*GeneLab Standard Operating Procedure: Setting up NovaSeq 6000 and iSeq 100 sequencers*

*Sep 2021*

*Version 1.1*

# Document Revisions

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| --- | --- | --- | --- |
| Document Number | Revision Number | Date | Description of Changes |
| GL-SOP-7.1 | 1.0 | May 2020 | Original document |
| GL-SOP-7.1 | 1.1 | Sep 2021 | Added UPX kit NovaSeq read set up table  Added note on custom primers  Added step 19, to select “custom primers” in the NovaSeq set up window |
| GL-SOP-7.1 | 1.2 |  | Correct all tables for use of new sequencing reagent version |
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# Scope and Purpose

This procedure describes the workflows and parameters to follow in order to set up NovaSeq 6000 and/or iSeq 100 sequencers.

* If setting up iSeq 100 sequencer refer to procedure A.
* If setting up NovaSeq 6000 sequencer refer to procedure B.

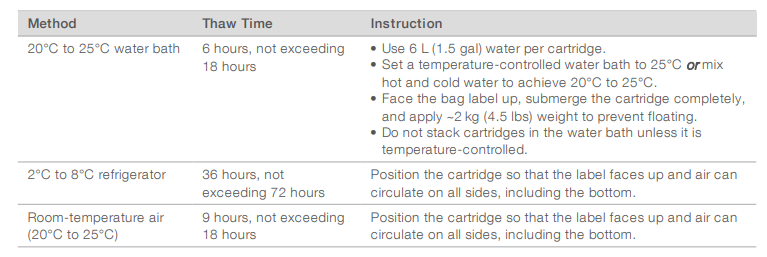
This SOP only describes the set-up of the sequencer, for all the other procedural details; make sure to read the most updated System Guides and the proper SOP’s.

# Procedure A: Reagents

1. iSeq 100 i1 Cartridge (Stored at -20°C)
2. iSeq 100 i1 Flow Cell (Stored at 4°C)
3. Consumables required for library dilution and spike in

# Procedure A: Setting up iSeq 100 sequencer

1. Thaw the iSeq cartridge and the flow cell according to the System Guide



Flow cell should be at room temperature at the time of sequencer loading.

1. Prepare SampleSheet file. Make sure to use the right configuration of i5 index (Index2).
2. Dilute the multiplexed library and spike the PhiX control according to the experimental design, dilute the spiked pool to loading concentration.
3. Load the library pool onto the cartridge following the System Guide.
4. Set up the sequencing run.

Local Run Manager mode:

* 1. Open iSeq Control Software.
  2. Select Local Run Manager 🡪 Open Local Run Manager.
  3. Create a sequencing run by Importing a sample sheet.
  4. Save run.
  5. In the control software, select “Sequence”. The Software will open the sequencer door and eject the tray.
  6. Place the sequencing cartridge with the flow cell and diluted, spiked library onto the tray.
  7. Select Close Door, the tray will retract and the door will close.
  8. If does not appear automatically, select the previously created run from the “Run Name” list. Try Refreshing if the run is missing.
  9. Make sure the read length parameters for Read 1, Index 1, Index 2, Read 2 fit with the planned experimental design:

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| **Kit** | **Number of Cycles** | **Requested read lenght** | **Adapters used** | **Read 1 Setting** | **Index 1(i7) Setting** | **Index 2(i5) Setting** | **Read 2 Setting** |
| i1 | 300 | PE150 | Illumina UDI | 151 | 8 | 8 | 151 |
| 300 | PE150 | IDT UDI + UMI | 148 | 17 | 8 | 148 |
| 300 | PE100 | Illumina UDI | 101 | 8 | 8 | 101 |
| 300 | PE100 | IDT UDI + UMI | 99 | 17 | 8 | 99 |
| 300 | PE50 | Illumina UDI | 51 | 8 | 8 | 51 |
| 300 | PE50 | IDT UDI + UMI | 49 | 17 | 8 | 49 |

**Note:** for 10X library sequencing, QIAgen QIAseq UPX 3’ Transcriptome kit, or Visium kits, follow the instructions provided by manufacturer for setting up the run on an iSeq instrument.

* 1. Make sure to record/take an image of the run set up window.
  2. Select Start Run.
  3. When run is finished, dispose of the used reagents according to the safety regulations.

Manual mode (without SampleSheet and Local Run Manager):

1. Open iSeq Control Software.
2. In the control software, select “Sequence”. The Software will open the sequencer door and eject the tray.
3. Place the sequencing cartridge with the flow cell and diluted, spiked library onto the tray.
4. Select Close Door, the tray will retract and the door will close.
5. In the “Run Name” field, enter a unique name, make sure to include the date, the project and any other information that might be useful in identifying this run.
6. Select “Single Read/Dual Read” option.
7. Enter the number of cycles to perform in each read following the planned experimental design.

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| **Kit** | **Number of Cycles** | **Requested read length** | **Adapters used** | **Read 1 Setting** | **Index 1(i7) Setting** | **Index 2(i5) Setting** | **Read 2 Setting** |
| i1 | 300 | PE150 | Illumina UDI | 151 | 8 | 8 | 151 |
| 300 | PE150 | IDT UDI + UMI | 148 | 17 | 8 | 148 |
| 300 | PE100 | Illumina UDI | 101 | 8 | 8 | 101 |
| 300 | PE100 | IDT UDI + UMI | 99 | 17 | 8 | 99 |
| 300 | PE50 | Illumina UDI | 51 | 8 | 8 | 51 |
| 300 | PE50 | IDT UDI + UMI | 49 | 17 | 8 | 49 |

**Note:** for 10X library sequencing, QIAgen UPX Transcriptome kit, or Visium kits, follow the instructions provided by manufacturer for setting up the run on an iSeq instrument.

1. Make sure to record/take an image of the run set up window.
2. Select Start Run.
3. When run is finished, dispose of the used reagents according to the safety regulations.

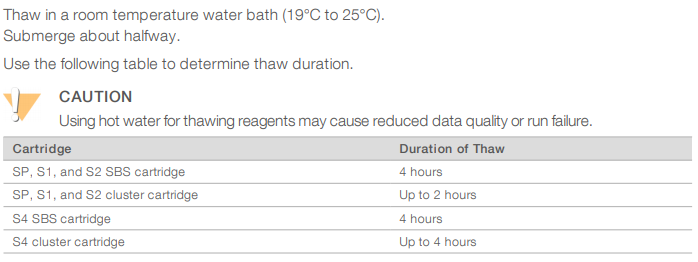
# Procedure B: Reagents

1. NovaSeq 6000 SBS Cartridge v1 (Stored at -20°C)
2. NovaSeq 6000 Cluster Cartridge v1 (Stored at -20°C)
3. NovaSeq 6000 Buffer Cartridge v1 (Stored at RT)
4. NovaSeq 6000 Flow Cell v1 (Stored at 4°C)
5. Consumables required for library dilution and spike in

* If using version 1.5 of sequencing reagents, longer read1 and read 2 is possible.

# Procedure B: Setting up NovaSeq 6000 sequencer

1. Thaw the two NovaSeq cartridges and the flow cell according to the System Guide:

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Set the flow cell package aside at room temperature for 10-15 minutes to allow it to reach room temperature.

1. Empty used reagent bottles from the NovaSeq, dispose according to laboratory safety regulations.
2. Change gloves after handling waste.
3. Prepare SampleSheet file. Make sure to use the right configuration of i5 index (Index2).
4. Dilute the multiplexed library and spike the PhiX control according to the experimental design, denature and dilute the spiked pool according to the guide.
5. Inspect the SBS and the Cluster cartridges, invert 10 times and tap the bottom on the bench.
6. Load the library into the library tube and insert into position #8 of the cluster cartridge.
7. In the control software, select “Sequence” and then select a single or dual flow cell run

The flow cell door will open.

1. With new powder free gloves, unpack the flow cell and inspect for defects.
2. Align the flow cell over the four raised clamps and place it on the flow cell stage.
3. Select “Close Flow Cell Door.”
4. Load the SBS, Cluster and Buffer Cartridges.
5. Select Checkbox acknowledging that the used reagent bottles have been emptied.
6. Select Run Setup button to enter run parameters.
7. Select workflow type (Xp/Standard).
8. In the “Run Name” field enter a unique name, make sure to include the date, the project and any other information that might be useful in identifying this run.
9. Enter the number of cycles to perform in each read following the planned experimental design and the reagents used to make the libraries and sequence:

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| **Kit** | **Number of Cycles** | **Requested read length** | **Adapters used** | **Read 1 Setting** | **Index 1(i7) Setting** | **Index 2(i5) Setting** | **Read 2 Setting** |
| SP | 100 | PE50 | Illumina UDI | 51 | 8 | 8 | 51 |
| PE50 | IDT UDI + UMI | 49 | 17 | 8 | 49 |
| 200 | PE100 | Illumina UDI | 101 | 8 | 8 | 101 |
| PE100 | IDT UDI + UMI | 99 | 17 | 8 | 99 |
| 300 | PE150 | Illumina UDI | 151 | 8 | 8 | 151 |
| PE150 | IDT UDI + UMI | 149 | 17 | 8 | 149 |
| 500 | PE250 | Illumina UDI | 251 | 8 | 8 | 251 |
| PE250 | IDT UDI + UMI | 247 | 17 | 8 | 247 |
| PE250 | Nextera Flex DNA | 249 | 10 | 10 | 249 |
| S1 | 100 | PE50 | Illumina UDI | 51 | 8 | 8 | 51 |
| PE50 | IDT UDI + UMI | 49 | 17 | 8 | 49 |
| 200 | PE100 | Illumina UDI | 101 | 8 | 8 | 101 |
| PE100 | IDT UDI + UMI | 99 | 17 | 8 | 99 |
| 300 | PE150 | Illumina UDI | 151 | 8 | 8 | 151 |
| PE150 | IDT UDI + UMI | 149 | 17 | 8 | 149 |
| S2 | 100 | PE50 | Illumina UDI | 51 | 8 | 8 | 51 |
| PE50 | IDT UDI + UMI | 49 | 17 | 8 | 49 |
| 200 | PE100 | Illumina UDI | 101 | 8 | 8 | 101 |
| PE100 | IDT UDI + UMI | 99 | 17 | 8 | 99 |
| 300 | PE150 | Illumina UDI | 151 | 8 | 8 | 151 |
| PE150 | IDT UDI + UMI | 149 | 17 | 8 | 149 |
| S4 | 200 | PE100 | Illumina UDI | 101 | 8 | 8 | 101 |
| PE100 | IDT UDI + UMI | 99 | 17 | 8 | 99 |
| 300 | PE150 | Illumina UDI | 151 | 8 | 8 | 151 |
| PE150 | IDT UDI + UMI | 149 | 17 | 8 | 149 |

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| 10X Single Cell | |  |  |  |  |  |  |
| **Kit** | **Number of Cycles** | **Requested read length** | **Adapters used** | **Read 1 Setting** | **Index 1(i7) Setting** | **Index 2(i5) Setting** | **Read 2 Setting** |
| Any | 200 | N/A | 10X adapters | 28 | 8 | 0 | 91 |
| Spatial Transcriptomics | | |  |  |  |  |  |
| **Kit** | **Number of Cycles** | **Requested read length** | **Adapters used** | **Read 1 Setting** | **Index 1(i7) Setting** | **Index 2(i5) Setting** | **Read 2 Setting** |
| Any | 200 | N/A | ST | 28 | 10 | 10 | 121 |
| QIAgen QIAseq UPX 3’ Transcriptome kit | | |  |  |  |  |  |
| **Kit** | **Number of Cycles** | **Requested read length** | **Adapters used** | **Read 1 Setting** | **Index 1(i7) Setting** | **Index 2(i5) Setting** | **Read 2 Setting** |
| Any | 100 | N/A | QIAgen sequencing adapters | 93 | 6 | 0 | 28 |

**Note:** QIAseq kit uses custom primers that are diluted, mixed and added to the sequencing cartridge. Make sure to follow QIAgen instructions for proper dilution of custom primers and NovaSeq user guide for setting up the sequencer to use custom primers.

1. Select output folder.
2. If custom primers are used, select “Custom primer”, the sequencer will use primers from a different well of the sequencing cartridge.
3. Confirm the run parameters on the Review screen.
4. Make sure to record/take an image of the run set up window.
5. Select “Start Run.”