



Product Sheet

Desulfitobacterium metallireducens (ATCC®) BAA-636™

Please read this FIRST



Intended Use

This product is intended for research use only. It is not intended for any animal or human therapeutic or diagnostic use.

Citation of Strain

If use of this culture results in a scientific publication, it should be cited in that manuscript in the following manner: *Desulfitobacterium metallireducens* (ATCC® BAA-636™)

Description

Designation: 853-15A

Deposited Name: *Desulfitobacterium metallireducens* Finneran et al.

Propagation

Medium

Modified 2260 for BAA-636

Growth Conditions

Temperature: 30.0°C

Propagation Procedure

2. If needed exchange the gas in the test tube for 80% N₂ 20% CO₂.
3. Add 0.2 ml Cysteine 3.0% stock solution) per 10 ml of medium. Let the medium sit at room temperature for 30 to 40 minutes before inoculating.
4. When the Balch tube is ready to inoculate, thaw the frozen vial at room temperature under a gentle stream of oxygen-free gas.
5. For inoculation, use an anaerobic (see c below) 1.0 ml syringe tipped with 22-gauge needle, withdraw the cell suspension from the vial and transfer it to the broth. Plate 0.1 ml of the inoculated culture onto a non-selective medium and incubate aerobically at 30°C. Use 0.5 ml of the inoculated culture to inoculate a nonselective aerobic broth and an additional tube of #2550 broth. Incubate the non-selective aerobic broth tubes at 30°C. Incubate the anaerobic tube at 30°C.

ANAEROBIC CONDITIONS:

- a. Resazurin is a commonly used redox indicator that is pink when the redox potential is above 50 mv., and colorless when the redox potential is below 110 mv. i.e. highly reducing. Most strict anaerobes require this low redox potential for optimum growth.
- b. To obtain a fully reduced medium, it is necessary that the medium be anoxic and that a reducing agent be added. Common reducing agents are sodium sulfide, cysteine, dithiothreitol, and titanium citrate.
- c. Syringes can be made anaerobic by one of two methods. 1. Displace the dead space in the syringe with a sterile
2. Displace the dead space in the syringe with a reducing agent.

Notes

When examined microscopically, the cells appear as slightly curved rods. Always use freshly prepared anaerobic medium. If there is any question about the anaerobic condition of the medium, the medium can be reduced with the addition of 1.5% cysteine (2.0 ml per 100 ml of medium). Cysteine is the reducing agent of choice since it does not cause the ferrous ammonium sulfate to precipitate.

References

References and other information relating to this product are available online at www.atcc.org.

Biosafety Level: 1

Appropriate safety procedures should always be used with this material. Laboratory safety is discussed in the current publication of the *Biosafety in Microbiological and Biomedical Laboratories* from the U.S. Department of Health and Human Services Centers for Disease Control and Prevention and National Institutes for Health.

ATCC Warranty

The viability of ATCC® products is warranted for 30 days from the date of shipment, and is valid only if the product is stored and cultured according to the information included on this product information sheet. ATCC lists the media formulation that has been found to be effective for this strain. While other, unspecified media may also produce satisfactory results, a change in media or the absence of an additive from the ATCC recommended media may affect recovery, growth and/or function of this strain. If an alternative medium formulation is used, the ATCC warranty for viability is no longer valid.

Disclaimers

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Please see the enclosed Material Transfer Agreement (MTA) for further details regarding the use of this product. The MTA is also available on our Web site at www.atcc.org

Additional information on this culture is available on the ATCC web site at www.atcc.org.

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