



## Homogenization protocol for Ginger roots using Bullet Blender

### Materials:

Samples: 25 mg of fresh roots

Bead lysis kit: 5 mL Eppendorf Lysis Kit (SKU: NAVY5E20 or NAVE5E100)

Buffer volume: 2 mL of buffer

### Method - Homogenization:

1. Determine the sample size, buffer (depends on the downstream application), buffer volume and the bead lysis kit. **Note:** Choose the correct [lysis kit](#) for optimal homogenization.
2. Cut the sample and place into the buffer-filled tubes (Figure 1).
3. Close the tubes tightly and place into the Bullet Blender. **Note:** Confirm the compatibility of the [contact plate](#) with the tubes (RINO/EPPENDORF) used.
4. Set the speed and time on the Bullet Blender (Table 1). Press “Start”, and wait for the run to complete.

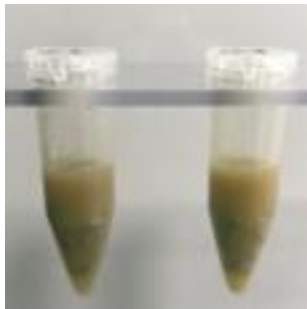
Bullet Blender Model	Settings
BB5E-Pro	Speed 12; Time 5

5. Remove the tubes and visually inspect the samples to confirm complete homogenization (Figure 2). **Note:** Foaming in the sample tubes may be observed after homogenization.
6. If homogenization is satisfactory, proceed with the downstream steps.

**Figure 1:** Pre-homogenization



**Figure 2:** Post-homogenization



*Homogenization verification*

