

Patient		Physician	
Name : xxxxxxxxxxx		Name : xx xxxxxxxxxxx	
Gender : M	Date of Birth : DD/MM/YYYY	Institute : xxxxxxxxxxx	

Diagnosis : Adenocarcinoma of Colon			
Sample Type : FFPE	Sample Collection Date : DD/MM/YYYY	Sample ID : PB_CG_SC_XXXX-XX_X	
Test : PositiveSelect CRC	Technology : Illumina NGS	Coverage : 1000x	Report Date : DD/MM/YYYY

Patient Tumor Type Specific Genes

Gene	Genetic Alteration	Result
BRAF Mutation [p.V600E]	No alteration detected	Negative
KRAS Mutation [Exon 2]	[c.35G>A, p.G12D]	Positive
KRAS Mutation [Exon 2, 3, 4]	No alteration detected	Negative
MLH1 Mutation	No alteration detected	Negative
MSH2 Mutation	No alteration detected	Negative
MSH6 Mutation	[c.467C>A, p.Ser156Ter]	Positive
NRAS Mutation [Exon 2, 3, 4]	No alteration detected	Negative

* Note: MSH6 [c.467C>A] mutation identified in this patient has been classified as pathogenic by ACMG [American College of Medical Genetics and Genomics]. Another variant at the same position MSH6 c.467C>G [p.Ser156Ter] has been regarded pathogenic as per ClinVar, however the identified mutation in the patient [c.467C>A, p.Ser156Ter] also results in premature stop codon. Thus the identified mutation may also result in similar effects as [c.467C>G] due to protein truncation.

Note: Genomic alterations in the 6 genes related to cancer type (Colorectal Cancer) as listed in NCCN guidelines are reported here.

Positive Biosciences Ltd.

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About PositiveSelect CRC

PositiveSelect CRC is a targeted-NGS based diagnostic test, which tests for all four categories of genomic alterations; single nucleotide variations, insertions, copy number variations, structural variations, in the 6 genes as listed above. It adds the advantage of making available a comprehensive molecular testing solution as indicated in the guidelines with specificity and sensitivity higher than the conventional platforms.

The PositiveSelect CRC test, has been designed to work both on FFPE and liquid biopsy, thus ensuring the criteria for sample procurement and requirement is made convenient. The blood collection tube that comes with the test, ensures low risk of sample degradation and extraction of adequate ctDNA for the test. The custom-designed assay enriches all the 6 recommended gene targets from the cfDNA and further is sequenced in a depth of 1000X on the Next-Seq Illumina NGS platform. The end result involves massive parallel sequencing of all the enriched cancer-specific target regions along with flanking introns.

Analyzed by:

Verified by:

Scientific Officer

Sr. Scientific Officer

Disclaimer: The information in this report is meant for medical professionals only. This report should not be construed as personal medical advice and is not intended to replace medical advice ordered by physicians. This document should not be used to establish any standard of care. Clinicians should use their own clinical judgment and not base clinical decisions solely on this document. Positive Bioscience will not be liable for any direct, indirect, consequential, special, exemplary, or other damages. Using this report means the person undertaking this test consents and fully agrees to all terms listed at <http://www.positivebioscience.com/terms.aspx>