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TRANSFORMATION OF BEAN CALLUS BY USING HIGH-VELOCITY MICROPROJECTILES

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Abstract:

By using hypocotyl explant and vertical culture on B5 medium containing I mg/l kinetin and 2 mg/l 2,4-D, we obtained a rapidly growing callus from bean. We found that 25 mg/l hygromycin(Hgm) was enough to differentiate transgenic cells from the non-transgenic ones. The GeneBooster particle delivery system was used for the bombardment of callus and the Hgm resistance gene was used as a selectable marker. Selective test was made by using medium containing 25 and 30 mg/l Hgm to screen for transformants calli. After selection on Hgm-containing media, several Hgm resistant growing calli were obtained. Selective pressure was maintained over a period of 13 month for transformation tests.