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Efficient Microorganisms for Bioethanol Production from the Natural Environment of Sri Lanka

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sustainable socio-economic development of the country is extensive. The vast microbial Sri Lanka is biologically diverse. The potential of utilizing this rich biodiversity for application of native microorganisms is still underutilized. The objective of this study was diversity is a key component in biological diversity. However, the industrial scale cellulase production. Yeasts were isolated from local fruits viz: grapes, oranges and ethanologenic yeast for application in bioethanol production. Fifty fungi were isolated analysis was conducted using Milli-Q water in 0.6 ml/minute flow rate as the solvent. Hifermentation medium. Detection and quantification of ethanol were done by High mangoes. Ethanol production by each yeast isolate was assessed in a glucose containing from soil. The total cellulase activity of fungal isolates was determined to compare significantly different from A. niger. Among six ethanologenic yeast, the highest ethanol cellulase activity of Penicillium oxallicum was, 0.438 FPU/ml, which was second most efficient cellulase producer with 0.464 FPU/ml total cellulase activity. Total total cellulase activity of 0.574 FPU/ml followed by the Aspergillus niger, being the Aspergillus and Penicillium. Trichoderma viridae was the most efficient isolate giving a results, the highest total cellulase activities were given by fungal genera Trichoderma detection. Both were maintained at 55 °C temperature for the analysis. According to the Performance Liquid Chromatography (HPLC) using ethanol standards. to release fermentable sugars for yeast. efficient isolates in bioethanol production because cellulolytic fungi can degrade cellulose isolates reported ethanol yields above 2%. There is a great potential of applying these concentration was given by Y3 isolate as 9.651% while Y5 showed 5.84 %. All the Plex H, 300×7.7 mm column was used with Refractive Index Detector for ethanol explore microbial flora of Sri Lanka to isolate efficient cellulolytic fungi The

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