

520. RUMINOCLOSTRIDIUM CELLULOLYTICUM (CM3) MEDIUM

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|---|---------|----|
| (NH ₄) ₂ SO ₄ | 1.30 | g |
| KH ₂ PO ₄ | 1.50 | g |
| K ₂ HPO ₄ x 3 H ₂ O | 2.90 | g |
| FeSO ₄ x 7 H ₂ O solution (0.1% w/v in 0.1 N H ₂ SO ₄) | 1.25 | ml |
| Trace element solution SL-10 (see medium 320) | 1.00 | ml |
| Yeast extract | 2.00 | g |
| Na-resazurin solution (0.1% w/v) | 0.50 | ml |
| MgCl ₂ x 6 H ₂ O | 0.20 | g |
| CaCl ₂ x 2 H ₂ O | 75.00 | mg |
| Cellobiose | 6.00 | g |
| Na ₂ CO ₃ | 1.50 | g |
| L-Cysteine-HCl x H ₂ O | 0.50 | g |
| Distilled water | 1000.00 | ml |

Dissolve ingredients except magnesium chloride, calcium chloride, cellobiose, cysteine and carbonate, then sparge medium with 80% N₂ and 20% CO₂ gas mixture for 30 – 45 min to make it anoxic. Dispense medium under the same gas atmosphere into anoxic Hungate-type tubes or serum vials and autoclave. After autoclaving add magnesium chloride, calcium chloride and cellobiose from anoxic stock solutions prepared under 100% N₂ gas and carbonate from a sterile anoxic stock solution prepared under 80% N₂ and 20% CO₂. Cellobiose has to be sterilized by filtration. Prior to inoculation add cysteine from a sterile anoxic stock solution prepared under 100% N₂ gas and adjust pH to 7.2.

Note: Some strains can be adapted to cellulose as substrate using 10.00 g/l cellulose powder MN 301 (MACHEREY-NAGEL).

For [DSM 5974](#), [DSM 9801](#) and [DSM 17427](#) use 5.00 g/l D-glucose as the substrate. Supplement medium after autoclaving with 0.50 g/l Na₂S x 9 H₂O added from a sterile anoxic stock solution prepared under 100% N₂ gas and adjust pH of complete medium to 6.8 - 7.0, if necessary.