

## TECHNICAL DATA SHEET

# EASY STAPH AGAR

### ENUMERATION OF COAGULASE POSITIVE *STAPHYLOCOCCI*

## 1 INTENDED USE

EASY STAPH is an alternative method for the enumeration of coagulase positive *Staphylococci* in food products, in animal feeds and in environmental samples. Devoid of the need for confirmation, the method allows count of pathogenic staphylococci in 22 hours instead of the usual 48 hours in the case of the standardized method NF EN ISO 6888-2. The media support the use of surface inoculation, in depth or by the Spiral method.

For all products destined for human consumption, the method is certified NF VALIDATION, under the Attestation N°BKR 23/10 – 12/15, of which the validity is fixed at 3 December 2019.



## 2 PRINCIPLES

The base media has been specially formulated to allow the development of coagulase positive *Staphylococci* in 22 hours.

Optimization of the selective system has allowed an improvement in the inhibition of secondary flora that are traditionally found on Baird Parker plates.

The freeze-dried supplement EASY STAPH corresponds to an optimized Rabbit Plasma Fibrinogen supplement. The composition has been adjusted in order to favor the development of fibrin halo in 22 hours.

Rabbit plasma was chosen in light of its excellent specificity towards staphylococcal coagulase and its ability to rapidly form a clot with staphylothrombin from prothrombin. It is reinforced in bovine fibrinogen. The staphylothrombin acts by cutting the fibrinopeptides A & B of fibrinogen, which initiates a polymerization process that terminates in the appearance of a fibrin halo around the colonies. A trypsin inhibitor extracted from soy avoids lysis of the fibrin halos and maintains them intact for plate reading.

## 3 TYPICAL COMPOSITION

The composition can be adjusted in order to obtain optimal performance.

For 1 L of complete media

- Peptones .....	19,4 g
- Activators .....	21,7 g
- Buffering system .....	1,4 g
- Selective system .....	4,5 g
- Bacteriological agar.....	14,4 g
- Bovine fibrinogen .....	5,3 g
- Rabbit plasma, EDTA.....	25 mL
- Trypsin inhibitor .....	25 mg
- Potassium tellurite .....	25 mg

pH of the complete ready-to-use media at 25 °C : 7,3 ± 0,2.

**For 68,2 g of dehydrated base (BK216)**

- Peptones .....	21,6 g
- Activators .....	24,1 g
- Buffering system .....	1,5 g
- Selective system .....	5,0 g
- Bacteriological agar .....	16,0 g

**For one vial of supplement BS090**

(Qs 100 mL)

- Bovine fibrinogen .....	0,53 g
- Rabbit plasma, EDTA .....	2,5 mL
- Trypsin inhibitor .....	2,5 mg
- Potassium tellurite .....	2,5 ma

**For 190 mL of ready-to-use base (BM185) / kit BT012**

- Peptones .....	4,3 g
- Activators .....	4,8 g
- Buffering system .....	0,3 g
- Selective system .....	1,0 g
- Bacteriological agar .....	3,2 g

**For one vial of supplement (BS086) / kit BT012**

- Bovine fibrinogen .....	1,06 g
- Rabbit plasma, EDTA .....	5 mL
- Trypsin inhibitor .....	5 mg
- Potassium tellurite .....	5 mg

## 4 PREPARATION

**Preparation from dehydrated media :**

- Dissolve 68,2 g of dehydrated media (BK216) in 1 L of distilled or demineralized water.
- Slowly bring to boiling with constant agitation and maintain throughout the time needed to achieve complete dissolution.
- Divide into vials, 90 mL per vial, or into multiples of 90 mL.
- Sterilize in an autoclave at 121 °C for 15 minutes.
- Cool and maintain in a molten state at 44-47 °C.

✓ Reconstitution :  
68,2 g/L

✓ Sterilization :  
15 min at 121 °C

**Rehydration of freeze-dried supplements**

- Rehydrate the lyophilisate by adding aseptically a volume of sterile distilled water as shown in the following table :
  - Qsp 100 mL of complete media (BS090) : 10 mL of sterile water
  - Qsp 200 mL of complete media, specific to the BT012 kit (BS086) : 10 mL of sterile water
  - Qsp 500 mL of complete media (BS092) : 50 mL of sterile water
- Dissolution will take place quicker if warm water is used for the preparation (use room temperature water or water that has been slightly heated, not going over 44°C).
- Mix the vial well in order to achieve complete dissolution of the supplement, while avoiding the formation of foam. The dissolution may not occur immediately and the process can be accelerated by using a Vortex mixer. The freeze-dried pellet must be totally dissolved before use.

**Utilization of the kit BT01208 (Qsp 200 mL)**

- Melt the solidified media provided in the kit for the minimum amount of time necessary to achieve total liquefaction. Cool and maintain in a molten state at 44-47 °C.
- Dissolve the lyophilsate with 10 mL of sterile water.
- Add aseptically 10 mL of the specific supplement EASY STAPH (BS086) to 190 mL of base media (BM185).
- Use immediately after preparation, for the inoculation of plates using a pour plate method or pour into empty, sterile Petri plates for use with surface inoculation.

**Utilization of the kit BT01308 (Qsp 100 mL)**

- Melt the solidified media provided in the kit for the minimum amount of time necessary to achieve total liquefaction. Cool and maintain in a molten state at 44-47 °C.
- Dissolve the lyophilsate with 10 mL of sterile water.
- Add aseptically 10 mL of the specific supplement EASY STAPH (BS091) to 90 mL of base media (BM189).
- Use immediately after preparation, for the inoculation of plates using a pour plate method or pour into empty, sterile Petri plates for use with surface inoculation.

**NOTE :**

The complete media cannot be held for an extended period at 44-47 °C.

## 5 INSTRUCTIONS FOR USE

Respect all good laboratory practices (refer to the standard NF EN ISO 7218).

### Pour plate inoculation

- Transfer 1 mL of the inoculum and its serial dilutions to empty, sterile 90mm Petri plates.
- Pour roughly 15 mL of complete media previously prepared and held at 44°C.
- Mix well.
- Let solidify on a cool, flat surface.
- Incubate at  $37 \pm 1$  °C for  $24 \pm 2$  hours.

✓ **Inoculation :**  
1 mL Pour plates  
  
✓ **Incubation :**  
24 h at 37°C

**NOTE :** For reasons of laboratory management, plates can be incubated for up to 72 hours.

### Surface inoculation

- On the surface of prepared plates or if using the pre-poured media (BM187) brought to room temperature, transfer 0,1 mL of the sample to test and its serial dilutions.
- Spread across the surface of the plate with a sterile bent glass rod or "hockey stick".
- Incubate at  $37 \pm 1$  °C for  $24 \pm 2$  hours.

✓ **Inoculation :**  
0,1 mL on surface  
  
✓ **Incubation :**  
24 h at  $37 \pm 1$  °C

### NOTES :

- The method is validated also with the Spiral inoculation technology.  
The inoculation can be in logarithmic mode from 50 or 100 µL.
- The counting limits can be decreased by a factor of 10 by inoculating 1,0 mL of the sample or initial suspension to the surface of three 90mm plates.
- For reasons of laboratory management and organization, plates can be incubated up to 72 hours using the Spiral technology and up to 48 hours using manual surface inoculation.

## 6 RESULTS

Coagulase positive staphylococci are characterized on the surface as having formed white, grey or black colonies surrounded by an opaque halo of fibrin that is net, stable and well visible. Enumerate the colonies taking into consideration the standard NF EN ISO 7218.

See ANNEX 1 : PHOTO SUPPORT.

## 7 QUALITY CONTROL

**Dehydrated base media :** white/cream powder, free-flowing and homogeneous.

**Supplement EASY STAPH :** white to pink pellet, giving rise after reconstitution to a solution that is amber, limpid and slightly opaque.

**Complete, prepared media :** amber agar.

Typical culture response after 24h incubation at 37 °C

Microorganisms	Growth (Productivity ration : $P_R$ )	Characteristics
<i>Staphylococcus aureus</i> WDCM 00034	$P_R \geq 50\%$	Colonies with an opaque halo
<i>Staphylococcus aureus</i> WDCM 00032	$P_R \geq 50\%$	Colonies with an opaque halo
<i>Staphylococcus saprophyticus</i> WDCM 00159	Slowed, score 0-1	Colonies without opaque halo
<i>Escherichia coli</i> WDCM 00013	Inhibited, score 0	-

## 8 STORAGE / SHELF LIFE

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**Dehydrated base media** : 2-30 °C.

**Supplements EASY STAPH** : 2-8 °C.

**Kits** : 2-8 °C.

**Pre-poured media in plates** : 2-8 °C.

The expiration dates are indicated on the labels.

**Prepared base media in vials (\*)** : 6 months at 2-8 °C.

**Reconstituted freeze-dried supplement (\*)** : 8 days at 2-8 °C. Reheat to between 25 and 37 °C before use. The white precipitate observed at 2-8 °C will disappear when heated to between 25 and 37°C.

**Prepared, complete media in plates (\*)** : 1 month at 2-8 °C, shielded from light.

**Prepared, complete media in vials (\*)** : Use immediately.

(\*) Benchmark value determined under standard preparation conditions, following manufacturer's instructions.

## 9 PACKAGING

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### **Dehydrated base media**

500 g bottle -----	BK216HA
5 kg drum -----	BK216GC

### **EASY STAPH Supplement**

8 vials qsp 100 mL -----	BS09008
1 vial qsp 500 mL -----	BS09208

### **Kit (6 x 200 mL)**

6 x 190 mL vials of base media plus 6 freeze-dried supplements -----	BT01208
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### **Kit (6 x 100 mL)**

6 x 90 mL vials de base media plus 6 freeze-dried supplements -----	BT01308
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### **Pre-poured, complete media in Petri plates (Ø 90 mm)**

20 plates -----	BM18708
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### **Pre-poured, complete media in Petri plates (Ø 90 mm)**

120 plates -----	BM19008
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## 10 BIBLIOGRAPHY

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NF EN ISO 6888-2. Octobre 1999. Microbiologie des aliments. Méthode horizontale pour le dénombrement des staphylocoques à coagulase positive (*Staphylococcus aureus* et autres espèces). Partie 2 : Technique utilisant le milieu gélosé au plasma de lapin et au fibrinogène.

## 11 ADDITIONAL INFORMATION

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Document code : EASY STAPH\_V2

Creation date : 12-2015

Updated : 11-2016

Origin of revision : Extension of the ISO 16140 validation to all human food products



## ANNEX 1 : PHOTO SUPPORT

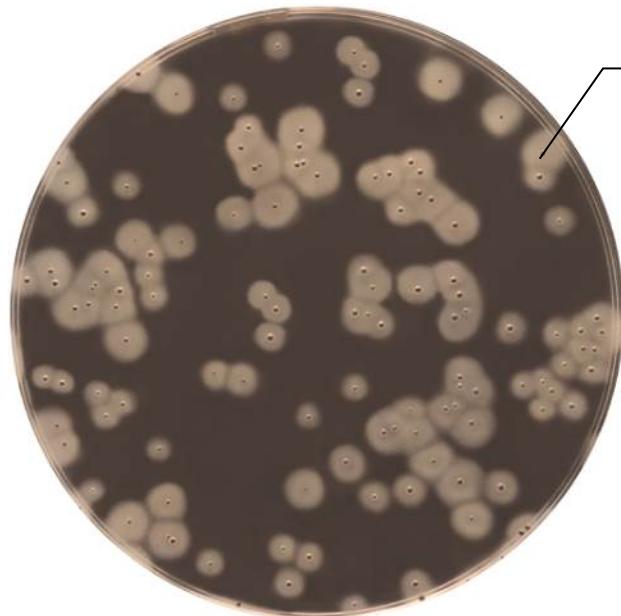
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### EASY STAPH Agar

Enumeration of coagulase positive *Staphylococci*.

#### Results :

Growth obtained after 24 hours of incubation at 37 °C.



***Staphylococcus aureus***

Characteristic colonies :  
surrounded by a halo of fibrin