

Study GO40782 Clinical Trial Assay

CAUTION—Investigational Device. Limited by Federal (or United States) law to investigational use.

For performance evaluation only - the results provided in this report are for investigational use only to determine eligibility for GO40782.

STUDY

Partner Name **Roche Pharma**Partner Study ID **GO40782**FMI Study ID **F1S-BPA-PRO-18-993**

TEST

FMI Test Order # ORD-0847341-01 Test Type FoundationOne DX1 Report Date 16 Jul 2020

PATIENT

Subject ID **15006-1308**

Site ID **15006**

Sex Male

Date of Birth 13MAY1956

Diagnosis Prostate acinar adenocarcinoma

SPECIMEN

Specimen ID **10022276**

Sample Type Slide Deck

Site Lymph Node

Collection Date 28MAY2020

Received Date 03JUL2020

Visit Type Archival/Pre-Treatment

Potential Enrollment Eligible Alterations

GENE ALTERATION

None Detected

GENOMIC FINDINGS

NOTE: This is a comprehensive list of cancer-related alterations detected in this patient's sample.

GENE ALTERATION

PTEN loss

PTEN A328fs*16 GNAS R160C TP53 T253N

GENOMIC SIGNATURES

NOTE: This section includes information for genomic signatures reported in this test.



Subject ID 15006-1308 Report Date
16 Jul 2020



Biomarker Result

Tumor Mutational Burden 3.78 mutations-per-megabase

Microsatellite Instability MS-Stable

VARIANTS OF UNKNOWN SIGNIFICANCE

Note: These variants may not have been adequately characterized in the scientific literature at the time this report was issued, and/ or the genomic context of these alterations makes significance unclear. FMI VUS are included here, in the event that they become clinically meaningful in the future.

GENE ALTERATION CALR amplification

SYK H506R CREBBP D7Y TSC2 L1137V PIK3CB N553I