Propagation of Camellia L. via in vitro culture H.HASSANABADI, E. MAJIJI and R. BOZORGIPOUR. seed and Plant Improvment Institute.

Starting materials for in vitro propagation of Camellia japonica were obtained from a sixty years old tree. Explants (shoot tips nodes) were selected directly from out door tree or from developed shoots in growth chamber. Contamination rate in the starting materials, developed in the growth chamber, was 55%.

These materials were decontaminated using 5% calcium Hypochlorite for 6 min. Best of three kind initial explants for developed shoots in growth chamber, were shoot tips and for out door shoots were nodal segments. Within five culture media used in this study.

WPM was better than the others. Most satisfactory shoot proliferation was achieved by 4 mgL-1 of BA. Whole harvested shoots longer than 15 mm with no apical bud produced more shoots.

Dipping the in vitro leaves in 1gl-1 solution of IBA for 30 min, and transfer to WPM medium with no browth regulatores and followed by 14 days darkness induced callus formation on the surface of the leaves.