



# ***Phaeospirillum molischianum* (Giesberger) Imhoff et al.**

**14031™**

## **Description**

**Strain designation:** NCIB 9957

**Deposited As:** *Rhodospirillum molischianum* Giesberger

**Type strain:** Yes

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## **Storage Conditions**

**Product format:** Freeze-dried

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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**

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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.

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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 1808: Yeast extract (ATCC medium 837) with 0.015% sodium sulfide

**Temperature:** 24-26°C

**Atmosphere:** Anaerobic

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## Handling Procedures

1. Open vial according to enclosed instructions.
2. This organism can tolerate brief exposure to oxygen, so it may be opened without anaerobic conditions. Aseptically transfer 0.5 ml of medium #1808 to the vial and rehydrate the pellet. Transfer this suspension to a single tube (5 to 6 ml) of

#1808 broth that has been pre-reduced by the addition of 0.015% sodium sulfide. To check for possible aerobic contamination, plate 0.2 ml of the culture on any non-selective media and incubate aerobically in the dark at 26°C. (There may be some slight growth). Fill the test tube to capacity, seal the tube with a screw cap, and incubate at 26°C (room temperature) under a tungsten lamp.

3. After four to seven days, growth should be evident as indicated by turbidity and red pigmentation through out the broth. When examined microscopically, the cells appear as spiral shaped rods, in singles and pairs that are motile. Once growth has been detected, the culture should be transferred to fresh broth. Subsequent growth should be detected within 48 to 72 hours.

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## Notes

When making additional transfers, it is not necessary to pre-reduce the medium if using a large inoculum (20% or greater).

Additional information on this culture is available 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: *Phaeospirillum molischianum* (Giesberger) Imhoff et al. (ATCC 14031)

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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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