Microsatellite Testing in Colon Cancer

Suzanne M. Mahon, RN, DNSc, AOCN®, APNG

Significant progress has been made in understanding the molecular genetic basis of colorectal cancer (CRC). That information is paving the way to understanding the genetic basis of other tumors, as well. Oncology nurses should anticipate the routine integration of this information and testing of CRC tumors to understand the molecular basis of the disease in clinical practice. Molecular testing can lead to the identification of families at risk for hereditary cancer syndromes, particularly Lynch syndrome, which sometimes is referred to as hereditary nonpolyposis colorectal cancer. Knowledge of the genetic basis of CRC also contributes valuable information aimed at selecting appropriate and effective targeted therapy.

Three pathogenetic pathways have been identified and implicated in the development of CRC: chromosomal instability (CIN), microsatellite instability (MSI), and CpG island methylator phenotype (CIMP). The characteristics of those molecular pathways are shown in Table 1.

Microsatellite Instability Defined

MSI is the condition in which genetic hypermutability (i.e., a state in which mutations are abnormally frequent) exists. MSI results from defective DNA mismatch repair (MMR) genes. Defective MMR, which can lead to MSI, occurs in two main situations: (a) an individual with Lynch syndrome who has a germline MMR mutation develops an acquired mutation in his or her working allele of the MMR gene, or (b) when an individual has acquired MLH1 promoter hypermethylation of one MMR gene and develops an acquired mutation of the other allele. Screening a colorectal tumor for MSI provides phenotypic evidence that MMR is not functioning properly but does not identify the underlying pathology.

MMR genes correct errors that spontaneously occur during DNA replication, including single-base mismatches or short insertions and deletions (Matloff, Lucas, Polydorides, & Itzkowitz, 2013). The proteins involved in MMR form a complex that binds to the mismatch, identifies the correct strand of DNA, and then subsequently excises the error and repairs the mismatch. Cells with abnormally functioning MMR tend to accumulate errors rather than correct them. As a result, gene sequences are not preserved consistently through DNA replication, and new microsatellite fragments are created. That repair system is mainly composed of four proteins (MLH1, MSH2, MSH6, and PMS2) interacting together to recognize mismatches and remove them (Buecher et al., 2013).

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Laboratory Techniques

MSI detects that MMR is defective, but does not imply the mechanism by which it is impaired. PCR technology can be used as a cost-effective screening tool for MMR gene mutations that can be confirmed by gene sequencing. MSI testing can be performed on fresh, frozen, or paraffin-embedded tumor material. PCR-based assays reveal defective microsatellites.

Five markers (often called Bethesda markers) have been recommended by the National Cancer Institute to screen for MSI in CRC tumors (Weissman et al., 2013). The Bethesda panel includes two mononucleotide repeats (BAT-25 and BAT-26) and three dinucleotide repeats (D2S123, D5S346, and D17S250). If the tumor has no instability (i.e., none of the repeat lengths has changed), the tumor is considered microsatellite stable. MSI detection in two of the markers (or 30% or more of unstable markers if a larger panel is used) is considered a positive result (Buecher et al., 2013). The MSI-low phenotype occurs with instability in only one marker, or 10%–30% of markers in larger panels.

The PCR method does not detect which protein in the MMR is deficient (Gibson, Lacy, Matloff, & Robert, 2014). PCR technology cannot distinguish between sporadic cancers or Lynch syndrome in MSI-high tumors. Immunohistochemical (IHC) analysis of MMR proteins is an alternative method to detect MSI and primarily is used to complement MSI genetic testing when Lynch syndrome is suspected (Buecher et al., 2013). The loss of expression of one or more of those proteins indicates an MMR defect and determines which gene is most likely to have a germline mutation. The interpretation of IHC must consider the dependent expression of specific MMR protein heterodimers: MSH2/MSH6 and MLH1/PMS2. PMS2 and MSH6 are considered minor MMR proteins that work with the two major MMR proteins, MLH1 and MSH2, respectively, and whose expression is dependent on their binding to the major partner. Therefore, the loss of expression of MSH2 is frequently associated with the loss of expression of MSH6, and this pattern is highly suggestive of an MSH2 germline mutation. In addition, loss of expression of MLH1 is frequently associated with loss of expression of PMS2, and this pattern may result either from MLH1 germline mutation or from acquired somatic hypermethylation of the MLH1 gene promoter (Power, Glogowski, & Lipkin, 2010). Loss of MSH2/MSH6 suggests Lynch syndrome, whereas loss of MLH1/PMS2, although seen in Lynch syndrome, is characteristic of sporadic CRC.
Clinical Implications of Microsatellite Instability Testing

MSI is found in about 15% of CRCs and has a key role in the diagnostic strategy of identifying individuals with Lynch syndrome, whose tumors are characterized by the presence of this phenotype (Buecher et al., 2013). About 25% of individuals with Lynch syndrome do not meet traditional clinical Amsterdam or Bethesda criteria for germline testing (Weissman et al., 2012) (see Figure 1). Lynch syndrome, which is an autosomal dominant condition caused by an inactivating germline mutation of one of the four genes involved in the MMR system (MLH1, MSH2, MSH6, PMS2), is the most frequent form of hereditary CRC and accounts for about 5% of all cases of CRC. The identification of individuals with Lynch syndrome is critical because the application of an early and intensive surveillance program combined with prophylactic surgery significantly reduces the incidence of colorectal and gynecologic cancers and other malignancies, as well as improves mortality rates, for both the patient and affected relatives. Therefore, accurate identification is beneficial in terms of cost-effectiveness and improved quality of life (Serrano et al., 2012).

Knowledge of MSI also provides important prognostic information. MSI tumors are associated with a good prognosis and are known to have a resistance to 5-fluorouracil (5-FU)–based adjuvant chemotherapy, which has a clinical application when selecting therapy (Kloor, Staffa, Ahadova, & von Knebel Doeberitz, 2014; Meguerditchian & Bullard Dunn, 2013). Patients with MSI-high CRC who receive 5-FU treatment do not have an advantage over those not receiving it, and this treatment might be harmful in MSI stage II CRC (Buecher et al., 2013; Hampel, 2010). Given the observation that MSI-high tumors display less aggressive behavior, in addition to the fact that MSI tumors respond poorly to 5-FU–based chemotherapy, MSI testing for stage II CRCs is becoming more routine (Gala & Chung, 2011) and is recommended by the National Comprehensive Cancer Network (Kelley, Van Bebber, Phillips, & Venook, 2011). A better understanding of somatic genetics and molecular pathways involved in MMR CRC is guiding continued research toward novel and tailored therapeutic strategies for this disease.

Table 1. Characteristics of the Molecular Pathways in CRC

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>CIN</th>
<th>MSI</th>
<th>CIMP</th>
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<tbody>
<tr>
<td>Definition</td>
<td>Loss or gain of chromosome arms, translocations, or gene amplifications</td>
<td>MSI CRCs have a better prognosis in general and a different response to the chemotherapeutic agent 5-FU. Sporadic MSI CRC usually occurs at an older age, and in these cases, no family history of cancer typically exists. Genetic MSI CRC usually occurs at a younger age (younger than 50 years) and with a family history of cancer.</td>
<td>Sometimes referred to as the serrated pathway</td>
</tr>
<tr>
<td>Precursor lesions</td>
<td>Tubular adenoma polyp or villous adenoma polyp</td>
<td>Polyps at an early age</td>
<td>Serrated adenoma</td>
</tr>
<tr>
<td>Mutation</td>
<td>Acquired somatic mutations in APC, KRAS, TP53, SMAD4, PIK3CA, SOX9, ARID1A, and FAM123B</td>
<td>Two-thirds of MSI tumors (10% of all CRCs) are sporadic and caused by somatic biallelic hypermethylation of the MLH1 promoter. One-third of MSI tumors (5% of all CRCs) are germline Lynch syndrome. MMR genes, MSH2, MSH6, MLH1, and PMS2, are involved in tumor initiation and progression.</td>
<td>DNA hypermethylation at specific regulatory sites, enriched in CpG islands in the promoter regions of tumor suppressor genes mainly in the BRAF mutation</td>
</tr>
<tr>
<td>Affected side of colon</td>
<td>Primarily left-sided</td>
<td>Primarily right-sided</td>
<td>Tend to be right-sided, but can be found throughout the colon</td>
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</tbody>
</table>

CIMP—CpG island methylator phenotype; CIN—chromosomal instability; CRC—colorectal cancer; 5-FU—5-fluorouracil; MMR—mismatch repair; MSI—microsatellite instability

Note. Percentage of CRCs is 50%–75% for CIN, 15% for MSI, and 20% for CIMP.

Note. Based on information from Gibson et al., 2014; Kalady, 2013; Rosner & Strul, 2014.
of inheritance but not MSI, and will not be identified with universal screening (Hall, 2010; Matloff et al., 2013). The advent of next-generation sequencing may help further identify families with hereditary predisposition and MSI tumors. Patients who receive care in an institution that performs universal MSI testing on CRC tumors do not necessarily have to give informed consent for MSI testing and may not realize that they might learn that they have Lynch syndrome and are at risk for the hereditary predisposition to colorectal and other cancers, which could be accompanied by negative psychosocial consequences (Hall, 2010). The ordering healthcare provider needs to ascertain that those with MSI-high tumors are referred to genetics professionals for complete evaluation and realize the clinical implications and potential risks of MSI testing (Weissman et al., 2012).

The MSI phenotype was first discovered in CRC, but since then, its detection has been regarded as indicative of a defective MMR system. MSI is not unique to CRCs, but also is observed in other tumor types, including gastric cancer, endometrial cancer, ovarian cancer, sebaceous carcinomas, glioblastoma, and lymphomas (Rosner & Strul, 2014). Oncology professionals should continue to expect more knowledge and research to emerge, which will provide insight into the best way to use the information gleaned from molecular testing, such as screening for hereditary predisposition syndromes and improved tailored treatment for malignancy.

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References


Figure 1. Amsterdam II and Bethesda Criteria

Note: Based on information from Burt et al., 2013; Umar et al., 2004; Weissman et al., 2012.