Microorganisms



1328: DEFLUVIITOGA MEDIUM

This recipe contains strain-specific modifications for Mesotoga infera DSM 25546 *

Final pH: 7.0

Final volume: 1001 ml

	KH ₂ PO ₄	0.30	g	
	K ₂ HPO ₄	0.30	g	
	NH ₄ Cl	1.00	g	
	NaCl	1.00	g	
	KCI	0.10	g	
	$MgCl_2 \times 6 H_2O$	0.50	g	
	CaCl ₂ x 2 H ₂ O	0.10	g	
	Trace element solution SL-10	1.00	ml	
	Yeast extract (OXOID)	2.00	g	
	Sodium resazurin (0.1% w/v)	0.50	ml	
	Sulfur (powdered)	10.00	g	
	L-Cysteine HCl x H ₂ O	0.50	g	
_	Trypticase peptone (BD BBL)	2.00	g	
_	Na ₂ -fumarate	3.20	g	
	Na ₂ CO ₃	1.00	g	
	$Na_2S \times 9 H_2O$	0.50	g	
	Na acetate	0.20	g	
	D-Xylose	3.00	g	
	Distilled water	1000.00	ml	

Dissolve ingredients (except sulfur, cysteine, peptone, fumarate, carbonate and sulfide) and sparge medium with $80\%~N_2$ and $20\%~CO_2$ gas mixture for 30 - 45 min to make it anoxic. Add and dissove cysteine, then dispense under $80\%~N_2$ and $20\%~CO_2$ gas atmosphere into anoxic Hungate-type tubes or serum vials containing already the appropriate amount of sulfur and autoclave at $121^{\circ}C$ for 20 min. Add peptone, fumarate and sulfide from sterile anoxic stock solutions prepared under $100\%~N_2$ gas and carbonate from a sterile anoxic stock solution prepared under $80\%~N_2$ and $20\%~CO_2$ gas mixture. Adjust pH of the complete medium to 7.0.

* Omit Trypticase peptone and fumarate. Increase amount of yeast extract to 2.0 g/l, supplement medium with 0.2 g/l Na-acetate and add 3.0 g/l D-xylose after autoclaving from anoxic stock solutions prepared under 100% N_2 gas and sterilized by filtration.

Trace element solution SL-10 (from medium 320)

HCI (25%)	10.00	ml
FeCl ₂ x 4 H ₂ O	1.50	g
ZnCl ₂	70.00	mg

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$MnCl_2 \times 4 H_2O$	100.00	mg
H_3BO_3	6.00	mg
CoCl ₂ x 6 H ₂ O	190.00	mg
CuCl ₂ x 2 H ₂ O	2.00	mg
NiCl ₂ x 6 H ₂ O	24.00	mg
$Na_2MoO_4 \times 2 H_2O$	36.00	mg
Distilled water	990.00	ml

First dissolve FeCl_2 in the HCl, then dilute in water, add and dissolve the other salts. Finally make up to 1000.00 ml.