04-2-22

An Efficient Clonal Propagation of *Eleutherococcus chiisanensis* via Somatic Embryogenesis using the Root Segments

Jae-hun Jeong*, Heung-Kyu Moon, Yong-Eui Choi¹, Seog Gu Son, Yong Wook Kim

Biotechnology Division, Korea Forest Research Institute,

¹Division of Forestry Resources, College of Forestry Science, Kangwon National University

Objectives

Clonal propagation of woody plants is an important tool for tree improvement, reforestation, and gene pool conservation etc. *Eleutherococcus chiisanensis* is an endangered medicinal woody plants belonging to *Araliceae*. Upto now somatic embryogenesis and plant regeneration has been achieved from the culture of zygotic embryos as initial explants in all *Eleutherococcus* plants. In this study, we developed an efficient micropropagation system via somatic embryogenesis of this species using different explants, especially focus on root explant.

Materials and Methods

- 1. Material: Eleutherococcus chiisanensis. in vitro plant, leaf, petiole, root
- 2. Methods

Callusing and embryogenic callus induction - MS medium with 1.0 mg/L 2.4-D, 0.01 mg/L TDZ, 1 g/L glutamine.

Somatic embryo induction: Hormone free 1/2 MS basal medium

Germination of somatic embryos - To induce germination, somatic embryos at cotyledonary stage were transferred to half-strength MS medium with 1 mg/L GA_3 + 0.02% activated charcol and cultured for three weeks.

Soil transfer - Converted plantlets were transferred to artificial soil mixture in a greenhouse.

Results and Discussion

Callus induction occurred on all of the explants, but embryogenic callus was observed on root explants only. The best frequency of embryogenic callus formation (15% in root explants) was achieved on MS medium supplemented with 1.0 mg/L 2,4-D, 0.01 mg/L TDZ, 1g/L glutamine. Embryogenic callus was friable and white in color. Somatic embryos could be induced on hormone free half-strength MS medium. To induce of somatic embryos germination, cotyledonary stage embryos were transferred to half-strength MS medium with 1 mg/L $GA_3 + 0.02\%$ activated charcoal and cultured for three weeks. *In vitro*-converted plantlets were acclimatized in soil mixture of vermiculite: perlite (1:1 v/v) and nurtured in a greenhouse. About 90% of the plantlets acclimatized, and successfully transferred to field condition.

^{*} Corresponding author: Heung-Kyu Moon, Tel: 031-290-1163 E-mail: hkmoon@foa.go.kr