



# Clinical characteristics of patients with serrated polyposis syndrome in Korea: comparison with Western patients

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**Background/Aims:** Serrated polyposis syndrome (SPS) has been shown to increase the risk of colorectal cancer (CRC). However, little is known about the characteristics of Asian patients with SPS. This study aimed to identify the clinicopathological features and risk of CRC in Korean patients with SPS as well as the differences between Korean and Western patients based on a literature review. **Methods:** This retrospective study included 30 patients with SPS as defined by World Health Organization classification treated at Samsung Medical Center, Korea, between March 1999 and May 2011. **Results:** Twenty patients (67%) were male. The median patient age at diagnosis was 56 years (range, 39–76 years). A total of 702 polyps were identified during a median follow-up of 43 months (range, 0–149 months). Serrated polyps were noted more frequently in the distal colon (298/702, 55%). However, large serrated polyps and serrated adenomas were mainly distributed throughout the proximal colon (75% vs. 25% and 81% vs. 19%, respectively); 73.3% had synchronous adenomatous polyps. The incidence of CRC was 10% (3/30 patients), but no interval CRC was detected. A total of 87% of the patients underwent esophagogastroduodenoscopy and 19.2% had significant lesions. **Conclusions:** The phenotype of SPS in Korean patients is different from that of Western patients. In Korean patients, SPS is more common in men, there were fewer total numbers of serrated adenoma/polyps, and the incidence of CRC was lower than that in Western patients. Korean patients tend to more frequently have abnormal gastric lesions. However, the prevalence of synchronous adenomatous polyps is high in both Western and Korean patients. (**Intest Res 2017;15:402-410**)

**Key Words:** Serrated polyposis syndrome; Serrated adenoma/polyp; Colorectal neoplasms

## INTRODUCTION

Serrated polyposis syndrome (SPS), previously known as hyperplastic polyposis, is a rare condition characterized by multiple serrated polyps (SPs) spread throughout the colon and rectum.<sup>1-3</sup> Since SPS is a somewhat complex syndrome

whose molecular basis is not yet clearly understood, the World Health Organization (WHO) developed consensus criteria for the clinical diagnosis of SPS.<sup>4</sup> SPS is associated with an increased personal and familial risk of developing colorectal cancer (CRC) despite endoscopic surveillance.<sup>1,5-10</sup> Boparai et al.<sup>6</sup> reported that the cumulative CRC risk was 7% at 5 years in patients with SPS undergoing endoscopic surveillance. Edelstein et al.<sup>8</sup> reported a high recurrence rate (68%) of sessile serrated adenomas/polyps (SSA/P) and/or adenomas as well as a significant risk of CRC. Therefore, close surveillance including an annual colonoscopy is recommended in patients with SPS.<sup>8,11-13</sup>

However, a considerable number of patients with symptoms suggestive of CRC including acute large bowel obstruction, diffuse abdominal pain or discomfort, weight loss, bowel habit changes, anemia, and rectal bleeding episodes were

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included in previous studies. These inclusions might have produced an overestimated CRC risk.<sup>2,12</sup> Furthermore, most previous studies were performed in Western countries. There is very little data regarding SPS patients in Asian countries.<sup>14</sup>

In the present study, we investigated the clinical characteristics of SPS and evaluated the CRC risk by assessing the cumulative incidence of CRC in patients with SPS in Korea. We also identified the differences between Western and Korean patients with SPS.

## METHODS

### 1. Patients

We retrospectively enrolled consecutive patients aged  $\geq 18$  years who fulfilled SPS WHO criterion 1 or 3 and were treated at Samsung Medical Center, Seoul, Korea, between March 1999 and May 2011.

Specific data including demographic factors such as age, sex, and personal history of CRC and other malignancies were obtained from the medical records. Colonoscopy reports with corresponding pathology reports during follow-up were collected to derive information regarding polyp size, number, location, morphology, and treatment method. Esophagogastroduodenoscopy (EGD) reports were ascertained to evaluate extracolonic cancer risks. A cumulative method was used to count the polyps; all that were diagnosed over the course of colonoscopies were counted. Interval CRC was defined as a CRC that developed under endoscopic surveillance after SPS diagnosis.

The follow-up period was measured from the date of the initial colonoscopy to that of the last surveillance colonoscopy. Patients were excluded if they had a known germline adenomatous polyposis coli mutation, Lynch syndrome, IBD, or total proctocolectomy.

This study was approved by the Institutional Review Board of Samsung Medical Center (IRB number: 2011-12-077). Because this study conducted a retrospective data review, consent was not required or obtained. However, patient records and information were anonymized and de-identified prior to the analysis.

### 2. Definition

The WHO criteria were used to identify SPS. The criteria included: (1) at least 5 SPs proximal to the sigmoid colon, of which 2 measured at least 10 mm in diameter (WHO criterion 1); (2) any number of SPs occurring proximal to the

sigmoid colon in an individual who has a first-degree relative with SPS (WHO criterion 2); and/or (3)  $\geq 20$  SPs spread throughout the colon (WHO criterion 3).<sup>4</sup> In the present study, none of the patients were diagnosed with SPS based on the second criterion alone because familial histories are not always reliable in a retrospective study. Adherence to the described criteria was assessed by analyzing endoscopy reports and the corresponding pathology reports as well as pathology reports of colonic surgical resection specimens. All of the counted polyps were biopsy proven and tallied once. Polypectomy or endoscopic mucosal resection for the polyps was performed once they were  $\geq 5$  mm in diameter.<sup>14-16</sup>

Polyps were classified as SP and conventional adenoma. SPs were subdivided as hyperplastic polyps (HPs) and serrated adenomas (SAs). The pathology reports showed SSA/P or traditional serrated adenoma (TSA). However, since the distinction between these polyps was not made throughout the early study period and they are both considered precursor lesions in the "serrated pathway," the SA category comprised both lesion types.<sup>16,17</sup>

The proximal colon was defined as the area proximal to the splenic flexure (transverse, ascending colon, and cecum), while the distal colon was defined as the descending colon and sigmoid colon.

## RESULTS

### 1. Clinical Characteristics of SPS Patients

A total of 120,095 patients (69,369 men and 50,726 women) underwent a colonoscopy at the Samsung Medical Center between March 1999 and May 2011. Among them, 30 (0.025%) met the WHO criteria for SPS. Table 1 presents the clinical characteristics of the patients with SPS. The median age at diagnosis was 56 years (range, 39–76 years), and 20 (66.7%) patients were male. Twenty-one of 30 patients (70%) met WHO criterion 1 and 12 patients (40%) fulfilled WHO criterion 3. Three patients initially diagnosed with SPS by WHO criterion I then met criterion III during surveillance. In addition, 22 patients (73.3%) had synchronous adenomatous polyps.

The reasons for the initial colonoscopy were screening in 18 patients (60%), colon polyp history in 6 patients (20%), bloody stool and/or anemia in 2 patients (6.7%), bowel habit changes in 2 patients (6.7%), abdominal pain in 1 patient (3.3%), and history of CRC in 1 patient (3.3%). The median overall follow-up period was 43 months (range, 0–149 months) and the median follow-up period from the diagnosis of SPS was 35 months (range, 0–128 months). A total

of 6 patients (20%) underwent only the initial colonoscopy. Among them, 2 patients could not undergo surveillance colonoscopy due to cancer progression (advanced gastric cancer in one, advanced rectal cancer in one). The other 4 patients (13.3%) were lost to follow-up after the initial colonoscopy. In the time period observed, 113 colonoscopies were performed for a median number of colonoscopies per

patient of 3 (range, 1–11). Three of the 30 patients underwent colectomy. Two patients were diagnosed with CRC at the initial colonoscopy. The other patient underwent colectomy because the polyps were too numerous to remove endoscopically. However, there was no evidence of hereditary polyposis syndrome on genetic testing.

A total of 26 patients (86.7%) underwent EGD. One patient (3.8%) had numerable HPs, 2 (7.7%) had gastric adenoma, and 2 (7.7%) were diagnosed with stomach cancer. Another 2 patients (7.7%) had other extracolonic malignancies that included pancreatic cancer and hepatocellular carcinoma (one each).

**Table 1.** Demographic and Clinical Characteristics of Patients with SPS

| Characteristic                                 | Value      |
|--|------------|
| No. of patients                                | 30         |
| Sex  |            |
| Male   | 20 (66.7)  |
| Female   | 10 (33.3)  |
| Age at SPS diagnosis (yr)                      |            |
| Mean±SD  | 57.4±9.9   |
| Median (range)                                 | 56 (39–76) |
| WHO criteria                                   |            |
| I  | 18 (60)    |
| III  | 9 (30)     |
| I & III <sup>a</sup>                           | 3 (10)     |
| Follow-up duration (mo)                        |            |
| Total  | 43 (0–149) |
| From SPS diagnosis                             | 35 (0–128) |
| Reason for colonoscopy                         |            |
| Screening                                      | 18 (60)    |
| Surveillance for history of colon polyp or CRC | 7 (23.3)   |
| Symptom <sup>b</sup>                           | 5 (16.7)   |
| Colonoscopy                                    |            |
| Total no.                                      | 113        |
| Per patients                                   | 3 (1–11)   |
| Patients with CRC                              |            |
| Past history of CRC                            | 1 (3.3)    |
| At initial colonoscopy                         | 2 (6.6)    |
| Interval CRC                                   | 0          |
| Patients with extracolonic malignancies        | 4 (13.3)   |
| Stomach cancer                                 | 2 (6.7)    |
| Pancreatic cancer                              | 1 (3.3)    |
| Hepatocellular carcinoma                       | 1 (3.3)    |

Values are presented as number (%) or median (range).

<sup>a</sup>Three patients initially diagnosed with SPS by WHO criterion I then met criterion III during surveillance.

<sup>b</sup>Symptoms included bowel habit changes, abdominal pain, bloody stool, and/or anemia.

SPS, serrated polyposis syndrome; WHO, World Health Organization; CRC, colorectal cancer.

**Table 2.** Polyp Characteristics of the Study Patients with SPS

| Characteristic                | Value        |
|-------------------------------|--------------|
| Patient                       |              |
| SA <sup>a</sup>               | 15 (50)      |
| SP <sup>b</sup> ≥10 mm        | 28 (93)      |
| Conventional adenoma          | 22 (73.3)    |
| Patient with recurrence       |              |
| SA                            | 3 (20)       |
| Conventional adenoma          | 15 (68.2)    |
| Recurrence interval in months |              |
| SA                            | 29 (3–50)    |
| Conventional adenoma          | 18 (3–35)    |
| Polyp                         |              |
| HP                            | 508 (72)     |
| SA                            | 31 (4.5)     |
| Conventional adenoma          | 137 (20)     |
| Other polyps <sup>c</sup>     | 26 (3.5)     |
| Distribution of polyps        |              |
| Proximal colon                |              |
| SP (HP/SA)                    | 241 (216/25) |
| Distal colon                  |              |
| SP (HP/SA)                    | 298 (292/6)  |
| Size in mm                    |              |
| Largest SP                    | 15 (8–45)    |
| Largest HP                    | 13 (5–25)    |
| Largest SA                    | 20 (10–45)   |

Values are presented as number (%) or median (range).

<sup>a</sup>SA includes sessile serrated adenoma and traditional serrated adenoma.

<sup>b</sup>SPs include hyperplastic polyp and serrated adenoma.

<sup>c</sup>Other polyps include inflammatory pseudopolyp, mucosal tag, and hamartomatous polyp.

SPS, serrated polyposis syndrome; SA, serrated adenoma; SP, serrated polyp; HP, hyperplastic polyp.

**2. Features of Polyps in SPS Patients**

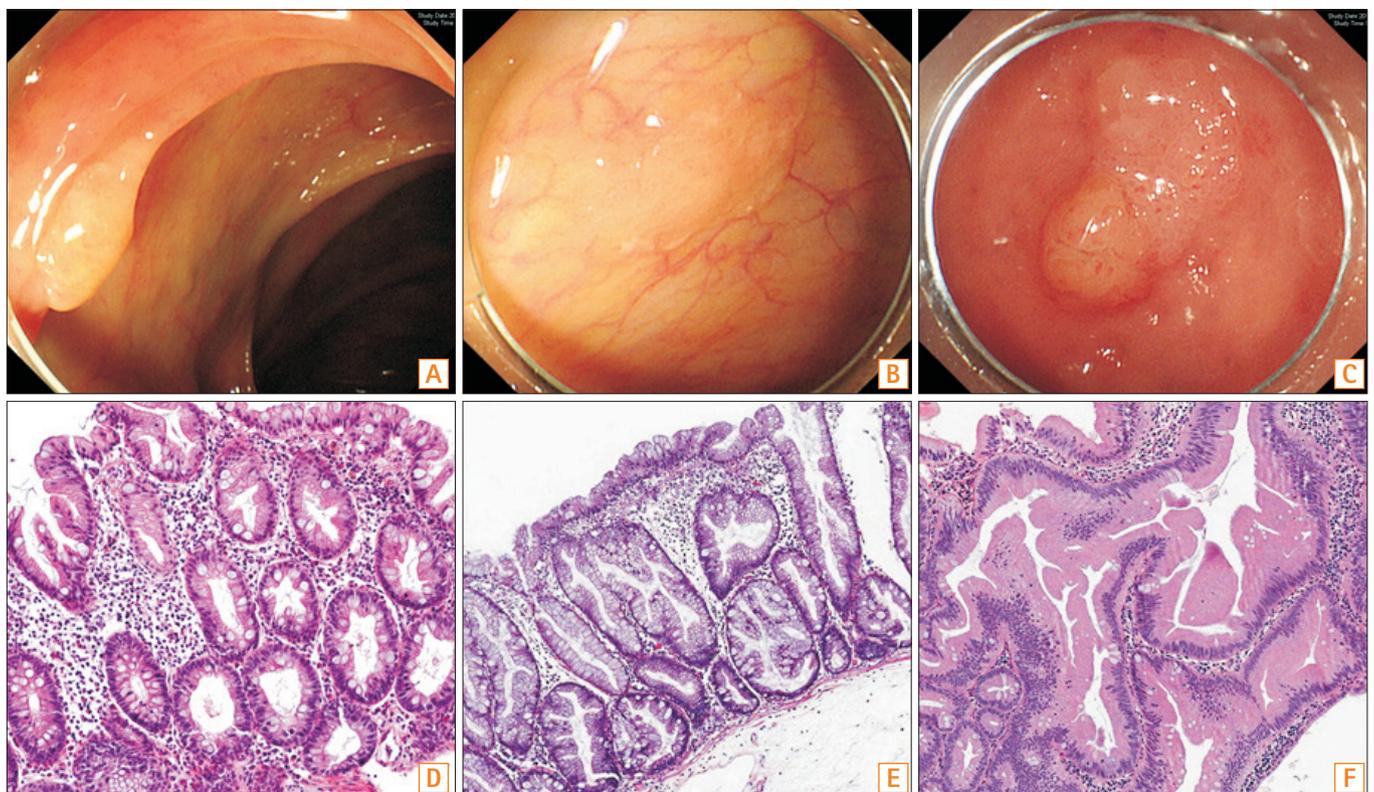
The characteristics of polyps in the patients with SPS are summarized in Table 2. A total of 702 polyps were detected: 508 (72%) HPs, 31 (4.5%) SAs, and 137 (19.5%) conventional adenomas (Fig. 1). Of the SPs, 241 (45%) were located in the proximal colon and 298 (55%) in the distal colon. However, SPs  $\geq 10$  mm in diameter were more found frequently in the proximal colon than in the distal colon. Similar to SPs  $\geq 10$  mm, the SAs were mainly distributed throughout the proximal colon (Fig. 2). The maximum size of the HPs and SAs were 25 mm and 45 mm, respectively; they were all located in the ascending colon. A total of 20% of patients (3/15) had recurrent SA, while 68% (15/22) had adenoma recurrence on surveillance colonoscopy. The median recurrence interval of SA and adenoma was 29 months (range, 3–50 months) and 18 months (range, 3–35 months), respectively. Additional polyps were discovered during surveillance colonoscopy in all patients (Table 3). During follow-up, the median cu-

mulative number of polyps per person was 19 (range, 5–82), 14 HPs (range, 5–55), 0.5 SAs (range, 0–5), and 2 adenomas (range, 0–29).

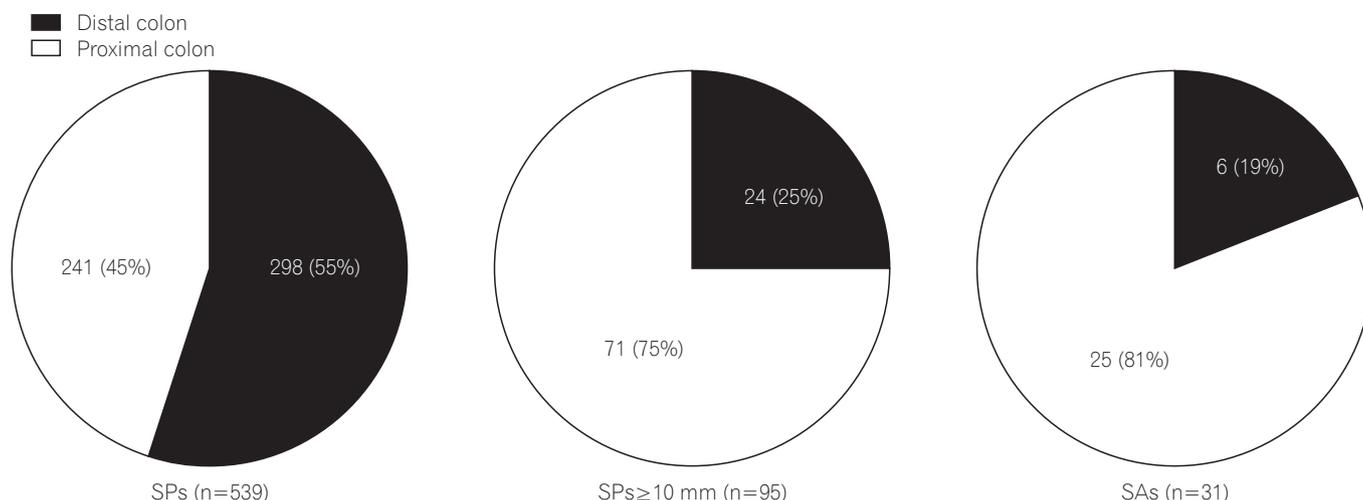
**3. Risk of CRC in SPS Patients**

Of the 30 patients with SPS included in this study, 3 (10%) were diagnosed with CRC. One was already diagnosed with CRC before inclusion, while 2 were diagnosed at the initial colonoscopy. The first patient underwent colon resection for rectal cancer (papillary adenocarcinoma, T2, N0, M0) at 39 years of age, 1 year before being included in this study. He was diagnosed with SPS at the surveillance colonoscopy. He subsequently had an additional 14 HPs located in the proximal colon and measured 3 to 15 mm at surveillance colonoscopy. During follow-up, no cases of SA or advanced adenoma were detected.

Two other patients (6.6%) were diagnosed with CRC at the time of diagnosis of SPS on the initial colonoscopy. One pa-



**Fig. 1.** Examples of the spectrum of lesions found in the colon. (A) Endoscopic finding of hyperplastic polyp (HP): a 6-mm transverse colon polyp with a smooth and pale appearance. (B) Endoscopic finding of sessile serrated adenoma (SSA): a 12-mm ascending colon polyp with a flat or sessile appearance and indistinct borders. It also has a characteristic rim of debris. (C) Endoscopic finding of traditional serrated adenoma (TSA): a 15-mm descending colon polyp showing a granulonodular and lobular appearance. (D) Histopathologic finding of HP: serrated crypts are confined to the upper crypt. (E) Histopathologic finding of SSA: serrated surfaces are enlarged and epithelial serration extends to the crypt bases. (F) Histopathologic finding of TSA: serrated with pseudostratified, elongated nuclei and abundant eosinophilic cytoplasm is characteristic (D–F, H&E stain).



**Fig. 2.** Distribution of serrated polyps (SPs) according to location. SPs ≥ 10 mm were found more frequently in the proximal colon, with the majority usually found in the distal colon. Similar to the distribution of SPs ≥ 10 mm, most serrated adenomas (SAs) were distributed throughout the proximal colon.

**Table 3.** Median Cumulative Number of Polyps per Person during Follow-Up

|                                       | HP          | SA <sup>a</sup> | Adenoma  | Total polyps |
|---------------------------------------|-------------|-----------------|----------|--------------|
| Initial colonoscopy                   | 5.0 (0–21)  | 0 (0–2)         | 1 (0–8)  | 6.0 (0–23)   |
| Colonoscopy at diagnosis <sup>b</sup> | 10.5 (4–26) | 0 (0–2)         | 1 (0–29) | 13.5 (5–56)  |
| Last surveillance colonoscopy         | 14.0 (5–55) | 0.5 (0–5)       | 2 (0–29) | 19.0 (5–82)  |

Values are presented as median (range).

<sup>a</sup>SA includes sessile serrated adenoma and traditional serrated adenoma.

<sup>b</sup>Colonoscopy at which the diagnosis of serrated polyposis was made.

HP, hyperplastic polyp; SA, serrated adenoma.

tient was a 75-year-old woman who underwent colonoscopy because of bowel habit changes that included persistent diarrhea. At the initial colonoscopy, she was diagnosed with CRC in the hepatic flexure. She also had multiple SPs including 1 SA throughout the proximal colon that measured 3 to 25 mm. The largest polyp was proven as HP. She underwent a right hemicolectomy, and examination of the resected specimen revealed an adenocarcinoma microsatellite stable tumor (T1, N0, and M0). The other patient was a 61-year-old man who complained of recurrent bloody stool. On colonoscopy, a rectal mass was apparent and was diagnosed as adenocarcinoma, microsatellite stable tumor. He also had 22 SPs including 1 SA and 1 adenoma located in all segments of the colon that measured 2 to 20 mm. Multiple hepatic metastases were detected during the preoperative work-up, and he subsequently underwent palliative chemotherapy. However, he died of cancer progression within 1 year. All 3 patients were diagnosed with CRC; however, advanced immunohistochemistry testing such as that for the *BRAF*/*KRAS* mutation was not conducted at the time of diagnosis.

During the follow-up of patients with SPS, except for 2 with SPS-synchronous CRC and 1 with a history of CRC, no cases of CRC were detected on the surveillance colonoscopy.

## DISCUSSION

To date, several studies and a case series have reported on SPS; however, they provided limited information on SPS because most were retrospective with small sample sizes. Scant data are available on the natural history of patients with SPS, and studies of Asian patients with SPS are nearly nonexistent. Here we evaluated the clinical characteristics and CRC risk of patients with SPS in Korea. Based on the present findings and data gleaned from the literature, we identified the differences between Western and Asian patients with SPS.

Table 4 summarizes the clinical characteristics of patients with SPS in recent studies reporting >10 cases of SPS. Only a few reports have addressed the prevalence of SPS. In Western studies, the prevalence of SPS was 0.08% to 0.66%.<sup>18-20</sup> One Korean study also reported the prevalence of SPS as

Table 4. Summary of Characteristics of Patients with Serrated (Hyperplastic) Polyposis in Recent Studies with More Than 10 Cases

| Author (year)                            | Nation                              | Case | Age <sup>a</sup> at diagnosis (yr) | Sex (M/F) | WHO criterion (I/II/III/IV) | FU duration <sup>a</sup> (mo) | Synchronous adenomatous polyp                     | CRC prevalence | Interval CRC | Extracolonic malignancy |
|--|-------------------------------------|------|------------------------------------|-----------|-----------------------------|-------------------------------|---|----------------|--------------|-------------------------|
| Leggett et al. (2001) <sup>5</sup>       | Australia                           | 12   | 56.8                               | 5/7       | NS                          | NS                            | 9 Patients (75.0)                                 | 7 (58.3)       | 2 (16.7)     | NS                      |
| Lage et al. (2004) <sup>10</sup>         | Portugal                            | 14   | 53.6                               | 7/7       | NS                          | NS                            | NS  | 6 (43.0)       | NS           | NS                      |
| Ferrández et al. (2004) <sup>21</sup>    | USA                                 | 15   | 52.6                               | 10/5      | NS                          | 36.5                          | 11 Patients (73.0)                                | 1 (6.7.0)      | 0            | NS                      |
| Rubio et al. (2006) <sup>9</sup>         | Sweden                              | 10   | 61.0                               | 8/2       | NS                          | NS                            | 8 Patients (80.0)                                 | 7 (70.0)       | NS           | NS                      |
| Chow et al. (2006) <sup>1</sup>          | Australia                           | 32   | 44.0                               | 15/17     | NS                          | NS                            | 26 Patients (81.0)                                | 10 (31.2)      | NS           | NS                      |
| Boparai et al. (2010) <sup>6</sup>       | The Netherlands                     | 77   | 56.0                               | 42/35     | NS                          | 48.0                          | 273/1,984 Polyps (14.0)                           | 27 (35.0)      | 5 (6.5)      | NS                      |
| Vemulapalli and Rex (2012) <sup>22</sup> | USA                                 | 20   | 63.8                               | 9/11      | 16/0/2/2                    | NS                            | 19 Patients (95.0)                                | 5 (25.0)       | 0            | NS                      |
| Edelstein et al. (2013) <sup>8</sup>     | USA                                 | 44   | 52.5                               | 21/23     | 10/0/34/NS                  | 24.0                          | NS  | 3 (7.6)        | 2 (4.5)      | NS                      |
| Guarinos et al. (2013) <sup>23</sup>     | Spain                               | 50   | 49.3                               | 27/23     | 15/NS/35/NS                 | NS                            | 29 Patients (58.0)                                | 9 (18.0)       | NS           | NS                      |
| Rosty et al. (2013) <sup>24</sup>        | Australia, USA, Canada, New Zealand | 100  | 50.0                               | 42/58     | 12/0/72/16                  | NS                            | 64/80 Patients (80.0)                             | 39 (42.0)      | NS           | NS                      |
| Jasperson et al. (2013) <sup>3</sup>     | USA                                 | 51   | 51.0                               | 24/27     | NS                          | 61.0                          | 42 Patients (82.0)                                | 7 (13.7)       | 1 (1.9)      | 12 (23.5)               |
| Kalady et al. (2011) <sup>25</sup>       | USA                                 | 115  | 62.0                               | 65/50     | NS                          | NS                            | 78 Patients (68.0)                                | 29 (25.0)      | 5 (4.3)      | 32 (28)                 |
| Hazewinkel et al. (2014) <sup>13</sup>   | The Netherlands                     | 41   | 56.5                               | 24/17     | 6/0/21/14                   | 43.0                          | 91/575 Polyps (16.0)                              | 13 (33.0)      | 0            | NS                      |
| Knabe et al. (2014) <sup>26</sup>        | Germany                             | 28   | 54.4                               | 15/13     | 11/0/17/NS                  | 21.5                          | 39/436 Polyps (8.9)                               | 2 (7.1)        | 0            | 4 (14.3)                |
| Hui et al. (2014) <sup>27</sup>          | USA                                 | 40   | 57.0                               | 16/24     | 16/3/32/NS                  | NS                            | 25 Patients (63.0)                                | 16 (40.0)      | 0            | NS                      |
| Elorza et al. (2014) <sup>19</sup>       | Spain                               | 23   | 51.0                               | 19/4      | 4/0/12/7                    | NS                            | 17 Patients (73.0)                                | 6 (26.0)       | 1 (4.3)      | 4 (17.3)                |
| Miwata et al. (2013) <sup>14</sup>       | Japan                               | 10   | 58.3                               | 6/4       | 5/0/5/NS                    | NS                            | 24/91 Polyps (26.4)                               | 3 (30.0)       | NS           | NS                      |
| Toyoshima et al. (2015) <sup>28</sup>    | Japan                               | 21   | 66.0                               | 15/6      | NS                          | NS                            | Advanced adenoma and/or CRC in 10 patients (47.6) | NS             | NS           | NS                      |
| Kim et al. (2015) <sup>29</sup>          | Korea                               | 11   | 55.6                               | 10/1      | 5/0/6/NS                    | NS                            | 7 Patients (64.0)                                 | 1 (9.0)        | NS           | 3 (27.0)                |
| Current study                            | Korea                               | 30   | 57.4                               | 20/10     | 18/0/9/3                    | 43.0                          | 22 Patients (73.3)                                | 3 (10.0)       | 0            | 4 (13.3)                |

Values are presented as number (%).

<sup>a</sup>Age and follow-up duration are shown as median value. If median value was unavailable, mean value was used instead. M, male; F, female; WHO, World Health Organization; FU, follow-up; CRC, colorectal cancer; NS, not specified or unknown.

0.06%.<sup>29</sup> In the present study, the prevalence was estimated as 0.025%. However, a recent Japanese study reported the prevalence of SPS as 8.4%,<sup>28</sup> which was higher than those of the previous studies. The key difference is that all the enrolled patients underwent initial colonoscopy using a magnifying videoscope with chromoendoscopy in the Japanese study. HPs and SPs are difficult to detect by conventional imaging, and the study included patients who were previously treated for a colorectal neoplasm at another hospital. Therefore, the prevalence of SPS is higher than previously known. The age at SPS diagnosis in both Western countries and Asia tended to be in the 50s. However, in the West, the age at diagnosis was slightly younger than that in Asia. Additionally, there was no sex predominance in Western countries, while males predominated in Asian studies.

In the Western studies, the majority of patients met WHO criterion 3 on the basis of a large number of SPs throughout the colon.<sup>8,13,23,24,26</sup> However, in the present study, the majority of patients who met WHO criterion 1 did not fulfill criterion 3. These findings correspond with those of a Japanese study and another Korean study in which half of the patients met WHO criterion 1. No patient underwent colectomy due to numerous polyps.<sup>14,29</sup>

Different lifestyle and dietary factors of Asians may cause these phenotypic differences between Western and Asian patients with SPS.<sup>30-32</sup> However, the distribution of polyps was similar in the Western and Asian studies. The polyps were distributed evenly throughout the colon or predominantly in the distal colon, while large polyps (>10 mm) or SAs tended to be distributed in the proximal colon except in a few Western studies.

Above all, this syndrome increases the risk of CRC. However, scant data exist about the natural history of patients with SPS, while almost all studies were performed in Western countries (Table 4). In the present study, we investigated the natural history of SPS patients in Korea during a median 43 months (range, 0–149 months). The incidence of CRC in the present study was 10% (3/30). This rate was similar to that of another Korean study.<sup>29</sup> In the Western studies, the incidence of CRC was 20% to 40% with the exception of a few small studies in which the incidence was much higher than those in the Korean studies.

There are several reasons for the difference in CRC incidence. First, the Western studies included more patients presenting with symptoms suggestive of CRC. For instance, in the study by Boparai et al.,<sup>6</sup> CRC was found in 35% of the patients (27/77). Among them, 96% (76/77) had reasons for undergoing colonoscopy that included colorectal polyp his-

tory, family history of CRC, and clinical symptoms suggestive of CRC. Because of this selection bias, the estimation of CRC risk is likely to be inflated and reflect a CRC risk associated with symptomatic patients.<sup>2,8,33,34</sup> On the contrary, more than half of the patients (60%) in the present study underwent colonoscopy for screening purposes. Actually, some of the studies that included a majority of asymptomatic patients diagnosed with SPS by screening colonoscopy showed similar CRC incidences to our results (Edelstein et al.,<sup>8</sup> 7.6% [3/44]; Orłowska et al.,<sup>35</sup> 6.6% [1/16]).

Second, the ethnic differences may cause a discrepancy in CRC incidence between Western and Korean SPS patients. Western countries have shown a higher incidence of CRC than Asia countries in general populations.<sup>36</sup> As mentioned above, fewer polyps were found in Asian patients with SPS than in Western patients.<sup>14</sup> Few polyps may cause a low CRC incidence. There may also be genetic difference in patients with SPS. Kim et al.<sup>30</sup> reported that SPs in Korean patients showed somewhat different molecular characteristics than those that occur in Americans.

Third, regardless of the discrepancy of CRC incidence between Western countries and Korea, the incidence of CRC in patients with SPS is higher than that in the general population. Accordingly, we must examine the prevalence of synchronous adenomatous polyps in patients with SPS, which was also higher than that in the general population.<sup>37-39</sup> Therefore, we do not know whether CRC in patients with SPS arose from the adenoma-carcinoma sequence or from the SA pathway. To date, there have been no reports of CRC origin in patients with SPS, so the CRC risk in patients with SPS may be exaggerated slightly.

The U.S. Multi-Society Task Force on Colorectal Cancer recommended annual colonoscopy surveillance in SPS patients.<sup>11</sup> This recommendation considered the high rate of interval cancer and rapid recurrence of colorectal polyps. A literature review<sup>34</sup> reported that 5 of 27 patients (18.5%) with SPS but without CRC on initial colonoscopy developed CRC during follow-up. However, interval cancer can develop from missed polyps or incompletely resected polyps, although some may grow rapidly due to biologic behavior.<sup>40</sup> Also, small polyps (<10 mm) can be easily missed during colonoscopy.<sup>15</sup> According to studies by Hazewinkel et al.<sup>13</sup> and Knabe et al.,<sup>26</sup> no CRC was found in SPS patients during annually surveillance colonoscopy after clearing colonoscopy. In addition, the number of newly detected polyps gradually decreased on sequential colonoscopies. These results support the notion that interval cancer mainly originates from missed polyps rather than from the rapid growth of polyps in

patients with SPS. Therefore, careful colonoscopic examination is more important than frequent surveillance to prevent missed polyps and decrease the CRC risk.

There have been few reports about extracolonic malignancies in patients with SPS. Although previous studies reported various types of extracolonic malignancies, data supporting an association between SP and extracolonic tumor risk is lacking.<sup>3,19,25,26,29</sup>

In the present study, 26 patients (87%) underwent EGD and 5 (19.2%) had significant lesions including multiple HPs (n=1), adenoma (n=2), and stomach cancer (n=2). Although few studies have evaluated the upper gastrointestinal tract, this finding was inconsistent with those of previous Western studies, which described no significant lesions detected on EGD.<sup>3,8,29</sup> Another Korean study also reported that no patients had significant lesions detected by EGD. However, that study enrolled only 10 patients with SPS. Therefore, the present study suggests that Asian patients with SPS may require an evaluation of the upper gastrointestinal tract.

This is the largest retrospective study to date on Korean patients with SPS. We identified the phenotypic differences between Western and Asian patients with SPS based on similar phenotypic characteristics described in other Asian studies. The current surveillance guideline about SPS does not reflect phenotypic differences between Western and Asian patients since it was made on the basis of Western patients. Therefore, large and long-term follow-up cohort studies on Asian patients with SPS and a surveillance guideline for Asian patients are needed.

Our approach has some limitations. First, it was retrospective. The study population could not be investigated under the same condition due to its retrospective nature. As in other retrospective studies, selection bias can influence the data. Second, we grouped SSA and TSA into a single SA category since a large proportion of our data was derived from a period before a distinction was made between SSA and TSA. Third, this study included a relatively short follow-up period. Therefore, the risk of interval cancer could be underestimated. It is difficult to draw a conclusion about the prognosis of SPS from the present study because of the small number of patients and relatively short-term follow-up period. However, most previous studies also had small study populations due to the low prevalence of SPS. Thus, the present study is still the largest observational study of Asian patients with SPS.

In conclusion, the phenotype of SPS in Korean patients is somewhat different from that of Western patients. In Korean patients, SPS is more common in men and the total number of SSA/P is lower than that in Western patients, while the

distribution of large SSA/P is similar. The incidence of CRC is lower in Korean patients, but this rate is higher than that in the general population. The high prevalence of synchronous adenomatous polyps is similar in Western and Korean patients. Although very few studies have evaluated the gastrointestinal tract, Korean patients tend to more frequently have abnormal gastric lesions. Our results show that Korean patients with SPS have a different phenotype from that of Western patients with SPS. Larger prospective studies in Asian SPS patients are needed to clarify the differences in phenotypes and formulate a surveillance guideline for Asian patients with SPS.

REFERENCES

1. Chow E, Lipton L, Lynch E, et al. Hyperplastic polyposis syndrome: phenotypic presentations and the role of MBD4 and MYH. *Gastroenterology* 2006;131:30-39.
2. Orlowska J. Serrated lesions and hyperplastic (serrated) polyposis relationship with colorectal cancer: classification and surveillance recommendations. *Gastrointest Endosc* 2013;77:858-871.
3. Jasperson KW, Kanth P, Kirchoff AC, et al. Serrated polyposis: colonic phenotype, extracolonic features, and familial risk in a large cohort. *Dis Colon Rectum* 2013;56:1211-1216.
4. Snover DC, Ahnen DJ, Burt RW, Odze RD. Serrated polyps of the colon and rectum and serrated polyposis. In: Bosman FT, Carneiro F, Hruban RH, Theise ND, eds. *WHO classification of tumours of the digestive system*. Lyon: IARC Press, 2010:160-165.
5. Leggett BA, Devereaux B, Biden K, Searle J, Young J, Jass J. Hyperplastic polyposis: association with colorectal cancer. *Am J Surg Pathol* 2001;25:177-184.
6. Boparai KS, Mathus-Vliegen EM, Koornstra JJ, et al. Increased colorectal cancer risk during follow-up in patients with hyperplastic polyposis syndrome: a multicentre cohort study. *Gut* 2010;59:1094-1100.
7. Boparai KS, Reitsma JB, Lemmens V, et al. Increased colorectal cancer risk in first-degree relatives of patients with hyperplastic polyposis syndrome. *Gut* 2010;59:1222-1225.
8. Edelstein DL, Axilbund JE, Hyland LM, et al. Serrated polyposis: rapid and relentless development of colorectal neoplasia. *Gut* 2013;62:404-408.
9. Rubio CA, Stemme S, Jaramillo E, Lindblom A. Hyperplastic polyposis coli syndrome and colorectal carcinoma. *Endoscopy* 2006;38:266-270.
10. Lage P, Cravo M, Sousa R, et al. Management of Portuguese patients with hyperplastic polyposis and screening of at-risk first-degree relatives: a contribution for future guidelines based on a clinical study. *Am J Gastroenterol* 2004;99:1779-1784.

11. Lieberman DA, Rex DK, Winawer SJ, et al. Guidelines for colonoscopy surveillance after screening and polypectomy: a consensus update by the US Multi-Society Task Force on Colorectal Cancer. *Gastroenterology* 2012;143:844-857.
12. East JE, Saunders BP, Jass JR. Sporadic and syndromic hyperplastic polyps and serrated adenomas of the colon: classification, molecular genetics, natural history, and clinical management. *Gastroenterol Clin North Am* 2008;37:25-46.
13. Hazewinkel Y, Tytgat KM, van Eeden S, et al. Incidence of colonic neoplasia in patients with serrated polyposis syndrome who undergo annual endoscopic surveillance. *Gastroenterology* 2014;147:88-95.
14. Miwata T, Hiyama T, Oka S, et al. Clinicopathologic features of hyperplastic/serrated polyposis syndrome in Japan. *J Gastroenterol Hepatol* 2013;28:1693-1698.
15. van Rijn JC, Reitsma JB, Stoker J, Bossuyt PM, van Deventer SJ, Dekker E. Polyp miss rate determined by tandem colonoscopy: a systematic review. *Am J Gastroenterol* 2006;101:343-350.
16. Pickhardt PJ, Choi JR, Hwang I, et al. Computed tomographic virtual colonoscopy to screen for colorectal neoplasia in asymptomatic adults. *N Engl J Med* 2003;349:2191-2200.
17. Torlakovic E, Skovlund E, Snover DC, Torlakovic G, Nesland JM. Morphologic reappraisal of serrated colorectal polyps. *Am J Surg Pathol* 2003;27:65-81.
18. Biswas S, Ellis AJ, Guy R, Savage H, Madronal K, East JE. High prevalence of hyperplastic polyposis syndrome (serrated polyposis) in the NHS bowel cancer screening programme. *Gut* 2013;62:475.
19. Elorza G, Enríquez-Navascués JM, Bujanda L, Larzábal M, Gil Lasa I, Martí L. Phenotype characteristics of patients with colonic serrated polyposis syndrome: a study of 23 cases. *Cir Esp* 2014;92:659-664.
20. Moreira L, Pellisé M, Carballal S, et al. High prevalence of serrated polyposis syndrome in FIT-based colorectal cancer screening programmes. *Gut* 2013;62:476-477.
21. Ferrández A, Samowitz W, DiSario JA, Burt RW. Phenotypic characteristics and risk of cancer development in hyperplastic polyposis: case series and literature review. *Am J Gastroenterol* 2004;99:2012-2018.
22. Vemulapalli KC, Rex DK. Failure to recognize serrated polyposis syndrome in a cohort with large sessile colorectal polyps. *Gastrointest Endosc* 2012;75:1206-1210.
23. Guarinos C, Sánchez-Fortún C, Rodríguez-Soler M, et al. Clinical subtypes and molecular characteristics of serrated polyposis syndrome. *Clin Gastroenterol Hepatol* 2013;11:705-711.
24. Rosty C, Walsh MD, Walters RJ, et al. Multiplicity and molecular heterogeneity of colorectal carcinomas in individuals with serrated polyposis. *Am J Surg Pathol* 2013;37:434-442.
25. Kalady MF, Jarrar A, Leach B, et al. Defining phenotypes and cancer risk in hyperplastic polyposis syndrome. *Dis Colon Rectum* 2011;54:164-170.
26. Knabe M, Behrens A, Ell C, Tannapfel A, Pohl J. Endoscopic management for patients with serrated polyposis syndrome is feasible and effective: a prospective observational study at a tertiary centre. *Z Gastroenterol* 2014;52:802-806.
27. Hui VW, Steinhagen E, Levy RA, et al. Utilization of colonoscopy and pathology reports for identifying patients meeting the World Health Organization criteria for serrated polyposis syndrome. *Dis Colon Rectum* 2014;57:846-850.
28. Toyoshima N, Sakamoto T, Makazu M, et al. Prevalence of serrated polyposis syndrome and its association with synchronous advanced adenoma and lifestyle. *Mol Clin Oncol* 2015;3:69-72.
29. Kim HK, Seo KJ, Choi HH, et al. Clinicopathological characteristics of serrated polyposis syndrome in Korea: single center experience. *Gastroenterol Res Pract* 2015;2015:842876.
30. Kim KM, Lee EJ, Ha S, et al. Molecular features of colorectal hyperplastic polyps and sessile serrated adenoma/polyps from Korea. *Am J Surg Pathol* 2011;35:1274-1286.
31. Chung SJ, Kim YS, Yang SY, et al. Prevalence and risk of colorectal adenoma in asymptomatic Koreans aged 40-49 years undergoing screening colonoscopy. *J Gastroenterol Hepatol* 2010;25:519-525.
32. Song KB, Atkinson C, Frankenfeld CL, et al. Prevalence of diadzein-metabolizing phenotypes differs between Caucasian and Korean American women and girls. *J Nutr* 2006;136:1347-1351.
33. Rosty C, Parry S, Young JP. Serrated polyposis: an enigmatic model of colorectal cancer predisposition. *Patholog Res Int* 2011;2011:157073.
34. Orłowska J. Hyperplastic polyposis syndrome and the risk of colorectal cancer. *Gut* 2012;61:470-471.
35. Orłowska J, Kiedrowski M, Kaminski FM, Regula J, Butruk E. Hyperplastic polyposis syndrome in asymptomatic patients: the results from the colorectal-cancer screening program. *Virchows Arch* 2009;455:47.
36. Jemal A, Bray F, Center MM, Ferlay J, Ward E, Forman D. Global cancer statistics. *CA Cancer J Clin* 2011;61:69-90.
37. Conteduca V, Sansonno D, Russi S, Dammacco F. Precancerous colorectal lesions (review). *Int J Oncol* 2013;43:973-984.
38. Brenner H, Altenhofen L, Kretschmann J, et al. Trends in adenoma detection rates during the first 10 years of the German screening colonoscopy program. *Gastroenterology* 2015;149:356-366.e1.
39. Kim HB, Lee YJ, Shim JY, Lee HR. The association between coronary calcification and adenomatous polyps of colon in Korean adults. *Clin Res Hepatol Gastroenterol* 2014;38:649-654.
40. Rex DK. Maximizing detection of adenomas and cancers during colonoscopy. *Am J Gastroenterol* 2006;101:2866-2877.