



"*Borrelia andersonii*" Marconi et al.

700555™

Description

"*Candidatus Borrelia andersonii*" strain 21038 is a bacterium that was isolated from tick larvae in Millbrook, New York. This strain is propagated under microaerophilic conditions.

Strain designation: 21038

Deposited As: "*Borrelia andersonii*" Marconi et al.

Type strain: Yes

Storage Conditions

Product format: Frozen

Storage conditions: -80°C or colder

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.

BSL 2

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

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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.

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 1914: Revised BSK medium

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

Temperature: 35°C**Atmosphere:** Microaerophilic

Handling Procedures

1. Open thawed vial.
2. Aseptically transfer the entire contents to a 5-6 mL tube of #1914 broth.
Additional test tubes can be inoculated by transferring 0.5 mL of the primary

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broth tube to these secondary broth tubes.

3. Use several drops of the primary broth tube to inoculate a #260 plate to check for purity.
4. Or, to obtain a biphasic culture, add several drops of the primary broth tube to an agar slant. Best practice is to incubate these slants at an angle.
5. Incubate at 35°C under microaerophilic conditions for 4-7 days. Use an anaerobe jar with an active catalyst and a microaerophilic gas generator pack or other acceptable method. All tubes and slants should be incubated with caps loosened.

Notes

A jump may be necessary to achieve sufficient growth. Does not grow on agar plates but it may grow with a biphasic slant.

Acid formation during growth will change the medium to a light or yellowish-orange color. Turbidity is not evident. Cells can be monitored under phase microscopy as long spiral rods, and their motility is observed by the twitching movement.

Borrelia species are fragile, sensitive organisms that must have the appropriate medium for growth. Rabbit serum is essential for the growth of this organism. Fresh medium enhances growth. Medium older than one month should not be used.

Referred to as the type strain of *Borrelia andersonii*, but this taxonomic name is not validly published at this time.

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

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: "*Borrelia andersonii*" Marconi et al. (ATCC 700555)

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References

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

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Contact Information

ATCC

10801 University Boulevard

***"Borrelia andersonii"* Marconi et al.**

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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
