



# *Treponema socranskii* subsp. *paredis* Smibert et al.

35535™

## Description

**Strain designation:** VPI D46CPE1

**Deposited As:** *Treponema socranskii* subsp. *paredis* Smibert et al.

**Type strain:** Yes

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## Storage Conditions

**Product format:** Frozen

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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 submerged 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 submerged 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 1494: Modified NOS medium

**Temperature:** 37°C

**Atmosphere:** Anaerobic

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## Handling Procedures

**1. Thaw the frozen vial at room temperature.**

2. Under anaerobic conditions, transfer the entire contents of the thawed vial into a single tube (5-6 ml) of #1494 broth. Mix well.

3. Additional tubes of broth may be inoculated with 0.5 ml each of the suspension. You may also use 0.5 ml to inoculate a #1494 slant. Inoculate two Trypticase Soy

Agar plates with 5% sheep blood (ATCC® Medium #260). Incubate one blood plate anaerobically to check for colony formation and incubate the second plate aerobically to check for purity.

4. Incubate tubes and plate under anaerobic conditions at 37°C. Incubate the second blood plate aerobically.

5. Growth occurs within 5 to 10 days, and should be evident by very slight turbidity in the broth and helical cells under phase contrast. The aerobic plate should show no signs of growth.

#### ANAEROBIC CONDITIONS:

Anaerobic conditions for transfer may be obtained by either of the following:

- Use of an anaerobic gas chamber, or
- Placement of test tubes under a gassing cannula system hooked to anaerobic gas.

Anaerobic conditions for incubation may be obtained by any of the following:

- Loose screw caps on test tubes in anaerobic chamber,
- Loose screw caps on test tubes in an activated anaerobic gas pack jar, or
- Use of sterile butyl rubber stoppers on test tubes so that an anaerobic gas headspace is retained.

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## Notes

Cells appear as motile, long, spiraled rods.

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

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## Material Citation

If use of this material results in a scientific publication, please cite the material in the

following manner: *Treponema socranskii* subsp. *paredis* Smibert et al. (ATCC 35535)

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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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## Revision

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## Contact Information

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35535

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