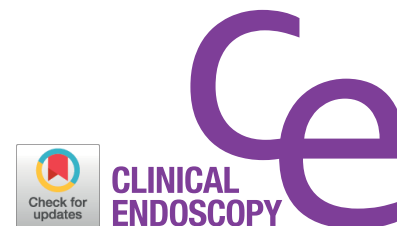


REVIEW

Clin Endosc 2019;52:235-238
https://doi.org/10.5946/ce.2018.112
Print ISSN 2234-2400 • On-line ISSN 2234-2443



Open Access

Screening Relevance of Sessile Serrated Polyps

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Conventional adenomas have historically been considered to be the only screening-relevant colorectal cancer (CRC) precursor lesion. The prevailing paradigm was that most CRCs arise along the chromosomal instability pathway, where adenomas accumulate incremental genetic alterations over time, leading eventually to malignancy. However, it is now recognized that this “conventional” pathway accounts for only about two-thirds of CRCs. The serrated pathway is responsible for most of the remainder, and is a disproportionate contributor to postcolonoscopy CRC. Hallmarks of the serrated pathway are mutations in the *BRAF* gene, high levels of methylation of promoter CpG islands, and the sessile serrated polyp (SSP). Accumulating evidence shows that SSPs can be considered adenoma-equivalent from the standpoint of CRC screening. SSPs have a higher prevalence than previously thought, and appear to have a relatively long dwell time similar to that of conventional adenomas. In addition, SSPs, whether sporadic or as part of the serrated polyposis syndrome, are associated with increased risk of synchronous and metachronous neoplasia. These features collectively support that SSPs are highly relevant to CRC prevention. **Clin Endosc** 2019;52:235-238

Key Words: Colorectal neoplasms; Screening; Colonoscopy

INTRODUCTION

Colorectal cancer (CRC) is amenable to primary prevention, which means that incidence and mortality can be reduced via screening and removal of precursor lesions before cancer occurs. Successful CRC screening fundamentally depends on the ability to detect and resect precursor lesions with relatively long dwell time. This paradigm has long been established for conventional adenomas, but the more recent emergence of serrated lesions in the CRC field has led to questions about its applicability to sessile serrated polyp (SSP). There are presently no prospective studies showing a measurable independent impact of SSP removal on CRC outcomes. This may be largely due to the relatively recent recognition of the importance of SSP in the CRC prevention field, compared to the very ex-

tensively studied conventional adenomas. Nevertheless, an extensive and growing literature provides several converging lines of evidence suggesting that SSP have a central role in CRC prevention. Recent studies describe the distinct biologic features of the serrated pathway to colorectal neoplasia, and reveal that SSP have high prevalence, a relatively long dwell time, and are associated with increased risk of both synchronous and metachronous colorectal neoplasia. A review of this literature is presented in the following sections.

THE SERRATED PATHWAY

Serrated polyps are distinct from conventional adenomas and are characterized histologically by a saw-tooth appearance of the crypt epithelium.¹ This class of lesions includes hyperplastic polyps, SSP, and traditional serrated adenomas (TSA). SSP and TSA are both considered premalignant lesions; however, TSA are very rare lesions and are less well-understood than SSP. Distortion of crypt growth pattern leading to abnormal-appearing crypt bases is characteristic of SSP. An activating mutation of the *BRAF* oncogene appears to be a fundamental step in the SSP-to-CRC serrated neoplasia pathway, and a characteristic feature is abnormal hypermeth-

Received: June 27, 2018 Accepted: July 10, 2018

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ylation of CpG islands in gene promoter regions, termed CpG island-methylator phenotype (CIMP). This hypermethylation can be extensive and lead to epigenetic silencing of genes, such as the mutation mismatch repair gene *MLH1* (also implicated in Lynch syndrome). This event leads to acceleration of development of cytological dysplasia (via the intermediate SSP with cytological dysplasia), and progression to CRC with microsatellite instability (MSI). Most CRCs with MSI and CIMP molecular signatures originate from the serrated pathway, with the SSP as the main precursor lesion.²

Additional evidence supporting the existence of a separate serrated pathway to CRC includes the observation that SSP may have risk factors distinct from those typically associated with adenomas. For example, calcium and vitamin D supplementation are associated with increased risk of SSP,³ while obesity, folate intake, fiber and fat intake are not.⁴

NATURAL HISTORY OF SSP

The natural history of SSP and rate of progression to CRC are not as clear compared to conventional adenomas and the adenoma-carcinoma sequence. However, the available evidence suggests comparable progression rates, with SSPs generally following an indolent or slowly progressive course, until a critical event (notably, *MLH1* silencing) which accelerates the progression to dysplasia and cancer.

A pathological series⁵ of 2,319 patients with 2,416 SSPs reported that the median age of patients increased as the histologic grade of the lesion worsened, from 61 years for SSP without dysplasia, to 76 years for CRC. A recent cross-sectional analysis⁶ of 137 SSPs with dysplasia or cancer showed that older age, female gender, proximal location, CIMP, and lack of aberrant p53 were all associated with loss of *MLH1* expression. The age of patients with dysplastic SSP was similar to those with cancer (about 76 years), and together, 17 years older than a historical cohort of patients with SSP without dysplasia.

Collectively, these data show that SSP without dysplasia generally have a long dwell time of at least 15 years, similar to conventional adenomas. This is a critically important observation, because uniformly rapid progression of serrated neoplasms would have threatened current CRC prevention paradigms which rely on relatively long latency of CRC and its precursor lesions.

PREVALENCE

Until recently, there has been uncertainty regarding the true

prevalence of SSP. Older studies addressing this question were limited because they reported aggregate data for endoscopists with wide ranges in individual detection rates, and because there is significant inter-observer variability among pathologists in the differentiation of serrated lesion subtypes. The true prevalence of SSP is important to define, because a rare lesion would not be a reasonable target to screen for. Recent colonoscopy studies^{7,8} involving endoscopists with high-level detection rates reported SSP prevalence of 8% to 9% (0.4% to 0.6% for SSP with cytological dysplasia). These studies show that SSP have higher prevalence than previously thought, adding further support to their screening relevance.

SERRATED NEOPLASIA AND POSTCOLONOSCOPY CRC

There is significant overlap in the molecular signatures of postcolonoscopy CRC (PCCRC) and the serrated pathway. Compared to sporadic CRC, PCCRC is more likely to demonstrate MSI and be associated with CIMP. In addition, PCCRC is more likely to be located in the proximal colon,⁹⁻¹¹ the area of predilection for SSP.

The majority of PCCRC are due to operator-dependent factors, and this is reflected in its inverse association with adenoma detection rate (ADR). The issues surrounding variable adenoma detection and its impact on the effectiveness of colonoscopy may also apply to serrated polyps, although there is currently no evidence that the variability in proximal serrated polyp detection is independently associated with PCCRC risk. Several studies have shown that the variability in detection of proximal serrated polyps exceeds that of adenomas among the same groups of endoscopists, and can range up to 18-fold (compared to 3- to 4-fold for the ADR).¹²⁻¹⁴ SSPs have subtle morphology which renders them more likely to be overlooked by operators not familiar with their endoscopic characteristics. Longer withdrawal time is significantly associated with improved proximal serrated polyp detection. An analysis¹⁵ of nearly 8,000 colonoscopies in the New Hampshire Colonoscopy Registry found that the detection of serrated polyps proximal to the sigmoid colon increased with each minute of withdrawal time above 6 minutes. Studies have generally shown a strong correlation between adenoma and serrated polyp detection,¹³ which is plausible because endoscopists who have meticulous examination technique would be expected to be proficient at detecting polyps of all types.

Their endoscopic features also render serrated polyps prone to incomplete resection, as shown in the Complete Adenoma Resection study.¹⁶ However, more recent studies show that dedicated endoscopic mucosal resection methods, with submuco-

sal injection with dye to highlight lesion borders, are associated with effective and complete resection of serrated polyps.^{17,18}

ASSOCIATION WITH SYNCHRONOUS NEOPLASIA

Several lines of evidence show that serrated polyps not only possess intrinsic risk of progression to CRC, but also alter the colorectal milieu in a manner which promotes neoplastic processes. A systematic review and meta-analysis¹⁹ of 9 studies with 34,000 subjects reported that individuals with serrated polyps had a 2-fold increased odds of advanced neoplasia; proximal location and large size were associated with even higher odds. The most extreme manifestation of serrated neoplasia is the serrated polyposis syndrome (SPS), which is defined as the presence of at least 5 serrated polyps proximal to the sigmoid colon with 2 or more measuring at least 10 mm in diameter, any number of serrated polyps proximal to the sigmoid colon in a person with a first-degree relative with SPS, or >20 serrated polyps of any size distributed throughout the colon.¹ SPS is the most common polyposis syndrome, with reported prevalence rates ranging from 1 in 238 to 1 in 111 in colonoscopy and fecal test-based screening programs, respectively.^{20,21} SPS is associated with increased risk of CRC, warranting close endoscopic surveillance of patients with SPS.²² Recent reports,²³ however, suggest that the risk of CRC in SPS is highest at the time of diagnosis: in a Spanish study of 296 SPS patients, CRC was found in 47 patients at mean age of 54 years, and only 4 cases were detected during surveillance (5-year incidence of 1.9%).

ASSOCIATION WITH METACHRONOUS NEOPLASIA

Convincing evidence shows that SSP are associated with

risk of metachronous neoplasia. A recent study²⁴ reported the long-term follow-up of 13,000 individuals in the Norwegian flexible sigmoidoscopy screening trial (NORCCAP), where patients with serrated polyps were re-invited for colonoscopy over a decade after initial enrollment. Compared with controls without polyps, the hazard ratio for CRC was 2.5 for patients with large serrated polyps, comparable to those who had an advanced adenoma (2.0). Interestingly, there were 23 large serrated polyps found at screening were left in situ for a median of 11 years, and none had progressed to CRC; this observation is further evidence that SSP exert a pro-neoplastic influence on the colorectal milieu, independent of their intrinsic risk of progression. A large population-based study²⁵ from Denmark assessed the risk of CRC associated with SSP. Among 272,342 individuals who had received colonoscopy between 1977 and 2009, there were 2,045 CRC cases and 8,105 age- and sex-matched controls without CRC. The investigators identified the first colorectal polyp(s) that had been resected or biopsied on the initial or subsequent colonoscopy, and polyps that were initially interpreted as hyperplastic were re-reviewed by expert pathologists. Patients with SSP are at increased risk for CRC, to a degree at least equivalent to that of patients with conventional adenomas (Table 1).²⁵

A recently published analysis²⁶ of the New Hampshire population-based colonoscopy registry showed that patients with large serrated polyps or SSP at baseline had higher risk of large serrated polyps at follow-up (odds ratio, 9.7 to 14.3), whereas patients with high-risk adenoma (HRA) plus SSP had risk of metachronous HRA which was 4 times higher than those with baseline HRA alone.

CONCLUSIONS

SSPs contribute significantly to the burden of CRC, and recognition of their importance has direct implications for CRC screening paradigms. The existence of a distinct biologic

Table 1. Odds of Colorectal Cancer according to Baseline Polyp Type²⁵

| | Adjusted OR (95% CI) | Estimated 10-year risk |
|---|----------------------|------------------------|
| SSP with synchronous conventional adenomas | 2.66 (1.70–4.16) | 2.47% |
| SSP without synchronous conventional adenomas | 3.40 (2.35–4.91) | 3.16% |
| SSP with cytologic dysplasia | 4.76 (2.59–8.73) | 4.43% |
| SSP without cytologic dysplasia | 2.75 (1.99–3.80) | 2.56% |
| Conventional adenomas without SSP | 2.50 (2.24–2.80) | 2.33% |
| TSA overall | 4.84 (2.36–9.93) | 4.50% |
| Hyperplastic polyps only | 1.30 (0.96–1.77) | 1.21% |

CI, confidence interval; OR, odds ratio; SSP, sessile serrated polyp; TSA, traditional serrated adenomas.

pathway, high prevalence, relatively long dwell time, relation to PCCRC, and association with synchronous and metachronous neoplasia render SSPs highly screening-relevant. Continued efforts to ensure that endoscopists achieve high levels of detection and complete resection are critical for effective CRC prevention at colonoscopy.

Conflicts of Interest

The author has no financial conflicts of interest.

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