Product Sheet

Haemophilus influenzae (Lehmann and Neumann) Winslow et al.

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Description

Haemophilus influenzae strain AMC 36-A-1 is a whole-genome sequenced bacterium that is typed as biotype I, serotype b. This product has applications in respiratory disease research and can be used as a quality control strain in media testing and susceptibility disc testing.
Strain designation: AMC 36-A-1 [572]
Deposited As: Haemophilus influenzae (Lehmann and Neumann) Winslow et al.
Type strain: No
Antigenic properties: Biotype I

Serotype: b

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.

BSL 2



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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.

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 2167: Haemophilus Test Medium ATCC Medium 814: GC Agar/Broth Medium Temperature: 37°C Atmosphere: 95% Air, 5% CO₂



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Handling Procedures

- 1. Open vial according to enclosed instructions or visit www.atcc.org for instructions.
- Rehydrate the entire pellet with approximately 0.5 mL of #2167 broth.
 Aseptically transfer the entire contents to a 5-6 mL tube of #2167 broth.
 Additional test tubes can be inoculated by transferring 0.5 mL of the primary broth tube to these secondary broth tubes.
- 3. Use several drops of the primary broth tube to inoculate a #814 plate and/or #814 agar slant.
- 4. Incubate at 37°C for 24-48 hours in an atmosphere of 5% CO₂.

Notes

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

Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: *Haemophilus influenzae* (Lehmann and Neumann) Winslow et al. (ATCC 10211)

References

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

Warranty

The product is provided 'AS IS' and the viability of $ATCC^{\circ}$ products is warranted for 30 days from the date of shipment, provided that the customer has stored and handled

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the product according to the information included on the product information sheet, website, and Certificate of Analysis. For living cultures, ATCC lists the media formulation and reagents that have been found to be effective for the product. While other unspecified media and reagents may also produce satisfactory results, a change in the ATCC and/or depositor-recommended protocols may affect the recovery, growth, and/or function of the product. If an alternative medium formulation or reagent is used, the ATCC warranty for viability is no longer valid. Except as expressly set forth herein, no other warranties of any kind are provided, express or implied, including, but not limited to, any implied warranties of merchantability, fitness for a particular purpose, manufacture according to cGMP standards, typicality, safety, accuracy, and/or noninfringement.

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Revision

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