**Chlamydia trachomatis Serovar E**

VR-348B™

**Description**

*Strain designation:* BOUR  
*Common name:* *Chlamydia trachomatis* Serovar E  
*Deposited As:* *Chlamydia trachomatis* (Busacca) Rake  
*Type strain:* No

**Storage Conditions**

*Product format:* Frozen  
*Storage conditions:* -70°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 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

**Host:** McCoy [McCoy B] (ATCC CRL-1696)

**Effects:** CPE; cytoplasmic inclusions

**Complete medium:**
DMEM (ATCC® 30-2002™) + 10% prescreened FBS + 10 mM HEPES + 2 µg/mL Cycloheximide (Sigma C-4859 Ready-Made)

**Temperature:** 35.5°C

**Recommendations for infection:** Plate cells 24-48 hours prior to infection and infect when cultures are 90-100% confluent. Prior to infection, disrupt cells by sonication or vortex with sterile glass beads. Remove medium and inoculate with a small volume of inoculum (e.g. 1 mL per 25 cm²) diluted to provide an optimal MOI (e.g. 1). Adsorb by centrifugation at 2100 x g for 1 hour at 20°C. End adsorption by adding growth medium.
**Incubation**: 3 days

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**Handling Procedures**

**Mycoplasma contamination**: Not detected

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**Notes**

FBS used to culture *Chlamydia* must be prescreened to verify that the serum does not contain antibodies to *Chlamydia* or other factors that would interfere with growth.

Rapid loss in titer when stored above -70°C. Activities with high potential for aerosol production require BSL 3 facilities and practices.

Next-generation sequencing (NGS) at ATCC on the McCoy cell line (ATCC® CRL-1696™) used as the host has shown the presence of *Mus Musculus* mobilized endogenous polytropic provirus and Murine leukemia virus.

**Key Abbreviations**: °C, Degrees Celsius; BSL, Biosafety level; CO₂, Carbon dioxide; DMEM, Dulbecco’s Modified Eagle’s Medium; FA, Fluorescent antibody assay; FBS, Fetal bovine serum; g, Acceleration of gravity; HEPES, N-(2-Hydroxyethyl)piperazine-N’-(2-ethanesulfonic acid); MOI, Multiplicity of infection

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**Material Citation**

If use of this material results in a scientific publication, please cite the material in the following manner: *Chlamydia trachomatis* Serovar E (ATCC VR-348B)

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

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Revision

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