



Product Sheet

Candida parapolymorpha (ATCC® 26012™)

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: *Candida parapolymorpha* (ATCC® 26012™)

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: DL-1 [NRRL Y-7560]

Deposited Name: *Hansenula polymorpha* de Morais et Maia, teleomorph

Product Description: An ampoule containing viable cells (may include spores and 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°C to 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 10 µL (or any amount desired up to all) 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.

Notes

type strain; produces 2-butanone alcohol dehydrogenase; produces formaldehyde; produces hydrogen peroxide; produces isopropanol dehydrogenase (NADP) secondary alcohol dehydrogenase; produces methyl ketones; transformation host; utilizes methanol; reduces 4-chloro-1-(4-fluorophenyl)butan-1-one; transformation host for expression of hirudin gene.

DNA Sequence

SSU rDNA of ATCC 26012

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AAAGATTAAGCCATGCATGTCTAAGTATAAGCAATTTATACAGTAACTGCGAATGGCTCATTAAAT
CAGTTATCGTTTATTTGATAGTTCCTTTACTACATGGATAACCGTGGTAATTCTAGAGCTAATACATGCTT
AAAGCCCCGACTTCTGGAAGGGGTGTATTTATTAGATAAAAAATCAATGCCTTCGGGCTTTTTGATGAT
TCATAATAACTTTTCGAAGCTCATGGCCTTGTGCTGGAGCTGGTTCATTCAAATTTCTGCCATCAACTT
TCGATGGTAGGATAGTGGCCTACCATGGTTTTACGGGTAACGGGGAATAAGGGTTTCGATTCGGGAGA
GGGAGCCTGAGAAACGGCTACCACATCCAAGGAAGGCAGCAGCGCGCAAAATTACCCAATCCTGAC
ACAGGGAGGTAGTGACAATATATAACGATACAGGGCCCTTTCGGGTCTTGTAAATGGAATGAGTACAA
TGTAATACCTTAACGAGGAACAATTGGAGGGCAAGTCTGGTGCCAGCAGCCGCGGAATTCAGCTC
CAATAGCGTATATTAAGTTGTTGCAGTTAAAAAGCTCGTAGTTGAACCTTGGGTCTGGTTAGCCGGTC
TGCCTTTTTGGCAGCTACTGGATTAACCGGGCCTTTCCTTCTGGCTAACCAACACTCTGTGTTGGCAA
ACCAGGACTTTTACTTTGAAAAAATTAGAGTGTTCAAAGCAGGCCCTTTCCTCGAATATATTAGCATGGA
ATAATAGAATAGGACGTTATGGTTCTATTTTGTGGTTTCTAGGACCATCGTAATGATTAATAGGGACG
GTCGGGGGCATCAGTATTCAGTTGTCAGAGGTGAAATCTTGGATTTACTGAAGACTAACTACTGCGA
AAGCATTTGGCAAGGACGTTTTTCATTAATCAAGAACGAAAGTTAGGGGATCGAAGATGATCAGATAC
CGTCGTAGTCTTAACCATAAACTATGCCGACTAGGGATCGGGTGGTGTTCCTTTTTTGACCCACTCGGCA
CCTTACGAGAAATCAAAGTTTTTGGGTTCTGGGGGAGTATGGTCGAAGGCTGAAACTTAAAGGAAT
TGACGGAAGGGCACCACCAGGAGTGAGCCTGCGGCTTAATTTGACTCAACACGGGGAACTCACCA
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GGTCCAGACACAATAAGGATTGACAGATTGAGAGCTCTTTCTTGATTTTGTGGGTGGTGGTGCATGGCC
GTTCTTAGTTGGTGGAGTGATTTGCTGCTTAATTGCGATAACGAACGAGACCTTAACCTACTAAATAG
TGATATTGGCTTTTGTGATTTTCACTTCTTAGAGGGACTATCGGTTTCAAGCCGATGGAAGTTTGAGGC
AATAACAGGTCTGTGATGCCCTTAGACGTTCTGGGCCGCACGCGCTACACTGACGGAGCCAACGAG
CTCTATCCTTGACCGAGAGGTCCGGTAATCTTGTGAAACTCCGTCGTGGGATAGAGCATTGCAA
TTATTGCTTTCAACGAGGAATTCCTAGTAAGCGCAAGTCATCAGCTTGCCTTGATTACGTCCCTGCCCT
TTGTACACACCGCCGCTGCTAGTACCGATTGAATGGCTTAGTGAGGCTTCAGGATTGGTTAGAGAA
GGGGGCAACTCCATCTTGAACCGAGAATCTAGTCAAACCTGGTCATTAGAGGAACTAAA

ITS of ATCC 26012

TTTCCGTAGGTGAACCTGCGGAAGGATCATTACAGTATTCTTCTAAGCTGGAGCTGTTAATTGATTGAA
CTGTTTGCCTGCGTGCTGAAACTGCGTGCGCGGTGATGGTTTTTTGGTTGCAGTTCTACTCAAGGC
TTAGAAATCTCCTACACACTTTGTTTTCTATTAGAGACAAATTTGCTTTGCTAGCGGTGTTAAACG
CGGCATGCCTGGCCAGAGGTTTTAACAAAACCTATTTTCGCTTGCCCTCGTGGGAAGATTGAAAACG
GTATTATTATTCGTCAAGTTATAAGAAGTTGCGTTTTGCAGCTTTTCTAAACTTTAAAACCTTCAACAACG
GATCTCTTGGTTCTCGCATCGATGAAGAACGACGAAATGCGATACGTAATGTGAATTGCAGATTTTC
GTGAATCATTGAATCTTGAACGCACATTGCGCCCTCTGGTATTCCAGAGGGCATGCCTGTTTGAGCGT
CATTCCCTCTCAAACCCCGGGTTTGGTATGGCAACTACTCTCGTCTGAGAGTTTCTTGAATGAAT
TGGCATGAGTTTTGATACGAACCTCAGCTGTTTATTCAATGTATTAGTTTTATCCATCGTTGATTGACA
ACTGCAAGCTTCTTCAAATTTGGCTCTGCCTTACAACCTTAATCATAAAGTTTGACCTCAAATCAGG
TAGGATTACCCGCTGAACTTAAGCATATCAAT

D1/D2 region of LSU rDNA of ATCC 26012

ATATCAATAAGCGGAGGAAAAGAAACCAACAGGGATTGCCTTAGTAGCGGCGAGTGAAGCGGCAAG
AGCTCAAATTTGAAATCTGGTACCTTCGGTGCCCGAGTTGTAATTTGAAGAAAGTAATCTTGAGGTTG
CCTTTGTCTATGTTCTTGAACAGGACGTCATGGAGGTGAGAAATCCCGTGTGATGAGGTGCCATCT
CCGTGTAAGATGCTTTCGAAGAGTGCAGTTGTTGGGAATGCAGCTCAAAGTGGGTGTTAAATCCAT
CTAAAGCTAAATTTGGCGAGAGACCGATAGCGAACAAGTACTGTGAAGGAAAGATGAAAAGAACT
TTGAAAAGAGAGTGAAGAAAGTACGTGAAATGTTGAAAGGGAAGGGTATTTGATCAGACTTGGTATT
TAGCTATCATCGCTCCTTGTGGGTGGTCTAGCTTTTACTGGCCAGCATCAGTTTTTGGTGGCAAGA
TAATGACAGTTGAATGTGGCTCCTCGGAGTGTTATAGCTTCTGTTGATATTGCCCTACCGGACTGAGGT
CTGCGCTTTTGCCTAGGATGCTGGCGTAATGATCCAATACCGCCGCT

Isolation

Soil, Massachusetts

References

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

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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