# **BAIRD-PARKER AGAR**

## **INTENDED USE**

Remel Baird-Parker Agar is a solid medium recommended for use in qualitative procedures for selective and differential isolation and presumptive identification of coagulase-positive staphylococci.

## SUMMARY AND EXPLANATION

Baird-Parker Agar was developed in 1962 for isolation and enumeration of coagulase-positive staphylococci from foods and other materials.<sup>1</sup> It is a modification of tellurite glycine medium developed by Zebovitz, Evan, and Nivan.<sup>2</sup> Baird-Parker Agar is recommended for use in the examination of foods and other materials by Food and Drug Administration (FDA), the American Public Health Association (APHA), and in the *Manual of Clinical Microbiology*.<sup>3-7</sup>

## PRINCIPLE

Beef extract, casein peptone, and yeast extract are sources of nitrogen, carbon, sulfur, vitamins, and trace minerals. Sodium pyruvate and glycine enhance the growth of *Staphylococcus aureus*. Lithium chloride and potassium tellurite are selective agents which inhibit most bacteria other than *S. aureus*. Potassium tellurite is also a differential agent which is reduced to metallic tellurium by *S. aureus* resulting in black colonies. Egg yolk emulsion provides for demonstration of the proteolytic action of coagulase-positive staphylococci, evidenced by a clear zone around the colonies. On further incubation, many strains of *S. aureus* form opaque zones within the clear zones as a result of lecithinase or lipase activity.

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

Glycine	g
Casein Peptone	g
Sodium Pyruvate10.0	ğ
Beef Extract	ğ
Lithium Chloride	g

Yeast Extract	1.0	g
Egg Yolk Suspension	50.0	mĬ
Potassium Tellurite 1%	10.0	ml
Agar	20.0	g
Demineralized Water	1000.0	mĬ

pH 7.0 ± 0.2 @ 25°C

\*Adjusted as required to meet performance standards.

## PROCEDURE

- 1. Inoculate and streak the specimen as soon as possible after it is received in the laboratory.
- 2. If the material is being cultured from a swab, roll the swab over a small area of the agar surface and streak for isolation.
- 3. Incubate plates aerobically at 33-37°C for 24-48 hours.
- 4. Observe plates for characteristic colonial morphology and color.

## INTERPRETATION OF THE TEST

Positive Test - Black colonies with clear to opaque zones Negative Test - White to brown colonies

# QUALITY CONTROL

All lot numbers of Baird-Parker Agar have been tested using the following quality control organisms and have been 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

Staphylococcus aureus ATCC<sup>®</sup> 25923 Staphylococcus epidermidis ATCC<sup>®</sup> 12228 Escherichia coli ATCC<sup>®</sup> 25922

## INCUBATION

Aerobic, 18-24 h @ 33-37°C Aerobic, 18-24 h @ 33-37°C Aerobic, 18-24 h @ 33-37°C

# RESULTS

Black colonies with clear to opaque zones Inhibition (partial to complete) Inhibition (partial to complete)

# LIMITATIONS

1. Coagulase-negative Staphylococcus may grow on Baird-Parker Agar, however, it does not form clear zones around colonies, which is a characteristic of S. aureus.<sup>8</sup>

## BIBLIOGRAPHY

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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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12076 Santa Fe Drive, Lenexa, KS 66215, USA General Information: (800) 255-6730 Website: <u>www.remel.com</u> Email: remel@remel.com Local/International Phone: (913) 888-0939 International Fax: (913) 895-4128