## **Ordering Information**

Cat. No	Description
SI-BG01 SI-BG05	Disruptor Beads 0.1mm Disruptor Beads 0.5mm
SI-D236 SI-D246 SI-D256 SI-D266 SI-D276 SI-D296 SI-D286	Disruptor Genie, 120V, 60Hz 0.65 amps, 1.5ml Disruptor Genie, 230V, 50Hz 0.5 amps, 1.5ml - No Plug Disruptor Genie, 230V, 50Hz 0.5 amps, 1.5ml - European Plug Disruptor Genie, 230V, 50Hz 0.5 amps, 1.5ml - British Plug Disruptor Genie, 230V, 50Hz 0.5 amps, 1.5ml - Swiss Plug Disruptor Genie, 230V, 50Hz 0.5 amps, 1.5ml - Australian Plug Disruptor Genie, 100V, 50/60Hz 1.0 amps, 1.5ml
SI-D237 SI-D247 SI-D257 SI-D267 SI-D277 SI-D297 SI-D287	Disruptor Genie, 120V, 60Hz 0.65 amps, 2.0ml Disruptor Genie, 230V, 50Hz 0.5 amps, 2.0ml - No Plug Disruptor Genie, 230V, 50Hz 0.5 amps, 2.0ml - European Plug Disruptor Genie, 230V, 50Hz 0.5 amps, 2.0ml - British Plug Disruptor Genie, 230V, 50Hz 0.5 amps, 2.0ml - Swiss Plug Disruptor Genie, 230V, 50Hz 0.5 amps, 2.0ml - Australian Plug Disruptor Genie, 100V, 50/60Hz 1.0 amps, 2.0ml
SI-0236 SI-0246 SI-0256 SI-0266 SI-0276 SI-0297 SI-0286	Vortex-Genie 2 (Model G560), 120V, 60Hz 0.65 amps Vortex-Genie 2 (Model G560E), 230V, 50Hz 0.5 amps - No Plug Vortex-Genie 2 (Model G560E), 230V, 50Hz 0.5 amps - European Plug Vortex-Genie 2 (Model G560E), 230V, 50Hz 0.5 amps - British Plug Vortex-Genie 2 (Model G560E), 230V, 50Hz 0.5 amps - Swiss Plug Vortex-Genie 2 (Model G560E), 230V, 50Hz 0.5 amps - Australian Plug Vortex-Genie 2, 100V, 50/60Hz 1.0 amps
SI-T236 SI-T246 SI-T256 SI-T266 SI-T276 SI-T296 SI-T286	Vortex-Genie 2T, 120V, 60Hz 0.65 amps Vortex-Genie 2T, 230V, 50Hz 0.5 amps - No Plug Vortex-Genie 2T, 230V, 50Hz 0.5 amps - European Plug Vortex-Genie 2T, 230V, 50Hz 0.5 amps - British Plug Vortex-Genie 2T, 230V, 50Hz 0.5 amps - Swiss Plug Vortex-Genie 2T, 230V, 50Hz 0.5 amps - Australian Plug Vortex-Genie 2T, 100V, 50/60Hz 1.0 amps
SI-A236 SI-A246 SI-A256 SI-A266 SI-A276 SI-A296 SI-A286	Digital Vortex-Genie 2, 120V, 60Hz 0.65 amps Digital Vortex-Genie 2, 230V, 50Hz 0.5 amps - No Plug Digital Vortex-Genie 2, 230V, 50Hz 0.5 amps - European Plug Digital Vortex-Genie 2, 230V, 50Hz 0.5 amps - British Plug Digital Vortex-Genie 2, 230V, 50Hz 0.5 amps - Swiss Plug Digital Vortex-Genie 2, 230V, 50Hz 0.5 amps - Australian Plug Digital Vortex-Genie 2, 100V, 50/60Hz 1.0 amps
SI-0563 SI-0562	TurboMix Attachment, 1.5ml Tubes TurboMix Attachment, 2.0ml Tubes

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# **DISRUPTOR BEADSTM**

# CELL DISRUPTION MEDIA REFERENCE GUIDE

Catalog No. SI-BG01 (0.1mm) & SI-BG05 (0.5mm)





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### Scientific Industries Disruptor Beads<sup>™</sup>

#### **OVERVIEW**

Spherical lead free soda lime glass beads are commonly used for mechanical disruption of many yeast, bacterial and soil samples. Glass beads of a pre-determined size and volume are placed in a 1.5ml or 2.0ml microtube along with a pre-determined sample amount. The closed tube is then shaken vigorously at high speed, causing collisions between the glass beads and sample material. Scientific Industries' Disruptor Genie<sup>®</sup> and TurboMix<sup>™</sup> attachment for the Vortex-Genie<sup>®</sup> 2 family of mixers are excellent choices for this process as they both simultaneously agitate and vortex at high speed, dramatically increasing cell or sample disruption. Each can hold up to twelve 1.5 ml or 2.0 ml microtubes at once. The disrupted cells may be removed after shaking for downstream processing.

Scientific Industries' Disruptor Beads are available in two sizes:

- 0.1 mm diameter beads (Catalog No. SI-BG01)— For use with Bacteria
- 0.5 mm diameter beads (*Catalog No. SI-BG05*)— For use with Yeast/Fungi

#### **CARE AND CLEANING**

Pre-preparation steps for Scientific Industries' Disruptor Beads are generally unnecessary. If desired, they may be soaked in a 1:8 dilution of household bleach for 20 minutes, rinsed with copious amounts of distilled or RO water, and baked at 50 to 65° C for a minimum of 2 hours, or until completely dry. If the glass beads do not pour freely, repeat the cleaning and drying process. Disruptor Beads may also be autoclaved after proper disinfecting or cleaning.

The Disruptor Beads may be reused, if desired, after proper disinfecting or cleaning and autoclaving. Subsequent uses and excessive handling of the beads may result in the creation of fines, which could adversely affect cell disruption efficiency. As such, it is not recommended to frequently reuse Disruptor Beads.

Disruptor Beads may be stored at room temperature or frozen in an airtight container prior to use. In addition, the Disruptor Genie and TurboMix attachment for the Vortex-Genie 2 and Vortex-Genie 2T may be used in cold rooms.

#### **SAMPLE APPLICATION METHODS**

NOTE: DETAILED DIRECTIONS FOR USE WILL DIFFER DEPENDING ON THE INDIVIDUAL PROTOCOL USED OR THE OUTCOME DESIRED. THE SAMPLE METHODS BELOW ARE EXAMPLES ONLY.

#### **Bacteria Disruption:**

Disruptor Beads, 0.1 mm diameter, are recommended for disruption of bacterial samples. A typical sample ratio would be 50% Disruptor Beads to 50% bacterial suspension by volume. This ratio may be adjusted as necessary. Allow head space (~20%) within the microtube to facilitate disruption action. It is recommended that beads and bacterial suspension be chilled prior to disrupting in order to offset any temperature rise within the microtube. Disruption at room temperature using chilled materials for 3 to 5 minutes at highest speed should be sufficient to recover 85% of the bacterial RNA. Disruption can be performed in a cold room as well. Samples should not be run for longer than 10 minutes consecutively to avoid any temperature rise.

#### Yeast/Fungi Disruption:

Disruptor Beads, 0.5 mm diameter, are recommended for disruption of yeast or fungi samples. A typical sample ratio would be 50% Disruptor Beads to 50% of yeast or fungus suspension by volume. This ratio may be adjusted as necessary. Allow head space (~20%) within the microtube to facilitate disruption action. It is recommended that beads and yeast or fungus suspension be chilled prior to disrupting in order to offset any temperature rise within the microtube. Yeast cells and fungi are generally more difficult to shear than bacterial cells, so increased disruption times may be necessary. Disruption in a cold room with chilled materials for 5 to 7 minutes at highest speed should be sufficient to disrupt the cell sample. Samples should not be run for longer than 10 minutes consecutively to avoid any temperature rise.

#### **Soil Sample Disruption:**

Either size of Disruptor Beads can be used for soil samples. A typical sample ratio would be 50% Disruptor Beads to 50% soil sample suspension by volume. Allow head space (~20%) within the microtube to facilitate disruption action. Samples should not be run for longer than 10 minutes consecutively to avoid any temperature rise.