

ORIGINAL RESEARCH

Surgical Site Infection in Major Lower Limb Amputation (SIMBA): an international multicentre audit: baseline unit survey

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Plain English Summary

Why we undertook the work: In 2022 over 3,000 people in the UK required an amputation of their leg. After amputation surgery there is a risk of wound infection. This can range from mild infections that can be treated with antibiotics to more serious problems including longer hospital stays, additional surgeries or even death. There are recommendations regarding prevention and treatment of wound infections; however, it remains unclear how effective these are and how closely hospitals follow this guidance.

What we did: We have designed an international audit: Surgical Site Infection in Major Lower Limb Amputation (SIMBA). Its aim is to evaluate infection rates, related complications and current care practices after amputation. As part of SIMBA, we conducted a survey to see how closely hospitals follow existing recommendations. It also looked to see which methods are most commonly used to prevent and treat wound infections.

What we found: We found that some practices were commonly used, such as using scans to plan surgery. However, there was significant variation in other areas. For example, not all hospitals routinely conduct pre-surgery assessments from specialists such as dieticians, psychologists and physiotherapists. Additionally, follow-up care, including rehabilitation and mental health support, varied widely between hospitals.

What this means: The results demonstrate that approaches to preventing wound infections after amputations vary and more specific evidence-based guidelines are needed. Better standardisation of practices could help to reduce infections and improve recovery for patients. More research focusing on amputation-specific guidelines could lead to better patient outcomes in the future.

Abstract

Introduction: Surgical site infection (SSI) is common after major lower limb amputation (MLLA) and is associated with significant morbidity and mortality. National and international guidelines and a best practice pathway aim to optimise care and prevent complications, but adherence is unknown.

Methods: Surgical Site Infection in Major Lower Limb Amputation (SIMBA) is an international, prospective, collaborative audit which compared current practice against national and international recommendations and evaluated equipoise regarding best practice. Each participating centre completed a baseline unit survey containing Likert scale questions regarding local MLLA pathways. Responses were compared with the Vascular Society of Great Britain and Ireland's best practice pathway, National Institute for Health and Care Excellence (NICE), Society of Vascular Surgery's Practice management guide and the European Journal of Vascular Surgery's Global Vascular Guidelines on the Management of Chronic Limb-Threatening Ischaemia guidelines.

Results: Forty centres (30 UK, 7 Europe, 2 Australasia and 1 Asia) completed the survey, yielding a response rate of 87% (40/46). MLLA was performed by vascular surgeons in all centres, with additional specialities also undertaking MLLA surgery including orthopaedic (n=10), plastic (n=4) and general surgery (n=3). Induction antibiotic prophylaxis was given in 32 (82.1%) of the centres. Prophylactic postoperative antibiotics were 'commonly' or 'always' given in 24 (61.5%) of the centres, typically comprising a 5-day intravenous course. Incise drapes were infrequently used (used 'never' for iodophor (39.5%, n=15) and non-iodophor

(44.7%, n=17) containing drapes). Routine follow-up was conducted in 27 centres (69.2%) and preoperative vascular imaging was 'commonly' or 'always' performed in 37 centres (92.5%). Preoperative assessment by physiotherapists and/or occupational therapists and diabetic specialists occurred 'commonly' or 'always' in 32 (82.1%) and 27 (71.1%) centres, respectively. Dietetic and psychological assessment only occurred 'commonly' or 'always' in 8 (21.6%) and 9 (25%) centres, respectively.

Conclusions: This audit highlights the variability in practice, underscoring the need for consensus on best practice. Future studies should focus on generating high quality evidence to refine recommendations and reinforce adherence to guidelines to reduce SSI and improve outcomes after MLLA.

Key words: major lower limb amputation, surgical site infection, chronic limb threatening ischaemia, wound breakdown

Introduction

Surgical site infections (SSIs) are a common complication following any surgical procedure, accounting for 20% of all hospital-associated infections.¹ The incidence of SSI following major lower limb amputation (MLLA) is particularly high, with a recent systematic review reporting an overall incidence of 7.2% and single-centre studies reporting rates up to 27%.² SSIs are a leading cause of in-hospital morbidity and mortality,³ and consequences of their development, including the substantial contribution to prolonged hospitalisation, result in SSIs being the costliest hospital-associated infection.¹ Furthermore, SSIs post MLLA increase the risk of stump dehiscence and need for revision amputations to the same or a higher level.² This may prevent a patient from independent ambulation,⁴ significantly affecting quality of life and negatively impacting mental health.⁵

The importance of this issue has been recognised by both clinicians and patients. The Priority Setting Partnership led by the Vascular Society of Great Britain and Ireland (VSGBI) in conjunction with the James Lind Alliance has highlighted improving wound healing and improving clinical outcomes after MLLA as two of the top 10 research priorities in amputation surgery.⁶ Furthermore, wound healing and stump infections have been highlighted in the core outcome set for MLLA.⁷ The VSGBI has established a best practice clinical care pathway designed to optimise quality of care and reduce complications after MLLA.⁸ The National Institute for Health and Care Excellence (NICE) have also published guidance relating to the prevention and treatment of SSI.³ However, the degree of adherence to these recommendations remains unclear. Various interventions, such as specialist dressings,⁹ negative pressure wound management systems¹⁰ and antimicrobial-coated sutures¹¹ have become increasingly available. Benefits have been demonstrated from prolonged prophylactic antibiotic courses to reduce the incidence of SSI in MLLA;^{12,13} however, aside from this, evidence of effective interventions to reduce the incidence of SSI in MLLA is sparse, and all adjuncts incur additional costs, contributing to variability in practice.

Surgical Site Infection in Major Lower Limb Amputation (SIMBA)

is an international collaborative audit comparing current practice against recommendations. It also aims to determine the incidence of SSIs and associated clinical sequelae, although these data are not part of this publication.¹⁴ The study consists of two parts: prospective data collection surrounding risk factors, interventions and outcome for patients undergoing MLLA, and an initial baseline unit survey completed once by each enrolled centre. This paper presents the results of the baseline unit survey. The primary aim of this survey was to assess adherence to published guidelines on reducing SSI. Secondary aims included assessing adherence to recommendations regarding optimising overall care and improving outcomes post MLLA, and evaluating equipoise regarding best practice for management of patients undergoing MLLA.

Methods

Study design

SIMBA is an international, prospective, collaborative audit designed to assess current clinical practice against established recommendations and to determine the incidence of SSI and associated clinical outcomes. A detailed protocol has been published in full.¹⁴ This audit is partially funded by the ROSSINI platform as part of the accelerator award scheme (Award ID: NIHR156728)¹⁵ and has been conducted in conjunction with the Birmingham Centre for Observational and Prospective Studies (BiCOPS) at the University of Birmingham. SIMBA is supported by the Vascular and Endovascular Research Network (VERN; <https://vascular-research.net/>), a multidisciplinary trainee-led vascular research collaborative.¹⁶

Centre enrolment began in October 2023, with data collection concluding on 1 May 2024. Any centre within the UK or internationally that provides emergency and/or elective MLLA under any speciality was eligible to participate. Centres were recruited through outreach by VERN using social media, email communications and professional networks. As part of the audit process, the lead consultant from each enrolled centre was required to complete a baseline unit survey detailing local pathways for managing patients undergoing MLLA.

Questionnaire development

Strategies for SSI prevention and management detailed within the NICE guidelines and recommendations for the care of those undergoing MLLA, including those within the VSGBI Best Practice clinical care pathway,⁸ the European Journal of Vascular Surgery Global Vascular Guidelines on the Management of Chronic Limb-Threatening Ischaemia¹⁷ and the Society of Vascular Surgery's Practice Management Guide,¹⁸ were identified and reviewed. Recommendations for preoperative, perioperative and

postoperative care were included. Based on these recommendations, a cross-sectional survey was created, including questions designed to assess centre compliance (Table 1). Additionally, questions were incorporated to explore variations in practice and areas of uncertainty to evaluate equipoise on best practice. Responses were collected using a Likert scale where possible, with a combination of multi-select and free-text options where required.

The survey underwent two rounds of internal validation by the

Table 1 Guideline adherence.

Guideline/recommendation (Grade of evidence where stated)	Recommending organisation				Relevant survey question	Percentage adherence
	NICE	EJVES	VSGBI	SVS		
Preoperative						
Patients should be assessed by the MDT prior to MLLA		X	X		7	42.5%*
Offer patients and carers clear, consistent information and advice through all stages of their care	X				35a	50%*
Be admitted under a named consultant in vascular surgery			X		14	92.3%*
Undergo diagnostic arterial imaging to determine revascularisation options			X	X	9	92.5%*
Have revascularisation options discussed at a vascular imaging MDT			X		11	75%*
Undergo assessment using TcPO2 to determine perfusion at a proposed amputation level				X	20	48.3%*
Involvement of clinical psychology				X	16a	25.0%*
Undergo assessment with OT/PT preoperatively (1C)		X	X		16b	82.1%*
Have nutritional assessment and receive dietician advice			X		16c	21.6%*
Have a member of diabetes team involved			X		16d	71.1%*
Have a venous thrombo-embolism risk assessment and prophylaxis as appropriate			X	X	18	97.4%*
Perioperative						
Antibiotic prophylaxis should not routinely be used for clean non-prosthetic uncomplicated surgery	X				21	17.9%
Do not use non-iodophor-impregnated incise drapes routinely for surgery	X				25	78.8%**
Do not use hair removal routinely	X				27	7.7%**
If hair has to be removed, use electric clippers	X				28	71.8%
Postoperative						
Antibiotic prophylaxis should not routinely be used for clean non-prosthetic uncomplicated surgery	X				36	15.4%**
Patients should be informed of the post-amputation care pathway		X			35b	50%*
Offer patients and carers information and advice about how to recognise a surgical site infection and who to contact if they are concerned	X				41	74.4%*
Day 1 postoperative review by acute pain team			X		33	61.5%*
Follow-up in clinic within a month after surgery				X	43	33.3%*
Outpatient review and rehabilitation follow-up with rehabilitation team (1C)		X	X	X	44	63.0%*
Referred to amputation support group				X	35d	36.8%*

* Percentage selecting 'commonly' or 'always'.

** Percentage selecting 'never' or 'rarely'.

EJVES, European Journal of Vascular and Endovascular Surgery; MDT, multidisciplinary team; NICE, National Institute for Health and Care Excellence; OT, occupational therapy; PT, physiotherapy; SVS, Society of Vascular Surgery; VSGBI, Vascular Societies of Great Britain and Ireland.

study management group. The survey was refined by consensus on major alterations (removing or adding questions) and minor alterations (wording or response modification). The first validation resulted in four major alterations (three questions added and one deleted) and five minor alterations, and the second yielded seven minor alterations.

The final survey included 29 questions; three were demographic questions, eight related to preoperative assessment and care, seven related to perioperative interventions and 11 assessed postoperative care and follow-up. The final version of the survey is provided in Appendix 1 online at www.jvsgbi.com.

Survey administration

The survey was built and published using the QualtricsXM Platform™ and was distributed via an email to the consultant leads for all centres enrolled in the SIMBA audit. Participants completed the survey by following the study URL link. Non-responding centres were followed up with reminder emails. Where duplicate responses were received from a single centre, the most complete response was retained; if all responses were equally complete, the first submitted response was kept.

Statistical analysis and reporting

Responses were exported to Microsoft Excel for cleaning and analysis. Non-response questionnaires were removed. Partially completed questionnaires were included. Dichotomous and Likert responses were reported as percentage of responses, and multiple selection questions were analysed to find a median or modal average. The Checklist for Reporting Of Survey Studies (CROSS)¹⁹ was followed during all steps of the study and a completed checklist is provided in Appendix 2 online at www.jvsgbi.com.

As many of the guidelines and recommendations selected as audit standard were UK based, sensitivity analysis was also performed using only the UK centres.

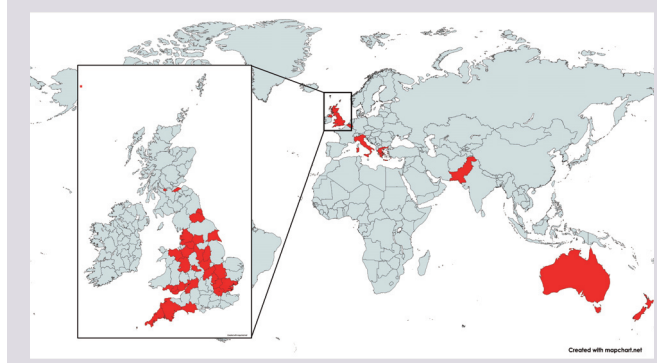
Results

Of the 46 SIMBA centres, 40 completed the survey giving an overall response rate of 87%. This included 30/34 in the UK (88%), 7/9 European centres (78%), 2/2 Australasian centres and 1/1 Asian centre (Figure 1).

In all centres MLLA was performed by vascular surgeons. In 30.0% (12/40) other specialties were also performing MLLA, including orthopaedic surgery (n=10), general surgery (n=4) and plastic surgery (n=3). Patients were 'commonly' or 'always' admitted under a named consultant in vascular or orthopaedic surgery in 92.3% (36/39) of centres.

Adherence to published guidelines and recommendations varied across participating centres (Table 1). The grade of evidence supporting these recommendations was rarely specified. A sensitivity analysis including only UK-based centres demonstrated broadly similar results (see Supplementary Table 1 - Appendix 3 online at www.jvsgbi.com).

Figure 1 Location of survey respondents (Ref. mapchart.net).



Adherence to SSI prevention and management guidelines

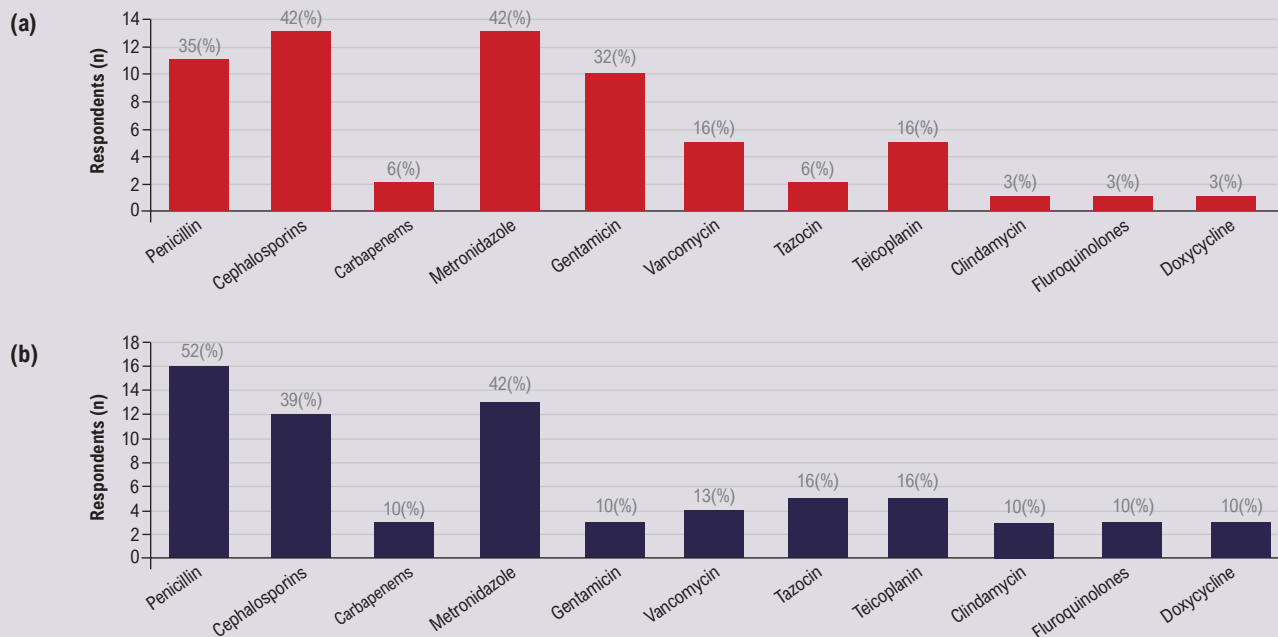
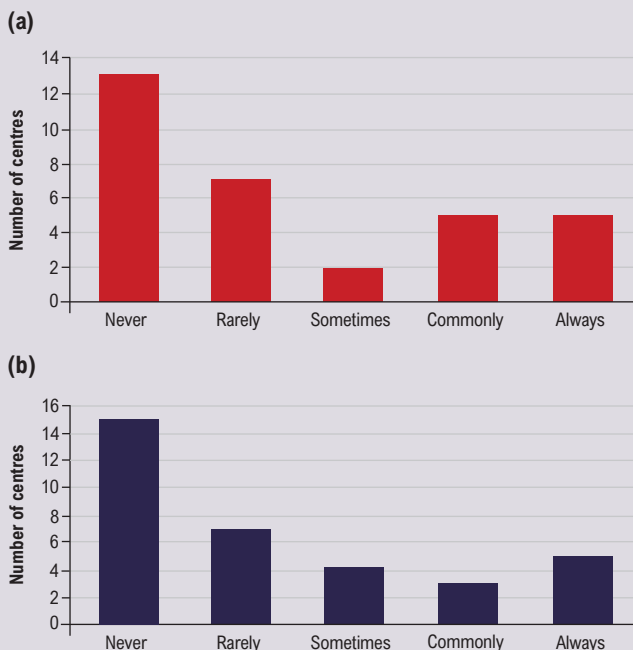
Antibiotic administration at the time of anaesthetic induction is routinely performed in 82.1% (32/39) of centres, with a median of two different antibiotics given intravenously. The most commonly used antibiotics are cephalosporins (13 centres) and metronidazole (13 centres) (see Figure 2a). Prophylactic postoperative antibiotics are 'commonly' or 'always' prescribed in 61.5% (24/39) of centres. The majority (22/24) administered exclusively intravenous antibiotics, whilst nine centres use either intravenous or oral routes and three centres routinely use oral antibiotics. Antibiotic choice varied between centres (see Figure 2b), with the most prevalent being penicillin (16/24), metronidazole (13/24) and cephalosporins (12/24) and the most common course duration ranging from 72 hours to 5 days. Adherence to the current NICE guidelines, which advise against the use of prophylactic antibiotics in clean, non-prosthetic, uncomplicated surgery, was low with 17.9% (7/39) of centres reporting they 'never' or 'rarely' administer antibiotics at induction and 15.4% (6/39) postoperatively.

Routine preoperative hair removal is 'commonly' or 'always' performed in (24/39) of centres, with 71.8% of these using electric clippers aligning with guideline recommendations. However, only 7.7% (3/39) of centres reported 'never' or 'rarely' performing hair removal, reflecting low adherence to guidance advising against routine hair removal. Just over half of the centres (60%, 20/32) 'commonly' or 'always' perform MLLA without incise drapes. When used, 78.8% of centres reported 'never' or 'rarely' using non-iodophor impregnated drapes, reflecting good adherence with guidelines (see Figure 3a and b).

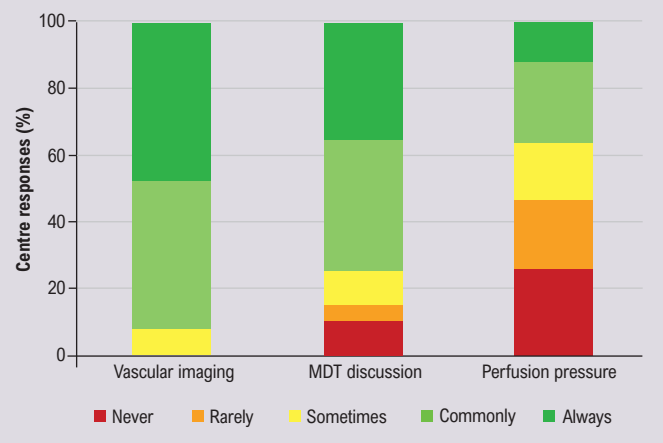
Adherence to guidelines on providing information about SSI recognition and management was high, with 74.4% (29/39) selecting 'commonly' or 'always' done. In comparison, leaflets detailing the procedure itself and expected postoperative recovery are 'commonly' or 'always' provided in 50% of centres (19/38).

Adherence to best practice clinical care recommendations

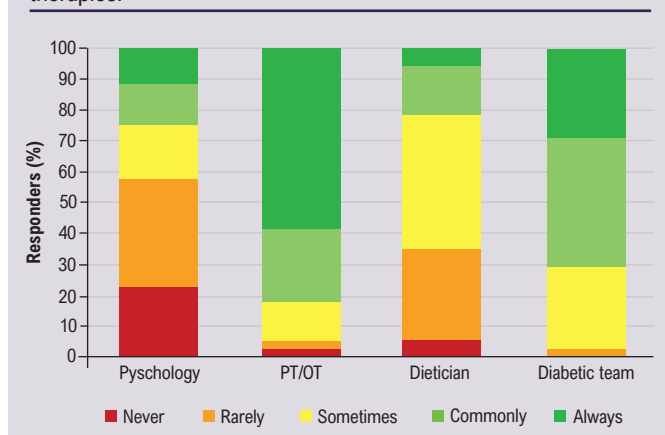
Diagnostic imaging to assess revascularisation options are 'commonly' or 'always' performed in 92.5% (37/40) of centres and 75% (30/40) routinely discuss these cases in vascular

Figure 2 Antibiotics used prophylactically for patients undergoing major lower limb amputation (a) at induction of anaesthesia and (b) postoperatively.**Figure 3** (a) Use of incise drapes and (b) use of iodophor-impregnated incise drapes.

multidisciplinary meetings, demonstrating strong adherence to recommendations. In contrast, use of preoperative perfusion pressure measurements such as TcPO₂ are only 'commonly' or 'always' implemented in 48.3% (14/29) of centres (Figure 4).

Figure 4 Preoperative assessment of revascularisation options.

Routine preoperative assessment with occupational therapy and/or physiotherapy is 'commonly' or 'always' conducted in 82.1% (32/39) of centres and diabetic assessments are 'commonly' or 'always' undertaken in 71.1% (27/38), indicating good adherence to recommendations for multidisciplinary evaluation. However, adherence to guidelines recommending involvement of a dietician and clinical psychology were poor, with only 21.6% (8/37) and 25% (9/36) of centres selecting 'commonly' or 'always', respectively (Figure 5). Almost all centres (97.4%; 38/39) 'commonly' or 'always' implement thromboembolism risk assessment and prescribe prophylactic anticoagulation according to their local protocols.

Figure 5 Frequency of preoperative assessment by supportive therapies.

In 61.5% (24/29) of centres, patients routinely receive input from the acute pain team on the first postoperative day, indicating moderate adherence to pain management recommendations.

Routine follow-up after MLLA is provided in 69.2% (27/39) of centres. Where routine follow-up was implemented, all centres (27/27) offered face-to-face appointments, with seven also offering telephone follow-up and two using video consultations. Follow-up is most commonly provided in consultant surgeon-led clinics (23/27), with 14 of these centres also offering a nurse-led and/or rehabilitation clinic follow-up. In four centres, follow-up is conducted solely in nurse-led clinics or by rehabilitation/artificial limb application clinic only, with no surgeon involvement. Overall, 63.0% had outpatient follow-up with the rehabilitation team, showing moderate compliance with recommendations. Adherence was notably low regarding referral to external (peer-to-peer) support groups such as the Limbless Association, with only 36.8% 'commonly' or 'always' offering this service.

Evaluating equipoise

Several domains of post-MLLA care demonstrated considerable variability. Marked differences were seen in the use of postoperative antibiotics, with 61.5% (24/39) of centres reporting routine use whilst 15.4% (6/39) reported that they 'never' or 'rarely' prescribe them. Among these, administration routes also varied with 91.7% (22/24) using intravenous antibiotics exclusively, 12.5% (3/24) using oral only, and 37.5% (9/24) reporting that they used either intravenous or oral depending on the case. Duration also varied with 12.5% (3/24) giving antibiotics for <24 hours, 29.2% (7/24) for 24–48 hours, 25% (6/24) for 48–72 hours, 50% (12/24) for 72 hours to 5 days and 58.3% (14/24) for >5 days.

Surgical skin preparation techniques also differed. Single skin preparation was 'commonly' or 'always' applied in 60.5% (23/38) of centres and double skin preparation in 33.3% (13/39), demonstrating an area of clinical equipoise. Use on incise drapes also showed further disparity, with 40.6% (13/32) of centres always avoiding them whilst 31.3% (10/32) still used them 'commonly' or 'always'.

Routine outpatient follow-up after MLLA was offered in 69.2% (27/39) of centres. Among these, 33.3% (9/27) occurred within 4–6 weeks, 33.3% (9/27) within the first month and the remainder at other time points. Format also differed: 85.2% (23/27) offered consultant-led review, 63.0% (17/27) included rehabilitation-led follow-up and 33.3% (9/27) offered nurse-led care. Notably, nine centres provided consultant-only follow-up while three relied solely on rehabilitation teams without surgical input. These variations highlight differing models of postoperative care delivery across centres.

Discussion

This audit provides insights into the current clinical practices surrounding MLLA and highlights the variability in adherence to current guidelines aimed at reducing SSI. The findings demonstrate that vascular surgeons are the primary specialists performing MLLA, although a notable proportion of centres also involved other specialities such as orthopaedics, general surgery and plastic surgery. The variability in parent speciality may reflect differences in surgical techniques, indication for procedures, patient demographics and subsequently risk factors for SSI development,²⁰ likely contributing to variability in practice. However, this also raises important questions about the potential challenges in developing policies and/or best practice pathways to improve patient outcomes, particularly in the context of SSI prevention.

According to the 1964 wound classification, MLLA wounds are typically classed as 'clean',²¹ and therefore prophylactic antibiotics are not routinely recommended in the NICE guidelines. However, neither the guidelines nor this classification system account for the range of procedures, specialities, incision sites, patient cohorts and the subsequent variability in SSI risk.²² Many believe the increased risk of bacterial contamination secondary to ischaemic or infected tissue in MLLA warrants prophylactic antibiotic use,^{23,24} a consensus that seems in line with the survey results, with most centres (82.1%) administering induction antibiotics. Additionally, evidence suggests benefits of prophylactic postoperative antibiotics following MLLA,^{13,23} a practice also adopted in most centres (61.5%), although there was no clear consensus on duration of therapy. A recent randomised controlled trial¹³ published in 2022, three years after the 2019 NICE guidelines, demonstrated benefits of an extended 5-day course of antibiotic prophylaxis, highlighting how emerging evidence can surpass existing guidelines. It is important to consider that variation in antibiotic choice and administration route is expected due to differing local antimicrobial policies. This remains consistent with existing guidelines, which specify that, where antibiotics are indicated, selection should be guided by local antibiotic formularies, resistance patterns and microbiological tests where available.³ Given the risks of prolonged antibiotic use and antimicrobial resistance, it remains essential to balance antimicrobial stewardship with the prevention of infection.

The intraoperative practices and guideline adherence varied between centres. Hair removal was routinely performed in 61.5% of centres. However, the use of electric clippers, recommended over

razors to minimise micro-abrasions, was not universal, with 25.6% of centres still employing razors often, which may increase the risk of SSIs.²⁵ Current guidelines recommend using an alcohol-based solution of chlorhexidine but do not specify whether single versus double preparation should be employed. However, it is interesting to note the variability in local protocols, with 60.5% routinely using a single skin preparation and 33.3% often employing double skin preparation. There are some data from other surgical specialities demonstrating a reduction in bacterial colonisation with double preparation,²⁶ including a randomised controlled trial of patients undergoing total joint arthroplasty which suggests that double preparation reduces SSI rates;²⁷ however, these are not specific to MLLA. Some centres (21.1%) routinely use iodophor-containing incise drapes, which in other specialities have been shown to reduce SSIs;²⁸ however, no studies have focused on MLLA and many centres commonly or always perform MLLA without the use of incise drapes.

The survey demonstrated widespread use of diagnostic imaging (92.5%) and multidisciplinary team (MDT) discussions (75%) to evaluate revascularisation options and suitability prior to MLLA. This is encouraging as vascular optimisation, when appropriate, reduces the rate of MLLA.^{29,30} Furthermore, imaging review and MDT discussion aid in the complex decision of selecting the appropriate amputation level, balancing functional outcomes against the risk of postoperative ischaemic wound breakdown. Recommendations advocate pre-procedural imaging and perfusion assessments; however, no single test is accepted as the gold standard to predict wound healing³¹ and the decision is often primarily based on clinical judgement.³² Whilst angiography is widely implemented, preoperative perfusion pressures such as TcPO₂ are routinely used in less than half (48.3%) of centres, indicating a lack of standardisation. Perfusion pressures may be used to detect viable tissue for the amputation site. Studies suggest that TcPO₂ values of >40 mmHg are associated with a higher percentage of successful healing whereas values of <20 mmHg may indicate an increased risk of non-healing.³³ However, factors including limb oedema, cardiac output, smoking and pain can reduce accuracy, limiting its reliability as a sole determinant of amputation level. Consequently, there is no consensus regarding a specific threshold value. Despite this, the evidence suggests that perfusion pressures still provide valuable information.³² Additionally, emerging technologies such as machine learning algorithms may enhance risk prediction models by integrating patient risk factors and objective measurements. A recent pilot study demonstrated that machine learning incorporating multispectral wound imaging alongside patient risk factors improved the prediction for amputation wound healing.³¹ As these technologies evolve they may become an integral component of preoperative planning.

Occupational and physiotherapy assessments were widely implemented both preoperatively and postoperatively. Early assessment for rehabilitation can help prepare the patient physically and psychologically for rehabilitation,³⁴ and evidence has shown us

that early postoperative physiotherapy has a significant effect on function.³⁵ However, fewer centres routinely engaged dieticians (21.6%) and psychiatrists (25%) preoperatively. This finding is concerning as malnutrition and psychological distress are risk factors for poor post-surgical recovery and SSI development.^{36,37} Recent studies have highlighted the potential value of integrated approaches such as enhanced recovery after surgery (ERAS) collaborative models³⁸ and surgeon-physician co-management³⁹ to ensure optimal prehabilitation and perioperative management,⁴⁰ working towards better patient outcomes.

Follow-up varies significantly between centres, with almost one-third of centres (30.2%) not routinely providing routine follow-up after MLLA, which could lead to wound complications such as SSI and wound breakdown being under-diagnosed and therefore under-treated. Furthermore, less than half of centres routinely provide follow-up with a rehabilitation clinic (42.5%) or refer patients to external peer-to-peer support groups (36.8%) such as the Limbless Association. This lack of routine referral to support services may reflect under-appreciation of the role these resources play in long-term recovery and rehabilitation, given the significant physical and psychosocial impact MLLA often has. Greater integration of support services could improve patient outcomes and quality of life.⁴¹

Although this audit highlights significant variability in guideline adherence, it is important to note the limitations of the guidelines themselves. Some recommendations are outdated and others are derived from low-quality evidence,^{42,43} and recent trials have challenged existing guidelines such as the FALCON trial which questioned the superiority of chlorhexidine preparation in clean surgery.^{44,45} Notably, a recent study on diabetic foot disease reported similar findings, highlighting inconsistent adherence to guidelines and a lack of robust randomised controlled trial evidence supporting their foundation.⁴⁶ More critically, the SSI prevention and management guidelines are designed for broader surgical contexts and do not specifically address MLLA. As a result, their applicability and effectiveness in this cohort are uncertain, given the risk of SSI is multifactorial, influenced by patient comorbidities, procedural techniques and perioperative care beyond the scope of current recommendations. Moreover, there is a lack of high-quality evidence supporting the efficacy of intervention 'bundles' (combinations of individually effective interventions) in reducing SSI rates when implemented concurrently.⁴⁷ This audit also identified several areas of clinical equipoise, where significant variation in practice reflects a genuine lack of consensus of optimal management. Notably, the use of postoperative antibiotics showed great variability, with significant differences in route of administration and duration, ranging from 24 hours to >5 days. Similarly, intraoperative techniques such as single versus double skin preparation and the use of incise drapes showed substantial disparity between centres. Variability in preoperative MDT assessments and postoperative follow-up further reflected uncertainty in the holistic care aspect for those undergoing MLLA.

This highlights the need for robust procedure-specific evidence to guide best practice in MLLA. Current works, including the ROSSINI-Platform trial designed to evaluate SSI prevention strategies across surgical specialities including MLLA,⁴⁸ offer promising opportunities to continue addressing these evidence gaps. Furthermore, the European Society for Vascular Surgery is commissioning a Clinical Practice Guidelines specific for MLLA, set for publication in 2027.⁴⁹

Study limitations

There are some limitations to this audit. The survey responses were self-reported, which may introduce recall or social desirability bias and affect the accuracy of reported adherence. However, the main SIMBA study includes prospective data collection, which will help validate these findings against actual clinical practice. Additionally, the survey was limited to centres that participated in the SIMBA audit, with an overall response rate of 87%. Non-responders included four UK sites and two European sites. This may have introduced response bias, and the relatively small sample size and geographical concentration within the UK may limit the generalisability of these findings. There are also some limitations in the survey design. The survey was designed and internally validated by the SIMBA study management group, composed primarily of vascular surgeons and researchers, without external validation or wider multidisciplinary input, which may have enhanced its comprehensiveness and applicability. As a result, some terms – for example, ‘MDT’ – may have been interpreted inconsistently. Additionally, we did not collect data on the availability of specific services (eg, psychology or rehabilitation services), which limits our ability to determine whether the absence of routine assessment reflects a lack of access or other factors such as surgical urgency. Similarly, the survey did not include qualitative fields or free-text options to explore reasons for ‘don’t know’ responses, which may reflect uncertainty or variation in terminology rather than true gaps in practice. Although some international guidelines were used, most recommendations assessed were based on UK-specific sources (eg, NICE and VSGBI), which may limit their applicability to international centres. Furthermore, while this audit provides a snapshot of current practice, it does not capture the real-time impact of these practices on SSI rates and patient outcomes. However, by evaluating adherence to established guidelines and identifying areas of clinical equipoise, this study highlights key areas for future research and focus points for future evidence-based guidelines tailored to this patient population.

Conclusion

This international multicentre audit highlights substantial variability in clinical practice and adherence to SSI prevention guidelines and best practice pathways in centres performing MLLA. Whilst there is good adherence to certain recommendations such as diagnostic imaging and multidisciplinary care, gaps remain, particularly in areas of preoperative nutritional and psychological evaluation,

KEY MESSAGES

- Surgical site infection is a common complication after amputation, potentially leading to prolonged hospital stays, revision amputations and even mortality.
- This survey revealed significant variation in practices including antibiotic use, preoperative assessments and postoperative care across 46 global centres.
- There is need for more tailored, evidence-based guidelines to reduce infection risk and improve patient outcomes following amputation.

intraoperative standardisation and postoperative support. However, given the lack of specificity of guidelines and the multifactorial nature of the risk of SSI, a more tailored evidence-based approach is needed. Future research should prioritise high-quality procedure-specific studies to evaluate the impact of different perioperative strategies on reducing the risk of SSI. This will facilitate the development of standardised care pathways, ultimately improving clinical outcomes for patients undergoing MLLA.

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Appendix 1 SIMBA Audit: Baseline Unit Survey

Start of Block: Default Question Block

Q1 The purpose of the survey is to understand the usual clinical care pathway and policies for managing patients undergoing a Major Lower Limb Amputation (MLLA) at your unit.

End of Block: Default Question Block

Start of Block: Block 1

Q2 Which country is your Unit based?



Q3 Which unit are you based?

▼ ... Other, please specify (70)

Q4 If unit is not listed, please specify:

End of Block: Block 1

Start of Block: Block 2

Page Break

Q5 Which specialty(s) at your unit performs MLLA (tick all that apply):

- ☐ Vascular
 - ☐ Orthopaedics
 - ☐ General Surgery
 - ☐ Other (please specify)
-

Q7 How often are patients who are being considered for MLLA referred to the MDT for assessment prior to surgery?

- ☐ Never
 - ☐ Rarely
 - ☐ Sometimes
 - ☐ Commonly
 - ☐ Always
 - ☐ Do not know
-

Q9 How often do patients with lower limb ischaemia who are being considered for MLLA undergo diagnostic imaging to determine revascularisation options?

- ☐ Never
- ☐ Rarely
- ☐ Sometimes
- ☐ Commonly
- ☐ Always
- ☐ Do not know
-

Q11 How often are the revascularisation options discussed at the Vascular Imaging MDT meeting (Vascular Surgeons plus Interventional Radiology) OR Orthopaedic MDT?

	Never	Rarely	Sometimes	Commonly	Always	Do not know
Vascular Imaging MDT	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Orthopaedic MDT	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

Q12 How often are the MDT decisions documented in the patient's medical notes (electronic or handwritten patient record)?

- ☐ Never
 - ☐ Rarely
 - ☐ Sometimes
 - ☐ Commonly
 - ☐ Always
 - ☐ Do not know
-

Q14 How often are patients undergoing MLLA admitted under a named Consultant/Attending in Vascular or Orthopaedic Surgery as appropriate for indication of MLLA?

- ☐ Never
 - ☐ Rarely
 - ☐ Sometimes
 - ☐ Commonly
 - ☐ Always
 - ☐ Do not know
-

Q16 How often are patients undergoing MLLA undergo routinely undergo assessment with:

	Never	Rarely	Sometimes	Commonly	Always	Do not know	Not applicable
Psychology (pre op/post op)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
OT/PT	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Dietician	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Diabetic team	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

Q18 How often do patients undergoing MLLA have venous thromboembolism risk assessment and prophylaxis according to local protocols?

- ☐ Never
 - ☐ Rarely
 - ☐ Sometimes
 - ☐ Commonly
 - ☐ Always
 - ☐ Do not know
-

Q20 How often in patients undergoing MLLA for lower limb ischaemia do you use objective assessment of perfusion pressure, i.e. TcPO₂, to determine level of amputation e.g. BKA vs TKA vs AKA?

- ☐ Never
 - ☐ Rarely
 - ☐ Sometimes
 - ☐ Commonly
 - ☐ Always
 - ☐ Do not know
-

Q21 Does your centre have a standard induction antibiotic regime for MLLA

- ☐ Yes
 - ☐ No
-

Q22 If yes, please select the antibiotic(s) in the antibiotic prophylaxis guideline:

- ☐ Penicillins (excluding Tazocin)
 - ☐ Cephalosporins
 - ☐ Carbapenems
 - ☐ Metronidazole
 - ☐ Clindamycin
 - ☐ Linezolid
 - ☐ Daptomycin
 - ☐ Fluoroquinolones
 - ☐ Vancomycin
 - ☐ Tigecycline
 - ☐ Gentamycin
 - ☐ Tazocin
 - ☐ Co-trimoxazole
 - ☐ Doxycycline
 - ☐ Teicoplanin
 - ☐ Other _____
 - ☐ Unknown
-

Q23 Administration method:

- ☐ IV
- ☐ IM
- ☐ PO
- ☐ Topical
- ☐ Other-specify _____
-

Q25 How often are incise drapes for MLLA surgery used at your centre?

	Never	Rarely	Sometimes	Commonly	Always	Do not know
Iodophor containing incise drapes	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Non-iodophor containing incise drapes	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
No incise drapes	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

Q27 How often is hair routinely removed pre-operatively for MLLA surgery?

- ☐ Never
- ☐ Rarely
- ☐ Sometimes
- ☐ Commonly
- ☐ Always
- ☐ Do not know
-

Q28 If hair removal is undertaken, which tool is used?

- ☐ Electric Clippers
- ☐ Razor
- ☐ Other-specify _____
- ☐ Not applicable
-

Q29 When applying skin preparation, is a single preparation or double preparation applied?

	Never	Rarely	Sometimes	Commonly	Always	Do not know
Single Prep	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Double Prep	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

Q33 How often do patients undergo their first review by the acute pain team on day 1 post-operatively?

- ☐ Never
- ☐ Rarely
- ☐ Sometimes
- ☐ Commonly
- ☐ Always
- ☐ Do not know
-

Q35 How often do patients undergoing MLLA routinely get support in the form of:

	Never	Rarely	Sometimes	Commonly	Always	Do not know
Pre-operative Information leaflets about the operation itself	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Post-operative information leaflets about recovery	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Psychologists	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
External support groups (e.g., Limbless Association)	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

Q36 How often do patients at your unit have a post-operative course of prophylactic antibiotics prescribed after undergoing MLLA?

- ☐ Always
- ☐ Commonly
- ☐ Sometimes
- ☐ Rarely
- ☐ Never
- ☐ Do not know

Q37 If yes, please select the antibiotic(s) in the antibiotic prophylaxis guideline:

- ☐ Penicillins (excluding Tazocin)
 - ☐ Cephalosporins
 - ☐ Carbapenems
 - ☐ Metronidazole
 - ☐ Clindamycin
 - ☐ Linezolid
 - ☐ Daptomycin
 - ☐ Fluoroquinolones
 - ☐ Vancomycin
 - ☐ Tigecycline
 - ☐ Gentamycin
 - ☐ Tazocin
 - ☐ Co-trimoxazole
 - ☐ Doxycycline
 - ☐ Teicoplanin
 - ☐ Other _____
 - ☐ Unknown
-

Q38 Administration method:

☐

IV

☐

IM

☐

PO

☐

Topical

☐

Other-specify

Q39 Length of course:

☐

0-24 hrs

☐

24-48 hrs

☐

48-72 hrs

☐

72hrs-5 days

☐

>5 days

☐

Other-specify:

Q41 How often do patients and carers receive information on wound care after discharge, including information on how to recognise a surgical site infection and who to contact if they are concerned?

- ☐ Never
 - ☐ Rarely
 - ☐ Sometimes
 - ☐ Commonly
 - ☐ Always
 - ☐ Do not know
-

Q42 Are patients routinely follow-up after MLLA?

- ☐ Yes
- ☐ No

Q43 When are patients routinely followed up following MLLA?

- ☐ 0-2 weeks
 - ☐ 2-4 weeks
 - ☐ 4-6 weeks
 - ☐ 6-8 weeks
 - ☐ 8-12 weeks
 - ☐ >12 weeks
 - ☐ Other, please specify _____
-

Q44 How are patients followed up? Tick all that apply

- ☐ Consultant led clinic
- ☐ Nurse led clinic
- ☐ Rehabilitation clinic / Artificial Limb Application Clinic
- ☐ Other: please specify

(_____)

Q45 What is the format of the follow-up appointment? Tick all that apply

- ☐ Face to face appointment
- ☐ Video appointment
- ☐ Telephone appointment
- ☐ Other: please specify

Page Break

End of Block: Block 2

Start of Block: Block 3

Q47 Thank you for completing this survey.

End of Block: Block 3

Appendix 2 Checklist for Reporting Of Survey Studies (CROSS)

Section/topic	Item	Item description	Reported on page #
Title and abstract			
Title and abstract	1a	State the word “survey” along with a commonly used term in title or abstract to introduce the study’s design.	Title Page
	1b	Provide an informative summary in the abstract, covering background, objectives, methods, findings/results, interpretation/discussion, and conclusions.	1
Introduction			
Background	2	Provide a background about the rationale of study, what has been previously done, and why this survey is needed.	4
Purpose/aim	3	Identify specific purposes, aims, goals, or objectives of the study.	5
Methods			
Study design	4	Specify the study design in the methods section with a commonly used term (e.g., cross-sectional or longitudinal).	6,7
Data collection methods	5a	Describe the questionnaire (e.g., number of sections, number of questions, number and names of instruments used).	6,7
	5b	Describe all questionnaire instruments that were used in the survey to measure particular concepts. Report target population, reported validity and reliability information, scoring/classification procedure, and reference links (if any).	7
	5c	Provide information on pretesting of the questionnaire, if performed (in the article or in an online supplement). Report the method of pretesting, number of times questionnaire was pre-tested, number and demographics of participants used for pretesting, and the level of similarity of demographics between pre-testing participants and sample population.	7
	5d	Questionnaire if possible, should be fully provided (in the article, or as appendices or as an online supplement).	Appendix 2
	6a	Describe the study population (i.e., background, locations, eligibility criteria for participant inclusion in survey, exclusion criteria).	6
Sample characteristics	6b	Describe the sampling techniques used (e.g., single stage or multistage sampling, simple random sampling, stratified sampling, cluster sampling, convenience sampling). Specify the locations of sample participants whenever clustered sampling was applied.	6
	6c	Provide information on sample size, along with details of sample size calculation.	6
	6d	Describe how representative the sample is of the study population (or target population if possible), particularly for population-based surveys.	6
Survey	7a	Provide information on modes of questionnaire administration, including the type and number of contacts, the location where the survey was conducted (e.g., outpatient	6,7

administration		room or by use of online tools, such as SurveyMonkey).	
	7b	Provide information of survey's time frame, such as periods of recruitment, exposure, and follow-up days.	6,7
		Provide information on the entry process:	7
	7c	→For non-web-based surveys, provide approaches to minimize human error in data entry.	7
		→For web-based surveys, provide approaches to prevent "multiple participation" of participants.	
Study preparation	8	Describe any preparation process before conducting the survey (e.g., interviewers' training process, advertising the survey).	7
Ethical considerations	9a	Provide information on ethical approval for the survey if obtained, including informed consent, institutional review board [IRB] approval, Helsinki declaration, and good clinical practice [GCP] declaration (as appropriate).	NA
	9b	Provide information about survey anonymity and confidentiality and describe what mechanisms were used to protect unauthorized access.	NA
	10a	Describe statistical methods and analytical approach. Report the statistical software that was used for data analysis.	7
	10b	Report any modification of variables used in the analysis, along with reference (if available).	NA
Statistical analysis	10c	Report details about how missing data was handled. Include rate of missing items, missing data mechanism (i.e., missing completely at random [MCAR], missing at random [MAR] or missing not at random [MNAR]) and methods used to deal with missing data (e.g., multiple imputation).	7
	10d	State how non-response error was addressed.	NA
	10e	For longitudinal surveys, state how loss to follow-up was addressed.	NA
	10f	Indicate whether any methods such as weighting of items or propensity scores have been used to adjust for non-representativeness of the sample.	NA
	10g	Describe any sensitivity analysis conducted.	NA

Results

Respondent characteristics	11a	Report numbers of individuals at each stage of the study. Consider using a flow diagram, if possible.	8
	11b	Provide reasons for non-participation at each stage, if possible.	NA
	11c	Report response rate, present the definition of response rate or the formula used to calculate response rate.	8

Descriptive results	11d	Provide information to define how unique visitors are determined. Report number of unique visitors along with relevant proportions (e.g., view proportion, participation proportion, completion proportion).	8
	12	Provide characteristics of study participants, as well as information on potential confounders and assessed outcomes.	8
	13a	Give unadjusted estimates and, if applicable, confounder-adjusted estimates along with 95% confidence intervals and p-values.	8-9
Main findings	13b	For multivariable analysis, provide information on the model building process, model fit statistics, and model assumptions (as appropriate).	NA
	13c	Provide details about any sensitivity analysis performed. If there are considerable amount of missing data, report sensitivity analyses comparing the results of complete cases with that of the imputed dataset (if possible).	NA
Discussion			
Limitations	14	Discuss the limitations of the study, considering sources of potential biases and imprecisions, such as non-representativeness of sample, study design, important uncontrolled confounders.	14,15
Interpretations	15	Give a cautious overall interpretation of results, based on potential biases and imprecisions and suggest areas for future research.	12-14
Generalizability	16	Discuss the external validity of the results.	14-16
Other sections			
Role of funding source	17	State whether any funding organization has had any roles in the survey's design, implementation, and analysis.	24
Conflict of interest	18	Declare any potential conflict of interest.	24
Acknowledgements	19	Provide names of organizations/persons that are acknowledged along with their contribution to the research.	Title Page

Appendix 3 Supplementary Table 1 - Guideline adherence UK centres only

Guideline/recommendation (Grade of evidence where stated)	Recommending organisation				Relevant survey question	Percentage adherence (UK)
	NICE	EJVES	VSGBI	SVS		
Preoperative						
Patients should be assessed by the MDT prior to MLLA		X	X		7	46.7%*
Offer patients and carers clear, consistent information and advice through all stages of their care	X				35a	48.3%*
Be admitted under a named consultant in vascular surgery			X		14	96.6%*
Undergo diagnostic arterial imaging to determine revascularisation options			X	X	9	96.6%*
Have revascularisation options discussed at a vascular imaging MDT			X		11	83.3%*
Undergo assessment using TcPO ₂ to determine perfusion at a proposed amputation level				X	20	31.0%*
Involvement of clinical psychology				X	16a	26.9%*
Undergo assessment with OT/PT preoperatively (1C).		X	X		16b	86.2%*
Have nutritional assessment and receive dietician advice.			X		16c	18.5%*
Have a member of diabetes team involved			X		16d	69.0%*
Have a venous thrombo-embolism risk assessment and prophylaxis as appropriate			X	X	18	100%*
Perioperative						
Antibiotic prophylaxis should not routinely be used for clean non-prosthetic uncomplicated surgery	X				21	17.2%
Do not use non-iodophor-impregnated incise drapes routinely for surgery	X				25	67.9%**
Do not use hair removal routinely	X				27	10.3%**
If hair has to be removed, use electric clippers.	X				28	85.7%

Appendix 3 Supplementary Table 1 - Guideline adherence UK centres only continued

Postoperative						
Antibiotic prophylaxis should not routinely be used for clean non-prosthetic uncomplicated surgery	X				36	20.7%**
Patients should be informed of the post-amputation care pathway		X			35b	48.3%*
Offer patients and carers information and advice about how to recognise a surgical site infection and who to contact if they are concerned	X				41	65.5%*
Day 1 postoperative review by acute pain team			X		33	69.0%*
Follow-up in clinic within a month after surgery				X	43	16.6%*
Outpatient review and rehabilitation follow up with rehabilitation team (1C)		X	X	X	44	63%*
Referred to amputation support group.				X	35d	37%*
<p>* Percentage selecting 'commonly' or 'always'.</p> <p>** Percentage selecting 'never' or 'rarely'.</p> <p>EJVES, European Journal of Vascular and Endovascular Surgery; MDT, multidisciplinary team; NICE, National Institute for Health and Care Excellence; OT, occupational therapy; PT, physiotherapy; SVS, Society of Vascular Surgery; VSGBI, Vascular Societies of Great Britain and Ireland.</p>						