CALLUS INDUCTION FROM NODAL SEGMENTS OF THE ANNONA SQUAMOSA L.

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The calogenesis induction and rapid cell multiplication are fundamental requirements in plant biotechnology. This work aimed to develop an efficient method of callus induction from explants of the Annona squamosa L. For the callus in vitro induction of A. squamosa L., nodal segments (0,5-1,0 cm) were disinfected, in laminar chamber, by standard methods using ethanol/sodium hypochloride and then rinsed three times with sterile water (10 min each) and inoculated in tubes containing 35 mL of MS medium, supplemented with 100 mg.L⁻ ¹ myo-inositol, 3% sucrose, 0,7% agar and various concentrations (mg.L⁻¹) and combinations of benzilaminopurine (BAP) and naphthaleneacetic acid (NAA), the treatments. The pH of medium was adjusted to 5,8 and autoclaving during 15 min at 120°C. The tubes were kept at 25±1°C in the dark. There were seven treatments (BAP+NAA), five replications, each replication being made up of one tube and four explants/tube, in completely randomized design. The effects of treatments were evaluated from Tukey test at 5%. After three weeks was analyzed who the best percentual callus formation and embryogenic callus were obtained using media with BAP 1,25 mg.L⁻¹+ NAA 1,50 mg.L⁻¹. Embryogenic callus obtained, are characterized by their white to cream colour and their nodular structure. These results suggest that the callus *in vitro* production is possible and will suport projects with A. squamosa L., especially micropropagation.

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Key words: Annona squamosa, Benzilaminopurine, Callus, Naphthaleneacetic acid.