

Quality of life after laparoscopic totally extraperitoneal repair of an asymptomatic inguinal hernia

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Abstract

Background When considering an asymptomatic inguinal hernia, surgeons must weigh the risks of watchful waiting against the risk of operative complications. Laparoscopy offers the benefit of reduced postoperative pain, which, for appropriate surgical candidates, may strengthen the case for repair. This study compares general and disease-specific quality of life following totally extraperitoneal (TEP) laparoscopic inguinal hernia repair (LIHR) of asymptomatic and symptomatic hernias.

Methods We summarize prospective data from 387 patients who underwent TEP LIHR between 2009 and 2015 by four surgeons at a single institution. Asymptomatic individuals were identified by pain scores of zero at pre-operative clinic visits. Validated quality of life (QOL) measurements were administered preoperatively and at 3 weeks, 6 months, and 1-year postop. Comparisons were made using Chi-square test, *t* test, or Mann–Whitney U

test. Changes over time were assessed using longitudinal mixed effects models.

Results A cohort of 79 asymptomatic cases were compared to 308 symptomatic individuals. The asymptomatic cohort had larger median hernia defects (2.5 vs 2 cm, $p < 0.01$), was older (mean 63.0 vs 58.9 years, $p = 0.03$), included fewer indirect hernias (57.7 vs 74.9%, $p < 0.01$), took pain medication for fewer days (mean 1.2 ± 1.5 vs 2.2 ± 3.0 days, $p = 0.02$), returned to baseline activities of daily living earlier (median 3 vs 5 days, $p < 0.01$), and reported decreased postoperative pain ($p = 0.02$). There was no significant difference in general QOL. There was one recurrence in the asymptomatic group and were two in the symptomatic cohort.

Conclusions Asymptomatic individuals undergoing TEP LIHR reported less postoperative pain, returned to baseline activities, and discontinued pain medication sooner than symptomatic patients. These results are encouraging and may inform patient-centered discussions about asymptomatic hernia repair.

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Symptoms in the presence of inguinal hernia vary widely from completely asymptomatic to disabling pain [1, 2]. Lifetime risk for developing an inguinal hernia is 27% for men and 3% for women, and inguinal hernia repair (IHR) is the most commonly performed operation in the United States, with an incidence of 28 per 100,000 individuals [3, 4]. IHR has a number of well-known early and late surgical complications. Early complications include urinary retention, hematoma, and surgical site infection. Among late complications are recurrence, chronic pain,

numbness, and fertility problems for men [5]. Chronic pain is generally defined as pain persisting beyond 3 months, and up to one-third of individuals will report some degree of chronic discomfort after IHR when asked on anonymous surveys. This figure drops to 10% when asked in clinic, with 3% experiencing pain significant enough to interfere with daily life [6]. Fertility problems can arise from injury to the spermatic cord in males. Unilateral injury to the vas deferens occurs in up to 0.3% of adult repairs. Injury of the testicular artery occurs in up to 0.5% and can lead to testicular atrophy [7]. As with any surgery, IHR carries a risk of death. A Danish study on over 26,000 hernia repairs found mortality rate to be 0.02% for those under 60 years of age, and 0.48% for those over [8].

The advent of laparoscopy brought new surgical approaches to IHR, and rigorous analysis comparing laparoscopic techniques to open has determined that the laparoscopic approach offers less postoperative pain, faster return to daily activities, and overall fewer complications [9, 10]. The laparoscopic repair has also been associated with less chronic pain than open repair [11]. On the other hand, the laparoscopic approach is more expensive and requires longer operative time [9, 10]. Both approaches are considered acceptable, and The National Institute for Health and Clinical Excellence recommends that patients should be educated to the risks and benefits carried by both techniques prior to surgery [12].

Given the rare but significant operative risks, recommending IHR must be approached on a case-by-case basis. Surgery is generally recommended for individuals experiencing symptoms when impact on quality of life seems to outweigh the small rate of complications. However, the question becomes more complicated when considering asymptomatic hernias. In the era of open surgery, a ‘watchful waiting’ approach was considered appropriate management for asymptomatic hernias. Watchful waiting carries its own risks; incarcerated and strangulated hernias are surgical emergencies, with delay being potentially fatal. While the risk of incarceration is low, at just 0.55% over 4 years, emergent repair is associated with increased morbidity and mortality, and watchful waiting is associated with a 1.59 odds-ratio of having emergency hernia surgery [13, 14]. Additionally, up to 68% of individuals with asymptomatic hernias eventually develop symptoms and end up having surgery at an older age [15]. With advancing technology and decreasing complication rates, the watchful waiting approach has been called into question, and several studies have compared watchful waiting to elective repair. Individual studies have failed to show a significant difference in chronic pain, and both approaches have been deemed safe forms of management [16–18]. This study is a review of an institutional database aimed to examine outcomes and quality of life (QOL) after laparoscopic inguinal

hernia repair (LIHR) using totally extraperitoneal (TEP) technique.

Methods

A prospectively collected database of all patients undergoing hernia repair between 2009 and 2015 was queried for those who underwent LIHR by TEP technique. Elective operations on primary, recurrent, unilateral, and bilateral inguinal hernias were included. Tacks were used for all cases of mesh fixation. General guidelines for fixation were direct hernias >2 cm and indirect hernias >6 cm, although some surgeons in the group use fixation more liberally. Hernia defect size was estimated intraoperatively using an open laparoscopic grasper as a reference. The asymptomatic cohort was populated with individuals who orally reported a pain score of zero to their surgeon at their preoperative clinic visit. Patients were asked to complete quality of life questionnaires preoperatively, and postoperatively at 3 weeks, 6 months and 1 year.

Health-related quality of life assessment tools

In total, three comprehensive quality of life instruments, encompassing both generic and surgery specific elements, were distributed to patients. The Short Form 36 Health Survey Version 2 (SF-36) commonly used to assess patients’ overall health status at any given time, consists of 36 total items divided amongst eight different categories. Responses are scored on a Likert scale [19]. Value of this scale in a surgical setting is limited, however, given the acute changes around the time of surgery are not reflected in this survey.

The Surgical Outcomes Measurement System (SOMS) is a relatively new comprehensive collection of measures specifically assessing patients in the postoperative state, regardless of the type of operation they had. Similar to SF-36, SOMS encompasses 34 items included within seven quality of life domains: pain on a visual analog scale (VAS), pain impact, pain quality, fatigue, physical functioning, body image, and satisfaction. Despite similarities to SF-36, SOMS is more specific to the surgical patient.

The Carolinas Comfort Scale (CCS), the most commonly used validated QOL instrument for hernia surgery, consists of 8 questions specific to pain, movement limitations, and the sensation of mesh experienced after hernia repair. Each question is scored on a 6-point Likert scale (0 = No symptoms, 2 = Mild and bothersome, and 5 = Disabling symptoms) where the best possible score is 0 [20].

Over the course of the study, the QOL instruments administered varied slightly as new measures were added.

Since 2009, SF-36 has been distributed to all patients pre- and postoperatively. In 2011 and 2012, CCS and SOMS were respectively added to the study protocol. CCS has only been administered postoperatively due to the nature of questions pertaining to mesh sensation. The SOMS tool, in its entirety, was given preoperatively and a modified version—SOMS Short Form, limited to the domains of Body Image, Pain Impact, Pain Quality, and Satisfaction, given postoperatively since being acquired.

Statistical methods

Data are presented as frequency count with percentage for categorical variables and mean with standard deviation or median with interquartile range for continuous variables. Patient demographics, intra-operative characteristics, postoperative outcomes, and QOL assessments were compared between asymptomatic and symptomatic hernia patients. Categorical variables were compared using Chi-square test or Fisher's exact test (for small cell size), and continuous variables were compared using *t* test or Mann–Whitney U test (nonparametric). Change in QOL assessments over time was assessed using mixed-effects linear regression models with an unstructured covariance structure controlling for random effects of intercept. An interaction term between group (symptomatic vs asymptomatic) and time was included to assess differential effects of QOL across time by group, with interactions considered significant at $p < 0.05$. Additional analyses adjusted for age. Statistical significance was established through at $p < 0.05$. All statistical analyses were performed using SAS 9.3 statistical software (SAS Inc. Cary, NC).

Results

Of 638 TEP LIHR patients in the database from 2009 to 2015, 387 had the necessary data available after chart review. A cohort of 79 patients (20.4%) reported a pain score of 0 at their preop clinic visit and were compared to 308 symptomatic individuals. Demographic comparisons are shown in Table 1. The groups had a similar gender distribution and incidence of comorbidities. The asymptomatic cohort was older than the symptomatic cohort (63.0 vs 58.9 years, $p = 0.03$). Intraoperative findings (Table 2) revealed the asymptomatic cohort had larger median hernia defects (2.5 vs 2 cm, $p < 0.01$) and included fewer indirect hernias (57.7 vs 74.9%, $p < 0.01$). Operative time and hernia laterality were similar. Mesh fixation was used in about 55% of cases, and tacks were the consistently used method. Analysis of postoperative variables (Table 3) showed that the asymptomatic patients took pain medication for fewer days on average (1.2 ± 1.5 vs

2.2 ± 3.0 days, $p = 0.02$), and returned to baseline activities of daily living earlier (median 3 vs 5 days, $p < 0.01$). There was one recurrence in the asymptomatic group, and two in the symptomatic group. All three of the recurrences occurred in individuals with indirect hernias, and two of the three had mesh fixation with tacks at the time of surgery.

46.5% of individuals who reported zero pain in clinic reported pain on the VAS pain score included in the SOMS questionnaire preoperatively, with an average score of 1.30. The average clinic pain score of the symptomatic group was higher than the SOMS reported score (3.35 vs 2.74, $p < 0.01$). The asymptomatic group's preop SOMS VAS score remained significantly different from the symptomatic group's (1.30 vs 2.74, $p < 0.01$). Over time, SOMS VAS score remained significantly different at 3 weeks, but not at 6 months or 1 year (Table 4).

There were no significant differences in postoperative complications between the cohorts. Rates of seroma, hematoma, and surgical site infection are within range of rates reported in the literature. Rate of urinary retention has been reported from 1.3 to 5.8% in prior studies [21]. In this analysis, the asymptomatic and symptomatic cohorts respectively experienced urinary retention at a rate of 8.6 and 6.0%. The higher rate seen in the asymptomatic cohort may also be explained by the older age of this group.

Longitudinal mixed-effects models indicated CCS Pain, Movement, and Total scores significantly improved over time. Asymptomatic patients had significantly more favorable CCS Pain scores compared to symptomatic patients [estimate (standard error): -1.84 (0.90), $p = 0.04$]; however, this significance was eliminated after adjusting for age [-1.37 (0.89), $p = 0.13$]. All SF-36 domains (physical functioning, role physical and emotional, energy and fatigue, well-being, social functioning, pain, and general health) were significantly improved over time, but there was no difference between asymptomatic and symptomatic patients. SOMS pain impact, pain quality, and fatigue were all significantly improved over time. Asymptomatic patients had significantly more favorable pain impact and pain quality scores compared to symptomatic patients, even after adjustment for age [estimate (se): -1.16 (0.49), $p = 0.02$; -1.03 (0.40), $p = 0.01$, respectively]. Table 5 presents results of mixed-effects models.

20% of patients who had TEP LIHR reported no pain at their preop clinic visit. Interestingly, nearly half of these reported some pain on the preop SOMS questionnaire, resulting in an average score of 1.3 on a 0–10 scale. The average pain score reported on SOMS by symptomatic individuals was significantly lower than the score reported in clinic. Figure 1 illustrates these discrepancies. Despite the preop SOMS pain scores being less divergent than the clinic scores, they remained significantly different over time until 6 months postoperatively (Fig. 2).

Table 1 Patient demographics and preoperative characteristics

Variables	Asymptomatic <i>N</i> (%)	Symptomatic <i>N</i> (%)	<i>p</i> value
Total # of patients	79 (20.4)	308 (79.6)	
Male gender	72 (91.1)	287 (93.2)	0.53
Age (years)—mean ± SD	63.0 ± 14.2	58.9 ± 15.6	0.03
BMI (kg/m ²)—mean ± SD	25.4 ± 3.7	26.1 ± 3.4	0.16
BMI <25	39 (50.0)	124 (41.3)	0.24
BMI 25–30	33 (42.3)	136 (45.3)	
BMI 30+	6 (7.7)	40 (13.3)	
Smoking status			
None	51 (64.6)	199 (64.8)	0.08
Current	1 (1.3)	24 (7.8)	
Former	27 (34.2)	84 (27.4)	
Previous hernia repair	15 (20.0)	79 (27.5)	0.19
Recurrent hernia repair	6 (7.6)	34 (11.0)	0.37
ASA class—median (Q1, Q3)	2 (2, 2)	2 (2, 2)	0.40

Table 2 Intraoperative variables

Variables	Asymptomatic <i>N</i> (%)	Symptomatic <i>N</i> (%)	<i>p</i> value
OR time (min)—median (Q1, Q3)	38 (30, 48)	38 (31, 47)	0.92
Bilateral hernia	21 (27.3)	106 (36.4)	0.10
Left hernia	28 (36.4)	72 (24.7)	
Right hernia	28 (36.4)	113 (38.8)	
Hernia type			
Femoral hernia	5 (5.8)	8 (2.9)	0.20
Direct	37 (43.5)	103 (34.5)	0.13
Indirect	49 (57.7)	224 (74.9)	<0.01
Pantaloon	9 (10.6)	51 (17.1)	0.14
Hernia size (cm)—median (Q1, Q3)	2.5 (2, 3)	2 (1.5, 2.5)	<0.01
Use of tacks	47 (61.0)	144 (49.5)	0.07

Table 3 Postoperative outcomes

Variables	Asymptomatic <i>N</i> (%)	Symptomatic <i>N</i> (%)	<i>p</i> value
LOS (hours)—median (Q1, Q3)	7 (6, 9)	7 (6, 9)	0.95
Pain at discharge—median (Q1, Q3)	2 (1, 3)	2 (0, 3)	0.29
30 day mortality	–	–	
Postop complications	15 (19.0)	58 (18.9)	0.98
Seroma	8 (11.4)	28 (10.5)	0.82
Hematoma	1 (1.4)	6 (2.2)	0.99
Urinary retention	6 (8.6)	16 (6.0)	0.42
Infection	0 (0.0)	4 (1.5)	0.58
ED visit	4 (6.9)	12 (6.2)	0.77
Readmission	0 (0.0)	7 (3.4)	0.35
Days until pain meds stopped—mean ± SD	1.2 ± 1.5	2.2 ± 3.0	0.02
Days return to ADL—median (Q1, Q3)	3 (2, 7)	5 (2, 7)	<0.01
Days return to work—median (Q1, Q3)	4 (3, 7)	4 (2, 7)	0.77
Hernia recurrence	1 (1.4)	2 (0.8)	0.52

The asymptomatic cohort discontinued pain medication and returned to normal activities of daily living more quickly than preoperatively symptomatic individuals. This is consistent with significant difference between the cohorts as measured by SOMS pain impact and quality. The longitudinal trends for these SOMS domains are shown in Figs. 3 and 4. It is interesting that pain impact for the asymptomatic group increases from preop to 3 weeks postop, before decreasing to well below preop levels by 6 months. As noted above, a mixed-effect model of pain by CSS over time demonstrated significantly lower pain in the asymptomatic group until adjustment for age. The CCS questionnaire data were rearranged to reflect proportion of response by time type for Figs. 5 and 6. There were significantly more “No symptoms” responses in the asymptomatic group at each time point postoperatively, remaining consistent with postoperative outcomes and SOMS trends.

Table 4 Average preoperative clinic and SOMS pain scores

	Asymptomatic	Symptomatic	<i>p</i> value
PreOp clinic	0.00	3.35	<0.01
SOMS			
PreOp	1.30	2.74	<0.01
3 weeks	1.28	1.96	0.02
6 months	1.00	1.17	0.77
1 year	0.55	1.44	0.08

Table 5 Longitudinal mixed-effects models for patient-reported outcomes (adjusted for age)

	Change over time		Asymptomatic (vs symptomatic)	
	Estimate (SE)	<i>p</i> value	Estimate (SE)	<i>p</i> value
SF-36 (higher scores better)				
Physical functioning	3.14 (0.57)	<0.01	−1.75 (2.79)	0.53
Role physical	4.42 (0.80)	<0.01	1.29 (3.57)	0.72
Role emotional	1.09 (0.56)	0.053	1.36 (2.86)	0.63
Energy/fatigue	1.00 (0.50)	0.05	−1.55 (2.72)	0.57
Well-being	0.54 (0.39)	0.17	0.19 (2.21)	0.93
Social functioning	1.73 (0.61)	<0.01	−1.43 (2.81)	0.61
Pain	4.27 (0.65)	<0.01	0.83 (3.20)	0.80
General health	0.75 (0.48)	0.12	−1.08 (2.52)	0.67
SOMS				
Physical functioning (higher scores better)	0.44 (0.10)	<0.01	1.07 (0.55)	0.051
Fatigue (lower scores better)	−0.21 (0.13)	0.11	−0.79 (0.63)	0.21
Pain impact (lower scores better)	−0.61 (0.11)	<0.01	−1.16 (0.49)	0.02
Pain quality (lower scores better)	−0.44 (0.09)	<0.01	−1.03 (0.40)	0.01
CCS total (lower scores better)				
Mesh	−1.02 (0.42)	0.02	−2.72 (2.27)	0.24
Pain	−0.07 (0.17)	0.68	−0.51 (0.76)	0.50
Pain	−0.51 (0.18)	<0.01	−1.37 (0.89)	0.13
Movement	−0.65 (0.18)	<0.01	−0.76 (0.78)	0.33

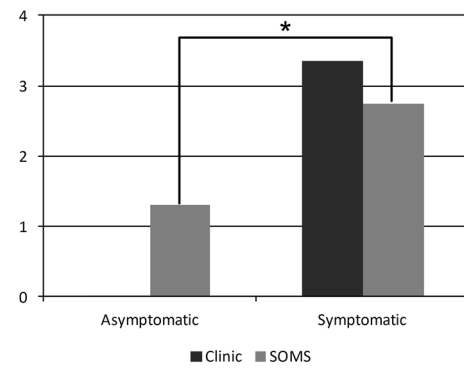


Fig. 1 Differences between preop clinic and preop SOMS pain scores between cohorts. Preop SOMS score between the cohorts was significantly different despite sorting cohorts by clinic scores

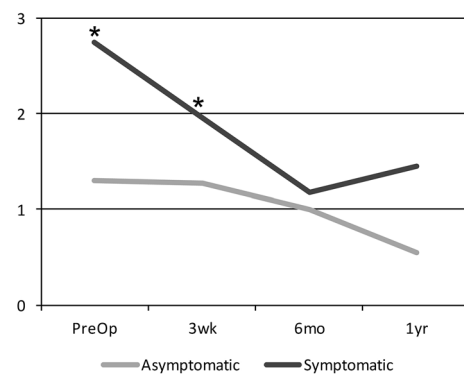


Fig. 2 SOMS VAS pain score over time. Difference between asymptomatic and symptomatic cohorts was significant at preop and 3 weeks postop

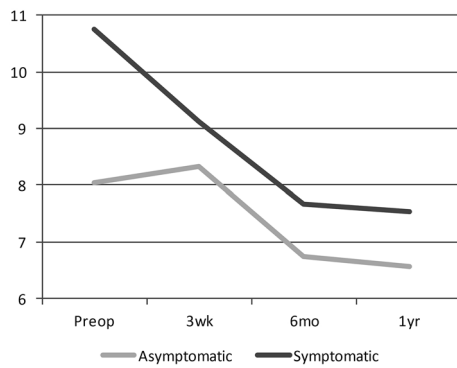


Fig. 3 Trend of SOMS pain impact domain over time between asymptomatic and symptomatic cohorts. Pain impact is scored from a minimum of 6 to a maximum of 30

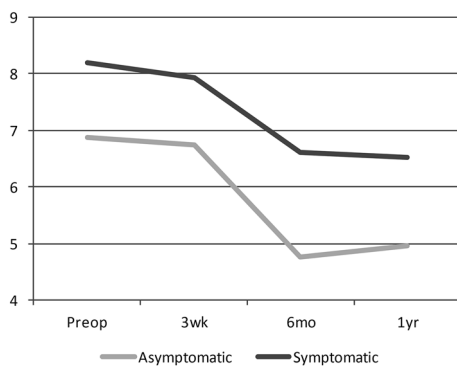


Fig. 4 Trend of SOMS pain quality domain over time between asymptomatic and symptomatic cohorts. Pain impact is scored from a minimum of 4 to a maximum of 21

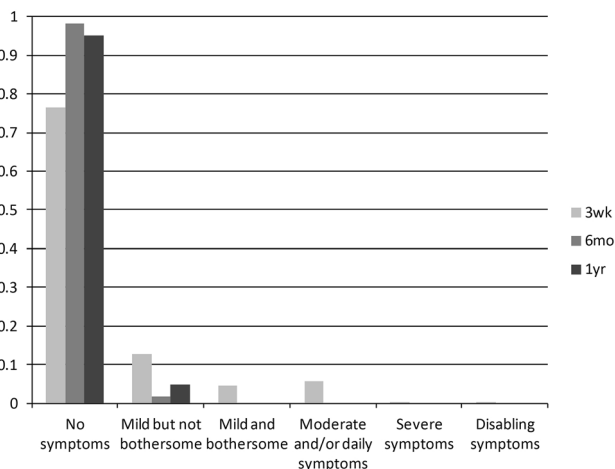


Fig. 5 Asymptomatic cohort: proportion of CCS score types by time

Discussion

In this review of 387 TEP LIHRs, patients in the asymptomatic cohort were older than their symptomatic counterparts. This may be explained by asymptomatic

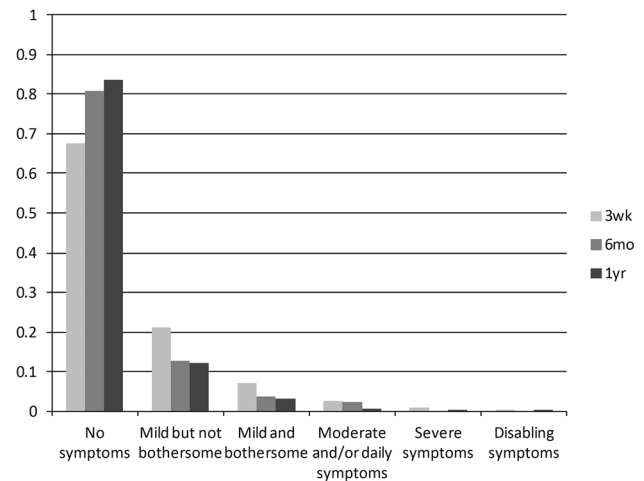


Fig. 6 Symptomatic cohort: proportion of CCS score types by time

individuals being diagnosed later or taking longer to seek surgical consultation. Intraoperatively, this cohort also was found to have larger hernia defects and a lower proportion of indirect hernias. This may indicate that smaller hernia defects and indirect hernias are more likely to cause symptoms. A review of literature was unable to identify other studies describing a difference in preop symptoms between direct and indirect or small and large hernia defects. Inguinal defects in the setting of athletic pubalgia or ‘sportsman’s hernia’ are documented, although findings vary, with some citing equal distribution of direct and indirect hernias, while others predominantly describing small, bilateral indirect defects [22, 23].

A clear limitation of this study resulting from its retrospective nature was that 251 of the 638 TEP LIHRs in the database were missing necessary parameters. The majority of these exclusions were due lack of orally reported pain score on a 0–10 scale from preoperative clinic. There was a consideration to run the analysis using the preoperative SOMS VAS score, which would have included more patients. However, since this measure is not routinely used clinically, an analysis from the perspective of oral patient reporting in clinic was felt to be more clinically relevant, especially given the significant difference found between oral pain score and the SOMS VAS preoperatively.

Groin pain in the presence of an inguinal hernia can be a challenging entity. Causality is often unclear and strong correlations between surgical findings and symptoms have not been established. However, it has been demonstrated that pain preoperatively is predictive of pain postoperatively [24, 25]. Likewise, this analysis suggests that lack of pain preoperatively predicts less pain postoperatively. Outcomes for asymptomatic patients were equivalent to symptomatic patients, and it may be concluded that elective, totally extraperitoneal laparoscopic repair is safe for asymptomatic, appropriate surgical candidates.

Compliance with ethical standards

Disclosures Dr. Ujiki reports personal fees from GORE and Medtronic, outside the submitted work. Dr. Linn reports personal fees and Industry sponsored research support from Medtronic, outside the submitted work. Dr. Haggerty reports personal fees from GORE and Medtronic, outside the submitted work. Drs. Hedberg, Butt, Linn, Denham, Lapin, Mr. Hall, Gitelis, and Ms. Carbray have no conflicts of interest or financial ties to disclose.

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