

Analysis Procedure – Multiplexed Microbial Assembly with SMRT® Link v6.0.0

For microbial multiplexing using SMRTbell® Template Prep Kit 1.0-SPv3 (PN 100-991-900), use the following Barcoded Adapter Kits (8A and 8B):

- Barcoded Adapter Kit 8A (PN 101-081-300): bc1001, bc1002, bc1004, bc1008, bc1009, bc1010, bc1012, bc1014
- Barcoded Adapter Kit 8B (PN 101-081-400): bc1015, bc1016, bc1017, bc1018, bc1019, bc1020, bc1021, bc1022

Set Up a Run on the Sequel® Instrument

Open the Run Design module in SMRT Link. Click **New Run Design**. Fill in the Sample Information section, then click the small arrow to open **Barcoded Sample Options**. Specify the following options:

1. Sample is Barcoded: **Yes**
2. Barcode Set: **Sequel_RSII_16_Barcodes_v2**
3. Same Barcodes on Both Ends of Sequence: **Yes**
4. Autofilled Barcode Name File: Click **Download File**
5. Barcoded Sample Name File: In the downloaded Autofilled Barcode Name File, fill in the desired values for Bio Sample Name, as shown below for the first 3 barcodes. Save the file and upload it using **Select CSV File**.

	A	B
1	Barcode Name	Bio Sample Name
2	bc1001_BAK8A--bc1001_BAK8A	E. coli
3	bc1002_BAK8A--bc1002_BAK8A	S. aureus
4	bc1004_BAK8A--bc1004_BAK8A	B. subtilis

6. Save the Run Design and launch the sequencing run.

Review the Resulting Data Sets

Once the run is complete, open the SMRT Link Data Management module. The barcoded samples are automatically demultiplexed. Click the number in the Demultiplexed Subsets column (**16** in this example) to view the barcoded Data Sets in a separate table:

Data Set Name	Demultiplexed Subsets	Bio Sample Name	Barcode Name
My microbial samples	16	[multiple]	[multiple]

Data Set Name	Bio Sample Name	Barcode Name
My microbial samples (E. coli)	E. coli	bc1001_BAK8A--bc1001_BAK8A
My microbial samples (S. aureus)	S. aureus	bc1002_BAK8A--bc1002_BAK8A
My microbial samples (B. subtilis)	B. subtilis	bc1004_BAK8A--bc1004_BAK8A

From the Data Management module home page, click the Data Set name of the parent sample (**My microbial samples** in the above example) to view the Data Set details. The Barcodes section displays the number of reads per barcode and several graphs to assess the quality of the demultiplexing results. Refer to the SMRT Link User Guide v6.0.0, Demultiplex Barcodes Application section, for further details on interpreting this report.

If review of the demultiplexing results reveals that any demultiplexing parameters need to be corrected (such as barcode set, barcode layout, or minimum barcode score), launch the **Demultiplex Barcodes** analysis application manually from the SMRT Analysis module, using the same Data Set as input.

Launch Assembly Analyses on the Microbial Data Sets

To launch one assembly analysis for each barcoded sample, open the SMRT Analysis module. Click **Create New Analysis**. Specify the following options:

1. Analysis Application: **Assembly (HGAP 4)**
2. Analysis Name: Enter an analysis name to persist across each analysis
3. Data Sets: Locate the parent Data Set of the demultiplexed set (**My microbial samples**, in the above example). Click the number in the Demultiplexed Subsets column (16) to view the child data sets. Click the checkbox in the column header to select all children.
4. Analysis Type: **One Analysis per Data Set – Custom Parameters**

Create New Analysis - Settings ✕ CANCEL ▶ START AND CREATE NEXT

Analysis Application: Assembly (HGAP 4)

Analysis Name: My microbial assembly - My microbial samples (E. coli)

Analysis of Multiple Data Sets: ← **Members of My microbial samples**

Analysis Type: One Analysis per Data Set - Custom Parameters

Data Sets: Advanced Search Search... ☰

<input checked="" type="checkbox"/>	Data Set Name	Bio Sample Name
<input checked="" type="checkbox"/>	My microbial samples (E. coli)	E. coli
<input checked="" type="checkbox"/>	My microbial samples (S. aureus)	S. aureus
<input checked="" type="checkbox"/>	My microbial samples (B. subtilis)	B. subtilis

Assembly

Genome Length: 5000000

ADVANCED ANALYSIS PARAMETERS

- For the first Data Set, edit the **Genome Length** parameter if desired.
- Click **Start and Create Next**. This launches the first assembly analysis on the first Data Set and proceeds to the launch page of the next Data Set, preserving any edits to the parameters made in the previous page.
- Edit the **Genome Length** parameter as appropriate for the second Data Set.
- Click **Start and Create Next**. Repeat the process for all Data Sets. After launching the last analysis, you are redirected to the SMRT Analysis module home page, which displays the status of each launched analysis.
- When an analysis completes, click its name to view the results.

▶ Analysis Overview	Polished Assembly	
▼ Polished Assembly		
Summary Metrics	Value	Analysis Metric
	1	Polished Contigs
	4,642,492	Maximum Contig Length
Contig Coverage vs Confidence	4,642,492	N50 Contig Length
	4,642,492	Sum of Contig Lengths
▶ Realignment to Draft Assembly	4,642,492	E-size (sum of squares / sum)
▶ Coverage		

Additional Troubleshooting Guidance

If the **Assembly (HGAP 4)** analysis generates a microbial genome assembly with >5 contigs, below are some additional suggestions to help improve the assembly, which can be tried individually or together.

In the Preassembly section of the analysis results page, check **Pre-Assembled Coverage** bases/genome_size).

1. If the preassembled coverage is below 20:
 - a. Launch a new **Assembly (HGAP 4)** analysis using the same input Data Set.
 - b. In the **Create New Analysis** page, click **Advanced Analysis Parameters** and change the **Seed coverage** parameter from the default of 30 to a level that will approximately compensate. For example, if your preassembled coverage was 10 instead of the desired 20, try changing the seed coverage to 60 to get double the preassembled coverage, since the preassembled yield tends to be similar as seed coverage increases. This is more frequently found in older sequencing chemistries prior to Sequel Sequencing Kit 3.0 and when library quality may be less than ideal.
 - c. Click **OK** to confirm the Advanced Analysis Parameters changes and run analysis.
2. If the preassembled coverage is above 20:
 - a. Launch a new **Assembly (HGAP 4)** analysis using the same input Data Set. In the **Create New Analysis** page, click **Advanced Analysis Parameters**.
 - b. In the **Assembly** section – **FALCON cfg overrides** type or copy/paste the following:
ovlp_DBsplit_option = -s50

Note that this setting does not always make an impact, but it can be helpful when there is a large difference in coverage across the genome, such as when the DNA was extracted from bacteria in an early log phase.
 - c. Click OK to confirm Advanced Parameters changes and run analysis.

Note: Be sure to use the troubleshooting guidance provided in this document for SMRT Link v6.0.0, as opposed to previous versions which are more relevant to SMRT Link v5.1.0. The guidance provided for earlier software versions are not forward compatible.

Circularizing Microbial Genomes

Circularizing the assembled genome is a step that can be performed after running **Assembly (HGAP 4)** to address issues with linear assemblies of circular genomes, such as removing common overlaps on each end of the contig. Circlator (<http://sanger-pathogens.github.io/circlator/>) is a tool to circularize genome assemblies that has been tested on PacBio data and that may be helpful to run downstream of **Assembly (HGAP 4)** analysis.

Revision History (Description)	Version	Date
Initial Release.	01	October 2018
Updates to consumables usage and troubleshooting guidance.	02	January 2019

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