(05-2-01)

## Plant regeneration from leaf and petiole derived callus of Pulsatilla koreana Nakai

Jeong J. H<sup>\*</sup>., Jung, S.J.<sup>1</sup>, Chakrabarty, D.<sup>2</sup>, Murthy, H.N.<sup>2</sup>, Choi, Y.E.<sup>1</sup>, Yun, E.S<sup>3</sup>., Kim Y.S. Dept. of the Development of Medicinal Resources and Horticulture, Namdo Provincial College of Jeonnam, Changhung, 529-850, Korea

<sup>1</sup>Division of Forest Resources, College of Forest Sciences, Kangwon National University, Chnchon 200-701, Korea.

<sup>2</sup>Research Center for the Development of Advanced Horticultural Technology, Chungbuk national University, Cheongju 361-763, Korea.

<sup>3</sup>Department of Biology, Kongju National University, Kongju 314-701, Korea.

## Abstract

An efficient protocol has been developed for mass propagation of *Pulsatilla koreana* from leaf and petiole derived callus. Optimal callus was developed from leaf and petiole explants on Murashige and Skoog (MS) medium supplemented with 4.52 µM 2,4- dichlorophenoxyaceticacid (2,4-D) and 2.22 µM 6-benzyladenine (BA). Adventitious shoots were regenerated (42.4%) from the surface of the callus on MS medium supplemented with 4.44 µM BA and 0.1 µM polyvinylpyrrolidone (PVP). Individual elongated shoots were rooted on half-strength MS medium containing different concentration of indole acetic acid (IAA). Regenerated plantlets with well developed shoots and roots were successfully transferred to soil. This *in vitro* propagation protocol might be useful for mass propagation as well as conservation of this plant.

<sup>\*</sup> Corresponding author: Jae-Hun Jeong, Tel: 061-860-8650, E-mail: jhjeong@namdo.ac.kr