

Parental preferences for brace weaning in developmental dysplasia of the hip: a discrete choice experiment

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Aims

To elicit and quantify parental preferences for brace weaning strategies in the treatment of developmental dysplasia of the hip (DDH) and explore how parents trade between treatment burden and the risk of further intervention.

Methods

A discrete choice experiment (DCE) was developed to assess preferences for timing and duration of weaning, alongside trade-offs related to the risk of further treatment. Parents of infants treated for DDH were recruited via STEPS Worldwide, a patient charity. Parents completed 16 hypothetical scenarios comparing different weaning regimens with immediate cessation. Data were analyzed using conditional logit models. A secondary analysis excluded participants with irrational or disengaged responses. Subgroup analysis explored whether preferences varied by experience.

Results

A total of 195 respondents completed the survey. In the primary analysis, night-time brace wear was preferred over immediate cessation, even when risk remained equal. In the secondary analysis, which excluded internally inconsistent responses, all weaning strategies were significantly less preferred than immediate cessation under equal risk conditions. Parents were willing to accept longer durations of treatment in exchange for reduced risk of subsequent intervention. The minimum acceptable risk reduction required for parents to accept four additional weeks of brace wear, compared with immediate cessation, was 5.8% (night-time bracing), 7.5% (night-time and naps bracing), 8.4% (gradual brace reduction), and 10.0% (daytime bracing). Preferences varied by experience, especially age at diagnosis and prior weaning.

Conclusion

Night-time-only weaning was the most acceptable weaning strategy. Parents were willing to trade longer brace treatment for a lower risk of further intervention. We were able to quantify the size of the benefit required by families, which may inform research investigating the effectiveness of weaning strategies.

Take home message

- Parents prioritize reducing the risk of further treatment and are willing to accept additional brace wear (especially night-time-only use) if the perceived benefit is sufficient.

- These findings inform the design of future research.

Introduction

Developmental dysplasia of the hip (DDH) is the most common orthopaedic condition in infants, typically managed with bracing

during early life.¹⁻³ Early intervention with devices such as the Pavlik harness is generally effective, particularly when initiated within the first few months of life.⁴ However, there is ongoing debate regarding the necessity and design of a 'weaning' period when discontinuing brace use.

In the UK, 65% of clinicians immediately remove the brace, while the remaining 35% wean.⁵ While weaning is practiced by a minority of clinicians in the UK, it is the predominant practice internationally. Surveys indicate that two-thirds of members of the Paediatric Orthopaedic Society of North America (POSNA) and half of those from the European Paediatric Orthopaedic Society (EPOS) use some form of weaning regimen when treating infants.⁶ Consensus exercises have produced mixed results about the necessity for weaning. The International Hip Dysplasia Institute (IHDI) supports weaning with night-time use only,⁷ whereas the British Society for Children's Orthopaedic Surgery (BSCOS) has not reached a consensus.⁸ There is limited evidence to guide practice, with no randomized controlled trials and only two relatively small observational studies investigating the impact of weaning.⁹⁻¹¹

A discrete choice experiment (DCE) is a stated preference method used to understand how individuals make decisions when faced with competing options. In DDH, a DCE can explore how parents weigh different features of weaning regimens against the risk of requiring further treatment. Including a default option of immediate cessation with a constant risk level allows comparison between weaning strategies and no weaning. By quantifying these trade-offs, a DCE provides insights into parental preferences that can support shared decision-making and inform the design of future clinical trials evaluating weaning strategies.

This study aims to elicit parental preferences regarding weaning from brace treatment in DDH using a DCE. It quantifies the importance of weaning attributes, examines trade-offs with the risk of further intervention, identifies the risk reduction needed to justify weaning, and explores variation by demographic and clinical factors.

Methods

The study was conducted in accordance with established methodological guidelines for the implementation of DCEs.^{12,13} Ethical approval was awarded on 7 January 2025 by the University of Liverpool, UK (Reference: 15575). Informed consent was obtained from all participants prior to their inclusion in the study.

Attribute and level selection

Attributes and levels were identified using a mixed-methods approach, incorporating prior work with families and clinicians and supported by a literature review to ensure comprehensive coverage of current practice. Weaning regimens were characterized by two key attributes: the timing of brace wear during the day (night-time only, night-time and naps, gradual reduction in total hours per day or daytime only); and the duration of brace wear, ranging from two to 12 weeks. 'Hours per day', although relevant to reduction-based regimens, was not included as a separate attribute due to its overlap with timing of wear. Its inclusion risked generating illogical combinations and would have compromised the orthogonality of the design. The risk of further intervention (e.g. bracing

or surgery) was chosen as the 'trade-off', as it was more relevant to parents than surrogate radiological measures. The attributes and levels included in the DCE are presented in [Table 1](#).

Survey development and design

A D-efficient fractional factorial design was generated using Ngene software, producing 16 optimized choice tasks.¹⁴ Scenarios were defined by two explanatory attributes (timing and duration of brace wear) and one trade-off attribute (the risk of further intervention including further bracing or surgery). The design was developed in consultation with the STEPS Worldwide¹⁵ charity and a parent representative.

Each scenario presented three options: two weaning regimens and a default alternative, immediate cessation of bracing. The weaning regimens varied in timing, duration, and associated risk of further intervention. Immediate cessation was assigned a fixed 20% risk, based on the highest plausible estimate reported in the literature, and was consistently associated with the highest or a comparable risk relative to the weaning strategies.^{16,17}

Participants completed all 16 hypothetical scenarios, selecting the option they would prefer for their child:¹ a weaning regimen;² an alternative weaning regimen; or immediate cessation of bracing.³ An example choice task is shown in [Figure 1](#).

In addition to completing the choice tasks, parents were asked to provide demographic and clinical information, including their role (e.g. mother or father), the age of their infant at diagnosis, whether treatment was successful, the duration of full-time brace wear, and whether their child underwent a weaning regime.

Pre-testing

Three parents were recruited via the STEPS charity to undertake a 'think-aloud' pretest.^{18,19} Parents received instructions for completing a think-aloud exercise, followed by the standard survey instructions. They completed the survey while verbalizing their thoughts. Feedback indicated that the survey was clear and manageable. As no major difficulties were identified for families, the pretest responses were retained in the final dataset.

Sample and recruitment

Guidance on sample size calculation for healthcare-related DCEs is limited, though evidence suggests smaller samples can still yield valuable insights.^{20,21} We used Orme's sample size calculation: $N \geq (500 \times C) / (t \times a)$, where C is the maximum number of attribute levels, t is the number of choice tasks per respondent, and a is the number of alternatives per task (excluding the 'none' option).²² With $C = 6$, $t = 16$, and $a = 2$, this yielded a recommended sample size of 94.

Participants were recruited in collaboration with the STEPS charity through social media and their email list. The DCE was open to all parents of children treated for DDH, irrespective of the treatment received or outcome.

Participant characteristics

Recruitment was conducted between 10 February 2025 and 17 March 2025. A total of 195 respondents completed the full

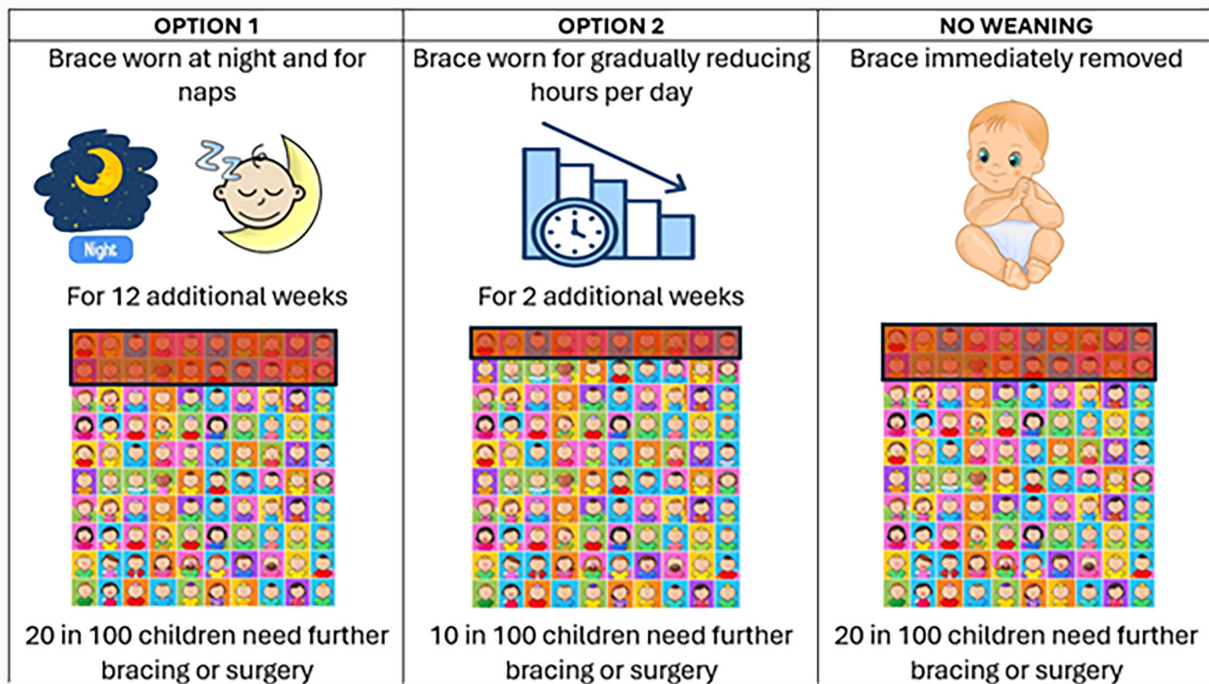


Fig. 1
An example choice task.

Table I. The attributes and levels included in the discrete choice experiment.

Attributes	Levels
Explanatory: timing of brace wear	Night-time only
	Night-time and naps
	Gradual reduction in total number of hours worn per day
	Daytime only
Explanatory: length of weaning regime, wks	2
	4
	6
	8
	10
	12
Trade-off: risk of requiring further intervention, %	5
	10
	15
	20

survey. Demographic and clinical characteristics are summarized in Table II. The primary analysis included all completed responses. To quantify parental preferences, a conditional logit model was used.

Statistical analysis

Responses were analyzed using a conditional logit model in Stata v. 18.0 (StataCorp, USA),²³ which estimates the probability of choosing an alternative based on the utility derived from its attributes. Grounded in random utility theory, the

model assumes that participants select the option offering the highest utility in each choice task.²⁴

This model was used to estimate preferences for individual weaning attributes and to quantify how participants weighed these preferences against the associated risk of requiring further intervention (e.g. bracing or surgery). The primary analysis included all responses. A secondary analysis was conducted in which participants who provided at least one 'irrational' response were excluded. An 'irrational' response was defined as selecting a weaning strategy over immediate cessation when both options carried the same level of risk. Participants who consistently selected the same weaning strategy across all choice tasks, a behaviour known as straightlining, were also excluded, as this pattern suggests a lack of engagement with the survey. In contrast, participants who consistently chose the opt-out alternative were retained, as this was considered a rational and intentional expression of preference.²⁵ The secondary analysis was used to estimate marginal rates of substitution (MRS) between attributes to evaluate how participants traded increased treatment burden, defined as additional weeks in a brace and the timing of wear, against reduced risk of further intervention. MRS values were calculated as the ratio of the relevant attribute coefficients. To contextualize these trade-offs, the total disutility of each weaning strategy over a four-week period was calculated by summing the attribute-specific coefficient with the disutility from four weeks of brace wear. Total disutilities were divided by the risk coefficient to estimate the percentage risk reduction required for each strategy to be equally acceptable to immediate cessation. Subgroup analysis was undertaken to assess whether late diagnosis, unsuccessful brace treatment, or weaning affected results. Statistical significance was set at $p < 0.05$.

Table II. Participant and infant characteristics.

Variable	N (%)
Participant role	
Mother	191 (97.9)
Father	4 (2.1)
Participant country	
UK	95 (48.7)
USA	58 (29.7)
Australia	17 (8.7)
Canada	11 (5.6)
Ireland	6 (3.1)
New Zealand	3 (1.5)
Chile, Croatia, Netherlands, Philippines, Portugal*	5 (2.5)
Care of infant, related to the participant completing the form	
Infant age at diagnosis, mnths	
≤ 3	162 (83.1)
4 to 6	20 (10.3)
7 to 9	6 (3.1)
10 to 12	7 (3.6)
Duration of treatment, wks	
≤ 6	27 (13.8)
7 to 12	72 (36.9)
> 12	61 (31.3)
Ongoing/interrupted	35 (17.9)
Weaning practices	
Ongoing full-time brace use	18 (9.2)
Completed a weaning regime	70 (35.9)
Immediate brace cessation	96 (49.2)
Early termination for alternative treatment	11 (5.6)
Treatment outcome	
Completed brace treatment without further intervention	126 (64.6)
Required further treatment post-bracing	46 (23.6)
Ongoing brace treatment	23 (11.8)

*One respondent each from Chile, Croatia, Netherlands, Philippines, and Portugal.

Results

Table III presents the estimated coefficients, standard errors, and p-values. In this analysis, coefficients represent the relative strength and direction of preference for each attribute level compared with immediate cessation, which was fixed at zero. Positive coefficients indicate a higher likelihood of selection compared with immediate cessation, whereas negative coefficients indicate a lower likelihood.

Participants' preferences for weaning strategies were assessed relative to immediate cessation, holding the risk of

further intervention constant. The most preferred strategy was night-time-only brace wear, followed by night-time and naps, gradual reduction, and daytime-only wear, which was the least preferred. Night-time-only brace wear was associated with a statistically significant preference (coefficient 0.432, standard error 0.090; $p < 0.001$). The preference for night-time and naps was positive but not statistically significant (coefficient 0.152; $p = 0.119$). Gradual reduction and daytime-only brace wear were both significantly less preferred than immediate cessation ($p < 0.001$ and $p = 0.008$, respectively).

Participants also showed a strong aversion to increasing the risk of further intervention (coefficient -0.189 per 1% increase; $p < 0.001$) and to longer weaning durations (coefficient -0.099 per additional week; $p < 0.001$), indicating that both higher risk and longer treatment duration reduced the likelihood of a weaning strategy being chosen.

A secondary analysis was conducted excluding participants who provided at least one irrational response or who displayed straightlining behaviour. In this analysis ($n = 90$), all weaning strategies were significantly less preferred than immediate cessation. Table IV presents the estimated coefficients, standard errors, and p-values from the conditional logit model. The most preferred weaning strategy relative to immediate cessation was night-time brace wear only ($p < 0.001$), followed by night-time and naps ($p < 0.001$), gradual reduction ($p < 0.001$), and daytime brace wear, which was the least preferred ($p < 0.001$).

As in the primary analysis, participants demonstrated a strong aversion to higher risk ($p < 0.001$) and longer treatment duration ($p < 0.001$), both of which reduced the likelihood of selecting a weaning strategy.

Parents demonstrated a willingness to accept increased treatment burden in exchange for reduced risk of further intervention. On average, respondents were willing to accept 1.6 additional weeks of bracing to achieve a 1% reduction in risk.

The values presented in Table V show the total disutility associated with each weaning strategy applied over a four-week period, calculated by combining the attribute-specific coefficient with the disutility of four weeks of brace wear. The corresponding risk reduction values indicate the percentage decrease in risk of further intervention required to make each strategy equally acceptable to immediate cessation. Based on the overall utility of night-time brace wear, a reduction in risk of approximately 5.8% would be required to make four weeks of night-time bracing equally acceptable to immediate cessation.

Subgroup analyses explored whether preferences for weaning strategies varied according to personal experience. Among participants whose infants were diagnosed after aged six months, lower coefficients were seen for all weaning strategies compared with the overall sample, indicating reduced acceptability of continued bracing in this group. Respondents whose infants had required further intervention following initial brace treatment showed similar preference patterns to the full cohort, with coefficients of comparable direction and magnitude. Differences were observed according to reported weaning experience. Participants who had weaned their infants demonstrated greater acceptance of night-time brace wear, compared with those who had stopped bracing immediately. In the weaning group, night-time brace

Table III. Primary analysis estimated coefficients, standard errors, and p-values.

Attribute	Coefficient	Standard error	p-value*
Night-time brace wearing	0.432	0.090	< 0.001
Night-time and naps brace wearing	0.152	0.098	0.119
Gradual reduction in brace wearing	-0.278	0.104	< 0.001
Daytime brace wearing	-0.611	0.107	0.008
Risk of requiring further intervention	-0.189	0.006	< 0.001
Length of weaning regime	-0.099	0.007	< 0.001

*Conditional logit model in Stata v. 18.0.

Table V. Estimated total disutility and required risk reduction for each weaning attribute (secondary analysis).

Weaning attributes	Total disutility	Risk reduction needed, %
Night-time brace wearing	-1.508	5.8
Night-time and naps brace wearing	-1.972	7.5
Gradual reduction in brace wearing	-2.195	8.4
Daytime brace wearing	-2.621	10.0

wear was not statistically significantly different from the opt-out alternative ($p = 0.468$), while in the non-weaning group, all weaning strategies remained significantly less preferred than cessation. These findings suggest that while the direction of preferences was largely consistent across subgroups, the strength of preferences was moderated by clinical context, particularly age at diagnosis and prior experience of weaning.

Discussion

In the primary analysis, night-time brace wear emerged as the most preferred weaning strategy, followed by night-time and naps, gradual reduction, and finally daytime wear. When the risk of further intervention was held constant, the overall cohort showed a preference for night-time and night-time-and-naps weaning strategies over immediate cessation, despite no associated reduction in risk. At face value, this response may appear illogical, as it involves continued brace use without additional clinical benefit. However, literature on stated preference methods consistently highlights that such responses are often not truly illogical; rather, they reflect underlying values or interpretations that are not immediately apparent from the choice data alone.²⁶⁻²⁸ One possible explanation is that parents may perceive a gradual transition out of bracing as less abrupt and more manageable than immediate cessation. This may provide a sense of control or emotional reassurance during what can be a stressful period. However, this interpretation is somewhat at odds with existing qualitative literature, which consistently emphasizes

Table IV. Secondary analysis estimated coefficients, standard errors, and p-values.

Attribute	Coefficient	Standard error	p-value*
Night-time brace wearing	-0.844	0.140	< 0.001
Night-time and naps brace wearing	-1.308	0.162	< 0.001
Gradual reduction in brace wearing	-1.531	0.167	< 0.001
Daytime brace wearing	-1.957	0.183	< 0.001
Risk	-0.262	0.011	< 0.001
Weeks	-0.166	0.013	< 0.001

*Conditional logit model in Stata v. 18.0.

the burden of brace wear on families.^{29,30} Further qualitative research would be valuable to understand whether weaning is perceived by some parents as a more acceptable or comforting approach, despite the additional treatment burden.

A second explanation may relate to how participants interpreted the choice tasks. Although the survey was pretested and found to be understandable, it is well-established that risk is a complex and often misunderstood concept in decision-making.³¹⁻³³ In this study, immediate cessation was associated with a fixed 20% risk of further intervention as the highest plausible estimate based on existing literature. Participants were therefore frequently presented with choices between options that all carried relatively high risk. In this context, some parents may have reasoned that limited continued bracing (such as night-time use) might offer even a small chance of benefit and therefore was worth attempting despite the fact that no actual reduction in risk was presented in those scenarios. This is consistent with the strong risk aversion observed throughout the DCE, where lower-risk options were consistently preferred, even when they involved greater treatment burden.

To address the potential influence of disengaged or irrational responses, a secondary analysis was conducted excluding participants who made at least one illogical choice or showed patterns consistent with straightlining. The findings remained consistent with the primary analysis in terms of the direction of preferences, though effect sizes were larger and all weaning strategies were significantly less preferred than immediate cessation. This refined model likely reflects improved data quality and highlights the importance of careful respondent screening in stated preference research. It could be perceived that the secondary analysis simply reflects the removal of participants who expressed a preference for weaning over immediate cessation. However, exclusions were based on predefined criteria of internal inconsistency, specifically selecting a more burdensome option over an equally risky but less burdensome alternative, rather than on stated preferences alone. This practice is well-supported in the stated preference literature as a means of enhancing the internal validity of model estimates.^{26,34} Moreover, the consistency in the direction of preferences across both analyses suggests that the refined model served to clarify,

rather than distort, the underlying decision-making patterns observed in the full sample.

The secondary analysis also enabled the estimation of trade-offs between treatment burden and risk, offering insight into how parents value additional bracing relative to reductions in the likelihood of further intervention. These trade-offs provide a useful framework for informing future clinical trial design, including the selection of outcome thresholds that reflect differences considered meaningful by parents. In particular, the quantified value parents place on risk reduction can be used to inform sample size calculations and the selection of non-inferiority or superiority margins that are both statistically and clinically relevant. Incorporating parent-derived thresholds in trial planning ensures that outcomes are not only statistically robust but also relevant to those directly affected by treatment decisions.

Subgroup analyses suggested that preferences were moderated by clinical experience. Parents of infants diagnosed after six months of age demonstrated consistently lower utility for all weaning strategies. Those whose infants required further treatment showed similar preferences to the overall sample, whereas parents who had experienced a weaning regime showed more favourable views of night-time weaning, suggesting their preferences may have been shaped by familiarity or perceived benefit.

Although clinician perspectives are undoubtedly important in the wider debate on brace weaning, their inclusion in this discrete choice experiment was intentionally avoided. Parents act as proxy decision-makers for infants and ultimately determine participation and adherence; their willingness to trade treatment burden against reductions in risk was therefore the primary focus. Incorporating clinicians would have risked conflating distinct perspectives, as attributes would need to be framed in ways meaningful to both groups, potentially reducing validity for each. By restricting the sample to caregivers, we were able to capture parental risk tolerance in a clear and undiluted manner. Clinician views on brace weaning have been explored separately.³⁵

While the overall sample size met recommended thresholds for DCE analysis, subgroup analyses were based on smaller numbers and should be interpreted with caution. The study sample was predominantly composed of mothers and largely represented families from the UK and other high-income western countries, and as such, the findings reflect the views and preferences of this demographic. Participants had varied experiences of DDH treatment, including differing treatment durations, outcomes, and approaches to brace weaning. The sample was also skewed toward highly engaged respondents, many of whom were recruited via a patient charity. This may limit the generalizability of findings to broader populations.

Additionally, as with all stated preference methods, hypothetical bias remains a concern, as responses may not fully reflect real-world decision-making.

Nonetheless, this study makes a significant contribution to a sparsely researched area. Brace weaning remains highly variable in clinical practice, with no consensus reached on its necessity or optimal design. By incorporating the voice of parents, those directly affected by treatment decisions, this study provides a foundation for patient-centred care.

Parents prioritized reducing the risk of further intervention and were willing to accept additional brace wear, particularly night-time-only use, if the perceived benefit was sufficient. These findings offer valuable insights to inform the design of future clinical trials and ensure they reflect family priorities.

Social media

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References

1. **Jacobsen KK, Laborie LB, Kristiansen H, et al.** Genetics of hip dysplasia - a systematic literature review. *BMC Musculoskelet Disord.* 2024;25(1):762.
2. **Burt JEA, AlKandari N, Campbell DM, MacLean JGB.** Who performs neonatal hip assessment: is there a cause for concern? *BMJ Paediatr Open.* 2024;8(1):e002490.
3. **Sioutis S, Kolovos S, Papakonstantinou ME, Reppas L, Koulalis D, Mavrogenis AF.** Developmental dysplasia of the hip: a review. *J Long Term Eff Med Implants.* 2022;32(3):39–56.
4. **Mulpuri K, Song KM, Goldberg MJ, Sevarino K.** Detection and nonoperative management of pediatric developmental dysplasia of the hip in infants up to six months of age. *J Am Acad Orthop Surg.* 2015;23(3):202–205.
5. **Westacott DJ, Perry DC.** The treatment of neonatal hip dysplasia with splints in the United Kingdom: time for consensus? *J Child Orthop.* 2020;14(2):112–117.
6. **Alves C, Truong WH, Thompson MV, et al.** Diagnostic and treatment preferences for developmental dysplasia of the hip: a survey of EPOS and POSNA members. *J Child Orthop.* 2018;12(3):236–244.
7. **Kelley SP, Feeney MM, Maddock CL, Murnaghan ML, Bradley CS.** Expert-based consensus on the principles of Pavlik harness management of developmental dysplasia of the hip. *JB JS Open Access.* 2019;4(4):e0054.
8. **Aarvold A, Perry DC, Mavrotas J, Theologis T, Katchburian M, BSCOS DDH Consensus Group.** The management of developmental dysplasia of the hip in children aged under three months: a consensus study from the British Society for Children's Orthopaedic Surgery. *Bone Joint J.* 2023;105-B(2):209–214.
9. **Dwan K, Kirkham J, Paton RW, Morley E, Newton AW, Perry DC.** Splinting for the non-operative management of developmental dysplasia of the hip (DDH) in children under six months of age. *Cochrane Database Syst Rev.* 2022;10(10):CD012717.
10. **Westacott DJ, Mackay ND, Waton A, Webb MSL, Henman P, Cooke SJ.** Staged weaning versus immediate cessation of Pavlik harness treatment for developmental dysplasia of the hip. *J Pediatr Orthop B.* 2014;23(2):103–106.
11. **Bram JT, Gohel S, Castañeda PG, Sankar WN.** Is there a benefit to weaning pavlik harness treatment in infantile DDH? *J Pediatr Orthop.* 2021;41(3):143–148.
12. **Weber S.** A Step-by-Step Procedure to Implement Discrete Choice Experiments in Qualtrics. *Soc Sci Comput Rev.* 2021;39(5):903–921.
13. **Ryan M, Gerard K.** Using discrete choice experiments to value health care programmes: current practice and future research reflections. *Appl Health Econ Health Policy.* 2003;2(1):55–64.
14. **No authors listed.** ChoiceMetrics. <https://www.choice-metrics.com> (date last accessed 24 November 2025).
15. **No authors listed.** STEPS Worldwide. <https://www.stepsworldwide.org/> (date last accessed 26 November 2025).
16. **Badrinath R, Orner C, Bomar JD, Upasani VV.** Narrative review of complications following DDH treatment. *JOJO.* 2021;55(6):1490–1502.
17. **Upasani VV, Bomar JD, Matheney TH, et al.** Evaluation of brace treatment for infant hip dislocation in a prospective cohort: defining the success rate and variables associated with failure. *J Bone Joint Surg Am.* 2016;98-A(14):1215–1221.
18. **Eccles DW, Arsal G.** The think aloud method: what is it and how do I use it? *Qual Res Sport Exerc Health.* 2017;9(4):514–531.
19. **Noushad B, Gerven P, de BA.** Twelve tips for applying the think aloud method to capture cognitive processes. *Med Teach.*

<https://www.tandfonline.com/doi/abs/10.1080/0142159X.2023.2289847> (date last accessed 24 November 2025).

20. Mannion L, Watson V, Mullassery V, et al. Treatment preferences of patients with muscle invasive bladder cancer: a discrete choice experiment. *BJUI Compass*. 2024;5(11):1059–1068.
21. Mangham LJ, Hanson K, McPake B. How to do (or not to do) ... Designing a discrete choice experiment for application in a low-income country. *Health Policy Plan*. 2009;24(2):151–158.
22. Orme B. Sample size issues for conjoint analysis studies. Sawtooth Software. 1998. <https://sawtoothsoftware.com/resources/technical-papers/sample-size-issues-for-conjoint-analysis-studies> (date last accessed 25 November 2025).
23. No authors listed. Stata: statistical software for data science. 2025. <https://www.stata.com/> (date last accessed 24 November 2025).
24. No authors listed. Random utility theory: an overview. ScienceDirect Topics. <https://www.sciencedirect.com/topics/economics-econometrics-and-finance/random-utility-theory> (date last accessed 24 November 2025).
25. Johnson FR, Yang JC, Reed SD. The internal validity of discrete choice experiment data: a testing tool for quantitative assessments. *Value Health*. 2019;22(2):157–160.
26. Lancsar E, Louviere J. Deleting “irrational” responses from discrete choice experiments: a case of investigating or imposing preferences? *Health Econ*. 2006;15(8):797–811.
27. Miguel FS, Ryan M, Amaya-Amaya M. “Irrational” stated preferences: a quantitative and qualitative investigation. *Health Econ*. 2005;14(3):307–322.
28. Ryan M, Watson V, Entwistle V. Rationalising the “irrational”: a think aloud study of discrete choice experiment responses. *Health Econ*. 2009;18(3):321–336.
29. Theunissen W, van der Steen MC, van Veen MR, van Douveren F, Witlox MA, Tolk JJ. Parental experiences of children with developmental dysplasia of the hip: a qualitative study. *BMJ Open*. 2022;12(9):e062585.
30. Poole C. Exploring the experiences of parents caring for their infant with developmental dysplasia of the hip (DDH): an interpretative phenomenological analysis. Edinburgh Napier University, 2019.
31. Lipkus IM, Peters E. Understanding the role of numeracy in health: proposed theoretical framework and practical insights. *Health Educ Behav*. 2009;36(6):1065–1081.
32. Wijohn TR, Newcombe RM, Reynolds J, El-Jack S, Armstrong GP. Informed consent-patients’ understanding of risk. *N Z Med J*. 2024;137(1590):14–21.
33. Barnes AJ, Hanoch Y, Miron-Shatz T, Ozanne EM. Tailoring risk communication to improve comprehension: Do patient preferences help or hurt? *Health Psychol*. 2016;35(9):1007–1016.
34. de Bekker-Grob EW, Ryan M, Gerard K. Discrete choice experiments in health economics: a review of the literature. *Health Econ*. 2012;21(2):145–172.
35. Craven J, Davies I, Perry DC. The role of weaning in brace treatment for developmental dysplasia of the hip: time to define best practice? *Bone Jt Open*. 2025;6(6):685–690.

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The data that support the findings for this study are available to other researchers from the corresponding author upon reasonable request.

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