

MYA-2950D-5<sup>™</sup>

## Description

Genomic DNA isolated from Nakaseomyces glabratus. This fungal strain is also available as ATCC MYA-2950.

**Organism:** Nakaseomyces glabratus (Anderson) Sugita et Takash **Derived from:** *Nakaseomyces glabratus* 303542 (ATCC MYA-2950)

Genome sequenced strain: Yes

Type strain: No Mass: 5 μg

Shipping information: Stored in 1X TE buffer

## **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<sub>1</sub>

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



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(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.

### Certificate of Analysis

For batch-specific test results, refer to the applicable certificate of analysis that can be found at www.atcc.org.

### Handling Procedures

#### Centrifuge tube prior to opening to prevent loss of pelleted material

- 1. Rehydrate contents of vial with molecular grade H<sub>2</sub>O. DNA is dried in 1X Tris buffer.
- 2. Place vial at 37°C for 1 hour or at +2°C to 8°C overnight.
- 3. For more complete rehydration and to fully recover DNA, incubate the sample overnight at 4°C while rocking; then incubate for 1 hour at 65°C. Resuspending the dried DNA in  $\geq$  250  $\mu$ L may give better results.

## **Quality Control Specifications**

**Electrophoresis - RNA content:** No RNA was detected by electrophoresis

Purity (A260/A280): 1.6 to 2.0

**Integrity:** Integrity of DNA was determined by electrophoresis on a 1% agarose gel stained with SYBR Safe<sup>™</sup>, and was found to be of high molecular weight.

**Functional tests:** Functional activity was confirmed by PCR amplification of approximately 1500 base pairs fragment of rRNA gene cluster including ITS1-5.8S-ITS2 region.

**Identity:** Identity confirmed by sequencing of ITS1, 5.8S gene and ITS2 regions of ribosomal RNA ( $\sim 500$  base pairs).

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Genomic DNA is appropriate for PCR and other molecular biology applications.

#### Material Citation

If use of this material results in a scientific publication, please cite the material in the following manner: Genomic DNA from *Nakaseomyces glabratus* strain VITEK 303542 (ATCC MYA-2950D-5)

#### References

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

## Warranty

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