

VIASURE CYSTIC FIBROSIS

NGS Solution

VS-CFD0148PHRUO*

For any lab within a working day.

VIASURE's solution for targeted amplicon-based gene sequencing offers a simplified workflow and exceptionally short assay time for detection of SNP and indels in the CFTR gene.

Comprehensive coverage of CF-causing SNVs and indels.

Variant filtering allows for flexible result analysis.

Dedicated and user friendly software for data analysis.

This minimizes the technical bioinformatics expertise required

Hands-on time

~40min

Total time required

<5h

Uniformity of coverage

100%

AMPLIFY YOUR RESEARCH to excellence



and userfriendliness.



Optimized, streamlined bioinformatic pipeline.



workflow.

Reduce the risk for sample mix up and contamination

Everything pre-dispensed and lyophilized





Modification rights reserved. All rights reserved. © Certest Biotec, S.L. The products, services and data set out in this document may suffer changes and/or variations on the texts and pictures shown.









