A UK-based pilot study of current surgical practice and implant preferences in lumbar fusion surgery

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Abstract

Lumbar fusion surgery is an established procedure for the treatment of low back pain. Despite the wide set of alternative fusion techniques and existing devices, uniform guidelines are not available yet and common surgical trends are scarcely investigated. The purpose of this UK-based study was to provide a descriptive portrait of current surgeons’ practice and implant preferences in lumbar fusion surgery.

A UK-based in-person survey was designed for this study and submitted to a group of consultant spinal surgeons (n = 32). Fifteen queries were addressed based on different aspects of surgeons’ practice: lumbar fusion techniques, implant preferences, and bone grafting procedures. Answers were analyzed by means of descriptive statistics.

Thirty-two consultant spinal surgeons completed the survey. There was clear consistency on the relevance of a patient-centered management (82.3%), along with a considerable variability of practice on the preferred fusion approach. Fixation surgery was found to be largely adopted (96.0%) and favored over stand-alone cages. With regards to the materials, titanium cages were the most used (54.3%). The geometry of the implants influenced the choice of lumbar cages (81.3%). Specifically, parallel-shape cages were mostly avoided (89.2%) and hyperlordotic cages were preferred at the lower lumbar levels. However, there was no design for lumbar cages which was consistently favored. Autograft bone graft surgeries were the most common (60.0%). Amongst the synthetic options, hydroxyapatite-based bone graft substitute was (87.6%) in injectable paste form (80.8%) were preferred.

Current lumbar fusion practice is variable and patient-oriented. Findings from this study highlight the need for large-scale investigative surveys and clinical studies aimed to set specific guidelines for certain pathologies or patient categories.

Abbreviations: ALIF = anterior lumbar interbody fusion, DLIF = direct lumbar interbody fusion, HA = hydroxyapatite, LIF = lumbar interbody fusion, LLIF = lateral lumbar interbody fusion, MIS = minimally invasive surgery, NHS = National Health Service, PEEK = polyetheretherketone, PLIF = posterior lumbar interbody fusion, TDR = total disc replacement, XLIF = extreme lateral interbody fusion.

Keywords: bone graft substitutes, fusion implant, interbody cage, lumbar fusion, spine surgery, surgical practice, survey

1. Introduction

Low back pain caused by spinal disorders such as degenerative disc diseases, congenital deformities and trauma affects over 80% of the population worldwide.\cite{11} Spinal fusion surgery is an established and widely adopted procedure for the treatment of low back pain and has recently experienced a substantial growth among the major developed countries. In the United States, the number of spinal fusion procedures increased by 77% between 2002 and 2011.\cite{21} In the United Kingdom, recent data highlighted a similar trend between 2005 and 2015 with an increase of 63% of spinal fusion procedures.\cite{11} Within National Health Service (NHS), spinal surgery was also reported as the largest single component of expenditure for the management of low back pain, with direct costs estimated to be over £1.6 billion per year.\cite{4,5} The spinal fusion market is expected to grow further from $4775m, in 2013, to $6982m by 2020 across United States, France, Germany, Italy, Spain, United Kingdom, Japan, Brazil, India, and China.\cite{6} Such worldwide increase might be partly explained by the recent advances in spinal implants and the development of less-invasive surgical methods.\cite{7} Moreover, novel solutions in terms of bio compatible materials and osteoconductive bone graft substitutes have been shown to reduce pseudarthrosis.\cite{8,9} Currently there are 4 main different approaches to lumbar interbody fusion: anterior lumbar interbody fusion (ALIF), posterior lumbar interbody fusion (PLIF), extreme lateral or direct lumbar interbody fusion (XLIF, DLIF), and transfemoral lumbar interbody fusion (TLIF). There is still no evidence that one surgical approach is clinically superior to the alternatives and indications vary based on the surgeon and the patient.\cite{10,12} The ALIF approach provides with a clear visualization and efficient access to the intervertebral space, thus
allowing for a complete discectomy and the placement of a large interbody fusion implant to maximize the contact area.\textsuperscript{[13,14]} Moreover, lumbar lordosis and sagittal balance can be efficiently restored with anterior hyperlordotic cages.\textsuperscript{[13,16]} These advantages have been shown to contribute to a reduced perioperative morbidity and higher fusion rates compared to other traditional approaches.\textsuperscript{[17]} ALIF is particularly favored for the treatment of degenerative disc disease, chronic pain, instability, and deformity at the lumbosacral junction; whilst contraindications include spondylolisthesis greater than grade II, severe vascular disease or inappropriate vascular anatomy.\textsuperscript{[10,18,19]} Stand-alone ALIF cages may be a suitable surgical option in cases with single-level degenerative disc disease; however, posterior fixation may be required for more difficult surgical cases.\textsuperscript{[14,20]} The PLIF approach is one of the most traditional and familiar techniques for spinal surgeons and have been related with an acceptable degree of fusion and a relatively low complication rate.\textsuperscript{[21,22]} Indeed, the posterior approaches avoid vascular complications associated with the anterior approach. However, because of the posterior surgical access, disadvantages of the technique include nerve root retraction and possible associated neurological complications.\textsuperscript{[23]} The traditional PLIF approach utilizes 2 small interbody implants and it is typically followed by posterior fixation. PLIF has been recommended for spondylolisthesis, retrolisthesis, spinal stenosis, and lumbar disc herniation.\textsuperscript{[24,25]} XLIF or DLIF, also referred as lateral lumbar interbody fusion (LLIF), is recognized as a less invasive technique compared to ALIF and PLIF, which has been correlated with reduced operation time and shorter hospital stays.\textsuperscript{[26,27]} Despite these advantages, XLIF has been also associated with a considerably higher frequency of neurological complications and other major morbidity.\textsuperscript{[28]} XLIF usually utilizes a single ovoid implant and is anatomically indicated from the lumbar levels L1/L2 to L4/L5, whilst it is avoided at the L5/S1 level.\textsuperscript{[7]} Indications for XLIF combined with posterior fixation include lumbar spinal stenosis and adjacent segment disease.\textsuperscript{[29,30]} Another posterior fusion approach is TLIF, which was proposed to reduce the extent of neural retraction required during the traditional PLIF approach. Accordingly, TLIF has been associated with less blood loss and reduced risk of injury to the neural and vascular structures, as well as shorter operating time.\textsuperscript{[31]} During TLIF, a unilateral access to the intervertebral space is achieved by sparing the unilateral laminectomy and a partial facetectomy to allow the insertion of a single “banana”-shaped or rectangular interbody fusion implant. Posterior fixation is usually performed to provide immediate segmental stability and potentially enhance the fusion outcomes.\textsuperscript{[32]} TLIF has been indicated for the treatment of low back pain associated with grade I or II spondylolisthesis, degenerative disc disease, recurrent disc herniation, and revision surgery.\textsuperscript{[13]} All the fusion approaches described above can now be performed using minimally invasive surgery (MIS) techniques, which have been shown to reduce intraoperative blood loss and shorten hospital stay.\textsuperscript{[34,35]} Long-term data are needed to examine the clinical effectiveness of MIS versus open LIF procedures.\textsuperscript{[16,27]} As an alternative to lumbar fusion, total disc replacement (TDR) is a motion-preserving technique which has been indicated for the treatment of symptomatic lumbar degenerative disc disease, although superiority compared to fusion has not yet been proved.\textsuperscript{[36,37]} While the number of available surgical options increases, the current practice related to lumbar fusion surgery has been scarcely investigated and no comprehensive information on fusion implant trends is available in the literature.\textsuperscript{[5,40,42]} With this UK-based study, we aim to provide a preliminary investigation of current surgeon practice and implant preferences in lumbar fusion surgery, by specifically focusing on the relevant factors which influence surgeons’ decision making process.

2. Methods

Data were obtained from consultant spinal surgeons to investigate the various factors defining the choice of a specific surgical procedure and interbody fusion implant. This was performed by designing a questionnaire survey which was submitted to a group of eligible participants.

2.1. Survey design and sample size

An in-person survey was conducted at BritSpine 2016 (April 6–8, 2016), in Nottingham, United Kingdom. Eligibility was limited to consultant spinal surgeons with at least 1 year experience as postholder. Eligibility was verified at a preliminary stage of the interviews before the submission of the survey.

The questionnaire was developed by the authors of this paper taking into account the perspectives of both academic researchers and clinicians. At the design stage, we incorporated feedbacks from researchers with survey experience and phrased the most pertinent questions to assure survey effectiveness. The survey was designed to be completed in approximately five minutes. The questionnaire included 15 questions (12 multiple-choice and 3 Likert scale) and consisted of 4 sections aiming to address the following aspects: (A) surgeon’s background and experience; (B) choice of surgical procedure for lumbar fusion; (C) factors influencing implant choice; (D) use of bone graft materials.

Section A included general demographic questions on surgeons’ specialty and experience as postholders.

Section B enquired about the number of spinal fusion surgeries performed in the last year of surgeons’ practice. A Likert scale (1–5 ranking) question was included to investigate the relevance of patient-specific evaluation and spinal segment for the choice of the surgical procedure. Questions on the preferred fusion surgeries were also included in this part of the survey.

Section C comprised 2 Likert scale queries (1–5 ranking) on the importance of clinical, market-related factors, and geometrical parameters for the choice of the fusion implant. A multiple-choice question addressed the correlation between cage lordotic angle and affected spinal segment. Queries on the preferred implant material, structure and design were also included. The latter was investigated by asking which design resembled surgeons’ favorite implant. Specifically, 6 different profiles of cage endplate geometry and 6 different designs for bone grafting area were shown.

Finally, section D aimed to investigate the current bone grafting procedures and the potential use of synthetic graft substitutes. The questionnaire is available as the Supplementary information to this manuscript, http://links.lww.com/MD/C301.

2.2. Data collection and analysis

An in-person methodology was selected for this survey to ensure the maximum response rate and minimize the chance of incomplete responses. During the conference, 32 eligible consultant spine surgeons were asked to complete a questionnaire presented in paper form. Data were collected anonymously and entered into a datasheet based on a randomized ID. Due to the nature of this study, no assessment by a medical ethics review
board was required. All data were analyzed using descriptive statistics using GraphPad Prism Software, by combining simple graphical and tabular descriptive summaries. Frequencies were depicted as percentages of valid responses.

3. Results

3.1. Demographics

Thirty-two questionnaires were returned: 22 fully completed, 10 with an average completion rate of 91.4%. Demographics of the participants were as follow: 87.5% of respondents were orthopedic spine surgeons, whereas only 12.5% of respondents were neurosurgeons. Surgeons’ medical specialties and experience as consultants are reported in Table 1.

3.2. Surgical procedures

3.2.1. Fusion approaches. The majority (58.1%) of surgeons performed 11 to 50 cases of TLIF. Accordingly, the most frequently applied fusion technique was TLIF (40.4%), followed by PLIF (28.1%), ALIF (15.8%), and XLIF or DLIF (14.0%). Table 2 shows the occurrences of surgical cases categorized according to each surgical procedure.

The majority of surgeons agreed that the choice of surgical procedure (84.4%) and fusion approach (87.5%) was based on a patient-specific evaluation. The spinal segment was also considered as a relevant parameter for the selection of the surgical procedure and fusion approach by the 78.1% and 81.3% of surgeons, respectively (Fig. 1).

3.2.2. Fixation procedures. Questions on the preferred fixation choices were completed by all the 32 respondents. The preferred fixation procedure involved the insertion of pedicle screws, according to the 46% of total responses. Conversely, the 24% and 16% of participants’ responses favored anterior screws fixation or a combination of anterior and posterior fixation, respectively. The use of fusion cages equipped with anterior fixation plates was reported as first choice for the 10% of responses. Stand-alone cages were selected in only 4% of responses.

3.3. Implant choice

3.3.1. Clinical and market-related factors. Patient specific evaluation and evidence of clinical outcomes turned out to be most relevant factors for the choice of fusion implants, according to the 75.0% and 81.3% of respondents, respectively. The majority of surgeons (81.3%) also considered implant geometry as a crucial parameter influencing implant choice, whilst product price and manufacturer were recognized as important by the 68.8% and 37.5% of surgeons, respectively (Fig. 2).

3.3.2. Material choice. Twenty-eight surgeons answered the questions related to the material structure. The most frequently used material for cages were titanium (68%), followed by polyether ether ketone (68%), and polyether ketone (28%).

Table 1

| Surgeons’ background and experience (n = 32). | Number of respondents (%) |
|---------------------------------------------|---------------------------|
| Medical specialty                            |                           |
| Orthopedic spine surgeons                   | 28 (87.5)                 |
| Neurosurgeons                               | 4 (12.5)                  |
| Years of experience as consultant           |                           |
| <5                                          | 11 (34.4)                 |
| 5–9                                         | 6 (18.8)                  |
| 10–14                                       | 6 (18.8)                  |
| 15–19                                       | 5 (15.6)                  |
| 20 or more                                  | 3 (12.5)                  |

Table 2

| Surgical procedure | None | 1–10 cases | 11–50 cases | 51–100 cases | >100 cases |
|--------------------|------|------------|-------------|--------------|------------|
| Anterior (ALIF)    | 8 (26.7) | 12 (40.0) | 10 (33.3) | 0 (0) | 0 (0) |
| Posterior (PLIF)   | 9 (28.1) | 10 (31.3) | 10 (31.3) | 2 (6.3) | 1 (3.1) |
| Lateral (XLIF/DLIF)| 13 (41.9) | 9 (29.0) | 8 (25.8) | 1 (3.2) | 0 (0) |
| Transforaminal (TLIF) | 4 (12.9) | 6 (19.4) | 18 (58.1) | 3 (9.7) | 0 (0) |
| Total disc replacement (TDR) | 26 (83.9) | 5 (16.1) | 0 (0) | 0 (0) | 0 (0) |

Figure 1. Relevance of a patient-specific evaluation and affected spinal segment for the selection of surgical procedure (e.g., fusion or disc replacement surgery) and fusion approach (e.g., anterior or posterior interbody fusion) (n = 32).
used material for fusion implants was found to be titanium, according to the 54.3% of responses, followed by tantalum and polyetheretherketone (PEEK) with 28.6% and 17.1% of responses, respectively. Overall, porous/trabecular implants were the first choice for 54.8% of responses, whilst 41.9% were nonporous materials with rough/threaded topology. Implants with smooth surface were consistently disregarded.

### 3.3.3. Implant geometry

Cage width (90.7%), length (87.5%), thickness (93.8%), and lordotic angle (93.8%) were all considered important variables by the majority of respondents. The majority of surgeons (56.8%) favored implants with double-side angle, whilst 32.4% and 10.8% of respondents chose one-side angled cages and parallel-shape cages, respectively. The correlation between cage lordotic angle and affected spinal segment is shown in Table 3. The most frequently used implant at the L1/L2 (56.7%), L2/L3 (51.7%), and L3/L4 (45.2%) levels was a 5° angled cage, whereas a 15° angled cage was preferred at the L5/S1 segment. At the L4/L5 level, surgeons similarly favored 5° (29.0%), 10° (32.3%), and 15° (32.3%) angle cages.

Cage endplate designs d and e were favored equally according to the 33.3% of responses, followed by design f at 19.4% (Fig. 3). For bone grafting area, design c with 2 bone graft voids was the preferred profile based on 35.5% of responses (Fig. 4). Instead,

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**Table 3**

| Lumbar segment | Number of occurrences (%) |
|----------------|---------------------------|
|                | 0° | 5°  | 10° | 15° | >15° |
| L1/L2          | 9  | 17  | 3   | 1   | 0    |
| L2/L3          | 6  | 15  | 7   | 1   | 0    |
| L3/L4          | 2  | 14  | 12  | 3   | 0    |
| L4/L5          | 1  | 9   | 10  | 10  | 1    |
| L5/S1          | 1  | 2   | 9   | 16  | 2    |

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**Figure 2.** Relevance of generic parameters, such as patient evaluation, clinical evidence, implant geometry, price, and manufacturer which may influence the selection of fusion implants (n=32).

**Figure 3.** Selection of an endplate geometry of implant by surgeons during fusion surgery procedures (n=31).

**Figure 4.** Selection of implant design based on available bone grafting area by spinal surgeons (n=26).
the 22.6% of participants’ responses favored a single-hole design b, whereas the 16.1% (5) of responses depicted a complex design f.

3.4. Bone graft materials

Most interviewed surgeons preferred autograft bone grafting procedures according to the 60.0% of responses, whilst the 33.3% of responses indicated synthetic bone graft substitutes (Table 4). Within the nonautograft bone graft solutions, the preferred materials were synthetic hydroxyapatite (HA)-based composites according to the 76.7% of responses. The preferred form for bone graft substitutes was injectable paste according to the 80.8% of responses.

4. Discussion

This UK-based pilot study provides a descriptive portrait of current surgical practice in lumbar fusion surgery. To-date, only few studies have described the current practice for the treatment of low back pain. However, none of them focused also on surgeons’ implant decision making process. Hence, the purpose of our survey was to provide not only a snapshot of the current treatment of choice in the United Kingdom, but also to explore the rationale behind each choice.

Amongst the possible surgical procedures, lumbar fusion is the most prevalent surgery. Indeed, the majority of respondents (83.9%) performed no cases of lumbar artificial disc replacement over their last year of clinical practice. There is some variability of practice related to the possible fusion approaches, which might confirm the agreement for a patient-centered care and individual patient management. Despite this variability, TLIF is the most performed fusion approach rather than PLIF, ALIF, or XLIF/DRIF. However, there is no convincing clinical evidence in the literature which supports the choice of a specific surgical technique for a certain population of patients. Our findings illustrate that spinal instrumentation is widely adopted as an adjunct to lumbar fusion. Accordingly, fixation surgery is largely favored (96.0%) over the use of stand-alone cages. Posterior fixation with pedicle screws is the most common procedure. However, there is no clear uniformity of opinion as it pertains to fixation surgery, which is in accordance with the variety of possible fusion approaches. Importantly, our findings consistently highlight the relevance of a patient-oriented management for the choice of the surgical procedure (84.4%), fusion approach (87.5%), and associated interbody fusion implant (75.0%).

With regards to the shape and materials of the device, there is limited evidence in the literature concerning the current fusion implant trends among surgeons’ practice. Interbody fusion cages are in fact available in a wide range varying materials, shapes, and architectures. However, it does not appear clear which parameter is currently the most important. According to a recent survey conducted in the Netherlands, neurosurgeons use cages made in polyether-ether-ketone (PEEK) more often than titanium cages, whereas orthopedic surgeons prefer titanium cages. In our study, the majority of respondents were orthopedic spinal consultants (87.5%) and, overall, titanium cages were the most common (54.3%). Additionally, implants with smooth surface are generally discarded, whilst porous/trabecular or rough/threaded implants seem to be the favorite choice. Overall, our findings suggest uniformity of opinion concerning the relevance of implant geometry for the choice of the fusion implant (81.3%). Design factors such as cage width, length, height, and lordotic angle are all considered relevant. Parallel-shape cages are avoided in most cases (89.2%) and hyperlordotic cages are preferred at the lumbosacral level. Indeed, in order to correct sagittal alignment and balance, a 5° to 6° angle was proven to be sufficient at L4–L5 level and above, but not enough at L5–S1 level. Furthermore, our results show that some implant designs are more common than others but there is uncertainty regarding which design should be preferred. A specific fusion technique limits the width of the surgical access and, in turn, the possible sizes and shapes of the implant are restricted. Hence, the established variety of practice well reflects the diversity of opinion on implant design. Several clinical and biomechanical studies have been focused on evaluating the performance of different profiles for lumbar cages. However, clinical evidence is still inadequate and comparisons are limited to few different designs. Based on this survey, surgeons have shown to value a patient-oriented care, nevertheless, there is limited scientific literature evidencing uniform guidelines for some patient categories or specific diseases. In this context, patient-specific technology has the potential to alter future spinal surgical practice by providing perfectly fitting implants tailored on a given situation. This approach has shown great popularity in orthopedics over the past 5 years, however, its applicability to spinal fusion surgery has only recently been explored. The feasibility of designing and manufacturing of anatomical-shaped fusion cages using 3D printing technology has been shown. However, clinically relevant studies in this area still need to be explored. The current stage of the spinal implant market is not yet patient-specific, but future industrial and academic investments will be likely focus towards the customization of spinal implants.

The last part of the survey aimed to investigate the current practice as it pertains the possible bone grafting procedures. The results of our survey highlight that autograft surgeries were the most common bone grafting procedures in the United Kingdom, still overcoming the use of synthetic substitutes. Contrarily, a slight shift from autologous to substitute grafts was reported in the United States during the past 16 years. Between the nonautograft options, surgeons favor the use of synthetic substitutes, preferable HA-based composite materials (76.7%), Injectable paste form is preferred (80.8%), perhaps for the ease of insertion and possibility to fill small bone cavities.

Table 4

| Preferred bone graft materials (multiple choice allowed) | Number of responses (%) |
|----------------------------------------------------------|-------------------------|
| Category of bone graft materials (n=28)                  |                         |
| Autograft                                                | 18 (60.0)               |
| Allograft                                                | 1 (3.3)                 |
| Synthetic                                                | 10 (33.3)               |
| Other                                                    | 1 (3.3)                 |
| Nonautograft bone graft materials (n=27)                 |                         |
| HA (hydroxyapatite)                                     | 9 (30.0)                |
| Si-HA (silicon-hydroxyapatite)                           | 2 (6.7)                 |
| HA-CC (hydroxyapatite–calcium carbonate)                 | 11 (36.7)               |
| Coraline hydroxyapatite                                  | 1 (3.3)                 |
| DBX, DBM (demineralized bone matrix)                     | 6 (20.0)                |
| Other                                                    | 1 (3.3)                 |
| Form of bone graft materials (n=25)                      |                         |
| Paste                                                    | 21 (80.8)               |
| Granules                                                 | 3 (11.5)                |
| Cement                                                   | 0 (0)                   |
| Other                                                    | 2 (7.7)                 |

4.2. Fusion approach

Most surgeons reported performing most common bone grafting procedures in the United Kingdom, still overcoming the use of synthetic substitutes. Contrarily, a slight shift from autologous to substitute grafts was reported in the United States during the past 16 years. Between the nonautograft options, surgeons favor the use of synthetic substitutes, preferable HA-based composite materials (76.7%), Injectable paste form is preferred (80.8%), perhaps for the ease of insertion and possibility to fill small bone cavities.

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The pilot nature of this study includes indeed several limitations. First, this survey was conducted at a single event (i.e., BritSpine 2016, April 6–8, 2016, Nottingham, United Kingdom). This event, which is held every 2 years, attracts participants from the British Association of Spinal Surgeons (BASS) which counted 381 memberships at the end of 2016, and includes a large part of the population of spinal consultants in the United Kingdom. The results of our study indeed include a sample of the UK consultant spinal surgeons. Because of the relatively small size of the interviewed group, we were unable to perform any more accurate statistical analysis of subpopulations or correlations among responses. Hence, this may affect the scalability of the results of this research. However, a strength of this study was the fact that we approached only expert spinal surgeons with at least 1 year experience as consultants. Therefore, only surgeons who undertook their own surgical decisions were recruited. Because of the face-to-face recruitment method, we could maximize the response and completion rates of the questionnaire, however, this may have interfered with the in-depth thinking of participants, thus increase the response bias. In addition, no specific distinction between open versus minimally invasive fusion surgeries was made in this study. In the context of such rapidly evolving techniques, it will be useful to continue this investigation by repeating the survey at future meetings and updating the questions accordingly. Such longitudinal analysis will allow us to monitor the progresses in this field. Moreover, surgeons’ indications for specific diseases such as lumbar degenerative disc disease, trauma, and deformity have not been assessed in details. Future investigations should therefore focus on identifying subgroups of cases for whom a specific lumbar fusion approach is contemplated as an effective treatment. As it pertains to the factors influencing implant choice, the correlation between distinctive designs with specific fusion approaches has not been fully addressed in the questionnaire, thus limiting the interpretation of the results. Indeed, the geometry of the interbody fusion implant is a direct function of the surgical approach that it is used for. Finally, a more detailed cost analysis of surgical procedures and implants would also be relevant for the description of surgeons’ decision making process. Despite these limitations, we believe that this pilot study can provide a snapshot of current surgeons’ practice and, at the same time, encourages bigger scale surveys.

5. Conclusions

The present survey attempts at investigating the variability in surgeons’ decision making process in spinal fusion surgery. Our findings suggest the relevance of a patient-oriented management, which is confirmed by a wide variability of choices among surgeons’ practice. There is a lack of consensus among spinal surgeons concerning the choice of the fusion approach, fixation surgery, and interbody fusion implant. Various parameters in terms of implant geometry might lead surgeons’ practice; however, current clinical evidence is still inadequate to justify the superiority of one implant over another. Given the importance of a patient-centered care, research needs to examine the outcomes of lumbar fusion surgery for certain pathologies and specific populations of patients.

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