

Complex Ventral Hernia Repair in Contaminated Fields: A Propensity-Score Matched Analysis of Long-term Quality-of-Life and Outcomes Between Different Prostheses.

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Disclosures

- **FM:** Intuitive, Medtronic, BD, Integra, Allergan and DeepBlue
- The rest of the authors have no relevant disclosures.

Background

- Optimal hernia repair in contaminated fields is controversial

Durability



Complications

- Unclear which approach better improves **patient satisfaction**



JAMA Surgery | **Original Investigation**

Biologic vs Synthetic Mesh for Single-stage Repair of Contaminated Ventral Hernias

A Randomized Clinical Trial

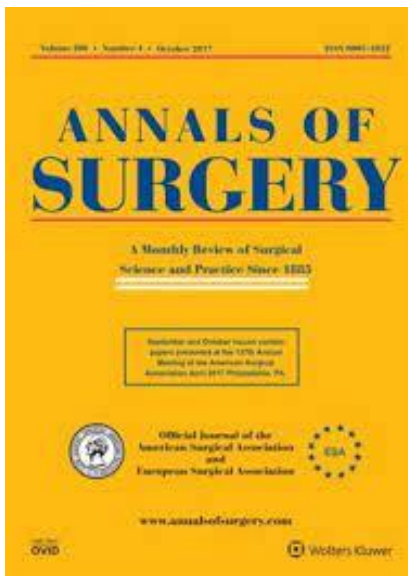
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RANDOMIZED CONTROLLED TRIAL

Preventing Recurrence in Clean and Contaminated Hernias Using Biologic Versus Synthetic Mesh in Ventral Hernia Repair

The PRICE Randomized Clinical Trial

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Permanent vs Absorbable Mesh for Ventral Hernia Repair in Contaminated Fields: Multicenter Propensity-Matched Analysis of 1-Year Outcomes Using the Abdominal Core Health Quality Collaborative Database


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2023 SAGES POSTER

Slowly absorbable mesh in sublay ventral hernia repair in contaminated fields

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Study Question

- Is there a long-term difference in patient-reported outcomes between different prostheses in contaminated ventral hernia repair?

Objective

- Evaluate long-term outcomes of complex ventral hernia repair in contaminated fields focusing on:

Quality of life

Pain scores

- Measured at baseline and 1, 3, 6, 12 months from baseline

Methods

- Abdominal Core Health Quality Collaborative (ACHQC)
- 118,868 patients
- 460 institutions from the United States
- Follow-up available up

Methods

- Inclusion criteria:
 - Ventral hernia repairs with mesh (2013–2023)
 - CDC class II–IV
 - Elective operations
 - 30-day follow up information

118,868 ACHQC patients
2013-2023

40,228 patients underwent ventral
hernia repair (VHR)

4,313 patients underwent VHR in
contaminated (CDC II-IV) fields

1,073 patients with available follow-up and
PRO information

Permanent mesh
N=920

Absorbable Synthetic
N=56

Biologic mesh
N=97

Outcomes

PROMIS 3A Pain
intensity

In the last 7 days: experienced?

- How would you rate your pain on average?
- What is the highest level of pain you experienced?
- How would you rate your current pain?

Outcomes

**PROMIS 3A Pain
intensity**

- NIH-Patient Reported Outcomes Measurement Information System
- 3 questions
- T distribution (Mean 50 / SD 10)
- Higher scores = increased pain
- Validated in hernia studies

Outcomes

HerQLes Questionnaire

- Hernia Related Quality of Life (QoL) Survey
- Score 0-100
- 15.6 points considered “minimal clinically relevant difference”
- Higher scores = increased QoL

Krpata et. Al JACS

(2015)
Renshaw et. Al JACS

Validated in hernia



For the following statements, please circle the number that is most appropriate for you.

	<i>Strongly Disagree</i>	<i>Moderately Disagree</i>	<i>Slightly Disagree</i>	<i>Slightly Agree</i>	<i>Moderately Agree</i>	<i>Strongly Agree</i>
1. My abdominal wall has a huge impact on my health	1	2	3	4	5	6
2. My abdominal wall causes me physical pain	1	2	3	4	5	6
3. My abdominal wall interferes when I perform strenuous activities, e.g. heavy lifting	1	2	3	4	5	6
4. My abdominal wall interferes when I perform moderate activities, e.g. bowling, bending over	1	2	3	4	5	6
5. My abdominal wall interferes when I walk or climb stairs	1	2	3	4	5	6
6. My abdominal wall interferes when I dress myself, take showers and cook	1	2	3	4	5	6
7. My abdominal wall interferes with my sexual activity	1	2	3	4	5	6
8. I often stay at home because of my abdominal wall	1	2	3	4	5	6
9. I accomplish less at home because of my abdominal wall	1	2	3	4	5	6
10. I accomplish less at work because of my abdominal wall	1	2	3	4	5	6
11. My abdominal wall affects how I feel every day	1	2	3	4	5	6
12. I often feel blue because of my abdominal wall	1	2	3	4	5	6

Results

- A total of 1,073 patients included
- Median age 62 years (IQR 53–70)
- 8% of active smokers
- Median BMI 32 (IQR 28–36)
- 48% recurrent hernias (median of 2 prior repairs)

General characteristics of the study cohort

Type of approach , N (%)

Open	899 (84)
Laparoscopic	49 (5)
Robotic	125 (12)

Wound class, N (%)

II	679 (63)
III	358 (33)
IV	36 (3)

Defect size in cm (median, IQR)

Width	20 (12-25)
Length	13 (8-16)

Underwent TAR, N (%)

Yes	716 (67)
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Mesh position, N (%)

Onlay	74 (7)
Inlay	27 (3)
Sublay	972 (91)

Type of sublay repair, N (%)

Retro-rectus	826 (85)
Preperitoneal	434 (45)
Intraperitoneal	116 (12)

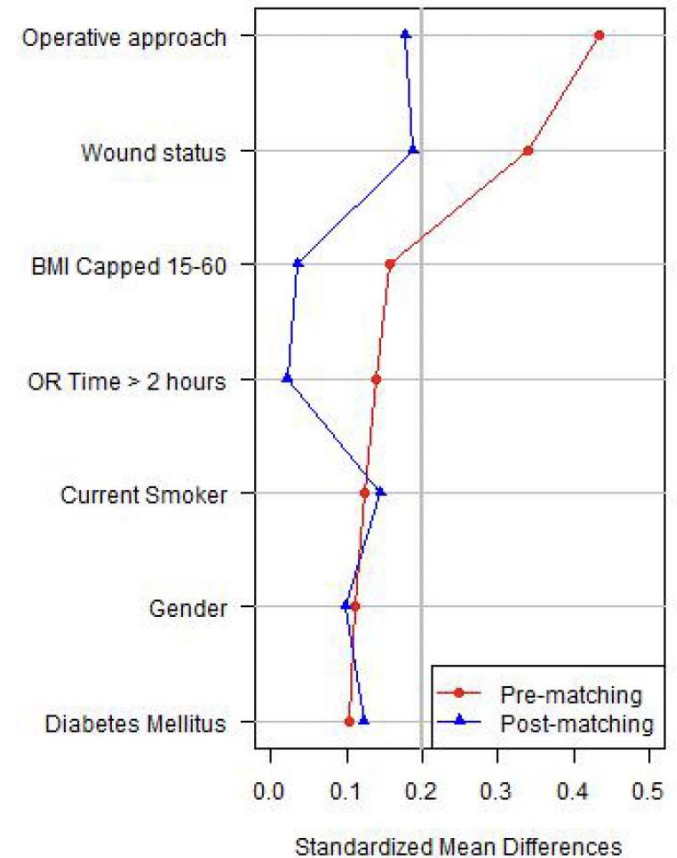
General characteristics of the study cohort

	Permanent (N=920)	Absorbable Synthetic (N=56)	Biologic (N=97)	p-value
Age (median, IQR)	62 (53-70)	60 (51-64)	59 (47-69)	0.009
Type of approach , N (%)				
Open	772 (84)	50 (89)	95 (98)	<0.001
MIS	148 (16)	6 (11)	2 (2)	
Wound class, N (%)				
II	597 (65)	35 (62)	47 (48)	<0.001
III	310 (34)	16 (29)	32 (33)	
IV	13 (1)	5 (9)	18 (19)	
Defect size in cm (median, IQR)				
Width	20 (13-25)	17 (7-25)	17 (12-23)	0.039
Length	13 (8-16)	10 (6-15)	12 (8-15)	0.44
Mesh position, N (%)				
Onlay	66 (7)	6 (11)	2 (2)	<0.001
Inlay	10 (1)	11 (20)	6 (6)	
Sublay	844 (92)	39 (70)	89 (92)	

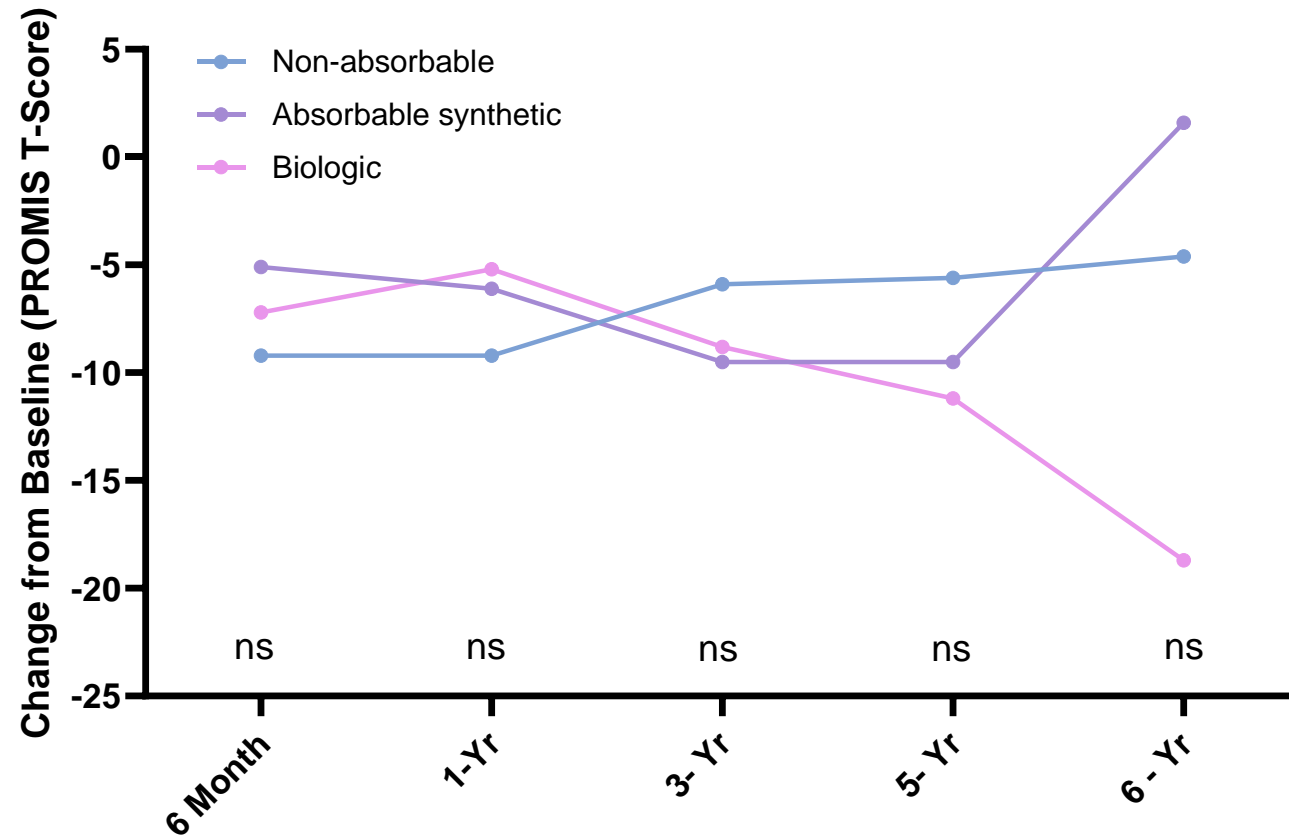
Propensity-score matching

- A total of 280 patients included

- Gender
- Diabetes
- BMI
- Smoking status
- Operative times
- Surgical approach



PROMIS 3A Pain



- No differences in patient reported pain at 6 months, 1-, 3-, 5-, and 6-year follow-up

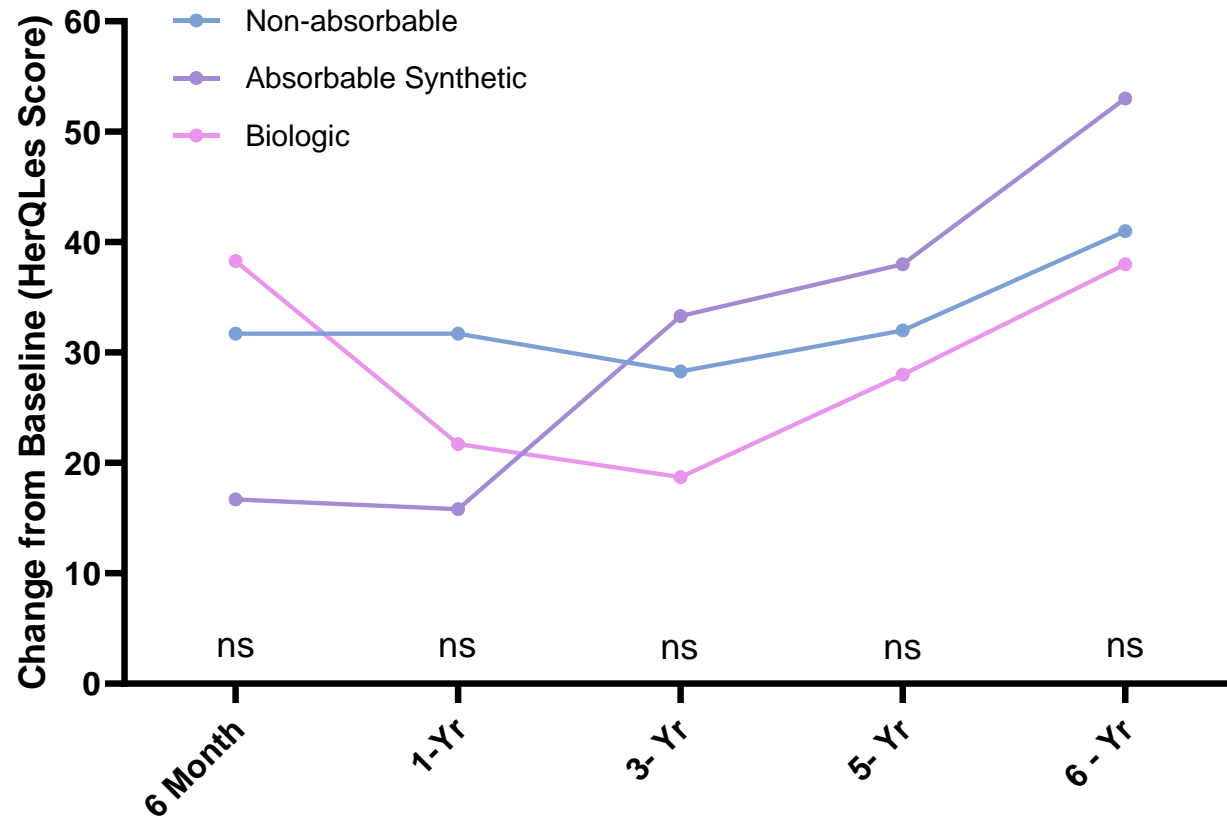
N=125
N=26

N=114

N=12

N=55

HerQLes Score



N=125
N=26

N=114

N=12

N=55

- No differences in quality of life at 6 months, 1-, 3-, 5-, and 6- year follow-up

Results

Other postoperative outcomes in the propensity-matched cohort.

	Permanent (N=168)	Absorbable Synthetic (N=56)	Biologic (N=56)	p-value
30-day readmission, N (%)	15 (9)	6 (11)	9 (16)	0.33
30-day SSI, N (%)	15 (9)	3 (5)	7 (12)	0.41
30-day SSO, N (%)	23 (14)	5 (9)	8 (14)	0.61
1-Year Recurrence, N (%)	13 (16)	7 (35)	13 (45)	0.006

Limitations

- Retrospective data
- Heterogeneity in patient characteristics
- Different grades of contamination among wound classes
- Limited number of patients with longer-term follow-up
- Abdominal wall specialists entering data into

Conclusions

- No differences in quality of life or patient-reported pain scores by type of mesh in contaminated ventral hernia surgery at up to 6-year follow-up
- No differences in other postoperative outcomes

Take home message

- This multicentric study, with extended follow-up data, continues to support that the use of mesh in contaminated fields is safe and effective in improving patient symptoms and quality of life

Thank you!

