

PARAMETER	BASIC SPECS OF SEQUENCING PLATFORMS <sup>1</sup>			
	NovaSeq6000	MiSeq	iSeq	PacBio SEQUEL IIe
Chemistry	SBS reverse terminators Two-color	SBS reverse terminators Four-color	SBS reverse terminators Two-color	Direct SBS, Single molecule seq, cleavable fluor ( <a href="#">gammaP</a> )
Seq Configurations	Four types of flow cells (S4, S2, S1, SP) SE or PE 35-250 cycles	Single loading port per flow cell run	One flow cell	8 independent SMRT cells per run
Most common error	Substitutions, G-C bias	Substitutions, G-C bias	Substitutions, G-C bias	Insertions GC deletions
% Error rate <sup>3</sup>	<0.1	<0.1	<0.1	CLR (12-15), CCS/Hifi <0.1
Seq Read length	35-250 bp PE 250 only on SP	35-300 bp	35-150 bp	Large-insert genomic: 10-30 kb Amplicon, <a href="#">IsoSeq</a> : 30-50 kb
SE Reads/full run	S4: 10 Billion S2: 4 Billion S1: 1.6 Billion SP: 800 Million	10-13 M (v2) 20-25 M (v3)	4 M	3-5 M
Max output/ PE run (in Gb)	S4: 3000 Gb S2: 1250 Gb S1: 500 Gb SP: 250 Gb PE150, 400 Gb PE250	V3: 4-15 Gb V2: 0.75-7.5 Gb Micro: 1.2 Gb Nano: 0.5 Gb	1.2 Gb	Library- and movie-dependent • Large-insert genomic: 60-120 Gb • Amplicon, <a href="#">IsoSeq</a> : 80-200 Gb
Run time	1-2 days	1-3 days	10-20 hr	10h- 30h/ SMRT cell
<sup>4</sup> Cost/Gb in USD	S4: ~5-8 (format dependent) S2: ~8-20 S1: ~12-26 SP: ~12-35	V3: 127-325 V2: 200-1500 Micro: 542 Nano: 1100-1650	550	Library- and movie- dependent 12-30
Most common app	<ul style="list-style-type: none"> <li>Ultra-high throughput</li> <li>Whole genome seq</li> <li>Variant Analysis, genotyping</li> <li>Transcriptome (<a href="#">RNAseq</a>)</li> <li>ChIP seq, <a href="#">SeqCap</a> (ExomeSeq)</li> <li>Methylation Anal (WGBS, RRBS)</li> <li>ATAC-Seq</li> <li>Single-cell Seq</li> </ul>	<ul style="list-style-type: none"> <li>Amplicon (e.g., 16S metagenomics)</li> <li>Targeted <a href="#">RNAseq</a></li> <li>Variant Analysis</li> </ul>	<ul style="list-style-type: none"> <li>Library QC for balancing pools</li> <li>Small amplicon pools</li> </ul>	<ul style="list-style-type: none"> <li><i>De novo</i> sequencing</li> <li>Isoform analysis (<a href="#">IsoSeq</a>)</li> <li>Amplicon Seq</li> <li>SNP validation</li> <li>Direct methyl-seq</li> <li>Microbial seq</li> <li>Variant analysis</li> </ul>