

Clustering Sheet for Flowcell

BTR99602-2506

<b>Name</b> Bartlomiej Gebarski	<b>Sequencer</b> iSeq1	<b>Run Type</b> iSeq PE150 (r1: 150, r2: 150)
<b>Clustering Date</b> 2024-04-24	<b>Sequencing Date</b> 2024-04-24	<b>Status</b> Analyzed
<b>1st Index Length</b> 10	<b>2nd Index Length</b> 10	
<b>Comments</b> no comment	<b>Reagents Checked Out</b> <input type="checkbox"/>	

▪ Iseq 100 i1 Reagents: 1

num	Id	Scientist	Custom	PR	Comments
1	M18371_R17017_1	<div></div>	<input type="checkbox"/>	Standard	RNA Seq

## Instructions

1. Check, if instrument is idle. If run just finished - remove used cartridge (follow instructions on the screen).
2. Prepare Cluster Sheet, Sample and all reagents.
3. Take out the flowcell from the fridge to equilibrate it to room temperature (5-10min).
4. Install Flowcell in the cartridge
5. Mix all reagents in the Eppendorf tube:
  - **1.25µl** of **0.1nM** PhiX;
  - **0.99µl** of **2.5nM** Library;
  - **47.76µl** of Dilution Buffer;
6. Vortex and spin down.
7. Transfer **20 µl** into iSeq cartridge in well marked 'sample'.
8. Load reagents into instrument, following instructions on the iSeq's screen.