


GENE PANEL

EXOME

GENOME

SEQUENCER

PRODUCTS	BIOINFORMATICS PIPELINE	FUNCTION
 DNaseq (Sentieon)  TNseq (Sentieon)  VS-CNV	FASTQ BAM VCF	<ul style="list-style-type: none"> ▶ Single nucleotide variation ▶ Copy number variation & loss of heterozygosity ▶ Chromosomal aberration
Annotations	Annotated VCF	<ul style="list-style-type: none"> ▶ Public & commercial annotations to enrich genomic data sets
 VarSeq  VSReports  VSPipeline	Clinical Report	<ul style="list-style-type: none"> ▶ Annotate & filter ▶ Visually inspect alignments ▶ Variant prioritization ▶ Clinical assessment
 VSClinical	Automated ACMG & AMP Guidelines	<ul style="list-style-type: none"> ▶ Clinical variant interpretation in concordance with ACMG Guidelines & AMP Guidelines
 VSWarehouse	Data Warehousing Web-Enabled Interface + Powerful API: RESTful, JSON, TSV, CSV, Direct SQL	<ul style="list-style-type: none"> ▶ Clinical assessment catalog ▶ Advanced data querying ▶ Versioning ▶ Interoperability ▶ Compliance with HIPPA, CLIA & CAP data discovery