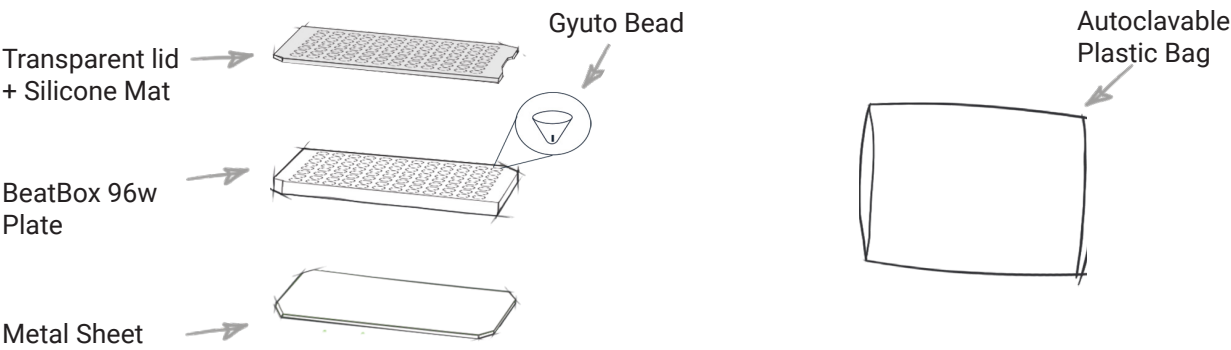


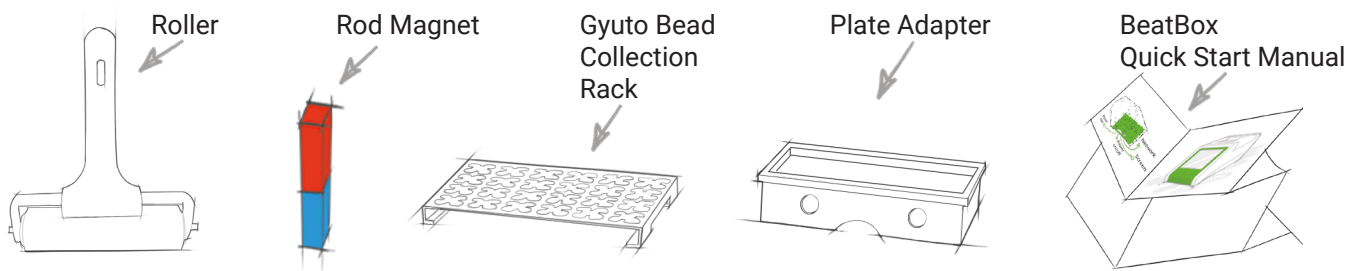
# BeatBox Tissue Kit 96x

Cell lysis protocol - Eucaryotic and procaryotic cells

## KIT CONTENT



## BEATBOX ACCESSORIES



## 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 \*NOTE1\*

- 1.1. Remove the **TRANSPARENT LID** and 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  $\mu\text{L}$  of cell suspension (equivalent to 100  $\mu\text{g}$  protein) into the well of the **BEATBOX 96w PLATE**. \*NOTE2\*
- 1.3. Add PreOmics LYSE buffer to the cell suspension as follows:
  - For  $\leq 10 \mu\text{L}$  cell suspension, add 100  $\mu\text{L}$  of PreOmics LYSE.
  - For 11  $\mu\text{L}$  - 50  $\mu\text{L}$  cell suspension, add 50  $\mu\text{L}$  of 2-fold concentrated PreOmics LYSE and fill up to 100  $\mu\text{L}$  with LC-MS water.
- 1.4. Cover the **BEATBOX 96w PLATE** with the **SILICONE MAT** and the **TRANSPARENT LID** and make sure that the plate is properly closed. Remove the **METAL SHEET** from the base of the **BEATBOX 96w PLATE**.



Want to know more? Visit the PreOmics website to learn more about BeatBox

PreOmics GmbH  
Martinsried | Germany

PreOmics Inc  
New York | USA

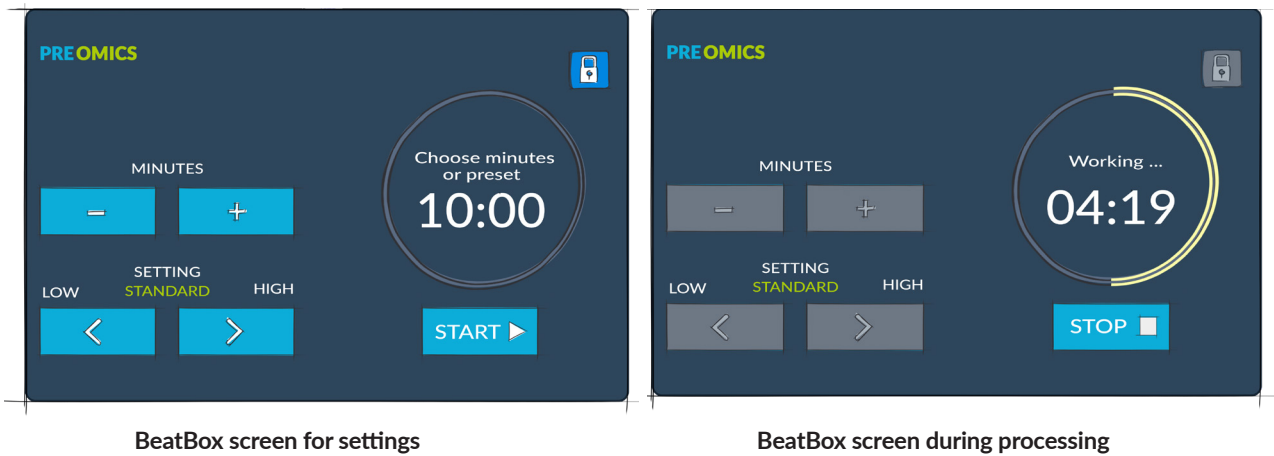
www.preomics.com  
beatboxsupport@preomics.com

## 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 **TRANSPARENT LID** and 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 protocol.
  - For SP3-iST kits: Please contact info@preomics.com.

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

When using the BeatBox for other sample types, please contact info@preomics.com.

*\*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).