

AGAR PLATES

INSTRUCTIONS FOR USE –

COLUMBIA HORSE BLOOD AGAR

REF 1033

SPML manufacture and supply a range of Pre-prepared media for the culture of microorganisms such as bacteria yeasts and moulds. SPML offer a range of media pre-poured culture media to meet the needs of its customers across a range of industries that include Clinical, food, water, pharmaceutical, and many more.

The range of plates cover selective and non-selective media and plate sizes and are intended for the use by professional parties in the isolation of microbial organisms.

INTENDED USE

A high quality non-selective, general purpose medium containing an enriched peptone mixture particularly adapted for the cultivation of nutritionally fastidious organisms. Supplemented with 6% defibrinated horse blood, typical alpha and beta haemolysis may be observed by haemolytic bacteria

• Micro Organism Reactions pH: 7.10 − 7.50

Organisms	Result, Colour of Colony
Streptococcus	Growth – Grey Alpha
Pneumoniae	Haemolytic
Streptococcus Pyogenes	Growth – Grey , Beta
	Haemolytic
Staphylococcus Aureus	Growth – White, Round
	Colony

Technique

Using a sterile loop inoculate the medium with 4±5 colonies and incubated in an aerobic atmosphere containing approximately 3 to 10 % CO2. Incubate plates at 35 ± 2 °C for 18 to 72 h. Read for the first time after 18 to 24 hours and re-incubate if necessary...

As methods and media preference may vary as to media types and test method due to jurisdictions, and personal preferences the customer should use the plates in accordance with their organisations stated methods and procedures however SPML product brochure does provide further product information

The end user should however take into consideration that selective media should, therefore, be compared with specimens/samples cultured on nonselective media to obtain additional information and help ensure recovery of potential pathogens and other significant organisms.

PRECAUTIONS

For professional use only.

Do not use plates if they show evidence of microbial contamination, discoloration, drying, cracking or other signs of deterioration.

Consult In house instructions to ensure correct application of product use is observed. Ensure GLP and aseptic handling procedures are followed, used plates should be treated as biohazards, and disposal of used product should be treated as such, and disposed of in accordance with local national regulations.

STORAGE AND SHELF LIFE

On receipt, store plates in the dark at 2 to 8° C, in their original sleeve wrapping until just prior to use. Avoid freezing and overheating. The plates may be inoculated up to the expiration date (see package label) and incubated for the recommended incubation times.

Plates from opened stacks of 10 plates can be used for one week when stored in a clean area at 2 to 8° C.

QUALITY CONTROL

SPML undertake batch release for all media checking pH, sterility, Inhibition and enhancement testing of all media batches based on the EN12322:19991¹, M22-A3², and ISO111133³,

For clinical purpose the product is **(**\(\epsilon\) marked in accordance with the requirements of 97/79/EC IVD Directive and registered with the Competent Authority MHRA in the United Kingdom.

FURTHER INFORMATION

SPML is ISO9001:2008 certified

For further information please contact your local SPML representative. Saudi Prepared Media Laboratory Company Ltd

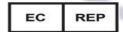
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References.

 $^{^3}$ ISO11133 $_$ Microbiology of food, animal feed and $\,$ water - Preparation, production storage and performance testing of culture media



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 $^{^{1}}$ EN 12322:1999 - In vitro diagnostic medical devices - Culture media for microbiology - Performance criteria for culture media.

²Clinical and Laboratory Standards Institute. 2004. Approved standard: M22-A3, Quality control for commercially prepared microbiological culture. media, 3rd ed





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Symbols Definition

REF	Product Reference Number
IVD	For in vitro diagnostic use
	Do not use if damaged
STERILE 1	Sterile by Moist / Dry heat
NON STERILE	Non Sterile
	Single Use
- ALL MAN	Manufacturer
LOT	Lot Number
&	Bio hazardous material
CE	CE Mark – Agai <mark>nst the Europe</mark> an In Vitro Diagnostic Medical Device Directive 98/79/EC
1	Storage temperature limitation
\sim	Manufacture date
[]i	Consult Instructions for use
类	Avoid direct sunlight
*	Keep Dry
$\mathbf{\Sigma}$	Expiration Date
EC REP	EU Representative
$\overline{\mathbb{V}}$	WARNING – Consult IFU





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