

Review Article

A review of experimental studies for available experimental evidence on the use of prosthetic material in diaphragmatic hiatal hernia repair

ABSTRACT

Aim: The benefits of prosthetic material in hiatal hernia repair have been well documented. However, the associated risks are substantial and they are related to the technique, ~~but and~~ also the choice of material. Experimental data are invaluable to understand and evaluate the interaction of different meshes with the host tissue. The purpose of this article is to summarize the available experimental evidence in the repair of hiatal hernias with the use of prosthetic materials in animal models.

Methods: A review of the literature from January 1990 to December 2014 was carried out for articles presenting experimental data on hiatal hernia repair.

Results: After discarding non relevant articles, 28 ~~35~~ articles were identified. A variety of synthetic and absorbable materials were studied. Review of the available studies showed that there is great variability between synthetic materials regarding tissue integration, shrinkage and adhesion formation, however they have greater mechanical strength when compared to ~~however~~ biological/absorbable materials, which have a tendency ~~for to~~ better integration ~~in host tissue~~. Biological adhesives seem to be an effective alternative method of mesh fixation.

Conclusions: Experimental data are essential in order to fully appreciate the process of repair of a hiatal hernia with a prosthetic material. The articles reviewed provide insight into the properties of different prosthetic materials. However, there were large variations in their quality and the methods used. Data from animal studies are an excellent way of evaluating the multitude of materials that have recently become available. Good quality, comparative animal studies are essential in an effort to further improve outcomes for patients who undergo hiatal hernia repair.

Keywords: hiatal, ~~hernia, diaphragmatic~~, mesh, animal, experimental, review

1. INTRODUCTION

The introduction of laparoscopic techniques in hiatal hernia repair resulted in a significant increase in the number of annually performed anti-reflux procedures in ~~the lastest than a~~ decade [1]. There are now randomized trials supporting the use of surgical management as a first-line treatment in selected patients [2]. In some patient subgroups, however, such as ~~the~~ patients with a large paraesophageal hernia, recurrence rates can reach 42% [3]. ~~Recurrence-It~~ usually occurs after disruption of the crural closure ~~and~~ as the tissues ~~being~~ approximated are frequently attenuated and sutured under tension [4].

In an effort to overcome these limitations, selective mesh use has been reported since the 1970s. In the first large series of patients published, Carlson *et al.* were able to achieve excellent results with polypropylene repair, without any clinical recurrences in long term follow up [5]. A number of clinical trials have ~~since~~ established the efficacy of prosthetic mesh in preventing recurrence in the hiatus [6], however, the emergence of relatively few, but in some cases devastating, complications such as mesh erosion, highlight the need for further research [7].

As new materials are continuously being developed it is important for surgeons to make an informed decision on which material to use. Animal studies are essential in evaluating the interaction between the different prosthetic materials and the host tissue and their relative safety and efficacy in hiatal hernia repair. We have performed a literature review in order to examine the contribution of the available experimental evidence towards selecting the optimal prosthetic material and surgical technique in mesh repair of ~~hiatal-diaphragmatic~~ hernia.

2. MATERIAL AND METHODS

We searched for articles on ~~hiatal-diaphragmatic~~ hernia repair meeting the criteria outlined below and analyzed them for specific outcomes using the PRISMA guidelines.

2.1 ELIGIBILITY CRITERIA

40 1) Type of study: Experimental animal (in vivo) study of repair of hiatal/paraesophageal or congenital
41 diaphragmatic hernia using prosthetic material (mesh). Models of congenital diaphragmatic hernia
42 were included in this review, because, although the mesh was not placed in the hiatus in these
43 models, they can be considered orthotopic models, usually involving creation of a hernia by excision
44 of part of the left hemidiaphragm, mimicking conditions like those found in a giant paraesophageal
45 hernia (large defect, attenuation of muscular tissue).

46 2) Language: English

47 3) Publication year: 1990-2014

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49 **2.2 LITERATURE SEARCH STRATEGY**

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51 Studies were identified by searching the PubMed/Medline and Scopus databases. The following
52 key words were used as search strings: hiatal, diaphragmatic, mesh, animal, experimental.

53 Potentially relevant articles were identified by the title and abstract and full papers were obtained
54 and assessed in detail by two of the authors (M.S. and P.T., both senior surgeons) prior to their inclusion
55 in the review. The reference list for each article was also screened to identify further relevant publications.

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57 **2.3 Study selection**

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59 Eligibility assessment was performed independently by 2 reviewers. Disagreements between
60 reviewers were resolved by consensus.

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62 **2.4 Data extraction**

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64 Data collection and analysis were carried out independently by 2 researchers. Studies were
65 classified into two experimental model groups which investigated mesh repair of either hiatal or congenital
66 diaphragmatic hernia. Articles were reviewed for a number of variables examining their design (number
67 and type of animals, mesh implantation time, use of comparative/control group,

69 biomechanical/histopathological analysis) and the technique used (Mesh type and shape, fixation type,
70 surgical technique).

71 Study results were specifically assessed for findings relevant to controversial topics in hiatal
72 diaphragmatic hernia repair with prosthetic mesh (Table 1).

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Table 1: Controversial topics in hiatal hernia repair with prosthetic mesh

1. Mesh shape	
2. Mesh type	a. Infection potential
	b. Handling characteristics
	c. Durability of repair
	d. Adhesion potential, tissue incorporation, fibrosis/stenosis/shrinkage potential
	f. Migration/erosion potential
3. Fixation method	
4. Sutured vs tension-free hiato-plasty	

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75 **3. RESULTS**

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77 **3.1 Literature search**

78 Our search strategy initially returned 924 studies which we evaluated based on title and
79 abstract and we selected 21 articles based on our inclusion criteria. The full text of these articles was
80 downloaded and another 9 studies were obtained from their reference lists. After excluding 2 articles
81 studying hiatal hernia repair in the context of fetal tissue engineering, 28 articles were assessed in detail
82 (Figure 1).

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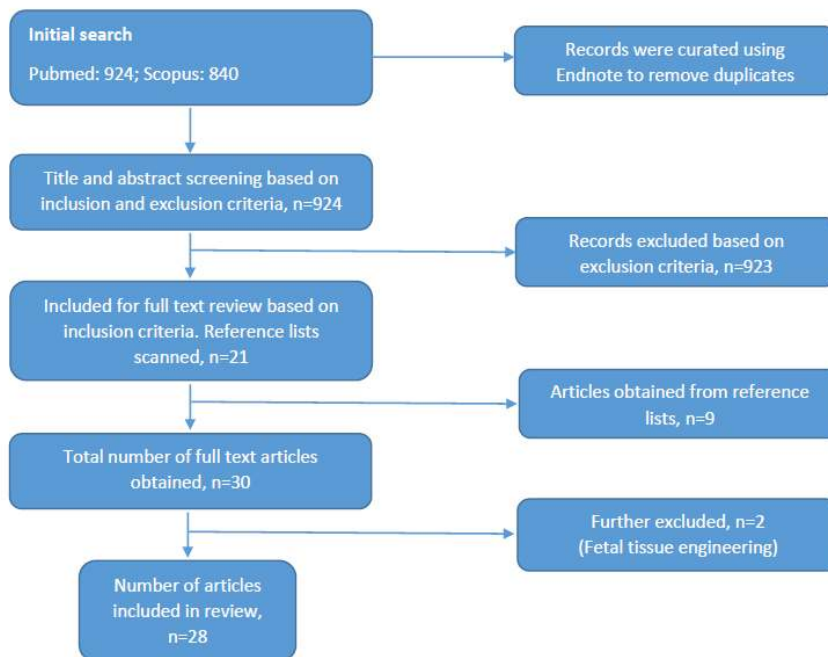
84 **3.2 Study design**

85 Large animals (swine, dogs) were used in most studies. The number of animals in each study was
86 small (6-36 animals). Implantation time ranged from 2 weeks to 12 months (Table 2). The majority of the

87 studies included a comparison or control group and histopathological analysis, however only a few
88 studies used endoscopic or radiological assessment or biomechanical analysis.

89 **3.3 Mesh characteristics and surgical technique**

90 A variety of meshes were evaluated, including conventional (polypropylene,
91 polytetrafluoroethylene - PTFE) and newer (polypropylene/ polyglactin 910 - PP-PG, poly(lactic-co-
92 glycolic acid) - PLGA) synthetic materials, biologically derived materials such as bovine pericardium and
93 newer biologic meshes (Small intestinal submucosa - SIS, acellular dermal matrix - Alloderm). Most
94 authors used a rectangular piece of mesh, but circular and U-shaped meshes were also used. The
95 surgical technique used in most studies was mesh fixation in the hiatus using an open technique, with or
96 without excision of part of the left hemidiaphragm, while in two studies an endoscopic approach was
97 utilized: laparoscopic creation of a defect in the left hemidiaphragm and repair in one study and
98 thoracoscopic creation of a paraesophageal hernia and subsequent laparoscopic repair in another.
99 Finally, mesh fixation was achieved with sutures in most cases, while a few of the authors used biological
100 adhesives, such as fibrin glue and polyethylene glycol (Table 3).



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102 **Figure 1: Flow diagram of literature search**

103 **3.4 Results of individual studies**

104 **3.4.1 Mesh shape**

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106 Although circular, rectangular and U-shaped meshes were used, no study directly compared meshes of
107 different shapes.

108 ~~In a study of polypropylene meshes of a circular shape fixed by sutures in a rabbit model, the meshes~~
109 ~~had usually moved from their implantation bed and had eroded into the esophagus [16]. However, in~~
110 ~~another study a circular polypropylene mesh was fixed in place using fibrin glue in a swine experimental~~
111 ~~model and in this case the authors reached conflicting results as they found the meshes stayed in~~
112 ~~position and their inner edge had retracted evenly from the esophagus [14].~~

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114 **3.4.2 Mesh type**

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117 *3.4.2.1 Infection potential*

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119 No study on mesh use in contaminated fields has been carried out.

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121 *3.4.2.2 Handling characteristics*

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123 The handling characteristics of each mesh i.e. the ease of its use in laparoscopic surgery was not
124 addressed in any study.

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126 *3.4.2.3 Durability of repair*

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128 Most of the studies ~~discussed~~ showed that the mesh repair remained successful during the observation
129 period of up to 12 months. SIS was shown to have equivalent strength to PTFE when applied on the
130 diaphragm [27, 32], although it was not as strong as polypropylene meshes [25, 26]. In another study
131 comparing two forms of SIS mesh in a dog model, the first comprised of 4-ply and the other from 8-ply,

132 the thicker version was shown to be stronger, while both showed more strength than native diaphragmatic
133 tissue [28]. Fascia lata was also shown to be equivalent to PTFE in mechanical strength [29].

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135 3.4.2.4 Adhesion potential, tissue incorporation, fibrosis/stenosis/shrinkage potential

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137 Polypropylene mesh consistently caused formation of strong adhesions [25, 26], which were less
138 pronounced with low-weight polypropylene [9]. Dualmesh showed less extensive adhesions than
139 polypropylene [17], while Surgisis showed less adhesions than PTFE in two studies [11, 32], but dense
140 adhesions were comparable to polypropylene in another [25].

141 Mesh shrinkage was shown to be around 50-70% of original size for polypropylene [9, 14, 16], while the
142 percentage of shrinkage was more higher for the low-weight mesh [9]. When PTFE, polyester and
143 polypropylene were compared, PTFE showed considerably more shrinkage that reached 34.9% of its
144 original size [AZF2][8].

145 Bohm et al compared two composite ~~synthetic~~ polypropylene meshes (Ultrapro, Proceed) to
146 Surgisis in a rabbit model [25, 26]. Inflammatory reaction at the border of the mesh was more pronounced
147 with Proceed, followed by Ultrapro and Surgisis. On the other hand Surgisis and Ultrapro showed better
148 tissue regeneration compared to Proceed. Collagen maturation was slower for Surgisis compared to the
149 synthetic meshes. [AZF3]A composite polypropylene mesh was compared to a conventional polypropylene
150 mesh and the composite mesh showed better integration and reduced inflammatory response, which
151 could be associated with a lower risk of erosion and postsurgical dysphagia [16]. Histological examination
152 and cross-polarization microscopy showed differences in cell proliferation rate, apoptosis and collagen
153 I/III ratio, which were statistically significant and show better tissue integration for the composite mesh
154 [15, 16] Another study showed excellent integration, for a titanium-polypropylene mesh [13].

155 Polytetrafluoroethylene (ePTFE/ Dualmesh) was evaluated and caused the formation of minimal
156 adhesions except in segments of the mesh where folding exposed its superior surface. There were no
157 erosions or migration noted. Microscopic evaluation showed only an unstable capsule encompassing the
158 mesh underlining the importance of a stable fixation [17, 24, 33].

159 Biologically-derived materials were evaluated in several studies. The authors reported complete
 160 mesh replacement by fibrovascular scar tissue with SIS mesh, with significant muscular regeneration,
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Table 2: Design of experimental studies in hiatal diaphragmatic hernia repair

Author	Pub year	Animal type	N	Implantation time	Comparative/control group	Biomechanical analysis	Histopathological analysis	Endoscopic/radiological assessment
<i>Hiatal hernia repair models</i>								
1	Muller-Stich [8]	2014	Swine	24	8 weeks	✓	✓	
2	Senft [9]	2014	Swine	24	8 weeks	✓		
3	Krpata [10]	2012	Swine	20	30 days			
4	Vereczkei [11]	2012	Dogs	3	1/3/6 months	✓	✓	
5	Jenkins [12]	2011	Swine	32	2 weeks			
6	Fortelny [13]	2010	Swine	7	4 weeks		✓	
7	Muller-Stich [14]	2008	Swine	9	6 weeks		✓	
8	Otto [15]	2008	Rabbits	20	3 months	✓	✓	
9	Jansen [16]	2007	Rabbits	20	3 months	✓		✓
10	Smith [17]	2007	Swine	18	3/28 weeks	✓	✓	
11	Desai [18]	2006	Dogs	6	12 months		✓	✓
<i>Congenital diaphragmatic hernia repair models</i>								
12	Brouwer [19]	2013	Lambs	7	6 months	✓	✓	
13	Zhao [20]	2013	Rats	52	1, 2, 4, and 6 months	✓	✓	
14	Brouwer [21]	2013	Rats	36	12 weeks	✓	✓	
15	Brouwer [22]	2013	Rats	36	2/12 weeks	✓	✓	
16	Brouwer [23]	2013	Rats	25	2/4/8/12/24 weeks	✓	✓	
17	Gonzalez [24]	2011	Swine	20	6 months	✓	✓	
18	Bohm [25]	2010	Rabbits	33	4 months	✓	✓	
19	Bohm [26]	2010	Rabbits	33	4 months	✓	✓	
20	Urita [27]	2008	Rats	24	1-3 months			
21	Sandovalb [28]	2006	Dogs	11	6 months	✓	✓	✓
22	Suzuki [29]	2002	Dogs	24	15/30 days			
23	Upadhyaya [30]	2001	rat	8	3 weeks	✓	✓	✓
24	Steinau [31]	2000	pigs	24	3/6 months	✓	✓	✓

25	Lantis II [32]	2000	Rabbits	32	6/12 weeks	✓	✓	✓
26	Kimber [33]	2000	Lambs	12	1,3,6 months	✓	✓	
27	Dalla Vecchia[34]	1999	Rats	87	2 weeks - 4 months	✓		✓
28	Lally [35]	1993	Rats	37	400 gr	✓		✓

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164 without any erosion in surrounding hollow viscera [18]. The 8-ply SIS mesh shows a slower rate of
 165 degradation compared to the 4-ply, which can in turn lead to better integration into host tissue [28]. SIS
 166 shows equivalent capillary ingrowth to Alloderm (acellular human cadaveric dermis), but a higher level of
 167 thinning [34]. When compared to PTFE, SIS shows better integration [11], more collagen deposition and
 168 skeletal muscle regeneration and neovascularization [24]. Finally, fascia lata showed superior integration
 169 and capillary ingrowth to ePTFE [29] and, in a separate study, excellent tissue integration and
 170 neovascularization, along with a mild to moderate inflammatory reaction [11].

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172 *3.4.2.5 Migration/erosion potential*

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174 The level of migration and the extent of foreign body reaction were higher when a conventional
 175 polypropylene mesh was used compared to a composite one [14, 15, 16]. The part of the mesh close to
 176 the diaphragm showed less mechanical stability compared to the one close to the esophagus. In a
 177 comparative study of PTFE and SIS in a pig model of congenital diaphragmatic hernia repair, the authors
 178 were able to demonstrate PTFE has a poorer integration into host tissue compared to SIS and tends to
 179 migrate and fold [24].

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181 **3.4.3 Fixation method (sutures/tacks/glue)**

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183 Biologically compatible adhesives like fibrin glue and polyethylene glycol were used with no evidence of
 184 migration, no evidence of any adverse effect to the incorporation of the mesh and equivalent strength to
 185 suture fixation. Krpata *et al.* used an acellular porcine dermal matrix and compared fibrin sealant to
 186 fixation with sutures [10]. Meshes fixed with fibrin glue showed no folding, while there was minimal folding

187 in the control group. Esophagograms did not exhibit any signs of strictures. The authors used a “peel” test
188 to compare the force needed to separate the mesh from the crura and found no difference between the
189 two techniques, whilst the introduction of glue between the crura and the mesh did not result in a
190 significantly different cellular response. Use of fibrin sealant resulted in a significant reduction in operative
191 time. Jenkins *et al.* compared two biological adhesives and found both equally effective in mesh fixation
192 [12]. In conclusion data from 6 experimental studies show that both adhesives seem very promising as an
193 alternative, safe, faster fixation method in hiatal hernia repair.

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195 **3.4.4 Sutured or tension free hiatoplasty**

196 In reviewing the available published studies we did not find any study comparing sutured to tension-free
197 hiatoplasty.

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199 **4. DISCUSSION**

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203 There are a number of controversial points regarding the best surgical technique in hiatal and
204 paraesophageal hernia surgery [36]; the most controversial of which concerns the placement of mesh in
205 the oesophageal hiatus [37-39]. There are reports of a significant reduction in recurrence rates when
206 mesh is used in the surgical repair of hiatal hernia [40, 41]. On the other hand, the surgical community
207 is now conscious that there are important drawbacks in the form of mesh-related complications, reports of
208 which were scarce for two decades and have now begun to appear in the literature [7, 42, 43].

209 **Polypropylene mesh in the hiatus can cause devastating complications including dense fibrosis,**
210 **oesophageal stenosis and intraluminal mesh erosion, the management of which may necessitate a**
211 **reoperation ranging from mesh removal to oesophagectomy [7].** A mesh placed in the diaphragm is

212 subjected to the constant movements of breathing, which are likely to affect its incorporation to the host
213 tissue. Therefore, although there are multiple articles available studying mesh use in animal models for a
214 variety of indications, it is important to evaluate results from animal models of **hiatal-diaphragmatic** hernia
215 repair.

216 Our literature review showed that there were large variations in the quality of experimental studies,
217 only a few of which incorporated histopathological, biomechanical, endoscopic and radiological

217 assessment. The number of animals was small and the implantation time was limited. The surgical
 218 technique used in most cases is a disadvantage, since
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Table 3: Mesh characteristics and surgical technique

Mesh type	Mesh shape	Surgical technique	Fixation method	
<i>Hiatal hernia repair models</i>				
1 [8]	PP/ PET/ PTFE	Circular	Open hiatoplasty and placement of patch in the oesophageal hiatus	Fibrin glue
2 [9]	heavyweight small-porous/heavyweight large-porous/lightweight large-porous PP	Circular	Open hiatoplasty and placement of patch in the oesophageal hiatus	Fibrin glue
3 [10]	acellular porcine dermal matrix	U shaped	Laparoscopic hiatal hernia repair	Sutures/Fibrin sealant
4 [11]	Pericardial and fascia lata patches	Rectangle	3x3 cm patches fixed on muscular part of diaphragm	Polypropylene 3/0
5 [12]	SIS	U shaped	Laparoscopic placement of patch in oesophageal hiatus	Fibrin glue/ polyethylene glycol
6 [13]	Titanium polypropylene mesh	Keyhole	Open placement of the patch without prior hiatoplasty	Fibrin glue
7 [14]	Heavy-weight polypropylene	Circular	Open hiatoplasty and placement of patch in the oesophageal hiatus	Fibrin glue
8 [15]	PP/ PP–polyglactone 25 composite	Circular	Open hiatoplasty and placement of patch in the oesophageal hiatus	Polypropylene 6/0
9 [16]	PP/ PP–polyglactone 25 composite	Circular	Open hiatoplasty and placement of patch in the oesophageal hiatus	Polypropylene 6/0
10 [17]	DualMesh	U shaped	Open transabdominal excision of left hemidiaphragm and open placement of patch in the oesophageal hiatus without prior hiatoplasty	Interrupted ePTFE
11 [18]	SIS	U shaped	Thoracoscopic creation of diaphragmatic hernia and subsequent laparoscopic repair, with hiatoplasty and placement of patch	Interrupted 2/0 polyester
<i>Congenital diaphragmatic hernia repair models</i>				
12 [19]	Collagen-Vicryl	Rectangle	Posterolateral 3x1.5 cm diaphragmatic defect	Running 4/0 prolene
13 [20]	poly(ε-caprolactone) and collagen type I	Rectangle	Excision of 70% of the left hemi-diaphragm (approximately 2-3 cm ²)	Interrupted 6/0 Prolene
14 [21]	Dual layered collagenous scaffolds	Rectangle	12 mm diameter right diaphragm defect	Interrupted 6/0 Prolene
15 [22]	Cross-linked collagenous scaffolds	Rectangle	12 mm diameter right diaphragm defect	Interrupted 6/0 Prolene/ interrupted 5/0 Vicryl
16 [23]	Cross-linked collagenous scaffolds	Rectangle	Excision of 1/3 of the right hemidiaphragm	Interrupted 6/0 Prolene
17 [24]	SIS, ePTFE	Rectangle	Excision of the left hemidiaphragm	Running 3/0 prolene
18 [25]	SIS, PP plus Polyglactone-25, and PP plus polydioxanone and cellulose plus Tachosil	Rectangle	A defect of 1cm in diameter was made into the lateral left diaphragm at the interface of tendon and muscle	Running 5/0 Prolene
19 [26]	SIS, PP plus Polyglactone-25, and PP plus polydioxanone and cellulose plus Tachosil	Rectangle	A defect of 1cm in diameter was made into the lateral left diaphragm at the interface of tendon and muscle	Running 5/0 Prolene
20 [27]	PLGA - collagen mesh	Rectangle	Open transabdominal left hemidiaphragm excision and repair	N/A
21 [28]	SIS	Rectangle	Open transabdominal left central hemidiaphragm excision and repair	N/A
22 [29]	Autologous fascia lata/ ePTFE	Rectangle	Left thoracotomy, left hemidiaphragm excision and repair	N/A
23 [30]	Integra	Rectangle	Open excision of left hemidiaphragm and patch repair	Interrupted 6/0 Vicryl
24 [31]	lyophilized dura/ transverse abdominal bovine	Rectangle	Open excision of left hemidiaphragm and patch repair	Polypropylene 3-0

	pericardial serosa			
25 [32]	SIS/ PTFE	Rectangle	Open transabdominal left hemidiaphragm excision and repair	N/A
26 [33]	PTFE/ fluoropolymer-coated PET	Rectangle	Laparoscopic creation of 2x2 cm defect in left hemidiaphragm and repair	3-0 braided polyester
27 [34]	SIS/AlloDerm	Rectangle	Open transabdominal left central hemidiaphragm excision and repair	N/A
28 [35]	ePTFE/oxidized cellulose/polyglactin 910	Rectangle	Excision of the left hemidiaphragm followed by repair with a patch	Running 4-0 silk.

SIS: small intestinal submucosa; PTFE: polytetrafluoroethylene; PP: polypropylene; PLGA: poly(lactic-co-glycolic acid); PET: polyester

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221 neither the creation of a hiatal hernia nor minimally invasive techniques were used, although similar
 222 experimental models have been described as in the article by Desai *et al.*, where a study incorporating
 223 the creation of a diaphragmatic hernia and its subsequent repair using laparoscopy is presented, in a
 224 model closely resembling the current clinical practice and enabling the surgeon to appreciate the handling
 225 characteristics of each mesh [18]. Finally, due to the heterogeneity of the studies quantitative analysis of
 226 the results was not possible.

227 We evaluated studies regarding specific topics and a number of these were not addressed at all
 228 (infection potential, handling characteristics, sutured/ tension free hiatoplasty), while there was limited
 229 data on the impact of mesh shape and the migration/erosion potential and durability of each mesh.

230 A test of the durability of the hiatoplasty should ideally compare biological/bioabsorbable
 231 prostheses to materials like polypropylene or PTFE, the efficacy of which has been demonstrated in
 232 randomized trials [40, 41]. Biological meshes made from small intestinal submucosa have been shown to
 233 reduce recurrence rates in a randomized trial with short-term follow-up [44] and can also be used as a
 234 control to evaluate newer biological materials. The ability of the mesh to prevent recurrence can be
 235 investigated at autopsy or radiologically. However, new recurrences have been known to occur for a long
 236 time after surgery. Indeed, long term observation of the patients in the previously mentioned trial showed
 237 no benefit in recurrence rates with SIS mesh [45]. The practical limitations of observation time in animal
 238 studies lead authors to perform biomechanical evaluation of mesh materials to evaluate the durability of
 239 the repair. Results confirm the better results obtained clinically with polypropylene compared to biologics,
 240 but are surprising since PTFE was weaker than expected [40].

241 Most of the authors focused on the potential of adhesion formation, biocompatibility and tissue
 242 integration of each mesh and adequate experimental data on several materials is available. The safety
 243 profile of each material i.e. its potential to adhere to and erode into viscera, to cause extended fibrosis

244 resulting in oesophageal stenosis, or to migrate from its position is the most pressing issue, since it is the
245 reason mesh-augmented hiatoplasty is not widely used in clinical practice. Polypropylene meshes
246 showed good integration, but also caused significant adhesions. Experimental studies will be very useful
247 for the comparison of the new generation lightweight meshes, so as to evaluate their advantages
248 compared with standard polypropylene meshes. PTFE resulted in less adhesions, but poor integration
249 with the host tissue. SIS mesh showed an excellent safety profile in experimental studies. These results
250 are in accordance with those obtained from clinical trials including a prospective randomized trial [44] but
251 a marked fibrous response was observed in a previous comparative experimental study, published in
252 abstract form, where significant esophageal stenosis was shown [46]. This finding is significant since
253 esophageal stenosis have been reported in clinical series of patients operated on with SIS mesh [7].
254 Compared to polypropylene, Surgisis induced a milder inflammatory reaction, with slower collagen
255 maturation [25,26]. Fascia lata has been used as a prosthetic material in the hiatus during the 1970s with
256 mixed results, showing efficacy but also some complications [47]. It is, however, in our opinion a very
257 interesting material because it is the only easily obtainable strong autologous patch and has been shown
258 to possess strength equal to PTFE and better incorporation on the diaphragm [28].^[AZF4]

259 The method used to fix the prosthesis to the crura presents a problem since laparoscopic suturing is
260 challenging and time consuming (and potentially risky for the inexperienced) and use of tacks, although
261 fast and effective, places large vessels and the heart at risk of serious injury with potentially catastrophic
262 results [48]. The stability of the mesh depends on both the fixation method but also the material itself and
263 the strength of its incorporation. T-peel testing is an elegant method of quantifying the strength of mesh
264 incorporation [10]. Directly observing the tendency of the mesh to migrate and cause adhesions is
265 tempting, however results must be interpreted with caution; failure of known complications to emerge in
266 these studies could be caused by the relatively short observation time (erosions occur up to nine years
267 after surgery) [7], but could also be interpreted to a lesser extent as proof of the importance of surgical
268 technique (i.e. to strengthen the argument that the reported complications are not inherent in the mesh
269 type but rather are a result of inadequate surgical technique). Indeed, there was a striking difference in
270 mesh migration of polypropylene mesh in the article by Fortelny *et al.* compared to the study of Jansen *et*
271 *al.* [13, 16]; the difference in the ratio of thickness between the mesh and the tissues of the two different

272 animal models was offered as an explanation. In the study by Jansen et al polypropylene meshes of a
273 circular shape were fixed by sutures in a rabbit model, but the meshes had usually moved from their
274 implantation bed and had eroded into the esophagus [16]. However, in another study a circular
275 polypropylene mesh was fixed in place using fibrin glue in a swine experimental model and in this case
276 the authors reached conflicting results as they found the meshes stayed in position and their inner edge
277 had retracted evenly from the esophagus [14].^[AZF5]
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Table 4: Summary of results^[AZF6]

1. Mesh shape : no differences
2. Mesh type
 - a. Infection potential : no differences
 - b. Handling characteristics : no differences
 - c. Durability of repair : Polypropylene stronger than PTFE, biologics
 - d. Adhesion potential, tissue incorporation, fibrosis/stenosis/*shrinkage* potential:
 - More adhesions with synthetic meshes, especially polypropylene. Significant amount of shrinkage for PTFE (34.9%), but also polypropylene.
 - Better integration for composite compared to conventional polypropylene, unstable integration for PTFE.
 - Better tissue integration and regeneration for SIS compared to synthetic materials.
 - Significant shrinkage for Mesh shrinkage was shown to be around 50-70% of original size for polypropylene [9, 14, 16], while the percentage of shrinkage was more higher for the low-weight mesh [9]. When PTFE, polyester and polypropylene were compared, PTFE showed considerably more shrinkage that reached 34.9% of its original size [8].
 - f. Migration/erosion potential : PTFE has a poorer integration compared to

SIS, while composite is better than conventional polypropylene,

3. Fixation method: Biologically compatible adhesives comparable to suture fixation

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281 **5. CONCLUSION**

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The introduction of the new collagen-based biomaterials and the preliminary encouraging results from their use raised great expectations for improved outcomes. The biomaterials from porcine small intestinal submucosa and porcine or human acellular dermis are already widely used in clinical practice (they were being used in 1/3 of all mesh-augmented hiatal diaphragmatic hernia repairs a few years ago [49]) and experimental data are invaluable to further our understanding of their incorporation in host tissue. Although, the reviewed articles study most of the types of meshes currently in clinical use, new biological and bioabsorbable materials are being introduced in clinical practice without any available published experimental data [50, 51].

There is an ever growing need for experimental studies, which should also be well-designed in order to also tackle ethical concerns with regards to animal sacrifice. Studies on mesh-augmented hiatoplasty should include a laparoscopic animal model, biomechanical evaluation and histopathological evaluation of no less than two different biomaterials at the very minimum. In the absence of good quality clinical trials, which are invariably difficult to put together due to the relatively small number of patients, good quality, comparative animal studies are essential in order to identify the mesh with the best safety/efficacy profile, determine the optimal shape and fixation method and enable surgeons to make an inform decision on the merits of using mesh in hiatal hernia

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REFERENCES

1. Finks JF, Wei Y, Birkmeyer JD. The rise and fall of antireflux surgery in the United States. *Surg Endosc.* 2006;20(11): 1698-1701.
2. Mehta S, Bennett J, Mahon D, Rhodes M. Prospective trial of laparoscopic nissen fundoplication versus proton pump inhibitor therapy for gastroesophageal reflux disease: Seven-year follow-up. *J Gastrointest Surg.* 2006;10(9): 1312-6; discussion 1316-7.
3. Hashemi M, Peters JH, DeMeester TR, Huprich JE, Quek M, Hagen JA et al. Laparoscopic repair of large type III hiatal hernia: objective followup reveals high recurrence rate. *J Am Coll Surg.* 2000;190(5): 553-560; discussion 560-551.
4. Soper, N. J. and D. Dunnegan. Anatomic fundoplication failure after laparoscopic antireflux surgery. *Ann Surg.* 1999; 229(5): 669-676; discussion 676-667.
5. Carlson MA, Condon RE, Ludwig KA, Schulte WJ. Management of intrathoracic stomach with polypropylene mesh prosthesis reinforced transabdominal hiatus hernia repair. *J Am Coll Surg.* 1998; 187(3): 227-230.
6. Granderath FA, Carlson MA, Champion JK, Szold A, Basso N, Pointner R et al. Prosthetic closure of the esophageal hiatus in large hiatal hernia repair and laparoscopic antireflux surgery. *Surg Endosc.* 2006;20(3): 367-379.

- 334 7. Stadlhuber RJ, Sherif AE, Mittal SK, Fitzgibbons RJ Jr, Michael Brunt L, Hunter JG et al. Mesh
335 complications after prosthetic reinforcement of hiatal closure: a 28-case series. *Surg Endosc.* 2009;23(6):
336 1219-26.
- 337
- 338 8. Müller-Stich BP, Senft JD, Lasitschka F, Shevchenko M, Billeter AT, Bruckner T et al. Polypropylene,
339 polyester or polytetrafluoroethylene-is there an ideal material for mesh augmentation at the esophageal
340 hiatus? Results from an experimental study in a porcine model. *Hernia.* 2014;18(6):873-81.
- 341
- 342 9. Senft J, Gehrig T, Lasitschka F, Linke GR, Shevchenko M, Bruckner T et al. Influence of weight and
343 structure on biological behavior of polypropylene mesh prostheses placed at the esophageal hiatus. *J*
344 *Laparoendosc Adv Surg Tech A.* 2014;24(6):383-90.
- 345
- 346 10. Krpata DM, Blatnik JA, Harth KC, Phillips MS, Novitsky YW, Rosen MJ. Evaluation of fibrin sealant for
347 biologic mesh fixation at the hiatus in a porcine model. *Surg Endosc.* 2012;Nov;26(11):3120-6.
- 348
- 349 11. Vereczkei A, Varga G, Tornoczky T, Papp A, Horvath ÖP. A new experimental method for hiatal
350 reinforcement using connective tissue patch transfer. *Dis Esophagus.* 2012;Jul;25(5):465-9.
- 351
- 352 12. Jenkins ED, Lerdsirisopon S, Costello KP, Melman L, Greco SC, Frisella MM et al. Laparoscopic
353 fixation of biologic mesh at the hiatus with fibrin or polyethylene glycol sealant in a porcine model. *Surg*
354 *Endosc.* 2011;25(10):3405-13.
- 355
- 356 13. Fortelny RH, Petter-Puchner AH, Glaser KS, Keibl C, Gruber-Blum S, Ohlinger W et al. Fibrin sealant
357 (Tisseel) for hiatal mesh fixation in an experimental model in pigs. *J Surg Res.* 2010;162(1):68-74.
- 358
- 359 14. Müller-Stich BP, Mehrabi A, Kenngott HG, Fonouni H, Reiter MA, Kuttymorotov G et al. Is a circular
360 polypropylene mesh appropriate for application at the esophageal hiatus? Results from an experimental
361 study in a porcine model. *Surg Endosc.* 2009;23(6):1372-1378.

362
363
364
365
366
367
368
369
370
371
372
373
374
375
376
377
378
379
380
381
382
383
384
385
386
387
388

15. Otto J, Kämmer D, Jansen PL, Anurov M, Titkova S, Ottinger A et al. Different tissue reaction of oesophagus and diaphragm after mesh hiatoplasty. Results of an animal study. BMC Surg, 2008;8: 7.

16. Jansen M, Otto J, Jansen PL, Anurov M, Titkova S, Willis S et al. Mesh migration into the esophageal wall after mesh hiatoplasty: comparison of two alloplastic materials. Surg Endosc. 2007;21(12): 2298-2303.

17. Smith GS, Hazebroek EJ, Eckstein R, Berry H, Smith WM, Isaacson JR et al. Evaluation of DualMesh for repair of large hiatus hernia in a porcine model. Surg Endosc. 2008;22(7): 1625-1631.

18. Desai KM, Diaz S, Dorward IG, Winslow ER, La Regina MC, Halpin V et al. Histologic results 1 year after bioprosthetic repair of paraesophageal hernia in a canine model. Surg Endosc 2006;20(11): 1693-1697.

19. Brouwer KM, Daamen WF, Hoogenkamp HR, Geutjes PJ, de Blaauw I, Janssen-Kessels W et al. Collagen-Vicryl scaffolds for reconstruction of the diaphragm in a large animal model. J Biomed Mater Res B Appl Biomater. 2014 May;102(4):756-63.

20. Zhao W, Ju YM, Christ G, Atala A, Yoo JJ, Lee SJ. Diaphragmatic muscle reconstruction with an aligned electrospun poly(ϵ -caprolactone)/collagen hybrid scaffold. Biomaterials. 2013 Nov;34(33):8235-40.

21. Brouwer KM, Daamen WF, Reijnen D, Verstegen RH, Lammers G, Hafmans TG et al. Repair of surgically created diaphragmatic defect in rat with use of a crosslinked porous collagen scaffold. J Tissue Eng Regen Med. 2013 Jul;7(7):552-61

389 22. Brouwer KM, Daamen WF, van Lochem N, Reijnen D, Wijnen RM, van Kuppevelt TH. Construction
390 and in vivo evaluation of a dual layered collagenous scaffold with a radial pore structure for repair of the
391 diaphragm. *Acta Biomater.* 2013 Jun;9(6):6844-51

392

393 23. Brouwer KM, Wijnen RM, Reijnen D, Hafmans TG, Daamen WF, van Kuppevelt TH. Heparinized
394 collagen scaffolds with and without growth factors for the repair of diaphragmatic hernia: construction and
395 in vivo evaluation. *Organogenesis.* 2013 Jul-Sep;9(3):161-7

396

397 24. Gonzalez R, Hill SJ, Mattar SG, Lin E, Ramshaw BJ, Smith CD. Absorbable versus nonabsorbable
398 mesh repair of congenital diaphragmatic hernias in a growing animal model. *J Laparoendosc Adv Surg*
399 *Tech A.* 2011;21(5):449-54.

400

401 25. Böhm G, Binnebösel M, Krähling E, Schumpelick V, Steinau G, Stanzel S et al. Influence of the
402 elasticity module of synthetic and natural polymeric tissue substitutes on the mobility of the diaphragm
403 and healing process in a rabbit model. *J Biomater Appl.* 2011;25(8):771-93.

404

405 26. Böhm G, Steinau G, Krähling E, Schumpelick V, Hermanns-Sachweh B, Stanzel S et al. Is
406 biocompatibility affected by constant shear stress?--comparison of three commercially available meshes
407 in a rabbit model. *J Biomater Appl.* 2011;25(7):721-41

408

409

410 27. Urita Y, Komuro H, Chen G, Shinya M, Saihara R, Kaneko M. Evaluation of diaphragmatic hernia
411 repair using PLGA mesh-collagen sponge hybrid scaffold: an experimental study in a rat model. *Pediatr*
412 *Surg Int.* 2006;24(9): 1041-1045.

413

414 28. Sandoval JA, Lou D, Engum SA, Fisher LM, Bouchard CM, Davis MM et al. The whole truth:
415 comparative analysis of diaphragmatic hernia repair using 4-ply vs 8-ply small intestinal submucosa in a
416 growing animal model. *J Pediatr Surg.* 2006;41(3): 518-523.

417

418 29 Suzuki K, Takahashi T, Itou Y, Asai K, Shimota H, Kazui T. Reconstruction of diaphragm using
419 autologous fascia lata: an experimental study in dogs. *Ann Thorac Surg.* 2002;74(1): 209-212.
420

421 30. Upadhyaya M, Orford JE, Smith N, Barker A, Gollow I. Incorporation of Integra in tissue defects: a
422 pilot study in the rat model. *Pediatr Surg Int.* 2007;23(7):669-73.
423
424

425 31. Steinau G, Dreuw B, Schleef J, Lawong G, Schumpelick V. Short-term absorbable material for
426 diaphragmatic replacement. *Pediatr Surg Int.* 2000;16(1-2):19-22.
427

428 32. Lantis JC 2nd, Gallivan EK, Hekier R, Connolly R, Schwaitzberg SD, Crombleholme T. A comparison
429 of collagen and PTFE patch repair in a rabbit model of congenital diaphragmatic hernia. *J Invest Surg.*
430 2000;13(6): 319-325.
431

432 33. Kimber CP, Dunkley MP, Haddock G, Robertson L, Carey FA, Cuschieri A. Patch incorporation in
433 diaphragmatic hernia. *J Pediatr Surg.* 2000;35(1):120-3.
434

435 34. Dalla Vecchia, Engum S, Kogon B, Jensen E, Davis M, Grosfeld J. Evaluation of small intestine
436 submucosa and acellular dermis as diaphragmatic prostheses. *J Pediatr Surg.* 1999;34(1): 167-171.
437

438 35. Lally KP, Cheu HW, Vazquez WD. Prosthetic diaphragm reconstruction in the growing animal. *J*
439 *Pediatr Surg.* 1993;28(1):45-7.
440

441 36. Neufeld, M., Graham A. Levels of evidence available for techniques in antireflux surgery. *Dis*
442 *Esophagus.* 2007;20(2): 161-7.
443

444 37. Targarona EM, Bendahan G, Balague C, Garriga J, Trias M. Mesh in the hiatus: a controversial issue.
445 *Arch Surg.* 2004;139(12): 1286-96; discussion 1296.
446

- 447 38. Smith, G. Mesh repairs in hiatal surgery. The case for mesh repairs in hiatal surgery. *Ann R Coll Surg*
448 *Engl.* 2007;89(5): 481-3.
449
- 450 39. Kelty, C. J., Falk, G. L. Mesh repairs in hiatal surgery. The case against mesh repairs in hiatal
451 surgery. *Ann R Coll Surg Engl.* 2007;89(5): 479-81.
452
- 453 40. Frantzides CT, Madan AK, Carlson MA, Stavropoulos GP. A prospective, randomized trial of
454 laparoscopic polytetrafluoroethylene (PTFE) patch repair vs simple cruroplasty for large hiatal hernia.
455 *Arch Surg.* 2002;137(6): 649-52.
456
- 457 41. Antoniou SA, Antoniou GA, Koch OO, Pointner R, Granderath FA. Lower recurrence rates after mesh-
458 reinforced versus simple hiatal hernia repair: a meta-analysis of randomized trials. *Surg Laparosc Endosc*
459 *Percutan Tech.* 2012;22(6):498-502.
460
- 461 42. Carlson, M. A., Frantzides, C. T. Complications and results of primary minimally invasive antireflux
462 procedures: a review of 10,735 reported cases. *J Am Coll Surg.* 2001;193(4): 428-39.
463
- 464 43. Tatum RP, Shalhub S, Oelschlager BK, Pellegrini CA. Complications of PTFE mesh at the
465 diaphragmatic hiatus. *J Gastrointest Surg.* 2008;12(5): 953-7.
466
- 467 44. Oelschlager BK, Pellegrini CA, Hunter J, Soper N, Brunt M, Sheppard B et al. Biologic prosthesis
468 reduces recurrence after laparoscopic paraesophageal hernia repair: a multicenter, prospective,
469 randomized trial. *Ann Surg.* 2006;244(4):481-90.
470
- 471 45. Oelschlager BK, Pellegrini CA, Hunter JG, Brunt ML, Soper NJ, Sheppard BC et al. Biologic
472 prosthesis to prevent recurrence after laparoscopic paraesophageal hernia repair: long-term follow-up
473 from a multicenter, prospective, randomized trial. *J Am Coll Surg.* 2011;213(4):461-8.
474

- 475 46. Halpin V, Meyers BF, Luttmann D. Laparoscopic paraesophageal hiatal hernia repair using
476 prosthetics in a canine model. *Surg Endosc.* 2002;16(Suppl 1): S977
477
- 478 47. Frantzides CT, Welle SN. Cardiac tamponade as a life-threatening complication in hernia repair.
479 *Surgery.* 2012 Jul;152(1):133-5.
480
- 481 48. Brain, R. H., Maynard, J. Fascia lata graft repair of esophageal hiatal hernia. *Am J Surg.*
482 1968;115(4): 488-501.
483
- 484 49. Frantzides CT, Carlson MA, Loizides S, Papafili A, Luu M, Roberts J, Zeni T, Frantzides A. Hiatal
485 hernia repair with mesh: a survey of SAGES members. *Surg Endosc.* 2010 May;24(5):1017-24.
486
- 487 50. Zehetner J, Lipham JC, Ayazi S, Oezcelik A, Abate E, Chen W et al. A simplified technique for
488 intrathoracic stomach repair: laparoscopic fundoplication with Vicryl mesh and BioGlue crural
489 reinforcement. *Surg Endosc.* 24(3): 675-9.
490
- 491 51. Powell BS, Wandrey D, Voeller GR. A technique for placement of a bioabsorbable prosthesis with
492 fibrin glue fixation for reinforcement of the crural closure during hiatal hernia repair. *Hernia.*
493 2013;17(1):81-4