Product Sheet

Genomic DNA from Saprolegnia diclina strain BA 810

56851D-2[™]

Description

Genomic DNA isolated from *Saprolegnia diclina* FBA 810 [203a]. This fungal strain is also available as ATCC[®] Catalog No.: 56851. **Organism:** *Saprolegnia diclina* Humphrey **Type strain:** No **Mass:** 2 μg **Shipping information:** Stored in 1X TE buffer

Storage Conditions Product format: Dried

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 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.

Certificate of Analysis

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

Quality Control Specifications

Electrophoresis - RNA content: No RNA was detected by electrophoresis Purity (A260/A280): 1.7 to 2.0 Integrity: Integrity of DNA was determined by electrophoresis on a 1% agarose gel stained with SYBR Safe[™], 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 (~ 500 base pairs).

Notes

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 *Saprolegnia diclina* strain BA 810 (ATCC 56851D-2)

References

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

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Page 3 of 5

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