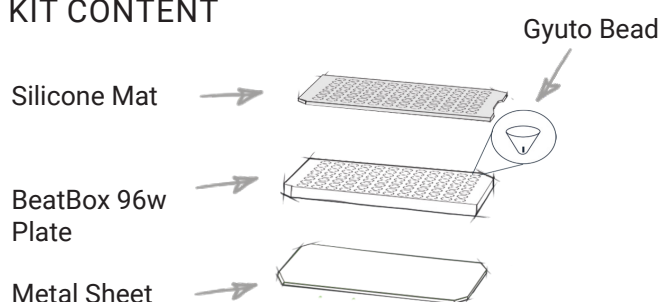
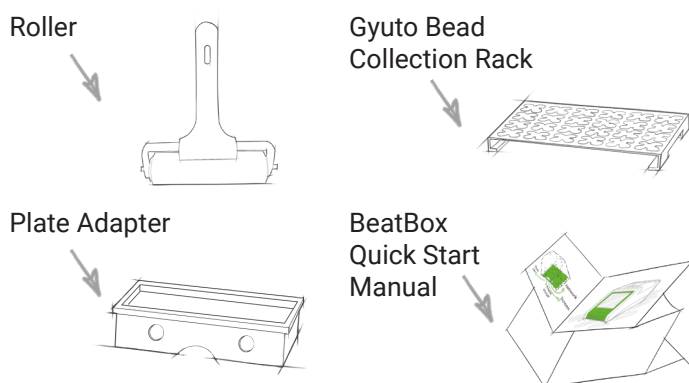


## KIT CONTENT



## BEATBOX ACCESSORIES



## Pre-Requisites

Common lab equipment is required for the sample preparation.

| Equipment              | Quantity and Description  |
|------------------------|---|
| CELL SAMPLE            | Various cell types can be processed, including mammalian, bacterial, and yeast cells. For other sample types contact PreOmics for adapted protocols.  |
| LYSIS BUFFER           | iST LYSE BUFFER from PreOmics' iST kits. For SP3-iST kits, contact info@preomics.com for an adapted protocol.   |
| 2-FOLD iST LYSE BUFFER | Optional: Required to continue with iST sample preparation, if larger volume of cell suspension is used. See protocol for details. 2-fold iST LYSE needs to be ordered in addition to the PreOmics' iST kits. |
| CENTRIFUGE             | Swing-bucket centrifuge for 96-well plate and adequate counterweight are required for spin-down of homogenate.  |

## Method

For a detailed description and graphical representation on how to use the BeatBox, please refer to the BeatBox Quick Start Manual 96x.

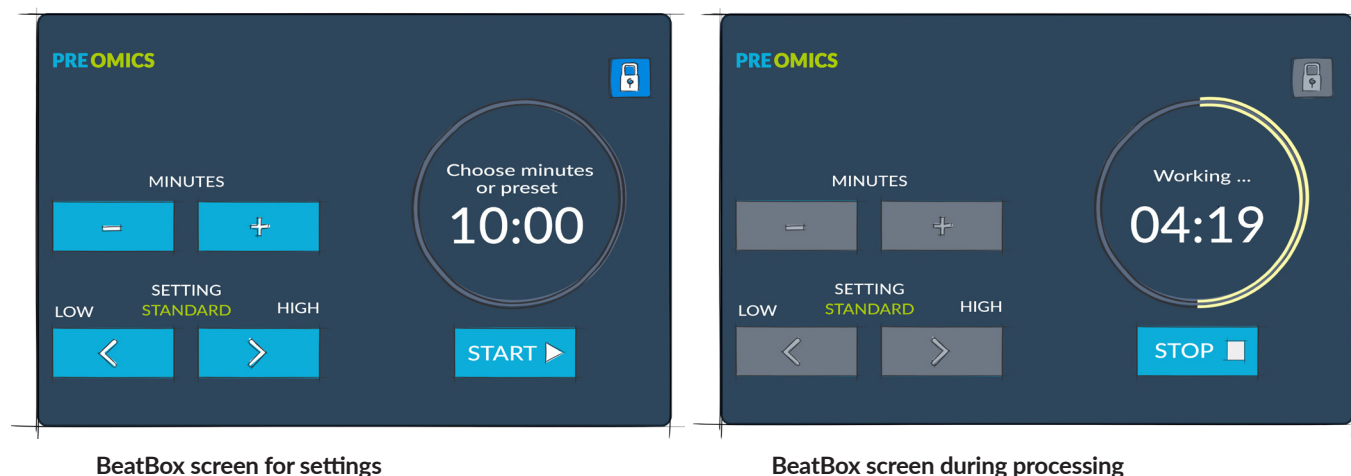
1. PLATE PREPARATION <sup>\*NOTE1\*</sup>

- 1.1. Remove the **SILICONE MAT** from the **BEATBOX 96w PLATE** while keeping the **METAL SHEET** attached to the base of the **BEATBOX 96w PLATE**.
- 1.2. Resuspend **CELL SAMPLE** in 1x PBS buffer. Transfer up to 50  $\mu$ L of cell suspension (equivalent to 100  $\mu$ g protein) into the well of the **BEATBOX 96w PLATE**. <sup>\*NOTE2\*</sup>
- 1.3. Add PreOmics' LYSE BUFFER to the cell suspension as follows:
  - For  $\leq 10$   $\mu$ L cell suspension, add 100  $\mu$ L of PreOmics' LYSE BUFFER.
  - For 11  $\mu$ L - 50  $\mu$ L cell suspension, add 50  $\mu$ L of 2-fold concentrated PreOmics' LYSE BUFFER and fill up to 100  $\mu$ L with LC-MS water.

- 1.4. Cover the **BEATBOX 96w PLATE** with the **SILICONE MAT** and make sure that the plate is properly closed by using the **ROLLER**. Remove the **METAL SHEET** from the base of the **BEATBOX 96w PLATE**.

## 2. BEATBOX LYSIS

- 2.1. Turn on the BeatBox, place the **BEATBOX 96w PLATE** on the PLATE ADAPTER and insert the PLATE and ADAPTER assembly into the GARAGE.
- 2.2. Use default configurations (SETTING: standard; MINUTES: 10 minutes) or optimize lysis conditions for your samples by adjusting SETTING and MINUTES in the BeatBox menu:



*SETTING: You can choose between LOW, STANDARD, HIGH. The power level increases from LOW to HIGH.*

*MINUTES: You can choose between 1 - 10 minutes (30 sec increments).*

- 2.3. Insert the GARAGE into the BeatBox and press START.
- 2.4. After the BeatBox run is completed, remove the GARAGE from the instrument, and the **BEATBOX 96w PLATE** from the ADAPTER.
- 2.5. Spin down the **BEATBOX 96w PLATE** (500 rcf; 30 - 60 sec).
- 2.6. Place the **BEATBOX 96w PLATE** on the GYUTO BEAD COLLECTION RACK and remove the **SILICONE MAT**. **\*NOTE3\***
- 2.7. Transfer the lysate into a new plate or tube for subsequent processing or analysis workflows.

## 3. CONTINUE WITH PREOMICS' KITS

- 3.1. Optional: Determine the protein concentration of the lysate.
- 3.2. Continue with PreOmics' kits:
- For iST kits, continue with the iST sample preparation workflow using up to 100 µg of extracted protein. Start with step "2. DIGEST" and follow the protocol.
  - For SP3-iST kits: Please contact [info@preomics.com](mailto:info@preomics.com).

**\*NOTE1\*** **SINGLE USE ONLY:** Kits components cannot be re-used.

**\*NOTE2\*** Various cell types can be processed on the BeatBox including mammalian, bacterial and yeast cells. Protein content varies considerably across distinct cell types and we recommend carrying out a protein concentration assay after the lysis step. A short overview of raw material amounts, as well as alternative buffers compatible with the iST workflow can be found in the FAQ (see [www.preomics.com/resources](http://www.preomics.com/resources)).

**\*NOTE3\*** In case of incomplete cell lysis, please repeat the BeatBox run (steps 2.1-2.6).