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Author/s:

Duffy, JMN;Adamson, GD;Benson, E;Bhattacharya, S;Bofill, M;Brian, K;Collura, B;Curtis, C;Evers, JLH;Farquharson, RG;Fincham, A;Franik, S;Giudice, LC;Glanville, E;Hickey, M;Horne, AW;Hull, ML;Johnson, NP;Jordan, V;Khalaf, Y;Knijnenburg, JML;Legro, RS;Lensen, S;MacKenzie, J;Mavrelos, D;Mol, BW;Morbeck, DE;Nagels, H;Ng, EHY;Niederberger, C;Otter, AS;Puscasiu, L;Rautakallio-Hokkanen, S;Sadler, L;Sarris, I;Showell, M;Stewart, J;Strandell, A;Strawbridge, C;Vail, A;van Wely, M;Vercoe, M;Vuong, NL;Wang, AY;Wang, R;Wilkinson, J;Wong, K;Wong, TY;Farquhar, CM;AlAhwany, H;Balaban, O;Barton, F;Beebeejaun, Y;Boivin, J;Bosteels, JJA;Calhaz-Jorge, C;D'Angelo, A;Dann, LF;De Jonge, CJ;du Mez, E;Ferriani, RA;Gerval, MO;Gingel, LJ;Greenblatt, EM;Hartshorne, G;Helliwell, C;Helliwell, C;Hughes, LJ;Jo, J;Jovanovic, J;Kiesel, L;Kietpeerakool, C;Kostova, E;Kucuk, T;Kumar, R;Lawrence, RL;Lee, N;Lindemann, KE;Loto, OM;Lutjen, PJ;MacKinven, M;Mascarenhas, M;McLaughlin, H;Mills, DJ;Mourad, SM;Nguyen, LK;Norman, RJ;Olic, M;Overfield, KL;Parker-Harris, M;Ramos, DG;Rendulic, A;Repping, S;Rizzo, R;Salacone, P;Saunders, CH;Sengupta, R;Sfontouris, IA;Silverman, NR;Torrance, HL

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Top 10 priorities for future infertility research: an international consensus development study^{† ‡}

J.M.N. Duffy^{1,2,*}, G.D. Adamson³, E. Benson⁴, S. Bhattacharya⁵, S. Bhattacharya⁵, M. Bofill⁶, K. Brian⁷, B. Collura⁸, C. Curtis⁹, J.L.H. Evers¹⁰, R.G. Farquharson¹¹, A. Fincham¹², S. Franik¹³, L.C. Giudice^{14,15}, E. Glanville¹⁶, M. Hickey¹⁷, A.W. Horne¹⁸, M.L. Hull¹⁹, N.P. Johnson¹⁹, V. Jordan⁶, Y. Khalaf²⁰, J.M.L. Knijnenburg²¹, R.S. Legro²², S. Lensen¹⁷, J. MacKenzie²³, D. Mavrelos²⁴, B.W. Mol²⁵, D.E. Morbeck^{6,26}, H. Nagels²⁷, E.H.Y. Ng^{28,29}, C. Niederberger³⁰, A.S. Otter³¹, L. Puscasiu^{32,33}, S. Rautakallio-Hokkanen¹², L. Sadler^{6,16}, I. Sarris¹, M. Showell²⁷, J. Stewart³⁴, A. Strandell³⁵, C. Strawbridge³⁶, A. Vail³⁷, M. van Wely³⁸, M. Vercoe²⁷, N.L. Vuong³⁹, A.Y. Wang⁴⁰, R. Wang²⁵, J. Wilkinson³⁷, K. Wong⁹, T.Y. Wong¹⁶, C.M. Farquhar^{6,27}, and the Priority Setting Partnership for Infertility[§]

¹King's Fertility, Fetal Medicine Research Institute, London, UK ²Institute for Women's Health, University College London, London, UK ³ARC Fertility, Cupertino, CA, USA ⁴Patient and Public Participation Group, Priority Setting Partnership for Infertility, University of Auckland, Auckland, New Zealand ⁵Institute of Applied Health Sciences, University of Aberdeen, Aberdeen, UK ⁶Department of Obstetrics and Gynaecology, University of Auckland, Auckland, New Zealand ⁷Women's Network, Royal College of Obstetricians and Gynecologists, London, UK ⁸Resolve: The National Infertility Association, VA, USA ⁹School of Psychology, University of Waikato, Hamilton, New Zealand ¹⁰Centre for Reproductive Medicine and Biology, University Medical Centre Maastricht, Maastricht, The Netherlands ¹¹Department of Obstetrics and Gynaecology, Liverpool Women's NHS Foundation Trust, Liverpool, UK ¹²Fertility Europe, Belgium ¹³Department of Obstetrics and Gynaecology, Münster University Hospital, Münster, Germany ¹⁴Center for Research, Innovation and Training in Reproduction and Infertility, Center for Reproductive Sciences, University of California, San Francisco, CA, USA ¹⁵International Federation of Fertility Societies, Mount Royal, NJ, USA ¹⁶Auckland District Health Board, Auckland, New Zealand ¹⁷Department of Obstetrics and Gynaecology, University of Melbourne, Victoria, Australia ¹⁸MRC Centre for Reproductive Health, University of Edinburgh, Edinburgh, UK ¹⁹Robinson Research Institute and Adelaide Medical School, University of Adelaide, Adelaide, Australia ²⁰Department of Women and Children's Health, Kings College London, London, UK ²¹Freya, Gorinchem, The Netherlands ²²Department of Obstetrics and Gynaecology, Penn State College of Medicine, PA, USA ²³Fertility Plus, Auckland, New Zealand ²⁴Reproductive Medicine Unit, University College Hospital, London, UK ²⁵Department of Obstetrics and Gynaecology, Monash University, Melbourne, Australia ²⁶Fertility Associates, Auckland, New Zealand ²⁷Cochrane Gynaecology and Fertility, University of Auckland, Auckland, New Zealand ²⁸Department of Obstetrics and Gynaecology, The University of Hong Kong, Hong Kong ²⁹Shenzhen Key Laboratory of Fertility Regulation, The University of Hong Kong-Shenzhen Hospital, China ³⁰Department of Urology, University of Illinois at Chicago College of Medicine, Chicago, IL, USA ³¹Osakidetza OSI, Bilbao, Spain ³²Pharmacy, Science, and Technology, University of Medicine, Targu Mures, Romania ³³Center for Reproductive Medicine, Amsterdam Reproduction and Development Institute, Amsterdam University Medical Centres, Amsterdam, The Netherlands ³⁴British Fertility Society, Middlesex, UK ³⁵Sahlgrenska Academy, Department of Obstetrics and Gynecology, University of Gothenburg, Sahlgrenska University Hospital, Göteborg, Sweden ³⁶Fertility Network UK, London, UK ³⁷Centre for Biostatistics, University of Manchester, Manchester Academic Health Science Centre, Manchester, UK ³⁸Center for Reproductive Medicine, Amsterdam Reproduction and Development Institute, Amsterdam University Medical Centres, Amsterdam, The Netherlands ³⁹Department of Obstetrics and Gynaecology, University of Medicine and Pharmacy at Ho Chi Minh City, Ho Chi Minh City, Vietnam ⁴⁰Australian Centre for Public and Population Health Research, Faculty of Health, University of Technology, Sydney, Australia

*Correspondence address. King's Fertility, Fetal Medicine Research Institute, 16-20 Windsor Walk, Denmark Hill, London SE5 8BB, UK. E-mail: james.duffy3@nhs.net, @jamesmnduffy

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[§]Members of the Priority Setting Partnership for Infertility are listed in the [Appendix](#).

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STUDY QUESTION: Can the priorities for future research in infertility be identified?

SUMMARY ANSWER: The top 10 research priorities for the four areas of male infertility, female and unexplained infertility, medically assisted reproduction and ethics, access and organization of care for people with fertility problems were identified.

WHAT IS KNOWN ALREADY: Many fundamental questions regarding the prevention, management and consequences of infertility remain unanswered. This is a barrier to improving the care received by those people with fertility problems.

STUDY DESIGN, SIZE, DURATION: Potential research questions were collated from an initial international survey, a systematic review of clinical practice guidelines and Cochrane systematic reviews. A rationalized list of confirmed research uncertainties was prioritized in an interim international survey. Prioritized research uncertainties were discussed during a consensus development meeting. Using a formal consensus development method, the modified nominal group technique, diverse stakeholders identified the top 10 research priorities for each of the categories male infertility, female and unexplained infertility, medically assisted reproduction and ethics, access and organization of care.

PARTICIPANTS/MATERIALS, SETTING, METHODS: Healthcare professionals, people with fertility problems and others (healthcare funders, healthcare providers, healthcare regulators, research funding bodies and researchers) were brought together in an open and transparent process using formal consensus methods advocated by the James Lind Alliance.

MAIN RESULTS AND THE ROLE OF CHANCE: The initial survey was completed by 388 participants from 40 countries, and 423 potential research questions were submitted. Fourteen clinical practice guidelines and 162 Cochrane systematic reviews identified a further 236 potential research questions. A rationalized list of 231 confirmed research uncertainties was entered into an interim prioritization survey completed by 317 respondents from 43 countries. The top 10 research priorities for each of the four categories male infertility, female and unexplained infertility (including age-related infertility, ovarian cysts, uterine cavity abnormalities and tubal factor infertility), medically assisted reproduction (including ovarian stimulation, IUI and IVF) and ethics, access and organization of care were identified during a consensus development meeting involving 41 participants from 11 countries. These research priorities were diverse and seek answers to questions regarding prevention, treatment and the longer-term impact of infertility. They highlight the importance of pursuing research which has often been overlooked, including addressing the emotional and psychological impact of infertility, improving access to fertility treatment, particularly in lower resource settings and securing appropriate regulation. Addressing these priorities will require diverse research methodologies, including laboratory-based science, qualitative and quantitative research and population science.

LIMITATIONS, REASONS FOR CAUTION: We used consensus development methods, which have inherent limitations, including the representativeness of the participant sample, methodological decisions informed by professional judgment and arbitrary consensus definitions.

WIDER IMPLICATIONS OF THE FINDINGS: We anticipate that identified research priorities, developed to specifically highlight the most pressing clinical needs as perceived by healthcare professionals, people with fertility problems and others, will help research funding organizations and researchers to develop their future research agenda.

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Introduction

The ultimate aim of infertility research is to improve clinical practice and optimize the chances of people with fertility problems achieving parenthood. For this to be possible, research needs to address questions that are pertinent to people with infertility, be conducted using appropriate methods, and be reported in a comprehensive, transparent and accessible manner (Duffy *et al.*, 2017). The first step in research production is to identify appropriate questions. Traditionally, research funding organizations and researchers have identified, refined and prioritized their own research agenda. It is unlikely that such prioritization has used formal consensus methods, engaged wider stakeholders, including people with fertility problems, and was independent of commercial interests. There has been modest improvement in some countries, including the Netherlands, the UK and the USA, which has emphasized the importance of including patients and the public in developing research priorities (Graham *et al.*, 2020).

Sir Iain Chalmers, founder of the Cochrane Collaboration, has advocated for research priorities to be jointly identified by healthcare professionals, patients and communities (Chalmers and Glasziou, 2009). He established the James Lind Alliance, which brings together healthcare professionals, patients and others, in priority setting partnerships. Using formal consensus methods, each priority setting partnership engages in an open and transparent process to identify and prioritize unanswered research questions, known as research uncertainties, in a particular area of health care (James Lind Alliance, 2018). The expectation is that prioritized research uncertainties will establish the future research agenda of funding organizations and researchers. As a result, it is hoped that the gap will close between what research is needed and what research is pursued (Wilkinson *et al.*, 2019a).

An international collaboration has brought healthcare professionals, people with fertility problems and others together within a Priority Setting Partnership for Infertility to develop future research priorities for male infertility, female and unexplained infertility, medically assisted reproduction and ethics, access and organization of care.

Materials and methods

An international multidisciplinary steering group, including healthcare professionals, people with fertility problems and researchers, was established to provide a diverse range of perspectives to inform key methodological decisions. The steering group was convened during the development of the study protocol, before the launch of the initial survey and interim prioritization survey, and before the consensus development meeting. A systematic review of registered, progressing and completed priority setting research settings was completed to assist with the planning and delivery of the study (Graham *et al.*, 2020).

Research uncertainties related to infertility associated with endometriosis, miscarriage and polycystic ovary syndrome were not considered because of other current or completed research prioritization initiatives (Home *et al.*, 2017; Prior *et al.*, 2017).

Research priorities were developed in a three-stage process using consensus methods advocated by the James Lind Alliance (2018). Potential research uncertainties were gathered through an online survey of healthcare professionals, people with fertility problems and others. Healthcare professionals, including embryologists, fertility specialists and gynecologists, were recruited through the British Fertility Society, Core Outcomes in Women's Health (CROWN) initiative, Cochrane Gynaecology and Fertility Group, Fertility and Sterility Forum, Reproductive Medicine Clinical Study Group and Royal College of Obstetricians and Gynecologists. People with fertility problems were recruited through Fertility Europe, an umbrella organization of more than 20 European patient organizations, including Fertility Network UK and Freya, Fertility New Zealand, RESOLVE: The National Infertility Association, and the Women's Voices Involvement Panel hosted by the Royal College of Obstetricians and Gynecologists. Other people could register to participate, including healthcare funders, healthcare regulators and researchers. Recruitment was supported by an active social media campaign. Potential participants received an explanatory video abstract, a plain-language summary and survey instructions. Before completing the survey, participants provided demographic details, including age, gender and geographical location, and information pertaining to their professional or personal experience of infertility. Participants were invited to suggest up to five research questions related to infertility that they considered unanswered.

After the survey had closed, the survey responses were examined in detail within an iterative process. Individual responses were reviewed by at least two members of the steering group. Responses were excluded if they included questions that did not fit the scope of the study, were not answerable by research, related to a specific person or situation or were ambiguous. Incomplete responses were also excluded. The remaining responses were formatted into appropriate research questions.

In addition, research recommendations were identified from a systematic review of clinical practice guidelines and Cochrane systematic reviews. Clinical practice guidelines relevant to infertility were identified by searching bibliographical databases, including Embase, International Guideline Library and MEDLINE, from 2007 to July 2017. Research recommendations were extracted verbatim from clinical practice guidelines. Using a data extraction tool available to the Cochrane Gynaecology and Fertility Group, research recommendations were extracted from individual Cochrane reviews evaluating potential fertility treatments. Research recommendations from clinical practice guidelines and Cochrane systematic reviews were reviewed by two members of the steering group and formatted into appropriate research questions. Differences in opinion were resolved by discussion with the steering group.

The long list of potential research questions was organized by allocating individual research questions in four categories: male infertility; female and unexplained infertility, including age-related infertility, ovarian cysts, uterine cavity abnormalities and tubal factor infertility; medically assisted reproduction including ovarian stimulation, IUI and IVF; and ethics, access and organization of care. These categories were identified in consultation with the steering group. Duplicate research questions were removed. Research questions were checked against the published research evidence, including clinical practice guidelines, Cochrane systematic reviews and randomized trials, and those questions considered to be already answered were removed.

The long list of confirmed research uncertainties was entered into an interim prioritization survey. Initial survey participants were invited to participate in the survey. In addition, healthcare professionals, people with fertility problems and others were recruited using the same methods as the initial survey. Before completing the survey, participants provided demographic details, including age, gender and geographical location, and information pertaining to their professional or personal experience of infertility. Participants were invited to select the research uncertainties they considered most important. After the survey had closed, questions were ranked based on the frequency they had been chosen by participants.

The top 15 research uncertainties in each category were discussed during a consensus development meeting (data are presented in the [Supplementary Table S1](#)). A formal consensus development method, the modified nominal group technique, was used to identify the top 10 research uncertainties for each category ([James Lind Alliance, 2018](#)). Healthcare professionals, people with fertility problems and others who had completed the initial or interim prioritization survey were invited to participate. The modified nominal group technique does not depend on statistical power. In consultation with the steering group, the aim was to recruit between 15 and 30 participants, as this number has yielded sufficient results and assured validity in other settings ([Murphy et al., 1998](#)).

Before the consensus development meeting, participants provided demographic details, including age, gender and geographical location, and information pertaining to their professional or personal experience of infertility. Following an introductory session, participants were assigned to one of two groups, each with a facilitator, to discuss the ranking of prioritized research uncertainties. The assignments were pre-specified to ensure a mixture of healthcare professionals, people with fertility problems and others. The groups were provided with a set of cards with an individual research uncertainty printed on each. Each participant was asked to contribute their opinions on the research uncertainties they felt most and least strongly about. Following this initial discussion, participants were invited to discuss the ordering of the research uncertainties. By the end of the session, the research uncertainties were placed in ranked order. The rankings from the two groups were aggregated into a single ranking order and presented to the entire group. Participants were invited to discuss the ordering of the research uncertainties. By the end of the discussion, the research uncertainties were placed in a final ranked order.

The National Research Ethics Service, UK, advised the study did not require formal review.

Results

The initial survey was completed by 179 healthcare professionals (46%), 153 people with fertility problems (39%) and 56 others (14%), from 40 countries ([Table 1](#)). Four hundred and twenty-three responses were submitted ([Fig. 1](#)). Following review, 136 responses (32%) were excluded. Clinical practice guidelines relevant to infertility were identified by searching bibliographical databases; the search strategy identified 3680 records. After excluding 731 duplicate records, 2949 titles and abstracts were screened. Thirty-two potentially relevant clinical practice guidelines were evaluated. Fourteen clinical practice guidelines met the inclusion criteria, including two guidelines related to infertility in general ([Loh et al., 2014](#); [National Institute for Health and Care Excellence, 2017](#)), five guidelines related to male infertility ([American Urological Association, 2010](#); [Jarvi et al., 2010](#); [Jungwirth et al., 2018](#)), five guidelines related to uterine anomalies ([Kroon et al., 2011](#); [American Association of Gynecologic Laparoscopists, 2012](#); [Carranza-Mamane et al., 2015](#); [Practice Committee of the American Society for Reproductive Medicine, 2016a, 2017](#)) and two guidelines related to medically assisted reproduction ([Practice Committee of the American Society for Reproductive Medicine, 2016b](#); [Penzias et al., 2017](#)). Thirteen research recommendations were extracted from the clinical practice guidelines. The Cochrane Gynaecology and Fertility Group provided research recommendations from 162 Cochrane systematic reviews. Two hundred and twenty-three potential research questions were extracted from these research recommendations. A long list of 533 potential research uncertainties was reviewed, 241 duplicate research uncertainties were removed and 51 research uncertainties which had been answered by research were also removed.

A rationalized list of 231 confirmed research uncertainties was developed, which included 34 research uncertainties related to male infertility, 48 research uncertainties related to female and unexplained infertility, 101 research uncertainties related to medically assisted reproduction and 48 research uncertainties related to ethics, access and organization of care. These confirmed research uncertainties were entered into an interim prioritization survey, which was completed by 143 healthcare professionals, 119 people with fertility problems and 55 others, from 43 countries.

Nineteen healthcare professionals, 14 people with personal experience of infertility and 8 others, from 11 countries, participated in the consensus development meeting. The modified nominal group technique was used to prioritize the top 10 research uncertainties for male infertility, female and unexplained infertility, medically assisted reproduction and ethics, access and organization of care. Fifteen highly prioritized research uncertainties for each category were discussed during the consensus development meeting ([Supplementary Table S1](#)). The 15 highly prioritized research uncertainties were initially discussed by two separate groups and at the end of the discussion, they ranked the research uncertainties. The first-round ranking is presented in [Supplementary Table S1](#). The rankings from the two groups were aggregated into a single ranking order and discussed by the entire group ([Supplementary Table S1](#)). Participants were encouraged to discuss and finalize the rank order of the research priorities. The top 10 research priorities are presented in [Fig. 2](#).

Table 1 Characteristics of the participants in a survey to identify the priorities for future infertility research.

	Survey 1 Initial survey n = 388	Survey 2 Interim prioritization n = 317	Consensus meeting Final prioritization n = 41
Stakeholder group, n			
People with fertility problems	153	119	14
Healthcare professionals	179	143	19
Embryologists	39	26	4
Fertility specialists	71	64	6
Gynecologists	44	28	6
Others	25	25	3
Researchers	28	28	7
Others	15	10	1
Prefer not to say	13	17	0
Gender, n			
Female	223	176	25
Male	129	119	16
Prefer not to say	36	22	0
Age (years), n			
Below 30	47	26	2
30–39	118	85	12
40–49	61	60	5
50–59	73	61	13
Over 60	42	29	5
Prefer not to say	47	56	4
Geographical location, n			
Africa	15	14	0
Asia	57	34	3
Australia and New Zealand	61	51	22
Europe	115	117	13
North America	82	54	3
South America	27	19	0
Prefer not to say	31	28	0

Discussion

The Priority Setting Partnership for infertility has brought together healthcare professionals, people with fertility problems and others to identify the top 10 research priorities for future infertility research. These research priorities are diverse and seek answers to questions regarding prevention, treatment and the longer-term impact, as well as wider contextual issues related to access and public health policy. They highlight the importance of pursuing research which has often been overlooked, including addressing the emotional and psychological impact of infertility, improving access to fertility treatment, particularly in lower resource settings, and securing appropriate regulation. Addressing these priorities will require diverse research methodologies, including laboratory-based science, qualitative and quantitative research and population science.

Strengths and limitations

The [James Lind Alliance \(2018\)](#) has published guidance to inform the design of research priority setting studies. This study has followed this guidance to ensure the research priorities were developed using a clear and transparent process using formal consensus development methods. The study design, development and delivery were also informed by a systematic review of research priority setting studies relevant to women’s health ([Graham et al., 2020](#)). With 388 respondents from 40 countries participating in the initial survey, 317 respondents from 43 countries participating in the interim prioritization survey, and 41 participants from 11 countries included in the consensus development meeting, the global participation achieved in this study should secure the generalizability of the results within an international context. The study included people with fertility problems and they were able to suggest potential research uncertainties during the initial survey, share their views regarding the importance of research uncertainties

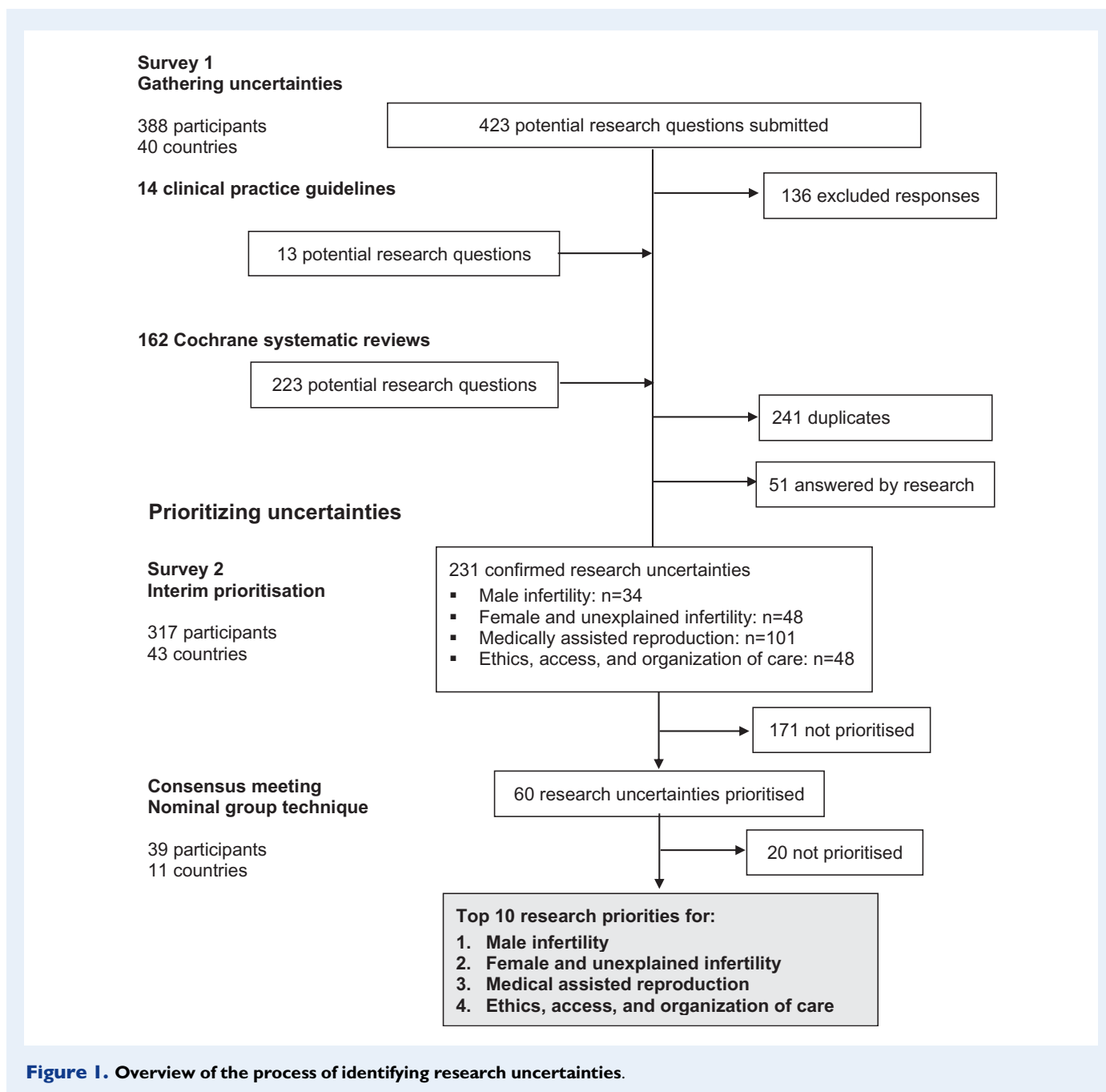


Figure 1. Overview of the process of identifying research uncertainties.

during the interim prioritization survey and participate fully in the consensus development meeting which prioritized the final research priorities.

This consensus study is not without limitations. Consideration should be given to the representativeness of the study's participants. For example, when considering the initial survey, there was a higher response from participants who identified as living in Europe (115 participants; 30%). To participate in the initial survey and interim prioritization survey, English proficiency and literacy, a computer and internet access were required. We appreciate that limitations in the representativeness of the sample could impact upon the

research uncertainties suggested and prioritized. There is uncertainty regarding the optimal consensus development method to prioritize research uncertainties, and methodological research is required to evaluate different approaches to priority setting and the use of different consensus methods. Further contextual information, including the number of people the research priority impacts upon, the feasibility of answering the research priority, and the resources required to address the research uncertainty could have assisted participants to prioritize research uncertainties. Future methodological research should evaluate the use of contextual information in research priority studies.

Top 10 research priorities for male infertility

1. Are sperm tests other than bulk parameters useful in evaluating male fertility? If so, which?
2. What is the emotional and psychological impact of male infertility? Can addressing it improve outcomes?
3. Do environmental factors cause male infertility? If so, which?
4. Does treating specific causes of male infertility improve outcomes?
5. Can we improve surgical sperm extraction outcomes by using endocrine stimulation protocols?
6. What modifiable risk factors cause male infertility?
7. Does treating modifiable risk factors improve outcomes?
8. What co-morbidities are associated with infertility?
9. Does treating co-morbidities improve outcomes?
10. Are nutraceuticals useful in improving male reproductive potential? If so, which?

Top 10 research priorities for female and unexplained infertility

1. Can age-related infertility be prevented?
2. Can a predictive model be developed, tested, and validated to compare the outcomes of different management strategies for couples with unexplained infertility?
3. In couples with unexplained infertility, what is the optimal ART?
4. Can a predictive model for fertility based upon ovarian reserve tests be developed, tested, and validated?
5. In women at risk of age-related infertility does standardized fertility assessment before attempting expectant management improve live birth rates?
6. What causes unexplained infertility?
7. In women with uterine fibroids what is the optimal management strategy to preserve fertility?
8. In women with otherwise unexplained infertility does hysteroscopic removal of an endometrial polyp increase live birth rates?
9. In women with mild intrauterine adhesions and otherwise unexplained infertility, does removal increase live birth rates?
10. In women with a uterine septum and otherwise unexplained infertility does hysteroscopic resection increase live birth rates?

Top 10 research priorities for medically assisted reproduction

1. What are the causes of implantation failure?
2. What is the optimal treatment for women who are poor responders undergoing IVF to increase live birth rates?
3. What is the optimal method of sperm selection in IVF cycles?
4. In couples with unexplained infertility does IUI increase live birth rates when compared with other ARTs, including IVF?
5. In couples with unexplained infertility what is the optimal number of IUI cycles before moving to IVF?
6. What is the optimal method of embryo selection in IVF cycles?
7. What are the factors which affect cycle to cycle variability in the number and quality of oocytes produced in an IVF cycle?
8. What is the optimal time interval between ovulation and IUI?
9. What is the emotional and psychological impact on children born using donor gametes?
10. What is the emotional and psychological impact of repeated fertility treatment failure?

Top 10 research priorities for ethics, access, and organization of care

1. Which public health interventions are effective in preventing infertility?
2. How can the cost of infertility treatment be reduced?
3. How can infertility treatment be made available in lower resource settings?
4. How should the information needs of people with infertility be met?
5. What age limit should be applied to women and men seeking infertility treatment?
6. What is the economic burden of infertility?
7. What is the minimum standard of care people with infertility should expect?
8. How should financial conflicts of interest be managed in clinical and research settings?
9. How should social egg freezing be regulated?
10. What are the optimal methods to report long term maternal and offspring outcomes across national and international settings?

Figure 2. The top 10 priorities for future infertility research in each of the four categories.

Reflections on the research priorities

Reproductive medical care for men has lagged behind that for women. Setting impactful and tractable priorities for male reproduction is consequently a critically important task. For diagnosis, the variation in morphology is extraordinary and counting sperm is challenging, severely limiting our ability to make predictions of male reproductive potential from the standard semen analysis, and begging the question: are there other, better tests of sperm? We need to explore how overall health affects male fertility and whether treating other diseases improves it. Because a man does not live in a vacuum, we need to understand how the environment affects male reproduction. When considering the treatment of male infertility, men often ask what they can do to improve their fertility, and well-conducted studies into diet and nutraceuticals are essential. The endocrine system drives the making of sperm and further evidence is required to understand if hormonal therapy could improve the production of sperm and improve live birth rates.

The priorities for unexplained infertility seek answers to several challenging and long-standing questions, including the prevention of age-related infertility and exploring the role of fibroids, polyps, intrauterine adhesions and uterine septa in unexplained infertility. It is also surprising that it remains unclear what the first-line treatment is for couples with unexplained infertility, IVF or IUI, and the timing of the superior treatment for that couple.

When considering medically assisted reproduction, new large prospective cohorts that consider all variables and use advance

methodology will be required to address casual relationships related to implantation failure. Similar complexity will exist when studying oocyte yield and quality over subsequent IVF cycles, even though similar stimulation protocols have been used. The three research priorities concerning the effectiveness of IVF are seeking to identify optimal ovarian stimulation protocols in poor responders, sperm selection techniques and embryo selection. These contrast with the research priorities which explore if, when, and how IUI should be used. To answer these effectiveness questions, well-designed randomized controlled trials will be required (Wilkinson *et al.*, 2019b). The psychological impact of fertility treatment is brought into sharper focus with research priorities related to the emotional and psychological impact of repeated fertility treatment failure and in children following gamete donation. Strong involvement of patient representatives, psychologists and behavioral scientists will be required to establish the appropriate qualitative and quantitative studies to address these important priorities.

The research priorities for ethics, access and organization of care broadly fall into two overarching themes: access and infertility as a public health issue. When considering access, cost is a major barrier to appropriate care, which is reflected in the research priorities aiming to explore interventions to reduce the cost of fertility treatment and increase the availability of fertility treatment in lower-resources settings. Turning to infertility as a public health issue, prevention of infertility should be a key priority for public health initiatives. We need to

determine the minimum standard of care that people with fertility problems should expect, especially if we are seeking reimbursements for this care.

Wider context

A prioritized list of research uncertainties, developed to specifically highlight the most pressing clinical needs as perceived by healthcare professionals, people with fertility problems and others, should help funding organizations and researchers to set their future research agenda. The selected list of research uncertainties should serve to focus a discussion regarding the allocation of limited resources.

Many of the research priorities will require national and international collaboration. Several countries, including China, the Netherlands, the UK and the USA, have developed national networks to undertake infertility research (Devall et al., 2020). Further development of national infrastructure is required. Collaboration should spread beyond national boundaries and develop within an international context. It is hoped the development of a prioritized research agenda could be an important enabler to deepen international collaboration. Development of generic infrastructure could help foster collaboration, including the use of minimum data sets, known as core outcome sets, low-cost data repositories and standardized approaches to the reporting of research. A core outcome set has recently been developed for future infertility trials (Duffy et al., 2018). Over 400 healthcare professionals, researchers and patients, from 40 countries, have used formal consensus development methods to identify a core outcome set for infertility (Duffy et al., 2020a). Consensus definitions have also been agreed for individual core outcomes (Duffy et al., 2020b). It is hoped the core outcome set will provide generic tools to collect outcomes during research, provide concise guidance regarding statistical analysis and standardize the approach to research reporting (Duffy et al., 2019).

Research priorities identified in this study correspond with research priorities identified by the Priority Setting Partnership for Miscarriage, including determining the emotional and psychological impact of miscarriage, investigating the modifiable risk factors which cause miscarriage and identifying specific comorbidities which cause miscarriage (Prior et al., 2017). Other similarities exist when considering the research uncertainties prioritized by the Priority Setting Partnership for Endometriosis and International Polycystic Ovary Syndrome Network (Home et al., 2017).

Answering the prioritized research questions would represent a significant step forward for our specialty. The steering group recognizes the important role of research which stems from the intellectual curiosity of individuals, fundamental research which does not have an immediate clinical application and research which is funded by special interest groups raising funding for the topic of their particular interest. A blended research strategy should offer the optimal pathway to improving clinical care and patient outcomes.

Perhaps the most important part of this process has been the strengthening of relationships between partner organizations, healthcare professionals and people with lived experience of infertility. The prioritized list of uncertainties that require research should help funding organizations and researchers to set their future research agenda. Our approach should ensure that future research has the necessary reach and relevance to inform clinical practice and to improve patient outcomes.

Supplementary data

Supplementary data are available at *Human Reproduction* online.

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Authors' roles

Study concept and design: J.M.N.D., S.B., B.C., C.C., J.L.H.E., R.G.F., S.F., L.C.G., A.W.H., N.P.J., Y.K., J.M.L.K., R.S.L., S.L., B.W.M., H.N., E.H.Y.N., C.N., A.S.O., L.P., S.R.H., M.S., J.S., A.S., C.S., A.V., M.v.W., M.A.V., N.L.V., A.Y.W., R.W., J.W. and C.M.F. Acquisition of data: J.M.N.D., S.B., K.B., C.B., C.C., J.L.H.E., R.G.F., A.F., S.F., L.C.G., A.W.H., N.P.J., Y.K., J.M.L.K., R.S.L., S.L., B.W.M., E.H.Y.N., C.N., A.S.O., L.P., S.R.H., M.S., J.S., A.S., C.S., A.V., M.v.W., M.A.V., N.L.V., A.Y.W., R.W., J.W. and C.M.F. Analysis and interpretation of data: J.M.D., G.D.A., E.B., S.B., S.B., M.B., K.B., B.C., C.C., J.L.H.E., R.G.F., A.F., S.F., L.C.G., E.G., M.H., A.W.H., M.L.H., N.P.J., V.J., Y.K., J.M.L.K., R.S.L., S.L., J.M., D.M., B.W.M., D.E.M., H.N., E.H.Y.N., C.N., A.S.O., L.P., S.R.H., L.S., I.S., M.S., J.S., A.S., C.S., A.V., M.v.W., M.V., N.L.V., A.Y.W., R.W., J.W., K.W., T.W. and C.M.F. Drafting of the manuscript: J.M.D., B.C., S.L., H.N., C.N., M.S., M.v.W., M.V., R.W., J.W. and C.M.F. Critical revision of the manuscript for important intellectual content: G.D.A., E.B., S.B., S.B., M.B., B.K., C.C., J.L.H.E., R.G.F., A.F., S.F., L.C.G., E.G., M.H., A.W.H., M.L.H., N.P.J., V.J., Y.K., J.M.L.K., R.S.L., J.M., M.M., D.M., B.W.M., D.M., E.H.Y.N., A.S.O., L.P., S.R.H., L.S., I.S., J.S., A.S., C.S., A.V., N.L.V., A.Y.W., K.W. and T.W. Statistical analysis: J.M.D., J.W. and A.V. Study supervision: C.M.F.

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Conflict of interest

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Appendix

Priority Setting Partnership for Infertility

Dr Hisham AlAhwany, University of Nottingham, UK; Ofra Balaban, CHEN: Patient Fertility Association, Israel; Faith Barton, UK; Dr Yusuf Beebeejaun, King's Fertility, Fetal Medicine Research Institute, UK; Professor Jacky Boivin, Cardiff University, UK; Professor Jan J. A. Bosteels, Imelda Hospital, Belgium; Professor Carlos Calhaz-Jorge, Faculdade de Medicina da Universidade de Lisboa, Portugal; Dr Arianna D'Angelo, Wales Fertility Institute, UK; Dr Leona F. Dann, Health Quality and Safety Commission, New Zealand; Professor Christopher J. De Jonge, University of Minnesota Medical Center, United States; Elyce du Mez, University of Auckland, New Zealand; Professor Rui A. Ferriani, University of Sao Paulo, Brazil; Dr Marie-Odile Gerval, Chelsea and Westminster Hospital NHS Foundation Trust, UK; Lynda J. Gingel, UK; Dr Ellen M. Greenblatt, Mount Sinai Fertility, University of Toronto, Toronto; Professor Geraldine Hartshorne, University of Warwick, UK; Charlie Helliwell, New Zealand; Charlotte Helliwell, New Zealand; Lynda J. Hughes, The Fertility Clinic, London Health Sciences Centre, Canada; Dr Junyoung

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