



## Product Sheet

# Quantitative Synthetic *Treponema pallidum* DNA (ATCC® BAA-2642SD™)

Please read this **FIRST**

Storage Temp.  
**-20°C or colder**

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Biosafety Level  
**1**

### Intended Use

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

This product is intended for research use only. It is not intended for any animal or human therapeutic or diagnostic use. The synthetically engineered sequence of the product constitutes intellectual property belonging to ATCC. Unauthorized use, including sequencing, modification, or reverse-engineering, of the product is expressly prohibited without prior ATCC consent.



### Citation of Strain

If use of this culture results in a scientific publication, it should be cited in that manuscript in the following manner: Quantitative Synthetic *Treponema pallidum* DNA (ATCC® BAA-2642SD™)

### Nucleic Acid Information

Specification range:  $1 \times 10^5$  to  $1 \times 10^6$  copies/ $\mu$ L  
100  $\mu$ L per vial with Biomatrix DNASTable

American Type Culture Collection  
PO Box 1549  
Manassas, VA 20108 USA  
[www.atcc.org](http://www.atcc.org)

800.638.6597 or 703.365.2700  
Fax: 703.365.2750  
Email: [Tech@atcc.org](mailto:Tech@atcc.org)

Or contact your local distributor

### Description

**Source:** Quantitative Synthetic *Treponema pallidum* DNA

**Description:** ATCC® Genuine Nucleics can be used for assay development, verification, validation, monitoring of day to day test variation and lot to lot performance of molecular-based assays. The quantitative format allows for the generation of a standard curve for quantitative PCR (qPCR) to determine bacterial load. Preparation includes fragments from the *poA*, *tpr*, 23S gene, *arp*, 16S gene, *flaA*, 47kDa protein gene, and *bmp*.

**Note:** Aliquotting is highly recommended to avoid multiple freeze-thaws.

### Batch-Specific Information

Refer to the Certificate of Analysis for batch-specific test results.

### Preparation Procedure

1. Thaw the vial at room temperature and immediately place on ice. Avoid exposing the synthetic DNA to repeated freeze-thaw cycles as it may result in degradation of the DNA and variation in copy number.
2. Gently mix the sample to ensure an even distribution of material.
3. Briefly centrifuge the tube before opening to ensure all liquid is at the bottom.

### Biosafety Level: 1

Appropriate safety procedures should always be used with this material. Laboratory safety is discussed in the current publication of the *Biosafety in Microbiological and Biomedical Laboratories* from the U.S. Department of Health and Human Services Centers for Disease Control and Prevention and National Institutes for Health.

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Please see the enclosed Material Transfer Agreement (MTA) for further details regarding the use of this product. The MTA is also available on our Web site at [www.atcc.org](http://www.atcc.org)

Additional information on this culture is available on the ATCC web site at [www.atcc.org](http://www.atcc.org).

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