



# ***Acytostelium anastomosans* Cavender et al.**

**MYA-3266™**

## **Description**

**Strain designation:** OC6BA

**Type strain:** Yes; paratype

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## **Storage Conditions**

**Product format:** Frozen

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## **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.

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## **BSL 1**

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ATCC highly recommends that appropriate personal protective equipment is always

used when handling vials. For cultures that require storage in liquid nitrogen, it is important to note that some vials may leak when submersed in liquid nitrogen and will slowly fill with liquid nitrogen. Upon thawing, the conversion of the liquid nitrogen back to its gas phase may result in the vial exploding or blowing off its cap with dangerous force creating flying debris. Unless necessary, ATCC recommends that these cultures be stored in the vapor phase of liquid nitrogen rather than submersed in liquid nitrogen.

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## Certificate of Analysis

For batch-specific test results, refer to the applicable certificate of analysis that can be found at [www.atcc.org](http://www.atcc.org).

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## Growth Conditions

**Medium:**

ATCC Medium 919: Non-nutrient agar

**Temperature:** 20-25°C

**Incubation:** Grown in two-member culture with *Escherichia coli* ATCC 23437

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## Handling Procedures

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 minus 20°C).** Storage of frozen material at this temperature will result in the death of the culture.

1. Prepare media by streaking center of agar surface with a large X of *Escherichia coli* (ATCC 23437) and incubating at 25°-30°C one day before inoculating organism. Several replicates are suggested for optimum results.
2. To thaw frozen ampoule, place in a 37°C water bath, immerse ampoule to depth of one millimeter above the level of frozen material in the ampoule. Keep ampoule
- 3.
4. immersed until material is thawed but no more than 3 minutes. Do not agitate or vortex the ampoule.
5. Immediately after thawing, wipe down ampoule with 70% ethanol and aseptically transfer one loop full of contents onto center of X grown out with *Escherichia coli*.
6. Incubate the plates at the temperature recommended.
7. Allow culture to incubate for 2-5 days. Visually inspect by inverting plate under 10X objective. Look for swarms of amoebae feeding on bacteria and the initials of fruiting bodies and/or fruiting bodies rising from the surface of the agar.

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## **Material Citation**

If use of this material results in a scientific publication, please cite the material in the following manner: *Acytostelium anastomosans* Cavender et al. (ATCC MYA-3266)

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## **References**

References and other information relating to this material are available at [www.atcc.org](http://www.atcc.org).

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### **Contact Information**

ATCC

10801 University Boulevard

Manassas, VA 20110-2209

USA

US telephone: 800-638-6597

Worldwide telephone: +1-703-365-2700

Email: [tech@atcc.org](mailto:tech@atcc.org) or contact your local distributor

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