

## 1597a. METHANONATRONARCHAEUM MEDIUM

### Solution A:

Na <sub>2</sub> CO <sub>3</sub>	92.50	g
NaHCO <sub>3</sub>	17.50	g
NaCl	8.00	g
K <sub>2</sub> HPO <sub>4</sub>	0.50	g
Distilled water	500.00	ml

### Solution B:

NaCl	120.00	g
K <sub>2</sub> HPO <sub>4</sub>	1.25	g
KCl	2.50	g
Distilled water	500.00	ml

### Solution C:

Trace elements solution (see medium 1369)	1.00	ml
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### Solution D:

Selenite-tungstate solution (see medium 385)	1.00	ml
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### Solution E:

NH <sub>4</sub> Cl	0.20	g
MgSO <sub>4</sub> x 7 H <sub>2</sub> O	0.25	g
Distilled water	5.00	ml

### Solution F:

Methanol	2.00	ml
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### Solution G:

Na-formate	3.40	g
Na-acetate	0.16	g
Distilled water	10.00	ml

### Solution H:

Yeast extract	0.02	g
Distilled water	5.00	ml

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**Solution I:**

2-Mercaptoethanesulfonic acid (coenzyme M)	0.15	g
Distilled water	10.00	ml

**Solution J:**

Vitamin solution (see medium 141)	10.00	ml
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**Solution K:**

Ferrous sulfide sludge (see medium 1267 <i>solution C</i> )	5.00	ml
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**Solution L:**

Na <sub>2</sub> S x 9 H <sub>2</sub> O	0.25	g
Distilled water	10.00	ml

Sterilize *solutions A* and *B* in closed thick-walled screw-top bottles (e.g., SCHOTT) for 20 min at 120°C. The pH of *solution A* after sterilization should be 10. There is some precipitate forming that settles at the bottom after 3 – 4 days. It is best to remove precipitates by decantation before using *solution A* for medium preparation.

Combine *solution A* with *solution B* and sparge medium with 100% N<sub>2</sub> gas for at least 30 – 45 min to make it anoxic, then dispense under same gas atmosphere into anoxic Hungate-type tubes or serum vials and autoclave. *Solutions C* to *L* are sterilized separately under 100% N<sub>2</sub> gas. Vitamins and coenzyme M should be sterilized by filtration. To complete the medium appropriate amounts of *solutions C* to *L* are added to the combined sterile solutions *A* and *B* in the sequence as indicated. Final pH of the medium should be 9.5.

*Note: Addition of 10 - 20 mg sodium dithionite per liter (e.g. from 5% (w/v) solution, freshly prepared under N<sub>2</sub> and filter-sterilized) may stimulate growth at the beginning. For transfers use 5 - 10% inoculum.*