

## ACCEPTED MANUSCRIPT

1       **Cryopreserved Homografts in Infected Infrainguinal Fields are Associated with Frequent**  
2                                   **Reinterventions and Poor Amputation Free Survival**

3  
4  
5       S. Keisin Wang MD, Ashley R. Gutwein MD, Natalie A. Drucker MD, Michael P. Murphy MD, Andres  
6           Fajardo MD, Michael C. Dalsing MD, Raghu L. Motaganahalli MD, and Gary W. Lemmon MD

7  
8                                   Indiana University School of Medicine  
9                                   Department of Surgery  
10                                  Division of Vascular Surgery  
11                                  Indianapolis, IN

12  
13  
14       **Corresponding Author:**

15       S. Keisin Wang MD  
16       Integrated Vascular Surgery Research Fellow  
17       Indiana University School of Medicine  
18       Department of Surgery  
19       Division of Vascular Surgery  
20       1801 N. Senate Blvd MPC2-3500  
21       Indianapolis, IN 46202  
22       Phone: 317-962-0282  
23       Fax: 317-962-0289  
24       Email: [wangkei@iupui.edu](mailto:wangkei@iupui.edu)

25  
26       **Keywords:** cryopreserved homografts, critical limb ischemia, infection, conduit

27  
28       **Presentation:**

29       The project outlined in this manuscript was submitted as an abstract for oral presentation to VESS  
30       2018 (Vail, Colorado).

---

This is the author's manuscript of the article published in final edited form as:

Wang, S. K., Gutwein, A. R., Drucker, N. A., Murphy, M. P., Fajardo, A., Dalsing, M. C., ... Lemmon, G. W. (2018). Cryopreserved Homografts in Infected Infrainguinal Fields Are Associated with Frequent Reinterventions and Poor Amputation-Free Survival. *Annals of Vascular Surgery*. <https://doi.org/10.1016/j.avsg.2017.10.032>

31 **Abstract**

32

33 *Objective:*

34           Single-length saphenous vein continues to be the conduit of choice in infected-field critical  
35 limb ischemia (CLI). However, half of these individuals have inadequate vein secondary to previous  
36 use or chronic venous disease. We reviewed our outcomes of infected-field infrainguinal bypasses  
37 performed with cryopreserved homografts (CH), a widely-accepted alternative to autogenous vein  
38 in this setting.

39

40 *Methods:*

41           This is a retrospective, institutional descriptive analysis of infected-field infrainguinal  
42 revascularizations between 2012-2015.

43

44 *Results:*

45           Twenty-four operations were performed in the same number of patients for limb ischemia  
46 with signs of active infection. The mean age of the cohort examined was  $62.5 \pm 14.4$  (standard  
47 deviation) years. Mean SVS risk score was 3.9 with a baseline Rutherford's chronic ischemia score  
48 of 4.3 at presentation. Emergent procedures constituted 29% of cases and the remainder were  
49 urgent. The CH bypass captured was a reoperative procedure in all but one of the patients. Culture  
50 positivity was present in 75% of cases with *S. aureus* (29%) the most commonly isolated organism.

51

52           30-day mortality and major adverse cardiovascular events were both 4%. Amputation free  
53 survival (AFS) was 75% at 30-days. Similarly, 30-day reintervention was 38% with debridement  
54 (43%) and bleeding (29%) the most common indications.

55

56 Average duration of follow-up was  $27.9 \pm 20.4$  months (range 0.5 – 60.4). Mean length of  
57 stay was 14.8 days. Reinfection requiring an additional procedure or antibiotic regimen separate  
58 from the index antibiotic course was 13%. Primary patency and AFS at 1-year was 50% and 58%,  
59 respectively. Primary patency and AFS at 2-years was 38% and 52%, respectively. Limb salvage at  
60 one and 2-years was 70% and 65%. Fifteen (63%) patients required reintervention during the  
61 follow-up period with 40% of those subjects undergoing multiple procedures.

62

63 *Conclusions:*

64 CHs remain a marginal salvage conduit in the setting of infection and no autogenous  
65 choices. Therefore, clinicians should individualize usage of this high-cost product in highly selected  
66 patients only.

**67 Introduction**

68

69           Autogenous single-length vein continues to remain the gold standard conduit for lower  
70 extremity infected-field revascularizations for critical limb ischemia (CLI).<sup>1</sup> However, up to 45% of  
71 patients who require bypass do not have adequate continuous vein secondary to chronic venous  
72 disease or previous vein harvest.<sup>2,3</sup> Unfortunately, the use of alternative synthetic conduits such as  
73 PTFE (polytetrafluoroethylene) puts the patient at increased risk for graft infection and limb loss.  
74 Therefore, cryopreserved autologous homografts (CHs) have become a popular alternative in the  
75 infected surgical field. The purpose of this retrospective analysis was to define outcomes for  
76 contemporary use of CHs in infected fields with respect to patency and limb salvage.

**77 Methods**

78

79 After obtaining Indiana University Institutional Review Board (IRB) approval, a single-  
80 center retrospective review was completed of all infected-field CHs implanted for infrainguinal  
81 arterial disease from 2012 to 2015. All procedures were performed at one institution by a group of  
82 nine academic surgeons. Patients not seen by a vascular surgeon in our system for 12-months were  
83 deemed lost to follow-up; for those, contact by phone was attempted.

84

85 Demographics captured included age, sex, disease severity, and relevant comorbidities.  
86 Presence of infection was defined as observation of cellulitis overlying a bypass graft or perigraft  
87 purulence/fluid/air on imaging. Operative characteristics captured included location of  
88 proximal/distal anastomosis, muscle flap usage, and intraoperative cultures. Post-operative  
89 management strategies queried included use of anticoagulation, antiplatelets, and antibiotics  
90 duration.

91

92 CHs implanted were kept on-site in a liquid nitrogen dewar. These conduits were prepared  
93 per manufacturer's instruction but not routinely seromatched to the host. Based on availability and  
94 surgeon preference, the choice of cryopreserved vein or artery was made on a case-by-case basis.  
95 CHs were used as the first choice in infected fields during this time over rifampin soaked prosthetic  
96 and spliced autogenous vein if continuous vein was not available. All infected fields were copiously  
97 irrigated with antibiotic and saline solution. No antibiotic impregnated beads were implanted in  
98 our series.

99

100 After surgery, all patients maintained IV or PO antibiotic use depending on the clinical  
101 severity of infection. In general, it was our practice to extend antibiotics to 4-weeks before a

102 decision on additional duration was made in the outpatient setting. Post-operative imaging,  
103 vascular labs, and overall management was left to the discretion of the individual attending  
104 surgeon. Most commonly, a post-operative wound check was scheduled two to four weeks after the  
105 index procedure. Graft surveillance was scheduled for every three months for the first year  
106 followed by every six months thereafter. After the second year of follow-up, patients were  
107 extended to annual visits if the bypass remained patent.

108

109 Events captured included one and 2-year primary patency by vascular labs or CTA, one and  
110 2-year amputation free survival (AFS), reinterventions, reinfection, anastomotic bleeding,  
111 mortality, and major adverse cardiovascular events (MACE). AFS was defined as freedom from all-  
112 cause mortality or above-ankle amputation. Reinfection was defined as any decline in clinical  
113 status secondary to a new or persistent infection resulting in escalation of antibiotics, drainage, or  
114 reoperation to revise the index bypass graft.

## 115 **Results**

116

117 From 2012 to 2015, 24 infrainguinal CH (33% vein) bypasses were performed secondary to  
118 an infected conduit (n=23) or native artery (n=1). The mean age of our population was  $62.5 \pm 14.4$   
119 years (**Table 1**). The most common comorbidities included previous bypass (96%), HTN (92%),  
120 active smoking (58%), HLD (54%), CAD (42%), and DM (25%).

121

### 122 *Indications*

123

124 Twelve percent of the patients were referred acutely after initial evaluation by an outside  
125 vascular surgeon for definitive management. Mean Rutherford's chronic limb ischemia score and  
126 mean SVS risk score were 4.3 and 3.9, respectively.<sup>4,5</sup> Emergent procedures (performed within 6  
127 hours of admission) constituted 29% of cases and the remainder were urgent (within 24 hours).  
128 All patients demonstrated signs of local infection on physical exam or imaging; however, only 8%  
129 were septic at the time of presentation. All but one of the procedures were performed as a repeat  
130 bypass. This exception was a male with a primary infection of the superficial femoral artery  
131 secondary to chronic IV drug use and accidental arterial injection.

132

### 133 *Intra-operative*

134

135 Most of the infections were located in the groin (66.7%). All proximal sites of anastomosis  
136 were distal to the external iliac artery. Distal targets were divided into tibioperoneal (17%), below-  
137 knee (4%), and above-knee (79%) categories. Three patients received an extranatomic bypass  
138 consisting of two obturator bypasses and a femoral to femoral bypass via a retrorectus tunnel.  
139 Upon exploration, 33% of the patients had a pseudoaneurysm at the presumed site of infection

140 (Table 2). The majority (92%) of the infected conduits were unincorporated into the soft tissue.  
141 Frank purulence was noted in 46% of limbs. Complete graft explantation was completed in 61%,  
142 and the remainder received a partial explant at the location of active infection. Rotational muscle  
143 flaps were utilized in 46% of cases. Seventy-five percent of cultures returned an identifiable  
144 organism (Table 3). The most common isolated organisms were *S. aureus* (29%), *P. aeruginosa*  
145 (24%), and coagulase negative Staphylococcus (24%). Eighteen percent of positive cultures further  
146 demonstrated extended spectrum antibiotic resistance.

147

#### 148 *Post-operative and 30-day Outcomes*

149

150 After the index operation, 38% of patients received therapeutic anticoagulation while 88%  
151 received antiplatelet therapy (Table 4). All subjects received either IV (92%) or PO antibiotics in  
152 the peri-operative period. The average duration of antibiotic coverage after surgery was 4.6 weeks.  
153 30-day AFS was 75% with a mortality rate of 4%. The lone death occurred in an individual  
154 presenting with peri-graft fluid and sepsis. Antibiotic sensitive *S. aureus* was isolated from  
155 cultures, but the patient continued to decline clinically resulting in multi-system organ failure and  
156 eventual withdrawal of care by the family. One patient experienced stroke/MI, suffering from an  
157 NSTEMI several days post-operatively. Three (13%) patients experienced anastomotic bleeding  
158 with two requiring takebacks for exploration. Reintervention at 30-days was 38% (n=7) most  
159 commonly for further debridement (4/7) or bleeding (2/7). Average length of stay was 14.8 days.

160

#### 161 *Overall Outcomes*

162

163 Mean follow-up for our population was  $27.9 \pm 20.4$  months (Table 5). Primary patency in  
164 our population at one and 2-years was 50% and 38%. AFS at one and 2-years was 58% (6



165 amputations, 5 deaths) and 50% (7 amputations, 7 deaths). Limb salvage at one and 2-years was  
166 70% and 65%, respectively. Reintervention rate during follow-up was 63% with 40% of these  
167 patients requiring repeat bypass. The most common cause of reintervention was for stenosis or  
168 occlusion; one third of reinterventions were for debridement or drainage. There were no  
169 additional episodes of anastomotic bleeding during long-term follow-up compared to the three  
170 observed within 30-days. Thirteen percent of patients had reinfection of the implanted CH. Seven  
171 patients (29%) required major amputation (3 BKA, 4 AKA) during follow-up. More than half, 54%  
172 of all treated patients died during the follow-up period (**Figure 1**). Of these 11 deaths, 4 were from  
173 unknown causes outside of our hospital system. The remainder of deaths occurred secondary to  
174 lung cancer (n=1), hepatic failure (1), pulmonary embolism (1), renal failure (2), and sepsis (2).

175 **Discussion**

176

177 The optimal management strategy for an infected lower extremity bypass graft or artery  
178 would be complete excision and in-line reconstruction with continuous autogenous vein.  
179 Unfortunately, availability of suitable vein having adequate caliber and length is lacking in many  
180 vascular patients with a previous history of bypass.<sup>6</sup> In our cause, this was 96% of the population  
181 studied. As such, we routinely employ the use of CHs if the operation involves a potentially infected  
182 field. However, we do not routinely implant CHs for sterile-field bypasses given their dismal  
183 patency and limb salvage rates.<sup>7-10</sup>

184

185 CHs are harvested from multi-organ donors and preserved in dimethyl sulfoxide (DMSO)  
186 before being frozen in liquid nitrogen (-196°C) for storage.<sup>11</sup> Additionally, each CH vendor employs  
187 a unique preservation process to decrease antigenicity. The complex harvest and preservation  
188 process does incur a significant financial burden to the patient when this conduit is selected for  
189 bypass.<sup>12</sup> When needed, grafts are thawed to room temperature and individually modified by the  
190 surgeon. After pressurization, the endothelial layer is slowly effaced and the tunica media  
191 infiltrated by leukocytes resulting in chronic fibrosis.<sup>13</sup> This smoldering inflammatory response  
192 likely has a large role in late graft failure characterized by intense fibroplasia.<sup>14</sup>

193

194 CHs seem to be more resistant to infection than prosthetic materials through an unclear  
195 mechanism. This effect has been postulated to be related to the presence of the conduit  
196 extracellular matrix allowing for the increased transfer of leukocytes and antibiotics into the  
197 perigraft space.<sup>15</sup> Alternatively, it may be related to vendor-unique methods of tissue processing  
198 including the storage of grafts in the presence of antibiotics.<sup>16</sup>

199

200 We report one and 2-year patencies of 50% and 38% corresponding to limb salvage rates of  
201 70% and 65%. AFS during the same time periods were 58% and 52%. Seven patients required  
202 amputation of the ischemic limb during the follow-up phase; however, amputation risk was  
203 frontloaded as all but one of the subjects lost their limb within 21 days of the index operation. This  
204 data clearly suggests a danger period for limb loss in the perioperative phase of infected-field  
205 repeat bypasses.

206  
207 Surprisingly, robust contemporary series describing CH conduits in infected fields have  
208 been few and small.<sup>10,17</sup> Brown *et al.* reviewed their experience with CHs in infected fields which  
209 included peripheral, but also, carotid and visceral non-aortic reconstructions. Their published  
210 experience described 39 total cases with a mean follow-up of 18 months. Mortality at 30-days was  
211 2.6%. Interestingly, graft reinfection did not occur in their population in contrast to our observed  
212 rate of 13%. Unfortunately, their 1-year patency was not published.<sup>7</sup> The largest series of  
213 cryopreserved vein bypasses was reported in 2003 of 240 consecutive cases in both clean and  
214 infected fields. The majority (89%) were performed for rest pain or tissue loss. The percentage of  
215 infected limbs were not published. The authors did note an overall 30% 1-year primary patency  
216 and 80% limb salvage for all comers.<sup>8</sup>

217  
218 We found a high reintervention rate of 63% in our study. Twenty-eight additional  
219 reinterventions following the index procedure were documented in our 24 patients. The most  
220 common indication was for stenosis, occlusion, or necrotizing soft tissue requiring a combination of  
221 angioplasty, thrombectomy, redo bypass, and debridement. It seems apparent that the index  
222 bypass for this indication cannot be considered the final and definitive operation. Therefore, before  
223 selecting the patient for limb salvage or primary amputation, it is imperative to disclose the risk of  
224 prolonged hospitalization and additional interventions. Based on our experience, we have adopted

225 the use of cryopreserved homografts in infected fields with concurrent placement of a muscle flap  
226 when possible for the sole purpose of limb salvage. After clearance of the infection, consideration  
227 should be made into reoperation with an alternative conduit to improve long-term outcomes.

228

229           Unfortunately, the retrospective nature of this study makes it impossible to be sure all  
230 adverse events were tracked and captured. As many patients were referred to us from outside  
231 vascular surgeons and hospital systems – their follow-up often occurred external to our records.  
232 Regardless, the limited adverse events abstracted in this study illustrates well the poor prognosis of  
233 this population. Another potential confounder present is inherent to a group practice, where  
234 multiple vascular surgeons perform operations per their expertise, often on the same patient. Thus,  
235 standard protocol and procedure were lacking.

236 **Conclusion**

237

238 CHs are an accepted alternative to continuous autogenous vein for redo bypasses in the  
239 setting of an infected field. However, the surgeon should be aware of the increased incidence of  
240 amputation, death, and reintervention prior to offering CH limb salvage for this difficult population.

MANUSCRIPT

241 **Disclosures**

242

243 The authors have no conflicts of interest to disclose.

MANUSCRIPT

244  
245  
246  
247  
248  
249  
250  
251  
252  
253  
254  
255  
256  
257  
258  
259  
260  
261  
262  
263  
264  
265  
266  
267  
268  
269  
270  
271  
272  
273  
274  
275  
276  
277  
278  
279  
280  
281  
282  
283  
284  
285  
286  
287  
288  
289  
290  
291

### References:

1. Mamode N, Scott RN. Graft type for femoro-popliteal bypass surgery. *Cochrane Database of Systematic Reviews*. 1999(2).
2. Kent KC, Whittemore AD, Mannick JA. Short-term and midterm results of an all-autogenous tissue policy for infrainguinal reconstruction. *Journal of vascular surgery*. 1989;9(1):107-114.
3. Taylor LM, Edwards JM, Brant B, Phinney ES, Porter JM. Papers of the North Pacific Surgical Association Autogenous reversed vein bypass for lower extremity ischemia in patients with absent or inadequate greater saphenous vein. *The American Journal of Surgery*. 1987;153(5):505-510.
4. Chaikof EL, Fillinger MF, Matsumura JS, Rutherford RB, White GH, Blankensteijn JD, et al. Identifying and grading factors that modify the outcome of endovascular aortic aneurysm repair. *Journal of vascular surgery*. 2002;35(5):1061-1066.
5. Hardman RL, Jazaeri O, Yi J, Smith M, Gupta R. Overview of Classification Systems in Peripheral Artery Disease. *Seminars in Interventional Radiology*. 2014;31(4):378-388.
6. Kreienberg PB, Darling Iii RC, Chang BB, Champagne BJ, Paty PSK, Roddy SP, et al. Early results of a prospective randomized trial of spliced vein versus polytetrafluoroethylene graft with a distal vein cuff for limb-threatening ischemia. *Journal of vascular surgery*. 2002;35(2):299-306.
7. Brown KE, Heyer K, Rodriguez H, Eskandari MK, Pearce WH, Morasch MD. Arterial reconstruction with cryopreserved human allografts in the setting of infection: A single-center experience with midterm follow-up. *Journal of vascular surgery*. 2009;49(3):660-666.
8. Farber A, Major K, Wagner WH, Cohen JL, Cossman DV, Lauterbach SR, et al. Cryopreserved saphenous vein allografts in infrainguinal revascularization: analysis of 240 grafts. *Journal of vascular surgery*. 2003;38(1):15-21.
9. Gentile AT, Lee RW, Moneta GL, Taylor Jr LM, Edwards JM, Porter JM. Results of bypass to the popliteal and tibial arteries with alternative sources of autogenous vein. *Journal of vascular surgery*. 1996;23(2):272-280.
10. Hartranft CA, Noland S, Kulwicki A, Holden CR, Hartranft T. Cryopreserved saphenous vein graft in infrainguinal bypass. *Journal of vascular surgery*. 2014;60(5):1291-1296.
11. Martin RS, Edwards WH, Mulherin JL, Edwards WH, Jenkins JM, Hoff SJ. Cryopreserved saphenous vein allografts for below-knee lower extremity revascularization. *Annals of surgery*. 1994;219(6):664-672.
12. Huber AJ, Brockbank K, Riemann I, Schleicher M, Schenke-Layland K, Fritze O, et al. Preclinical evaluation of ice-free cryopreserved arteries: structural integrity and hemocompatibility. *Cells Tissues Organs*. 2012(1422-6421 (Electronic)).
13. Calhoun A, Baur G, Porter J, Houghton D, Templeton JW. Fresh and cryopreserved venous allografts in genetically characterized dogs. *J Surg Res*. 1977(0022-4804 (Print)).
14. Carpenter JP, Tomaszewski JE. Human saphenous vein allograft bypass grafts: immune response. *Journal of vascular surgery*. 1998(0741-5214 (Print)).
15. Vogt PR, Brunner-LaRocca H-P, Lachat M, Ruef C, Turina MI. Technical details with the use of cryopreserved arterial allografts for aortic infection: Influence on early and midterm mortality. *Journal of vascular surgery*. 2002;35(1):80-86.
16. Camiade C, Goldschmidt P, Koskas F, Ricco J-B, Jarraya M, Gerota J, et al. Optimization of the Resistance of Arterial Allografts to Infection: Comparative Study with Synthetic Prostheses. *Annals of vascular surgery*. 2001;15(2):186-196.
17. Castier Y, Francis F, Cerceau P, Besnard M, Albertin J, Foulhe L, et al. Cryopreserved arterial allograft reconstruction for peripheral graft infection. *Journal of vascular surgery*. 2005;41(1):30-37.

### Legends

**Table 1:** Comorbidities

**Table 2:** Operative characteristics

**Table 3:** Culture results

**Table 4:** Peri-operative management and results

**Table 5:** Extended outcomes

**Figure 1:** Kaplan-Meier analysis for AFS over time for patients receiving cryopreserved homografts in an infected surgical field. The inputs displayed are all-cause mortality and major (above-ankle) amputation.



<b>Comorbidity</b>	<b>Incidence</b>
HTN	92%
Active Smoker	58%
HLD	54%
CAD	42%
Obesity	33%
DM	25%
CRI (Cr > 1.5)	21%
CVD	17%
Arrhythmia	17%
HD	13%
COPD	8%
Rutherford's Ischemia Score	4.3 ± 0.6
SVS Risk Score	3.9 ± 2.1

<b>Operative Characteristics</b>	<b>Incidence</b>
Emergent	29%
Septic	8%
Loss of Incorporation	92%
PSA	33%
Purulence	46%
Wound Culture Positive	75%
Muscle Flap	46%

Patient #	Graft Material	Culture Results	ESBL/MRSA
1	Synthetic	S. aureus, S. marascens	No
2	Synthetic	Coagulase <sup>-</sup> Staph	No
3	Synthetic	Coagulase <sup>-</sup> Staph, P. aeruginosa, Citrobacter	No
4	Synthetic	P. aeruginosa	No
5	Synthetic	S. aureus	Yes
6	Synthetic	Klebsiella	No
7	Vein	P. aeruginosa	Yes
8	Synthetic	S. aureus, Enterococcus	No
9	Synthetic	Coagulase <sup>-</sup> Staphylococcus	No
10	Synthetic	S. aureus	No
11	Synthetic	Corynebacterium	No
12	Native Artery	P. aeruginosa	No
13	Synthetic	Corynebacterium	No
14	Synthetic	S. aureus	Yes
15	Synthetic	Coagulase <sup>-</sup> Staphylococcus	No
16	Synthetic	Enterococcus	No
17	Synthetic	Corynebacterium	No

<b>Post-Operative Regimen</b>	<b>Incidence</b>
Antiplatelets	88%
Anticoagulation	38%
Antibiotics	100%
Antibiotic Duration	4.6 ± 2.2 Weeks
<b>30-day Outcomes</b>	<b>Incidence</b>
MACE	4%
Major Amputation	21%
Mortality	4%
AFS	75%
Bleeding	13%
Reintervention	38%

<b>Long-Term Outcomes</b>	<b>Incidence</b>
Follow-up	27.9 ± 20.4 Months
LOS	14.8 ± 16.3 Days
Primary Patency	17.4 ± 18.2 Months
1-yr Primary Patency	50%
1-yr AFS	54%
2-yr Primary Patency	38%
2-yr AFS	52%
Reintervention Rate	63%
Reinfection	13%
Bleeding	13%
Major Amputation	29%
Death	54%

