

Technical Product Report

For Research Use Only; Not for use in Diagnostic Procedures

Product Description: Seraseq® ctDNA Complete Reference Material WT
 Material Number: 0710-0674 Batch Number: 10612741
 Material Description: A ctDNA-like mixture of human genomic DNA from the reference cell line, GM24385
 Concentration (Qubit dsDNA BR Assay): Nominal value: 25 ng/mL; Average measured value after extraction using Qiagen QIAamp Circulating Nucleic Acid Kit: 30.5 ng/mL
 Fill Volume: 5.0 mL
 Date of Manufacture: 06 APR 2022 Expiration Date: 06 APR 2025
 Storage: 2-8°C

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

Gene ID	COSMIC Identifier	Amino Acid Change	Average AF% ¹
AKT1	COSM33765	p.E17K	0.000
BRAF	COSM476	p.V600E	0.000
EGFR	COSM6224	p.L858R	0.000
EGFR	COSM6240	p.T790M	0.000
ERBB2	COSM20959	p.A775_G776insYVMA	0.000
KIT	COSM1314	p.D816V	0.000
KRAS	COSM521	p.G12D	0.000
NCOA4/RET	NA	Translocation	0.000
NRAS	COSM584	p.Q61R	0.000
PIK3CA	COSM775	p.H1047R	0.000
PIK3CA	COSM12464 ²	p.N1068fs*4	0.000
EML4-ALK	NA	Translocation	0.000
ALK	COSM144250	p.G1202R	0.000
ALK	COSM28055	p.F1174L	0.000
BRCA1	COSM1383519	p.K654fs*47	0.000
BRCA2	COSM1738242	p.R2645fs*3	0.000
EGFR	COSM12370	p.L747_P753>S	0.000
EGFR	COSM6256	p.S752_I759delISPKANKEI	0.000
EGFR	COSM6223	p.E746_A750delELREA	0.000
KRAS	COSM516	p.G12C	0.000
CD74/ROS1	NA	Translocation	0.000
KRAS	COSM554	p.Q61H	0.000

NA = not applicable

¹Variant allele frequencies > 0.00% for this wild-type negative control are within the expected range for stochastic positive dPCR reactions.

²As of June 2019, this mutation is no longer listed in the COSMIC database.

Technical Product Report

For Research Use Only; Not for use in Diagnostic Procedures

<p>Product Description:</p> <p>Material Number:</p> <p>Next Generation Sequencing testing using Archer® Reveal ctDNA™ 28 Kit run on an Illumina® MiSeq™ using v2 (2x150 bp) PE chemistry reagents¹</p> <p>=</p>	<p>SeraSeq® ctDNA Complete Reference Material WT</p> <p>0710-0674 Batch Number: 10612741</p> <p>NGS was performed as an orthogonal verification step. Results confirm no variants were detected above 0.1%.</p> <p><u>NGS Parameters:</u> DNA input = 50ng # of samples / flow cell = 1 # of total reads / sample = 14.6M Average read depth = 7500X On-target reads = 96% Q30 score = 91% Analysis = Archer Analysis Suite v6.2.2 (default settings except for: default settings were used.)</p>
--	---

¹Please see the poster from NIST for more information about assay sensitivity:
<https://digital.seracare.com/multilab-assessment-reference-materials-ctdna-poster2018>

NOTE: Copy numbers of ERBB2 and MET were assayed by dPCR and found to be normal. They were not assayed by NGS as the wild-type sample is used as a normal control for determining copy number of genes in other samples.

Approval:



09 MAY 2022

Prepared By

Date