SampleSheet Generation for PG-Seq[™] Rapid v2 kit

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Loading PG-Seq[™] Barcodes in Illumina[®] Experiment Manager*

*This has to be done only when using PG-seq for the first time

1. Go to C:\Program Files (x86)\Illumina \Illumina Experiment Manager

2. In the "SamplePrepKits" folder paste the PGSeq Adapters 12 nt SetA.txt and PGSeq Adapters 12 nt SetB.txt files

Computer	Local Disk (C:) Program Files (;	86) 🕨 Illumina 🕨 Illumina Expe	riment Manager
t View Tools I	Help		
e 🔻 🛛 Include in l	brary 👻 Share with 👻 New	folder	
prites	Name	Date modified	Туре
esktop	Applications	11/11/2018 20:14	File folder
ownloads	퉬 Genomes	11/11/2018 20:14	File folder
cent Places	퉬 IndexKits	06/07/2019 15:41	File folder
-epben	퉬 SamplePrepKits	06/07/2019 15:41	File folder

3. Go to "Applications" folder	Applications Genomes	View Tools	« Local Disk (C:) > Program Files (x86) > Illumina > Illumina Experiment Manager > Applications > ew Tools Help Include in library < Share with < New folder						
	IndexKits SamplePrepKits	Include in tes tcop nloads ent Places pbox Drive ies uments	tes ktop inloads ent Places pbox Drive Enrichment.bt es Figure GenerateFASTQ.bt		Type File folder Text Document Text Document Text Document Text Document Text Document Text Document Text Document				
4. Within this folder open the Resequencing.txt file		iic ures :05 iuter al Disk (C:) p (\\emeamilf t Generation S ro (165.88.37.	LibraryQC.txt Metagenomics.txt NextSeqGenerateFASTQ.txt NovaSeqGenerateFASTQ.txt PCRAmplicon.txt Plasmids.txt Resequencing.txt RNA-Seq.txt SmallRNA.txt Targeted RNA.txt	20/08/2018 9:22 21/05/2019 20:05 28/05/2019 12:30 20/08/2018 9:22 20/08/2018 9:22 20/08/2018 9:22 03/06/2019 14:27 20/08/2018 9:22 10/07/2019 15:46 20/08/2018 9:22	Text Document Text Document Text Document Text Document Text Document Text Document Text Document Text Document Text Document Text Document				

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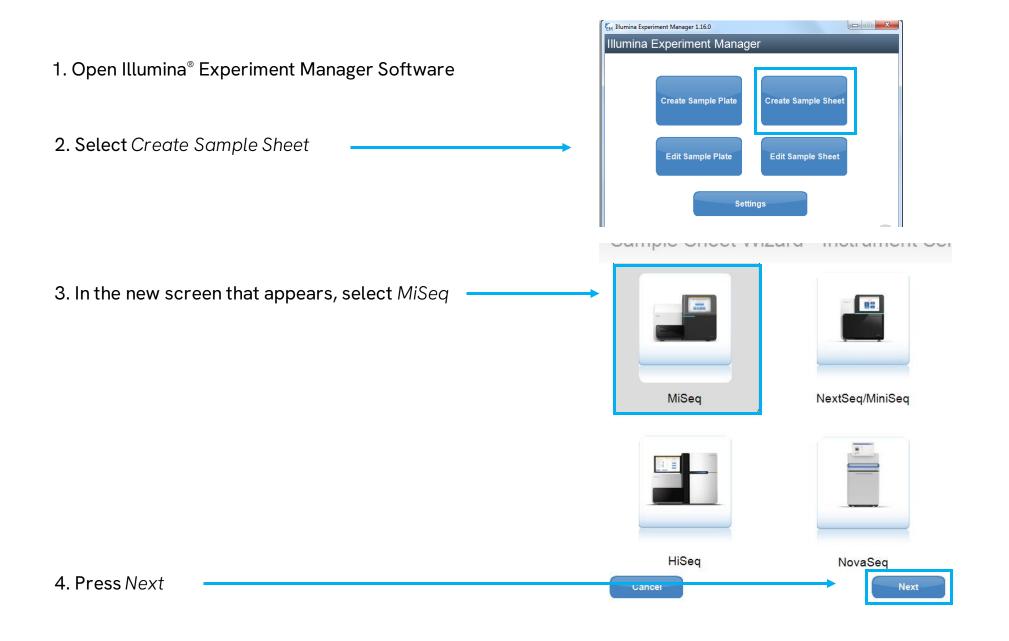
5. In the list of kits that appears add the txt files relevant to you

Resequencing.txt - Notepad	🗐 *Resequencing - Notepad — 🗆 🗙
	File Edit Format View Help
File Edit Format View Help	[Version]
[Version] 1 [Workflow Name] Resequencing [Display Name] Resequencing [Category] Small Genome Sequencing [Compatible Sample Prep Kits] Nextera DNA Nextera DNA Nextera DNA Nextera Mate Pair Nextera XT TruSeq DNA PCR-Free TruSeq Nano DNA	1 Workflow Name] Resequencing [Display Name] Resequencing [Category] Small Genome Sequencing [Compatible Sample Prep Kits] Nextera DNA Illumina DNA Prep Nextera Mate Pair Nextera XT TruSeq DNA PCR-Free TauSeq Mape DMA PGSeq Adapters 12 nt SetA PGSeq Adapters 12 nt SetB [Settings] [SampleSheetFields]

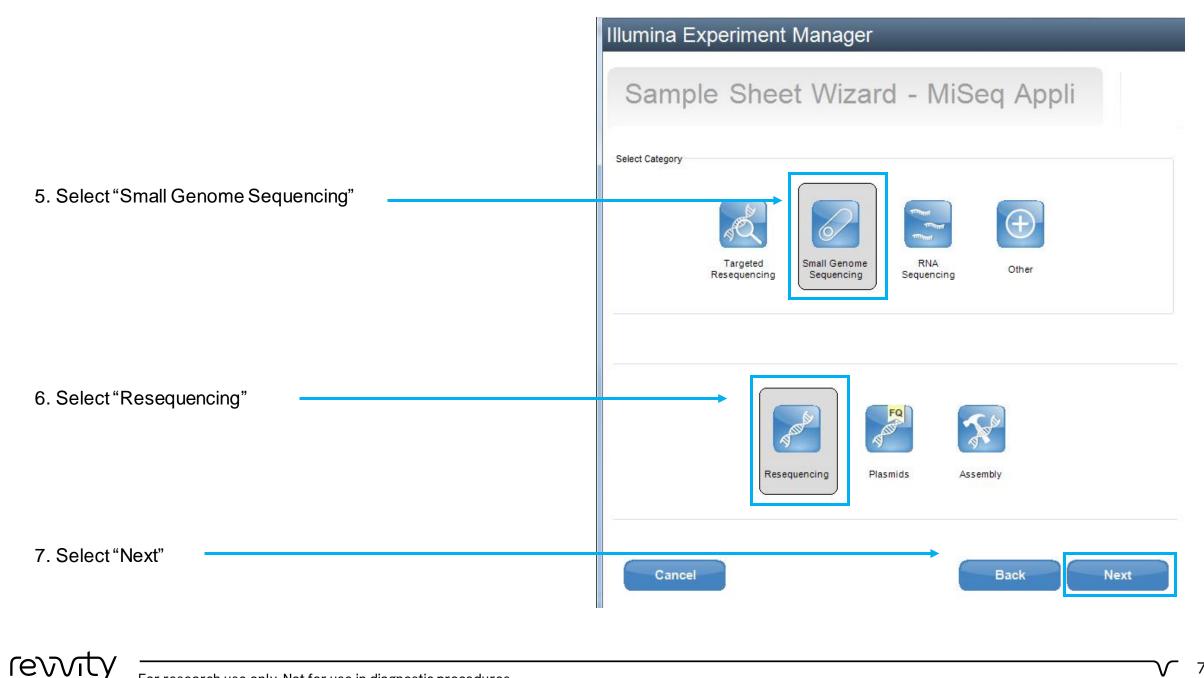
6. Save changes

Notepad	X
Files (x86)	ant to save changes to C:\Program \Illumina\Illumina Experiment Applications\Resequencing.txt?
	Save Don't Save Cancel

Creating a SampleSheet for PG-Seq[™] Barcodes



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8. Make sure screen looks EXACTLY like this (putting corresponding Reagent cartridge barcode and experiment name). Select the right Set according to the barcodes used

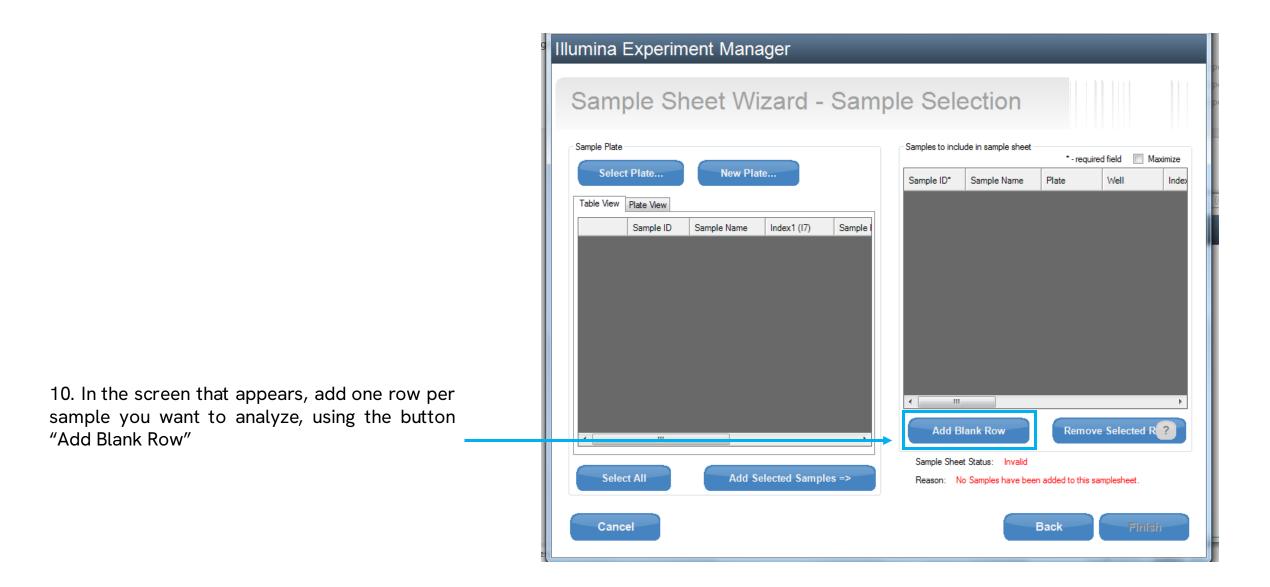
Set A \rightarrow Barcodes 1-96 Set B \rightarrow Barcodes 97-192

9. Press Next

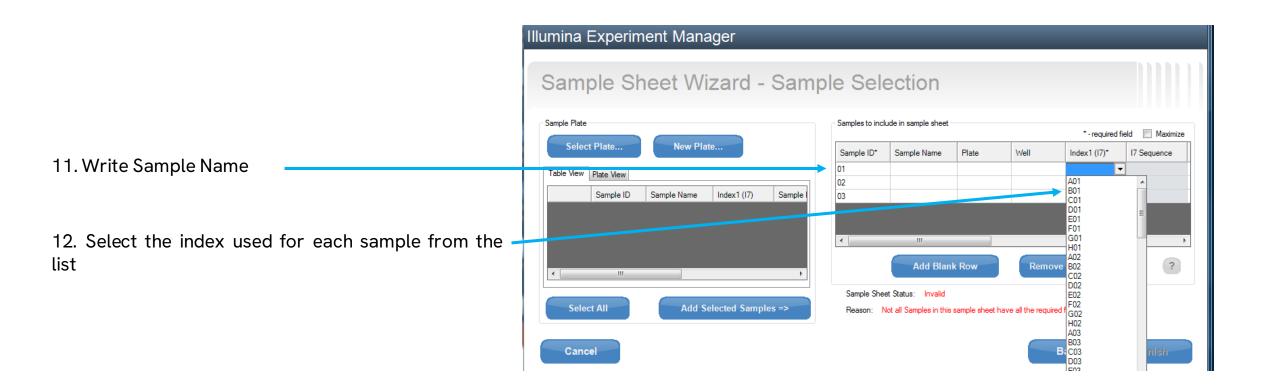
Illumina Experiment Manager

Sample Sheet Wizard - Workflow Parameters

Resequencing Run Settings		Resequencing Workflow-Specific Settin	ngs	
Reagent Cartridge Barcode*	Cartridge barcodes	Custom Primer for Read 1		^
Library Prep Workflow	PGSeq Adapters 12 nt SetA $\qquad \lor$	Custom Primer for Index		
Index Adapters	PGSeq Adapters 12 nt SetA $\qquad \lor$	Custom Primer for Read 2		
Index Reads	0 (None) 1 (Single) 2 (Dual)			
Experiment Name*	Experimental name here	Use Somatic Variant Caller		
Description		✓ Flag PCR Duplicates		
Date	21/03/2024	Reverse Complement		
Read Type	O Paired End	Indel Realignment GATK		
Cycles Read 1	76	Variant Quality Filter 30	÷	
				~
* - required field				
Cancel			Back	Next



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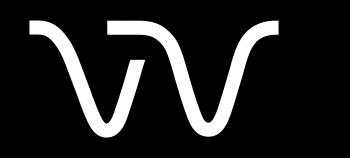


	Samples to include in sample sheet *- required field Maximize						
13. In the Genome Folder field, select Homo_sapiens	Sample Name	Plate	Well	Index1 (17)*	17 Sequence	Genome Folder*	
genome for all samples				A01	GGCCGGCTAG	Homo_sapiens\UCS(
6				B01	AAGGAAGAGA		
				C01	GGACGGCATC		

Sample Sheet Wizard - Sample Selection

	Sample Plate					- Samples to	include in samp	ole sheet		* - required field [Maniatian
	Selec	t Plate	New Plat	ie			Index1 (I7)*	17 Sequence	Genome Folder*	- required field	Sample
								-	Homo_sapiens\UCSC\hg19\Seq		
	Table View	Plate View					301		Homo_sapiens\UCSC\hg19\Seq		
	-	Sample ID	Sample Name	Index1 (I7)	Sample I		:01		Homo_sapiens\UCSC\hg19\Seq		
14. Press "Finish"	< Sele	ct All	Add Se	elected Samp	> les =>	 Sample Reason 	Sheet Status:	Add Blank Row Valid	Remove Selec	-	>
	Cano	el							Back		inish

15. Save the sample sheet



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