



### Clinical Test Report : Oncomine™ Dx Target Test US v10.0

Sample ID: ADF10\_PRZ\_Run41\_Sample7\_20211117165500\_ET3JO462 Date Of Birth: 01 AUG 1980

Date: 13 DEC 2021

#### Draft Report

### Sample Details

Cancer Type:	<b>Non-small Cell Lung Cancer</b>	Ordering Physician:	<b>Dr. Jane Smith</b>	Sample Type:	<b>FFPE,Block</b>
Patient ID:	<b>Steve Snow</b>	%Necrosis:		Sample ID:	<b>ADF10_PRZ_Run41_Sample7_20211117165500_ET3JO462</b>
Gender:	<b>Male</b>	%Cellularity:		Collection Date:	<b>01 JUL 2016</b>
Date Of Birth:	<b>01 AUG 1980</b>	Reference Interval:		Created On:	<b>17 NOV 2021 16:55</b>
Sample Condition:	<b>Moderate</b>			Sample Source:	<b>Tissue</b>

### Results for Sequence Variations for Therapeutic Use

#### Gene Fusions (RNA) for Therapeutic Use

Gene	Display Name	Test Result	Therapy
RET	RET Fusion	<b>ABSENT</b>	None Indicated
ROS1	ROS1 Fusion	<b>ABSENT</b>	None Indicated

#### DNA Sequence Variants for Therapeutic Use

Note: Results for positive variants are listed first to provide therapy information, followed by variants with no calls, followed by variants that were reported as negative results.

Gene	Display Name	Amino Acid Change	Nucleotide Change	Test Result	Hotspot ID	Therapy
EGFR	EGFR Exon 20 Insertion	p.Ala767_Ser768insSerValGly	c. 2308_2309insGCAGCG TGG	<b>POSITIVE</b>	COSM18429	EXKIVITY™ (mobocertinib), RYBRENT™ (amivantamab-vmjw)

Gene	Display Name	Amino Acid Change	Nucleotide Change	Test Result	Hotspot ID	Therapy
BRAF	BRAF V600E	p.Val600Glu	c.1799T>A	<b>NEGATIVE</b>	COSM476	None Indicated
BRAF	BRAF V600E	p.Val600Glu	c.1799_1800delTGinsAA	<b>NEGATIVE</b>	COSM475	None Indicated
EGFR	EGFR Exon 19 deletion	p.Glu746_Ala750del	c.2235_2249delGGAATT AAGAGAAGC	<b>NEGATIVE</b>	COSM6223	None Indicated
EGFR	EGFR Exon 19 deletion	p.Glu746_Ala750del	c.2236_2250delGAATTA AGAGAAGCA	<b>NEGATIVE</b>	COSM6225	None Indicated
EGFR	EGFR Exon 19 deletion	p.Glu746_Arg748del	c.2239_2247delTTAAGA GAA	<b>NEGATIVE</b>	COSM6218	None Indicated
EGFR	EGFR Exon 19 deletion	p.Glu746_Glu749del	c.2235_2246delGGAATT AAGAGA	<b>NEGATIVE</b>	COSM28517	None Indicated
EGFR	EGFR Exon 19 deletion	p.Glu746_Ser752delinsAsp	c.2238_2255delAATTAAG AGAAGCAACATC	<b>NEGATIVE</b>	COSM6220	None Indicated
EGFR	EGFR Exon 19 deletion	p.Glu746_Ser752delinsVal	c.2237_2255delAATTAAGAGAAGCAACATCinsT	<b>NEGATIVE</b>	COSM12384	None Indicated
EGFR	EGFR Exon 19 deletion	p.Glu746_Thr751del	c.2236_2253delGAATTA AGAGAAGCAACA	<b>NEGATIVE</b>	COSM12728	None Indicated
EGFR	EGFR Exon 19 deletion	p.Glu746_Thr751delinsAla	c.2237_2251delAATTAAGAGAAGCAA	<b>NEGATIVE</b>	COSM12678	None Indicated

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EGFR	EGFR Exon 19 deletion	p.Glu746_Thr751delinsIle	c.2235_2252delGGAATT AAGAGAAGCAACinsA AT	NEGATIVE	COSM13551	None Indicated
EGFR	EGFR Exon 19 deletion	p.Glu746_Thr751delinsValAla	c.2237_2253delAATTAA GAGAAGCAACinsTTG CT	NEGATIVE	COSM12416	None Indicated
EGFR	EGFR Exon 19 deletion	p.Leu747_Ala750delinsPro	c.2239_2248delTTAAGA GAAGinsC	NEGATIVE	COSM12382	None Indicated
EGFR	EGFR Exon 19 deletion	p.Leu747_Ala750delinsPro	c.2238_2248delATTAAG AGAAGinsGC	NEGATIVE	COSM12422	None Indicated
EGFR	EGFR Exon 19 deletion	p.Leu747_Pro753delinsGln	c.2239_2258delTTAAGA GAAGCAACATCTCins CA	NEGATIVE	COSM12387	None Indicated
EGFR	EGFR Exon 19 deletion	p.Leu747_Pro753delinsSer	c.2240_2257delTTAAGA GAAGCAACATCTC	NEGATIVE	COSM12370	None Indicated
EGFR	EGFR Exon 19 deletion	p.Leu747_Ser752del	c.2239_2256delTTAAGA GAAGCAACATCT	NEGATIVE	COSM6255	None Indicated
EGFR	EGFR Exon 19 deletion	p.Leu747_Thr751del	c.2240_2254delTTAAGA GAAGCAACAT	NEGATIVE	COSM12369	None Indicated
EGFR	EGFR Exon 19 deletion	p.Leu747_Thr751delinsGln	c.2238_2252delATTAAG AGAAGCAACinsGCA	NEGATIVE	COSM12419	None Indicated
EGFR	EGFR Exon 19 deletion	p.Leu747_Thr751delinsPro	c.2239_2251delTTAAGA GAAGCAAinsC	NEGATIVE	COSM12383	None Indicated
EGFR	EGFR Exon 19 deletion	p.Leu747_Thr751delinsSer	c.2240_2251delTTAAGA GAAGCAA	NEGATIVE	COSM6210	None Indicated
EGFR	EGFR Exon 19 deletion	p.Lys745_Ala750delinsThr	c.2234_2248delAGGAA TTAAGAGAA	NEGATIVE	COSM1190791	None Indicated
EGFR	EGFR Exon 19 deletion	p.Lys745_Glu749del	c.2233_2247delAAGGA ATTAAGAGAA	NEGATIVE	COSM26038	None Indicated
EGFR	EGFR Exon 20 Insertion	NA	NA	NEGATIVE	COSM26720	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Ala767_Ser768insSerValAsp	c.2311_2312insGCGTGG ACA	NEGATIVE	COSM13428	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Ala767_Ser768insSerValGly	c.2308_2309insGGAGCG TGG	NEGATIVE	COSM1235344	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Ala767_Ser768insTyrValMet	c.2301_2302insTACGTG ATG	NEGATIVE	COSM1651740	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Ala767_Val769dup	c.2309_2310delACinsCC AGCGTGGAT	NEGATIVE	COSM13558	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771_Pro772insArgHis	c.2314_2315insGGCACC	NEGATIVE	COSM166390	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771_Pro772insHis	c.2314_2315insACC	NEGATIVE	COSM1238031	None Indicated

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EGFR	EGFR Exon 20 Insertion	p.Asn771_Pro772insHisHis	c.2314_2315insACCACC	NEGATIVE	COSM6931207	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771_Pro772insLeu	c.2314_2315insTCC	NEGATIVE	N771_P772insL	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771_Pro772insLeu	c.2313_2314insTTG	NEGATIVE	c.2313_2314insTTG	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771_Pro772insProHis	c.2319_2320insCCCCAC	NEGATIVE	COSM12380	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771_Pro772insProHisVal	c.2322_2323insCCCCACGTG	NEGATIVE	COSM6845098	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771_Pro772insProThrHis	c.2315_2316insGACACACCC	NEGATIVE	COSM48923	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771_Pro772insThr	c.2313_2314insACA	NEGATIVE	c.2313_2314insACA	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771_Pro772insVal	c.2313_2314insGTC	NEGATIVE	COSM6922328	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771delinsGlyPhe	c.2311_2312delAAinsGGTT	NEGATIVE	COSM18431	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771delinsGlyTyr	c.2311_2311delAinsGGTT	NEGATIVE	COSM53189	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771delinsLysGly	c.2313_2313delCinsGGGG	NEGATIVE	N771delinsKG	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771delinsLysHis	c.2312_2313insACA	NEGATIVE	OMINDEL1123	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771delinsLysLeu	c.2312_2313insACT	NEGATIVE	COSM6438147	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771delinsProHis	c.2311_2311delAinsCCC	NEGATIVE	MATNV04	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771delinsSerGlyHis	c.2311_2312insGTGGCC	NEGATIVE	COSM1651744	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771delinsSerHis	c.2311_2312insGTC	NEGATIVE	COSM24434	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771delinsSerThrHis	c.2311_2312insGCACCC	NEGATIVE	COSM6920147	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771delinsThrHis	c.2311_2312insCAC	NEGATIVE	COSM22946	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asn771delinsValHis	c.2311_2311delAinsGTC	NEGATIVE	COSM5023007	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771delinsAlaGlyGly	c.2309_2312delACAAinsCTGGTGG	NEGATIVE	COSM12737	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771delinsAlaGlyHis	c.2309_2311delACAinsCTGGCC	NEGATIVE	MATNV09	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insAlaProTrp	c.2310_2311insGCACCGTGG	NEGATIVE	COSM20886	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insAsn	c.2313_2314insAAC	NEGATIVE	COSM13003	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insAsnHis	c.2314_2315insACAACC	NEGATIVE	OMINDEL1084	None Indicated

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Gene	Display Name	Amino Acid Change	Nucleotide Change	Test Result	Hotspot ID	Therapy
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insAsnPro	c.2316_2317insAACCCC	NEGATIVE	MAN123	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insAsnProGly	c.2316_2317insGGCAACCCC	NEGATIVE	c.2316_2317insGGCAACCCC	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insAsnProGly	c.2316_2317insGGAAACCCC	NEGATIVE	P772_H773insGNP	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insAsnProHis	c.2319_2320insAACCCCAC	NEGATIVE	COSM12381	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insAsnProHisGly	c.2320_2321insGCAACCCCACG	NEGATIVE	COSM51544	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insAsnProPro	c.2317_2318insCCAACCCC	NEGATIVE	P772_H773insPNP	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insAsnProThrPro	c.2316_2317insACACCCAACCCC	NEGATIVE	COSM6977296	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insGlnArgGly	c.2310_2311insCAGCGTGGC	NEGATIVE	COSM4970107	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insGly	c.2310_2311insGGC	NEGATIVE	COSM13004	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insGly	c.2310_2311insGGG	NEGATIVE	MATNV05	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insGly	c.2310_2311insGGT	NEGATIVE	COSM12378	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insGlyLeu	c.2310_2311insGGGTTA	NEGATIVE	COSM48921	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insGlyPhe	c.2310_2311insGGGTTT	NEGATIVE	COSM655155	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insGlyThr	c.2310_2311insGGCACA	NEGATIVE	COSM1238029	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insHis	c.2310_2311insCAC	NEGATIVE	OMINDEL1081	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insMetAlaThrPro	c.2311_2312insTGGCCACCCCA	NEGATIVE	COSM26719	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insSerValGlu	c.2311_2312insGCGTCGAAA	NEGATIVE	COSM1651743	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insThr	c.2311_2312insCCA	NEGATIVE	COSM5023008	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770_Asn771insTyr	c.2310_2311insTAC	NEGATIVE	COSM1238030	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770delinsAsnAsnProHis	c.2308_2308delGinsAACACCCCC	NEGATIVE	OMINDEL1078	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770delinsGlyThrHis	c.2308_2309insGCACAC	NEGATIVE	COSM6983510	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Asp770delinsGlyTyr	c.2308_2309insGTT	NEGATIVE	COSM12427	None Indicated
EGFR	EGFR Exon 20 Insertion	p.His773_Val774insGln	c.2319_2320insCAG	NEGATIVE	COSM131552	None Indicated
EGFR	EGFR Exon 20 Insertion	p.His773_Val774insThrGlnProPro	c.2319_2320insACACAAACCC	NEGATIVE	COSM3727813	None Indicated

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EGFR	EGFR Exon 20 Insertion	p.His773delinsArgTyr	c.2317_2318insGTT	NEGATIVE	MATNV08	None Indicated
EGFR	EGFR Exon 20 Insertion	p.His773delinsAsnProTyr	c.2317_2317delCinsAACCCCT	NEGATIVE	H773delinsNPY	None Indicated
EGFR	EGFR Exon 20 Insertion	p.His773delinsProAsnProTyr	c.2317_2318insCTAACCCT	NEGATIVE	COSM1735761	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Met766_Ala767insAlalle	c.2302_2303insTAGCCA	NEGATIVE	COSM13559	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Met766_Ala767insAlaSerVal	c.2308_2309insCCAGCGTGG	NEGATIVE	COSM12376	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Met766_Ala767insAlaSerVal	c.2303_2304insTGTGGCCAG	NEGATIVE	COSM20884	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Met766_Ala767insAlaThrLeu	c.2302_2303insCGCTGGCCA	NEGATIVE	COSM12425	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Pro772_His773insGlyThr	c.2316_2317insGGCACC	NEGATIVE	MATNV02	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Pro772_His773insHis	c.2319_2320insCAC	NEGATIVE	COSM12377	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Pro772_His773insHisAla	c.2320_2321insCCCACG	NEGATIVE	COSM1238028	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Pro772_His773insHisAsn	c.2319_2320insAACCCAC	NEGATIVE	COSM5023006	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Pro772_His773insHisVal	c.2321_2322insCCACGT	NEGATIVE	COSM18432	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Pro772_His773insHisVal	c.2322_2323insCACGTG	NEGATIVE	COSM22948	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Pro772_His773insThrPro	c.2316_2316delCinsAACCCCT	NEGATIVE	COSM12388	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Pro772_His773insVal	c.2316_2317insGTT	NEGATIVE	COSM255205	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Ser768_Val769delinsIleLeu	c.2303_2305delGCGinsTCC	NEGATIVE	COSM6984779	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Ser768_Val769delinsIleLeu	c.2303_2305delGCGinsTCT	NEGATIVE	COSM85750	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Ser768_Val769insVal	c.2308_2309insTGG	NEGATIVE	COSM6506514	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Ser768_Val769insValAlaAsn	c.2303_2304insTGTGGCCAA	NEGATIVE	COSM1651741	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Ser768_Val769insValAspAsn	c.2313_2314insGTGGACAAC	NEGATIVE	COSM20885	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Ser768_Val769insValAspAsn Pro	c.2316_2317insGTGGACAACCCC	NEGATIVE	V769_P772dup	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Ser768_Val769insValCys	c.2307_2308insTGCGTG	NEGATIVE	COSM12379	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Ser768_Val769insValGlyVal	c.2308_2309insGGGTCTGG	NEGATIVE	COSM18430	None Indicated

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EGFR	EGFR Exon 20 Insertion	p.Val769_Asp770insAspAsnPro	c.2315_2316insGGACAA CCC	<b>NEGATIVE</b>	COSM6845099	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Val769_Asp770insAspAsnPro	c.2316_2317insGACAAC CCC	<b>NEGATIVE</b>	COSM1651745	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Val769_Asp770insAspGly	c.2310_2311insGGGGAC	<b>NEGATIVE</b>	COSM85795	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Val769_Asp770insAspGly	c.2310_2311insGGCGAC	<b>NEGATIVE</b>	COSM22955	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Val769_Asp770insAspLys	c.2312_2313insGGACAA	<b>NEGATIVE</b>	D770_N771insKD	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Val769_Asp770insGluArgGly	c.2309_2310insGCGTGG AGA	<b>NEGATIVE</b>	COSM1651742	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Val769_Asp770insGlyGly	c.2308_2309insGGGGGG	<b>NEGATIVE</b>	MATNV07	None Indicated
EGFR	EGFR Exon 20 Insertion	p.Val774_Cys775insProArg	c.2322_2323insCCACGT	<b>NEGATIVE</b>	COSM4170223	None Indicated
EGFR	EGFR L858R	p.Leu858Arg	c.2573T>G	<b>NEGATIVE</b>	COSM6224	None Indicated

The following reference files are used:hg19, Oncomine™ Dx Target Panel US v10.0 Fusion Reference. Associated therapy is within Non-Small Cell Lung Carcinoma (NSCLC).

For additional questions on "no call" results please contact testing service laboratory.

Example Report



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### Results for Analytical Sequence Variations Detected

#### Analytical DNA Sequence Variants Detected

Gene	Amino AcidChange	NucleotideChange	Test Result	Hotspot ID
PIK3CA	p.Arg93Gln	c.278G>A	POSITIVE	COSM86041
ERBB2	p.Arg678Gln	c.2033G>A	POSITIVE	COSM436498

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Lab Director: Max Smith CLIA number: 03C1021009

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Example Report



**Clinical Test Report : Oncomine™ Dx Target Test US v10.0**

Sample ID: ADF10\_PRZ\_Run41\_Sample7\_20211117165500\_ET3J0462 Date Of Birth: 01 AUG 1980

Date: 13 DEC 2021

**Draft Report**
**Intended Use**

The Oncomine™ Dx Target Test is a qualitative in vitro diagnostic test that uses targeted high throughput, parallel-sequencing technology to detect single nucleotide variants (SNVs), deletions, and insertions in 23 genes from DNA and fusions in ROS1 and RET from RNA isolated from formalin-fixed, paraffin-embedded (FFPE) tumor tissue samples from patients with non-small cell lung cancer (NSCLC), and IDH1 SNVs from FFPE tumor tissue samples from patients with cholangiocarcinoma (CC) using the Ion PGM™ Dx System.

The test is indicated to aid in selecting NSCLC and CC patients for treatment with the targeted therapies listed in Table 1 in accordance with the approved therapeutic product labeling.

Table 1. List of Variants for Therapeutic Use

Tissue Type	Gene	Variant	Targeted Therapy
Non-small Cell Lung Cancer (NSCLC)	BRAF	BRAF V600E mutation	TAFINLAR® (dabrafenib) in combination with MEKINIST® (trametinib)
	EGFR	EGFR L858R mutation, EGFR Exon 19 deletions	IRESSA® (gefitinib)
	EGFR	EGFR Exon 20 Insertions	EXKIVITY™ (mobocertinib), RYBREVANT™ (amivantamab-vmjw)
	RET	RET fusions	GAVRETO™ (pralsetinib)
	ROS1	ROS1 fusions	XALKORI® (crizotinib)
Cholangiocarcinoma (CC)	IDH1	IDH1 R132C, IDH1 R132G, IDH1 R132H, IDH1 R132L, IDH1 R132S	TIBSOVO® (ivosidenib)

Safe and effective use has not been established for selecting therapies using this device for the variants other than those in Table 1.

Results other than those listed in Table 1 are indicated for use only in patients who have already been considered for all appropriate therapies (including those listed in Table 1). Analytical performance using NSCLC specimens has been established for the variants listed in Table 2.

Table 2. List of Variants with Established Analytical Performance Only

Gene	Variant ID/ Type	Amino Acid Change	Nucleotide Change
KRAS	COSM512	p.Gly12Phe	c.34_35delGGinsTT
KRAS	COSM516	p.Gly12Cys	c.34G>T
MET	COSM707	p.Thr1010Ile	c.3029C>T
PIK3CA	COSM754	p.Asn345Lys	c.1035T>A

The test is not indicated to be used for standalone diagnostic purposes, screening, monitoring, risk assessment, or prognosis.

**Warnings and Precautions**

For NSCLC, the Oncomine Dx Target Test assay definition file includes prevalent but not all rare or newly identified RET isoforms, ROS1 isoforms, and EGFR exon 20 insertions. The Oncomine Dx Target Test may miss rare or newly identified:

- RET isoforms carried by a subset of patients who may derive benefit from GAVRETO™ (pralsetinib)
- ROS1 isoforms carried by a subset of patients who may derive benefit from XALKORI® (crizotinib)
- EGFR exon 20 insertions carried by a subset of patients who may derive benefit from EXKIVITY™ (mobocertinib) or RYBREVANT™ (amivantamab-vmjw)

## Clinical Test Report : OncoPrint™ Dx Target Test US v10.0

Sample ID: ADF10\_PRZ\_Run41\_Sample7\_20211117165500\_ET3J0462 Date Of Birth: 01 AUG 1980

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## Limitations

- The OncoPrint™ Dx Target Test has only been validated for use with FFPE tumor slide specimens.
- DNA and/or RNA from a single sample extraction must meet the concentration requirements specified in the procedure. Do not use DNA from one extraction with RNA from a different extraction.
- The effects of potential variations in FFPE specimen fixation have not been evaluated.
- Extraction from FFPE sample curls has not been evaluated.
- A potential source of contamination in the procedure is nucleic acid from previous sample processing steps. Follow good laboratory practices and all precautions and guidelines in these user guides to avoid cross-contamination between samples.
- The OncoPrint™ Dx Target Test is a qualitative test. The test is not for quantitative measurements of percent mutation.
- The safe and effective use of the variants reported in the Analytical Sequence Variations Detected section has not been established for selecting therapy using this device. The variants for KRAS (COSM512/p.Gly12Phe/c.34\_35delGGinsTT and COSM516/p.Gly12Cys/c.34G>T), MET (COSM707/p.Thr1010Ile/c.3029C>T) and PIK3CA (COSM754/p.Asn345Lys/c.1035T>A) have been analytically validated. Performance of all other variants identified by the test, other than clinically validated therapeutic variants and analytically validated variants, has not been directly demonstrated.

## Test Description

The OncoPrint™ Dx Target Test reports the following genes in each indication:

Non-small cell lung cancer (NSCLC):

- Single nucleotide variants (SNVs), deletions, and insertions from DNA: AKT1, ALK, BRAF, CDK4, DDR2, EGFR, ERBB2, ERBB3, FGFR2, FGFR3, HRAS, KIT, KRAS, MAP2K1, MAP2K2, MET, MTOR, NRAS, PDGFRA, PIK3CA, RAF1, RET and ROS1.
- Fusions from RNA: RET and ROS1.

Cholangiocarcinoma (CC):

- Single nucleotide variants (SNVs) from DNA: IDH1.

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**Results for Analytical Sequence Variations Not Detected**

## Analytical DNA Sequence Variants Not Detected

Note: Results for no calls are listed first, followed by variants that were reported as negative results.

Gene	Amino Acid Change	Nucleotide Change	Test Result	Hotspot ID
ALK	p.Leu1152Arg	c.3455T>G	NO CALL	COSM97185
ALK	p.Leu1152Pro	c.3455T>C	NO CALL	COSM1407659
FGFR3	p.Arg248Cys	c.742C>T	NO CALL	COSM714

Gene	Amino Acid Change	Nucleotide Change	Test Result	Hotspot ID
AKT1	p.Glu17Lys	c.49G>A	NEGATIVE	COSM33765
ALK	p.Arg1275Gln	c.3824G>A	NEGATIVE	COSM28056
ALK	p.Arg1275Leu	c.3824G>T	NEGATIVE	COSM28060
ALK	p.Cys1156Tyr	c.3467G>A	NEGATIVE	COSM99136
ALK	p.Gly1128Ala	c.3383G>C	NEGATIVE	COSM98475
ALK	p.Gly1202Arg	c.3604G>A	NEGATIVE	COSM144250
ALK	p.Ile1171Asn	c.3512T>A	NEGATIVE	COSM28498
ALK	p.Ile1171Thr	c.3512T>C	NEGATIVE	COSM4381100
ALK	p.Leu1196Gln	c.3587T>A	NEGATIVE	COSM1169447
ALK	p.Leu1196Met	c.3586C>A	NEGATIVE	COSM99137
ALK	p.Phe1174Cys	c.3521T>G	NEGATIVE	COSM28059
ALK	p.Phe1174Ile	c.3520T>A	NEGATIVE	COSM28491
ALK	p.Phe1174Leu	c.3522C>G	NEGATIVE	COSM28061
ALK	p.Phe1174Leu	c.3522C>A	NEGATIVE	COSM28055
ALK	p.Phe1174Leu	c.3520T>C	NEGATIVE	COSM28057
ALK	p.Phe1174Ser	c.3521T>C	NEGATIVE	COSM53063
ALK	p.Phe1174Val	c.3520T>G	NEGATIVE	COSM28054
ALK	p.Phe1245Cys	c.3734T>G	NEGATIVE	COSM28500
ALK	p.Phe1245Ile	c.3733T>A	NEGATIVE	COSM28492
ALK	p.Phe1245Leu	c.3735C>G	NEGATIVE	COSM28062
ALK	p.Phe1245Leu	c.3735C>A	NEGATIVE	COSM28493
ALK	p.Phe1245Val	c.3733T>G	NEGATIVE	COSM28499
ALK	p.Ser1206Tyr	c.3617C>A	NEGATIVE	COSM144251
ALK	p.Val1180Leu	c.3538G>C	NEGATIVE	COSM4381101
BRAF	p.Asp594Asn	c.1780G>A	NEGATIVE	COSM27639
BRAF	p.Asp594Gly	c.1781A>G	NEGATIVE	COSM467
BRAF	p.Gly466Glu	c.1397G>A	NEGATIVE	COSM453
BRAF	p.Gly466Val	c.1397G>T	NEGATIVE	COSM451
BRAF	p.Gly469Ala	c.1406G>C	NEGATIVE	COSM460
BRAF	p.Gly469Arg	c.1405G>A	NEGATIVE	COSM457
BRAF	p.Gly469Val	c.1406G>T	NEGATIVE	COSM459
BRAF	p.Lys601Glu	c.1801A>G	NEGATIVE	COSM478
BRAF	p.Val600_Lys601delinsGlu	c.1799_1801delTGA	NEGATIVE	COSM1133
BRAF	p.Val600Arg	c.1798_1799delGTinsAG	NEGATIVE	COSM474
BRAF	p.Val600Lys	c.1798_1799delGTinsAA	NEGATIVE	COSM473
CDK4	p.Arg24Cys	c.70C>T	NEGATIVE	COSM1677139
CDK4	p.Arg24His	c.71G>A	NEGATIVE	COSM1989836

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CDK4	p.Arg24Leu	c.71G>T	NEGATIVE	COSM363684
CDK4	p.Arg24Ser	c.70C>A	NEGATIVE	COSM3463914
CDK4	p.Lys22Arg	c.65A>G	NEGATIVE	COSM232013
CDK4	p.Lys22Gln	c.64A>C	NEGATIVE	OM3153
CDK4	p.Lys22Met	c.65A>T	NEGATIVE	COSM3463915
DDR2	p.Arg124Leu	c.371G>T	NEGATIVE	COSM400880
DDR2	p.Arg124Trp	c.370C>T	NEGATIVE	COSM4024594
EGFR	p.Ala289Asp	c.866C>A	NEGATIVE	COSM21685
EGFR	p.Ala289Thr	c.865G>A	NEGATIVE	COSM21686
EGFR	p.Ala289Val	c.866C>T	NEGATIVE	COSM21687
EGFR	p.Arg108Gly	c.322A>G	NEGATIVE	COSM1451536
EGFR	p.Arg108Lys	c.323G>A	NEGATIVE	COSM21683
EGFR	p.Cys797Ser	c.2389T>A	NEGATIVE	COSM6493937
EGFR	p.Cys797Ser	c.2390G>C	NEGATIVE	COSM5945664
EGFR	p.Glu709Ala	c.2126A>C	NEGATIVE	COSM13427
EGFR	p.Glu709Gly	c.2126A>G	NEGATIVE	COSM13009
EGFR	p.Glu709Lys	c.2125G>A	NEGATIVE	COSM12988
EGFR	p.Glu709Val	c.2126A>T	NEGATIVE	COSM12371
EGFR	p.Gly598Ala	c.1793G>C	NEGATIVE	COSM3412196
EGFR	p.Gly598Val	c.1793G>T	NEGATIVE	COSM21690
EGFR	p.Gly719Ala	c.2156G>C	NEGATIVE	COSM6239
EGFR	p.Gly719Asp	c.2156G>A	NEGATIVE	COSM18425
EGFR	p.Gly719Cys	c.2155G>T	NEGATIVE	COSM6253
EGFR	p.Gly719Ser	c.2155G>A	NEGATIVE	COSM6252
EGFR	p.Leu858Met	c.2572C>A	NEGATIVE	COSM12366
EGFR	p.Leu861Arg	c.2582T>G	NEGATIVE	COSM12374
EGFR	p.Leu861Gln	c.2582T>A	NEGATIVE	COSM6213
EGFR	p.Ser492Arg	c.1474A>C	NEGATIVE	COSM236671
EGFR	p.Ser492Arg	c.1476C>A	NEGATIVE	COSM236670
EGFR	p.Ser768Ile	c.2303G>T	NEGATIVE	COSM6241
ERBB2	p.Arg896Cys	c.2686C>T	NEGATIVE	COSM14066
ERBB2	p.Arg896His	c.2687G>A	NEGATIVE	COSM119971
ERBB2	p.Asp769His	c.2305G>C	NEGATIVE	COSM13170
ERBB2	p.Asp769Tyr	c.2305G>T	NEGATIVE	COSM1251412
ERBB2	p.Gly776Val	c.2327G>T	NEGATIVE	COSM18609
ERBB2	p.Leu755Met	c.2263T>A	NEGATIVE	COSM1205571
ERBB2	p.Leu755Pro	c.2263_2264delTTinsCC	NEGATIVE	COSM683
ERBB2	p.Ser310Phe	c.929C>T	NEGATIVE	COSM48358
ERBB2	p.Ser310Tyr	c.929C>A	NEGATIVE	COSM94225
ERBB2	p.Thr733Ile	c.2198C>T	NEGATIVE	COSM14059
ERBB2	p.Val777Leu	c.2329G>T	NEGATIVE	COSM14062
ERBB2	p.Val842Ile	c.2524G>A	NEGATIVE	COSM14065
ERBB3	p.Ala232Thr	c.694G>A	NEGATIVE	COSM4043440
ERBB3	p.Ala232Val	c.695C>T	NEGATIVE	COSM1242239
ERBB3	p.Asp297Tyr	c.889G>T	NEGATIVE	COSM160822
ERBB3	p.Asp297Val	c.890A>T	NEGATIVE	COSM941490
ERBB3	p.Glu332Lys	c.994G>A	NEGATIVE	COSM254677

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ERBB3	p.Met60Arg	c.179T>G	NEGATIVE	COSM941484
ERBB3	p.Met60Leu	c.178A>T	NEGATIVE	COSM1606366
ERBB3	p.Met60Lys	c.179T>A	NEGATIVE	COSM254678
ERBB3	p.Met91Ile	c.273G>A	NEGATIVE	COSM122890
ERBB3	p.Met91Ile	c.273G>C	NEGATIVE	COSM1299636
ERBB3	p.Val104Leu	c.310G>C	NEGATIVE	COSM160824
ERBB3	p.Val104Leu	c.310G>T	NEGATIVE	COSM191840
ERBB3	p.Val104Met	c.310G>A	NEGATIVE	COSM172423
FGFR2	p.Ala314Asp	c.941C>A	NEGATIVE	COSM49171
FGFR2	p.Asn549His	c.1645A>C	NEGATIVE	COSM250083
FGFR2	p.Asn549Lys	c.1647T>G	NEGATIVE	COSM36902
FGFR2	p.Asn549Lys	c.1647T>A	NEGATIVE	COSM36912
FGFR2	p.Asn549Ser	c.1646A>G	NEGATIVE	COSM3665553
FGFR2	p.Cys382Arg	c.1144T>C	NEGATIVE	COSM36906
FGFR2	p.Cys382Tyr	c.1145G>A	NEGATIVE	COSM915493
FGFR2	p.Lys659Asn	c.1977G>T	NEGATIVE	COSM49173
FGFR2	p.Lys659Asn	c.1977G>C	NEGATIVE	COSM683054
FGFR2	p.Lys659Glu	c.1975A>G	NEGATIVE	COSM36909
FGFR2	p.Lys659Met	c.1976A>T	NEGATIVE	COSM49175
FGFR2	p.Pro253Arg	c.758C>G	NEGATIVE	COSM49170
FGFR2	p.Pro253Leu	c.758C>T	NEGATIVE	COSM537801
FGFR2	p.Ser252Trp	c.755C>G	NEGATIVE	COSM36903
FGFR2	p.Tyr375Cys	c.1124A>G	NEGATIVE	COSM36904
FGFR2	p.Tyr375His	c.1123T>C	NEGATIVE	COSM1560916
FGFR3	p.Gly697Cys	c.2089G>T	NEGATIVE	COSM24802
FGFR3	p.Lys650Asn	c.1950G>T	NEGATIVE	COSM1428730
FGFR3	p.Lys650Gln	c.1948A>C	NEGATIVE	COSM726
FGFR3	p.Lys650Glu	c.1948A>G	NEGATIVE	COSM719
FGFR3	p.Ser249Cys	c.746C>G	NEGATIVE	COSM715
HRAS	p.Gln61Arg	c.182A>G	NEGATIVE	COSM499
HRAS	p.Gln61His	c.183G>T	NEGATIVE	COSM502
HRAS	p.Gln61His	c.183G>C	NEGATIVE	COSM503
HRAS	p.Gln61Leu	c.182A>T	NEGATIVE	COSM498
HRAS	p.Gln61Lys	c.181C>A	NEGATIVE	COSM496
HRAS	p.Gln61Pro	c.182A>C	NEGATIVE	COSM500
HRAS	p.Gly12Ala	c.35G>C	NEGATIVE	COSM485
HRAS	p.Gly12Arg	c.34G>C	NEGATIVE	COSM482
HRAS	p.Gly12Asp	c.35G>A	NEGATIVE	COSM484
HRAS	p.Gly12Cys	c.34G>T	NEGATIVE	COSM481
HRAS	p.Gly12Ser	c.34G>A	NEGATIVE	COSM480
HRAS	p.Gly12Val	c.35G>T	NEGATIVE	COSM483
HRAS	p.Gly13Arg	c.37G>C	NEGATIVE	COSM486
HRAS	p.Gly13Asp	c.38G>A	NEGATIVE	COSM490
HRAS	p.Gly13Cys	c.37G>T	NEGATIVE	COSM488
HRAS	p.Gly13Ser	c.37G>A	NEGATIVE	COSM487
HRAS	p.Gly13Val	c.38G>T	NEGATIVE	COSM489
KIT	p.Arg796Lys	c.2387G>A	NEGATIVE	COSM1600411
KIT	p.Asn822Lys	c.2466T>A	NEGATIVE	COSM1321

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KIT	p.Asn822Lys	c.2466T>G	NEGATIVE	COSM1322
KIT	p.Asp419_Arg420del	c.1255_1260delGACAGG	NEGATIVE	COSM1578132
KIT	p.Asp419del	c.1255_1257delIGAC	NEGATIVE	COSM29014
KIT	p.Asp579del	c.1735_1737delIGAT	NEGATIVE	COSM1294
KIT	p.Asp816His	c.2446G>C	NEGATIVE	COSM1311
KIT	p.Asp816Tyr	c.2446G>T	NEGATIVE	COSM1310
KIT	p.Asp816Val	c.2447A>T	NEGATIVE	COSM1314
KIT	p.Leu576Pro	c.1727T>C	NEGATIVE	COSM1290
KIT	p.Lys642Glu	c.1924A>G	NEGATIVE	COSM1304
KIT	p.Trp557_Lys558del	c.1669_1674delTGGAAG	NEGATIVE	COSM1217
KIT	p.Trp557_Val559delinsPhe	c.1670_1675delGGAAGG	NEGATIVE	COSM1226
KIT	p.Trp557Arg	c.1669T>A	NEGATIVE	COSM1216
KIT	p.Trp557Arg	c.1669T>C	NEGATIVE	COSM1219
KIT	p.Trp557Gly	c.1669T>G	NEGATIVE	COSM1221
KIT	p.Val559Ala	c.1676T>C	NEGATIVE	COSM1255
KIT	p.Val559Asp	c.1676T>A	NEGATIVE	COSM1252
KIT	p.Val559del	c.1679_1681delTTG	NEGATIVE	COSM1247
KIT	p.Val559Gly	c.1676T>G	NEGATIVE	COSM1253
KIT	p.Val560Asp	c.1679T>A	NEGATIVE	COSM1257
KIT	p.Val654Ala	c.1961T>C	NEGATIVE	COSM12706
KIT	p.Val825Ala	c.2474T>C	NEGATIVE	COSM1323
KRAS	p.Ala146Pro	c.436G>C	NEGATIVE	COSM19905
KRAS	p.Ala146Thr	c.436G>A	NEGATIVE	COSM19404
KRAS	p.Ala146Val	c.437C>T	NEGATIVE	COSM19900
KRAS	p.Ala59Glu	c.176C>A	NEGATIVE	COSM547
KRAS	p.Ala59Gly	c.176C>G	NEGATIVE	COSM28518
KRAS	p.Ala59Thr	c.175G>A	NEGATIVE	COSM546
KRAS	p.Gln61Arg	c.182A>G	NEGATIVE	COSM552
KRAS	p.Gln61Glu	c.181C>G	NEGATIVE	COSM550
KRAS	p.Gln61His	c.183A>T	NEGATIVE	COSM555
KRAS	p.Gln61His	c.183A>C	NEGATIVE	COSM554
KRAS	p.Gln61Leu	c.182A>T	NEGATIVE	COSM553
KRAS	p.Gln61Lys	c.181C>A	NEGATIVE	COSM549
KRAS	p.Gln61Lys	c.180_181delTCinsAA	NEGATIVE	COSM87298
KRAS	p.Gln61Pro	c.182A>C	NEGATIVE	COSM551
KRAS	p.Gly12Ala	c.35G>C	NEGATIVE	COSM522
KRAS	p.Gly12Arg	c.34G>C	NEGATIVE	COSM518
KRAS	p.Gly12Asp	c.35G>A	NEGATIVE	COSM521
KRAS	p.Gly12Cys	c.34G>T	NEGATIVE	COSM516
KRAS	p.Gly12Phe	c.34_35delGGinsTT	NEGATIVE	COSM512
KRAS	p.Gly12Ser	c.34G>A	NEGATIVE	COSM517
KRAS	p.Gly12Val	c.35G>T	NEGATIVE	COSM520
KRAS	p.Gly13Ala	c.38G>C	NEGATIVE	COSM533
KRAS	p.Gly13Arg	c.37G>C	NEGATIVE	COSM529
KRAS	p.Gly13Asp	c.38_39delGCinsAT	NEGATIVE	COSM531
KRAS	p.Gly13Asp	c.38G>A	NEGATIVE	COSM532
KRAS	p.Gly13Cys	c.37G>T	NEGATIVE	COSM527
KRAS	p.Gly13Ser	c.37G>A	NEGATIVE	COSM528

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KRAS	p.Gly13Val	c.38G>T	NEGATIVE	COSM534
KRAS	p.Lys117Asn	c.351A>T	NEGATIVE	COSM28519
KRAS	p.Lys117Asn	c.351A>C	NEGATIVE	COSM19940
MAP2K1	p.Glu203Lys	c.607G>A	NEGATIVE	COSM232755
MAP2K1	p.Glu203Val	c.608A>T	NEGATIVE	COSM3386991
MAP2K1	p.Lys57Asn	c.171G>C	NEGATIVE	COSM5520914
MAP2K1	p.Lys57Asn	c.171G>T	NEGATIVE	COSM1235478
MAP2K1	p.Lys57Met	c.170A>T	NEGATIVE	MAN124
MAP2K1	p.Lys57Thr	c.170A>C	NEGATIVE	COSM4756761
MAP2K1	p.Phe53Ile	c.157T>A	NEGATIVE	COSM3503329
MAP2K1	p.Phe53Leu	c.157T>C	NEGATIVE	COSM555604
MAP2K1	p.Phe53Leu	c.159T>A	NEGATIVE	COSM1725008
MAP2K1	p.Phe53Leu	c.159T>G	NEGATIVE	COSM2257208
MAP2K1	p.Phe53Val	c.157T>G	NEGATIVE	COSM1562837
MAP2K1	p.Pro124Gln	c.371C>A	NEGATIVE	COSM1167912
MAP2K1	p.Pro124Leu	c.371C>T	NEGATIVE	COSM1315861
MAP2K1	p.Pro124Ser	c.370C>T	NEGATIVE	COSM235614
MAP2K2	p.Gln60Pro	c.179A>C	NEGATIVE	COSM145610
MAP2K2	p.Phe57Leu	c.171T>G	NEGATIVE	OM3158
MAP2K2	p.Phe57Leu	c.171T>A	NEGATIVE	COSM3389034
MAP2K2	p.Phe57Leu	c.169T>C	NEGATIVE	COSM1235618
MAP2K2	p.Phe57Val	c.169T>G	NEGATIVE	COSM3534171
MET	NA	NA	NEGATIVE	COSM29633
MET	NA	NA	NEGATIVE	COSM24687
MET	NA	NA	NEGATIVE	COSM35468
MET	p.His1112Arg	c.3335A>G	NEGATIVE	COSM703
MET	p.His1112Leu	c.3335A>T	NEGATIVE	COSM698
MET	p.His1112Tyr	c.3334C>T	NEGATIVE	COSM696
MET	p.Met1268Ile	c.3804G>A	NEGATIVE	COSM694
MET	p.Met1268Thr	c.3803T>C	NEGATIVE	COSM691
MET	p.Thr1010Ile	c.3029C>T	NEGATIVE	COSM707
MET	p.Tyr1021Asn	c.3061T>A	NEGATIVE	COSM48564
MET	p.Tyr1021Phe	c.3062A>T	NEGATIVE	COSM339515
MET	p.Tyr1248Cys	c.3743A>G	NEGATIVE	COSM699
MET	p.Tyr1248His	c.3742T>C	NEGATIVE	COSM690
MET	p.Tyr1253Asp	c.3757T>G	NEGATIVE	COSM700
MTOR	p.Cys1483Arg	c.4447T>C	NEGATIVE	COSM3747775
MTOR	p.Cys1483Phe	c.4448G>T	NEGATIVE	COSM462616
MTOR	p.Cys1483Trp	c.4449C>G	NEGATIVE	OM3149
MTOR	p.Cys1483Tyr	c.4448G>A	NEGATIVE	COSM462615
MTOR	p.Glu1799Lys	c.5395G>A	NEGATIVE	COSM180789
MTOR	p.Leu2427Arg	c.7280T>G	NEGATIVE	COSM2119114
MTOR	p.Leu2427Gln	c.7280T>A	NEGATIVE	COSM1185313
MTOR	p.Phe1888Ile	c.5662T>A	NEGATIVE	COSM3358968
MTOR	p.Phe1888Leu	c.5664C>G	NEGATIVE	COSM462604
MTOR	p.Phe1888Leu	c.5664C>A	NEGATIVE	COSM893813
MTOR	p.Phe1888Leu	c.5662T>C	NEGATIVE	COSM3358967
MTOR	p.Phe1888Val	c.5662T>G	NEGATIVE	COSM893814

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MTOR	p.Ser2215Phe	c.6644C>T	NEGATIVE	COSM1686998
MTOR	p.Ser2215Pro	c.6643T>C	NEGATIVE	COSM1560108
MTOR	p.Ser2215Tyr	c.6644C>A	NEGATIVE	COSM20417
MTOR	p.Thr1977Arg	c.5930C>G	NEGATIVE	COSM462602
MTOR	p.Thr1977Lys	c.5930C>A	NEGATIVE	COSM462601
MTOR	p.Thr1977Ser	c.5929A>T	NEGATIVE	COSM1289945
MTOR	p.Val2006Ile	c.6016G>A	NEGATIVE	COSM893804
MTOR	p.Val2006Leu	c.6016G>C	NEGATIVE	COSM1134662
MTOR	p.Val2006Phe	c.6016G>T	NEGATIVE	COSM249481
NRAS	p.Ala146Thr	c.436G>A	NEGATIVE	COSM27174
NRAS	p.Ala146Val	c.437C>T	NEGATIVE	COSM4170228
NRAS	p.Ala59Thr	c.175G>A	NEGATIVE	COSM578
NRAS	p.Gln61Arg	c.182A>G	NEGATIVE	COSM584
NRAS	p.Gln61Glu	c.181C>G	NEGATIVE	COSM581
NRAS	p.Gln61His	c.183A>T	NEGATIVE	COSM585
NRAS	p.Gln61His	c.183A>C	NEGATIVE	COSM586
NRAS	p.Gln61Leu	c.182A>T	NEGATIVE	COSM583
NRAS	p.Gln61Lys	c.181C>A	NEGATIVE	COSM580
NRAS	p.Gln61Pro	c.182A>C	NEGATIVE	COSM582
NRAS	p.Gly12Ala	c.35G>C	NEGATIVE	COSM565
NRAS	p.Gly12Arg	c.34G>C	NEGATIVE	COSM561
NRAS	p.Gly12Asp	c.35G>A	NEGATIVE	COSM564
NRAS	p.Gly12Cys	c.34G>T	NEGATIVE	COSM562
NRAS	p.Gly12Ser	c.34G>A	NEGATIVE	COSM563
NRAS	p.Gly12Val	c.35G>T	NEGATIVE	COSM566
NRAS	p.Gly13Ala	c.38G>C	NEGATIVE	COSM575
NRAS	p.Gly13Arg	c.37G>C	NEGATIVE	COSM569
NRAS	p.Gly13Asp	c.38G>A	NEGATIVE	COSM573
NRAS	p.Gly13Cys	c.37G>T	NEGATIVE	COSM570
NRAS	p.Gly13Ser	c.37G>A	NEGATIVE	COSM571
NRAS	p.Gly13Val	c.38G>T	NEGATIVE	COSM574
NRAS	p.Lys117Asn	c.351G>T	NEGATIVE	MAN13
PDGFRA	p.Asn659Lys	c.1977C>A	NEGATIVE	COSM22415
PDGFRA	p.Asn659Lys	c.1977C>G	NEGATIVE	COSM22414
PDGFRA	p.Asn659Tyr	c.1975A>T	NEGATIVE	COSM22416
PDGFRA	p.Asp842_His845del	c.2526_2537delCATCATGCATGA	NEGATIVE	COSM737
PDGFRA	p.Asp842_Met844del	c.2524_2532delGACATCATG	NEGATIVE	COSM12401
PDGFRA	p.Asp842Tyr	c.2524G>T	NEGATIVE	COSM12396
PDGFRA	p.Asp842Val	c.2525A>T	NEGATIVE	COSM736
PDGFRA	p.Ile843_Asp846del	c.2527_2538delATCATGCA TGAT	NEGATIVE	COSM12400
PDGFRA	p.Ile843_Ser847delinsThr	c.2528_2539delTCATGCATGATT	NEGATIVE	COSM12407
PDGFRA	p.Val561Asp	c.1682T>A	NEGATIVE	COSM739
PIK3CA	p.Arg108His	c.323G>A	NEGATIVE	COSM27497



**Clinical Test Report : Oncomine™ Dx Target Test US v10.0**

Sample ID: ADF10\_PRZ\_Run41\_Sample7\_20211117165500\_ET3JO462 Date Of Birth: 01 AUG 1980

Date: 13 DEC 2021

**Draft Report**

Gene	Amino Acid Change	Nucleotide Change	Test Result	Hotspot ID
PIK3CA	p.Arg38Cys	c.112C>T	NEGATIVE	COSM744
PIK3CA	p.Arg38Gly	c.112C>G	NEGATIVE	COSM40945
PIK3CA	p.Arg38His	c.113G>A	NEGATIVE	COSM745
PIK3CA	p.Arg38Ser	c.112C>A	NEGATIVE	COSM87310
PIK3CA	p.Arg88Gln	c.263G>A	NEGATIVE	COSM746
PIK3CA	p.Arg93Trp	c.277C>T	NEGATIVE	COSM27493
PIK3CA	p.Asn1044Lys	c.3132T>A	NEGATIVE	COSM12592
PIK3CA	p.Asn345Ile	c.1034A>T	NEGATIVE	COSM94978
PIK3CA	p.Asn345Lys	c.1035T>A	NEGATIVE	COSM754
PIK3CA	p.Cys378Arg	c.1132T>C	NEGATIVE	COSM756
PIK3CA	p.Cys378Phe	c.1133G>T	NEGATIVE	COSM21450
PIK3CA	p.Cys378Tyr	c.1133G>A	NEGATIVE	COSM1041478
PIK3CA	p.Cys420Arg	c.1258T>C	NEGATIVE	COSM757
PIK3CA	p.Cys901Arg	c.2701T>C	NEGATIVE	COSM1420899
PIK3CA	p.Cys901Phe	c.2702G>T	NEGATIVE	COSM769
PIK3CA	p.Cys901Tyr	c.2702G>A	NEGATIVE	COSM1420901
PIK3CA	p.Gln546Arg	c.1637A>G	NEGATIVE	COSM12459
PIK3CA	p.Gln546Glu	c.1636C>G	NEGATIVE	COSM6147
PIK3CA	p.Gln546Lys	c.1636C>A	NEGATIVE	COSM766
PIK3CA	p.Gln546Pro	c.1637A>C	NEGATIVE	COSM767
PIK3CA	p.Glu365Gly	c.1094A>G	NEGATIVE	COSM1420797
PIK3CA	p.Glu365Lys	c.1093G>A	NEGATIVE	COSM86044
PIK3CA	p.Glu365Val	c.1094A>T	NEGATIVE	COSM1484860
PIK3CA	p.Glu39Lys	c.115G>A	NEGATIVE	COSM30625
PIK3CA	p.Glu542Lys	c.1624G>A	NEGATIVE	COSM760
PIK3CA	p.Glu542Val	c.1625A>T	NEGATIVE	COSM762
PIK3CA	p.Glu545Ala	c.1634A>C	NEGATIVE	COSM12458
PIK3CA	p.Glu545Asp	c.1635G>C	NEGATIVE	COSM27374
PIK3CA	p.Glu545Asp	c.1635G>T	NEGATIVE	COSM765
PIK3CA	p.Glu545Gln	c.1633G>C	NEGATIVE	COSM27133
PIK3CA	p.Glu545Gly	c.1634A>G	NEGATIVE	COSM764
PIK3CA	p.Glu545Lys	c.1633G>A	NEGATIVE	COSM763
PIK3CA	p.Glu547Lys	c.1639G>A	NEGATIVE	COSM29315
PIK3CA	p.Glu726Gly	c.2177A>G	NEGATIVE	COSM1420887
PIK3CA	p.Glu726Lys	c.2176G>A	NEGATIVE	COSM87306
PIK3CA	p.Glu81Lys	c.241G>A	NEGATIVE	COSM27502
PIK3CA	p.Gly1049Arg	c.3145G>C	NEGATIVE	COSM12597
PIK3CA	p.Gly1049Ser	c.3145G>A	NEGATIVE	COSM777
PIK3CA	p.Gly106Val	c.317G>T	NEGATIVE	COSM748
PIK3CA	p.His1047Arg	c.3140A>G	NEGATIVE	COSM775
PIK3CA	p.His1047Leu	c.3140A>T	NEGATIVE	COSM776
PIK3CA	p.His1047Tyr	c.3139C>T	NEGATIVE	COSM774
PIK3CA	p.His701Arg	c.2102A>G	NEGATIVE	COSM1420881
PIK3CA	p.His701Pro	c.2102A>C	NEGATIVE	COSM778
PIK3CA	p.Lys111Glu	c.331A>G	NEGATIVE	COSM13570
PIK3CA	p.Met1043Ile	c.3129G>A	NEGATIVE	COSM29313
PIK3CA	p.Met1043Ile	c.3129G>T	NEGATIVE	COSM773

**Clinical Test Report : Oncomine™ Dx Target Test US v10.0**

Sample ID: ADF10\_PRZ\_Run41\_Sample7\_20211117165500\_ET3J0462 Date Of Birth: 01 AUG 1980

Date: 13 DEC 2021

**Draft Report**

Gene	Amino Acid Change	Nucleotide Change	Test Result	Hotspot ID
PIK3CA	p.Met1043Val	c.3127A>G	NEGATIVE	COSM12591
PIK3CA	p.Pro539Arg	c.1616C>G	NEGATIVE	COSM759
PIK3CA	p.Thr1025Ala	c.3073A>G	NEGATIVE	COSM771
PIK3CA	p.Tyr1021Cys	c.3062A>G	NEGATIVE	COSM12461
PIK3CA	p.Val344Ala	c.1031T>C	NEGATIVE	COSM86951
PIK3CA	p.Val344Gly	c.1031T>G	NEGATIVE	COSM22540
RAF1	p.Ser257Leu	c.770C>T	NEGATIVE	COSM181063
RAF1	p.Ser257Trp	c.770C>G	NEGATIVE	COSM581519
RAF1	p.Thr421Met	c.1262_1263delCCinsTG	NEGATIVE	MAN9
RET	p.Ala883Phe	c.2646_2648delAGCinsTTT	NEGATIVE	COSM981
RET	p.Ala883Ser	c.2647G>T	NEGATIVE	COSM133167
RET	p.Asp898_Glu901del	c.2694_2705delTGTTTATG AAGA	NEGATIVE	COSM962
RET	p.Cys618Arg	c.1852T>C	NEGATIVE	COSM29803
RET	p.Cys618Tyr	c.1853G>A	NEGATIVE	COSM980
RET	p.Cys620Arg	c.1858T>C	NEGATIVE	COSM29804
RET	p.Cys634Arg	c.1900T>C	NEGATIVE	COSM966
RET	p.Glu768Asp	c.2304G>C	NEGATIVE	COSM21338
RET	p.Glu768Gly	c.2303A>G	NEGATIVE	COSM1347811
RET	p.Met918Thr	c.2753T>C	NEGATIVE	COSM965
ROS1	p.Gly2032Arg	c.6094G>C	NEGATIVE	MAN11
ROS1	p.Gly2032Arg	c.6094G>A	NEGATIVE	MAN10
ROS1	p.Leu1951Met	c.5851C>A	NEGATIVE	COSM1072521

Example Report