



LJMU Research Online

Rosala-Hallas, A, Jones, AP, Williamson, PR, Bedson, E, Compton, V, Fernandes, RM, Lacy, D, Lyttle, MD, Peak, M, Thorburn, K, Woolfall, K, Van Miert, C and McNamara, PS

Which outcomes should be used in future bronchiolitis trials? Developing a bronchiolitis core outcome set using a systematic review, Delphi survey and a consensus workshop

<https://researchonline.ljmu.ac.uk/id/eprint/18740/>

Article

Citation (please note it is advisable to refer to the publisher's version if you intend to cite from this work)

Rosala-Hallas, A, Jones, AP, Williamson, PR, Bedson, E, Compton, V, Fernandes, RM, Lacy, D, Lyttle, MD, Peak, M, Thorburn, K, Woolfall, K, Van Miert, C ORCID logoORCID: <https://orcid.org/0000-0003-0287-9832> and McNamara. PS (2022) Which outcomes should be used in future



LJMU has developed **LJMU Research Online** for users to access the research output of the University more effectively. Copyright © and Moral Rights for the papers on this site are retained by the individual authors and/or other copyright owners. Users may download and/or print one copy of any article(s) in LJMU Research Online to facilitate their private study or for non-commercial research. You may not engage in further distribution of the material or use it for any profit-making activities or any commercial gain.

The version presented here may differ from the published version or from the version of the record. Please see the repository URL above for details on accessing the published version and note that access may require a subscription.

For more information please contact researchonline@ljmu.ac.uk

<http://researchonline.ljmu.ac.uk/>

BMJ Open Which outcomes should be used in future bronchiolitis trials? Developing a bronchiolitis core outcome set using a systematic review, Delphi survey and a consensus workshop

A Rosala-Hallas,¹ Ashley P Jones ¹, Paula R Williamson,² Emma Bedson,¹ Vanessa Compton,³ Ricardo M Fernandes,^{4,5} David Lacy,⁶ Mark David Lyttle ^{7,8}, Matthew Peak,⁹ Kentigern Thorburn,³ Kerry Woolfall,¹⁰ Clare Van Miert,¹¹ Paul S McNamara¹²

To cite: Rosala-Hallas A, Jones AP, Williamson PR, *et al.* Which outcomes should be used in future bronchiolitis trials? Developing a bronchiolitis core outcome set using a systematic review, Delphi survey and a consensus workshop. *BMJ Open* 2022;**12**:e052943. doi:10.1136/bmjopen-2021-052943

► Prepublication history and additional supplemental material for this paper are available online. To view these files, please visit the journal online (<http://dx.doi.org/10.1136/bmjopen-2021-052943>).

CVM and PSM are joint senior authors.

Received 25 May 2021

Accepted 27 January 2022



© Author(s) (or their employer(s)) 2022. Re-use permitted under CC BY-NC. No commercial re-use. See rights and permissions. Published by BMJ.

For numbered affiliations see end of article.

Correspondence to

Professor Paul S McNamara; Mcnamp@liverpool.ac.uk

ABSTRACT

Objectives The objective of this study was to develop a core outcome set (COS) for use in future clinical trials in bronchiolitis. We wanted to find out which outcomes are important to healthcare professionals (HCPs) and to parents and which outcomes should be prioritised for use in future clinical trials.

Design and setting The study used a systematic review, workshops and interviews, a Delphi survey and a final consensus workshop.

Results Thirteen parents and 45 HCPs took part in 5 workshops; 15 other parents were also separately interviewed. Fifty-six items were identified from the systematic review, workshops and interviews. Rounds one and two of the Delphi survey involved 299 and 194 participants, respectively. Sixteen outcomes met the criteria for inclusion within the COS. The consensus meeting was attended by 10 participants, with representation from all three stakeholder groups. Nine outcomes were added, totalling 25 outcomes to be included in the COS.

Conclusion We have developed the first parent and HCP consensus on a COS for bronchiolitis in a hospital setting. The use of this COS will ensure outcomes in future bronchiolitis trials are important and relevant, and will enable the trial results to be compared and combined.

Trial registration number ISRCTN75766048.

INTRODUCTION

Background and objectives

Bronchiolitis, an acute viral lower respiratory tract infection which predominantly affects infants, is a major cause of morbidity and mortality worldwide.¹ Typical clinical features include a coryzal prodrome lasting approximately 3 days, persistent cough, increased respiratory rate, chest recession and wheeze or crackles on auscultation.² While most children with bronchiolitis have mild symptoms

Strengths and limitations of this study

- With stakeholder input from medically qualified staff, nurses and other clinical staff, and parents of children with bronchiolitis, we have developed a preliminary core outcome set (COS) for paediatric bronchiolitis trials.
- A mixed-method approach was used to determine which outcomes would be included in the COS.
- This COS can be considered when designing paediatric bronchiolitis trials as a minimum for outcomes to be collected and reported.
- While we followed guidance within the COMET (Core Outcome Measures in Effectiveness Trials) handbook in regards to Delphi processes and face-to-face consensus meeting format, one could argue that parental input during the consensus meeting had a relatively large effect on which items were included in the final COS.

and can be managed at home,³ in the UK and USA approximately 3% are hospitalised, most commonly between 3–6 months of age.^{1,4}

Although multiple therapeutic interventions for bronchiolitis have been assessed in clinical trials, treatment remains supportive.² Oxygen therapy and the use of oximetry are the only interventions that have significantly impacted survival over the last 40 years, contributing to a reduction in mortality from approximately 20% in some studies to <1%.^{5,6} Interventional clinical trials in bronchiolitis have increased in number over the past decade, investigating novel ways of administering oxygen nasally and non-invasively with varying levels of positive airway pressure or flow,^{7–9} and novel antiviral medications against respiratory syncytial virus, the

principal viral cause of bronchiolitis.^{10 11} More interventions are in the therapeutic pipeline.¹²

Randomised controlled trials (RCTs) are regarded as the gold standard for evaluating healthcare interventions.¹³ Although they produce high quality evidence that inform clinical care through practice guidelines, their clinical impact is often diminished by variations in outcome measurement and reporting. Systematic reviews in many different branches of medicine have consistently demonstrated the large number and heterogeneity of outcome reporting in trials and other research studies.^{14–16} This makes clinically relevant comparisons between trials and pooling of results in meta-analyses difficult. Furthermore, multiplicity of outcome measurement can lead to the selective reporting of significant findings, referred to as outcome reporting bias.¹⁷

A proposed solution is to develop and use a ‘Core Outcome Set’ (COS).¹⁸ This consists of a minimum set of outcomes that key stakeholders agree are important, the measurement of which should be considered for all trials in a particular field.¹⁸ This has the potential to improve the efficiency with which research can answer clinical questions. The benefits of COS have been embraced internationally by funding bodies,¹⁸ regulatory bodies^{19 20} and journal editors,²¹ all of which recommend their use where available. As a result, the development of COS is increasingly common. The COMET (Core Outcome Measures in Effectiveness Trials) initiative has recorded approximately 700 published or ongoing studies into COS in many branches of medicine. Until now there has been no COS for trials of children with bronchiolitis although the need for one has previously been identified.²²

The aim of this study was to obtain consensus from key stakeholders on which outcomes should be included in a COS for use in future bronchiolitis trials and other studies. This study formed part of a larger research project (Non-Invasive Ventilation for the Management of Children with Bronchiolitis: a feasibility study (NOVEMBR)).²³

METHODS

The COS was developed in three phases using methods recommended by COMET and COSMIN (CONsensus-based Standards for the selection of health Measurement INSTRUMENTS).¹⁸ The aim of the first phase was to generate a list of initial outcomes for consideration within the COS; this comprised a systematic review, stakeholder workshops and interviews. The second phase was a two-round Delphi survey, and the third phase, a face-to-face consensus meeting where results from the Delphi were presented.

COS-STAR (Core Outcome Set-STAndards for Reporting) guidelines for reporting were followed.²⁴

Patient and public involvement

Parents were involved in writing the original study protocol, information sheets and Delphi questionnaires. Discussion regarding non-invasive ventilation (NIV)

modalities was part of a separate exercise designing a protocol of NIV for infants with bronchiolitis. A parent was also a member of the trial management group.

Phase I: systematic review of the literature and stakeholder perspectives

Lists of outcomes were generated separately by both systematic review of the literature and stakeholder workshops/interviews.

The systematic review identified outcomes to assess efficacy and safety of interventions (pharmacological and non-pharmacological) used to treat children with acute bronchiolitis in published clinical trials since 2000 (search strategy provided in online supplemental appendix A). Details of this systematic review are not contained within this publication and will be published at a later date.

Parents or guardians were eligible to participate in a workshop or interview if their child had been admitted to hospital with bronchiolitis within the previous year. Eligible parents were invited to participate by research nurses at seven study sites (Alder Hey Children’s NHS Foundation Trust (Liverpool); Wirral University Teaching NHS Foundation Trust; Countess of Chester NHS Foundation Trust; Derby Teaching Hospital NHS Foundation Trust; Darlington Memorial Hospital; Cambridge University Hospitals NHS Trust and Royal Alexandra Hospital, Brighton. The lead centre will be Alder Hey Children’s NHS Foundation Trust (Liverpool)), through advertisements on social media. Healthcare professionals (HCPs) were eligible to participate in a workshop or interview if they had at least 6 months experience in managing children with bronchiolitis. Study sites emailed invitations to eligible HCPs to register interest in a workshop. Parent and HCP selection aimed to ensure variance (eg, child’s age 0–24 months, gender and severity of illness and HCP role and hospital geographical location).

For both HCP and parent workshops, interview topic guides exploring aspects of trial design were developed based on previous research.^{25–27} To identify prioritised outcomes, participants were first asked to reflect on their personal experiences including what they would consider as a sign that a child was getting better. To inform discussion, they were then shown a list of 34 outcomes (online supplemental appendix B) extracted from Cochrane Systematic Reviews and National Institute for Health and Care Excellence (NICE) guidance on bronchiolitis,²⁸ asked to consider which were important to them, and identify outcomes not included on the list. This process was adapted for parent interviews (eg, list of outcomes emailed to interview participants).

A professional transcription company (Voicescript, Bristol, UK) transcribed verbatim digital audio recordings. Transcripts were anonymised and checked for accuracy. NVivo V.10 software (QSR International, Melbourne, Australia) was used to assist in the organisation and coding of outcomes identified both through responses to direct questioning, and referred to by participants during interview and workshop discussions.

A conceptual outcomes framework was developed based on previous exploratory work in which domains and subdomains were defined and outcomes categorised accordingly.^{29 30} Similar outcomes identified by systematic literature review and workshops/interviews were collapsed together—following discussions between study management group members.

Phase II: Delphi process and outcome scoring

Participants

Key stakeholders who participated in the Delphi survey included parents/legal representatives of children hospitalised with bronchiolitis defined as per UK NICE Bronchiolitis Guidelines (2015).²⁸ These were identified through Phase I workshops and interviews, and by the seven study sites by research nurses on the wards and asked whether they would like to participate, they were then contacted at a later date. Parents of children who had died during their hospital admission were not approached. Non-English speakers were not eligible to participate.

Also included in the Delphi survey were HCPs (and nurses/other clinical staff) with experience of caring for children with bronchiolitis. These were identified via emails sent to professional organisations, and distributed via global email address lists or associated social media sites. HCPs who had previously completed the NOVEMBR National Survey of Current Practice were also approached if they had expressed an interest in participating.³¹ Participants were also invited to pass on details of the study to any of their own contacts who met eligibility requirements. The Delphi process was conducted and managed by DelphiManager software (www.comet-initiative.org/delphimanager). Access to the Delphi survey was via a hyperlink distributed by email. In round one, participants confirm their eligibility, and assigned a score (using a Likert scale of 1–9) to each of 56 outcomes, listed alphabetically, based on their opinion of its importance in the management of children with bronchiolitis. The Grading of Recommendations, Assessment, Development and Evaluations (GRADE) guidelines scale was used for scoring: scores of 1–3 indicated that the outcome was ‘of low importance’, 4–6 indicated ‘equivocal—important but not critical for decision making’ and 7–9 indicated ‘critical for decision-making’.³² Participants were also given the option to respond, ‘do not know’. Lastly, there was the opportunity to suggest any additional outcomes along with a score for importance. All additional outcomes were reviewed by the Study Management Group (SMG) (which contained clinicians, research nurses, trials unit representatives and parents) and where applicable carried forward to round two.

Those who completed round one were invited to participate in round two. In round two participants were shown the distribution of scores given by each stakeholder group and then asked to review and re-score each outcome. At the end of the first-round participants were shown their own scores for and asked to re-score based on the same question as the first round.

Table 1 Definition of consensus

Consensus classification	Description	Definition
Consensus in	Consensus that outcome should be included in the core outcome set	70% or more participants scoring as 7–9 AND <15% participants scoring as 1–3 in each group
Consensus out	Consensus that outcome should not be included in the core outcomes set	≤50% of participants scoring as 7–9 in each group
No consensus	Uncertainty about importance of outcome	Anything else

Data were presented as counts and percentages for categorical data and mean and SD for continuous data.

Consensus for inclusion of an outcome in the COS was achieved if 70% or more participants gave the outcome a score of 7–9 and less than 15% gave a score of 1–3 in each stakeholder group for round two of the survey (table 1) in all three stakeholder groups. The rationale for these levels of agreement was based guidance in the published literature.^{30 33} Consensus for exclusion of an outcome followed if 50% or fewer participants scored 7–9 in all of the three stakeholder groups. Any other outcome was classified as ‘no consensus’. Reminder emails were sent to participants to encourage completion. Participants who completed both Delphi survey rounds received a certificate of completion and were entered into a prize draw to win an iPad. Completion of the surveys was deemed as consent to participate.

Phase III: consensus meeting

The results from each round of the Delphi survey were presented at a face-to-face consensus meeting involving a representative group of stakeholders; members of this group were those who participated in the survey or workshops and expressed an interest in attending the consensus meeting. Participants were not specifically invited based on their views in the survey and were not required to have completed the Delphi survey in order to participate. An independent medical professional, with expertise in COS development and not a member of the SMG, chaired the meeting. Meeting attendees were asked to review the full list of outcomes and were given the opportunity to discuss whether they agreed with the outcome consensus classifications from the Delphi process in Phase II. They were then asked to discuss the outcomes classified as ‘no consensus’ from the Delphi process. The Chair ensured that all participants had equal opportunity to give their views on each of the outcomes prior to voting taking place. Voting was carried out with the same 9-point Likert scale

used in the Delphi (described in Phase II) and conducted anonymously using TurningPoint software and handsets (Turning Technologies LLC, Youngstown, USA). At the end of the meeting the outcomes that met the criteria for 'consensus in' were presented to the stakeholders.

Registration

The study protocol was registered retrospectively with the COMET initiative and is available online,²³ and on the ISRCTN Registry (18 December 2017).

RESULTS

Phase I: literature review and stakeholder perspectives

The systematic review identified 154 studies for inclusion (online supplemental figure 1). Within these studies, 923 individual outcome measures were identified. These outcomes were grouped and tabulated under appropriate outcome domains using a predefined conceptual framework as described previously.^{29 30} This work is being written up as a separate manuscript.

Between April 2016 and March 2017, 13 parents and 45 practitioners took part in five workshops (three for HCPs and two for parents); 15 other parents were also interviewed by telephone. No outcomes were identified during these workshops and interviews that had also not been identified as part of the systematic review.

All outcomes identified from the systematic review, workshops and interviews were reviewed by for similarity. Outcomes considered sufficiently similar with regards to what they measured were collapsed and merged together. Furthermore, this process was discussed and reviewed with the SMG to agree the final list of 56 outcomes to be included into the Delphi survey (online supplemental table 1).

Phase II: Delphi process

Round one of the survey was conducted between 19 February 2018 and 23 March 2018. Round two was conducted between 29 March 2018 and 13 April 2018.

Online supplemental table 2 shows the breakdown of stakeholder group and their participation in both rounds. In total, 299 participants (from the UK) registered on the online system and 286 (96%) scored at least one outcome in round one. Sixty-eight per cent (194/286) were medically qualified, 28% (81/286) were nurses and other clinical staff, and the remaining 4% (11/286) were parents. Participants were invited to round two if they had scored at least one outcome in round one. Sixty-eight per cent (194/286) participated in round two by scoring at least one outcome. Mean round one scores for those who participated in both rounds and those who participated in round one only were similar (online supplemental table 3).

Following round one, one item was added to the list of outcomes for consideration in round two: length of time spent on oxygen.

Table 2 Final outcomes to be included in a core outcome set from the Delphi process

Domain (subdomain)	Outcome
Physiological and clinical (general symptoms)	Appearance Level of consciousness Non-respiratory physiological parameters/vital signs Worsening illness
Physiological and clinical (feeding; nutrition and hydration)	Feeding Need for feeding tube Inhalation (breathing in) of milk; fluids or solids
Physiological and clinical (respiratory distress)	Apnoea Oxygen saturation Cyanosis Effort of breathing Paediatric Early Warning score
Physiological and clinical (respiratory interventions and support)	Need for respiratory support
Resource use (hospital related short term)	Critical care admission
Death	Death
Adverse events	Serious adverse events

Online supplemental table 4 shows the results from round two. Sixteen outcomes met the criteria for 'consensus in' for inclusion in the COS, 8 outcomes met the criteria for 'consensus out' and 32 outcomes were classified as 'no consensus'.

The final set of 16 outcomes that were included from the Delphi process are included in [table 2](#).

Phase III: consensus meeting

The consensus meeting took place on the 14 June 2018 and was attended by 10 participants: 4 were medically qualified staff, 3 were parents and the remaining 3 were nurses and other clinical staff.

When round two classifications of 'consensus in' outcomes were reviewed, the inclusion of Paediatric Early Warning (PEW) score was discussed as not every hospital uses it and even in those that do, the composite measures can vary. However, participants agreed it should remain in the COS as a standardised, widely used objective PEW score has the potential to be a quick indicator of changing health and healthcare needs. No further comments were provided on 'consensus in' outcomes. Discussion around 'consensus out' outcomes was centred on the economic cost outcome; one participant believed it should be included since it impacts on whether the intervention is adopted by some healthcare funders. After some discussion, it was agreed that other core outcomes can be used

Table 3 Outcomes that were added to the core outcome set during the consensus meeting

Domain (subdomain)	Outcome
Physiological and clinical (general symptoms)	Pain and discomfort Parent report of symptoms and/or resolution of illness
Physiological and clinical (feeding; nutrition and hydration)	Reduced urine output
Physiological and clinical (respiratory interventions and support)	Length of time spent on oxygen
Life impact (health-related quality of life)	Quality of life
Resource use (hospital related short term)	Hospital length of stay Time until ready for discharge from hospital
Adverse events	Adverse events Long-term effects of illness or treatment interventions

for a health economic evaluation, without the need for a separate outcome.

Following discussion and re-vote, nine outcomes were added to the 'consensus in' list (table 3). The final COS contains 25 outcomes across eight domains (tables 2 and 3). In the Delphi, need for fluids given through a drip (intravenously) was scored 7–9 by 80%, 85% and 67% of medically qualified staff, nurses and parents, respectively. However, in the consensus meeting only 20% of meeting attendants scored it 7–9; discussions at the time centred on the relative subjectivity of the assessment of need for fluids through a drip and that it was not a critical outcome. Likewise, bronchiolitis severity score and additional chest infections/pneumonia were close to being included as 'consensus in' in the Delphi but were voted 'consensus out' during the consensus meeting. There were differing views in the room on whether or not a disease specific score should be included in the COS but after discussion, only 60% voted 7–9. Although all agreed additional chest infection/pneumonia was important, it was decided that it was not critical. A full report from the consensus meeting detailing discussion behind each outcome has been included in online supplemental appendix C. At the close of the meeting the final COS was agreed by all the participants.

Protocol changes

The definition of 'Consensus Out' was changed from '70% or more participants scoring 1–3 and <15% of participants scoring 7–9 in each group' to '≤50% of participants scoring 7–9 in each group'. This change was made for practical reasons only, in order to refine and reduce the list of outcomes to be discussed at the consensus meeting. Meeting attendees were also given

the opportunity to comment on and discuss any of the 'consensus out' outcomes.

DISCUSSION

We have developed a COS of 25 outcomes of importance to medically qualified staff, nurses and other clinical staff, and parents of children with bronchiolitis, for use in RCTs of interventions for children with a clinical diagnosis of bronchiolitis in a hospital setting.

On the COMET database, 10% of the approximate 700 listed published and ongoing COS are in child health with most being for chronic paediatric diseases. Those for acute conditions include sepsis, infantile colic, appendicitis, head injury, acute diarrhoea and asthma. Two other studies on the COMET database mention bronchiolitis, one a systematic review of outcome measures and measurement instruments used in bronchiolitis RCTs,³⁴ and the other a European Respiratory Society guideline on 'Endpoints in respiratory diseases' based solely on expert views of HCPs and published in 2010.³⁵ This is the only COS developed for use in for bronchiolitis, one of the most common acute causes of childhood admission to hospital, developed using recommended consensus based methods.¹⁸

One of the strengths of this study was the involvement of both parent and HCPs in each phase and the necessity for agreement by each stakeholder group before consensus could be reached; this ensured all views were incorporated throughout. For example, at the final consensus meeting, discussions were carefully 'managed' by an independent chair who ensured all participants could voice their opinions and have their opinions heard. The use of 'anonymous' TurningPoint software also allowed peer/social pressures to be minimised when it came to voting. This enabled the views of all participants to be heard, and particularly those of parents who might otherwise have been reluctant to contribute when surrounded by experienced HCPs. As a result, parental 'championing' of four 'no consensus' items led directly to their inclusion in the COS (pain and discomfort, parent report of symptoms and/or resolution of illness, quality of life and reduced urine output). Three of these are relatively subjective measures. For changes in urine output, parents argued that this provided a tangible way for them to know whether a treatment or intervention was working and their child was 'on the mend'.

There were a number of limitations to this study. Since the Delphi survey was disseminated via mailing lists, it was not possible to ascertain the total number of individuals approached and therefore the proportion who took part. Also, although reminders for completion were circulated, not all participants who took part in round one also took part in round two. However, comparing scores for round one for those who completed both rounds with scores for those who completed round one only, did not highlight any attrition bias. Indeed, the overall number of participants who took part in both rounds of this study (194)

was above the median of 111 from a recent review of 31 published and ongoing studies that used a Delphi to develop a COS.³⁵ In our study, having fewer outcomes to assess may have increased the response rate from round one to round two.³⁶

We would have liked to include a larger number of parents in the Delphi survey. Parents were approached during their child's admission to hospital with bronchiolitis and many agreed to take part in the study and gave their contact details. However, only 11 of these subsequently completed the two surveys. We speculate that by the time parents of otherwise healthy infants and young children were contacted some months after the acute bronchiolitis episode, time pressures perhaps meant that their priorities lay elsewhere. Of note, 30% of attendees at the consensus meeting were parents, which ensured their voice was heard at this crucial part of the process and that the consensus meeting comprised a representative group of stakeholders.

Responders were primarily UK based which raises the question of whether the COS would be applicable to trials overseas. An extension of this work would be to repeat the process including international participants, to determine whether the choice of outcomes changes depending on location. It is likely that this would indeed be so, given that respiratory support or nasogastric feeding are not universally available in all settings. Future work on the COS could also include pharmaceutical representatives and regulators.

Burden of measurement should also be taken into consideration when developing a COS. Although the COS we have developed contains 25 outcomes, it can be seen that several of the outcomes are related, for example, adverse events and serious adverse events would both be captured when measuring safety. Similarly, feeding and need for feeding tube, hospital length of stay and time until ready for discharge from hospital are related and would require minimal time to capture these data.

We believe the COS should be used by future trialists as the minimum set of outcomes that should be collected in a bronchiolitis trial. Importantly, this COS provides an essential step towards increasing and improving much-needed evidence synthesis in this disease area.^{22 37} There was some discussion during the consensus meeting on how potential outcomes would be measured, for example appearance, however the chair reminded participants that the aim of the meeting was to agree on 'what' should be measured rather than 'how' to be measured. Now that we have established which outcomes should be reported, there is a need for future research to ascertain the best methods to operationalise these outcomes, particularly some of the more subjective ones such as pain and discomfort, parent report of symptoms and inhalation of milk/feeds.

CONCLUSIONS

This was an important study identifying the outcomes of importance to key stakeholder groups which will provide guidance to future trialists in paediatric bronchiolitis. This is particularly pertinent with anti-viral treatments in the therapeutic pipeline, and the need to clearly define how some existing interventions (such as high flow nasal cannula oxygen) are best used in clinical practice. We recommend researchers designing bronchiolitis trials consider this preliminary COS as a minimum for outcomes collected and reported in trials. Future work is now needed to test the validity and generalisability of the COS and ongoing work should include routine updating of the COS.

Author affiliations

¹Liverpool Clinical Trials Centre, University of Liverpool, Liverpool, Merseyside, UK

²Health Data Science, Institute of Population Health, University of Liverpool, Liverpool, UK

³Paediatric Intensive Care Unit, Alder Hey Children's NHS Foundation Trust, Liverpool, UK

⁴Department of Pediatrics, Santa Maria Hospital, Lisbon Academic Medical Centre, Lisbon, Portugal

⁵Clinical Pharmacology and Therapeutics, Faculty of Medicine, Instituto de Medicina Molecular, Universidade de Lisboa, Lisboa, Portugal

⁶Department of Paediatrics, Wirral University Teaching Hospital NHS Foundation Trust, Wirral, Merseyside, UK

⁷Emergency Department, Bristol Royal Hospital for Children, Bristol, UK

⁸Faculty of Health and Applied Science, University of the West of England, Bristol, UK

⁹Clinical Research Division, Alder Hey Children's NHS Foundation Trust, Liverpool, UK

¹⁰Institute of Population, Health and Society, University of Liverpool, Liverpool, UK

¹¹School of Nursing and Allied Health, Liverpool John Moores University, Liverpool, UK

¹²Department of Child Health (University of Liverpool), Institute in the Park, Alder Hey Children's Hospital, Liverpool, UK

Twitter Mark David Lytle @mdlytle

Acknowledgements The study team would like to thank all those who took part in the Delphi survey and to those who attended and contributed to the consensus meeting. We would like to acknowledge Dr Steven Lane for assisting in the design of the study and performing the preliminary analysis. We would also like to thank the independent chair of the consensus meeting, Associate Clinical Professor Ian Sinha.

Contributors PM and CVM conceived the study. PM, CVM, RMF, MDL, PRW and KW designed the study. CVM undertook the systematic review. CVM and KW facilitated healthcare and parent workshops, and CVM conducted parent interviews. EB, MP, KT, VC, KW and DL were members of the SMG. AR-H performed the final analysis of the data. APJ supervised data analysis. AR-H, APJ, KW and PM drafted the manuscript. All authors reviewed and provided comments on the manuscript. PM accepts full responsibility for this work and acts as guarantor.

Funding This study formed part of a larger research project (Non-Invasive Ventilation for the Management of Children with Bronchiolitis: a feasibility study (NOVEMBR)) which was funded by the National Institute for Health Research (NIHR), Research for Patient Benefit (RfPB) Programme (Grant Reference Number: PB-PG-1014-35081). Views expressed are those of the authors and not necessarily those of the NHS, the NIHR or the Department of Health.

Competing interests The authors declare that they have no competing interests.

Patient consent for publication Not applicable.

Ethics statement Ethics approval was provided by the NHS Research Ethics Service (REC: 17/EM/0471; IRAS Project ID 238056). Written informed consent was sought prior to interviews and workshops during Phase I. A statement was included on the Delphi registration page highlighting that completion of the questionnaire is regarded as consent. On arrival for Phase III, all participants completed a consent form to confirm they were happy to take part.

Provenance and peer review Not commissioned; externally peer reviewed.

Data availability statement Data can be made available on reasonable request from the corresponding author.

Supplemental material This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

Open access This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>.

ORCID iDs

Ashley P Jones <http://orcid.org/0000-0001-5253-730X>

Mark David Lyttle <http://orcid.org/0000-0002-8634-7210>

REFERENCES

- Florin TA, Plint AC, Zorc JJ. Viral bronchiolitis. *Lancet* 2017;389:211–24.
- Ricci V, Delgado Nunes V, Murphy MS, et al. Bronchiolitis in children: summary of NICE guidance. *BMJ* 2015;350:h2305.
- Viswanathan Met al. Management of bronchiolitis in infants and children. *Evid Rep Technol Assess* 2003;69:1–5.
- Murray J, Bottle A, Sharland M, et al. Risk factors for hospital admission with RSV bronchiolitis in England: a population-based birth cohort study. *PLoS One* 2014;9:e89186.
- Deshpande SA, Northern V. The clinical and health economic burden of respiratory syncytial virus disease among children under 2 years of age in a defined geographical area. *Arch Dis Child* 2003;88:1065–9.
- Panickar JR, Dodd SR, Smyth RL, et al. Trends in deaths from respiratory illness in children in England and Wales from 1968 to 2000. *Thorax* 2005;60:1035–8.
- Luo J, Duke T, Chisti MJ, et al. Efficacy of high-flow nasal cannula vs standard oxygen therapy or nasal continuous positive airway pressure in children with respiratory distress: a meta-analysis. *J Pediatr* 2019;215:199–208.
- Kepreotes E, Whitehead B, Attia J, et al. High-Flow warm humidified oxygen versus standard low-flow nasal cannula oxygen for moderate bronchiolitis (HFWHO RCT): an open, phase 4, randomised controlled trial. *Lancet* 2017;389:930–9.
- Franklin D, Babl FE, Schlapbach LJ, et al. A randomized trial of high-flow oxygen therapy in infants with bronchiolitis. *N Engl J Med* 2018;378:1121–31.
- DeVincenzo JP, Whitley RJ, Mackman RL, et al. Oral GS-5806 activity in a respiratory syncytial virus challenge study. *N Engl J Med* 2014;371:711–22.
- DeVincenzo JP, McClure MW, Symons JA, et al. Activity of oral ALS-008176 in a respiratory syncytial virus challenge study. *N Engl J Med* 2015;373:2048–58.
- Xing Y, Proesmans M. New therapies for acute RSV infections: where are we? *Eur J Pediatr* 2019;178:131–8.
- Pocock SJ, Elbourne DR. Randomized trials or observational tribulations? *N Engl J Med* 2000;342:1907–9.
- Hirsch BR, Califf RM, Cheng SK, et al. Characteristics of oncology clinical trials: insights from a systematic analysis of ClinicalTrials.gov. *JAMA Intern Med* 2013;173:972–9.
- Meher S, Alfrevic Z. Choice of primary outcomes in randomised trials and systematic reviews evaluating interventions for preterm birth prevention: a systematic review. *BJOG* 2014;121:1188–94.
- Rodgers S, Brealey S, Jefferson L, et al. Exploring the outcomes in studies of primary frozen shoulder: is there a need for a core outcome set? *Qual Life Res* 2014;23:2495–504.
- Kirkham JJ, Dwan KM, Altman DG, et al. The impact of outcome reporting bias in randomised controlled trials on a cohort of systematic reviews. *BMJ* 2010;340:c365.
- Williamson PR, Altman DG, Blazeby JM, et al. Developing core outcome sets for clinical trials: issues to consider. *Trials* 2012;13:132.
- Administration FaD. CDER pilot grant program: standard core clinical outcome assessments (CoAs) and their related endpoints, 2020
- The European Agency for the Evaluation of Medicinal Products, E.o.M.f.H.U. Points to consider on clinical investigation of medicinal products other than NSAIDs for treatment of rheumatoid arthritis, 2003
- Khan K. The crown initiative: Journal editors invite researchers to develop core outcomes in women's health. *Midwifery* 2014;30:1147–8.
- Fernandes RM, Oleszczuk M, Woods CR, et al. The Cochrane library and safety of systemic corticosteroids for acute respiratory conditions in children: an overview of reviews. *Evid Based Child Health* 2014;9:733–47.
- van Miert C, Fernandes RM, Eccleson H, et al. Non-Invasive ventilation for the management of children with bronchiolitis (NOVEMBR): a feasibility study and core outcome set development protocol. *Trials* 2018;19:627.
- Kirkham JJ, Gorst S, Altman DG, et al. Core outcome Set-STAndards for reporting: the COS-STAR statement. *PLoS Med* 2016;13:e1002148.
- Woolfall K, Frith L, Gamble C, et al. How parents and practitioners experience research without prior consent (deferred consent) for emergency research involving children with life threatening conditions: a mixed method study. *BMJ Open* 2015;5:e008522.
- Woolfall K, Young B, Frith L, et al. Doing challenging research studies in a patient-centred way: a qualitative study to inform a randomised controlled trial in the paediatric emergency care setting. *BMJ Open* 2014;4:e005045.
- Knapp P, Raynor DK, Silcock J, et al. Can user testing of a clinical trial patient information sheet make it fit-for-purpose?--a randomized controlled trial. *BMC Med* 2011;9:89.
- NICE, National Institute for Health and Care Excellence guideline. Bronchiolitis in children: diagnosis and management (NG9), 2015
- Fernandes RMRMC. *Interventions and outcomes in bronchiolitis clinical trials*. Universidade de Lisboa (Portugal), ProQuest Dissertations Publishing, 2015.
- Williamson PR, Altman DG, Bagley H, et al. The comet Handbook: version 1.0. *Trials* 2017;18:280.
- Rosala-Hallas A, Jones AP, Bedson E, et al. National survey of feasibility of NIV trials for management of children with bronchiolitis. *BMJ Paediatr Open* 2020;4:e000780.
- Grade. Working group.
- Harman NL, Bruce IA, Callery P, et al. MOMENT--Management of Otitis Media with Effusion in Cleft Palate: protocol for a systematic review of the literature and identification of a core outcome set using a Delphi survey. *Trials* 2013;14:70.
- Fernandes RM. Outcomes in clinical trials of bronchiolitis 2009.
- de Benedictis FM, Guidi R, Carraro S, et al. Endpoints in respiratory diseases. *Eur J Clin Pharmacol* 2011;67 Suppl 1:49–59.
- Gargon E, Crew R, Burnside G, et al. Higher number of items associated with significantly lower response rates in COS Delphi surveys. *J Clin Epidemiol* 2019;108:110–20.
- Schuh S, Babl FE, Dalziel SR, et al. Practice variation in acute bronchiolitis: a pediatric emergency research networks study. *Pediatrics* 2017;140.