

ABOUT THE TEST FoundationOne®Liquid is a next generation sequencing (NGS) assay that identifies clinically relevant genomic alterations in circulating tumor DNA.

PATIENT

TUMOR TYPE Breast cancer (NOS) REPORT DATE

CRF# XXXXXXXXX

PATIENT

DISEASE Breast cancer (NOS) NAME Not Given DATE OF BIRTH Not Given SEX Not Given MEDICAL RECORD # Not Given

PHYSICIAN

ORDERING PHYSICIAN Not Given MEDICAL FACILITY Not Given ADDITIONAL RECIPIENT Not Given MEDICAL FACILITY ID Not Given PATHOLOGIST Not Given

SPECIMEN

SPECIMEN ID Not Given SPECIMEN TYPE Blood DATE OF COLLECTION Not Given SPECIMEN RECEIVED Not Given

Biomarker Findings MSI Status Undetermined

Genomic Findings

For a complete list of the genes assayed, please refer to the Appendix. ESR1 D538G, E380Q, Y537N FGFR1 amplification NF1 G751fs*9 MDM2 amplification MYC amplification TP53 R248Q

6 Therapies with Clinical Benefit

30 Clinical Trials

3 Therapies with Lack of Response

30 Clinical Iri

ACTIONABILITY

BIOMARKER FINDINGS

MSI Status Undetermined

GENOMIC FINDINGS	MAF %	THERAPIES WITH CLINICAL BENEFIT (IN PATIENT'S TUMOR TYPE)	THERAPIES WITH CLINICAL BENEFIT (IN OTHER TUMOR TYPE)
ESR1 - D538G	3.0%	Fulvestrant	none
E380Q	0.09%	▲ Anastrozole ¹	
Y537N	0.3%	A Exemestane ¹	
9 Trials see p. 12		▲ Letrozole ¹	
FGFR1 - amplification	n/a	none	Pazopanib
6 Trials see p. 14			Ponatinib
NF1 - G751fs*9	0.45%	none	Cobimetinib
			Trametinib
9 Trials see p. 15			Binimetinib
MDM2 - amplification	n/a	none	none
1 Trials see p. 16			
		Definition he resistant to	

▲ 1. Patient may be resistant to indicated therapy



p. 5

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GENOMIC FINDINGS	MAF %	THERAPIES WITH CLINICAL BENEFIT (IN PATIENT'S TUMOR TYPE)	THERAPIES WITH CLINICAL BENEFIT (IN OTHER TUMOR TYPE)
MYC - amplification	n/a	none	none
5 Trials see p. 17			

GENOMIC FINDINGS WITH NO REPORTABLE THERAPEUTIC OR CLINICAL TRIALS OPTIONS

For more information regarding biological and clinical significance, including prognostic, diagnostic, germline, and potential chemosensitivity implications, see the Genomic Findings section.

TP53 - R248Q

NOTE Genomic alterations detected may be associated with activity of certain FDA-approved drugs; however, the agents listed in this report may have varied clinical evidence in the patient's tumor type. Neither the therapeutic agents nor the trials identified are ranked in order of potential or predicted efficacy for this patient, nor are they ranked in order of level of evidence for this patient's tumor type.



CRF# XXXXXXXX

	25%		
	2%		
Mutant Allele Frequency Percentage (MAF%)			
	Not Detected	Liquid biopsy 01 Jan 2018	
HISTORIC PATIENT FINDINGS	5	TEST 1	
ESR1	• D538G*2	3.0%	
	• E380Q	0.09%	
	• Y537N	0.3%	
FGFR1	amplifcation	n/a	
NF1	G 751fs*9	0.45%	
MDM2	amplifcation	n/a	
TP53	amplifcation	n/a	
ТР53	R248Q	21.0%	

NOTE This comparison table refers only to genes and biomarkers assayed by prior FoundationOne® Liquid or FoundationOne® tests. Up to five previous tests may be shown.

For some genes in FoundationOne Liquid only select exons are assayed. Therefore, an alteration found by a previous test may not have been confirmed despite overlapping gene lists. Please refer to the Appendix for the complete list of genes and exons assayed. The gene and biomarker list will be updated periodically to reflect new knowledge about cancer biology.

As new scientific information becomes available, alterations that had previously been listed as Variants of Unknown Significance (VUS) may become reportable.