



Product Sheet

Acetobacterium fimetarium (ATCC® 51795™)

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: *Acetobacterium fimetarium* (ATCC® 51795™)

Description

Designation: Z-4290 [DSM 8238]

Deposited Name: *Acetobacterium fimetarium* Kotsyurbenko et al.

Propagation

Medium

ATCC® Medium 1019: Acetobacterium medium

Growth Conditions

Temperature: 20.0°C

Atmosphere: Under a gas mixture of 80% N₂, 20% H₂

Propagation Procedure

1. Open vial according to enclosed instructions.
2. Under anaerobic conditions, withdraw 0.5 ml of the recommended broth from a single tube (5 to 6 ml) and rehydrate the vial contents.
4. Incubate tubes under an anaerobic atmosphere at 20°C. Incubate one agar plate anaerobically for colony formation, and one aerobically for aerobic contamination check.
5. In 48 hours, growth should be evident by turbidity that settles to the bottom of the tube. No growth should occur on agar plate incubated aerobically.

ANAEROBIC CONDITIONS:

- a. **Balch tubes (available from Bellco Glass, Vineland, NJ; are specially designed for anaerobic work and use an aluminum crimp cap to hold a rubber stopper in place. Needles can easily be inserted through the stopper, and the tubes can be pressurized to 2 atm. Alternatively, serum vials may be used, or screw cap tubes with butyl rubber stoppers, in the latter case the stopper may be removed and the tube placed under a cannula system that dispenses sterile, oxygen free gas for addition of reducing agents or inoculation.**
- b. **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.**
- c. **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.**
- d. **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**

Notes

Additional information on this culture is available on the ATCC web site at www.atcc.org. While every effort is made to insure authenticity and reliability of strains on deposit, ATCC is not liable for damages arising from the misidentification or misrepresentation of cultures.

ATCC recommends that individuals contemplating commercial use of any culture first contact the originating investigator to negotiate an agreement. Third party distribution of this culture is discouraged, since this practice has resulted in the unintentional spreading of contaminated cultures.

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

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effective for this product. 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 product. If an alternative medium formulation is used, the ATCC warranty for viability is no longer valid.

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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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