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ABSTRACT

Group B streptococci (GBS) are causative agents in a

wide variety of human infections. Neonatal sepsis and meningitis are of particular concern because of their high incidence and

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Rapid Identification of Group B Streptococcus

by Using Purified Sida Tomato Lectin Resgent

Compared with Streptex B Reagent

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mortality rates. Reliable and rapid identification of GBS in primary culture from colonized mothers are needed for the protection of neonatal infections; and in this study used a tomato lectin reagent to identify GBS. From chemical and biological studies, an affinity purified lectin from variety Tomato Sida was found to be a glycoprotein with molecular weight 71,000 daltons and consisted of 63% protien and 37% carbohydrate (arabinose 75%; fructose 25%). Amino acid analysis revealed an high abundance of serine. Its properties were closely related to, but not identical with the common tomato, a different variety reported by other investigators. This lectin was coupled with polystyrene latex particles to yield an agglutination reagent for the identification of GBS. A total of 368 bacterial strains were tested, their grouping identity was previously

determined by the standard methods; and in a blind study, the results were comparable to the grouping results obtained by Streptex B reagent as reference method of detecting GBS. It was shown that 151 of 160 GBS gave strong and rapid agglutination reactions with the lectin-latex reagent (94.4 % sensitivity) and none of the 150 other serogroups of beta-hemolytic streptococci caused agglutination (100% specificity). However, cross-reaction was observed in all strains of Streptococcus pneumoniae with alpha-hemolysis; thus it was to use suggested this reagent for detecting only isolates presumptively identified as betahemolytic streptococcus. In addition, 9 of 160 GBS showed weakto - moderate (1+ to 2+) agglutination reactions, thus it should be confirmed by serological grouping.

The test using tomato lectin-latex reagent had similar efficacy with the other three conventional tests, i.e. Christie, Atkins, Munch-Petersen (CAMP), hippurate hydrolysis, and pigment production, for differentiating group B from other serogroups of beta-hemolytic streptococci. The benefit over the conventional tests is in its ease, rapidity, and low cost. When combined with colonial morphology and hemolytic reaction, this test reagent appears to provide a highly rapid, simple, specific and cost-effective means and can be used as an alternative method of identifying GBS from primary cultures for small clinical laboratories.