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Contamination of bone allografts

Analysis of incidence and predisposing factors

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ABSTRACT

We analysed the bacterial contamination of 1999 allografts retrieved from 200 cadaver donors under sterile operating conditions. The effect of various factors on the relative risk of contamination was estimated using a multiple logistic regression model.

Organisms of low pathogenicity were cultured from 50% of the grafts and of high pathogenicity from 3%. The risk of contamination with low pathogenic organisms (mainly skin commensals) increased by a factor of 1.6 for each member added to the procurement team. The risk of contamination with high pathogenic organisms (mainly contaminants from the gastrointestinal tract) was 3.4 times higher in donors with a traumatic cause of death and 5.2 times higher in those with a positive blood culture. Preceding organ procurement did not significantly influence the risk of contamination. Rinsing the graft with an antibiotic solution was not an effective decontamination method.

The major source of contamination is exogenous and is strongly influenced by the procurement team. Contamination from endogenous sources can be controlled by donor selection. We discuss methods that can be used to decrease contamination and the rate of discarding of bone allografts.

INTRODUCTION

There is always a risk of bacterial contamination when harvesting bone allografts.¹⁻⁵ Grafts may be contaminated during the process of procurement from the surgeon or the donor. The risk of infecting the recipient can be reduced by adequate microbiological monitoring. If contamination is detected the choices are additional sterilisation or discarding the allograft. Sterilisation can alter the biomechanical and biological properties.⁶⁻⁸ For osteochondral grafts, it is unsuitable as it kills cartilage cells. Such grafts should be preserved fresh frozen without additional sterilisation. To prevent the wastage of time and grafts, contamination should be minimised.

The aim of our study was to determine the incidence and potential causes of contamination and to assess the effectiveness of antibiotic rinsing of allografts.

MATERIALS AND METHODS

Potential cadaver donors were carefully selected and screened according to the guidelines of the American Association of Tissue Banks (AATB)⁹ and the European Association of Musculo Skeletal Transplantation (EAMST).¹⁰ They were not considered for tissue donation if there was a clinical history of sepsis at the time of death. The methods of procurement were those described by Tomford.^{1,11} Using aseptic techniques bone and soft-tissues were procured in an operating theatre within 24 hours of death. Blood from the heart or subclavian vein was taken for culture before the procurement.

After removal the grafts were rinsed with an antibiotic solution (bacitracin 50 000 U/L, Apotekernes Lab, Oslo, and polymyxin-B 500 000 U/L, Pfizer, Connecticut, USA). Allografts from the last 50 donors were rinsed with sterile saline only. Subsequently, swabs were taken of the entire graft surface, and if possible the medullary canal, using a 15 cm polyester-tipped applicator (Becton Dickinson and Company, Cockeysville, Maryland) and placed in a transport medium (Port-A-Cul, Becton Dickinson). They were then inoculated on to blood agar and chocolate agar plates and cultured under aerobic and anaerobic conditions. The entire swabstick was placed in an brain heart infusion (BHI) broth (Oxoid; Unipathe, Basingstoke, UK). After two to five days, the broth was subcultured on

agar plates which were cultured aerobically and anaerobically for 72 hours. Blood samples were cultured aerobically and anaerobically for seven days. The level of contamination (microbial load) was considered low if micro-organisms grew only in the broth and high when they grew directly on the plates.

Between December 1988 and June 1994 we procured bone grafts from 200 cadaver donors. The cause of death was cardiac failure in 87, cerebral haemorrhage in 50, trauma in 42 and other causes in 21. The mean duration between death and procurement of tissue was 8.2 hours (1.0 to 23.8). The bone and soft-tissue procurements were preceded by an organ procurement in 58 and by a heart-valve procurement in 42 cases. The number of staff members performing the procurement varied from three to six (mean, 4.6). The mean duration of the procurement process was 1.7 hours (0.75 to 3.0).

A total of 1999 separate bone and related soft-tissue grafts was obtained with each cadaver contributing six to eighteen. These were; humerus 192, radius 24, ilium/hemipelvis 141, femoral fragments 655, tibial fragments 399, Achilles tendon 238, fascia lata 48, others 19.

Statistical analysis

We performed a multivariate analysis, using a logistic regression model. As one donor produced several bone grafts, the correlations between grafts within one donor was accounted for using a random-effects model. We used the package EGRET.¹² We considered $p < 0.05$ as significant. The effect of various factors on the relative risk of contamination was estimated by odds ratios; p values were calculated on the basis of the likelihood ratio tests.

RESULTS

Graft contamination

From the 1999 allografts procured, micro-organisms of low pathogenicity were cultured from 999 (50%) and micro-organisms of high pathogenicity from 57 (3%) (Table I). From 943 grafts (47%) no organisms were cultured. Contamination with organisms of low pathogenicity was of low microbial load in 69% (689/999) of the cases.

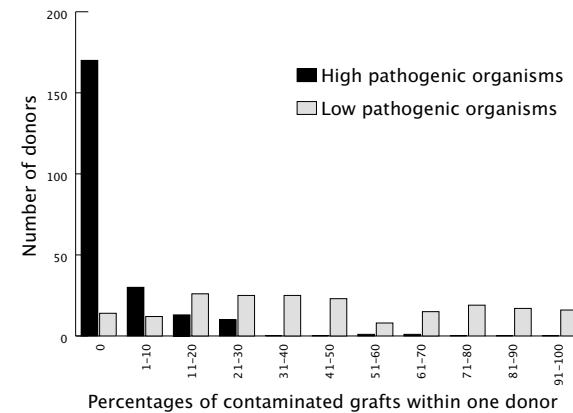


Fig. 1. Frequency histogram for percentages of contaminated grafts within one donor. Grafts can be contaminated with organisms of high and low pathogenicity.

Table I. Details of micro-organisms cultured from bone grafts

Micro-organisms	Number of times cultured*	
<i>Low pathogenicity</i>		
Coagulase-negative staphylococcus	959	
<i>Corynebacterium</i> species	97	
<i>Propionibacterium acnes</i>	29	
<i>Bacillus</i> species	15	
Diphtheroids	4	
Micrococcus	2	
<i>Sarcina</i>	3	
<i>High pathogenicity</i>		
<i>Streptococcus</i> species	36	
<i>Staphylococcus aureus</i>	9	
<i>Clostridium</i> species	5	
<i>Bacteroides</i>	4	
<i>Escherichia coli</i>	4	
Yeast	3	
<i>Pseudomonas aeruginosa</i>	1	
Candida	1	
<i>Grafts contaminated with</i>		
Micro-organisms of low pathogenicity	999	(50%)
Micro-organisms of high pathogenicity	57	(3%)
No growth	943	(47%)
Total cultures	1999	

* multiple organisms were cultured from 114 graft

For any one donor, the proportion of grafts which were contaminated with micro-organisms of low pathogenicity varied from 0% to 100% and with micro-organisms of high pathogenicity primarily from 0% to 30% (Fig. 1).

A single organism was cultured from 942 grafts and more than one from 114. The most frequently cultured organisms was coagulase-negative staphylococcus (82%). Of organisms of high pathogenicity the most frequently seen were *Streptococcus* species (Table I).

Blood contamination

Blood for culture was obtained from 186 of the 200 donors. Seventeen cultures contained organisms of low pathogenicity originating from the skin but identical organisms were not cultured from their allografts. We considered these cultures to be falsely positive because of external contamination due to needle perforation. Twenty-six blood cultures were truly positive. The pre-dominant organisms cultured were *Streptococcus* species.

Predisposing factors for graft contamination

Multiple logistic regression analysis showed that several risk factors were simultaneously significant. Furthermore, the correlation of grafts within one donor (random effect) was highly significant ($p < 0.001$). To account for this correlation when assessing the significance of the risk factors, a random effect term was added to the logistic regression model. If this correlation is not taken into account unwarranted levels of significance may arise. For the analysis a distinction was made between micro-organisms of high pathogenicity and those of low pathogenicity.

Micro-organisms of low pathogenicity

The number of staff members performing the procurement, the type of graft and the time since death were risk factors associated with contamination (Table II). Each staff member added to the procurement team increased the risk of contamination by a factor of 1.6 ($p = 0.001$). For each hour increase between the time of death and procurement, the risk of contamination decreased by a factor of 0.93 ($p = 0.002$). The type of graft also influenced the risk of contamination ($p = 0.004$). The contamination rate was low for the radii and high for the Achilles tendons. Trauma as

the cause of death, preceding organ/heart-valve procurement and bone procurement duration were not associated with increased risk of contamination with organisms of low pathogenicity (Table II).

Table II. The effect of various factors on the relative risk of graft contamination with low pathogenic organisms (see Table I) estimated by odds ratios

Factors	Contamination of Grafts*		Logistic Regression†		
	No	Yes	Odds Ratio	95% CI	p value
Traumatic cause of death					0.39
No (n=158)	767	802	(1)		
Yes (n=42)	176	197	1.3	0.7 to 2.2	
Preceding procurement					0.27
No (n=100)	490	461	(1)		
Organ (n=58)	263	326	0.9	0.5 to 1.5	
Heart valve (n=42)	190	212	1.5	0.9 to 3.1	
Time since death					0.002
Per 1 hr increase			0.93	0.9 to 1.0	
Team members					0.001
3 (n=5)	34	13	(1)		
4 (n=90)	456	391	1.6	1.2 to 2.5	
5 (n=90)	387	483	2.4		
6 (n=15)	66	112	3.7		
Procurement duration					0.42
Per 1 hr increase			1.3	0.7 to 2.5	
Type of grafts					0.004
Humerus	84	97	(1)		
Radius	18	6	0.4	0.1 to 1.2	0.29
Femur	320	313	0.9	0.6 to 1.4	0.81
Tibia	190	204	1.0	0.6 to 1.6	0.68
Fibula	150	126	0.8	0.5 to 1.2	0.36
Pelvis	65	73	1.1	0.6 to 2.0	0.30
Achilles tendon	87	145	1.6	1.3 to 3.4	0.004
Blood contamination‡					0.85
No (n=160)	755	814	(1)		
Yes (n=26)	121	112	0.9	0.6 to 1.6	
Antibiotic rinse					0.02
No (n=50)	191	260	(1)		
Yes (n=150)	752	739	0.5	0.2 to 0.8	

* 57 of 1999 grafts were contaminated with high pathogenic organisms; these were not included in this analysis

† with random-effect to account for graft-within-donor structure; all estimated effects (expressed by the odds ratios) are adjusted for each other in a multivariate analysis

‡ blood sampling not performed for 14 donors, is 140 grafts

Micro-organisms of high pathogenicity

Two factors were associated with an increased risk of contamination with organisms of high pathogenicity (Table III). A traumatic cause of death increased the risk by a factor of 3.4 ($p = 0.007$) and a positive blood culture by a factor of 5.2 ($p = 0.005$). In nine out of 26 donors with a positive blood culture, identical organisms were found in one or more cultures taken from the grafts.

Predisposing factors for blood contamination

A traumatic cause of death increased the risk of blood contamination by a factor of 14.6 ($p < 0.001$). Time from death and preceding organ/heart-valve procurement did not alter the risk of blood contamination (Table IV).

Effectiveness of antibiotic rinse

All the allografts procured from the first 150 donors (1544 grafts) were rinsed with an antibiotic solution, but in the last 50 donors (455 grafts) only saline was used. Multiple logistic regression analysis of the data showed that rinsing with an antibiotic solution decreased the risk of contamination with organisms of low pathogenicity by a factor of 0.5 ($p = 0.02$) (Table II). Further analysis showed that this decrease was caused by a reduction in the number of grafts with a low microbial load (31% *v* 45%), but the risk of contamination with organisms of high pathogenicity did not alter (Table III).

Table III. The effect of various factors on the relative risk of graft contamination with high pathogenic organisms (see Table I) estimated by odds ratios

Factors	Contamination of Grafts*		Logistic Regression†		
	No	Yes	Odds Ratio	95% CI	p value
Traumatic cause of death					0.007
No (n=158)	1569	31	(1)		
Yes (n=42)	373	26	3.4	2.0 to 5.5	
Preceding procurement					0.76
No (n=100)	951	26	(1)		
Organ (n=58)	589	22	1.0	0.4 to 3.0	
Heart valve (n=42)	402	9	0.9	0.3 to 3.5	
Time since death					0.86
Per 1 hr increase			1.0	0.9 to 1.2	
Team members					0.93
3 (n=5)	47	0	(1)		
4 (n=90)	847	23	1.0	0.5 to 1.9	
5 (n=90)	870	28	1.0		
6 (n=15)	178	6	1.0		
Procurement duration					0.65
Per 1 hr increase			1.3	0.4 to 4.3	
Type of grafts					
Humerus	181	11	-		
Radius	24	0	-		
Femur	633	22	-		
Tibia	394	5	-		
Fibula	276	7	-		
Pelvis	138	3	-		
Achilles tendon	323	6	-		
Blood contamination‡					0.005
No (n=160)	1593	9	(1)		
Yes (n=26)	233	24	5.2	1.6 to 17	
Antibiotic rinse					0.30
No (n=50)	411	4	(1)		
Yes (n=150)	1491	53	1.9	0.5 to 11	

* with random-effect to account for graft-within-donor structure; all estimated effects (expressed by the odds ratios) are adjusted for each other in a multivariate analysis

† logistic regression on "type of graft" was invalid because of small size of the (sub-)populations

‡ blood sampling not performed for 14 donors; is 140 grafts

Table IV. The effect of various factors on the relative risk of blood contamination estimated by odds ratios.

Factors	Contamination of Grafts*		Logistic Regression†		
	No	Yes	Odds Ratio	95% CI	p value
Traumatic cause of death					<0.001
No (n=147)	137	10	(1)		
Yes (n=39)	23	16	14.6	5.1 to 43	
Preceding procedure					0.42
No (n=98)	83	15	(1)		
Organ (n=50)	46	4	0.4	0.1 to 1.5	
Heart valve (n=38)	31	7	0.6	0.2 to 2.2	
Time since death‡					0.08
Per 1 hr increase			1.2	1.0 to 1.5	

* a total of 186 blood cultures were analysed, in 14 out of 200 donors blood samples were not obtained due to technical reasons
† with random-effect to account for graft-within-donor structure; all estimated effects (expressed by the odds ratios) are adjusted for each other in a multivariate analysis
‡ if there was a preceding procedure; the samples were taken at the time of the preceding procurement to exclude bias, we only analysed procurement without preceding organ/heart-valve procurement

DISCUSSION

Despite careful donor selection and aseptic tissue procurement, we frequently cultured micro-organisms from bone and soft-tissue allografts. Those of low pathogenicity, which we considered to be skin commensals, were cultured from 50% of the grafts. Two-thirds of these grafts (69%) had what we termed low microbial load. Micro-organisms of high pathogenicity were cultured from only 3% of the allografts. Others have reported rates of bacterial contamination between 5% and 44%.¹⁵ Comparison of the contamination rates in these reports is not possible because of differences in culture techniques or lack of information about the technique used. Blood cultures from 14% of the allograft donors were positive, *Streptococcus* species being most frequently grown. This agrees with another report¹³ in which a comparable percentage of blood was contaminated with organisms other than skin commensals.

We are of the opinion that it is important to distinguish between allograft contamination with micro-organisms of low and high pathogenicity, as they are likely to originate from different sources. Most organisms of low pathogenicity are skin commensals (e.g. coagulase-negative

staphylococci, *Corynebacterium* species and *Propionibacterium acnes*) and probably represent external contamination at the time of procurement. Organisms of high pathogenicity probably originate from endogenous sources in the donor. They are usually contaminants from the upper respiratory or gastrointestinal tract (e.g. streptococci, *Staphylococcus aureus* and *Escherichia coli*) and are more likely to cause infection in the recipient of the allograft.

For each staff member added to the procurement team the risk of contamination with organisms of low pathogenicity increased by a factor of 1.6. Micro-organisms shed from the skin of the staff are considered the major source of airborne bacteria, and are deposited by gravity on to the surface of the grafts.¹⁴ In conventionally ventilated operating rooms around 95% of the wound bacteria come from the air.^{15,16} We recommend that allograft procurement should be performed by a small and experienced surgical team.

The procurement of Achilles tendons was associated with an increased risk of contamination with skin commensals. Adhesive drapes used at this site were frequently found to detach. As the tendons were the last to be removed there was a greater risk of their contact with heavily contaminated surgical drapes or instruments.

We found it interesting that preceding procurement of organs or heart valves did not influence the risk of contamination with low pathogenic organisms. We suggest two explanations. First, micro-organisms shed from the different staff performing the preceding procurement are a relatively small risk for contamination, already having settled to the floor and other surfaces. Particles settle (move downwards) at a speed of 20 to 30 cm/min.¹⁴ Once settled on the floor or another surface, organisms are only resuspended into the air with great difficulty.¹⁷ Secondly, during donor and back table preparation, the number of organisms is increased substantially. Our measurement of colony-forming units (CFU) in the operating theatre showed a mean count of 5 CFU/m³ (0 to 16) in an empty operating theatre just before the procurement with a sharp rise to 65 CFU/m³ (8 to 235) during donor and backtable preparation. This was independent of any preceding procurement. By the time the surgical excision of the allografts occurred the number had dropped to 25 CFU/m³ (0 to 125).

Contamination with organisms of high pathogenicity can be reduced by careful donor selection. If there had been trauma to the maxillofacial area or abdomen bacteraemia is likely to occur with haematogenous spread

to the bones. We found that after a traumatic cause of death the risk of graft contamination with organisms of high pathogenicity increased by a factor of 3.4 ($p = 0.007$), and the risk of blood contamination by a factor of 14.6 ($p < 0.001$). Haematogenous spread was emphasised by the strong relationship between a positive blood culture and the contamination of bone with high pathogenic organisms (odds ratio = 5.2; $p = 0.005$). Thus, caution must be exercised before selecting as allograft donors those who have had severe trauma.

It has been reported that transmigration of micro-organisms from their areas of colonisation to new sites is not frequent after death if tissues are excised within a 24-hour period.^{13,17,18} We agree with this finding, the risk of contamination with high pathogenic organisms from an endogenous source in the donor having an odds ratio of 1.0 ($p = 0.86$). We did not notice an effect of overgrowth of micro-organisms on the skin of the donor after a prolonged time. By contrast, an increased time between death and procurement reduced the risk of contamination with low pathogenic skin contaminants by a factor of 0.93 per hour. This occurred in the allografts with and without preceding organ procurement. We cannot explain this phenomenon but do not consider it to be of practical importance.

Rinsing with solutions of antibiotics is commonplace procedure. In an experimental model²⁰ the microbial load was reduced by a factor of two. Our data showed that after rinsing with an antibiotic solution the risk of contamination with organisms of low pathogenicity was reduced. This effect was caused by a decrease in a number of allografts contaminated with a low microbial load, but there was no decrease in allografts with a high microbial load. The risk of contamination with organisms of high pathogenicity was not affected. We support the view² that pathogenic organisms persist despite washing in antibiotic solutions. We found no clear advantage in rinsing allografts with the antibiotic solution, possibly because the exposure time was too short for the antibiotics to be effective.

The demand for bone and cartilage allografts is increasing and tissue banks must meet this demand. Tissue procurement must be performed under conditions that minimise the risk of bacterial contamination. Our study has shown that contamination with low pathogenic skin commensals is frequent and the rate can be influenced by the procurement team. Organisms shed from the staff and transferred directly to the graft before reaching the floor and other surfaces are a much greater cause of contami-

nation than resuspended organisms. The exclusion of donors with severe trauma in highly colonised areas should diminish the risk of implanting a fresh frozen graft contaminated with organisms of high pathogenicity. A swab used to culture tissue is a method with low sensitivity.²¹ We believe that constant stringent monitoring of the level of contamination is necessary to identify and eliminate predisposing factors for contamination.

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