

853b. FUSIBACTER BIZERTENSIS MEDIUM

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|--|---------|----|
| NH ₄ Cl | 1.00 | g |
| K ₂ HPO ₄ | 0.30 | g |
| KH ₂ PO ₄ | 0.30 | g |
| MgCl ₂ x 6 H ₂ O | 0.50 | g |
| CaCl ₂ x 2 H ₂ O | 0.10 | g |
| KCl | 0.10 | g |
| NaCl | 3.00 | g |
| Yeast extract (OXOID) | 1.00 | g |
| Trypticase peptone (BD BBL) | 1.00 | g |
| Trace element solution (see medium 141) | 10.00 | ml |
| Na-resazurin solution (0.1% w/v) | 0.50 | ml |
| Na ₂ S ₂ O ₃ x 5 H ₂ O | 3.16 | g |
| L-Cysteine-HCl x H ₂ O | 0.50 | g |
| NaHCO ₃ | 4.00 | g |
| D-Glucose | 3.60 | g |
| Na ₂ S x 9 H ₂ O | 0.30 | g |
| Distilled water | 1000.00 | ml |

Dissolve ingredients, except thiosulfate, cysteine, bicarbonate, glucose and sulfide, then sparge medium for 30 - 45 min with 80% N₂ and 20% CO₂ gas mixture to make it anoxic. Add thiosulfate and cysteine, then adjust pH to 7.0 and dispense medium under 80% N₂ and 20% CO₂ gas atmosphere into anoxic Hungate-type tubes or serum vials and autoclave. After autoclaving, add glucose and sulfide from sterile anoxic stock solutions prepared under 100% N₂ gas and bicarbonate from a sterile anoxic stock solution prepared under 80% N₂ and 20% CO₂ gas atmosphere. Adjust pH of complete medium to 7.2 – 7.4, if necessary.