

## GN BROTH (Hajna) (7218)

### Intended Use

**GN Broth (Hajna)** is used for the selective enrichment of Gram-negative organisms in a laboratory setting. GN Broth (Hajna) is not intended for use in the diagnosis of disease or other conditions in humans.

### Product Summary and Explanation

Hajna formulated Gram Negative (GN) Broth as an enrichment medium for enteric Gram-negative bacilli, especially *Salmonella* spp. and *Shigella* spp.<sup>1,2,3</sup> Croft and Miller demonstrated improved recovery of *Shigella* spp. using GN Broth enrichment compared to direct inoculation of agar.<sup>4</sup> Taylor and Schelhart reported improved recovery of *Salmonella* spp. and *Shigella* spp. when using GN Broth enrichment compared to direct inoculation of media.<sup>5</sup> Taylor and Schelhart found GN Broth to be superior to selenite enrichment medium for recovering *Shigella* spp.<sup>6</sup>

This broth is used as a nonselective enrichment to recover *Salmonella* spp. and *Shigella* spp. from food.<sup>7</sup>

### Principles of the Procedure

Enzymatic Digest of Casein and Enzymatic Digest of Animal Tissue are used as a nitrogen and vitamin source in this medium. Dextrose and Mannitol are the fermentable carbohydrates. The higher concentration of Mannitol over Dextrose favors growth of mannitol-fermenting *Salmonella* spp. and *Shigella* spp. over mannitol non-fermenting species, such as *Proteus*. The Phosphates are buffering agents. Sodium Citrate and Sodium Deoxycholate inhibit growth of Gram-positive bacteria and coliforms. Sodium Chloride maintains the osmotic balance of the medium.

### Formula/Liter

Enzymatic Digest of Casein .....	10 g
Enzymatic Digest of Animal Tissue .....	10 g
Dextrose .....	1 g
Mannitol .....	2 g
Sodium Citrate .....	5 g
Sodium Deoxycholate .....	0.5 g
Dipotassium Phosphate .....	4 g
Monopotassium Phosphate .....	1.5 g
Sodium Chloride .....	5 g

Final pH: 7.0 ± 0.2 at 25°C

Formula may be adjusted and/or supplemented as required to meet performance specifications.

### Precautions

1. For Laboratory Use Only.
2. HARMFUL. May be harmful if absorbed through the skin or ingested. Irritating to eyes, respiratory system, and skin.

### Directions

1. Dissolve 39 g of the medium in one liter of purified water.
2. Mix thoroughly.
3. Autoclave at 121°C for 15 minutes.

### Quality Control Specifications

**Dehydrated Appearance:** Powder is homogeneous, free flowing, and light beige.

**Prepared Appearance:** Prepared medium is gold to light amber, brilliant to clear with none to light precipitate.

**Expected Cultural Response:** Cultural response in GN Broth (Hajna) under aerobic conditions at 35 ± 2°C and examined for growth after 18 – 24 hours incubation.

Microorganism	Approx. Inoculum (CFU)	Expected Growth
<i>Escherichia coli</i> ATCC® 25922	10 - 300	Good to excellent
<i>Enterococcus faecalis</i> ATCC® 29212	300 - 1000	None to poor
<i>Salmonella typhimurium</i> ATCC® 14028	10 - 300	Good to excellent
<i>Shigella sonnei</i> ATCC® 25931	10 - 300	Good to excellent

The organisms listed are the minimum that should be performed for quality control testing.

### Test Procedure

Refer to appropriate references for specific procedures.

### Results

Growth of Gram-negative organisms, especially *Salmonella* spp. and *Shigella* spp. is enhanced.

### Storage

Store dehydrated medium at 2 - 30°C. Once opened and recapped, place container in a low humidity environment at the same storage temperature. Protect from moisture and light by keeping container tightly closed.

### Expiration

Refer to expiration date stamped on the container. The dehydrated medium should be discarded if not free flowing, or if appearance has changed from the original color. Expiry applies to medium in its intact container when stored as directed.

### Limitation of the Procedure

Due to nutritional variation, some strains may be encountered that grow poorly or fail to grow on this medium.

### Packaging

GN Broth (Hajna)	Code No.	7218A	500 g
		7218B	2 kg
		7218C	10 kg

### References

1. MacFaddin, J. F. 1985. Media for isolation-cultivation-identification-maintenance of medical bacteria, vol. 1, p. 357-359. Williams & Wilkins, Baltimore, MD.
2. Hajna, A. A. 1955. A new specimen preservative for gram-negative organisms of the intestinal group. Public Health Lab. **13**:59-62.
3. Hajna, A. A. 1955. A new enrichment broth medium for gram-negative organisms of the intestinal group. Public Health Lab. **13**:83-89.
4. Croft, C. C., and M. J. Miller. 1956. Isolation of *Shigella* from rectal swabs with Hajna "GN" broth. Am. J. Clin. Path. **26**:411-417.
5. Taylor, W. I., and D. Schelhart. 1967. Isolation of shigellae, IV. Comparison of plating media with stools. Am. J. Clin. Path. **48**:356-362.
6. Taylor, W. I., and D. Schelhart. 1968. Isolation of shigellae, V. Comparison of enrichment broths with stools. Appl. Microbiol. **16**:1383-1386.
7. Vanderzant, C. and D.F. Splittstoesser (eds.). 1992. Compendium of methods for the microbiological examination of foods, 3<sup>rd</sup> ed. American Public Health Association, Washington, D.C.

### Technical Information

Contact Acumedia Manufacturers, Inc. for Technical Service or questions involving dehydrated culture media preparation or performance at (517)372-9200 or fax us at (517)372-2006.