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| B060 | WILSON BLAIR AGAR BASE | | | | | |
| Formula | | | | | | |
| Ingredients : | | gms/lit. | | | | |
| Peptone special | | 10.00 | | | | |
| Meat Extract B# | | 5.00 | | | | |
| Dextrose | | 10.00 | | | | |
| Sodium chloride | | 5.00 | | | | |
| Agar | | 30.00 | | | | |
| #- Equivalent to Beef extract | | | | | | |
| Final pH (at 25°C) : | | 7.3 ± 0.2 | | | | |
| Directions : | | | | | | |
| Suspend 60 grams in 1000 ml distilled water. Heat to boiling to dissolve the medium completely. Sterilize by autoclaving at 15 lbs pressure (121°C) for 15 minutes. To sterile melted base, aseptically add 4 ml of 1% brilliant green solution and 70 ml of selective reagent | | | | | | |
| Selective Reagent | | | | | | |
| Solution 1: 40 gm sodium sulphite in 100 ml distilled water. | | | | | | |
| Solution 2: 21 gm dibasic sodium phosphate in 100 ml distilled water. | | | | | | |
| Solution 3: 12.5 gm bismuth ammonium citrate in 100 ml distilled water. | | | | | | |
| Solution 4: 0.96 gm ferrous sulphate in 20 ml distilled water with 2 drops of hydrochloric acid. | | | | | | |
| Prepare each solution separately and then combine. Boil the combined solution until a slate grey colour develops. | | | | | | |
| Principle : | | | | | | |
| Peptone special and Meat Extract B provide nitrogenous, carbonaceous compounds and other growth nutrients. Brilliant green dye inhibits all gram-positive bacteria. Dextrose is the fermentable carbohydrate. Ferrous sulphate aids in H ₂ S production. Bismuth is a heavy metal, which is inhibitory to most gram-negative enteric bacilli other than Salmonella. Ferrous sulphate is reduced by Salmonella species in presence of bismuth sulphite and dextrose to form iron sulphide, indicated by black coloured colonies. Disodium hydrogen phosphate buffers the medium well. Sodium chloride balances the osmotic equilibrium. Do not store the medium in refrigerator (4°C) for longer than 2 days, as the medium changes to green colour and reduces its selectivity. | | | | | | |
| QC Tests – (I) Dehydrated Medium | | | | | | |
| Colour : | | Cream to yellow | | | | |
| Appearance : | | Homogeneous Free Flowing powder | | | | |
| (II) Rehydrated medium | | | | | | |
| pH (post autoclaving/heating) : | | 7.3 ± 0.2 | | | | |
| Colour (post autoclaving/heating) : | | Basal Medium : Light yellow coloured After addition of the selective reagent and 1% Brilliant green, greenish yellow coloured. | | | | |
| Clarity (post autoclaving/heating) : | | Basal Medium: Clear to slightly opalescent After addition of the selective reagent and 1% Brilliant green, opaque gel forms in Petri plates. | | | | |
| (III) Q.C. Test Microbiological | | | | | | |
| Cultural characteristics observed with added 1% Brilliant green and selective reagents after an incubation at 35-37°C for 24-48 hours. | | | | | | |
| MICROORGANISM (ATCC) | | GROWTH | COLOUR OF COLONY | | | |
| Proteus mirabilis (25933) | | Luxuriant | Green | | | |
| Salmonella typhi (6539) | | Luxuriant | Black with sheen | | | |
| Salmonella typhimurium (14028) | | Luxuriant | Black with sheen | | | |
| Escherichia coli (25922) | | Inhibited | -- | | | |
| Precautions : | | 1. For Laboratory Use. 2. Follow proper, established laboratory procedures in handling and disposing of infectious materials. | | | | |
| Limitations : | | 1. Since the nutritional requirements of organisms vary, some strains may be encountered that fail to grow or grow poorly on this medium. | | | | |
| Use : | | For isolation of Salmonella particularly Salmonella typhi after addition of selective reagents. | | | | |
| Storage : | | Dehydrated medium- below 30°C Prepared medium – Use freshly prepared medium. | | | | |
| Packing : | | 500 gm. bottle | | | | |
| Product profile: | | Reconstitution | Quantity on Preparation (500g) | pH (25°C) | Supplement | Sterilization |
| B060 | | 60.0 g/l | 8.33 L | 7.3 ± 0.2 | 1% brilliant green & selective reagents | 121°C for 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 BIOMARK LABORATORIES publications.

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