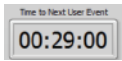


HMW DNA Extraction Kit

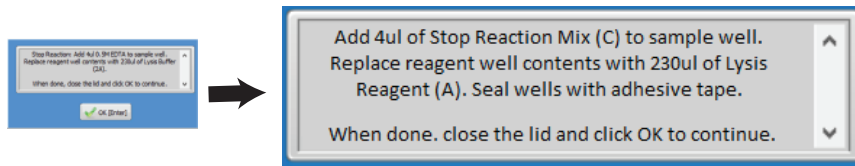
2. Prepare the Enzyme Reaction Mix:

- Remove **NEB Fragmentase** from the freezer, briefly vortex to mix
- To 800µl of **Enzyme Buffer**, add 2µl of **NEB Fragmentase** (1:400 dilution), vortex to mix

- Open the lid
- Remove the adhesive seals from the cassette(s)
- Replace the contents of the **sample wells** with **Fragmentase Reaction Mix, 80µl**
- Replace the contents of the **reagent wells** with **Enzyme Buffer (C), 230µl**
- Seal the wells with new adhesive tapes
- Close the lid.
- In the pop-up window, press "OK" to resume the workflow



- The User Event timer will countdown approximately 30 minutes. At the end of the count down, the instrument will pause and a pop-up window will appear:



- Open the lid
- Remove the adhesive seals from the cassette(s)
- To the **sample wells**, add 4µl of **Stop Reaction mix (D)**
- Replace the contents of the **reagent wells** with **Lysis Reagent (A), 230µl**
- Seal the wells with new adhesive tapes
- Close the lid.
- In the pop-up window, press "OK" to resume the workflow

H. Stage 3: Collection

- The User Event timer will countdown based on the collection waveform that was selected. At the end of the count down, the run will end, and samples may be removed. This step can take several hours.

Caution! Use wide-bore pipette tips to prevent shearing of DNA fragments.

Product: HEX-0004/0012
Revision A: March 27, 2017

Materials supplied with Cassette Kits:

4 /12	ea. Agarose Gel Cassettes	
20 /60	ea. Adhesive Tape Strips	
1	ea. HLS Lysis Reagent, 10/30 ml;	A
1	ea. Enzyme Buffer, 15/40 ml;	C
1	ea. Stop Reaction Mix, 250/250 µl;	D
1	ea. Running Buffer, 40/115 ml	E

Materials supplied or prepared by user:

NEBNext® dsDNA Fragmentase® (store at -20°C) **Cat# M0348S**

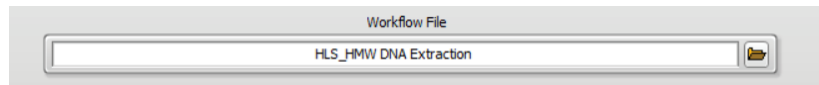
A. Prepare a cell suspension (refer to the guide provided with the cell suspension kit)

B. Prepare the Gel Cassette (refer to Section 4 in the SageHLS Operations Manual)

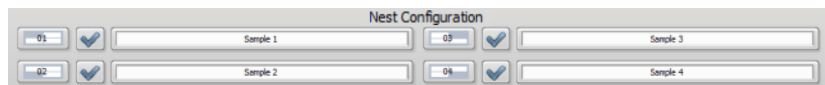
- Clear air bubbles around the perimeter of the gel columns
- Clear air bubbles from behind the elution wells
- Place cassette(s) on instrument nest(s) and remove adhesive tape(s)
- Replace the contents of the elution wells with **80µl** of **Running Buffer (E)**
- Add **Running Buffer (E)** to upper buffer chambers (see the schematic, next page) of each sample column until the buffer level is flush with the cassette cover

C. Prepare for the Run in the SageHLS Software

1. Go to the Main Screen
2. Press the folder icon and select "HLS_HMW DNA Extraction" from the file folder pop-up:



3. Select the sample lanes to be used by clicking the check boxes, and enter sample IDs in the text fields (optional):

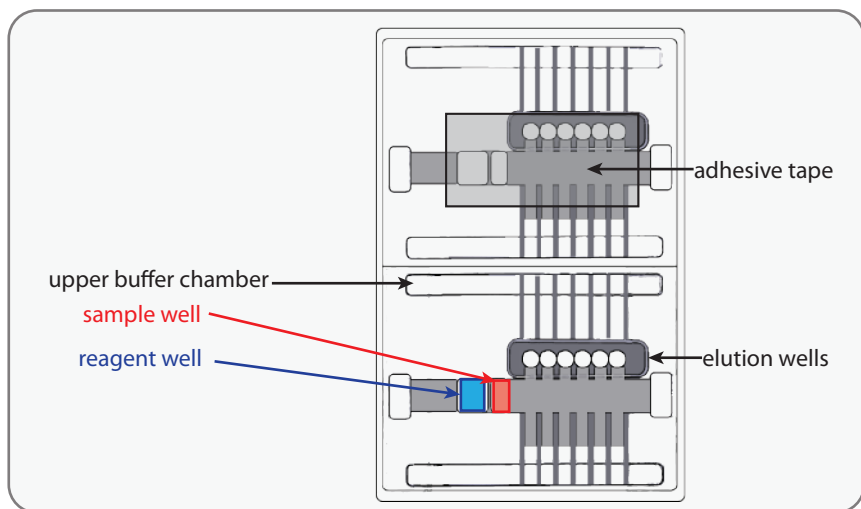


D. Run the Check Current Test

1. Close the lid
2. Press the "Check Current" button in the Command Menu:

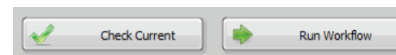


3. A pop-up window will appear. Press "Start" in the window.
4. At the completion of a successful Check Current test, press "Return":

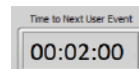


E. Begin the Run

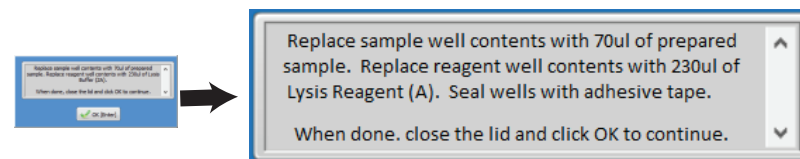
1. Press the "Run Workflow" button in the Command Menu:



F. Stage 1: Extraction



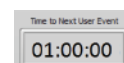
1. The User Event timer will countdown approximately 2 minutes. At the end of the count down, the instrument will pause and a pop-up window will appear:



2. Open the lid.
3. Replace the contents of the **sample wells** with **Cell Suspension, 70µl**
4. Replace the contents of the **reagent wells** with **Lysis Reagent (A), 230µl**
5. Seal the wells with adhesive tape
6. Close the lid.
7. In the pop-up window, press "OK" to resume the workflow

Caution! Without hurrying, users should minimize the length of the pause. It is important to load all sample wells first, and reagent wells second.

G. Stage 2: Treatment



1. The User Event timer will countdown approximately 1 hour. At the end of the count down, the instrument will pause and a pop-up window will appear:

