

Protocol for Malanga Root Tuber Homogenization in the Bullet Blender®

The protocol described in this document is for the use of the Bullet Blender® for the homogenization of malanga (*Xanthosoma sagittifolium*) root tubers. This protocol does not specify a particular buffer - you may choose which is most appropriate for your downstream application (nucleic acid isolation, protein extraction, etc.).

Materials Required: Malanga tuber, Bullet Blender®, homogenization buffer,

pipettor, microcentrifuge tubes, and 0.9-2.0mm stainless steel

bead blend or 1.0mm zirconium oxide beads (SSB14B or

ZROB10)

Instructions

- **1. OPTIONAL:** Wash malanga 3x with ~1mL PBS to remove soil and other surface contaminants and debris.
- **2.** Cut malanga into long, thin slices of 200mg or less and place each slice into a microcentrifuge tube.
- **3.** Add a a volume of beads equal to the mass of the malanga. **NOTE:** $100 \text{mg} \cong 100 \mu \text{L}$.
- **4.** Close the microcentrifuge tubes and place them into the Bullet Blender[®]. **NOTE:** There should be no buffer in the tubes at this point.
- 5. Set controls for SPEED 8 and TIME 4.
- **6.** Remove the samples from the Bullet Blender. The malanga should be finely pulverized into a thick paste. If not, run for another three minutes at speed 10.
- 7. Add 2 volumes of buffer to the tube for every mass of sample (ex. for 100 mg malanga add 200 μ L buffer).
- **8.** Close the microcentrifuge tubes and place them back into the Bullet Blender[®].
- 9. Set controls for SPEED 8 and TIME 3 minutes. Press Start.
- **10.** After the run, remove tubes from the instrument.
- **11.** Visually inspect samples. If homogenization is unsatisfactory, run for another three minutes at speed 10.
- **12.** Proceed with your downstream application.

SAFETY NOTE!!!

When using a centrifuge to separate your homogenate from the debris and beads, make sure your tubes are balanced.



Before



Pulverized



After

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