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Parental perspective on important health outcomes of extremely preterm infants

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ABSTRACT

Background and objective Neonatal outcome research and clinical follow-up principally focus on neurodevelopmental impairment (NDI) after extremely preterm birth, as defined by the scientific community, without parental input. This survey aimed to investigate parental perspectives about the health and development of their preterm children.

Methods Parents of children aged 18 months to 7 years born <29 weeks' gestational age presenting at a neonatal follow-up clinic over a 1-year period were asked to evaluate their children's health and development. They were also asked the following question: 'if you could improve two things about your child, what would they be?' Responses were analysed using mixed methods. Logistic regressions were done to compare parental responses.

Results 248 parents of 213 children (mean gestational age 26.6±1.6 weeks, 20% with severe NDI) were recruited. Parents evaluated their children's health at a median of 9/10. Parental priorities for health improvements were (1) development, mainly behaviour, emotional health and language/communication (55%); (2) respiratory health and overall medical fragility (25%); and (3) feeding/growth issues (14%). Nineteen per cent explicitly mentioned 'no improvements'. Parents were more likely to state 'no improvements' if child had no versus severe NDI OR 4.33 (95% CI 1.47 to 12.75) or if parents had no versus at least a high school diploma (OR 4.01 (95% CI 1.99 to 8.10)).

Conclusions Parents evaluate the health of their preterm children as being very good, with positive perspectives. Parental concerns outside the developmental sphere should also be addressed both in clinical follow-up and research.

INTRODUCTION

Preterm birth is associated with increased risk of neurosensory impairment and developmental delay.¹⁻³ The systematic assessment of these aspects of development are combined and categorised by levels of neurodevelopmental impairment (NDI).⁴ To monitor neurodevelopmental outcomes, neonatal follow-up programmes have become the standard of care.⁴⁻⁶ Neonatal follow-up research has thus been shaping the landscape of prematurity outcomes for several decades, determining which outcomes are reported. Classification of the outcomes of preterm children into NDI categories is now the norm in research.⁷ However, less interest is directed towards whether these outcomes are also of importance to families.⁸ This often

What is already known on this topic?

- Preterm outcome research generally focuses on adverse health and neurodevelopmental rates.
- Research results are used to council parents yet parental opinions regarding what outcomes are important for them has not been studied.

What this study adds?

- Parents of extremely preterm children report a balanced perspective on their child.
- This includes positive traits as well as concerns which encompass development and medical fragility, nutrition and respiratory health.
- Concerns vary with child's neurodevelopment, age and parental education.

leads to the erroneous perception that some children have a 'bad' or 'good' outcome, only based on NDI. Furthermore, children categorised as having 'normal' outcomes/no disability may have problems that are overlooked.⁹⁻¹⁰ The reporting of outcomes pertaining to the health and development of preterm children is undoubtedly essential, but could benefit from adequate representation and framing when it comes to families.¹¹⁻¹²

Researchers across medical fields are increasingly reporting patient-important outcomes. Initiative such as those reported by the Patient-Centered Outcomes Research Institute are thought to help determine outcomes that patients find most important, as they represent their lived experience with illness.¹³ The aim of the current study was to expand the knowledge of parental/family perspectives regarding the health of their children born preterm.

METHODS

Design and study participants

The Parents' Voices Project is a cross-sectional survey of parents of extremely preterm children seen at a level-III university hospital neonatal follow-up clinic. All infants born <29 weeks' gestational age (GA) are eligible for follow-up after discharge from the neonatal intensive care unit (NICU) for neurodevelopmental monitoring. All children are seen at 18 months corrected age (CA), at 36 months CA,



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then at 5 years and 7 years, according to standardised follow-up protocols (online supplemental table s1).¹

Between 1 July 2018 and 31 July 2019, parents of preterm children who were scheduled for a neonatal follow-up clinic visit were approached to participate. Foster parents were excluded.

Study procedure

At least one parent per eligible child was sent an email invitation to participate in the study. If parents did not respond, a research assistant attempted to contact them by telephone or in person at the follow-up visit. They were offered to respond to the questionnaire prior to the follow-up visit either online, on paper or by interview.

The questionnaire was developed and reviewed in partnership with parent stakeholders. Parents were asked to rate their child's health on a scale of 0 (very poor) to 10 (excellent). They were then asked to answer the following question: 'If you could improve any two things about your child's health and/or development, what would they be?' When possible, both parents were invited to participate separately. Parents of twins or triplets were asked to complete one questionnaire per child.

Demographic data and developmental outcomes

Parent characteristics including self-reported race, highest level of education and family composition were retrieved from questionnaires administered to all families during clinical care. Neonatal characteristics and demographics were collected from the infant's medical record by trained research personnel using Canadian Neonatal Network abstractor manual definitions.¹⁴ Neurodevelopmental outcome was determined for each child at the standardised 18-month CA visit following the Canadian Neonatal Follow-Up Network protocol.⁴ In summary, all children were administered the Bayley Scales of Infant and Toddler Development, 3rd edition (Bayley III). A neurological examination was performed by a paediatrician to identify signs of cerebral palsy and determine functional level using the Gross Motor Functional Classification System (GMFCS).¹⁵ Finally, data on visual and hearing function were retrieved from the medical chart. A level of neurodevelopmental impairment (NDI) was assigned for each child based on the 18-month assessment (table 1).⁴

Data analyses

Descriptive statistics were used to describe the study population and parental evaluations of their children's health. When both parents provided a rating for a single child, the mean score was

calculated. Comparison of parental rating score in relation to NDI status was done using the Kruskal-Wallis test.

Answers to the open-ended question were analysed using descriptive content analysis. Themes were developed simultaneously and independently by a team of eight investigators representing neonatology, general paediatrics, developmental paediatrics, rehabilitation care, sociology and parents. During the initial exploration of data, codes, subcodes, and coding definition and structure were developed gradually until consensus was reached. Then, systematic coding of all parental answers was done in an independent fashion by rotating teams of two investigators and discrepancies were resolved with a third coder. This brought multiple perspectives to the analysis to prevent mistakes and individual biases; all data were reviewed by a parent in initial or final coding steps. Codes of the qualitative content were introduced in the database as dichotomised variables (Yes/No). Statistical analyses were conducted with SPSS (IBM SPSS Statistics V.24). Logistic regressions were done to calculate the frequency of each theme in relation to the child's age, gestational age, level of NDI (none, mild to moderate, severe) and parental characteristics. As these analyses were exploratory in nature, we did not adjust for multiple comparisons.

RESULTS

During the study period, at least one parent for the 258 children scheduled for follow-up were contacted either by email, telephone or in person (figure 1); 248 parents of 213 children (83% of eligible children) participated and provided 285 individual responses. Both parents answered the questionnaire for 71 children; there were only maternal answers for 128 children and only paternal answers for 14. Among responding mothers and fathers, 63% and 82%, respectively, self-identified as Caucasian and 82% and 87%, respectively, reported having at least a high school diploma (table 2).

The 213 children were born at a mean gestational age of 26.6 weeks with mean birth weight of 907 g (table 3). For the majority (59%) of participants, parents answered the questionnaire at the 18-month or 36-month CA visit. Based on the 18-month assessment, 55% met criteria for no NDI, 25% had a mild to moderate NDI and 20% a severe NDI.

Evaluation of health

For the 213 children, the median score for health as reported by parents was 9/10 (IQR 7.5–10, range 3–10). When both parents responded for the same child, agreement was strong (Pearson's $r=0.668$). For children with no NDI, mild to moderate NDI,

Table 1 Definition of neurodevelopmental impairment

Impairments	Severe neurodevelopmental impairment (any one or more of the following)*	Mild-moderate neurodevelopmental impairment (any one or more of the following)†	No neurodevelopmental impairment
Motor	CP with GMFCS 3, 4 or 5 Bayley III Motor Composite <70	CP with GMFCS 1 or 2 Bayley III Motor Composite 70–84	No cerebral palsy Bayley III Motor Composite ≥85
Cognitive	Bayley III Cognitive Composite <70	Bayley III Cognitive Composite 70–84	Bayley III Cognitive Composite ≥85
Language	Bayley III Language Composite <70	Bayley III Language Composite 70–84	Bayley III Language Composite ≥85
Hearing	Hearing aid or cochlear implant	Sensorineural/mixed hearing loss without requirement for hearing devices	No hearing loss as per audiology report
Vision	Bilateral visual impairment	Unilateral visual impairment	No visual impairment as per ophthalmology report

*Includes children who could not be tested using the Bayley-III but who had a Bayley-III Adaptive Behaviour score <70 or who were assessed to have a significant developmental delay.

†Children who could not be tested using the Bayley-III but who had a Bayley-III Adaptive Behaviour score between 70 and 84. CP, cerebral palsy; GMFCS, Gross Motor Function Classification System.

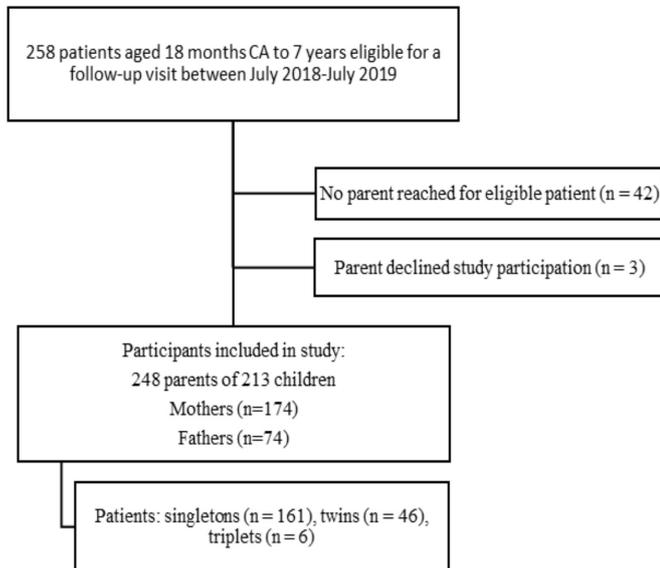


Figure 1 Flowchart of study population.

and severe NDI, the median health score was 9 (IQR 8.5–10), 8 (IQR 7–9) and 7 (IQR 6–9), respectively ($p < 0.001$). We examined children whose parent(s) rated their health as 6 or less (≤ 10 th percentile of the cohort, $n = 28$): 4 had no NDI, 9 mild to moderate NDI, and 15 severe NDI.

Parental priorities for health improvements

The main themes identified included (1) development (55%), (2) respiratory health and fragility (25%), (3) nutrition/feeding and growth problems (14%), with (4) 19% ($n = 52$) of parents reporting there was nothing to improve (figure 2). When parents described their child's development as suboptimal, language and communication (19%), behaviour and emotional health (18%), motor development and movements (15%) as well as cognitive and learning skills (14%) were most often cited as top concerns. Table 4 shows parents describing a wide array of problems.

Association between parent and child characteristics and parental priorities

Figure 3A shows a comparison of themes reported by parents of preterm children by NDI category (online supplemental tables s2 and s3). Development was mentioned equally by all parents. Growth and feeding were more concerning to parents of children in the severe NDI group, as compared with the no-NDI group (OR 2.86; 95% CI 1.27 to 6.45). Respiratory illness and

Table 3 Child characteristics ($n = 213$)

Mean gestational age (SD), weeks	26.6 (1.6)
Mean birth weight (SD), g	907 (247)
Male, n (%)	106 (50)
Multiple births, n (%)	65 (31)
Severe brain injury**, n (%)	44 (21)
Bronchopulmonary dysplasia††, n (%)	130/203 (64)
Median duration of NICU stay (IQR), days	105 (78, 130)
NDI at 18 months CA, n (%)	
None	114/209 (55)
Mild/moderate	53/209 (25)
Severe	42/209 (20)
Mean Bayley-III scores at 18 months CA (SD)	
Cognition	94 (14)
Language	91 (16)
Motor	91 (14)
Current age, n (%)	
18 months CA	65 (31)
36 months CA	59 (28)
5 years	56 (26)
7 years	33 (16)

*Severe brain injury defined as grade 3–4 intraventricular haemorrhage, cystic periventricular leucomalacia, cerebellar haemorrhage.

†Bronchopulmonary dysplasia defined as oxygen use at 36 weeks of postmenstrual age.

CA, corrected age; NDI, neurodevelopmental impairment; NICU, neonatal intensive care unit.

fragility were more frequently reported as priorities by parents of children in the mild-moderate NDI category, as compared with the severe NDI category (OR 2.50; 95% CI 1.04 to 6.00). Finally, parents of children without NDI were more likely to report having nothing to improve compared with parents in

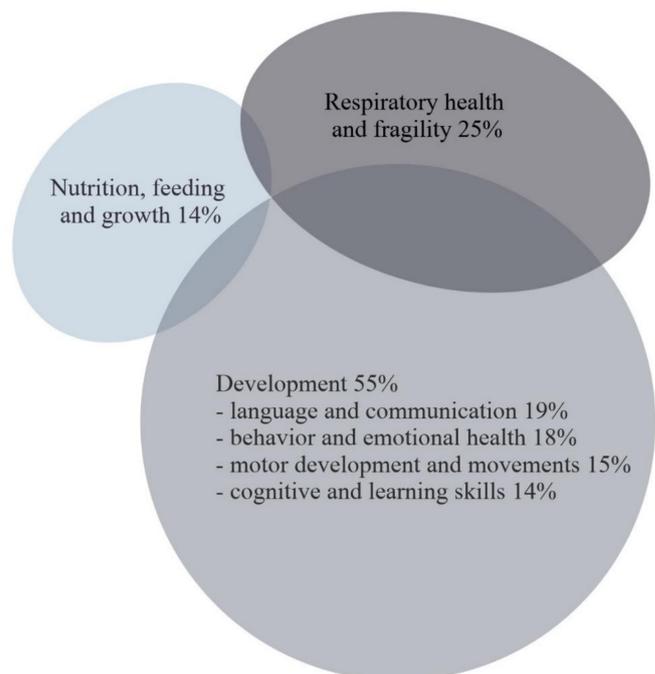


Figure 2 Venn diagram of parental priorities for improvements. Some parents reported on several themes, which is illustrated by the overlapping areas.

Table 2 Parental characteristics

	Mothers n=174	Fathers n=74
Mean age at birth (SD), years	32 (5)	NA
Lone parent, n (%)	14/172 (8)	1/74 (1)
Self-reported race, n (%)		
Caucasian	108/171 (63)	59/72 (82)
Black	42/171 (25)	7/72 (10)
Other	21/171 (12)	6/72 (8)
Education \geq high school, n (%)	138/168 (82)	62/71 (87)
No of responses provided	199	86
Singleton	139 (70%)	60 (70%)
Twin or triplet	60 (30%)	26 (30%)

Table 4 Examples of quotes provided by parents

Development	<p>"Her language, I wish she would express herself more clearly."</p> <p>"A better management of his crises (when he gets out of control)."</p> <p>"She acts like a baby, I'm worried about school."</p> <p>"His gross motor skills, including running."</p>	<p>Mother of a 3-year-old girl born at 26 weeks</p> <p>Father of 5-year-old born at 27 weeks</p> <p>Mother of a 3-year-old girl born at 26 weeks</p> <p>Mother of a 3-year-old boy born at 27 weeks</p>
Respiratory health and fragility	<p>"His lungs are still fragile."</p> <p>"His immune system, his pneumonias, he is sick all the time."</p> <p>"Inflammation of his lungs every time he catches a virus."</p>	<p>Mother of a 5-year-old boy born at 25 weeks</p> <p>Mother of an 18-month-old boy born at 27 weeks</p> <p>Father of 18-month-old boy born at 28 weeks</p>
Growth and nutrition	<p>"I hope she will learn how to eat."</p> <p>"Remove her gastrostomy."</p> <p>"His reflux and frequent vomiting. He does not eat much."</p>	<p>Father of an 18-month-old girl born at 25 weeks</p> <p>Mother of an 18-month-old girl born at 23 weeks</p> <p>Father of 5-year-old boy born at 25 weeks</p>

the two other groups. **Figure 3B** compares parental responses according to their child's age (online supplemental tables s2 and s3). Parents of 18-month-old children were less likely to mention development as an area for improvement. Parents of children aged 18–36 months were more likely to express concerns regarding respiratory health, compared with the 7-year-old

group, and growth and nutrition issues compared with parents of older children.

There was no difference between mothers and fathers or by gestational age groups (22–24 weeks, 25–26 weeks, 27–28 weeks) with respect to the frequency of themes invoked by parents (online supplemental tables s2 and s3). When examining

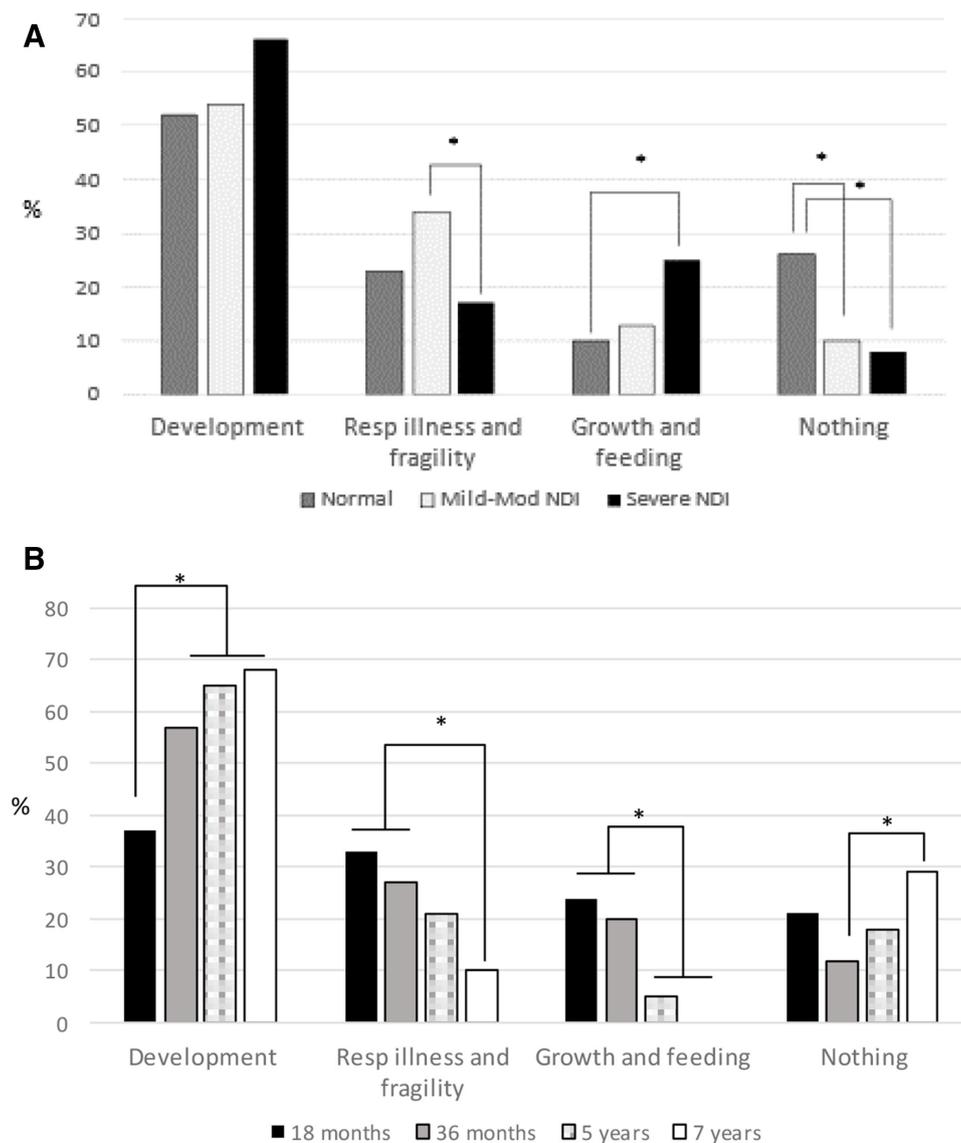


Figure 3 Areas of improvements by (A) neurodevelopmental impairment, (B) child age. NDI, neurodevelopmental impairment. * $p < 0.05$ by bivariate logistic regression.

responses in relation to level of education, parents who had not completed high school were less likely to report wishing to improve their child's development compared with parents with at least a high school diploma (OR 0.46, 95% CI 0.24 to 0.88). They were also more likely to respond that nothing needed to change (OR 4.01; 95% CI 1.99 to 8.10).

DISCUSSION

Measuring and classifying the outcomes of preterm infants is important for a number of reasons: identifying health and developmental concerns to ensure that children receive optimal healthcare support, to improve healthcare practices for all children (audit and quality improvement), for research purposes and to inform parents about prognostic elements regarding their preterm children.⁴⁻⁶ This is the first study investigating how parents rate their children's health and also their concerns and priorities: what they wished they could improve in an open-ended fashion (without an a priori). Parents rated their child's overall health as being high, despite disability, even if the presence and severity of NDI influenced parents' rating. In addition, parents did acknowledge the challenges faced by their children, mentioning several areas where improvements were desired.

These high ratings, despite the hardship of the NICU experience, overall fragility and, for some children, disability, potentially reflect the resilience of families and positive transformations.¹⁶ In other studies, parents reported a high quality of life for their children despite disabilities or described their child's happiness and good personality.¹⁷⁻¹⁸ These findings may indicate that for families, the well-being of their child, as opposed to disability, is what is truly important when health is evaluated.

Because we wanted to capture parents' subjective evaluation of their child, we did not use standardised health-related quality of life questionnaires. Other authors have previously reported the high parent-perceived and self-perceived quality of life ratings.¹⁷⁻¹⁹ In our cohort, perceived health remains high, but decreases with level of NDI, suggesting that neurodevelopmental outcomes do have an impact on parental subjective evaluations. In previous studies, authors have found that adults born premature rate their quality of life highly, independent of NDI. However, they are aware of their health and functional limitations.²⁰ Perhaps the parents in our study had a similar viewing of their child's health: lucidly assessing their child's function and what their children could do, and not only their diagnoses and limitations.

Even though parents generally rate their child's health with high scores, there is always room for improvement. This should be the basis for ongoing neonatal follow-up research as well as clinical research at large. In the clinical setting, health practitioners screen for difficulties early on, and the focus is mainly on child development. Development (especially behaviour and emotions) and respiratory health are the main areas parents wished they could improve. Some adverse outcomes considered to be mild by the medical literature, such as behavioural problems, hyperactivity and emotional health, were often considered as major problems by families. Whether these problems have more negative impacts on the child as well as the whole family, as opposed to deafness or using a walker due to cerebral palsy (classified as severe NDI), is a possibility. Interestingly, in terms of development, as children grew older, parents were more likely to express concerns about language and communication, socio-emotional problems and learning difficulties. Yet, standardised follow-up in most high-income countries often stops after 18–36 months CA, which might be too early to uncover more subtle

neurodevelopmental dysfunctions, such as those reported by parents, that are likely to interfere with school achievement.²¹ In a recent European survey, only 27% of very preterm children were receiving follow-up care at 5 years.²²

Furthermore, as previously shown, children of lower socioeconomic status were less likely to attend follow-up.²³⁻²⁴ Despite the association between lower socioeconomic status and neurodevelopmental vulnerabilities, especially in children at higher biological risk,²⁵ a lower level of parental education, in our study, was associated with parents explicitly or implicitly responding that nothing needed to be improved with their child. This could be interpreted as accepting the child as is, regardless of their challenges. This could also be associated with parental expectations as related to their own achievements, which may impact the value they place on educational achievements.²⁶ This raises the importance of engaging parents across educational levels to ensure proper developmental surveillance and screening so timely referral to intervention services is done when needed.

Our study also underlines the importance of reporting and investigating other outcomes, such as respiratory health and feeding/nutrition difficulties, as these are very important to many families, especially with younger children. These items should be systematically included and addressed in neonatal follow-up research, as they have a major impact on families' daily lives and function and for some families, they represent the main disability their children have.²⁷

This study did not have a comparison group of full-term children, whose parents would certainly also have concerns regarding health and development. However, we wished to examine the health and developmental outcomes which are of specific importance to parents of children born extremely premature. Also, we did not use standardised health-related quality of life questionnaires. This would have not allowed us to identify outcomes of interest to parents in their own words—what is most important to them—and can only be achieved by asking open-ended questions. Among parents of children who came for follow-up, the response rate was high (98%), but these responses represented a selected group. Given that preterm children attending follow-up are usually at higher biological risk of medical or developmental problems,²⁸ it is possible that we have underestimated the percentage of parents who actually did not wish to improve anything with their children. We are confident that parent and family important outcomes in this article represent the perspectives of parents in our centre, and possibly in places where children have access to universal healthcare and systematic follow-up. On the other hand, parent/family important outcomes may be different in centres where systematic follow-up or access to early intervention are not available.

CONCLUSION

Parents evaluate the health of their preterm children as being very good, with positive perspectives. Many express concerns outside the developmental sphere, such as medical fragility, nutrition and respiratory health. Many also express concerns about emotions and behaviour, which are not addressed nor categorised optimally in follow-up studies. Priorities differ according to the child's level of NDI, age and parental level of education. Incorporating measures reflecting what parents view as important can ensure that research findings are meaningful to families and that clinical follow-up targets relevant issues.

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Patient consent for publication Not applicable.

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