

MESHES IN CONTAMINATED FIELDS

OVERVIEW

JP Palot, Y Renard (Reims)







WHICH TYPE OF MESH? A HEAVILY DEBATED TOPIC ! NO CONSENSUS



Wound	Class definition
l (clean)	An uninfected operative wound in which no inflammation is encountered and the respiratory, alimentary, genital, or uninfected urinary tract are not entered.
II (clean- contaminated)	An operative wound in which the respiratory, alimentary, genital, or urinary tracts are entered under controlled conditions and without unusual contamination
III (contaminated)	Open, fresh, accidental wounds. Operations with major breaks in sterile technique or gross spillage from the gastrointestinal tract, and incisions in which acute, non-purulent inflammation are encountered.
IV (dirty-infected)	Old traumatic wounds with retained devitalized tissue and those that involve existing clinical infection or perforated viscera.

• Very few surgeons were using meshes

• Very few meshes

polypropylene



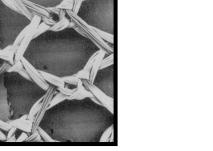
- Few solutions : repair without mesh or with an absorbable mesh

Our experience of Vicryl mesh: SSI=45.6% (superficial 26.3% deep=19.3%) 24 month recurrences: 68.5%

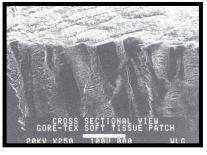








polyester



ePTFE







OFTEN A CATASTROPHIC SITUATION

Length of hospital stay, pain, re-admissions, re-interventions and sometimes death

RATHER A RECURRENCE THAN A MESH INFECTION



- SURGERY
- BETTER KNOWLEDGE OF MESH BIOLOGICAL BEHAVIOR
- NEW MESHES :
 - lightweight synthetic
 - biologic
 - biosynthetic absorbable

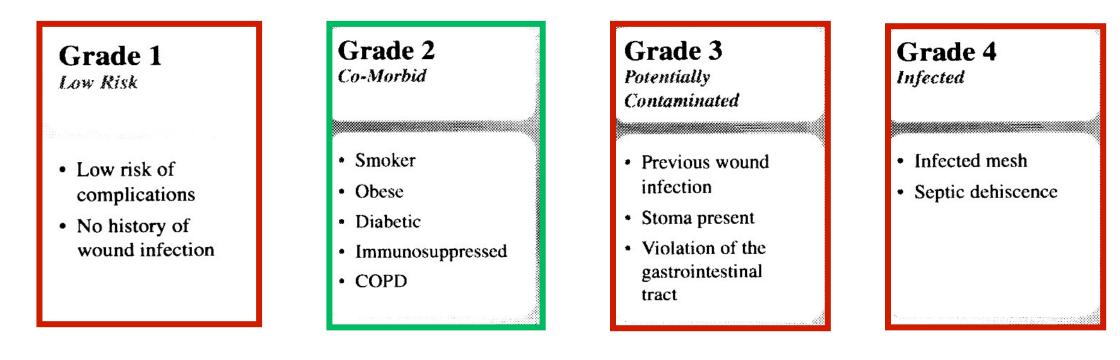


• NEW TOOLS for the treatment of SSI

Incisional ventral hernias: Review of the literature and recommendations regarding the grading and technique of repair Surgery 2010

The Ventral Hernia Working Group

The Ventral Hernia Working Group: Karl Breuing, MD,^a Charles E. Butler, MD, FACS,^b Stephen Ferzoco, MD, FACS,^a Michael Franz, MD,^c Charles S. Hultman, MD, MBA, FACS,^d Joshua F. Kilbridge,^e Michael Rosen, MD,^f Ronald P. Silverman, MD, FACS,^g and Daniel Vargo, MD, FACS,^h Boston, MA, Houston, TX, Ann Arbor, MI, Chapel Hill, NC, San Francisco, CA, Cleveland, OH, Baltimore, MD, and Salt Lake City, UT



SYNTHETIC MESH

SYNTHETIC MESH?

NO SYNTHETIC MESH

Surg Infect (Larchmt). 2011 Jun;12(3):205-10. Epub 2011 Jul 18.

Mesh infection in ventral incisional hernia repair: incidence, contributing factors, and treatment.

Sanchez VM, Abi-Haidar YE, Itani KM.

Department of Surgery, Veterans Affairs Boston Healthcare System and Boston University, Boston, Massachusetts.

• review of literature

• risk factors :

- obesity
- COPD
- Aortic aneurysm
- microporous mesh or ePTFE
- length of operation
- concomitant procedure
- violation of gastro-intestinal tract
- entero cutaneous fistula

Use of Mesh During Ventral Hernia Repair in Clean-Contaminated and Contaminated Cases: Outcomes of 33,832 Cases.

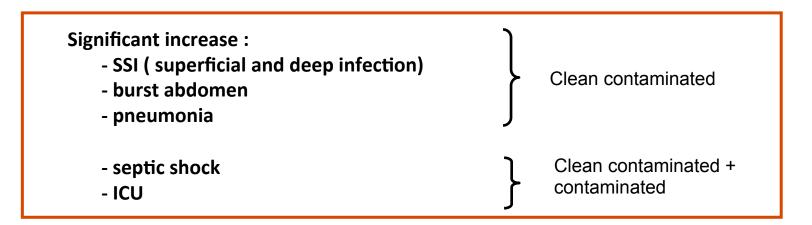
Choi JJ, Palaniappa NC, Dallas KB, Rudich TB, Colon MJ, Divino CM.

Source

Ann Surg 2011

From Division of General **Surgery**, Department of **Surgery**, Mount Sinai School of Medicine, New York.

National Surgical Quality Improvement Program



CONCLUSION:

There is a significant increase in risk of postoperative occurrences following VHRs using mesh in **clean-contaminated** and contaminated cases relative to **clean** cases. We recommend avoiding the use of mesh in any level of contamination.

BIOLOGIC ACELLULAR XENOGRAFTS

Holl-Allen RT (1984) Porcine dermal collagen repair of inguinal hernias. J R Coll Surg Edinb 29(3):154–157

Sarmah BD (1984) Porcine dermal collagen repair of incisional herniae. Br J Surg 71(7):524–525

LATE 90'S : ABDOMINAL WALL REPAIR IN POTENTIALLY CONTAMINATED / CONTAMINATED FIELDS

- very very (too much) expensive
- no (not yet) strong evidence
- published studies generally very heterogeneous : several types of meshes, lot of different techniques of repair, different indications

Different biologic meshes :

- cross linking or not
- different industrial processes
- variable quality of meshes

Biologic mesh in ventral hernia repair: Outcomes, recurrence, and charge analysis Surgery 2016

Ciara R. Huntington, MD, Tiffany C. Cox, MD, Laurel J. Blair, MD, Samuel Schell, BS, David Randolph, BS, Tanushree Prasad, MA, Amy Lincourt, PhD, MBA, B. Todd Heniford, MD, FACS, and Vedra A. Augenstein, MD, FACS, *Charlotte, NC*

	$\begin{array}{l} Alloderm\\ n = 40 \end{array}$	AlloMax n = 23	FlexHD n = 70	Strattice n = 68	$\begin{array}{l} Xenmatrix\\ n = 22 \end{array}$	
	% or means ± SD	% or means ± SD	% or means ± SD	% or means ± SD	% or means ± SD	P value
Hematoma	3.45	0	0	1.5	0	.58
Intraabdominal abscess	6.9	9.1	4.6	7.6	14.3	.61
Prolonged incisional pain	0	0	2.9	4.4	18.2	.02
Acute kidney injury	7.5	4.4	10.0	7.4	13.6	.81
Sepsis	10.0	8.7	4.3	4.4	13.6	.35
Seroma	37.9	13.6	34.9	15.2	31.8	.02
Wound dehiscence	3.5	13.6	21.2	10.6	9.5	.17
Postoperative SBO	0	4.6	4.8	3.1	0	.84
Postoperative wound vac	20.0	13.6	33.3	20.9	23.8	.29
Wound requires intervention	37.9	13.6	33.9	19.7	23.8	.12
Wound infection	44.8	27.2	43.9	29.9	33.3	.33
Mesh infection	0	0	1.4	1.5	0	1
Readmission within 30 d	20	13	24.3	20.6	22.7	,84
Reoperation within 30 d	15	8.7	12.9	16.2	13.6	.92
Inpatient mortality	2.6	0	1.4	1.5	0	1
Hernia recurrence	35.0	34.8	37.1	14.7	59.1	.001
Duration of stay (d)	9.3 ± 8.9	12.4 ± 11.4	13.1 ± 18.2	10.3 ± 8.6	16.1 ± 21.8	.47
Average follow-up (mo)	26.5 ± 37.0	17.0 ± 19.8	18.4 ± 22.1	17.6 ± 15.7	11.0 ± 13.1	.74
lean hospital charges (US dollars ± SD)						
Mesh charge	25,825 ± 23,535	13,311 ± 11,438	$22,981 \pm 30,709$	$31,875 \pm 17,960$	$48,955 \pm 28,580$	<.0001
Operating room charge*	$5,407 \pm 6,395$	$10,672 \pm 6,115$	$9,929 \pm 6,140$	$19,939 \pm 13,934$	$24,520 \pm 26,971$	<.001
Total hospital charge	$100,797 \pm 94,330$	$111,775 \pm 103,547$	$113,819 \pm 154,947$	$140,394 \pm 80,709$	$221,966 \pm 263,494$.02

*Operating room charge does not include mesh charge. Numbers represent % unless otherwise specified.

ASA, American Society of Anesthesiology; COPD, chronic obstructive pulmonary disease; GERD, gastroesophageal reflux disease; SBO, small bowel obstruction; SD, standard deviation; wound vac, wound vacuum negative pressure device.

A 5-Year Clinical Experience With Single-Staged Repairs of Infected and Contaminated Abdominal Wall Defects Utilizing Biologic Mesh

Michael J. Rosen, MD, David M. Krpata, MD, Bridget Ermlich, RN, and Jeffrey A. Blatnik, MD 201

Long-term outcomes (>5-year follow-up) with porcine acellular dermal matrix (PermacolTM) in incisional hernias at risk for infection

M. M. Abdelfatah · N. Rostambeigi · E. Podgaetz · M. G. Sarr

2013

Prospective study of single-stage repair of contaminated hernias using a biologic porcine tissue matrix: The RICH Study

Kamal M. F. Itani, MD, FACS,^a Michael Rosen, MD, FACS,^b Daniel Vargo, MD, FACS,^c 2012 Samir S. Awad, MD, FACS,^d George DeNoto III, MD, FACS,^e Charles E. Butler, MD, FACS,^f and the RICH Study Group, Boston, MA, Cleveland, OH, Salt Lake City, UT, Houston, TX, and Lake Success, NY

- 128 patients
- non cross-linked porcine matrix (Strattice)
- SSI : 47% no explantation
- follow up :26 months
- recurrences : 31,3%
- 65 patients
- cross linked porcine matrix (Permacol)
- different procedures (bridging+++)
- 59 patients followed > 5 ans
- explantation :25%
- recurrences : 46 à 80%

		Incidence of recurrences, % (n/N)		
80 patients	Type of repair	12 months	24 months	
Strattice	Fascial closure with	14 (9/64)	23 (15/64)	
SSI : 30%	and without CST (defect area: 203 ± 150 cm ²)			
no explantation	With CST (defect area: $220 \pm 150 \text{ cm}^2$)	17 (9/52)	27 (14/52)	
follow up: 2 years Recurrences 28%	Without CST (defect area: 126 ± 121 cm ²)	0 (0/12)	8 (1/12)	
	No fascial closure with and without CST (defect area: 355 ± 141 cm ²)	38 (6/16)	44 (7/16)	
	All intent-to-treat patients	19 (15/80)	28(22/80)	

STUDY PROTOCOL

Open Access

Use of biological mesh versus standard wound care in infected incisional ventral hernias, the SIMBIOSE study: a study protocol for a randomized multicenter controlled trial

Christophe Mariette^{1,2,3,6*}, Nicolas Briez^{1,2}, Fanette Denies⁴, Benoît Dervaux^{2,5}, Alain Duhamel^{2,5}, Marie Guilbert¹, Emilie Bruyère¹, William B Robb¹, Guillaume Piessen^{1,2,3} and on behalf of FRENCH

inclusions closed on 7/01/2016

Outcomes of Synthetic Mesh in Contaminated VentralHernia RepairsJ Am Coll Surg 2013

Alfredo M Carbonell, DO, FACS, Cory N Criss, MD, William S Cobb, MD, FACS, Yuri W Novitsky, MD, Michael J Rosen, MD, FACS

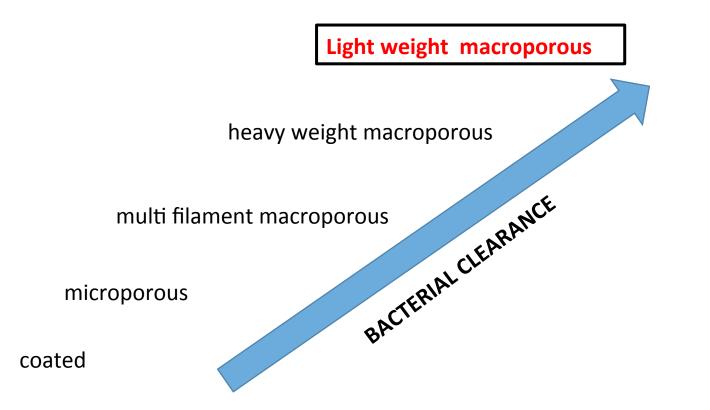
- Retrospective study on prospective databases
- Clean-contaminated/contaminated cases
- Bioburden-reduction
- Lightweight polypropylene mesh
- Sublay technique 100%
- Complete fascial closure 91%

Table 4. Surgical Site Occurrence, Surgical Site Infection, and Hernia Recurrence Rates of Patients Undergoing Clean-Contaminated and Contaminated Hernia Repairs

SSO			30-Day SSI		이 이 가장에서 있는 것은 이렇지 않아야 한 것을 가 많다. 것이 가지 않는 것이 있다.	Recurrence			
CDC wound classification	n	%	Frequency, %	n	%	Frequency, %	n	%	Frequency, %
Clean contaminated $(n = 42)$	11	26.2	11.0	3	7.1	3.0	3	7.1	3.0
Contaminated $(n = 58)$	20	34	20.0	11	19.0	7.0	4	6.8	4.0

SSI, surgical site infection; SSO, surgical site occurrence.

WHAT ABOUT LIGHT-WEIGHT SYNTHETIC MESHES ?



Central rupture and bulging of low-weight polypropylene mesh following recurrent incisional sublay hernioplasty.

Zuvela M¹, Galun D, Djurić-Stefanović A, Palibrk I, Petrović M, Milićević M. Hernia 2014

Central failures of lightweight monofilament polyester mesh causing hernia recurrence: a cautionary note

C. C. Petro · E. H. Nahabet · C. N. Criss · S. B. Orenstein · H. A. von Recum · Y. W. Novitsky · M. J. Rosen

Hernia 2015

Active *Staphylococcus aureus* infection: Is it a contra-indication to the repair of complex hernias with synthetic mesh? A prospective observational study on the outcomes of synthetic mesh replacement, in patients with chronic mesh infection caused by *Staphylococcus aureus*

Claudio Birolini^{*}, Jocielle Santos de Miranda, Edivaldo Massazo Utiyama, Samir Rasslan, Dario Birolini

General and Trauma Surgery, Department of Surgery, University of São Paulo, School of Medicine, São Paulo, Brazil Int J Surg 2016

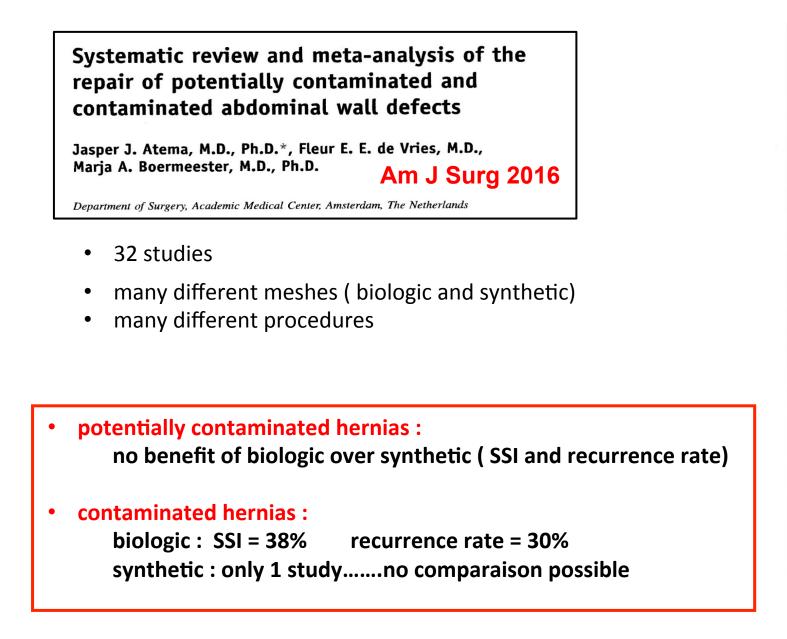
- Prospective observational study (2006-2014)
- 22 patients with chronic mesh infection
- Positive culture for Staphylococcue Aureus
- Mesh removal
- Restoration of the midline
- Heavy-weight large pore polypropylene mesh
- Onlay
- SSO : 8 = 36.7%
- SSI : 6 = 27.3%
- Deaths : 2 = 9%

• Recurrence : 1 = 5%

• Chronic sinus: 1 = 5%

5. Conclusion

Synthetic mesh replacement in patients suffering from CMI caused by SA has an acceptable incidence of postoperative wound infection and prevents hernia recurrence. Large-pore mono-filament polypropylene mesh is a suitable material to be used in the infected surgical field as an onlay graft.



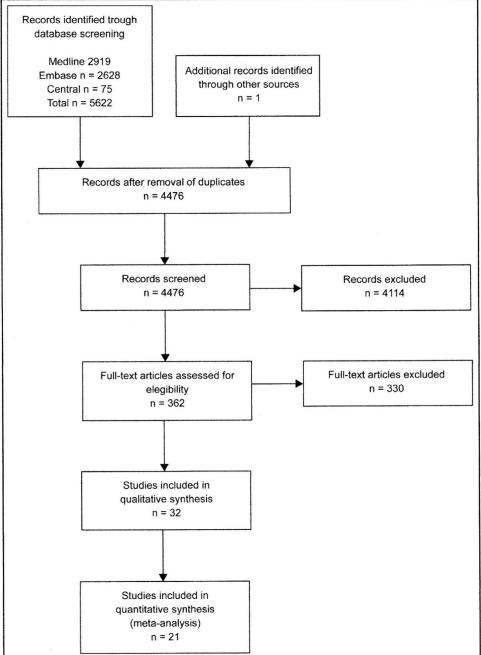
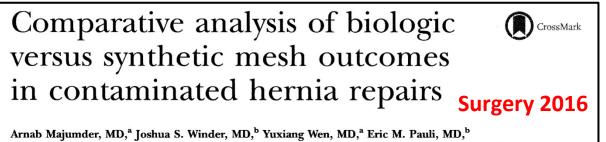


Figure 1 Flow diagram depicting search and study selection.



Igor Belyansky, MD,^c and Yuri W. Novitsky, MD,^a Cleveland, OH, Hershey, PA, and Annapolis, MD

- multicentric retrospective study
- clean contaminated/contaminated
- sublay implantation in > 98%cases

«Finally, given a significant
 variability of patient and surgeon
 treatment goals in contaminated
 repairs as well variation of
 technique and mesh selection,our
 findings may not be applicable to all
 clinical situations

	Biologic (n = 69)	Synthetic ($n = 57$)	P value
Wound morbidity			
SSE	29 (42.0%)	13 (22.8%)	.024
Seroma	3 (4.3%)	2 (3.5%)	1.000
Hematoma	1(1.4%)	1 (1.8%)	1.000
Soft tissue breakdown	2 (2.9%)	1(1.8%)	1.000
Cellulitis	1(1.4%)	2 (3.5%)	1.000
SSI	22 (31.9%)	7 (12.3%)	.010
Superficial	4 (5.7%)	3 (5.3%)	1.000
Deep	15 (21.7%)	3 (5.3%)	.009
Organ space	3 (4.3%)	1 (1.8%)	626
Postoperative outcomes			
Mesh explantation	2 (2.9%)	1 (1.8%)	1.000
Hernia recurrence	15 (26.3%)	4 (8.9%)	.039
Follow-up (mo)	21.5 ± 10.5	18.4 ± 9.6	.265
Follow-up duration >12 mo	57 (82.6%)	45 (78.9%)	.602
Duration of stay (days)	10.8 ± 6.4	7.7 ± 3.1	.002
90-day readmission	15 (21.7%)	4 (7.0%)	.025
Wound morbidity	13 (87%)	3 (75%)	.530
Other	2 (13%)	1 (25%)	.530

Values are mean ± standard deviation (SD) or number (percentage)

BIO-SYNTHETIC ABSORBABLE MESHES

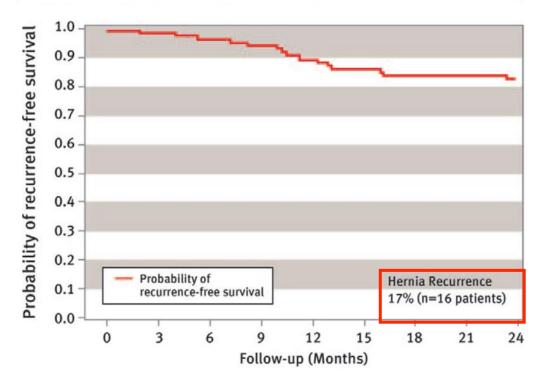
Multicenter, Prospective, Longitudinal Study of the Recurrence, Surgical Site Infection, and Quality of Life After Contaminated Ventral Hernia Repair Using Biosynthetic Absorbable Mesh: The COBRA Study

Rosen, Michael J. MD; Bauer, Joel J. MD; Harmaty, Marco MD; Carbonell, Alfredo M. DO; Cobb, William S. MD; Matthews, Brent MD; Goldblatt, Matthew I. MD; Selzer, Don J. MD, MS; Poulose, Benjamin K. MD, MPH; Hansson, Bibi M. E. MD, PhD; Rosman, Camiel MD; Chao, James J. MD; Jacobsen, Garth R. MD

Ann Surg 2017

- 104patients with ventral hernia repair
- clean contaminated : 23%
- contaminated : 77%
- clean and dirty excluded
- GORE BIO A
- sublay or underlay
- midline closure
- Follow up : 24 months

Variables	n = 104
Wound events*, n (%)	33 (28)
Surgical site infection [†]	21 (18)
Seroma	6 (6)
Fistula	2 (2)
Bowel obstruction	2 (2)
Wound dehiscence	1 (1)
Hematoma	1 (1)
Postoperative infections [†] , n (%)	21 (18)
Superficial incisional infections	9 (9)
Deep incisional infections	10 (10)
Organ space infections	2 (2)



CONCLUSIONS ?

Centers for Disease Control (CDC) wound classification

Wound	Class definition
l (clean)	An uninfected operative wound in which no inflammation is encountered and the respiratory, alimentary, genital, or uninfected urinary tract are not entered.
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contaminated)	contamination
III (contaminated)	Open, fresh, accidental wounds. Operations with major breaks in sterile technique or gross spillage from the gastrointestinal tract, and incisions in
	which acute, non-purulent inflammation are encountered.
IV (dirty-infected)	Old traumatic wounds with retained devitalized tissue and those that involve existing clinical infection or perforated viscera.

- **CLEAN CONTAMINATED** : macroporous synthetic meshes may probably be used preferably in sublay position
- CONTAMINATED / DIRTY NO CONSENSUS
 - two-step procedure : no mesh or absorbable mesh.....
 - biological mesh expecting a 20 30% recurrence rate
 - synthetic (lightweight?) mesh
 - biosynthetic absorbable mesh.....the future?