ABSTRACT

Amaryllis (Crinum asiaticum) has a prominent place among ornamental bulbous species, which are quite valuable in the floriculture economy. Constant culture is required; to meet the year-round market demand for amaryllis, hence, an effective micro propagation technique is required. Twin scales were used as explants to establish optimization and in vitro propagation for amaryllis (Crinum asiaticum ev. Giant crinum or lily). The twin scaling method was used to achieve the maximum number of explants (45–50) from a single bulb in the shortest period. Explants were surface sterilized with 30% bleach solution for 10 minutes and 70 % ethanol for 45 seconds and thoroughly rinsed with autoclaved distilled water. Twin scales were surface sterilized with 70% ethanol for 5 seconds and 10 % sodium hypochlorite solution for 2 minutes and rinsed with sterilized distilled water. These twin scales were cultured on MS medium fortified with different concentrations of plant growth regulators. Cultures were incubated under optimal conditions of $25 \pm 2^{\circ}$ C temperature, 2800 lux illumination and 16/8 h light and dark period respectively. Growth of Crinum asiaticum was observed under different concentration of PGRs individually and in combinations i.e., 2, 4-D, BAP, NAA TDZ, IBA and KIN. The successful growth was noticed in the combinations of 0.4 mg L -1 BAP + 0.1 mg L-1 2, 4-D, 2.0 mg L-1 BAP + 0.5 mg L-1 NAA, 5.0 TDZ + 0.3 IAA, 3.0 mgL-1 KIN + 0.8 mgL-1 IAA, 1.0 mgL-1 KIN + 4.0 mg L-1 NAA and under 6.0 mgL-1 BAP + 0.5 mg L-1 IBA as far as the shoot induction is concerned, but the best shoot induction was observed in 0.4 mg/l BAP with 0.1 mg/l 2, 4-D. The best root growth was observed under 0.4 mg/l BAP with 0.2 mg/l 2, 4-D. Viable plantlets were transferred to pots after 2 months of culture. In 4-5 months. Plantlets has grown to the size of a matured *crinum* asiaticum. Bulb had a 98% survivability.