

Establishment of Embryogenic Cell Suspension Culture of Chilli [*Capsicum annum* L. var. *accuminatum* Fingerh] for Somatic Embryogenesis

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Abstract

Chilli [*Capsicum annum* L. var. *accuminatum* Fingerh] is one of the important spice crops of Sri Lanka. Embryogenic suspension cultures of chilli were developed with an objective to induce somatic embryogenesis. Successful callus induction was obtained from both leaves and cotyledons of two weeks old seedlings in MS medium containing 1 ppm 2, 4-D, after incubation in the dark for two to three weeks. A combination of Kinetin (0.1 ppm) and 2,4-D (1 ppm) promoted callus proliferation at a high rate. Cell suspension cultures were established using 2 g of four week old leaf and cotyledon calli in 20 ml of liquid MS medium with 1 ppm 2, 4-D in 100 ml Erlenmeyer flasks. Weekly sub culturing was performed. MS medium with 2,4-D (1 ppm) stimulated embryogenesis on cotyledon callus after 12 weeks in culture. Embryogenic calli formed are pale yellow to brown, compact, organized and nodular in appearance. It comprised of small, richly cytoplasmic cells without large vacuoles. Both initiation of embryogenic cells and the subsequent development of these cells into embryoids occurred in the same MS (2, 4-D 1 ppm) medium. Within a period of five to seven days, 12 week old, 20 ml of embryogenic cell suspension produced 14 proembryoids. After 7-14 days they developed into heart stage and to mature embryoids. Plantlet development has not been observed until now in the tested MS media,

containing activated charcoal, zeatin, IBA and GA3.