



Product Sheet

Aspergillus fischeri var. *fischeri* (ATCC[®] 1020[™])

Please read this **FIRST**



Storage Temp.
Frozen: -80°C or colder
Freeze-Dried: 2°C to 8°C
Live Culture: See Propagation Section



Biosafety Level
1

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: *Aspergillus fischeri* var. *fischeri* (ATCC[®] 1020[™])

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
Email: Tech@atcc.org

Or contact your local distributor

Description

Strain Designation: NRRL 181 [4651.2, CBS 101.12, CBS 544.65, DSM 3700, IMI 211391, QM 1983, WB 181]
Deposited Name: *Aspergillus fischeri* Wehmer var. *fischeri*

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 336: Potato dextrose agar (PDA)
ATCC[®] Medium 325: Malt extract agar (Blakeslee's formula)

Growth Conditions

Temperature: 24°C to 26°C
Atmosphere: Typical aerobic

Recommended Procedure

For **freeze-dry (lyophilized)** ampoules:

1. Open an ampoule according to enclosed instructions.
2. From a single test tube of **sterile distilled water** (5 to 6 mL), withdraw approximately 0.5 to 1.0 mL with a sterile pipette and apply directly to the pellet. Stir to form a suspension.
3. Aseptically transfer the suspension back into the test tube of sterile distilled water.
4. Let the test tube sit at room temperature (25°C) undisturbed for **at least 2 hours**; longer (e.g., overnight) rehydration might increase viability of some fungi.
5. Mix the suspension well. Use several drops (or make dilutions if desired) to inoculate recommended solid or liquid medium. Include a control that receives no inoculum.
6. Incubate the inoculum at the propagation conditions recommended.
7. Inspect for growth of the inoculum/strain regularly. The sign of viability is noticeable typically after 2-3 days of incubation. However, the time necessary for significant growth will vary from strain to strain.

Notes

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

DNA Sequence

18S ribosomal RNA gene, partial sequence; internal transcribed spacer 1, 5.8S ribosomal RNA gene, and internal transcribed spacer 2, complete sequence; and 28S ribosomal RNA gene, partial sequence
AAGGATCATTACCGAGTGAGGGCCCTCTGGGTCCAACCTCCCACCCGTGTCTATCGTACCTTGTGCTTC
GGCGGGCCCGCCGTTTCGACGGCCGCGGGGAGGCCTCGCGCCCCGGGGCCCGCGCCCGCGAAGAC
CCCAACATGAACGCTGTTCTGAAAGTATGCAGTCTGAGTTGATTATCATAATCAGTAAAACTTTCAAC
AACGGATCTCTTGGTTCCGGCATCGATGAAGAACGCAGCGAAAATCGGATAAGTAAATGTGAATTGCAG
AATTCAGTGAATCATCGAGTCTTTGAACGCACATTGCGCCCCCTGGTATTCCGGGGGCGATGCCTGTCC
GAGCGTCATTGCTGCCCTCAAGCACGGCTTGTGTGTTGGGCCCCCGTCCCCTCTCCCGGGGACGGGCC
GAAAGGCAGCGGGCCACCGCTCCGGTCCCGAGCGTATGGGCTTTGTACCCCGCTCTGTAGGCC
GGCCGGCGCCAGCCGACACCACTTTATTTCTAAGGTTGACCTCGGATCAGGTAGGGATACCCGCTG
AACTTAAG

D1D2 region of the 28S ribosomal RNA gene

CATATCAATAAGCGGAGGAAAAGAAACCAACAGGGATTGCCTCAGTAACGGCGAGTGAAGCGGCAA
GAGCTCAAATTTGAAAGCTGGCCCCCTCGGGGTCCGCGTTGTAATTTGCAGAGGATGCTTCGGGTGCA
GCCCCCGTCTAAGTGCCTGGAACGGCCGTCATAGAGGTGAGAATCCCGTCTGGGACGGGGTGTCT
GCGTCCGTGTGAAGCTCCTTCGACGAGTCGAGTTGTTGGGAATGCAGCTCTAAATGGGTGGTAAATTT
CATCTAAAGCTAAATACTGGCCGGAGACCGATAGCGCACAAAGTAGAGTGATCGAAAGATGAAAAGC
ACTTTGAAAAGAGAGTTAAACAGCACGTGAAATTGTTGAAAGGGAAGCGTTTTCGACCCAGACTCGCC
CGCGGGGTTACGCGGCCATTCGTGCGGTTACTTCCCCTGGGCGGGCCAGCGTCCGTTTGGCGGGC
CGGTCAAAGGCCCTCGGAATGTATCACCTCTCGGGGTGCTTATAGCCGAGGGTGAATGCGGCCTGC
CCGACCGAGGAACGCGCTTCGGCTCGGACGCTGGCGTAATGGTCGCAATGA

Isolation

Canned apples



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References

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Biosafety Level: 1

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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