



#### Also known as

BP Agar; Egg Yolk Tellurite Glycine Piruvate Agar; ETGP Agar

#### **Specification**

Selective culture medium for the screening of staphylococci from a variety of samples, according to Pharmacopeial Harmonized Methods, ISO and DIN standards.

# Formula \* in g/L

Tryptone	10.0
Sodium pyruvate	10.0
Glycine	12.0
Meat extract	5.0
Lithium chloride	5.0
Yeast extract	1.0
Agar	17.0

Final pH 7.2 ±0.2 at 25 °C

#### Directions

Suspend 60 g in 950 mL of distilled water. Allow to soak and bring to the boil stirring constantly. Sterilize in the autoclave at 121°C for 15 minutes. Cool to 50°C and add 50 mL of Egg Yolk Tellurite Sterile Emulsion (Art. No. 06-026 or 064-BA1018). Homogenize and distribute into plates. Once prepared, the medium must not be reheated nor sterilized again.

# **Description**

Baird Parker Agar Base is recommended for the detection and enumeration of staphylococci in food and other material, since it allows a good differentiation of coagulase-positive strains. The growth of the accompanying bacteria is usually suppressed by the high concentration in lithium, glycine and pyruvate. Lithium and glycine enhances the growth of staphylococci. Occasionally the medium may grow some *Bacillus species*, yeast and very rarely, *Proteus*. The growth of *Proteus species* can be suppressed by adding 50 mg/L of sulphamethazine.

The presence of tellurite and egg yolk, which must be added to the medium after sterilisation, allows the differentiation of presumptive pathogenic staphylococcal colonies. There is a high correlation between the coagulase test and the presence of clear zones of lypolysis in this medium, which is due to the staphylococcal lecithinase. Studies show that almost 100% of coagulase-positive staphylococci are capable of reducing tellurite, which produces black colonies, whereas other staphylococci can not always do so.

For the Coagulase test, add 1 vial of RPF Lyophilized Supplement (064-TA0155) in 90 ml of the Baird Parker Agar Base (01-030), and proceed according to the instructions for use of the supplement.

# **Technique**

The inoculation is carried out by spreading 0.1 ml of sample over each plate with a Drigalsky loop. After 24-48 hours of incubation à 37±1°C, select the colonies which are black, shiny and convex with regular margins surrounded by a clear zone. These can be presumbly identified as coagulase-positive Staphylococcus aureus.

Colonial appearance after 24-48 hours à 37±1 °C:

- Staphylococcus aureus: Black, shiny, convex, regular margins, 1.0-1.5 mm diameter, surrounded by a clear zone of lipolysis 2-5 mm in width. Wide opaque zones of precipitate extending into the cleared medium may occur after 48 hours.
- Other species of Staphylococcus: Black, usually dull, with regular margins. Sometimes brown with zones of clearing but these present as wide opaque zones.
- Micrococcus spp: Brown, very small and without clearing zones.
- Bacillus spp: Various shades of brown, big. May produce clearing zones after 48 hours.
- Yeasts: White, big and smooth.

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<sup>\*</sup> Adjusted and /or supplemented as required to meet performance criteria



Reference: 01-030

Product :

**BAIRD PARKER AGAR BASE** 

# Scharlau Microbiology - Technical data sheet

# **Quality control**

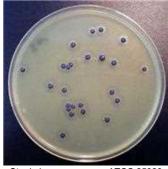
Incubation temperature:  $37 \, ^{\circ}\text{C} \pm 1$  Incubation time:  $24\text{-}48 \pm 2 \, \text{h}$ 

**Inoculum:** Practical range 100  $\pm$  20 CFU. min. 50 CFU (productivity)/ 10 $^4$  -10 $^-$  CFU (selectivity)/  $\ge$  10 $^3$  CFU

(specificity), according to ISO 11133:2014/Amd 1:2018.

### Microorganism

Escherichia coli ATCC® 8739
Staphylococcus aureus ATCC® 25923
Staphylococcus aureus ATCC® 6538
Staphylococcus epidermidis ATCC® 12228
Staphylococcus saprophyticus ATCC® 15305



Staphylococcus aureus ATCC 25923

# Growth

Inhibited
Productivity > 0.50
Productivity > 0.50
Poor to good (Specificity)
Poor to good (Specificity)

Staphylococcus aureus ATCC 25923 (Lecithinase Halos)

# Remarks

Selectivity

Black colonies; Lecithinase (+)
Black colonies; Lecithinase (+)
Black colonies; Lecithinase (-)
Black colonies; Lecithinase (-)



Staphylococcus aureus ATCC 6538

# References

- · ATLAS R.M. & L.C. PARKS (1993) Handbook of Microbiological Media. CRC Press. London.
- · BAIRD-PARKER, A.C. (1962) An improved diagnostic and selective medium for isolating coagulase-positive staphylococci. J. Appl. Bact. 25:12.
- · COLIPA (1997) Guidelines on Microbial Quality Management (MQM). Brussels.
- · DOWNES, F.P. & K. ITO (2001) Compendium of Methods for the Microbiological Examination of Foods. 4<sup>th</sup> ed. APHA. Washington. USA.
- · EUROPEAN PHARMACOPOEIA (2007) 5<sup>th</sup>ed. Suppl. 5.6 § 2.6.13 Microbiological examination of non-sterile products. EDQM. Council of Europe. Strasbourg.
- · FIL-IDF 60:2001 Standard. Lait et produits à base de lait Detection des staphylocoques à coagulase positive Technique du nombre le plus probable. Brussels.
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- · ISO 6888-1:1999/Adm.2:2018. Standard. Microbiology of food and animal feeding stuffs Horizontal method for the enumeration of coagulase-positive staphylococci ( *Staphylococcus aureus* and other species)- Part 1 Technique using Baird-Parker Agar medium. Adment 2: Inclusion of an alternative confirmation test using RPFA stab method.
- · ISO 6888-2:1999 Standard. Microbiology of food and animal feeding stuffs Horizontal method for the enumeration of coagulase-positive staphylococci Part 1 Technique using rabbit plasma fibrinogen agar medium. Geneva.
- . ISO 11133:2014/ Adm 1:2018. Microbiology of food, animal feed and water. Preparation, production, storage and performance testing of culture media.
- · ISO 22718 Standard (2015) . Cosmetics Microbiology Detection of Staphylococcus aureus.
- · USP 31 NF 26 (2008) <61> Microbial Limit Tests. US Phamacopoeial Conv. Inc. Rockville. MD. USA.
- · ZANGERL, P. & H. ASPERGER (2003) Media used in the detection and enumeration of *Staphylococcus aureus*. In Handbook.

#### Storage

For laboratory use only. Keep tightly closed, away from bright light, in a cool dry place (+4 °C to 30 °C).