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Top 10 research priorities for people with obstructive sleep apnoea, families and clinicians: James Lind Alliance Priority Setting Partnership

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ABSTRACT

Obstructive sleep apnoea (OSA) is associated with significant comorbidity, preventable accidents and reduced quality of life. Little is known about the research priorities of patients with OSA, family members and clinicians. A James Lind Alliance research priority setting partnership was conducted. An initial survey (690 respondents who generated 1110 questions), a prioritisation survey (250 respondents), and a final workshop were used to identify the top 10 research priorities. Consensus was achieved on the top-ranked research priorities. Our results will inform the efforts of funders,

researchers and policy-makers to align directly with stakeholder priorities related to OSA.

Worldwide, obstructive sleep apnoea (OSA) is estimated to affect nearly 1 billion people aged 30–69 years, with most cases remaining undiagnosed and untreated.¹ Rising rates of obesity, chronic disease and population ageing increase the likelihood that OSA prevalence will continue to increase over time.²

It has been well recognised that patients make important decisions about how to best self-manage chronic conditions such as OSA

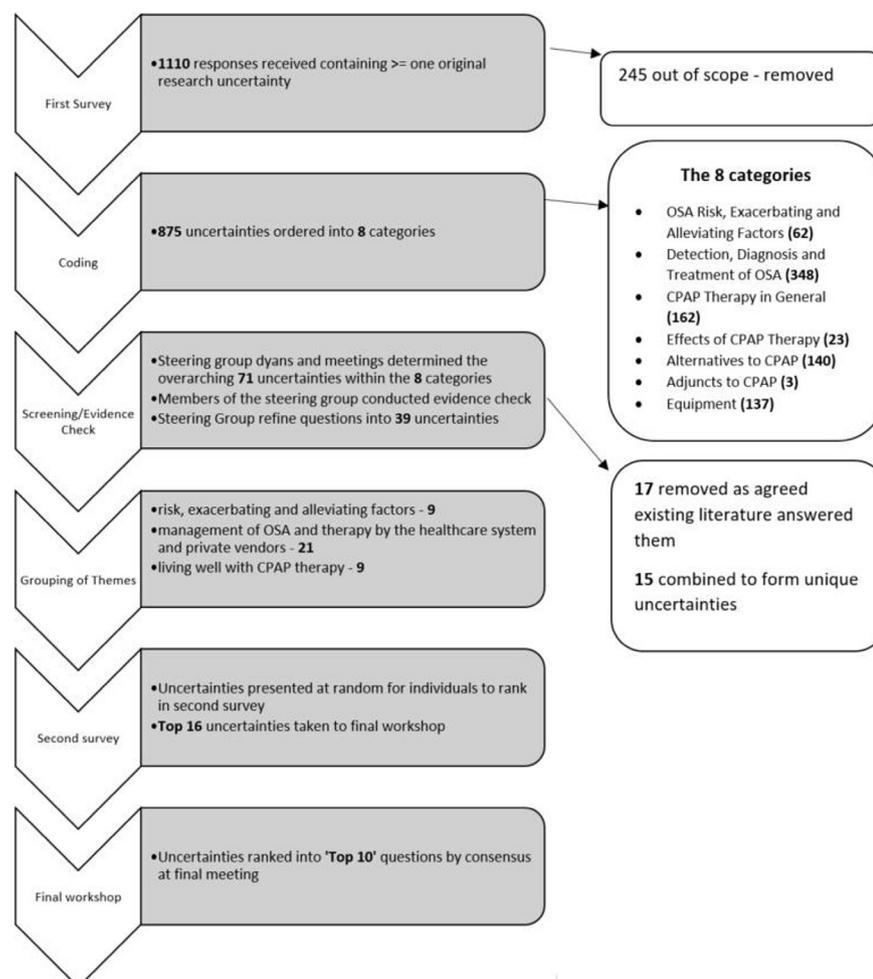


Figure 1 James Lind Alliance process overview. CPAP, continuous positive airway pressure; OSA, obstructive sleep apnoea.

Table 1 Characteristics of patient and family member respondents

Characteristic (n, %)	Identifying priorities questionnaire (N=630)	Prioritisation questionnaire (N=218)
Gender		
Male	271 (43.0)	86 (39.4)
Female	353 (56.0)	104 (47.7)
Non-binary	2 (0.3)	1 (0.5)
No response/prefer not to answer	7 (1.1)	27 (12.4)
Age		
18–39	147 (23.3)	20 (10.5)
40–59	269 (42.5)	90 (47.1)
60–69	141 (22.3)	58 (30.4)
70–79	63 (10.0)	19 (9.9)
80–89	10 (1.6)	4 (2.1)
Missing	3 (0.5)	27 (12.4)
Ethnicity		
Caucasian	548 (86.6)	176 (80.7)
Metis	24 (3.8)	10 (4.6)
South-East Asian	18 (2.8)	1 (0.5)
First Nations	16 (2.5)	3 (1.4)
Middle Eastern	4 (0.6)	1 (0.5)
Black	2 (0.3)	0
Other	17 (2.7)	
Missing	4 (0.6)	
Respondent category		
Diagnosed with OSA	466 (73.9)	196 (89.9)
On waiting list for assessment	7 (1.1)	
Suspects OSA but has not sought evaluation	27 (4.3)	
Family member of patient diagnosed with OSA	94 (14.8)	22 (10.1)
Family member of person with suspected OSA	36 (5.7)	
Agricultural producers	45 (7.2)	13 (6.0)
Ever prescribed treatment of OSA		
Yes	566 (89.4)	NA
No	62 (9.8)	
Missing	5 (0.8)	
Currently using CPAP		
Yes, 4 or more nights per week	420 (66.4)	NA
Yes, 1–3 nights per week	16 (2.5)	
Yes, intermittently	52 (8.2)	
No	141 (22.3)	
Missing	4 (0.6)	
Duration of CPAP use		
Under 1 year	49 (7.7)	NA
1–5 years	260 (41.1)	
6–10 years	93 (14.7)	
>10 years	85 (13.4)	
Missing/NA	146 (23.1)	
Reasons for going without treatment		
Unsure where to go for assessment	73 (11.5)	NA

Continued

Table 1 Continued

Characteristic (n, %)	Identifying priorities questionnaire (N=630)	Prioritisation questionnaire (N=218)
Cost of equipment	195 (30.8)	
Travel distance to assessment	42 (6.6)	
On waiting list	47 (4.7)	
Other reasons (free-text responses)	340 (53.7)	

CPAP, continuous positive airway pressure; NA, not available; OSA, obstructive sleep apnoea.

and key risk factors such as obesity.³ For example, while effective continuous positive airway pressure (CPAP) use has been associated with significant improvements in many of the sequelae of OSA,⁴ up to 30% of patients diagnosed with OSA fail to accept CPAP therapy,⁵ and between 29% and 83% are estimated to be non-adherent.⁶ Understanding the unanswered questions of those with lived experience of OSA can make a significant contribution to the relevance of future research in this area.

Patient-oriented research, also referred to as coproduction of research,⁷ contributes value by engaging patients, their caregivers and families as partners in the research process in order to focus on patient-identified priorities and lead to improved patient outcomes. The James Lind Alliance (JLA) Priority Setting Partnership (PSP) brings patients, caregivers and clinicians together to identify their unanswered questions. We conducted a JLA PSP to identify the key diagnostic and treatment uncertainties related to OSA and produce the top 10 research questions in these areas.

METHODS

Our PSP took place between October 2019 and January 2021. Guided by a JLA facilitator and following the prescribed methodology,⁸ we adapted the process in several ways. First, in order to engage populations such as older, rural and indigenous persons, as well as those with limited access to technology, we augmented the first online elicitation survey with in-person data collection (prior to COVID-19). Second, the final workshop was conducted virtually due to COVID-19 restrictions. The Reporting guideline for PRiority Setting of health research guidelines⁹ ensured comprehensive and transparent reporting of this PSP.

Our steering group included: six patient partners; a Knowledge Keeper working with Indigenous Services at the Saskatchewan Health Authority; two sleep specialist physicians; a nurse who facilitated an OSA support group; a nurse manager from the local sleep lab; a nurse specialising in rural health; a patient engagement specialist from the Saskatchewan Centre from Patient-Oriented Research; a nurse researcher and a research coordinator.

The initial Elicitation Survey was conducted between November 2019 and March 2020 in both online and paper formats. The survey was widely advertised through posters, broadcast media interviews and social media. In addition to demographic questions, the survey asked ‘What questions on Obstructive Sleep Apnoea would you like to see answered by research?’ Respondents submitted up to five free-text questions.

Two team members (BD and DG) independently reviewed all questions submitted, consolidating similar questions. Steering group members reviewed the consolidated questions to determine whether they adequately reflected the submitted

questions and revised the items. Questions unrelated to diagnosis or treatment of OSA were removed. Evidence checking consisted of examining systematic reviews and clinical guidelines to determine whether the consolidated questions had been fully, partially or not answered by existing research. Questions considered fully addressed were eliminated from the pool of items for the prioritisation survey, but formed the basis for several knowledge translation products for OSA patients aimed at improving knowledge, including three animated videos. The revised questions were reviewed by the JLA facilitator (KC) for quality assurance.

An online only (due to COVID-19 restrictions) prioritisation survey (September–December 2020) of the 39 retained questions allowed patients, family members and clinicians to rank their Top 10 questions. The highest-ranked questions chosen for each group respectively in the prioritisation survey were then discussed in an online workshop (due to COVID-19 restrictions) held in January 2021. Led by three independent JLA facilitators, patient partners and professional participation selected the final top 10 research priorities through small group discussion and plenary voting.

Figure 1 provides an overview of the JLA process for this PSP. The Elicitation survey was completed by 663 members of the public and 57 clinicians, generating a total of 1110 questions. The remaining 875 questions were combined into 71 questions, which were categorised into three themes: (1) risk, exacerbating and alleviating factors; (2) management of OSA and therapy by the healthcare system and private vendors; and (3) living well with CPAP therapy.

Table 2 Final workshop rankings of questions

Rank	Question
1	How can a diagnosis of Obstructive Sleep Apnoea (OSA) be made easier to obtain?
2	How could improved affordability and insurance coverage for OSA diagnosis and treatment affect patient outcomes?
3	How can access, coordination, and quality of public services be improved for persons with OSA?
4	What are important considerations when treating a patient with OSA who has other chronic health conditions?
5	How can OSA services to rural areas be improved?
6	How can OSA services to indigenous peoples be improved?
7	Are there other therapies that could be used along with continuous positive airway pressure (CPAP) to improve OSA and under what circumstances?
8	How can the treatment and care of OSA in long-term and residential care settings be improved?
9	How do lifestyle changes (such as weight loss, exercise and stress reduction) affect the need for ongoing CPAP treatment?
10	How often should CPAP therapy be re-evaluated by a healthcare provider?
11	What should be done if the patient does not notice any difference after beginning/receiving CPAP treatment?
12	What can be done to improve sleep quality for people with OSA?
13	Can OSA be permanently eliminated?
14	Can CPAP therapy ever be stopped and under what circumstances?
15	Apart from CPAP, what other ways (including alternative therapies and physical or breathing exercises) could effectively treat OSA?
16	How do bed surfaces, pillows and sleeping position affect the symptoms of OSA?

Demographics of patient and family members respondents to both surveys are shown in table 1.

Priorities were identified for patient respondents, for family member respondents and for provider respondents. Within each of these three groups, questions were given points based on the number of times they were selected, resulting in a ranked order for the questions for each respondent group. A shortlist was created by taking the top 10 questions from each group's ranked order. With overlap, where there was consensus between the groups, this resulted in a shortlist of 16 questions. The JLA recommended that this was a manageable number to navigate in an online discussion. The 16 top-ranked questions were short-listed for the final workshop which included nine patients and seven clinicians. The final top 10 questions are listed in table 2.

This JLA PSP in OSA engaged patients with OSA, their caregivers and healthcare professionals who treat those with OSA to develop a Top 10 list of patient-oriented research questions. Given the integral role patients with OSA play in self-managing their conditions and the limited resources available to conduct research, the priorities identified in this JLA PSP can serve to guide funders and researchers to topics that are most relevant and of primary importance to patients and healthcare professionals.

Contributors DG, BD and KC conceptualised the project. DG and BD prepared the manuscript. All authors were involved in the design, analysis of data and approved the final manuscript.

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