

# **Technical Information**

# **Campylobacter Agar Plate**

Product Code: PM 1994

**Application:** Campylobacter Agar Plate recommended for selective isolation of *Campylobacter* species from faecal specimens, food and environmental specimens.

Composition**
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Ingredients	Gms / Litre	
Proteose peptone	15.000	
HML extract #	2.500	
Yeast extract	5.000	
Sodium chloride	5.000	
Agar	12.000	
MS2006/MS2008		
Blood	100ml	
Final pH ( at 25°C)	7.4±0.2	
**Formula adjusted standardized to suit norformance narameter	•	

<sup>\*</sup>Formula adjusted, standardized to suit performance parameters

# **Principle & Interpretation**

Campylobacter species are ubiquitous in the environment inhabiting a wide variety of ecological niches (8). Infection with a Campylobacter species is one of the most common causes of human bacterial gastroenteritis (8). Most species are found in animals (cattle, swine) and cause infertility and abortion (7). C. jejuni was originally isolated on a blood-containing media with antibiotics (3). Skirrow described a selective medium for Campylobacter species consisting of Blood Agar Base No. 2 supplemented with horse blood and antibiotics (9). Subsequently, Blaser et al isolated *C.jejuni* on Brucella Agar supplemented with sheep blood and four antibiotics (2). Later on, a fifth antibiotic, cephalothin was added to improve the selectivity of the medium by inhibition of accompanying faecal bacteria (12). Campylobacter Agar Base is used by APHA for selective isolation of *Campylobacter* species (10). Campylobacter Agar Base is well supplemented to support luxuriant growth of *Campylobacter* species. Osmotic equilibrium of the medium is maintained by sodium chloride. Blood serves as an additional source of nutrients including X-factor. The antibiotic supplements namely Blaser-Wang (MS2006) and Skirrow (MS2008) markedly reduce the growth of normal enteric bacteria while enhancing the growth and recovery of *C.jejuni* from faecal specimens. Amphotericin B in Blaser- Wang supplement greatly or completely inhibits growth of fungi. C.jejuni colonies appear nonhaemolytic, flat and gray with an irregular edge or raised and round with a mucoid appearance. Some strains may appear tan or slightly pink. Swarming may be observed on moist surfaces. Incubation at 35-37°C may show a delayed growth of *C.jejuni* cultures. Incubating the plates at 42°C can fasten this. The contaminated food sample (10 to 25 grams) is enriched in Campylobacter Enrichment Broth Base (DM1899 + MS2042). The broth is incubated with agitation under a micro aerobic atmosphere for 16-18 hrs. The enrichment culture is then plated onto the selective media i.e. Campylobacter Agar Base (DM1994) (10).

### Type of specimen

Clinical samples - Faeces; Food and dairy samples; Environmental samples.

## **Specimen Collection and Handling:**

For clinical samples follow appropriate techniques for handling specimens as per established guidelines (5,6).
For food and dairy samples, follow appropriate techniques for sample collection and processing as per guidelines (1,4,11).
After use, contaminated materials must be sterilized by autoclaving before discarding.

#### Warning and Precautions:

In Vitro diagnostic Use. Read the label before opening the pack. Wear protective gloves/protective clothing/eye protection/ face protection. Follow good microbiological lab practices while handling specimens and culture. Standard precautions as per established guidelines should be followed while handling clinical specimens. Safety guidelines may be referred in individual safety data sheets.



<sup>#</sup> Equivalent to Liver digest



### **Limitations:**

1. Due to nutritional variations, some strains may show poor growth.

## Performance and Evaluation

Performance of the medium is expected when used as per the direction on the label within the expiry period when stored at recommended temperature

## Methodology

Either streak, inoculate or surface spread the test inoculum (50-100 CFU) aseptically on the plate.

## **Quality Control**

#### Physical Test

#### Appearance

Sterile Campylobacter Agar in 90 mm disposable plates.

На

7.20-7.60

#### Quantity of medium

25 ml of medium in 90 mm disposable plates.

#### Colour of medium

Red coloured medium

#### Sterility Test

Passes release criteria

#### **Cultural Response**

Cultural characteristics observed under reduced oxygen atmosphere after incubation at 35-37°C for 24-48 hours.

Oragnism	Growth
Candida albicans	none - poor
ATCC 10231 (00054*)	
Campylobacter jejuni	good-luxuriant
ATCC 29428 (00156*)	
Escherichia coli	none - poor
ATCC 25922 (00013*)	
Enterococcus faecalis	none - poor
ATCC 29212 (00087*)	

# Storage and Shelf Life

- On receipt store between 20-30°C.
- Use before expiry date on the label.
- Product performance is best if used within stated expiry period.

# Disposal

User must ensure safe disposal by autoclaving and/or incineration of used or unusable preparations of this product.

Follow established laboratory procedures in disposing of infectious materials and material that comes into contact with clinical sample must be decontaminated and disposed of in accordance with current laboratory techniques (5,6).





## **Further Reading**

- 1.American Public Health Association, Standard Methods for the Examination of Dairy Products, 1978, 14th Ed., Washington D.C. 2.Blaser M. J., Cravens B. W., Powers and Wang W. L., 1978, Lanect (ii): 979
- 3. Dekeyser P., Hossuin-Detrain M, Butzler J. P. Sterron J., 1972, J. In fect. Dis., 125: 390
- 4. Downes F. P. and Ito K., (Ed.), 2001, Compendium of Methods for the Microbiological Examination of Foods, 4th Ed.,American Public Health Association, Washington, D.C.
- 5. Isenberg, H.D. Clinical Microbiology Procedures Handbook. 2nd Edition.
- 6. Jorgensen, J.H., Pfaller, M.A., Carroll, K.C., Funke, G., Landry, M.L., Richter, S.S and Warnock., D.W. (2015) Manual of Clinical Microbiology, 11th Edition. Vol. 1.
- 7. Koneman E. W, Allen S. D., Janda W. M, Schreckenberger P. C., Winn W. C. Jr, 1992, Colour Atlas and Textbook of Clinical Microbiology, 4th Edition, J. B. Lippincott Company.
- 8. Manning H., Duim B., Wassenaar T., Wagenaar A., Ridley A., Newell D.G., 2001, Appl. Environ. Microbiol., 67:1185 9.Skirrow M. D., 1977, Br. Med. J. 2:9
- 10. Vanderzant C., and Splittstoesser D. F., (Eds.), 1992, Compendium of Methods for the Microbiological Examination of foods, 3rd Ed., APHA, Washington, D.C.
- 11. Wehr H. M. and Frank J. H., 2004, Standard Methods for the Microbiological Examination of Dairy Products, 17th Ed., APHA Inc., Washington, D.C.
- 12. Wilson and Wang, 1979, Information flier, Campylobacter Laboratory, Veterans Administration Hospital, Denver. Co.

### Disclaimer :

- User must ensure suitability of the product(s) in their application prior to use.
- The product conform solely to the technical information provided in this booklet and to the best of knowledge research and developmentwork carried at **CDH** is true and accurate
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