## Protocol for Leek Leaf Homogenization in the Bullet Blender<sup>®</sup>

The protocol described in this document is for the use of the Bullet Blender<sup>®</sup> for the homogenization of leek (*Allium ampeloprasum*) leaves. 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:

red: leek leaf, Bullet Blender<sup>®</sup>, homogenization buffer, pipettor, microcentrifuge tubes, and 0.9-2.0mm stainless steel bead blend (part number SSB14B)

## Instructions

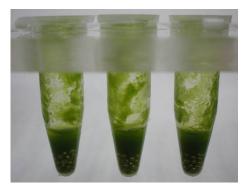
- **1.** Cut leaf into long, thin slices of 200mg or less and place each slice into a microcentrifuge tube.
- **2.** Add a volume of beads equal to the mass of tissue. **NOTE:**  $100 \text{mg} \approx 100 \mu \text{L}$ .
- **3.** Add 2 volumes of buffer to the tube for every mass of sample.
- **4.** Close the microcentrifuge tubes and place them into the Bullet Blender<sup>®</sup>.
- 5. Set controls for SPEED 9 and TIME 4 minutes. Press Start.
- **6.** After the run, remove tubes from the instrument.
- **7.** Visually inspect samples. If homogenization is unsatisfactory, run for another two minutes at speed 10.
- **8.** 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



After