Our Chocolate Agar (Modified Martin-Lewis) is used for the selective isolation and cultivation of Neisseria species.

The first formulation of chocolate agar was devised in 1927 by McLeod, et al, which contained a combination of yeast extract and various peptones. Numerous modifications have since been made to chocolate agar to improve the recovery of Neisseria species and to increase its selectivity. Several key researchers which include Thayer, Martin, Lester, and Lewis devised formulations of chocolate agar which contained selective agents to help suppress the normal flora found in clinical specimens taken from the throat, vagina, rectum, and urethra. Our current formulation is based on the work of Martin and Lewis who improved the Thayer-Martin formulation by replacing nystatin with anisomycin. Anisomycin has superior inhibitory properties on yeasts when compared to nystatin, and possesses selective properties similar to amphotericin B.

Our formulation contains an improved casein and animal tissue digest that provide the organism with nitrogen, amino acids, and other elements essential for growth. Neisseria species are highly sensitive to toxic substances such as fatty acids; therefore the addition of cornstarch helps neutralize possible toxic metabolites, while potassium phosphate helps maintain an uniform pH during growth. Hemoglobin provides X-factor (hemin) required for growth of Haemophilus species on normal Chocolate Agar, and isovitox enrichment (Dalynn Catalogue No. VI85) provides V-factor (nicotinamide dinucleotide), cocarboxylase, and other complex compounds which enhance the growth of Neisseria species. Our VCAT supplement contains four antibiotics: vancomycin which inhibits gram-positive organisms, colistin which inhibits gram-negative bacteria, amphotericin B which inhibits yeasts and molds, and trimethoprim which is primarily added to inhibit swarming Proteus species.

**Formula per Litre of Medium**

Casein/Meat Peptone................................. 15.0 g  
Corn Starch........................................... 1.0 g  
Potassium Phosphate, Dibasic..................... 4.0 g  
Potassium Phosphate, Monobasic............... 1.0 g  
Sodium Chloride................................... 5.0 g  
Agar.................................................. 10.0 g  
Hemoglobin Solution (2%)..................... 500.0 mL  
Isovitox Enrichment.............................. 10.0 mL  
VTAC Supplement................................. 10.0 mL  

pH 7.2 ± 0.2

**Recommended Procedure**

1. Allow medium to reach room temperature.
2. Using an inoculum from the specimen, perform a four-quadrant streak to obtain well-isolated colonies. Alternatively, if the specimen is contained on a swab, roll the swab across part of the medium in a Z-pattern so that an adequate amount of the sample is transferred. Then taking a sterile loop streak the plate to disperse the sample throughout the medium.
3. Incubate at 35°C in a 5 to 10% CO₂ atmosphere.
4. Examine after 18-24 hours and again at 48 and 72 hours.
Interpretation of Results

*Neisseria* species grow while the majority of other organisms are inhibited. *N. gonorrhoeae* produces small, grey to white, mucoid colonies. *N. meningitidis* produces larger bluish-grey, mucoid colonies. Modified Martin-Lewis Chocolate Agar is a selective, primary plating medium, therefore a subculture of potential *Neisseria* colonies onto a non-selective medium is necessary so that additional biochemical and/or serological tests can be performed from pure culture.

- *Chocolate Agar contains less agar than other solid media therefore streaking should be done carefully to avoid gouging into the agar*
- *Some non-pathogenic species of Neisseria such as N. sicca are inhibited on Modified Martin-Lewis Chocolate Agar*

Quality Control

After checking for correct pH, colour, depth, and sterility, the following organisms are used to determine the growth performance of the completed medium.

<table>
<thead>
<tr>
<th>Organism</th>
<th>Expected Results</th>
</tr>
</thead>
<tbody>
<tr>
<td><em>Neisseria gonorrhoeae</em></td>
<td>Growth</td>
</tr>
<tr>
<td>ATCC 43069</td>
<td></td>
</tr>
<tr>
<td><em>Staphylococcus epidermidis</em></td>
<td>Inhibition</td>
</tr>
<tr>
<td>ATCC 12228</td>
<td></td>
</tr>
<tr>
<td><em>Proteus mirabilis</em></td>
<td>Inhibition</td>
</tr>
<tr>
<td>ATCC 12453</td>
<td></td>
</tr>
<tr>
<td><em>Escherichia coli</em></td>
<td>Inhibition</td>
</tr>
<tr>
<td>ATCC 25922</td>
<td></td>
</tr>
<tr>
<td><em>Candida albicans</em></td>
<td>Inhibition</td>
</tr>
<tr>
<td>ATCC 10231</td>
<td></td>
</tr>
</tbody>
</table>

Storage and Shelf Life

Our Chocolate Agar (Modified Martin-Lewis) should be stored away from direct light at 4°C to 8°C. The medium side should be uppermost to prevent excessive accumulation of moisture on the agar surface. Under these conditions this medium has a shelf life of 8 weeks from the date of manufacture.

References

6. Thayer JD, Lester A. Transgrow, a medium for transport and growth of *Neisseria gonorrhoeae* and *Neisseria meningitidis*. HSMHA Health Rep 1971; 86:30-3

Original: January 2001
Revised / Reviewed: October 2014