text{ mg/L IAA} + 5 \text{ mg/L GA}_3$	
			,
Origanum vulgare L.			
	seeds	BM	no germination

BM

seeds