

# Packages for Population Genomics

### Ultra-high Throughput with Flexible and Customizable Automation



✓ Automation from sample to report

## $\odot$ DNBSEQ-T10×4RS \* Introduction

DNBSEQ-T10 × 4RS is an ultra high-throughput sequencer based on DNBSEQ<sup>™</sup> sequencing technology. It is designed to meet the requirement of population genomics sequencing market.

Read length	PE100*
Effective reads**	360G
Average data output	72Tb
Run time***	96h
Data quality****	Q30 ≥ 85%

#### **Sequencing Specification**

One DNBSEQ-T10 × 4RS supports the operation of 8 sequencing slides simultaneously, producing up to 20Tb data per day(about 200 whole human genomes sequenced at 30 × ). A single set of DNBSEQ-T10 × 4RS can produce more than 50,000 WGS per year.

\* PE100 sequencing is recommended, PE150 will be available by 2021 Q2;

- \*\* The sequencing data was measured by running 8 sequencing slides;
- \*\*\* Run time includes DNB loading and sequencing. FASTQ file generation does not take up sequencer time
- \*\*\*\* ≥Q30 base ratio obtained by sequencing WGS standard library using DNBSEQ-T10×4RS.





The dip-immersion biochemistry, which ensures more uniform biochemical reaction of sequencing slide with larger size, as well as significantly saves the cost of reagents. Sequencing slide DNB loading density is up to 4 million/mm<sup>2</sup>, and one slide can load 70G DNB spots

#### **Innovative Sequencing Technologies**

Different from ordinary sequencing slide and closed reaction systems used in most platforms, DNBSEQ-T10 × 4RS uses **dip-immersion biochemistry** and open systems firstly to achieve the best balance among sequencing read length, throughput, data quality and cost.

	PE100				
Sample	NA12878	NA24694	NA24695		
Clean data rate	99.97	99.98	99.98		
Clean Q30(%)	91.565	91.625	91.62		
Mapping rate(%)	99.81	99.81	99.81		
Duplicate rate(%)	1.03	0.83	0.88		
Mismatch rate(%)	0.3	0.3	0.3		
Average sequencing depth(×)	30.66	30.76	30.71		
Coverage(%)	99.2	99.89	99.21		
Coverage at least 20×(%)	94.39	90.91	94.39		
SNP_Precision	0.9994	0.9991	0.9992		
SNP_Sensitivity	0.9908	0.9929	0.9927		
INDEL_Precision	0.9901	0.9918	0.9917		
INDEL_Sensitivity	0.9807	0.9803	0.9802		

#### **Sequencing Data Quality**

Human reference genomes NA12878, NA24694 and NA24695 were sequenced using DNBSEQ-T10 × 4RS. The bioinformatic analysis of 30 × WGS data showed that the accuracy and sensitivity of SNPs exceeded 99%, and the accuracy and sensitivity of Indel also exceeded 98%, reaching industry-leading level.

- Sample Human Cell Line
- Library Prep MGIEasy FS PCR-Free Kit DNA Library Prep Set
- Sequencing PE100
- O Data analysis MegaBOLT

## ©Total Package - Customizable and Automated



Calculation is based on human WGS sequenced at 30 imes depth.



#### Demo Planning for Annual Sequencing of 50,000 and 500,000 WGS\*

	<b>Zone</b> Biobank	Library center				
		Biobank	Library preparation	DNB preparation	Sequencing center	Data center
	Main Function	Whole blood separated into plasma and buffy coat for gDNA extraction	From gDNA to library construction	From library to make DNB	DNB loading and sequencing	Write FQ, data analysis, laboratory management system
50,000 WGS per year	Area(m <sup>2</sup> )**	50	50	50	100	20
	Key Equipment	MGISP-960	MGISP-96XL	MGISP-96XL	DNBSEQ-T10×4RS	ZLIMS/ZTRON/ MegaBOLT Pro
500,000 WGS per year	Area(m <sup>2</sup> )	100	150	150	500	50
	Key Equipment	MGIGLab-S MGISP-960	MGIGLab-L	MGISP-96XL	DNBSEQ-T10×4RS	ZLIMS Elite/ZTRON/ MegaBOLT Pro

Recommended equipment quantity, configuration and site area for annual sequencing capacity of 50,000~500,000 WGS. The recommended area is only for core functional zone. The actual area should be determined by the on-site conditions.

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