**Product Sheet** 

# Butyrivibrio fibrisolvens Bryant and Small 19171<sup>™</sup>

#### Description

Type strain **Strain designation:** [ATCC 12560] **Deposited As:** *Butyrivibrio fibrisolvens* Bryant and Small **Type strain:** Yes

# **Storage Conditions**

Product format: Freeze-dried Storage conditions: 2°C to 8°C

# Intended Use

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

#### **BSL1**

ATCC determines the biosafety level of a material based on our risk assessment as guided by the current edition of *Biosafety in Microbiological and Biomedical Laboratories (BMBL)*, U.S. Department of Health and Human Services. It is your responsibility to understand the hazards associated with the material per your organization's policies and procedures as well as any other applicable regulations as enforced by your local or national agencies.



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ATCC highly recommends that appropriate personal protective equipment is always used when handling vials. For cultures that require storage in liquid nitrogen, it is important to note that some vials may leak when submersed in liquid nitrogen and will slowly fill with liquid nitrogen. Upon thawing, the conversion of the liquid nitrogen back to its gas phase may result in the vial exploding or blowing off its cap with dangerous force creating flying debris. Unless necessary, ATCC recommends that these cultures be stored in the vapor phase of liquid nitrogen rather than submersed in liquid nitrogen.

# **Certificate of Analysis**

For batch-specific test results, refer to the applicable certificate of analysis that can be found at www.atcc.org.

# **Growth Conditions**

Medium: ATCC Medium 602: E medium for Anaerobes ATCC Medium 260: Trypticase soy agar/broth with defibrinated sheep blood Temperature: 37°C Atmosphere: Anaerobic

# Handling Procedures

- 1. Open vial according to enclosed instructions or visit www.atcc.org for instructions.
- Under anaerobic conditions aseptically rehydrate the entire pellet with approximately 0.5 mL of #602 broth. Aseptically transfer the entire contents to a 5-6 mL tube of #602 broth. Additional test tubes can be inoculated by



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transferring 0.5 mL of the primary broth tube to these secondary broth tubes. Best practice dictates the use of pre-reduced media.

- 3. Use several drops of the primary broth tube to inoculate a #260 plate and/or #260 agar slant.
- 4. Incubate in an anaerobic atmosphere at 37° C for 2-3 days. Incubate one agar plate aerobically at 37°C to check for contamination. Once growth has been established, the culture should be transferred to fresh broth every 24 to 48 hours.
- 5. This culture is very sensitive to oxygen, therefore steps should be taken to avoid exposure to oxygen. When the culture exhibits good growth it will remain viable for up to 1 week if stored at 4°C under anaerobic condition.

#### ANAEROBIC CONDITIONS:

Anaerobic conditions for transfer may be obtained by the use of an anaerobic gas chamber or placement of test tubes under a gassing cannula system connected to anaerobic gas.

Anaerobic conditions for incubation may be obtained by any of the following:

- Loose screw caps on test tubes in an anaerobic chamber
- Loose screw caps on test tubes in an activated anaerobic gas pack jar
- Use of sterile butyl rubber stoppers on test tubes so that an anaerobic gas headspace is retained

#### Notes

Growth should be detected within 24 to 72 hours indicated by turbidity throughout the broth.

Once growth has been establish the culture should be transferred every 24 hours when maintained at 37°C. The culture can be maintained at 4°C for up to 1 week.

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

#### **Material Citation**

If use of this material results in a scientific publication, please cite the material in the following manner: *Butyrivibrio fibrisolvens* Bryant and Small (ATCC 19171)

#### References

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

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#### Revision

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# **Contact Information**

ATCC 10801 University Boulevard Manassas, VA 20110-2209 USA US telephone: 800-638-6597 Worldwide telephone: +1-703-365-2700 Email: tech@atcc.org or contact your local distributor

