

Distribution: Central File

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OXOID QUALITY ASSURANCE

PRODUCT SPECIFICATION

BRILLIANCE™ UTI AGAR (OXOID)

CM0949

Formula

Peptone	grams per litre	15.0
Chromogenic mix		26.3
Agar		15.0

Directions

Suspend 56.3 g in 1 litre of distilled water and mix well. Sterilize by autoclaving at 121°C for 15 minutes. Cool to approximately 50°C. Mix well to resuspend and pour into sterile Petri dishes.

Physical Characteristics

Straw, free flowing powder
 Colour on reconstitution - pale buff
 Moisture level - less than 7%
 pH - 6.8 ± 0.2 at 25°C
 Clarity - opaque
 Gel strength - firm, comparable to 15g/litre Agar

Bacteriological Tests Using Optimum Inoculum Dilution

Control Medium: Tryptone Soya Agar

Reactions after incubation at 37°C for 18 hours

Medium is challenged with 10-100 colony forming units

<i>Escherichia coli</i>	ATCC® 25922	1-2mm pink colonies
<i>Enterobacter aerogenes</i>	ATCC® 13048	1-2mm dark blue/purple colonies
<i>Proteus mirabilis</i>	NCTC 10975	0.5-1.5mm straw colonies, brown halo
<i>Enterococcus faecalis</i>	ATCC® 29212	0.5-1mm blue/green colonies
<i>Staphylococcus aureus</i>	ATCC® 25923	0.5-1mm white colonies
<i>Citrobacter freundii</i>	NCTC 8581	1-1.75mm blue/purple colonies

A satisfactory result is represented by recovery of equal to or greater than 70% of the control medium.

Indole may be detected by removing a few colonies, spreading onto filter paper and adding 1-2 drops of DMACA Indole reagent (dimethylamino cinnamaldehyde). *Escherichia coli* should be positive (blue/green) and *Enterobacter aerogenes* negative (colourless/pink).