

DeepChek® Assay Whole Genome SARS-CoV-2 Genotyping

REF 159A48 **(€ IVD**



Flexible & Innovative End-to end Solution for SARS-CoV-2 Whole Genome Sequencing & Genotyping for Routine Diagnostic Settings











RNA Extraction

RT-PCR

NGS sequencing

Data analysis

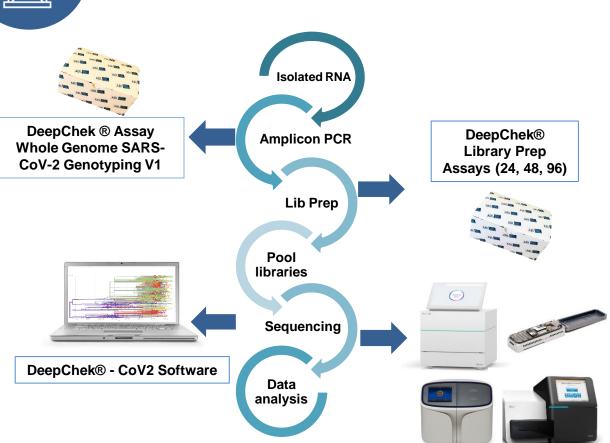
Automation (lab & IT)

- Next Generation Sequencing (NGS) Assay & Software
- Compatible with most NGS platforms (iSeq-100, MiSeq, MinIon...)
- Multiplex targeting SARS-CoV-2 Whole Genome (>99% genome covered)
- Flexible design for small to large sequencing series (<24, 48 or 96)
- Sample to result in 2 3 days
- Pooling of SARS-CoV-2 samples with other applications (HIV...)
- Downstream analysis software (variants & mutations detections, QC...)
- Ability to identify all VOC (Omicron, Delta, Alpha...) as well as well-characterized mutations (like E484K, D614G...)
- Up-to-date knowledge databases (viral infectivity, vaccine efficacy...) including
 Pangolin & NextClade
- Reporting capacity suited to routine diagnostics (PDF reports...), consensus..
- Automation capacity (lab robot & IT integrations LIMS) LargeScale Module
- Secured data hosting (healthcare-level compliance, local installations...)





WORKFLOW



NUMBER OF SARS-CoV-2 SAMPLES PER RUN

(can be pooled with other samples: HIV, HCV, 16s...)

NGS platform	Samples/run	
iSeq-100	24	
MiniSeq	24 – 96	
MiSeq	24 – 96	
NextSeq	24 – 96	
lon S5	24 – 96	
Minlon	>24	



REPORTING



DEEPCHEK® CoV-2 Software



READY-TO-USE GENOTYPING REPORT

Increased infectivity & transmissibility

s K417N s S477N s N501Y s D614G s P681H s H69_V70del

The Omicron variant has been classified as a Variant Of Concern (VOC) due to the presence of several mutations that may have an impact on transmissibility, severity of infection and vaccine effectiveness and due to preliminary evidence of an increased risk of reinfection.

Sample assigned to the Omicron variant (lineage B.1.1.529, clade 21K, first detected in South Africa on November 2021).

s G339D s S371L s K417N s N440K s G446S s S477N s T478K s E484A s Q493R s G496S s Q498R s N501Y s T547K s D614G s H655Y s A67V s N679K s P681H s N764K s D796Y s N856K s T951 s Q954H s N969K

Mutations characterizing the lineage were chosen according to the SARS-CoV-2 VOC/I by ASIL based on the SFM and Constellations (Version S.3 26-11-2025



REPORTING

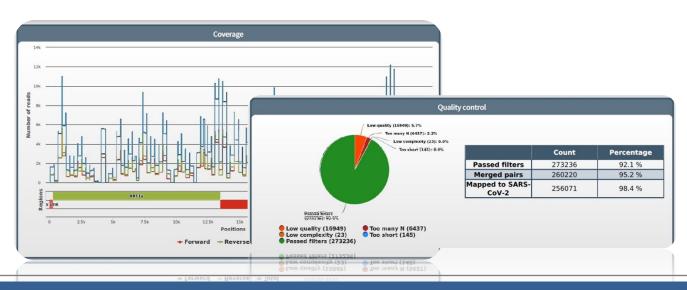


DEEPCHEK® CoV-2 Software

SARS-COV-2 MUTATIONS DETECTION

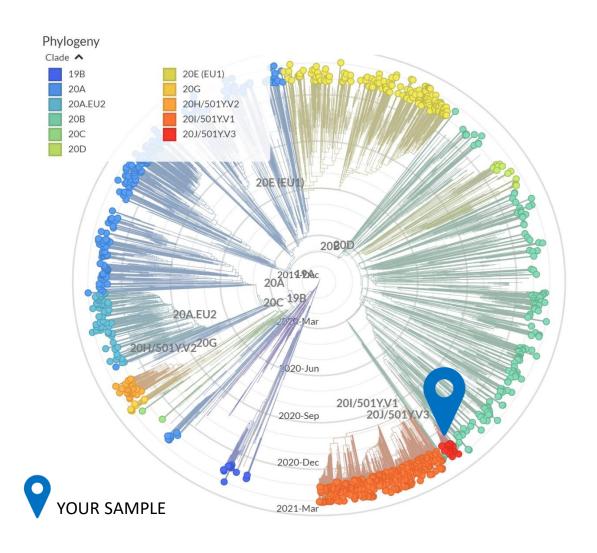
Region	Mutation	Frequency	Comments
	T265I	99.03 %	
	K1655N	98.95 %	
ORF1a	N2596S	98.93 %	
	K3353R	97.94 %	
	A3456V	99.54 %	
	S3675del3	99.36 %	
ORF1b	P314L	99.41 %	
	L18F	99.56 %	
	D80A	99.02 %	
	D215G	99.52 %	
	T240fs	98.73 %	
	L241del2	98.73 %	
	A243fs	98.73 %	
	K417N	98.76 %	
S	E484K	98.52 %	Escape to some mAbs Escape to some convalescent sera
	N501Y	98.52 %	Fast growing lineage and sits in the Recept Binding Motif (RBM) Has been described to increase binding affinity to hACE2 recepto
	D614G	98.93 %	Moderate effect on transmissibility (increasinfectivity)
	A701V	99.44 %	
ODE2-	Q57H	99.09 %	
ORF3a	S171L	100 %	
E	P71L	99.09 %	
N	T205I	99.57 %	

INTEGRATED EXPERT SYSTEM FOR QUALITY CONTROL



DEEPCHEK® CoV-2 Software

PHYLOGENETIC TREES THROUGH CONSENSUS SEQUENCES





Product References



ASSAYS

•	DeepChek Assay W	/hole Genome SARS	-CoV-2 Genotyping V1	(48 tests) (CE IVD)	159A48	C € IVD
---	------------------	-------------------	----------------------	---------------------	--------	---------

■ DeepChek Assay Whole Genome SARS-CoV-2 Genotyping V2 (48 tests) (RUO). . . . 159B48

SEQUENCING REAGENTS

-	DeepChek® NGS LIB PREP (95 indexes)	116B96 + 124B96
-	DeepChek® NGS LIB PREP (48 indexes)	116B48 + 124B48
•	DeepChek® NGS LIB PREP (24 indexes)	116B24 + 124B24

SOFTWARE

•	DeepChek® CoV-2 Software	S-12-023 ((CVL)	
---	--------------------------	------------	-------	--

■ DeepChek® - CoV2 LargeScale Premium Module...... S-12-023 (CVS)



Laboratories

https://www.ablsa.com

TEL: (+352) 26 38 96 761 FAX: (+352) 26 38 96 76 31

