## ATCC medium 2487: Broth: MG Medium

MS-OCM Base Medium (see Appendix I) with 2.5 g of NaCl per liter and 5 mM sodium acetate. Add formate to a final concentration of 50 mM.

## MS-OCM Base Medium

Many methanogens at the ATCC are grown in either MS medium or its derivatives that contains high salt concentrations or an alkaline pH. The medium uses a bicarbonate buffer. We recommend having the following stock solutions on hand.

**Solution A:** (100X solution; use 10 ml of solution A per liter of medium)

- 100.0 g Ammonium chloride
- 100.0 g Magnesium chloride hexahydrate
- 40.00 g Calcium chloride dihydrate

Dissolve in deionized water, adjust pH to 4.0 with HCl and bring up to one liter.

**Solution B:** (500X solution; use 2 ml of solution B per liter of medium)

• 200.0 g Potassium dibasic phosphate trihydrate

Dissolve in deionized water then bring the volume up to one liter.

**Solution C:** (500X; use 2 ml of solution C per liter of medium)

• 0.5 g Resazurin

Dissolve in deionized water, then bring the volume up to one liter.

**Trace Mineral Supplement**, Catalog No. MD-TMS: (Available from ATCC as a sterile ready-to-use liquid.)

• 10.0 ml per liter of medium

To prepare the medium dissolve 8.4 gm sodium bicarbonate in 976.0 ml of deionized water (100 mM  $Na_2HCO_3$ ). The solution is bubbled with an oxygen-free gas mixture of  $N_2$ - $CO_2$  (70:30). While gassing, the stock solutions A and B are added. The high initial pH of this solution may cause the minerals to precipitate. As the pH decreases as a result of equilibration of the  $CO_2$ :bicarbonate buffering system the minerals will go back into solution; at the same time  $O_2$  is driven out of the medium. Dry ingredients are also added and dissolved in the order listed.

- 2.0 g Yeast extract
- 2.0 g Trypticase peptones

The medium is allowed to gas until it has become clear, usually 30 to 60 minutes depending on the gas flow rate. The pH of the cleared medium should be between pH 7 and pH 7.3. At this point the solution C and trace minerals supplement can be added to the medium.

The medium is then ready to be transferred to culture tubes. For this purpose, a gas-cannula system with at least three separate gassing ports should be used. A rack of Balch tubes should be made ready. The flask of medium is continuously gassed with one gas cannula; the other two cannula are placed, one each, in two sequential Balch tubes. A 10 ml pipette is used to pipette 10 ml of medium from the flask to the 1<sup>st</sup> Balch tube. The cannula is then removed from this tube and at the same time a butyl rubber stopper is pushed into the top; it is important to do this all in one motion to minimize contamination of the headspace of the tube with air. This cannula is then moved to the 3<sup>rd</sup> tube in sequence and the 2<sup>nd</sup> tube is filled with media and capped, and so on. After the all tubes are filled, capped, and crimped, the medium can be autoclaved. Following autoclaving a precipitate may form; with periodic mixing this should go back into solution within 48 hours.

Note: In our experience, the resazurin is often colorless after autoclaving MS medium, probably because the yeast extract provides enough reducing power to cause complete reduction of the dye. We still recommend adding a reducing agent to the medium.