



# ***Enterococcus mundtii*** **Collins et al.**

**882™**

## **Description**

**Strain designation:** A

**Deposited As:** *Streptococcus faecalis*. Formerly known as *Enterococcus faecium*

**Type strain:** No

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## **Storage Conditions**

**Product format:** Freeze-dried

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## **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.

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## **BSL 1**

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.

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.

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## Certificate of Analysis

For batch-specific test results, refer to the applicable certificate of analysis that can be found at [www.atcc.org](http://www.atcc.org).

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## Growth Conditions

### Medium:

ATCC Medium 260: Trypticase soy agar/broth with defibrinated sheep blood

**Temperature:** 37°C

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## Handling Procedures

1. Open vial according to enclosed instructions.
  2. Using a single tube of #44 broth (5 to 6 ml), withdraw approximately 0.5 to 1.0 ml with a Pasteur or 1.0 ml pipette. Rehydrate the entire pellet.
  3. Aseptically transfer this aliquot back into the broth tube. Mix well.
  4. Use several drops of the suspension to inoculate a #260 agar slant and/or plate.
  5. Incubate the tubes and plate at 37°C for 24 hours.
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## Notes

Colonies on #260 plates are entire, glistening, circular, and smooth. Yellow pigment is not produced on #44 or #260 agar.

This culture was re-identified as *Enterococcus mundtii* based on 16s sequence data and biochemical reactions. This strain utilizes xylose, sorbitol, rhamnose, melibiose, and does not grow at 50°C which supports the *E. mundtii* identification. Variable results have been reported for raffinose utilization.

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## Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: *Enterococcus mundtii* Collins et al. (ATCC 882)

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## References

References and other information relating to this material are available at [www.atcc.org](http://www.atcc.org).

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