

1490. DESULFOHALOTOMACULUM PECKII MEDIUM

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|--|--------|----|
| NaCl | 20.0 | g |
| NH ₄ Cl | 1.0 | g |
| K ₂ HPO ₄ | 0.3 | g |
| KH ₂ PO ₄ | 0.3 | g |
| KCl | 0.1 | g |
| Na ₂ SO ₄ | 2.0 | g |
| CaCl ₂ x 2 H ₂ O | 0.1 | g |
| Trace elements solution SL-10 (see medium 320) | 1.0 | ml |
| Yeast extract | 2.0 | g |
| Na-resazurin solution (0.1% w/v) | 0.5 | ml |
| NaHCO ₃ | 2.5 | g |
| Na ₂ S ₂ O ₃ x 5 H ₂ O | 3.0 | g |
| MgCl ₂ x 6 H ₂ O | 3.0 | g |
| L-Cysteine-HCl x H ₂ O | 0.5 | g |
| Na ₂ S x 9 H ₂ O | 0.5 | g |
| Distilled water | 1000.0 | ml |

Dissolve ingredients (except bicarbonate, thiosulfate, magnesium chloride, cysteine and sulfide), then sparge medium with 80% H₂ and 20% CO₂ gas mixture for 30 - 45 min to make it anoxic. Dispense medium under same gas atmosphere into anoxic Hungate-type tubes or serum vials to 30% of their volume and autoclave. Add magnesium chloride, cysteine 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 mixture. Adjust pH of the complete medium to 6.8 - 7.0, if necessary. After inoculation pressurize cultivation vials to 1 bar overpressure with sterile 80% H₂ and 20% CO₂ gas mixture.