

Sessile serrated lesion detection rates continue to increase: 2008–2020





Authors

Nicholas Edwardson¹, Prajakta Adsul^{2,3}, Zorisadday Gonzalez², V. Shane Pankratz^{2,3}, Gulshan Parasher^{2,4}, Kevin English⁵, Shiraz Mishra^{3,6}

Institutions

- University of New Mexico, School of Public Administration, Albuquerque, New Mexico, United States
- 2 University of New Mexico, Department of Internal Medicine, Albuquerque, New Mexico, United States
- 3 University of New Mexico, Comprehensive Cancer Center, Cancer Control and Population Sciences, Albuquerque, New Mexico, United States
- 4 University of New Mexico, Department of Gastroenterology, Albuquerque, New Mexico, United States
- 5 Albuquerque Area Indian Health Board Inc., Albuquerque Area Southwest Tribal Epidemiology Center, Albuquerque, New Mexico, United States
- 6 University of New Mexico, Department of Pediatrics, Albuquerque, New Mexico, United States

submitted 13.7.2022 accepted after revision 17.11.2022

Bibliography

Endosc Int Open 2023; 11: E107–E116 DOI 10.1055/a-1990-0509 ISSN 2364-3722 © 2023. The Author(s).

This is an open access article published by Thieme under the terms of the Creative Commons Attribution-NonDerivative-NonCommercial License, permitting copying and reproduction so long as the original work is given appropriate credit. Contents may not be used for commercial purposes, or adapted, remixed, transformed or built upon. (https://creativecommons.org/licenses/by-nc-nd/4.0/)

Georg Thieme Verlag KG, Rüdigerstraße 14, 70469 Stuttgart, Germany

Corresponding author

Nicholas Edwardson, University of New Mexico, School of Public Administration, MSC05 3100 Albuquerque, Albuquerque, NM 87131, United States Fax: +505-277-2529 nedwardson@unm.edu

ABSTRACT

Background and study aims We assessed sessile serrated lesion detection rate (SSLDR) at a large academic medical center from 2008 to 2020 and modeled a local, aspirational target SSLDR. We also assessed SSLDRs among all gastroenterology fellows to better understand the relationship between SSLDRs and total colonoscopies performed.

Patients and methods SSL-positive pathology results were flagged from a dataset composed of all screening colonoscopies for average-risk patients from 2008 to 2020. Unadjusted SSLDRs were calculated for individual endoscopists by year. A mixed effects logistic regression was used to estimate the log odds of SSL detection, with one model estimating division-wide predictors of SSL detection and a second model focused exclusively on colonoscopies performed by fellows. Model-adjusted SSLDRs were estimated for all 13 years and across both categories of all endoscopists and fellows only.

Results Adjusted SSLDRs showed a consistent improvement in SSLDR from a low of 0.37% (95% confidence interval [CI]: 0.10–0.63) in 2008 to a high of 7.94% (95% CI: 6.34–9.54) in 2020. Among fellows only, the odds of SSL detection were significantly lower during their first year compared to their second year (OR: 0.80, 95% CI: 0.66–0.98) but not significantly higher in their third year compared to their second year (OR: 1.09, 95% CI: 0.85–1.4).

Conclusions SSLDR increased steadily and significantly throughout our study period but variance among endoscopists persists. The peak SSLDR from 2020 of 7.94% should serve as the local aspirational target for this division's attendings and fellows but should be continuously reevaluated.

Introduction

Despite a decades-long reduction in colorectal cancer (CRC) incidence and mortality rates, CRC is still the third most commonly diagnosed cancer and the third leading cause of cancer mortality in the US [1,2]. Population-wide improvement in CRC screening adherence is believed to be one of the major drivers of recent decreases in CRC incidence and mortality rates [3]. However, for this downward trajectory of CRC incidence and mortality rates to continue, the quality, and not just the quantity, of CRC screening must improve. A growing body of evidence suggests that the current recommended CRC screening modalities are not equally effective at detecting all types of precursor lesions. Of particular interest are serrated lesions, which account for approximately one-third of all CRC and an even greater percentage of post-colonoscopy interval cancers [4–6]. These lesions are more likely to be missed than conventional adenomas by high-sensitivity guaiac fecal occult blood tests (HS-gFOBT) [7], fecal immunochemical tests (FIT) [7–9], computed tomographic (CT) colonography when diminutive (≤5 mm) [10], and, given serrated lesions' preponderance for the proximal colon [4, 11], flexible sigmoidoscopies [12].

This leaves colonoscopy as the best-suited screening modality for detecting serrated lesions [3]. However, even colonoscopy exhibits disadvantages for detection relative to conventional adenomas. Serrated lesions are generally flat-or sessile-and are often covered with a mucous cap. Combined with their common location in the proximal colon, these lesions can be easily missed by endoscopists. Even when detected, incomplete resection is more common with serrated lesions given the difficulty in accurately detecting borders [13]. Furthermore, even upon successful detection and resection, the pathological criteria for defining serrated lesions have evolved over the past 15 years [14–17]. These endoscopic and pathological detection characteristics, combined with the relatively nascent knowledge and training base related to the serrated pathway, have contributed to highly variable detection rates among endoscopists [18, 19]. Two recent meta-analyses of serrated lesion detection included studies with sessile serrated lesion detection rates (SSLDRs) as low as 0.6% and as high as 10.3% [11, 20]. Both meta-analyses were conducted to begin the process of approximating an aspirational SSLDR, which does not yet exist, similar to the adenoma detection rate (ADR) benchmark of at least 25% (men and women combined) [3].

While the field continues to move toward determining the true prevalence of serrated lesions across diverse patient populations, a call has been made to focus on local SSLDR and to use the highest rate from a unit as the local aspirational target [21]. Doing so incorporates nuances of the local patient population while continuing to call attention to this less understood cancer pathway. In this study, we retrospectively examined the SSLDR of the Gastroenterology Division at the University of New Mexico Health Sciences Center from 2008 to 2020 and model a local, aspirational target SSLDR per Kahi & Rex's recommendation [21]. We also assessed the progression of SSLDR among gastroenterology fellows to better understand the relationship be-

tween SSLDR and total number of colonoscopies performed through the course of a multi-year fellowship.

Patients and methods

After receiving approval from the institutional review board of the University of New Mexico Health Sciences Center (Ref. 18-355), we conducted a retrospective, longitudinal analysis of all average-risk colonoscopies conducted by the Division of Gastroenterology and Hepatology at the University of New Mexico Health Sciences Center from January of 2008 through December of 2020. The Albuquerque, New Mexico-based Division of Gastroenterology and Hepatology is an academic research center that cares for more than 10,000 patients annually across numerous clinics. The program offers a three-year fellowship program with an optional fourth year for advanced endoscopy specialization. At the time of this analysis, the Division used white light with high-definition (HD) endoscopes with the ability to switch to narrow band imaging on selected cases. The Division switched to HD endoscopes in 2008. As such, no cases included in this study were conducted using standard definition colonoscopes. No ancillary techniques such as underwater immersion colonoscopies were performed in the Division during the study period. Similarly, no distal attachments or caps were used. For withdrawal technique besides standard withdrawal, second forward view was used in most cases. The practice of retroflexion in the right colon to examine proximal folds has increased during the past five years but is still not widely adopted by gastroenterologists across the country and was used only in a minority of our study cases. Documentation of these technologies and techniques were sporadic or were not machine-readable in colonoscopy reports and were therefore excluded from this analysis.

Study sample and variable selection

Average-risk patient colonoscopies were identified through billing data and included codes: G0121 (Average risk screening); 45378-33 (colonoscopy with modifier 33 indicating a preventive service); ICD-9 code V76.51; or ICD-10 code Z12.11. Billing data were joined to electronic health record (EHR) and pathology record data using a clinical encounter number. Pathology records for the Division are created by attending pathologists, most of whom, but not all, are gastrointestinal pathologists. Pathology records were queried for any observations of SSL to capture the evolution of the pathological nomenclature used over the 13-year study period. This search included any mention in a pathology report for: sessile serrated lesion, sessile serrated adenoma, sessile serrated polyp, SSA, SSL, or sessile serrated polyp (SSP). Proximal hyperplastic polyps were not included in the query. Although the evidence base now suggests that many SSLs were historically misclassified as hyperplastic polyps, histopathological distinction did not emerge until later in our study period. Reviewing and reclassifying all previously diagnosed hyperplastic polyps in our data warehouse from 2008 onward as either SSL or confirmed hyperplastic polyps was outside of the scope of this study. SSLpositive pathology records were then manually reviewed to

confirm positive SSL identification and to rule out instances of negation (e.g. "not SSL"). EHR data were used to obtain patient demographics including sex, age, race, and ethnicity. Only patients aged 50 years and older were included in the analysis. Division administrative data were used to identify if the performing endoscopist was an attending alone or a fellow. (All fellows perform colonoscopies under the supervision of an attending.) Due to a change in billing record vendors, fellows could not be identified prior to 2014 and were excluded from the analysis from 2008 through 2013, apart from a single fellow who became an attending in 2013. Any endoscopist with fewer than 50 qualified screening colonoscopies over the course of 1 calendar year was excluded from that year's data. Due to numerous changes in software and EHR vendors during the 13-year study period, withdrawal times were not available or were not machine-readable for all eligible colonoscopies and were, therefore, excluded from this analysis.

Statistical analysis

We began with unadjusted, descriptive analyses of SSLDR at the endoscopist and Division levels over the 13-year study period. Unadjusted SSLDRs were calculated for individual endoscopists by year. Next, given the binary outcome variable of SSL detection and the longitudinal and hierarchical nature of the dataset (i.e., colonoscopies could be performed by a particular attending or a fellow or an attending who had been a fellow during the 13-year study period), a mixed effects logistic regression was used. The model estimated the log odds of SSL detection, exponentiated to odds ratios, as a linear combination of fixed and random predictor variables. One key fixed variable was year, and the included random effect term estimated a per-physician effect in order to account for their performing multiple colonoscopies over the study period. Two models were utilized. The first model used all colonoscopies and endoscopists to estimate Division-wide predictors of SSL detection. The second model focused exclusively on colonoscopies performed by fellows to determine if and to what degree SSLDR improved during their multi-year fellowships. Model-adjusted SSLDRs were then estimated for all years and across both categories of all endoscopists and fellows. All analyses were performed using Stata Statistical Software: Release 17 (College Station, TX).

Results

Our pooled data for all 13 years of the study included 36,467 average-risk screening colonoscopies. Table 1 presents the patient and endoscopist characteristics for these colonoscopies. The number of colonoscopies performed annually were relatively stable from 2010 through 2019 but notably decreased by nearly one-third in 2020, likely a result of patients seeking fewer elective procedures during the SARS-CoV-2 pandemic. The average age and percentage of patients who were male were relatively stable throughout the study period, hovering around 60 years and 45%, respectively. Finally, the percentage of colonoscopies performed by fellows versus attendings varied significantly during the period where data on fellows were

available (2014–2020), from a low of 47.1% in 2017 to a high of 67.6% in 2019, with no apparent trend (**►Table 1**).

Total SSLs detected on an annual basis increased steadily throughout the study period, from a low of eight in 2008 to 255 in 2017. SSLs detected appear to drop off to 150 in 2020, but taking into account the significantly lower denominator of total colonoscopies performed that year due to SARS-CoV-2, the unadjusted SSLDR is the second highest in 2020 at 7.37%. ► Fig. 1, which presents unadjusted SSLDR at the physician level, shows significant SSLDR variance across providers throughout the study period but also illustrates general improvement in SSLDR at both the physician and division levels. Most fellows, generally identifiable in ► Fig. 1 as those with only 2 or 3 years of data, show consistent year-over-year improvement in SSLDR.

▶ Table 2 presents the odds ratios (ORs) for SSL detection using all colonoscopies and endoscopists from the mixed effects logistic regression model. The odds of SSL detection were lower for first-year fellows compared to non-first-year fellows and all attendings, with first-year fellows having a 19% reduction in the odds of detecting SSL (OR: 0.81, 95% confidence interval [CI]: 0.67-0.96). Male patients had a 14% reduction in the odds of having SSL detected than females (OR: 0.86, 95% CI: 0.78–0.96). Each additional year of age for a patient yielded a moderately higher likelihood of SSL detection (OR: 1.01, 95% CI: 1.001–1.01). Compared to white patients, American Indian patients had a 61% reduction in the odds of having SSL detected, Asian patients, a 43% reduction, and Black patients, a 60% reduction. Non-Hispanics/Non-Latinos had 69% higher odds of an SSL detected than Hispanics/Latinos (OR: 1.69, 95% CI: 1.51–1.89). Finally, the probability of SSL detection increased significantly in every year compared to 2008. Most notably, the odds of having an SSL detected in 2020 were nearly 24 times higher than in 2008, controlling for all other variables (OR: 23.81, 95 % CI: 11.17-50.75). The non-zero random effect supports our utilization of a mixed model and suggests heterogeneity among endoscopists in detecting SSL. The reported random effects variance of 0.099 suggests that the best endoscopist had a roughly 2.6 higher odds of detection than the average endoscopist ($e^{(3 \times \sqrt{0.099})} = 2.6$) when controlling for all other predictors in the model.

Using the model's fixed and physician random effects, we created an adjusted, unit-level SSLDR by year with 95% confidence intervals (**> Fig. 2**). The adjusted SSLDR shows a consistent improvement in SSLDR from a low of 0.37% (95% CI: 0.10–0.63) in 2008 to a high of 7.94% (95% CI: 6.34–9.54) in 2020.

Fig. 3 presents changes in SSLDR for each fellow between their first and second year and between their second and third year. Fellows with teal plots improved their SSLDR from the lower bound in the preceding year to the upper bound in the following year. Fellows with blue plots decreased their SSLDR from the upper bound in the preceding year to the lower bound in the following year. These unadjusted results indicate more year-over-year improvement between the first and second year of the gastroenterology fellowship than the second and third year. In comparing all eligible fellows between their first and second years, 28 of 34 fellows (82.4%) had higher SSLDRs

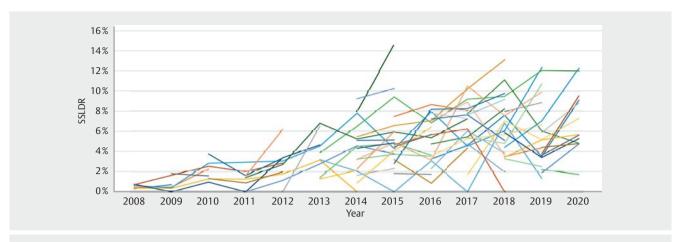


	2008	2009	2010	2011	2012	2013	2014	2015	2016	2017	2018	2019	2020	Total
Total screening colonoscopies	2,077	1,960	3,207	3,340	2,673	2,992	3,173	2,724	3,088	3,288	2,838	3,072	2,035	36,467
Colonoscopies with SSLs detected	8	20	80	50	77	140	157	159	186	255	204	215	150	1,701
SSL detection rate	0.4%	1.0%	2.5%	1.5%	2.9%	4.7%	4.9%	5.8%	6.0%	7.8%	7.2%	7.0%	7.4%	4.7 %
Patient characteristi	CS													
 Mean age (SD) 	60.0 (8.0)	59.7 (7.8)	58.8 (7.6)	59.3 (7.7)	59.6 (7.6)	59.5 (7.5)	59.4 (7.5)	60.0 (7.7)	60.1 (7.6)	61.1 (7.7)	61.7 (8.0)	61.5 (7.8)	62.1 (8.1)	60.2 (7.8)
 Percentage male 	41.4	42.6	41.0	44.6	42.8	44.6	45.4	45.7	44.0	45.6	46.2	47.6	46.0	44.5
Race [count (column	ı %)]													
 American Indian/Alaska Native 	131 (6.3)	115 (5.9)	185 (5.8)	231 (6.9)	167 (6.2)	184 (6.1)	200 (6.3)	231 (8.5)	232 (7.5)	229 (7.0)	208 (7.3)	219 (7.1)	164 (8.1)	2,496 (6.8)
Asian	46 (2.2)	59 (3.0)	87 (2.7)	110 (3.3)	71 (2.7)	109 (3.6)	98 (3.1)	83 (3.0)	123 (4.0)	125 (3.8)	95 (3.3)	111 (3.6)	57 (2.8)	1,174 (3.2)
Black/African American	40 (1.9)	51 (2.6)	86 (2.7)	99 (3.0)	68 (2.5)	91 (3.0)	102 (3.2)	74 (2.7)	90 (2.9)	93 (2.8)	73 (2.6)	74 (2.4)	56 (2.8)	997 (2.7)
 Unknown 	324 (15.6)	321 (16.4)	630 (19.6)	606 (18.1)	487 (18.2)	540 (18.0)	493 (15.5)	377 (13.8)	524 (17.0)	517 (15.7)	415 (14.6)	480 (15.6)	336 (16.5)	6,050 (16.6)
• White	1,536 (74.0)	1,414 (72.1)	2,219 (69.2)	2,294 (68.7)	1,880 (70.3)	2,068 (69.1)	2,280 (71.9)	1,959 (71.9)	2,119 (68.6)	2,324 (70.7)	2,047 (72.1)	2,188 (71.2)	1,422 (69.9)	25,750 (70.6)
Ethnicity [count (col	umn %)]													
Hispanic/ Latino	634 (30.5)	693 (35.4)	1,315 (41.0)	1,332 (39.9)	1,128 (42.2)	1,265 (42.3)	1,325 (41.8)	1,109 (40.7)	1,341 (43.4)	1,352 (41.1)	1,145 (40.3)	1,215 (39.6)	838 (41.2)	14,692 (40.3)
 Not Hispanic/ Latino 	877 (42.2)	829 (42.3)	1,329 (41.4)	1,548 (46.3)	1,391 (52.0)	1,563 (52.2)	1,645 (51.8)	1,443 (53.0)	1,556 (50.4)	1,741 (53.0)	1,509 (53.2)	1,622 (52.8)	1,030 (50.6)	18,083 (49.6)
 Unknown 	566 (27.3)	438 (22.3)	563 (17.6)	460 (13.8)	154 (5.8)	164 (5.5)	203 (6.4)	172 (6.3)	191 (6.2)	195 (5.9)	184 (6.5)	235 (7.6)	167 (8.2)	3,692 (10.1)
Endoscopist rank [co	unt (colu	mn %)]												
 Performed by attending 	2,077 (100)	1,960 (100)	3,207 (100)	3,240 (100)	2,618 (97.9)	2,992 (100)	1,659 (52.3)	1,147 (42.1)	1,439 (46.6)	1,740 (52.9)	1,111 (39.2)	996 (32.4)	858 (42.2)	25,144 (69.0)
 Performed by fellow 	0 (0.0)	0 (0.0)	0 (0.0)	0 (0.0)	55 (2.1)	0 (0.0)	1,515 (47.7)	1,577 (57.9)	1,649 (53.4)	1,548 (47.1)	1,727 (60.9)	2,077 (67.6)	1,177 (57.8)	11,323 (31.1)

in their second year. Average unadjusted SSLDR for eligible fellows grew from 4.34% in the first year of their fellowship to 5.79% in their second year, an absolute improvement of 1.45%. In comparing all eligible fellows between their second and third years, 13 out of 19 fellows (68.4%) had higher SSLDRs in their third year. Average unadjusted SSLDRs for all eligible fellows grew from 5.24% in the second year of their fellowship to 6.04% in their third year, an absolute improvement of only 0.80%. ▶ Fig. 3 also illustrates the relatively high variance of SSLDRs across fellows at all points of time.

▶ Table 3 presents the ORs for SSL detection using only fellow-performed colonoscopies from a second mixed effects lo-

gistic regression model. Results confirmed the analysis of the unadjusted trends across years of fellowship (▶ Fig. 3), with the odds of SSL detection being significantly lower during a fellow's first year compared to their second year (OR: 0.80, 95% CI: 0.66–0.98) but not significantly higher in their third year compared to their second year (OR: 1.09, 95% CI: 0.85–1.4). With fellows only, male patients had a 16% reduction in the odds of having an SSL detected (OR: 0.84, 95% CI: 0.71–0.99), American Indians and Black patients were less likely than Whites to have SSL detected (OR: 0.43, 95% CI: 0.30–0.62, and OR: 0.48, 95% CI: 0.27–0.85, respectively), and Non-Hispanics/Non-Latinos had 94% higher odds to have SSL detected



▶ Fig. 1 Unadjusted physician-level sessile serrated lesion detection rate, 2008–2020. Each line represents the sessile serrated lesion detection rate of an individual physician.

than Hispanics/Latinos (OR: 1.94, 95% CI: 1.61–2.33). Using 2014 as a referent, fellows were no more or less likely to detect SSLs during any of the subsequent years.

Using the model's fixed and physician random effects, we created an adjusted SSLDR by year of fellowship with 95% CIs (▶ Fig. 4). The adjusted SSLDR for fellows for each year of the fellowship program showed significant improvement in SSLDR from 5.05% (95% CI: 4.24–5.86) during the first year of fellowship to 6.73% (95% CI: 5.24–8.22) during the third year of fellowship (95% CI of difference: −0.033 − −0.001; *P*=0.038).

Discussion

To continue the downward trajectory of CRC incidence and mortality rates, the quality, and not just the quantity, of CRC screening colonoscopies must improve. A conspicuous area for quality improvement is the proper detection and removal of serrated lesions, which account for approximately one-third of all CRC and a disproportionately high percentage of post-colonoscopy interval cancers [4,5]. However, guidelines and training for proper detection and removal of serrated lesions have proven more difficult and uniquely different than those of conventional adenomas. While the field continues to move toward determining the true prevalence of serrated lesions across diverse patient populations, a call has been made to focus on local SSLDR and use the highest rate from a unit as the local aspirational target [21]. In this study, we examined the SSLDR of a large, academic gastroenterology division from 2008 to 2020 and modeled such a local, aspirational target for SSLDR. We also assessed the progression of SSLDR among gastroenterology fellows to better understand the relationship between SSLDR and total colonoscopies performed during a multi-year fellowship.

We found that SSLDR increased steadily and significantly throughout our study period, from a low of 0.37% in 2008 to a high of 7.94% in 2020. The SSLDR from 2020 of 7.94% should serve as the local aspirational target for the Gastroenterology Division at the University of New Mexico Health Sciences Center per Kahi & Rex's recommendation.[21] Our SSLDRs by year are

likely similar to results from other SSLDR studies when taking the year of observation into consideration. In two recent metaanalyses, the authors found lower SSLDR (estimated across all qualifying studies) than our results. However, their analyses included over a dozen SSLDR studies that were published as far back as 2010 and used even earlier data [11, 20]. Our model notably estimated that the odds of detecting an SSL in 2020 were nearly 24 times higher than in 2008, controlling for all other covariates. Our pooled SSLDR (averaged across all years of our study) of 4.66% is closer to Huang et al.'s SSLDR of 2.5% (95% CI: 1.5-3.8) [20] and Desai et al.'s SSLDR of 2.5% (95% CI: 1.8-3.4) [11], though this pooled figure is arguably of less significance for targeting purposes than our SSLDRs observed in, say, 2019 or 2020. This finding, combined with the observation that our study's SSLDR trend does not appear to be flattening by 2020, serves as an important consideration for policymakers and authors of future meta-analyses that the year of observation is a possible, even likely, driver of SSLDR and weighting by year ought to be considered. Our study's local aspirational target SSLDR of 7.94% falls well within the range of SSLDR observed in other individual studies [22-25] and is remarkably similar to the 8.1% observed in a study that focused solely on an individual endoscopist and pathologist with notably high SSLDR [16].

Our year indicator variable also likely serves as a proxy, catch-all measure for the true drivers of the observed, sustained improvement of SSLDR during our study period. Although this study was unable to isolate and assign singular dates for all the minor and major changes related to fellowship training, attending awareness, histopathological distinctions, or technological advancement, it is likely that each of these contributed in some part to the observed improvements in SSLDR. Indeed, studies with much narrower study periods have linked numerous endoscopic interventions to improved SSLDR [26].

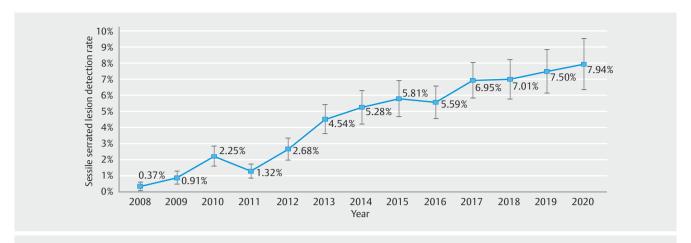
Our study also found that SSLDR for gastroenterology fellows increased significantly between the first and second year of the gastroenterology fellowship program but did not further

▶ Table 2 Mixed effects logistic regression results: odds ratios for sessile serrated lesion detection.

Fixed effects	Odds ratio	Standard error	P>z	95% CI	
Attending vs. fellow					
Attending-performed	REF	REF	REF	REF	REF
Fellow-performed	0.934	0.094	0.497	0.768	1.137
First-year fellow	0.805	0.074	0.018	0.673	0.964
Patient sex					
• Female	REF	REF	REF	REF	REF
• Male	0.859	0.044	0.003	0.777	0.949
Patient age	1.008	0.003	0.017	1.001	1.014
Patient race					
• White/Anglo	REF	REF	REF	REF	REF
American Indian	0.385	0.051	0.000	0.296	0.500
• Asian	0.568	0.090	0.000	0.416	0.775
Black/African American	0.399	0.082	0.000	0.266	0.597
 Unavailable 	0.918	0.074	0.287	0.785	1.074
Patient ethnicity					
Hispanic/Latino	REF	REF	REF	REF	REF
Not Hispanic/Latino	1.687	0.098	0.000	1.505	1.890
 Unavailable 	1.256	0.135	0.033	1.018	1.550
Year					
2008	REF	REF	REF	REF	REF
2009	2.495	1.060	0.031	1.085	5.737
2010	6.262	2.410	0.000	2.945	13.31
2011	3.626	1.436	0.001	1.668	7.882
2012	7.523	2.917	0.000	3.518	16.08
2013	13.030	4.959	0.000	6.180	27.47
2014	15.299	5.867	0.000	7.215	32.44
2015	16.961	6.504	0.000	7.999	35.96
2016	16.256	6.209	0.000	7.690	34.36
2017	20.573	7.797	0.000	9.788	43.24
2018	20.765	7.915	0.000	9.837	43.83
2019	22.371	8.552	0.000	10.575	47.32
• 2020	23.813	9.193	0.000	11.174	50.74
Constant	0.002	0.001	0.000	0.001	0.005
Physician random effect (Variance of random intercepts)	0.099	0.032		0.052	0.186

increase in the third year. Alternatively, our initial model with all endoscopists suggested that first-year fellows are less likely to detect SSL than all other endoscopists. Although we can find no similar studies that focus solely on SSLDR among gastroenterol-

ogy fellows or similar training programs, our findings generally align with other research that shows adenoma and polyp detection rates positively correlate with colonoscopy volume [24, 27–31]. This finding highlights the importance of high-quality



▶ Fig. 2 Adjusted sessile serrated lesion detection rate by year with 95% confidence intervals. Each marker represents the model-adjusted, division-level sessile serrated lesion detection rate including its 95% confidence interval.



▶ Fig. 3 Sessile serrated lesions detection rate by fellow: first year vs. second year (top) and second year vs. third year (bottom). Each column represents the year-over-year percentage change in sessile serrated lesion detection rate (SSLDR) of an individual fellow. Striped columns represent positive year-over-year change (improvement) is SSLDR while checkered columns represent negative year-over-year change (diminishment).

colonoscopy training, particularly for new gastroenterology fellows. Our results suggest that at least during a fellow's first year

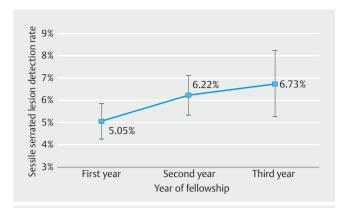
of performing colonoscopy, additional "at-the-elbow" guidance by SSL-attuned attendings may be warranted.

► Table 3 Mixed Effects Logistic Regression Results: Odds Ratios for Sessile Serrated Lesion Detection among Fellows Only.

Fixed effects	Odds ratio	Standard error	P>z	95 % CI	
Fellowship year					
First year	0.800	0.082	0.030	0.654	0.97
 Second year 	REF	REF	REF	REF	REF
Third year	1.089	0.141	0.509	0.845	1.40
Patient sex					
• Female	REF	REF	REF	REF	REF
• Male	0.838	0.069	0.032	0.714	0.98
Patient Age	1.002	0.005	0.705	0.992	1.01
Patient Race					
• White/Anglo	REF	REF	REF	REF	REF
American Indian	0.428	0.080	0.000	0.297	0.61
• Asian	0.700	0.153	0.103	0.456	1.07
Black/African American	0.480	0.139	0.011	0.272	0.84
 Unavailable 	1.038	0.139	0.782	0.799	1.34
Patient ethnicity					
Hispanic/Latino	REF	REF	REF	REF	REF
Not Hispanic/Latino	1.937	0.181	0.000	1.613	2.32
 Unavailable 	1.125	0.226	0.558	0.758	1.66
Year					
2014	REF	REF	REF	REF	REF
2015	1.138	0.206	0.475	0.798	1.62
2016	1.108	0.211	0.589	0.764	1.60
2017	1.190	0.240	0.389	0.801	1.76
2018	0.979	0.203	0.919	0.652	1.47
2019	0.982	0.202	0.928	0.656	1.46
• 2020	1.068	0.241	0.771	0.686	1.66
Constant	0.052	0.019	0.000	0.025	0.10
Physician random effect (variance of random intercepts)	0.076	0.035		0.031	0.18

While there have been numerous evaluations of SSLDR, few have included the total number of years and colonoscopies that we included in this study. In addition, to our knowledge, our study marks the first time that SSLDR has been studied in the context of a gastroenterology fellowship program to better understand the relationship between SSLDR and colonoscopy training during a three-year fellowship program. However, this study is not without its limitations. The lengthy study period, which served as a feature for reasons stated above, also prevented the inclusion of numerous variables. During the 13 years studied, UNM HSC and the Gastroenterology Division imple-

mented and replaced numerous information technology systems. Changes in data standards, clinical terminology, and billing requirements resulted in a mélange of variables, few of which were available in each of the study's 13 years. As a result, we were unable to include key indicator variables related to the patient that are known to correlate with CRC including family history, patient weight, and smoking status, among others. Additionally, due to changes in the Gastroenterology Division's clinical software, clinical observation variables related to the colonoscopy such as withdrawal time, bowel preparation, and cecal intubation were not uniformly available or were not ma-



▶ Fig. 4 Adjusted sessile serrated lesion detection rate by fellowship year with 95% confidence interval. Each marker represents the model-adjusted, division-level sessile serrated lesion detection rate among fellows only including its 95% confidence interval.

chine-readable. Similar changes in clinical software for the pathology department precluded us from being able to track the ever-changing composition of the pathology department and its review processes. Finally, the technologies used by the Gastroenterology Division were upgraded numerous times during the study period (e.g., zoom magnification and narrow band imaging). Many of these technologies undoubtedly improved the ability to detect and resect SSLs [26, 32]. However, newer scopes and monitors were never replaced en masse, but were instead replaced on a rolling basis across months and even years in some cases. As a result, we were unable to account for these changes in the model. However, due to the continuous rotation of fellows and attendees across clinics and endoscopy suites within the division, we believe that these differing technologies were likely equally diffused across endoscopists over time. Future research should incorporate more of these patient- and technology-level variables to control for potential confounding more comprehensively.

Implications for practice

SSLDR is becoming an increasingly recognized colonoscopy quality indicator making for important healthcare implications in the screening and detection rates of these lesions. It is obvious now that national benchmarks for SSLDRs need to be established. High-quality randomized prospective studies, including patient demographics, indication, standardized prep, withdrawal time, and physician training will help identify valid benchmarks eventually. Fellowship and endoscopy training programs should also continuously reappraise the evidence base for various endoscopic advances including add-on devices and endocuff vision [26]. Our study demonstrated an increasing SSLDR with over time and within a three-year fellowship experience. Previous studies have similarly shown a stepwise increase in adenoma detection rate with advancing years of gastroenterology fellowship training compared to procedures performed by an attending physician alone [30, 31]. Trainee education at all levels should include the importance of detection

of SSLs and their essential role in colorectal cancer pathogenesis.

Educational initiatives for new gastroenterology fellows should be implemented to improve SSLDR such as endoscopic image dataset and atlas image review, close one-to-one fellow to attending supervision, education of endoscopy nurses about SSLDRs, video tape analysis of entire colonoscopy, and mandating longer withdrawal times based on level of training [30, 31]. SSLDR-specific educational initiatives should also include training of non-gastroenterology physicians performing colonoscopies, including surgeons and family physicians. Future improvement in endoscopic visualization technology along with artificial intelligence-based algorithms for SSL image base recognition will also help improve the detection of SSLs.

Acknowledgement

The authors would like to thank the following individuals for their numerous contributions to this study: Rosa Matonti, DNP, RN; Daniel De Francisco Cabral; Sergio A. Sánchez-Luna, MD, ABOM-D; David R, Martin, MD; and Richard Feddersen, MD.

Competing interests

The authors declare that they have no conflict of interest.

Funding

This work was supported by the National Cancer Institute of the National Institutes of Health, grants R01CA192967 (Mishra, PI) and 3P30CA118100-16S4 (Tomkinson PI, Mishra, PD), and the Biostatistics Shared Resource of the University of New Mexico Comprehensive cancer Center. USA

References

- [1] Siegel RL, Miller KD, Fuchs HE et al. Cancer statistics, 2021. CA: Cancer | Clinicians 2021; 71: 7–33
- [2] National Cancer Institute. Surveillance, Epidemiology, and End Results Program Cancer Stat Facts: Colorectal Cancer. In: Surveillance E, and End Results Program ed. 2021
- [3] Shaukat A, Kahi CJ, Burke CA et al. ACG Clinical Guidelines: Colorectal Cancer Screening 2021. Am J Gastroenterol 2021; 116: 458–479
- [4] Kahi CJ. Screening relevance of sessile serrated polyps. Clin Endosc 2019; 52: 235
- [5] Kim SY, Kim TI. Serrated neoplasia pathway as an alternative route of colorectal cancer carcinogenesis. Intest Res 2018; 16: 358
- [6] Karnes WE, Johnson DA, Berzin TM et al. A polyp worth removing: a paradigm for measuring colonoscopy quality and performance of novel technologies for polyp detection. J Clin Gastroenterol 2021; 55: 733–739
- [7] Shapiro JA, Bobo JK, Church TR et al. A comparison of fecal immunochemical and high-sensitivity guaiac tests for colorectal cancer screening. Am J Gastroenterol 2017; 112: 1728
- [8] Chang L-C, Shun C-T, Hsu W-F et al. Fecal immunochemical test detects sessile serrated adenomas and polyps with a low level of sensitivity. Clin Gastroenterol Hepatol 2017; 15: 872–879.e871

- [9] Anderson JC, Robertson DJ. Serrated polyp detection by the fecal immunochemical test: an imperfect FIT. Clin Gastroenterol Hepatol 2017; 15: 880–882
- [10] Kim DH, Matkowskyj KA, Lubner MG et al. Serrated polyps at CT colonography: prevalence and characteristics of the serrated polyp spectrum. Radiology 2016; 280: 455–463
- [11] Desai M, Anderson JC, Kaminski M et al. Sessile serrated lesion detection rates during average risk screening colonoscopy: A systematic review and meta-analysis of the published literature. Endosc Int Open 2021; 9: E610–E620
- [12] Rameshshanker R, Purchiaroni F, Ana W et al. PTH-039 Prevalence of sessile serrated adenomas/polyps in distal colon during screening colonoscopy/flexible sigmoidoscopy: a single bowel cancer screening experience from uk. Gut 2017; 66: A225
- [13] Pohl H, Srivastava A, Bensen SP et al. Incomplete polyp resection during colonoscopy – results of the complete adenoma resection (CARE) study. Gastroenterology 2013; 144: 74–80.e71
- [14] Torlakovic E, Skovlund E, Snover DC et al. Morphologic reappraisal of serrated colorectal polyps. Am J Surg Pathol 2003; 27: 65–81
- [15] Bettington M, Walker N, Clouston A et al. The serrated pathway to colorectal carcinoma: current concepts and challenges. Histopathology 2013; 62: 367–386
- [16] Abdeljawad K, Vemulapalli KC, Kahi CJ et al. Sessile serrated polyp prevalence determined by a colonoscopist with a high lesion detection rate and an experienced pathologist. Gastrointest Endosc 2015; 81: 517–524
- [17] Kanth P, Boylan KE, Bronner MP et al. Molecular biomarkers of sessile serrated adenoma/polyps. Clin Translat Gastroenterol 2019; 10: e00104
- [18] Hetzel JT, Huang CS, Coukos JA et al. Variation in the detection of serrated polyps in an average risk colorectal cancer screening cohort. Am J Gastroenterol 2010; 105: 2656–2664
- [19] Kahi CJ, Hewett DG, Norton DL et al. Prevalence and variable detection of proximal colon serrated polyps during screening colonoscopy. Clin Gastroenterol Hepatol 2011; 9: 42–46
- [20] Huang J, Chan PS, Pang TW et al. Rate of detection of serrated lesions at colonoscopy in an average-risk population: a meta-analysis of 129,001 individuals. Endosc Int Open 2021; 9: E472–E481

- [21] Kahi CJ, Rex DK. Sessile serrated lesions: Searching for the true prevalence. Endosc Int Open 2021; 9: E635–E636
- [22] Ross WA, Thirumurthi S, Lynch PM et al. Detection rates of premalignant polyps during screening colonoscopy: Time to revise quality standards? Gastrointest Endosc 2015; 81: 567–574
- [23] IJspeert J, Nolthenius CT, Kuipers E et al. CT-colonography vs. colonoscopy for detection of high-risk sessile serrated polyps. Am J Gastroenterol 2016; 111: 516–522
- [24] Shaukat A, Gravely AA, Kim AS et al. Rates of detection of adenoma, sessile serrated adenoma, and advanced adenoma are stable over time and modifiable. Gastroenterol 2019; 156: 816–817
- [25] Buda A, De Bona M, Dotti I et al. Prevalence of different subtypes of serrated polyps and risk of synchronous advanced colorectal neoplasia in average-risk population undergoing first-time colonoscopy. Clin Translat Gastroenterol 2012; 3: e6
- [26] Aziz M, Fatima R, Lee-Smith W et al. Comparing endoscopic interventions to improve serrated adenoma detection rates during colonoscopy: a systematic review and network meta-analysis of randomized controlled trials. Eur J Gastroenterol Hepatol 2020; 32: 1284–1292
- [27] Qayed E, Vora R, Levy S et al. Colonoscopy procedural volume increases adenoma and polyp detection rates in gastroenterologytrainees. World | Gastrointest Endosc 2017; 9: 540–551
- [28] Pace D, Borgaonkar M, Lougheed M et al. Effect of colonoscopy volume on quality indicators. Can J Gastroenterol Hepatol; 2016; 2580894
- [29] Zwink N, Stock C, Birkner B et al. Screening colonoscopy volume and detection of colorectal neoplasms: a state-wide study from Bavaria, Germany. Europ | Cancer Prev 2017; 26: 181–188
- [30] Peters SL, Hasan AG, Jacobson NB et al. Level of fellowship training increases adenoma detection rates. Clin Gastroenterol Hepatol 2010; 8: 439–442
- [31] Rogart JN, Siddiqui UD, Jamidar PA et al. Fellow involvement may increase adenoma detection rates during colonoscopy. Am J Gastroenterol 2008; 103: 2841–2846
- [32] Aziz M, Mehta TI, Weissman S et al. Do water-aided techniques improve serrated polyp detection rate during colonoscopy? A systematic review with meta-analysis | Clin Gastroenterol 2021; 55: 520–527