

Result Summary Positive Result- The following heterozygous alteration was identified:
Amino Acid change: p.R497PfsX6 (Arg497ProfsX6) DNA change: c.1489dupC (g.37070354)
Classification: PATHOGENIC. Interpretation - The c.1489dupC (p.R497PfsX6) alteration is
a known pathogenic mutation. This result is consistent with a diagnosis of Lynch syndrome for
this individual. Method a³¹P-Bi-directed sequencing analysis was performed to test for the
presence of a mutation in all coding regions and intron/exon boundaries of the MLH1 gene. Array
comparative genomic hybridization (aCGH) was used to test for the presence of large deletions
and duplications in this gene.