



To mesh or not to mesh for hiatal hernias: what does the evidence say

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Contributions: (I) Conception and design: Both authors; (II) Administrative support: None; (III) Provision of study materials or patients: None; (IV) Collection and assembly of data: None; (V) Data analysis and interpretation: None; (VI) Manuscript writing: Both authors; (VII) Final approval of manuscript: Both authors.

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Abstract: This review article discusses the history and evidence for outcomes from synthetic, biologic, and absorbable synthetic mesh reinforcement of the hiatus during paraesophageal hernia repair. Topics of discussion also include the use of mesh for the repair of type I hiatal hernias, as well as the use of relaxing incisions to close the difficult hiatus. The available literature suggests that use of synthetic mesh may reduce recurrence rates compared to primary closure alone. However, synthetic mesh placed at the hiatus has also been associated with complications that can be highly morbid, even resulting in a gastrectomy or esophagectomy. In contrast, the absorbability of biologic mesh is thought to minimize complications related to the presence of a permanent foreign body at the hiatus. There is evidence that hiatal reinforcement with biologic mesh reduces short-term recurrence rates after paraesophageal hernia repair compared to primary repair alone, but the rate reduction does not persist over long-term follow-up. While absorbable synthetic mesh remains another option, there is limited evidence on whether it effectively reduces recurrence rates after hiatal hernia repair. Due to the lack of high-quality comparative data between different types of mesh for paraesophageal hernia repairs, there is little evidence to guide whether to reinforce the hiatus with synthetic, biologic, or absorbable synthetic mesh.

Keywords: Recurrence; hiatal hernia; paraesophageal hernia; mesh; paraesophageal hernias (PEH)

Received: 03 December 2019; Accepted: 29 May 2020; Published: 20 October 2021.

doi: 10.21037/ales-19-249

View this article at: <http://dx.doi.org/10.21037/ales-19-249>

A brief history

The laparoscopic approach for paraesophageal hernia repair (LPEHR) began in the 1990s but was plagued by a recurrence rate as high as 33–42% (1,2). Prior to this, the laparoscopic approach was less frequently performed, and outcomes were more heavily focused on the technique and morbidity of repair rather than on anatomic recurrence (2-5). Once it was clear that laparoscopic repair was associated with lower morbidity, the focus shifted toward long-term anatomic success.

Large hiatal hernias have an intrinsically high recurrence rate attributed to tension at the hiatus, either axial along

the length of esophagus or laterally at the crural repair (6,7). Sources of tension at the hiatus include esophageal shortening, the natural pressure gradient between the abdomen and thorax, large hiatal defects, attenuated crural tissue, and the dynamic function of the diaphragm, which is in constant motion from respiration and also subject to sudden increased tension from coughing, laughing, sneezing, or straining (2,6-9). Factors that have been proposed to contribute to the higher rates or recurrence after laparoscopic repair include inaccurate assessment of intraabdominal esophageal length due to elevation of the diaphragm by pneumoperitoneum, decreased adhesions, less tactile feedback to determine tension of the crural closure,

and lack of deep bites at the crura when using laparoscopic suturing devices (7,10).

To address the high recurrence rates of LPEHR, surgeons began to examine the use of mesh to reinforce the hiatal repair given the success of using mesh for inguinal and ventral hernias (8). In the late 1990s to early 2000s, several randomized controlled trials (RCTs) compared recurrence rates after simple suture repair versus suture repair with mesh reinforcement of the hiatus during laparoscopic repair (9,11,12). All three RCTs found decreased short-term recurrence rates with the use of mesh reinforcement for hiatal hernia repair.

Still, there was concern with the use of mesh at the hiatus, including complications such as erosion, dense fibrosis, and esophageal stenosis (8,13-16). These complications have resulted in serious morbidity including esophageal perforation and need for reoperation, including major esophageal or gastric resection (16,17). Concern for complications from synthetic mesh has since led to the investigation of other types of intraperitoneal mesh for reinforcement of the hiatus, but to this day, the practice of whether to use mesh and what kind of mesh to use remains highly variable among surgeons (18).

In this article, we will review the evidence for outcomes from synthetic, biologic, and absorbable synthetic mesh reinforcement of the hiatus for paraesophageal hernias (PEH). We will briefly discuss mesh repair of type I hiatal hernias and the difficult hiatus, and then conclude with directions for future research.

Synthetic mesh

Early reports of using synthetic mesh for LPEHR involved the use of polypropylene mesh, which was associated with visceral adhesions to the mesh, prompting the search for another option for intraperitoneal mesh (8). One of the first reports of using polytetrafluoroethylene (PTFE) mesh to reinforce the crural closure in hiatal hernias was published by Frantzides and Carlson in 1997 (19). In their report, the authors described their experience using a PTFE mesh to circumferentially reinforce the posterior cruroplasty in a “keyhole” fashion in three patients with a large (≥ 8 cm) hiatal hernia. There were no complications or recurrences within the first 11 months of follow-up, prompting the authors to perform a preliminary comparison between 16 cases of primary cruroplasty to 15 cases of cruroplasty with PTFE mesh reinforcement (20). At follow up of 12–36 months, they found an 18.8% recurrence in the

primary repair group versus no recurrence in the mesh group ($P=0.08$), suggesting that there may be some benefit to PTFE mesh reinforcement of the crural repair in large hiatal hernias.

Therefore, Frantzides *et al.* published a subsequent RCT that enrolled over twice as many patients with a hiatal hernia defect ≥ 8 cm undergoing laparoscopic repair (9). In their analysis of 36 patients undergoing suture repair versus 36 patients undergoing suture repair with PTFE mesh reinforcement, patients were followed with an upper endoscopy and esophagram 3 months postoperatively and every 6 months thereafter to document any anatomic recurrence. The authors reported no recurrences in the mesh group compared to 22% recurrence in the non-mesh group at a median of 2.5 years (range, 6 months to 6 years) of follow-up ($P<0.006$). There were no differences in complications between the two groups and specifically no mesh-related complications.

The association between synthetic mesh reinforcement and decreased recurrence rates was affirmed by a subsequent RCT by Granderath *et al.* comparing 50 patients who had polypropylene mesh reinforcement of the hiatus versus 50 patients who underwent suture cruroplasty without mesh (11). In this study, patients with gastroesophageal reflux disease undergoing laparoscopic Nissen fundoplication were randomized to either hiatal suture repair versus suture repair with a 1×3 cm² polypropylene mesh sutured across the posterior crural repair. The authors found a statistically significant decrease in the rate of recurrence in the mesh group at one-year follow-up (26% versus 8%; $P<0.001$). Of note, patients with mesh reinforcement also experienced a statistically significant higher rate of postoperative dysphagia at 3-month follow-up (12% versus 4%; $P<0.05$), but this difference disappeared over time with 4% residual dysphagia in both groups by 1-year follow-up. Similar to the results published by Frantzides *et al.* above, there were no reported mesh-related complications in this study, which the authors attributed to keeping the mesh away from the esophagus, buffered by the fundoplication.

The most recent RCT comparing synthetic mesh use in LPEHR was published by Oor *et al.*, who examined reinforcement for hiatal defects >5 cm. The authors compared 36 patients undergoing laparoscopic hiatal repair with mesh versus 36 patients undergoing repair with suture alone (21). In contrast to the previously mentioned RCTs, the authors of this study found no statistically significant difference in rates of recurrence between the two groups at 6-month follow-up with endoscopy and/or esophagram

(25% versus 19.5%, respectively; $P=0.581$). A possible explanation for the different outcomes may be the type of fundoplication used in Oor *et al.*'s study compared to the other two RCTs. In Oor *et al.*'s study, although patients were randomized, statistically significantly more patients underwent a posterior partial fundoplication (Toupet) in the mesh versus non-mesh group (83.3% versus 16.7%, respectively; $P<0.001$), and the rest of the patients underwent an anterior 180-degree partial (Dor) fundoplication. Choice of fundoplication was left to the surgeon's discretion and was thus unblinded to the use of mesh. No patients underwent a full, 360-degree (Nissen) fundoplication, whereas in both Frantzides *et al.*'s (9) and Granderath *et al.*'s (11) studies, all patients underwent a complete Nissen fundoplication. It is unclear how the type of fundoplication affects recurrence rates for PEHs in combination with or without mesh reinforcement, as such a study has yet to be published. However, the nonrandomized use of Toupet fundoplication in 83.3% of mesh cases versus only 16.7% of non-mesh cases should certainly be regarded as a significant confounder in Oor *et al.*'s study, and results of the study should be interpreted with caution.

Despite level 1 evidence from two RCTs demonstrating that hiatal reinforcement with synthetic mesh reduced recurrence rates after LPEHR with no difference in rates of complications or postoperative symptoms, routine adoption of the technique was limited by reports of significant complications resulting from placement of synthetic mesh at the hiatus such as erosion, stricture, and significant dysphagia (8,13-15). More recent evidence has reinforced these concerns. In Müller-Stich *et al.*'s systematic review of 124 studies on mesh use in LPEHR, the authors found an overall 0.9–1.9% rate of mesh-associated complications, the majority (71.5%) of which involved synthetic mesh, either polypropylene or PTFE, with a synthetic mesh-related complication rate of 0.8–2.5% (22).

Although mesh-related complications are rare, they can have disastrous consequences. In their report of 28 hiatal hernia mesh-related complications, Stadlhuber *et al.* described 21 complications specifically related to use of synthetic mesh, including 16 cases of mesh erosion, three cases of dense fibrosis, and one case of esophageal stenosis (16). Major resection (partial or total esophagectomy or gastrectomy) was required in 45% of the reoperations for synthetic mesh complications and in 32% of the mesh complications overall. This is consistent with the finding that 30% of patients undergoing revisional foregut surgery end up requiring major resection if they have

a history of mesh placement at the hiatus, compared to only 4% incidence of major resection without mesh (23). As a result, most foregut surgeons avoid using synthetic mesh during hiatal hernia repair with rare exception, as demonstrated in a Society of American Gastrointestinal and Endoscopic Surgeons survey reporting that 65% of surgeons use biologic material when mesh is used for crural reinforcement during hiatal hernia repair (18).

Biologic mesh

In an effort to avoid the complications related to synthetic mesh at the hiatus, when biologic mesh materials hit the market, surgeons began using them as an alternative to synthetic mesh. The theoretic advantage of a biologic prosthesis was that provides a temporary collagen matrix to allow native tissue ingrowth at the hiatal repair with a resulting repair that is stronger than native tissue (24). Since the biologic scaffold dissolves over time as it is incorporated by the body, the thought was that the mesh could be used to support the hiatal repair during healing while also avoiding complications related to having a permanent foreign body at the hiatus.

In the only published RCT focused exclusively on biologic mesh use in LPEHR, we compared outcomes of patients with PEH who had their hiatal repair reinforced with mesh made from porcine small intestinal submucosa ($n=51$) versus primary repair without mesh ($n=57$) (12). We found a statistically significant decrease in anatomic recurrence with the use of mesh reinforcement at six months postoperatively (9% versus 24%; $P=0.04$), which encouraged the use of biologic mesh reinforcement among the surgical community. However, long-term follow-up at five years demonstrated no statistically significant difference in recurrence rates between the two groups (54% versus 59%, respectively; $P=0.7$) (25). Nevertheless, there were no mesh-related complications and no reoperations in the mesh group throughout the long-term follow-up of median 58 months (range, 42–78 months). There were also no statistically significant differences in postoperative symptom or quality-of-life scores between the two groups either at short-term or long-term follow-up. In other words, although there appeared to be no downside to biologic mesh, the short-term reduction in recurrence rate did not appear durable.

A more recent RCT by Watson *et al.* compared outcomes between patients undergoing LPEHR with mesh reinforced by either synthetic ($n=42$) or biologic ($n=41$) mesh versus patients with no mesh reinforcement ($n=43$), and found no

difference in the 6-month recurrence rates between mesh reinforcement (either synthetic or biologic) versus no mesh (21.8% versus 23.1%, respectively; $P=1$) (26). The authors' definition of recurrence was any intrathoracic stomach as determined by esophagram or endoscopy, whereas our study limited recurrence to a hiatal hernia >2 cm on upper gastrointestinal series. However, even when Watson *et al.* limited their analysis of recurrence to cases in which the hiatal hernia was ≥ 2 cm, there was still no statistically significant difference between the mesh versus no-mesh groups (2.6% versus 7.7%, respectively; $P=0.329$). There were no reported cases of mesh-related erosion or stricture. At one-year follow-up, there were no differences in dysphagia rates or satisfaction scores among the three groups, but biologic mesh was associated with increased nausea (27.5% versus 15% no mesh and 4.9% synthetic mesh; $P=0.0197$) and decreased ability to relieve bloating (72.5% versus 92.5% no mesh and 97.5% synthetic mesh; $P=0.0017$). The authors point out that the magnitude of these differences was small and unlikely to be clinically significant. They further reiterated that the overall outcomes and adverse effects in all groups were similar, and that long-term analysis with more extended follow-up is needed.

The natural next step was to compare recurrence rates between synthetic versus biologic mesh for LPEHR, however the lack of comparative studies renders making any comparison very difficult. The only randomized trial published to date that compares biologic to synthetic mesh for hiatal hernia repair is Watson *et al.*'s study discussed above (26). Recurrence of any degree of intrathoracic stomach was 30.8% in the absorbable mesh group versus 12.8% in the synthetic mesh group, and recurrence limited to a hiatal hernia ≥ 2 cm was 5.9% versus 0%, respectively. While these differences were not statistically significant, the results suggest that synthetic mesh may decrease recurrence rates more than biologic mesh, but perhaps the study groups were too small to reach statistical significance.

However, a recently published systematic review of nonrandomized studies by Castelijns *et al.* did find a statistically significant decrease in the rate of recurrence with the use of synthetic mesh reinforcement compared to biologic mesh (27). The authors pooled 16 studies on the use of mesh in laparoscopic hiatal hernia repair to compare 704 patients who received synthetic mesh with 385 patients who received biologic mesh and found a recurrence rate of 6.8% in the synthetic mesh group versus 16.1% in the biologic mesh group ($P<0.05$). However, given the heterogeneity of the included studies, the authors

highlighted the need for more RCTs to directly compare outcomes from biologic and synthetic meshes. Fortunately, there is an ongoing RCT that will compare recurrence rates and symptomatic/quality-of-life assessments between synthetic and biologic mesh reinforcement of LPEHR, but the study completion date is scheduled for August 2020 (ClinicalTrials.gov Identifier NCT02242526) and thus results are still unavailable.

While there are a number of meta-analyses of case series and clinical trials on the use of mesh for LPEHR and the comparison of synthetic versus biologic mesh, interpretation of the results is limited by the significant diversity of the included studies (22,28-33). These studies compared cases in which there were variable indications for hernia repair, different types/sizes of hiatal hernias, various types/shapes/positions of mesh, diverse methods of securing mesh, and different types of fundoplication. In addition, some studies included cases that involved a lengthening procedure (e.g., Collis gastroplasty) and/or relaxing incisions, whereas these cases were excluded in other studies. Furthermore, the definition of recurrence (e.g., any intrathoracic stomach versus only >2 cm intrathoracic stomach) and duration of follow-up were highly variable among studies. The extreme heterogeneity among the studies used in these meta-analyses limits the reliability of any of the conclusions drawn from the analyses.

However, even if synthetic mesh is ultimately found to have lower recurrence rates compared to biologic mesh, observational studies seem to suggest that there are fewer mesh-related complications from biologic mesh compared to synthetic mesh. In a paper summarizing mesh-related complications after hiatal hernia repair, synthetic mesh was associated with 75% of the reported complications and all 16 cases of erosion related to a specific type of mesh (16). There was one additional case of mesh erosion associated with biologic mesh, but it was unclear whether the biologic mesh or the coexisting synthetic mesh was the underlying cause of the complication. Li and Cheng likewise reported that the vast majority of mesh-related erosion at the hiatus was related to synthetic mesh, with only 2% of the 50 reported cases occurring after a biologic mesh (34). The only case of erosion related to biologic mesh in their study was the same one that was combined with a synthetic mesh as reported by Stadlhuber *et al.* Of particular concern is that 26% of the patients with mesh-related erosion at the hiatus ultimately require major resection including distal esophagectomy or partial/total gastrectomy (34). In contrast, long-term follow-up out to a median of

4–5 years after biologic mesh placement has revealed no mesh-related complications (25,35). Thus, it is important to consider whether the potential decreased recurrence rate offered by synthetic mesh reinforcement is worth the added risk of significant morbidity. While the short-term benefit of biologic mesh reinforcement does not appear to be durable at 5 years, there are also limited if any long-term complications from the use of biologic mesh at the hiatus.

Absorbable synthetic mesh

Given the outcomes reported from biologic mesh use in LPEHR, the absorbable properties that make biologic mesh appealing for reducing mesh-related complications may also impair its ability to provide a durable hiatal hernia repair. In addition, the relatively high cost of biologic mesh material has limited its routine use for even abdominal wall hernias (36). Thus, absorbable synthetic mesh material was developed as a potentially more cost-effective option for hernia repair (37). Although data are still lacking, the use of absorbable synthetic mesh in LPEHR may also be a promising option that combines the more resilient crural reinforcement provided by synthetic mesh with the absorptive properties of biologic mesh. Indeed, comparison of an absorbable synthetic versus biologic mesh for ventral hernia repair in a rat model demonstrated that the absorbable synthetic mesh is rapidly incorporated with less mesh shrinkage compared to biologic mesh, and the repair was stronger than native tissue at 60 days (38). These biophysical properties of absorbable synthetic mesh are encouraging, but whether the results translate to clinical outcomes remains to be seen. However, there are a few published case series that suggest positive results when absorbable synthetic mesh is used for reinforcement during hiatal hernia repair (39–42).

In a retrospective cohort study, Asti *et al.* compared 43 patients who had LPEHR with no mesh reinforcement to 41 patients who had hiatal reinforcement with Bio-A mesh (polyglycolic acid/trimethylene carbonate) (40). At a median 24-month (minimum 1-year) follow-up, there was a 9.7% recurrence rate in the mesh group versus an 18.6% recurrence rate in the non-mesh group, although this result was not statistically significant. There were no differences in complications and no reoperations in either group. Asti *et al.* later reported outcomes of their first 100 cases of any hiatal hernia repair (90% PEHs) with absorbable synthetic mesh and reported a recurrence rate of only 9% at a median 30 months of follow-up (41). This recurrence rate was

similar to the 9.5% recurrence at median follow-up of 14 months (range, 11–34 months) reported by Zehetner *et al.* in their review of 35 patients undergoing LPEHR with Vicryl (polyglactin) absorbable synthetic mesh reinforcement (39).

In a recent report of 50 consecutive cases of LPEHR with Phasix-ST absorbable synthetic mesh, Abdelmoaty *et al.* reported an 8% recurrence rate at 1-year follow-up with no reported mesh complications (42). Phasix-ST mesh is composed of poly-4-hydroxybutyrate with a Septra-Technology coating that allows for placement next to viscera, with slow resorption over 12–18 months (43,44). While the results appear promising, the authors noted that they were unable to determine whether the relatively low recurrence rate was due to the mesh itself, due to the use of Collis gastroplasty (used in 42% of cases) or relaxing incisions (4% of cases), or due to a combination of these techniques.

Currently there have been no reported cases of mesh-related complications from the use of absorbable mesh during LPEHR, but the meshes are relatively new and long-term outcomes have yet to be published.

Type I hiatal hernias

The discussion thus far has focused on repair of PEHs, with typically an irreducible portion of intrathoracic stomach and/or with a hiatal defect of >5 cm. However, given a recurrent hiatal hernia is one of the most common reasons for symptomatic failure after a fundoplication for antireflux surgery (45–47), some surgeons have extrapolated the potential benefit of mesh reinforcement in PEH to type I (sliding) hiatal hernias (45,46,48–50).

There are no RCTs evaluating the effect of mesh reinforcement of the hiatus during repair for exclusively type I hiatal hernias. Although Granderath *et al.*'s (11) study likely included a number of patients with type I hiatal hernias, each group had about 60% of patients with a defect >5 cm and thus likely included a majority of PEHs.

There are limited published case series comparing the effect of mesh reinforcement of the hiatus for type I hiatal hernias. Kamolz *et al.* examined the outcomes of 100 patients with synthetic mesh reinforcement of the crural repair during laparoscopic antireflux surgery and compared them to outcomes of 100 patients without mesh reinforcement (45). At 1-year follow-up, there were no recurrences in the mesh group compared to 9% in the non-mesh group based on endoscopy. The mesh group was associated with increased rates of dysphagia at 3 months

postoperatively (35.4% versus 20.8%), but there were no differences by 1 year after surgery (4.8% versus 5.3%). Similarly, Jacobs *et al.* found decreased recurrence rates in patients who underwent laparoscopic antireflux surgery with biologic mesh reinforcement of the hiatus compared to repair without mesh (3.3% versus 20%, respectively) (48). Note that neither of these studies specified the size or type of hiatal hernias repaired, and it is possible that some PEHs were included with variable frequency in either group.

Turkcapar *et al.* specifically excluded all PEHs in their case series examining 511 patients who underwent laparoscopic fundoplication with synthetic mesh (n=176) versus without mesh (n=335) reinforcement of the crural repair (46). The authors found a recurrence rate of 1.8% in the mesh group versus 6% in the group without mesh based on endoscopy and/or esophagram at a follow-up of a least 2 years and a mean follow-up of 75 months (P=0.036). There were no differences in dysphagia between the two groups and no reported mesh-related complications.

Schmidt *et al.* examined the effect of biologic mesh reinforcement on recurrence rates for small hiatal defects, defined as measuring 1–5 cm on either preoperative esophagram or endoscopy (50). The authors compared laparoscopic fundoplication with mesh (n=38) versus without mesh reinforcement and found that at 1-year follow-up with esophagram and/or endoscopy, the rate of recurrence in the mesh group was 0% compared to 16% in the group without mesh (P=0.017).

Of note, none of these studies were randomized and all represented case series in which mesh reinforcement was introduced and implemented as standard practice after a specific time point. Thus, the non-mesh group comprises patients operated on earlier in the series, whereas patients in the mesh group were operated on later in the surgeons' experience. The sequential collection of data for one group followed by the comparison group can confound the results given that recurrence rates have been observed to decrease over time as surgeons overcome their learning curve (51).

There is currently no high-quality evidence that mesh reinforcement of the hiatus reduces recurrence rates in type I hiatal hernias, although results from a few case series suggest a possible benefit. Nonetheless, few surgeons believe mesh should be used for type I hiatal hernias until a benefit is clearly shown for larger PEHs.

The difficult hiatus

When the crura cannot be approximated at all or will only

do so under excessive tension, a diaphragmatic relaxing incision with mesh reinforcement is recommended to decrease tension at the hiatus (6,52,53) (Figure 1A,B). Diaphragmatic relaxing incisions can be made either to the left or the right of the hiatus, or even bilaterally when needed to allow approximation of the crural pillars and primary closure of the hiatal defect.

Typically, a right-sided diaphragmatic relaxing incision is preferred as the first maneuver to close a difficult hiatus (Figure 1C,D,E), and can reduce average tension at the hiatus by 46% (52). The right-sided relaxing incision is preferred as it typically involves just a small incision parallel to the right crus. In addition, the caudate and left lateral segment of the liver may help cover the defect to minimize the occurrence of a diaphragmatic hernia. In contrast, a left diaphragmatic relaxing incision is not only more exposed, but it requires a large lateral incision to avoid injury to the pericardium and phrenic nerve (6). However, if the right crus is excessively attenuated, fibrotic, or too close to the inferior vena cava, a left-sided diaphragmatic relaxing incision can certainly be used as the initial maneuver to reduce radial tension at the hiatus (6,53). On rare occasions, bilateral relaxing incisions may be needed to decrease the radial tension enough to approximate the crus. Some authors also advocate intentional entry into the pleural space to improve pliability of the diaphragm and reduce any tension from an adhered lung (6,7,52).

Any diaphragmatic defect from a relaxing incision should be covered with mesh to prevent herniation of intraabdominal contents (Figure 1E). On the right side, it is debatable whether to use synthetic or biologic mesh. There are no studies specifically comparing synthetic versus biologic mesh for a right-sided diaphragmatic relaxing incision. Those who advocate for synthetic mesh repair believe biologic materials should never be used to bridge a gap, while biologic proponents point to liver protection and lack of published evidence on right crural hernias. Regardless, the type of mesh at this location is unlikely to have great significance since the mesh is placed away from the esophagus and is protected from the viscera by the caudate and left lateral lobe of the liver. For a left-sided relaxing incision, we recommend the use of synthetic mesh only, since absorbable mesh can result in a diaphragmatic hernia once the mesh absorbs in this exposed position (6,53).

Conclusions

The literature to date supports the potential of mesh to

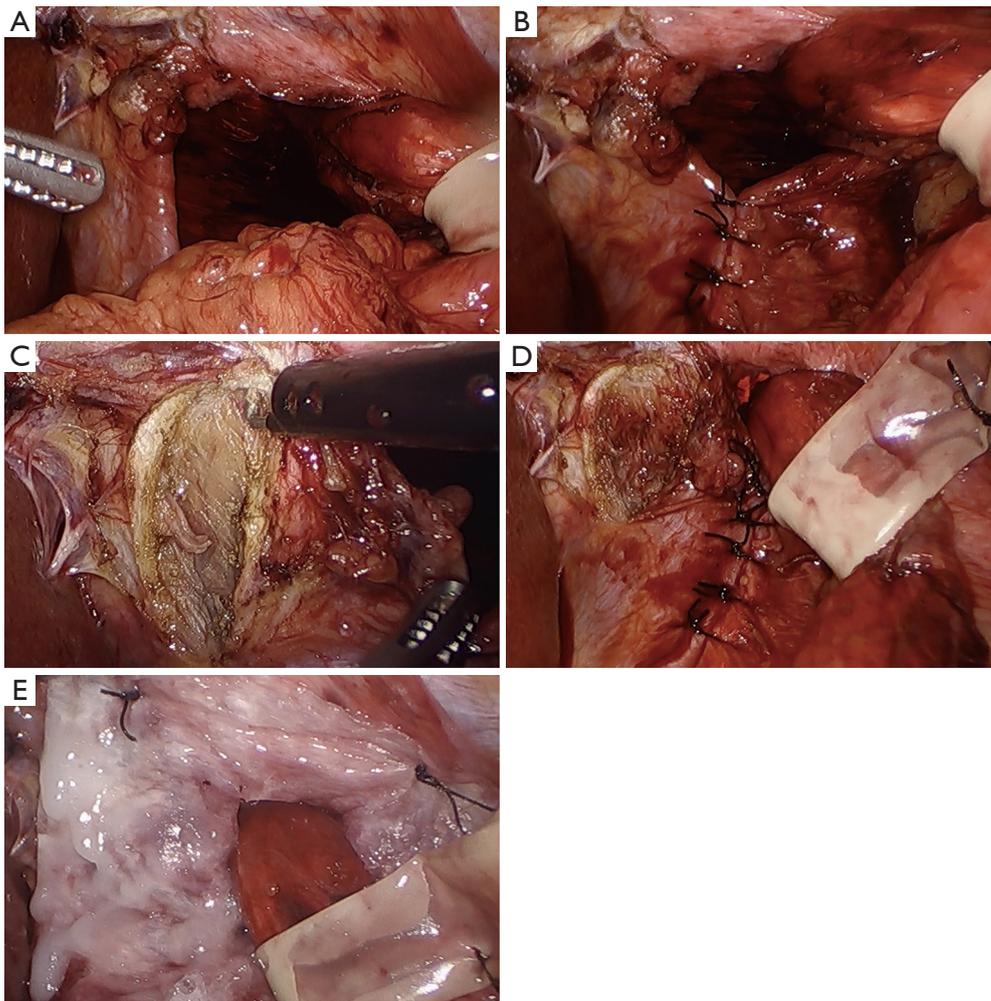


Figure 1 The difficult hiatus. There is a wide hiatal defect (A) and excessive tension is noted as the defect is repaired (B). A right diaphragmatic relaxing incision is performed (C), which allows for complete primary closure of the defect (D). The repair is reinforced with a biologic mesh that has been secured with both fibrin glue and sutures (E).

decrease recurrence after LPEHR compared to primary suture cruroplasty. The evidence is great for synthetic materials, but these are also associated with devastating, albeit rare, complications when placed at the hiatus. On the other hand, there are fewer reports of complications related to biologic mesh, but little evidence for preventing long-term recurrence 5 years and beyond. Newer absorbable synthetic mesh may reduce the cost associated with biologic materials but data on efficacy compared to synthetic or biologic mesh are lacking.

There is no strong evidence to either support or oppose the use of mesh reinforcement at the hiatus, although mesh should certainly be used to cover a diaphragmatic defect resulting from a relaxing incision for a difficult hiatus.

Unfortunately, there is a lack of high-quality comparative data between different types of mesh for LPEHR, so when mesh is needed, there is little evidence to guide which type to use within the main three categories of mesh.

Future research should focus on RCTs to determine long-term outcomes for different types of mesh compared to each other, as well as compared to non-mesh LPEHR. This topic has many unanswered questions deserving of further investigation.

Acknowledgements

Funding: None.

Footnote

Provenance and Peer Review: This article was commissioned by the Guest Editors (Lee L. Swanstrom and Steven G. Leeds) for the series “Hiatal Hernia” published in *Annals of Laparoscopic and Endoscopic Surgery*. The article has undergone external peer review.

Conflicts of Interest: Both authors have completed the ICMJE uniform disclosure form (available at <http://dx.doi.org/10.21037/ales-19-249>). The series “Hiatal Hernia” was commissioned by the editorial office without any funding or sponsorship. The authors have no other conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. This study did not require approval by an ethics board as there was no involvement of patients or patient data.

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doi: 10.21037/ales-19-249

Cite this article as: Inaba CS, Oelschlager BK. To mesh or not to mesh for hiatal hernias: what does the evidence say. *Ann Laparosc Endosc Surg* 2021;6:40.