



Mast Group Ltd.
Mast House, Derby Road,
Bootle, Merseyside, L20 1EA
United Kingdom

Tel: + 44 (0) 151 472 1444 Fax: + 44 (0) 151 944 1332 email: sales@mast-group.com Web: www.mast-group.com



Mast Diagnostica GmbH Feldstrasse 20 DE-23858 Reinfeld

Tel: + 49 (0) 4533 2007 0 Fax: + 49 (0) 4533 2007 68 email: mast@mast-diagnostica.de Web: www.mast-group.com

Germany



12 rue Jean-Jacques Mention CS91106, 80011 Amiens, CEDEX 1 France

Tél: + 33 (0) 3 22 80 80 67 Fax: + 33 (0) 3 22 80 99 22 email: info@mast-diagnostic.fr Web: www.mast-group.com



MAST® *ID* ADH Agar

IDM22

Intended use

For the demonstration of amino-acid dihydrolase production.

Contents

See pack label.

Formulation*

Material:	Concentration in medium:
Peptone mixture	5.0g/litre
Yeast extract	3.0g/litre
Glucose	5.0g/litre
L-arginine hydrochloride	10.0g/litre
m-cresol purple	0.1g/litre
Agar	24.0g/litre
Final pH: 7.8 ± 0.2	

Storage and shelf life

All dehydrated culture media containers should be kept tightly closed and stored in a dry place at 10 to 25°C until the expiry date shown on the pack label.

Precautions

For *in vitro* diagnostic use only. Observe approved hazard precautions and aseptic techniques. To be used only by adequately trained and qualified laboratory personnel. Sterilise all biohazard waste before disposal. Refer to Product Safety Data sheet (available on request or via MAST® website).

Materials required but not provided

Standard microbiological supplies and equipment such as loops, MAST® selective supplements, swabs, applicator sticks, incinerators and incubators, etc., as well as serological and biochemical reagents and additives such as blood.

Procedure

- Refer to pack label for quantities and volumes required. Prepare MAST® /D ADH Agar (IDM22/A) by suspending the powder in distilled or deionised water. For sachet packs, dissolve the entire contents of the sachet in the volume shown on the label.
- Sterilise by autoclaving at 121°C (15 p.s.i.) for 15 minutes. Do not overheat any carbohydratecontaining medium.
- Mix well and pour culture plates (15 to 20ml per plate) into Petri dishes which have been labelled using the self-adhesive labels provided. Self-adhesive labels are provided in each box of preweighed sachets.
- 4. Prepared culture plates may be used immediately or stored in plastic bags at 2 to 8°C for up to one week.

- Prepare a suspension of each organism equivalent in density to a 0.5 McFarland standard. Inoculate the surface of a well-dried plate using a replicating device, e.g. the SCANURIDOT Multipoint Inoculator, to deliver each inoculum onto the agar surface.
- Allow the inoculum drops to dry before disturbing and incubate plates aerobically for 18 to 24 hours at 35 to 37°C (or alternative temperatures according to the methodology followed).

Interpretation of results

After incubation record growth and colour development in the medium. A positive result is indicated by a red/brown colour and a negative result by a yellow colour.

Quality control

Check for signs of deterioration. Quality control must be performed with at least one organism to demonstrate expected performance. Do not use the product if the result with the control organism is incorrect. The list below illustrates a range of performance control strains which the end user can easily obtain.

Test Organisms	Result
Escherichia coli	Positive
ATCC® 25922	
Pseudomonas aeruginosa	Positive
ATCC® 27853	
Shigella sonnei	Positive
ATCC® 25931	
Proteus mirabilis	Negative
ATCC® 29906	
Klebsiella pneumoniae	Negative
ATCC® 13883	

References

Bibliography available on request.