Ordered By

Patient

Name: Jessica Moore Date of Birth: 1941-07-06

Sex: Male

Case Number: 1636

Diagnosis: Follicular lymphoma

Specimen Information

Primary Tumor Site: Lymphoma

Specimen Site: Spleen Specimen ID: 9720

Specimen Collected: 2023-07-01

Test Initiated: 2023-07-04

Biomarker	Method	Analyte	Result		Biomarker level	
ER	IHC	protein	Positive 3+, 100%	BENEFIT	abemaciclib, palbociclib, ribociclib, endocrine, therapy, everolimus	level 2
PR	IHC	protein	Positive 2+, 95%	BENEFIT	abemaciclib, palbociclib, ribociclib, endocrine therapy	level 2
ТМВ	seq	DNA tumor	6 m/Mb Low	BENEFIT	pembrolizumab	level 2
ERBB2	IHC	Protien	Negative 0	LACK OF BENEFT	trastuzumab, ado-trastuzumab emtansine, pertuzumab, fam-trastuzumab deruxtecan-nxki, lapatinib, neratinib, tucatinib	level 1

Cancer-Type Relevant Biomarkers

BioMarker	Method	Analyte	Result
HBA2	Seq	RNA-Tumor	Fusion not detected
Mismatch repair status	IHC	Protien	Positive 3+, 76%
PD-L1(SP142)	IHC	Protien	Positive 1+, 25%
CSF1R	Seq	DNA-Tumor	Mutation not detected
JAK3	Seq	DNA-Tumor	Stable

Genomic Signatures

BioMarker	Method	Analyte	Result
Microsatellite instability	Seq	DNA tumor	Equivocal
Tumor mutational burden	Seq	DNA tumor	6 mutations/Mb Low
Genomic loss of heterozygosity (LOH)	Seq	DNA tumor	Low - 8% of tested genmoic segments exhibit LOH

Genes Tested with Pathogenic Alterations or likely Pathogenic Alterations

Gene	Method	Analyte	Variant Interpretation	Protien Alteration	Exon	DNA Alteration	Allele Frequency %
EGFR	Seq	DNA tumor	Likely Pathogenic	p.H1038R	6	c.3113A>G	20.25
NT5C2	Seq	DNA tumor	Likely Benign	p.G503A	8	c.1508G>C	11.15
APC	Seq	DNA tumor	Benign	p.G388R	7	c.1162G>A	21.7
PKLR	Seq	DNA tumor	Likely Benign	p.R367Q	6	c.1100G>A	16.25

Gene Variants of Unknown Significance

Gene	Method	Analyte	Variant Interpretation	Protien Alteration	Exon	DNA Alteration	Allele Frequency %
PTEN	Seq	DNA tumor	Variant of uncertain significance	p.M1V	7	c.1A>G	21.64
KRAS	Seq	DNA tumor	Variant of uncertain significance	p.Y736fs*4	8	c.2207_2212delinsTAGATTC	13.75
RNF43	Seq	DNA tumor	Variant of uncertain significance	p.T1195S	15	c.3583A>T	3.9

Immunohistochemistry Results

Biomarker	Result
PR	Positive 2+, 3%
AR	Negative 1+, 7%
PD-L1(SP142)	Positive 1+, 25%
PTEN	Negative 3+, 72%
ER	Negative 2+, 21%

Genes Tested with Indeterminate Results by Tumor DNA Sequencing

HDAC1 KIT MYOD1

Specimen Information

Specimen ID: 9720 Specimen Collected: 2023-07-01 Specimen Recieved: 2023-07-04 Testing Initiated: 2023-07-04

Gross Description: 9720

Pathological Diagnosis:

Left breast, central, 12:00, suspicious mass, 12-gauge core needle biopsy: Infiltrating moderately-differentiated mammary carcinoma, grade 2, Nottingham score 6 (architectural grade 3, nuclear grade 2, mitotic figures 1).

Dissection Information:

Molecular testing of this specimen was performed after harvesting of targeted tissues with an approved manual microdissection technique. Candidate slides were examined under a microscope and areas containing tumor cells (and separately normal cells, when necessary for testing) were circled. A laboratory technician harvested targeted tissues for extraction from the marked areas using a dissection microscope.

Clinical Trials Connector

CHEMOTHERAPY CLINICAL TRIALS							
Drug class Biomarker Method Analyte Investigational agents		Investigational agents					
Anti hormonal therapy	ER	IHC	protein	anastrazole, letrozole, exemestane, fulvestrant, tamoxifen, goserelin, leuprolide			
Anti hormonal therapy	PR	IHC	protein	anastrazole, letrozole, exemestane, fulvestrant, tamoxifen, goserelin, leuprolide			
Anti inflammatory agents	PIK3CA	NGS	DNA tumor	aspirin			

TARGETED THERAPY CLINICAL TRIALS						
Drug class Biomarker Method		Analyte	Investigational agents			
Akt inhibitors	ARID1A	NGS	DNA tumor	AZD5363, MK-2206, ipataserib		
immunomodulatory agents	TMB	NGS	DNA tumor	avelumab, atezolizumab, durvalumab, ipilimumab, nivolumab, pembrolizumab		
PARP inhibitors	NBN	NGS	DNA tumor	BGB-290, BMN-673, olaparib, rucaparib, talazoparib		
Akt/mTor inhibitors	PIK3CA	NGS	DNA tumor	AZD5363, BYL719, MK-2206, ipataserib, everolimus, temsirolimus		