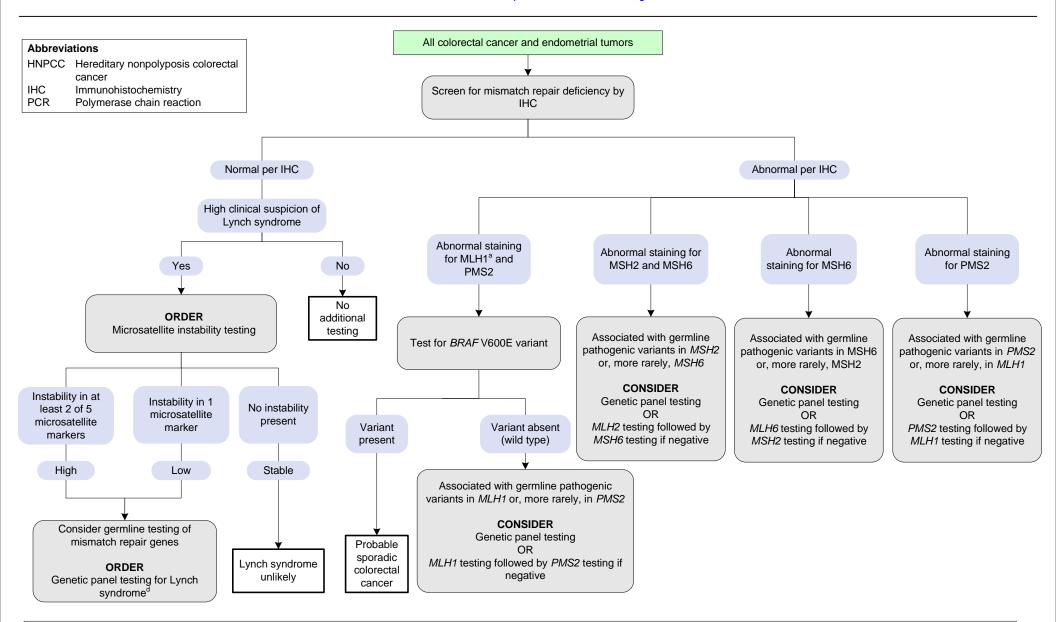


Lynch Syndrome (HNPCC) Testing

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^aLoss of MLH1 may be due to either acquired hypermethylation (in sporadic tumors) or a germline mutation (in Lynch syndrome).

^bPanel (reflex) tests are available (Mismatch Repair by Immunohistochemistry with Reflex to *BRAF* Codon 600 Mutation and *MLH1* Promoter Methylation; Mismatch Repair by Immunohistochemistry with Reflex to *MLH1* Promoter Methylation).

^cNot applicable to endometrial cancers; order only *MLH1* Promoter Methylation.

^dConsider targeted testing if a specific variant has been previously identified in a family member.