



Product Sheet

Cryptococcus gattii (ATCC® MYA-4071™)

Please read this **FIRST**



Intended Use

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

Citation of Strain

If use of this culture results in a scientific publication, it should be cited in that manuscript in the following manner: *Cryptococcus gattii* (ATCC® MYA-4071™)

American Type Culture Collection
PO Box 1549
Manassas, VA 20108 USA
www.atcc.org

800.638.6597 or 703.365.2700
Fax: 703.365.2750
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Or contact your local distributor

Description

Strain Designation: WM 276

Deposited Name: *Cryptococcus bacillisporus*

Product Description:

An ampoule containing viable cells (e.g. yeast cells, spores, or agar cubes with mycelia) suspended in cryoprotectant.

Propagation

The information recommended in this section is to assist users in obtaining living culture(s) for their studies. The recommendation does not imply that the conditions or procedures provided below are optimum. Experienced researchers may initiate the growth of a culture in their own way.

ATCC® Medium 28: Emmons' modification of Sabouraud's agar

ATCC® Medium 200: YM agar or YM broth

ATCC® Medium 1245: YEPD

Growth Conditions

Temperature: 24°C to 26°C

Atmosphere: Typical aerobic

Recommended Procedure

Frozen ampoules packed in dry ice should either be thawed immediately or stored in liquid nitrogen. If liquid nitrogen storage facilities are not available, frozen ampoules may be stored at or below -70°C for approximately one week. **Do not under any circumstance store frozen ampoules at refrigerator freezer temperatures (generally -20°C).** Storage of frozen material at this temperature will result in the death of the culture.

1. To thaw a frozen ampoule, place in a **25-30 °C** water bath, until just thawed (**approximately 5 minutes**). Immerse the ampoule just sufficient to cover the frozen material. Do not agitate the ampoule.
2. Immediately after thawing, wipe down ampoule with 70% ethanol and aseptically transfer at least 50 µl (or 2-3 agar cubes) of the content onto a plate or broth with medium recommended.
3. Incubate the inoculum/strain at the temperature and conditions recommended.
4. Inspect for growth of the inoculum/strain regularly. The sign of viability is noticeable typically after 1-2 days of incubation. However, the time necessary for significant growth will vary from strain to strain.

Colony and Cell Morphology: After 5 days on Emmons' medium at 25°C, colony is cream-colored, smooth, mucoid. Cells are globose, single or with bud.

Notes

Genome sequencing strain (Canada's Michael Smith Genome Sciences Centre, Canada; University of British Columbia, Canada); For multigene phylogeny and phenotypic characterization, see Findley K. et al.; This isolate is infertile under Lab conditions. For related strains, see ATCC 32609, ATCC 208821, and ATCC MYA-4093. Additional, updated information on this product may be available on the ATCC web site at www.atcc.org.

DNA Sequence

>Cryptococcus gattii ATCC MYA-4071 18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 26S ribosomal RNA gene, partial sequence

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AAGGATCAGTAGAGAATGCTGGGCTTCGGTCCATTTATCTACCCATCTACACCTGTGAACTGTTTATGT  
GCTTCGGCACGTTTTACACAAACTTCTAAATGTAATGAATGAATCTTATTATAACAATAATAAACTTT  
CAACAACGGATCTCTTGGCTTCCACATCGATGAAGAACGCAGCGAAATGCGATAAGTAATGTGAATTG  
CAGAATTCAGTGAATCATCGAGCTTTGAACGCAACTTGCGCCCTTTGGTATTCCGAAGGGCATGCCTG  
TTTGAGAGTCATGAAAATCTCAATCCCTCGGGTTTTATTACCTGTTGGACTTGGATTGGGTGTTTGCCG  
CGACCTGCAAAGGACGTCGGCTCGCCTTAAATGTGTTAGTGGGAAGGTGATTACCTGTCAGCCCGGCG  
TAATAAGTTTCGCTGGGCCTATGGGGTAGTCTTCGGCTTGCTGATAACAACCATCTCTTTTTTTGTTTGACC  
TCAATCAGGTAGGGCTACCCGCTGAACCTAA
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>Large subunit ribosomal RNA gene, partial sequence.

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ATATCAATAAGCGGAGGAAAAGAACTAACAAGGATTCCCTTAGTAACGGCGAGTGAACCGGGAAG  
AGCTCAAATTTGAAATCTGGCGTCCCGGGCGTCCGAGTTGTAATCTACAGAACGTTTTCCGTGCTG  
GACCGTGTCTAAGTCCCTTGGAAATAGGGTATCAAAGAGGGTGACAATCCCGTACTTGACACGATCACC  
AGTGCTCTGTGATACGTTTTCTACGAGTCGCGTTACTTGGGAGTGTAGCGCAAATGGGTGGTAAACTC  
CATCTAAAGCTAAATATTGGTGAAGACCGATAGCGAACAAGTACCGTGAGGGAAAGATGAAAAGC
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ACTTTGAAAGAGAGTAAACAGTACGTGAAATTGTTGAAAGGAAACGATTGAAGTCAGTCGTGTC
TATTGGGTTTCAGCCAGCTCTGCTGGTGTATTCCCTTTAGACGGGTCAACATCAGTTCTGATCGGGTGGATA
AGGGCTGGAGGAATGTGGCACTCTTCGGGGTGTGTTATAGCCTCCTGTCCATACACTGGTTGGGACTG
AGGAATGCAGCTCGCCTTTATGGCCGGGGTTCGCCACGTTTCGAGCTTAGGATGTTGACAAAATGGCTT
TAAACGA



Eucalyptus tereticornis debris
Australia
Isolation date: December, 1993



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



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

Additional information on this culture is available on the ATCC web site at www.atcc.org.
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