



Original

Applicability of consensus methods in hip arthroscopy: a systematic review

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ABSTRACT

Objective: consensus methods may represent a useful tool when the available evidence is scarce. The increase in the indications of hip arthroscopy makes it necessary to carry out studies with consensus methodology. The present study was carried out to identify the consensus methods used in the hip arthroscopy setting.

Material and methods: a systematic review was made consulting MEDLINE and PreMEDLINE (Ovid), Embase, Scopus, Web Of Science databases and healthcare technology evaluation resources up until September 2021. Inclusion was made of studies using the Delphi technique, the RAND/UCLA method, nominal group, consensus conferences or informal discussions, and which involved patients with hip disease of any kind subjected to hip arthroscopy. Two reviewers selected the studies on an independent basis. Relevant data were extracted by a reviewer and checked by a second reviewer. Quality assessment was performed on a paired basis and using a checklist.

Results: a total of 13 studies were selected, corresponding to 10 original articles, two healthcare technology evaluation reports and one appropriate use criteria document. Of the 13 studies, four used non-structured methods and 9 used structured methods such as the consensus conference, Delphi technique, nominal group and RAND/UCLA methodology. The studies involving formal consensus methods yielded higher methodological quality assessment scores.

Conclusions: the consensus methods evaluating the use of arthroscopy in hip disease were predominantly formal consensus methods. In most cases, the use of these structured methods

RESUMEN

Aplicabilidad de los métodos de consenso en la artroscopia de cadera: revisión sistemática

Objetivo: los métodos de consenso pueden convertirse en una herramienta útil cuando la evidencia disponible es escasa. El aumento de las indicaciones de la artroscopia de cadera hace necesario el uso de estudios con metodología de consenso. El objetivo de este estudio fue identificar los métodos de consenso empleados en el ámbito de la artroscopia de cadera.

Material y método: se realizó una revisión sistemática mediante consulta a las bases de datos MEDLINE y PreMEDLINE (Ovid), Embase, Scopus, Web Of Science y recursos de evaluación de tecnologías sanitarias, hasta septiembre de 2021. Se incluyeron estudios que emplearan la técnica Delphi, el método RAND/UCLA, el grupo nominal, las conferencias de consenso o discusiones informales, e implicaran a pacientes con cualquier patología de cadera sometidos a una artroscopia de cadera. Dos revisores llevaron a cabo la selección de los estudios de forma independiente. La extracción de los datos relevantes fue realizada por un revisor y comprobada por un segundo revisor. La evaluación de la calidad de realizó por pares y mediante una lista de comprobación.

Resultados: se seleccionaron 13 estudios, correspondientes a 10 artículos originales, 2 informes de evaluación de tecnologías sanitarias y 1 documento de criterios de uso apropiado. De los 13 estudios, 4 emplearon métodos no estructurados y 9 utilizaron métodos estructurados como la conferencia de consenso, la técnica Delphi, el grupo nominal y la metodología RAND/UCLA. Los estudios con métodos formales de consenso obtuvieron



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provided the criteria needed to establish consensus among the professionals.

Key words: Consensus. Arthroscopy. Hip. Systematic review.

Introduction

In some cases, when answers to very concrete questions are needed, or when addressing relatively novel issues, the scientific evidence may be lacking, or the existing evidence may be of suboptimal quality. In such cases it is necessary to resort to the opinion and experience of experts, adopting a systematic approach based on consensus methodology, in order to explore the level of agreement or disagreement on a given subject. Thus, the need for consensus arises from the lack of consensus⁽¹⁾.

Among the different consensus methods, informal consensus is based on open and non-systematised discussion that often takes place in the context of a single physical meeting. It affords recommendations and very little information on the basis sustaining the consensus. Informal consensus is obviously very susceptible to bias and group effects^(2,3), and since the anonymity of the participants is not guaranteed, some opinions are often not expressed or are diluted among those of other more reputed experts⁽⁴⁾. In contrast, formal consensus combines scientific evidence with methodological techniques and structured processes in the making of decisions. In the healthcare setting, use fundamentally has been made of three formal consensus methods⁽⁵⁾: the Delphi technique was introduced in the 1960s⁽⁶⁾, followed by the nominal group technique in the 1970s⁽⁷⁾ and, in 1977, by the consensus conference method developed by the United States National Institute of Health Consensus Development Program⁽⁸⁾.

A fourth strategy, the RAND/UCLA method, was developed in the 1980s by the RAND Corporation and the

puntuaciones más elevadas en la evaluación de la calidad metodológica.

Conclusiones: los métodos de consenso que evaluaron la utilización de la artroscopia en patología de cadera fueron mayoritariamente métodos formales de consenso. La utilización de estos métodos estructurados permitió conseguir, en su mayoría, los criterios para establecer el consenso entre los profesionales.

Palabras clave: Consenso. Artroscopia. Cadera. Revisión sistemática.

University of California at Los Angeles (UCLA), and constitutes a hybrid of the Delphi and nominal group methods⁽⁹⁾.

Table 1 highlights some of the differences between these approaches⁽⁵⁾.

Although there are variations in the consensus methods, they all follow highly formalised protocols and share a number of fundamental principles that distinguish them from informal consensus: anonymity, iteration, controlled feedback, group statistical response and structured interaction^(10,11).

Although these methods have recently been used in traumatology⁽¹²⁻¹⁵⁾, it would be advisable to specifically examine their use in the context of hip arthroscopy. Based on the use of these consensus techniques, it has been attempted to synthesise the collective opinions in a current state of uncertainty (differences in opinion on different aspects of the approach to hip arthroscopy). For this reason, we raised the hypothesis that there are quality publications related to consensus methods in hip arthroscopy which could guide clinical decisions in the event of the lack of an adequate body of evidence.

Table 1. Characteristics of the formal and informal consensus methods

Method	Submission of questionnaires	Private votes	Formal feedback of the group votes	In-person contact	Structured interaction	Summing method
Informal method						
Informal or non-structured	No	No	No	Yes	No	Implicit
Formal methods						
Delphi	Yes	Yes	Yes	No	Yes	Explicit
Nominal group technique	No	Yes	Yes	Yes	Yes	Explicit
Consensus conference	No	No	No	Yes	No	Implicit
RAND	Yes	Yes	Yes	Yes	Yes	Explicit

The general objective of the present study was to identify the consensus methods used in the hip arthroscopy setting.

Material and methods

A systematic review was made of the literature using formal methods to ensure a pertinent and precise search and retrieval process. The present study was carried out following the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) statement^(16,17). Thus, before starting the literature search and subsequent data extraction, a review protocol was developed, describing each step of the systematic review, including the exclusion criteria. This protocol was reviewed and approved by three of the authors.

Information sources and search strategy

Systematic searches were made in September 2021 (without time restrictions) in the following electronic databases: MEDLINE and PreMEDLINE (Ovid), Embase, Scopus and Web Of Science (WOS), combining terms such as *consensus*, *Delphi*, *RAND*, *nominal*, *hip* or *arthroscopy*. A manual review was made of the references of all the selected articles in order to locate other studies not appearing in the first search. We consulted the Centre for Reviews and Dissemination (CRD) and International Network of Agencies for Health Technology Assessment (INAHTA) databases. In addition, we consulted the website of the Spanish Network of Agencies for the Evaluation of Healthcare Technologies and Services of the National Healthcare System (*Red Española de Agencias de Evaluación de Tecnologías Sanitarias y Prestaciones del Sistema Nacional de Salud* [RedETS]). An example of the MEDLINE search is provided in **Figure 1**. The rest of the

literature searches are available upon request addressed to the corresponding author.

Selection of studies

On an independent and paired basis, two reviewers selected the studies by reading the titles and abstracts located through the scientific literature search. The full-text versions of the selected articles were reviewed and classified as included or excluded by the two reviewers, based on the established screening criteria. When doubts or discrepancies were found, they were resolved by consensus. The following study selection criteria were applied:

- Types of studies: consensuses using the Delphi technique, the RAND method, nominal group, consensus conferences or informal discussions.
- Participants: patients with hip disease of any kind.
- Intervention: any disease condition implying hip arthroscopy.
- Comparator: any.
- Outcome measures: data were extracted related to identification of the study, with the design and methodology. Communications at congresses, letters to the editor, editorials and comments were excluded. We also excluded studies not written in Spanish, English, French, Portuguese or Italian.

Data extraction

Data extraction from the included studies was carried out by a reviewer and checked by a second reviewer. Doubts or discrepancies were resolved by consensus. The data were entered on customised electronic sheets.

The data extraction procedure was carried out in two phases. The first phase compiled information referred to identification of the study, such as the year of conduction / publication of the study, country and organising entity. The second phase of the data extraction procedure included the consensus method used, the purpose of the study, the number of participants and the setting (national or international). In addition, we determined whether a literature review was made, whether the participants received prior information, whether the survey method was described (survey sent by conventional mail, e-mail or conducted on an in-person basis), the number of rounds and the number of participants that responded in each round, whether voting was private and whether anonymity was preserved, whether there was a predetermined definition of consensus and if so, what was the definition. Lastly, in the event of consensus, we evaluated whether it constituted forced consensus. The reviewers evaluated not only the presence or absence of each of these parameters in the studies but also whether they had been explicitly declared and described in sufficient detail.

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Database: Ovid MEDLINE(R) and Epub Ahead of Print, In-Process, In-Data-Review
& Other Non-Indexed Citations, Daily and Versions(R) <1946 to September 03, 2021>
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1 (consensus development conference or consensus development conference nih).pt.
2 consensus/
3 consensus development conferences as topic/ or consensus development
4 conferences, nih as topic/
5 delphi technique/
6 ((rand adj2 method) or (modified adj2 delphi) or (nominal adj2 group) or
7 (delphi or Rand or consens*)).ti.
8 appropriateness.ti.
9 1 or 2 or 3 or 4 or 5 or 6
10 Arthroscopy/
11 Arthroscopes/
12 arthroscop*.ti,ab.
13 8 or 9 or 10
14 Hip/
15 Hip Joint/ or Hip Injuries/
16 exp Pelvis/
17 (hip or acetabulofemoral or femoroacetabular or acetabul* or pelvi*).ti,ab,kw,sh.
18 12 or 13 or 14 or 15
19 11 and 16
20 7 and 17

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Figure 1. Example of MEDLINE literature search.

Quality assessment

The recommendations of Humphrey-Murto *et al.*⁽¹⁸⁾ were followed to guarantee methodological rigour in using the consensus group methods. These recommendations afford a checklist with the purpose of providing all the information that is essential for writing, interpreting and correctly using the results of a consensus. In general terms, the greater the compiled evidence in support of the development or choice of a method, the more reliable the findings derived from its use will be.

Data synthesis

A narrative synthesis was made, with tabulation of the information collected from the included studies.

Results

Figure 2 shows the study selection process. The search in the aforementioned electronic databases resulted in the identification of 313 literature references; this figure was reduced to 212 after removing duplicates. Following the selection process, we finally included 13 studies corresponding to 10 original papers⁽¹⁹⁻²⁸⁾, two healthcare technology evaluation reports^(29,30) and one appropriate use criteria document adopted by the steering committee of the American Academy of Orthopaedic Surgeons (AAOS)⁽³¹⁾.

Table 2 shows the studies discarded after full-text evaluation⁽³²⁻⁵⁶⁾ and the reasons for exclusion.

Of the 13 finally included studies, four used non-structured methods^(19,20,24,30), while 9 used structured methods such as consensus conference⁽²¹⁾, Delphi^(22,23,25,28), Delphi together with nominal group⁽²⁶⁾ and RAND/UCLA methodology^(27,29,31). The number of consulted specialists ranged between 9-869, and most of them were specialised in traumatology. The principal characteristics of the studies are described in Table 3.

Most of the studies addressed degenerative diseases such as osteoarthritis^(19,27,29,31) or femoroacetabular impingement^(21,24-26,28,29). Other addressed aspects were referred to hip dysplasia⁽²⁴⁾, infections^(22,23), thromboembolism⁽²⁰⁾ and clinical trial screening criteria⁽³⁰⁾.

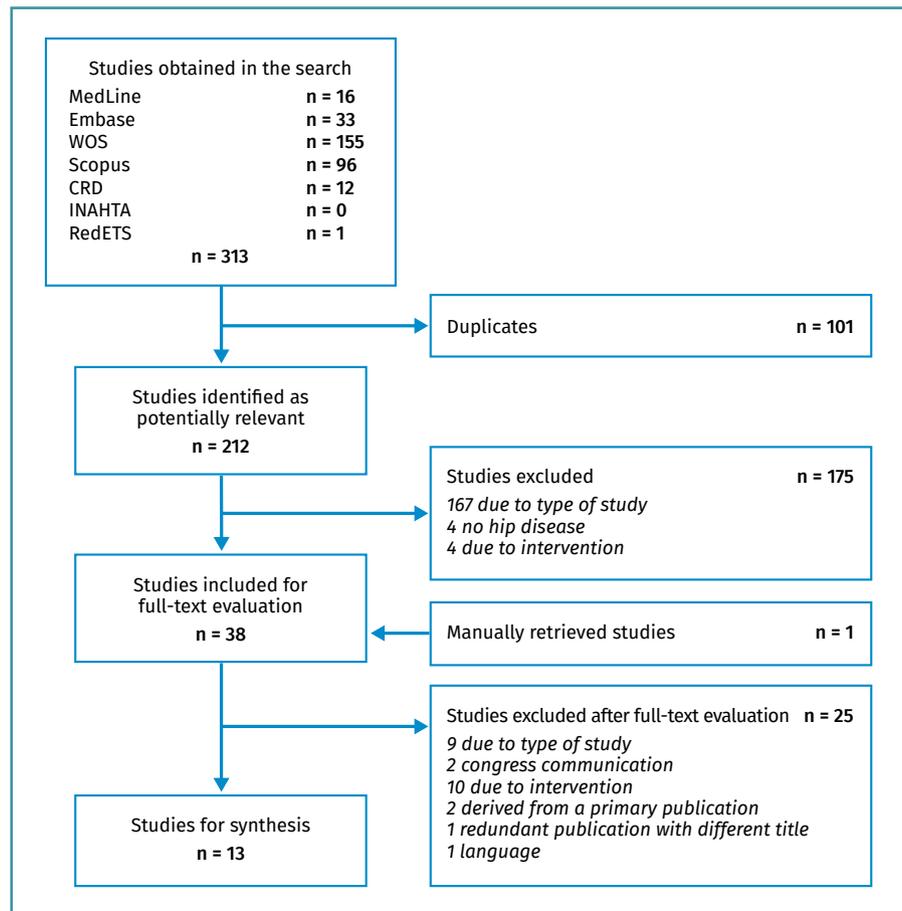


Figure 2. Flowchart of the study selection process.

The study published by Altman *et al.*⁽¹⁹⁾ was designed to describe the best available measurement method for detecting the progression of hip osteoarthritis, particularly in therapeutic trials. Consensus determined that radiography is an adequate primary assessment method for changes in hip osteoarthritis. The progression of osteoarthritis can be calculated by measuring the width of the joint space.

The report by Molina *et al.*⁽²⁹⁾ developed criteria for the appropriate or adequate use of hip arthroscopy in osteoarthritis and femoroacetabular impingement. Hip arthroscopy was considered generally inadequate as surgical treatment for osteoarthritis and adequate for femoroacetabular impingement, based on the presence of the following criteria: joint clinical manifestations, duration of the symptoms, functional alteration and patient age.

The AAOS consensus⁽³¹⁾, also contemplated in the study of Riddle *et al.*⁽²⁷⁾, included pharmacological and non-pharmacological aspects, and surgical procedures, for symptomatic hip osteoarthritis. The suitability of hip preservation surgery was based exclusively on patient age and the radiographic evaluation of hip osteoarthritis.

Table 2. Articles excluded after full-text evaluation

Article	Main reason for exclusion
Bick et al. (2003) ⁽³²⁾ Bounameaux et al. (2010) ⁽³³⁾ García et al. (2020) ⁽³⁴⁾ Griffin et al. (2012) ⁽³⁵⁾ Kemp et al. (2016) ⁽³⁶⁾ Lall et al. (2020) ⁽³⁷⁾ Maldonado et al. (2019) ⁽³⁸⁾ Pohlemann et al. (2007) ⁽³⁹⁾ Prisco et al. (2014) ⁽⁴⁰⁾	Type of study
Bick et al. (2019) ⁽⁴¹⁾ Rillo et al. (2015) ⁽⁴²⁾	Conference communication
Aalirezaie et al. (2019) ⁽⁴³⁾ Cardiovascular Disease Educational and Research Trust et al. (2006) ⁽⁴⁴⁾ Conrozier et al. (2020) ⁽⁴⁵⁾ McClincy et al. (2021) ⁽⁴⁶⁾ Misiolek et al. (2018) ⁽⁴⁷⁾ Mosler et al. (2020) ⁽⁴⁸⁾ Reiman et al. (2017) ⁽⁴⁹⁾ Reiman et al. (2020) ⁽⁵⁰⁾ Takla et al. (2020) ⁽⁵¹⁾ Zhang et al. (2008) ⁽⁵²⁾	Intervention different from that proposed in the inclusion criteria
Molina-Linde et al. (2018) ⁽⁵³⁾ Molina-Linde et al. (2018) ⁽⁵⁴⁾	Derived from a primary publication, Molina Linde (2017)
Della Rocca et al. (2013) ⁽⁵⁵⁾	Duplicate in the body of the article with Randelli (2013)
Hospach et al. (2018) ⁽⁵⁶⁾	Language

The study carried out by Griffin *et al.*⁽²¹⁾ aimed to reach multidisciplinary agreement with international experts on the diagnosis and treatment of femoroacetabular impingement. Consensus determined that in order to establish the diagnosis, the patients must present consistent symptoms, positive clinical signs and imaging findings. Adequate treatments in turn comprised conservative management, rehabilitation and arthroscopic or open surgery.

The consensus published by Marín-Peña *et al.*⁽²⁴⁾ addressed the indications of hip arthroscopy in degenerative disease of the hip and in hip dysplasia, even contributing surgical "tricks".

The study carried out by Radha *et al.*⁽²⁵⁾ addressed hip preservation in femoroacetabular impingement, in particular, intraoperative management of the capsule, the *labrum*, cartilage defects, the round ligament and bone impingement.

The purpose of the study by Lynch *et al.*⁽²⁶⁾ was to develop pre-, intra- and postoperative recommendations for femoroacetabular impingement through evidence-based consensus via a meta-analysis, a systematic review and a group of arthroscopists. Consensus was reached to the effect that hip arthroscopy should be the standard of care

for the surgical treatment of classical or arthroscopically accessible femoroacetabular impingement.

The main objective of the study carried out by Winiger *et al.*⁽²⁸⁾ was to identify the key variables for performing arthroscopic treatment in femoroacetabular impingement syndrome. Consensus emphasised that treatment of the *labrum* and correction of the cam-type deformity are the two key elements in hip arthroscopy for the management of femoroacetabular impingement syndrome.

The international consensus of orthopaedic infections addressed the prevention and reduction of risks in the study of Aalirezaie *et al.*⁽²³⁾ and the treatment and surgical techniques in the study of Abouljoud *et al.*⁽²²⁾. However, the information on the methodology of these studies appears in the editorial of the monograph dedicated to the international consensus on orthopaedic infections⁽⁵⁷⁾. The consensus indicated that there is no evidence that prior arthroscopy increases the risk of subsequent periprosthetic joint infections.

Randelli *et al.*⁽²⁰⁾ aimed to establish agreement upon recommendations for the management of thromboembolism in orthopaedic and trauma surgery. They reported that, in all patients requiring pharmacological antithrombotic preventive measures, it is advisable to evaluate both thrombotic risk and bleeding risk – identifying high risk patients and those who will need careful evaluation.

The healthcare technologies evaluation report published by Griffin *et al.*⁽³⁰⁾ developed screening criteria for randomised clinical trials. In a survey of clinicians, the latter expressed their reserves about being able to conduct a clinical trial in patients with femoroacetabular impingement, though they were in favour of being able to randomise.

Quality assessment

With regard to quality assessment, of the 13 included studies, those that made use of formal methods obtained better scores, in general terms. In this respect, three of them could be considered of high quality^(25,26,29), four of moderate quality^(21,27,28,31) and two of low quality^(22,23). In contrast, the four studies that used informal or non-structured methods were assessed as being of low quality^(19,20,24,30). In all but the studies of Abouljoud *et al.*⁽²²⁾ and Aalirezaie *et al.*⁽²³⁾, the purpose or objective of the investigation was clearly defined. With the exception of the studies by Radha *et al.*⁽²⁵⁾, Lynch *et al.*⁽²⁶⁾, Winiger *et al.*⁽²⁸⁾ and Molina *et al.*⁽²⁹⁾, none of the publications explained how anonymity was maintained. The details referred to the evaluation of the quality of the studies are found in **Table 4**.

Discussion

The credibility and usefulness of the results of a consensus are directly proportional to the rigour applied

Table 3. Characteristics of the studies included in the review

	Altman et al. ⁽⁵⁹⁾	Randelli et al. ⁽⁶⁰⁾	Griffin et al. ⁽⁶¹⁾	Griffin et al. ⁽⁶⁰⁾	Molina et al. ⁽⁵⁹⁾	Abouljoud et al. ⁽²²⁾ and Aal-irezaie et al. ⁽²²⁾	Marin-Peña et al. ⁽²⁴⁾	Radha et al. ⁽²⁵⁾	Lynch et al. ⁽²⁶⁾	AAOS ⁽¹⁾ and Riddle et al. ⁽²⁷⁾	Winiger et al. ⁽²⁸⁾
Country	Spain	Italy	United Kingdom	United Kingdom	Spain	United States	Spain	United Kingdom	United States	United States	United States
Panellist nationality (context)	International	National	International	International	National	International	International	International	National	National	National
Consensus method used	Discussion panel (non-structured)	Inter-societies consensus in plenary sessions (non-structured)	Consensus conference	Discussion panel (non-structured)	Delphi modified (RAND/UCLA)	Delphi	Debate among panellists	Delphi	Delphi and nominal group technique	Delphi modified (RAND/UCLA)	Delphi
Number of participants	ND	9 clinical experts plus 2 reviewers (not specified, compiled from author and affiliation)	22 clinical and academic experts plus 1 patient	16 clinical experts	11 clinical experts	869 potential delegates for participation	10 panellists	165 panellists	15 panellists	16 panellists	6 panellists
Specialty of the participants	Members of the Osteoarthritis Research Society International (OARS) and International Cartilage Repair Society (ICRS)	Traumatologists, family physicians, haematologists and anaesthetists (not specified, compiled from author and affiliation)	Specialists in sports medicine, physiotherapists, traumatology surgeons and radiologists	Surgeons specialised in femoroacetabular impingement	Traumatologists with different interventional activity	Traumatologists	Traumatologists (members of the Spanish Association of Traumatology)	Surgeons with an interest in hip preservation surgery	Surgeons dedicated to hip arthroscopy	Orthopaedic surgeons, physiotherapist, radiologist and rheumatologist	Hip arthroscopy surgeons
Described form of survey (in-person, e-mail, conventional mail)	In-person meeting	ND	In-person meeting	ND	E-mail plus in-person meeting	E-mail plus in-person meeting	In-person meeting	E-mail	Online and in-person vote during a meeting	Electronic system plus in-person meeting	E-mail
Forced consensus	ND	ND	ND	ND	No	No	ND	No	No	No	No
Private votes	ND	ND	ND	ND	Yes	Yes	ND	Yes	Yes	Yes	Yes

ND: not described

in preparing and conducting the consensus. Consensus methods inevitably must be performed with great methodological rigour and complying with a series of quality requirements. In this respect, we have seen that most of the studies that used a formal or structured consensus method presented greater methodological quality, while in contrast those methods based on informal consensus strategies lacked this high expected quality.

In simpler terms and taking into account that each formal consensus technique can exhibit numerous variants, the main characteristics of the four methods presented can be described as follows. In the Delphi method the participants are surveyed in different rounds. They receive a questionnaire, and individual and/or group feedback is provided on the scores between rounds, specifying their positions and the global positions of the group. Consensus is obtained by means of a mathematical procedure involving the simple summing of individual judgements and the elimination of extreme (outlier) positions. The participants never meet or interact directly^(5,10). The number of modifications implemented in the Delphi method has led to considerable confusion about its application^(58,59).

In the nominal group technique, the participants physically come together in a meeting directed by an experienced

moderator⁽⁶⁰⁾. In this meeting, and in an extremely formalised manner, they present their ideas, individually define their points of view, explain their differences, and individually vote each proposed solution^(5,10,61). As in the previous case, consensus is obtained by means of a mathematical procedure involving the simple summing of individual judgements.

Consensus conferences involve the evaluation of the available evidence referred to some diagnostic or therapeutic intervention before a jury composed of experts and non-experts that are required to issue a report with recommendations on the use of the intervention. The process simulates an oral hearing in court. During the session, the experts defend the conclusions drawn from the evidence and interact with the public invited to the conference. In 2013, the Office of Disease Prevention withdrew the consensus conferences programme⁽⁸⁾, though it is still conducted by other investigators.

Lastly, the RAND/UCLA method begins in a first phase or round with the submission of a questionnaire, while in a second round an in-person meeting is held to clarify or discuss the appropriate or inappropriate use of a medical or surgical procedure^(9,61).

The findings of this review show that most of the 13 studies synthesised in the tables of our results were car-

Table 4. Assessment of the methodological rigour of the included consensus studies

Recommendation	Altman et al. ⁽¹⁹⁾	Randelli et al. ⁽²⁰⁾	Griffin et al. ⁽²⁰⁾	Griffin et al. ⁽²⁰⁾	Molina et al. ⁽²⁹⁾	Abouljoud et al. ⁽²²⁾ and Aalirezaie et al. ⁽²³⁾	Marín-Peña et al. ⁽²⁴⁾	Radha et al. ⁽²⁵⁾	Lynch et al. ⁽²⁶⁾	AAOS ⁽²⁷⁾ and Riddle et al. ⁽²⁷⁾	Winiger et al. ⁽²⁸⁾
1. Study objective or purpose is defined	Yes	Yes	Yes	Yes	Yes	Partially	Yes	Yes	Yes	Yes	Yes
2. Each step of the process is summarised. If modifications were made, a well founded reason is provided for the choices made	Partially	No	Partially	Partially	Yes	Partially	No	Yes	Yes	Yes	Yes
3. Selection and preparation of the scientific evidence for the participants is described.	Partially	No	Yes	Partially	Yes	No	Partially	Partially	Yes	Yes	Partially
4. Description is given of how the items / variables were selected for inclusion in the initial questionnaire	No	No	Yes	Partially	Yes	No	No	Partially	Yes	Partially	Yes
5. Description is given of how the participants were selected	No	No	Partially	Partially	Yes	Yes	Partially	Yes	Yes	No	Yes
6. The planned number of rounds and/ or criteria for terminating the process is described	No	No	Yes	Partially	Yes	No	No	Yes	Yes	Yes	Yes
7. Clear description is given of how consensus was defined	No	No	Yes	No	Yes	No	No	Yes	Partially	Yes	Yes
8. Response percentage and the results are reported after each round	No	No	Yes	No	Yes	Yes	No	Yes	Yes	Yes	No
9. The type of comments provided after each round is described	Partially	No	Yes	No	Yes	Yes	No	Yes	No	Yes	No
10. Description is given of how anonymity was maintained	No	No	No	No	Yes	No	No	Yes	Yes	No	Yes
11. Possible methodological issues in the discussion are addressed	Partially	No	No	No	Yes	No	No	Yes	Yes	No	Partially

ried out in the United States (6 publications)^(22,23,26-28,31), with Spain^(19,24,29) and the United Kingdom^(21,25,30) being the European countries with the greatest scientific production (3 publications), followed by Italy⁽²⁰⁾ (1 publication). This circumstance could be attributed to the existence of a greater tradition in the use of consensus methods over the years, and since the 1940s, on the part of the United States Army and Air Force⁽⁶²⁾.

It should be noted that the setting in which the identified consensus were developed was predominantly at national level in the United States – comprising all the

identified publications^(26-28,31), except two of the same study with an international character^(22,23) – while in Europe we located three publications in the United Kingdom^(21,25,30) and two in Spain^(19,24) involving an international setting. Only one publication in Spain⁽²⁸⁾ and another in Italy were characterised by a national setting⁽²⁰⁾. This could be due to the interest in obtaining consensus applicable to a broader setting and not circumscribed to a single country (in the case of Europe).

Likewise, the identified publications included experts whose number ranged widely from 6 to 869 profession-

als. Most of them were traumatologists with experience in hip surgery, i.e. from a single discipline. In only four publications^(20,21,27,31) did the professionals participating in the consensus have different types of training (multidisciplinary).

Furthermore, most of the publications^(19,21-24,26,27,29,31) identified the holding of at least one in-person meeting, thereby reflecting the importance of discussion among experts in establishing useful conclusions and consensus.

The results of this systematic review also show that all the studies that employed informal or non-structured consensus methods^(19,20,24,30) in relation to the use of hip arthroscopy, and two of those that used the Delphi technique^(22,23), were carried out without clearly defining how consensus was agreed. Therefore, when the authors conclude that the results of the study reflect consensus-based opinion, it seems that the achievement of consensus was assumed as an integral part of the method used. Although consensus may be the expected result of applying a consensus method, we believe that it is necessary to better define the criteria for reaching such consensus and to document the degree of agreement along with the results obtained.

Even though most of the studies included in our systematic review had consensus as an objective, only some of them defined consensus with a specific criterion^(21,25,27-29,31). Furthermore, this criterion was the reason for termination of the process, normally on the basis of a definition established *a priori*^(21,25-29,31). However, we believe that an adequate approach would be to establish an *a priori* formal definition of the criteria used for consensus, instead of assuming the latter as an automatic outcome due to the intrinsic fact of making use of a consensus method. Furthermore, the investigators should also specify alternative criteria for termination of the process, including possibly a maximum number of contemplated rounds. If the studies are to be made in the course of a certain number of rounds, the authors should specify how the degree of agreement is going to be quantified at the end of the study.

To the best of our knowledge, there are no validated quality indicators for studies involving consensus methods. We therefore resorted to the recommendations of Humphrey-Murto *et al.*⁽¹⁸⁾. These indicators were selected on the basis of those which we believe would allow the study to have both internal and external validity. According to these indicators, the quality of the reviewed studies was generally moderate or high in the publications involving formal consensus. However, it is important to recognise that this scoring is based more on what is reported in the study than on the quality of the study as such. Therefore, we propose that these or other similar criteria should constitute a set of suggested elements to be included in all publications involving consensus meth-

odology. We consider that the applicability of these criteria in the publications would be useful for the diffusion of quality protocols for clinical practice.

These considerations acquire importance due to the fact that level V evidence (expert opinion) remains a necessary component in the methodological repertoire used to determine the response to a clinical question, particularly in situations characterised by the absence of high quality evidence (and by the difficulty of obtaining such evidence), and by clinical variability.

Medicine based on evidence classifies randomised clinical trials and meta-analyses as the highest-ranking evidence, while less relevance is attributed to expert opinion, which is classified as corresponding to the lowest category. Nevertheless, randomised clinical trials and meta-analyses have weaknesses and strengths (since no research method is perfect), and they cannot always be applied or used as a design to obtain investigational results, due to type of patients involved (such as frail individuals or children), or the type of intervention under study (e.g. surgeries) — since doing so would not be acceptable from the ethical perspective.

Indeed, "no study design is perfect, and contradictory findings may arise from all types of studies"⁽⁶³⁾. Having said this, the practical alternatives to studies where strong confidence has been placed on their results (randomised clinical trials, cohorts, observational studies, etc. with good designs) range from the current observational studies (since we are now in the era of big data in large health registries) to the traditional methods — including expert opinions (more feasible and accessible in some cases).

Whichever design is considered more appropriate for achieving the objective of our research, it must be accompanied by quality and methodological rigour in order to be able to rely upon and extrapolate the results with the lowest risk of biases or limitations. All scientific research is fundamentally dependent upon the use of adequate and rigorously detailed investigational methods — and studies based on expert opinions are no exception to this⁽⁶⁴⁾. It may be pertinent to present studies that use consensus methods in accordance with certain indicators similar to those of the CONSORT statement, as used for example in randomised controlled trials.

This systematic review of the literature offers an overview of the different consensus methods used in hip arthroscopy. However, the limitations of the present study are those inherent to the application of its methodology, including publication bias derived from the fact that many scientific studies are not ultimately published, or selection bias, which depends on the objectiveness of the inclusion and exclusion criteria used in the studies. We have minimised the risk of such biases by including several literature sources and working with broad criteria for the inclusion of studies.

Conclusions

The consensus methods analysed in this review and which evaluated the use of arthroscopy in hip disease were predominantly formal consensus protocols. In most cases, the use of these structured methods provided the criteria needed to establish consensus among the professionals.

Ethical responsibilities

Conflicts of interest. The authors state that they have no conflicts of interest.

Financial support. This study has received no financial support.

Protection of people and animals. The authors declare that this research has not involved human or animal experimentation.

Data confidentiality. The authors declare that the protocols of their centre referred to the publication of patient information have been followed, and that all the patients included in the study have received sufficient information and have given written informed consent to participation in the study.

Right to privacy and informed consent. The authors declare that no patient data appear in this article.

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