

Revised Technical Product Report

For Research Use Only; Not for use in Diagnostic Procedures

Product Description: Seraseq® ctDNA Complete Mutation Mix AF1%

Material Number: 0710-0530 Batch Number: 10670286

Material Description: A ctDNA-like mixture of human genomic DNA from the reference cell line, GM24385, and synthetic DNA constructs

Concentration
(Qubit dsDNA BR Assay): Nominal value: 10 ng/μL; Average measured value: 15 ng/μL

Fill Volume: 25 μL

Date of Manufacture: 16 JUN 2023 Expiration Date: 16 JUN 2025

Storage: -20°C

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Digital PCR testing
using BioRad
QX200™ Droplet
Digital™ PCR
System:

Gene ID	COSMIC Identifier	Amino Acid Change	Average AF%
AKT1	COSM33765	p.E17K	0.97
BRAF	COSM476	p.V600E	1.00
EGFR	COSM6224	p.L858R	1.06
EGFR	COSM6240	p.T790M	1.00
ERBB2	COSM20959	p.A775_G776insYVMA	0.55
KIT	COSM1314	p.D816V	1.03
KRAS	COSM521	p.G12D	1.10
NCOA4/RET	NA	Translocation	0.97
NRAS	COSM584	p.Q61R	1.10
PIK3CA	COSM775	p.H1047R	1.09
PIK3CA	COSM12464 ¹	p.N1068fs*4	1.09
EML4-ALK	NA	Translocation	0.98
ALK	COSM144250	p.G1202R	0.95
ALK	COSM28055	p.F1174L	0.95
BRCA1	COSM1383519	p.K654fs*47	0.90
BRCA2	COSM1738242	p.R2645fs*3	1.01
EGFR	COSM12370	p.L747_P753>S	1.23
EGFR	COSM6256	p.S752_I759delSPKANKEI	1.09
EGFR	COSM6223	p.E746_A750delELREA	1.06
KRAS	COSM516	p.G12C	1.00
CD74/ROS1	NA	Translocation	0.96
KRAS	COSM554	p.Q61H	1.06

Gene ID	Average CNV in ctDNA ²	Average Additional Copies (per cell) in ctDNA
ERBB2	2.92	0.92
MET	2.82	0.82
MYC	2.89	0.89

NA = not applicable

¹As of June 2019, this mutation is no longer listed in the COSMIC database.

²Compare to a normal CN of 2.00.

Next Generation Sequencing testing using Archer® Reveal ctDNA™ 28 Kit run on an Illumina® MiSeq™ using v2 (2x150 bp) PE chemistry reagents¹:

Gene ID	COSMIC Identifier	Amino Acid Change	AF%
AKT1	COSM33765	p.E17K	1.06
BRAF	COSM476	p.V600E	1.01
EGFR	COSM6224	p.L858R	0.62
EGFR	COSM6240	p.T790M	1.1
ERBB2	COSM20959	p.A775_G776insYVMA	0.34
KIT	COSM1314	p.D816V	1.13
KRAS	COSM521	p.G12D	1.13
NCOA4/RET	NA	Translocation	NA
NRAS	COSM584	p.Q61R	0.75
PIK3CA	COSM775	p.H1047R	0.80
PIK3CA	COSM12464 ³	p.N1068fs*4	0.67
EML4-ALK	NA	Translocation	NA
ALK	COSM144250	p.G1202R	0.80
ALK	COSM28055	p.F1174L	0.91
BRCA1	COSM1383519	p.K654fs*47	NA
BRCA2	COSM1738242	p.R2645fs*3	NA
EGFR	COSM12370	p.L747_P753>S	0.88
EGFR	COSM6256	p.S752_I759delSPKANKEI	0.85
EGFR	COSM6223	p.E746_A750delELREA	0.74
KRAS	COSM516	p.G12C	0.63
CD74/ROS1	NA	Translocation	NA
KRAS	COSM554	p.Q61H	0.73

Gene ID	CNV in ctDNA ⁴	Additional Copies (per cell) in ctDNA
ERBB2	3.06	1.06
MET	3.04	1.04
MYC	NA	NA

NA = not applicable; AF% and CNV marked NA were not targeted by the panel.

¹NGS was performed as an orthogonal verification step. Parameters used:

DNA input = 50 ng
 # of samples / flow cell = 3
 # of total reads / sample = 5.7 M
 Average read depth = 8740
 On-target reads = 96.2%
 Q30 score = 85.1%
 Analysis = Archer Analysis Suite v6.2.7 (default settings)

²Please see the poster from NIST for more information about assay sensitivity:

<https://digital.seracare.com/multilab-assessment-reference-materials-ctdna-poster2018>

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Note: The MET gene is amplified using two synthetic constructs with a small region of overlap between the constructs (see package insert for genomic coordinates). Assays which target this region of overlap may report higher amplification levels.

Approval:

BSL

04/03/2024

Prepared By

Date