

380. MAGNETOSPIRILLUM MEDIUM

KH ₂ PO ₄	0.68	g
NaNO ₃	0.12	g
L(+)-Tartaric acid	0.37	g
Succinic acid	0.37	g
Na-acetate	0.05	g
Vitamin solution (see medium 141)	10.00	ml
Trace element solution (see medium 141)	5.00	ml
Fe(III) quinate solution (see below)	2.00	ml
Agar (BD Bacto, for semi-solid medium)	1.30	g
Na-resazurin solution (0.1% w/v)	0.50	ml
Na-thioglycolate	0.05	g
Distilled water	1000.00	ml

Dissolve ingredients (except thioglycolate) in the order given and adjust pH to 6.75 with NaOH.

Preparation of liquid medium: Sparge medium with 100% N₂ gas for 30 -45 min and dispense under the same gas atmosphere into anoxic Hungate-type tubes to 50% of their volume. Seal vials with screw caps and gas tight butyl rubber closures. Autoclave at 121°C for 15 min. Before inoculation add thioglycolate from a 0.5% (w/v) stock solution, freshly prepared under 100% N₂ gas and filter-sterilized. Then add sterile air (with hypodermic syringe through the rubber closure) to a concentration of ca. 1% (v/v) O₂ in the vial (e.g., add 1 ml air to a Hungate-type tube of 16 ml total volume).

Preparation of semi solid medium: Supplement medium with agar, bring medium to the boil and cool under 100% N₂ gas atmosphere. Dispense under same gas atmosphere aliquots of 10 ml semi-solid medium into Hungate-type tubes. Prior to inoculation add thioglycolate from a 0.5% (w/v) stock solution, freshly prepared under 100% N₂ gas and filter-sterilized. Then add sterile air (with hypodermic syringe through the rubber closure) to a concentration of ca. 1% (v/v) in the vial.

*Note: Prior to inoculation media should be slightly pink in color. Strongly reduced conditions will not support growth of microaerophilic Magnetospirillum species. Use as inoculum 10% (v/v). Incubate tubes with medium without agitation in an inclined position. During growth O₂ will be consumed and the pH will increase. If higher densities of magnetic cells are wanted, feed oxygen (by adding air), succinic acid and ferric quinate from sterile stock solutions (maintain pH below 7). **For cultivation of magnetic cells we recommend preparation of liquid medium, while semi-solid medium is more suitable for demonstration of microaerophilic band formation and storage.***

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Ferric Quinate Solution, 0.01 M :

FeCl ₃ x 6 H ₂ O	4.50	g
Quinic acid	1.90	g
Distilled water	1000.00	ml

Sterilize by filtration under 100% N₂ gas atmosphere.

For DSM 6361 increase the amount of added O₂ to a concentration of 5% (v/v) in the vial (e.g., add 4 ml sterile air to a Hungate-type tube of 16 ml total volume).