BIOMARK Laboratories-INDIA

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TECHNICAL SHEET

| B1183 | GELATIN PHOSPHATE BUFFER | | | | | | | |
|---|--|--------------------------|--------------------------------|-----------------|-------|------------------|-----------------------|--|
| Formula | | | | | | | | |
| Ingredients : gms/lit. | | | | | | | | |
| Sodium dihydrogen phosphate 4.00 | | | | | | | | |
| Gelatin 2.00 | | | | | | | | |
| Final pH (at 25°C) : 6.2 <u>+</u> 0.2 | | | | | | | | |
| Directions : | | | | | | | | |
| Suspend 6.0 grams in 1000 ml distilled water. Boil to dissolve the medium completely. Sterilize by | | | | | | | | |
| autoclaving at 15 lbs pressure (121°C) for 15 minutes. Mix well and dispense as desired. | | | | | | | | |
| Principle : | | | | | | | | |
| Botulinum toxin (botox) types A-G are produced by heterogeneous stains of Clostridium botulinum. Botox | | | | | | | | |
| types A,B,E and F have caused serious and sometimes fatal, cases of food borne illness in humans. The | | | | | | | | |
| vast majority of botulinum outbreaks in red meat and poultry products have involved either toxin A or B. | | | | | | | | |
| The current botulinum toxin test method is the mouse bioassay procedure. Gelatin Phosphate Buffer is one of the reagent used in this test method. | | | | | | | | |
| | | | | | | | | |
| | sts – (I)Dehydrated Medium | | | | | | | |
| | Colour : | | | Cream to yellow | | | | |
| Appearance : | | | Homogeneous coarse powder | | | | | |
| (II)Rehydrated m | 6.2 ± 0.2 | | | | | | | |
| pH (post autoclaving/heating) : Color (post autoclaving/heating) : | | | Colourless | | | | | |
| Clarity(postautoclaving/heating) : | | | clear solution forms in tubes. | | | | | |
| (III)Q.C. Test Microbiological | | | | | | | | |
| Cultural characteristics observed after 24 –48 hrs. at 35-37°C. | | | | | | | | |
| | GROWTH GELATINASE REACTION | | | | | | | |
| MICROORGANISM (ATCC) Clostridium botulinum (25723) | | | - | - | - | | | |
| | Lux | uriant Positive reaction | | | | | | |
| Precautions : 1. For Laboratory Use. | | | | | | | | |
| 2. Follow proper, established laboratory procedures in ha | | | | | | dures in handlir | ng and disposing of | |
| infectious materials. | | | | | | | | |
| Limitations : 1. Since the nutritional requirements of organisms vary, some strains may b | | | | | | | | |
| encountered that fail to grow or grow poorly on this medium. | | | | | | | atridium batulinum is | |
| Use : It isrecommended for toxin detection in food products when Clostridium botul | | | | | | | striaium potulinum Is | |
| Storage : | suspected. Dehydrated medium- below 30°C Prepared medium- Between 2 to 8°C. | | | | | | | |
| Packing : | 500 gm bottle | | | | | | ο C. | |
| | | | | | | Sterilization | | |
| | Preparatio | | | | | Supplement | Stermzation | |
| B1183 | 6.0 g/l | | 333L | 6.2 ± | 0.2 | nil | 121ºC / 15 minutes | |
| 51105 | 0.0 9/1 | 05. | 555L | 0.2 1 | . 0.2 | 1111 | 121 C/ 15 minutes | |

Disclaimer:

User must ensure suitability of the product(s) in their application prior to use. Products conform solely to the information contained in this and other related BIOMARKLABORATORIES publications.

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