PRINCIPLE / DISCUSSION:

The "Streptococcus milleri group," long recognized as more virulent than oth.er viridans streptococci and often associated with abscess formation, has gone through numerous name changes and taxonomic challenges. Recently, however, they have been recognized as a legitimate taxonomic group of three species that appear to be associated with distinctive clinical syndromes (2). They share several phenotypic characteristics that differentiate them from other streptococci; all Lancefield group F streptococci are "S. Milleri group," but the converse is not true (4). Members of the group may possess Lancefield group antigens other than F. "S. milleri" can exhibit any type of hemolysis on blood agar. A distinctive caramel-like odor may be observed in some isolates, which serves as a preliminary clue to their identity. All viridans streptococcal isolates from sterile body sites and those associated with abscesses, pneumonia, and other serious disease should be identified. Fluo-Milleri substrates are designed to differentiate the three Streptococcus milleri group species based on the enzymes produced by each (1,3,5). The last three substrates are all 4 methylumbelliferyl (4-MU) fluorogenic compounds. Enzymes produced by the organisms attack and degrade the target substrate bound to 4-MU, which releases blue fluorescent 4-methylumbelliferone. This blue fluorescence can be observed under a hand-held long- wave-length UV lamp (i.e. Wood's Lamp). Colonies must be identified as belonging to the "S. milleri group" before they can be tested. The following minimal characteristics are necessary to assign an isolate to this group: Gram positive cocci (may be elongated) in pairs and chains; Catalase negative; Voges-Proskauer positive; Arginine positive; Sorbitol negative and PYR negative (first well.)

MATERIALS REQUIRED:

FLUO-CARD Milleri test is provided in sets of 10 or 20 trays. FLUO-CARD Milleri requires fresh 24 hour growth of the organism on appropriate agar media, usually a nutrient agar base with blood or other comparable enrichment additive. A loop or stick for harvesting the organisms and distilled water, pH 6.5 to 7.5 are needed but not provided.

LIMITATIONS / COMMENTS:

Only viridans streptococci of the "*milleri* group" should be tested using this system. Other organisms may yield positive results in any of the test wells. Occasionally other colors of fluorescence occur. These reactions are not due to the respective enzyme activity and should be disregarded. If in doubt, hold the product label under the Wood's lamp as a color guide.

STORAGE:

Store at 2-8C in original packaging. It is not necessary to thaw before use. Do not allow the reagent to freeze. **PROCEDURE:**

(1) Put 2 drops of water in each circle.

(2) Smear 5-10 isolated colonies (a visible paste) of the test organism on each circle using a fresh loop or stick for each circle.

(3) Incubate covered for 15-20 minutes at 35-37^oC.

(4) After incubation, add 1 drop of developer to well one. Observe for the formation of a bright pink color which indicates a positive PYR. A positive test will also fluoresce a bright orange color under the Wood's lamp. If the PYR is positive, the organism is not any strain of *Strep milleri* and no guidelines are presented here for other organisms. Do not proceed.

(5) Observe for fluorescence under a woods lamp stopping at the first positive well as indicated on the reaction chart.

QUALITY CONTROL:

Dispose of all used material in a manner appropriate for bacterial contamination. We recommend testing each lot. Some suggested strains are listed on the flow chart.

REFERENCES:

(1) Whiley, R.A., K. Fraser, J. M. Hardie, and K. Beighton. 1990. Phenotypic differentiation of Streptococcus intermedius, Streptococcus constellatus and Streptococcus anginosus strains within the "Streptococcus milleri" group. J. Clin. Microbiol. 28:1497-1501.

(2) Whiley, R.A., D. Beighton, T.G. Winstanley, et al. 1992. Streptococcus intermedius, Streptococcus constellatus and Streptococcus anginosus (the "Streptococcus milleri" group): association with different body sites and clinical infections. J. Clin. Microbiol. 30:243--244.

(3) Beighton, D., and R. Whiley. 1990. Sialidase activity of the "Streptococcus milleri group" and other viridans group streptococci. J. Clin. Microbiol. 28: 1431-1433.

(4) Ruoff, K. and M. J. Ferraro. 1986. Presumptive identification of "Streptococcus milleri" in 5 h. J. Clin. Microbiol. 24:495-497.

(5) Flynn, Cynthia E. and Ruoff, Kathryn L.. 1995. Identification of "Streptococcus milleri" Group isolates to the Species Level with a Commercially Available Rapid Test System. J. Clin. Micro Vol 33:10.

FLOW CHART							
1.(-) ⇔	2.(-) ⇔	3.(-) ⇒	4.(-)				
(+) sто Д	DP (+)STOP	(+ _п) stop	_(+) stop				
₹5	₹	₹5	₹ ≻				
	Strep.	Strep.	Strep.				
Not Milleri	Intermedius	anginosus	constellatus				

EXPECTED REACTIONS: % POSITIVE

ORG/Strain 1.Pyr	2. ß-fuco.	3. ß-gluco.	4. α -gluco.
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S. Intermedius 27335	0	90	47	100
S. Anginosus 33397	0	0	96	19
S. Constellatus 27823	0	0	4	90



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