# UREA BROTH FOR AFB

## **INTENDED USE**

Remel Urea Broth for AFB is a liquid medium recommended for use in qualitative procedures to differentiate Mycobacterium species based on urea hydrolysis.

## **SUMMARY AND EXPLANATION**

In 1979, Steadham developed a urease test to aid in the identification of mycobacteria. This broth test was formulated to achieve more consistent results when testing for urease activity. The determination of urea hydrolysis (release of ammonia) is an aid in the characterization of mycobacterial strains, in particular, for distinguishing scotochromogens and nonphotochromogens.

## **PRINCIPLE**

Urea Broth for AFB is a highly buffered medium which aids in the differentiation of mycobacterial strains after 7 days incubation. Peptone and dextrose supply essential nutrients required for growth of mycobacteria. Polysorbate 80 is a surfactant which aids in the dispersion of aggregates. The low pH of 5.8 serves to facilitate a clear-cut differentiation of positive and negative results. When urea is hydrolyzed ammonia is released. The resulting alkalinity is evidenced by the phenol red indicator changing from yellow to dark pink or red.

## **REAGENTS (CLASSICAL FORMULA)\***

Urea	g	Monopotassium Phosphate
Sodium Chloride5.0	g	Phenol Red 1%10.0 mg
Dextrose1.0	g	Polysorbate 800.1 ml
Gelatin Peptone	g	Demineralized Water1000.0 ml

pH 5.8 ± 0.2 @ 25°C

- Inoculate a spadeful of growth from a young, actively growing culture of mycobacteria into Urea Broth for AFB and vigorously mix against the bottom and sides of the tube. Positive and negative control tubes should be inoculated with each test run.
- Incubate tubes aerobically (not in CO<sub>2</sub>) at 33-37°C for up to 7 days. Incubate an uninoculated control tube with each test run.
- Observe the broth for a pink-red color development at 1, 3, and 7 days.

### INTERPRETATION OF THE TEST

Positive Test - A dark pink to red color development Negative Test - No color change, broth remains yellow

## **QUALITY CONTROL**

All lot numbers of Urea Broth for AFB have been tested using the following quality control organisms and found to be acceptable. Testing of control organisms should be performed in accordance with established laboratory quality control procedures. If aberrant quality control results are noted, patient results should not be reported.

CONTROL **INCUBATION RESULTS** Mycobacterium fortuitum ATCC® 6841 Aerobic, 3-7 days @ 33-37°C Positive Aerobic, 3-7 days @ 33-37°C Negative

Mycobacterium gordonae ATCC® 14470

## **BIBLIOGRAPHY**

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- Isenberg, H.D. 2004. Clinical Microbiology Procedures Handbook. 2<sup>nd</sup> ed., Vol. 2. ASM Press, Washington, D.C.

Refer to the front of Remel Technical Manual of Microbiological Media for General Information regarding precautions, product storage and deterioration, specimen collection, storage and transportation, materials required, quality control, and limitations.

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<sup>\*</sup>Adjusted as required to meet performance standards.