

Systematic Review

Factors related to motor outcomes in adolescents and adults with cerebral palsy: A systematic review

Rayane Félix Lôbo Monteiro^{a,*}, Camila Araújo Santos Santana^a,
Paula Silva de Carvalho Chagas^b, Ana Carolina de Campos^a

^a Universidade Federal de São Carlos, São Carlos, SP, Brazil

^b Universidade Federal de Juiz de Fora, Juiz de Fora, MG, Brazil



ARTICLE INFO

Keywords:

Cerebral palsy

Motor decline

Physical therapy

Transition to Adult Care

ABSTRACT

Background: Although the brain injury caused by cerebral palsy (CP) is not progressive, there is often a worsening of physical conditions during the transition to adolescence and adulthood. The aim of this systematic review was to summarize factors related to decline in motor function in adolescents and adults with CP.

Methods: This review was registered in PROSPERO (CRD42022311783). Studies met the following criteria: participants diagnosed with CP, ages 10 to 59 years; scientific studies with any methodological design and emphasis on motor function, reporting quantitative analysis of the effect of age on motor function or factors relevant that declined or improved motor capacity; published in English; any publication year, in electronic databases: Pubmed/Medline, Scopus, Cinahl. Data selection and extraction used the Covidence software. Factors related to motor outcomes that were statistically significant in at least one study and present in more than one study were reported. The outcomes were grouped within major categories (gross motor function, neuromusculoskeletal functions, and gait) for summary.

Results: 23 studies met the inclusion criteria. Relevant factors for changes in gross motor function (n = 16 studies) were: pain, neuromusculoskeletal aspects, altered gait parameters, Gross Motor Function Classification System (GMFCS) level, age, and history of surgery; for gait (n = 11): mobility performance, altered gait parameters, age, GMFCS level, pain and neuromusculoskeletal aspects; and for neuromusculoskeletal functions (n = 4): altered gait parameters, neuromusculoskeletal aspects, age, pain, and GMFCS level.

Conclusion: Age, GMFCS level, and presence of pain stood out as significant for motor decline across categories. Monitoring these factors is relevant for planning interventions and transition programs.

Introduction

The life expectancy of individuals with cerebral palsy (CP) has been catching up with that of the general population as a result of the increasing survival rate of low and very low birth weight infants and also due to advances in neonatal health care.^{1,2} Improvements in health care have also led to a higher occurrence of individuals with CP going through the transition from childhood to adolescence and adult life, thus creating new challenges for healthcare planning.^{3,4}

Although the brain injury caused by CP is not progressive, it is known that worsening of physical conditions often occurs during the transition to adolescence and adult life,⁵ including decreased mobility and increased impairments in body structures and functions.^{5,6} A previous review showed that gait deterioration generally occurs in early

adulthood, and that 25 % or more of young adults with CP might experience a decline or gait deterioration.⁷ The topography of motor impairment, the presence of comorbidities, secondary musculoskeletal issues, and increased frequency and intensity of pain have been related to gait decline.⁷ Pain is considered the most common comorbidity among individuals with CP, and is often associated with reduced quality of life and well-being, with an increase in intensity during the transition to adulthood.⁸

Even though the decline in gait and associated factors have been previously explored,⁷ no reviews were found investigating motor decline more broadly, i.e., not only focusing on gait. This is relevant considering that a significant portion of individuals with CP do not walk and may be at risk for decline in their other motor functions. There is also a lack of studies summarizing the factors that should be monitored

* Corresponding author.

E-mail address: rayanemonteiro@estudante.ufscar.br (R. Félix Lôbo Monteiro).

<https://doi.org/10.1016/j.bjpt.2025.101225>

Received 16 April 2024; Received in revised form 9 December 2024; Accepted 28 April 2025

Available online 31 May 2025

1413-3555/© 2025 Associação Brasileira de Pesquisa e Pós-Graduação em Fisioterapia. Published by Elsevier España, S.L.U. All rights are reserved, including those for text and data mining, AI training, and similar technologies.

or addressed during the development of individuals with CP to prevent or manage motor decline. This information is crucial to support health-care practices and to guide future research about health development of young people in transition to adult life.^{3,9}

Thus, this systematic review aims to identify factors related to overall motor outcomes in adolescents and adults with CP.

Methods

Study design

Review procedures were based on recommendations from the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) Guidelines.¹⁰ The study question was based on the PECO strategy: Participants: Adolescents and adults with CP; Exposure: exposure to any factors shown to influence motor outcomes; Comparator: non-applicable; Outcome: any motor-related outcomes. The research question was defined as: Which factors are relevant for the motor outcomes of adolescents and adults with cerebral palsy?

This systematic literature review was registered in the International Prospective Registries of Systematic Reviews (PROSPERO): registration number CRD42022311783.

Eligibility criteria

Inclusion and exclusion criteria for the studies are presented in Table 1.

Information sources and search strategy

The search string comprised five concepts: Cerebral Palsy; Musculoskeletal development; Mobility limitation; Musculoskeletal system; and Transition to adult care. A systematic literature search was performed in three electronic databases: Pubmed/Medline, Scopus, Cinahl, including only studies in English, covering all years of publication until June 2023, with adjustments on the search string being made for each database. A manual search was also performed on the reference lists of the included articles to extract additional studies. The last search update was made in June 2023. [Supplementary material 1](#) provides an example of a search strategy from one of the databases.

Data collection and analysis

The Covidence software was used to manage the data. Studies were screened based on titles and abstracts, followed by reading of full texts.

Table 1
Inclusion and exclusion criteria.

Inclusion criteria	Exclusion criteria
Participants diagnosed with CP	Articles that evaluated other populations (participants under 10 years of age or 60 years and above exclusively), or including adolescents or adults with disabilities other than CP
Age: 10 to 59 years	Publications that were not research articles, such as abstracts, book chapters, theses, expert opinions, among others
Initial entry in the study could occur at a younger age	Studies that did not quantitatively analyze risk factors for motor decline
Studies analyzing the effects of age on motor function or analyzing risk factors leading to decline in functional capacity, with emphasis on motor function	
Published in English	
Any year of publication;	
Research articles	
With any methodological design, as long as it included a quantitative analysis of risk factors related to motor decline	

Study selection and data extraction were performed independently by two reviewers, and a third reviewer solved conflicts.

A data collection form developed by the research team and previously tested in the first studies was used to establish the consistency of the data with the study objective. The following data were extracted: (1) Characteristics of the study; (2) Characteristics of the participants; (3) Measurement tools and motor-related outcomes. The factor was considered relevant and included in the review when it was identified as statistically significant in at least one study that had a low to moderate risk of bias based on the checklist of choice for this study (description presented next) and if the same outcome was identified as statistically significant in at least two studies with a moderate to high risk of bias.¹¹

Even though the study initially focused on risk factors, protective factors for motor decline were also summarized if they met the criteria to be considered relevant factors. For easier presentation, the factors affecting motor outcomes were analyzed within the following major categories identified during data summary: 1) Gross motor function (e.g., measured by the Gross Motor Function Measure [GMFM-66] or Gross Motor Function Classification System [GMFCS]); 2) Gait; and 3) Neuromusculoskeletal functions (here defined as those related to the functions of joints, bones, reflexes, and muscles,¹² e.g., decrease in muscle strength or joint range of motion [(ROM])).

Risk of bias assessment

The checklist to assess study bias was scored based on the clear description of several relevant items, assigning 1 point for studies reporting the item and 0 for those that did not clearly report this item. The maximum score for each study is 16 points. Studies with scores between 12 and 16 were considered to have few methodological limitations, being classified as having good quality; scores between 7 and 11 points indicate moderate methodological limitations, with the study being classified as fair; a score below 7 points represents significant methodological limitations, and thus the study quality was classified as poor.^{13,14}

The heterogeneity of the outcomes did not allow meta analyses, therefore the results were analyzed descriptively. The results were summarized based on the factors related to each category of motor-related outcomes.

Results

The search resulted in 1691 articles, of which 23 met all inclusion criteria and were included in the review and proceeding with data extraction (Fig. 1).

As shown in Table 2, the most frequent study design was longitudinal (n = 11); followed by cross-sectional (n = 4); cohort (n = 4); observational (n = 1); and retrospective (n = 1). The countries of the selected studies were: United States (n = 8); Netherlands (n = 3); Canada (n = 3), Australia (n = 2), and Taiwan (n = 2). Norway, United Kingdom, and Japan presented one study each. The sample size ranged from 25 to 13,271 participants. The age of the participants ranged from 3 to 41 years, GMFCS levels were I-II (n = 20 studies), III (n = 19); IV (n = 18), and V (n = 17). Six studies did not report the GMFCS level. Participants' motor impairment was unilateral in 17 studies and bilateral in 30 studies. In total, 33 factors were identified in the studies, seven of which were considered relevant.

Table 3 presents the relevant factors influencing motor outcomes identified in this review.

Regarding gross motor function, factors relevant were identified in 16 studies and included: age, GMFCS level, neuromusculoskeletal aspects, pain, history of surgeries, and altered gait parameters. History of orthopedic surgeries was identified as a protective factor for gross motor function in three studies. For gait deterioration, 11 studies identified more severe GMFCS level, advancing age, pain, neuromusculoskeletal aspects (muscle strength, ROM, worsening of hip condition among

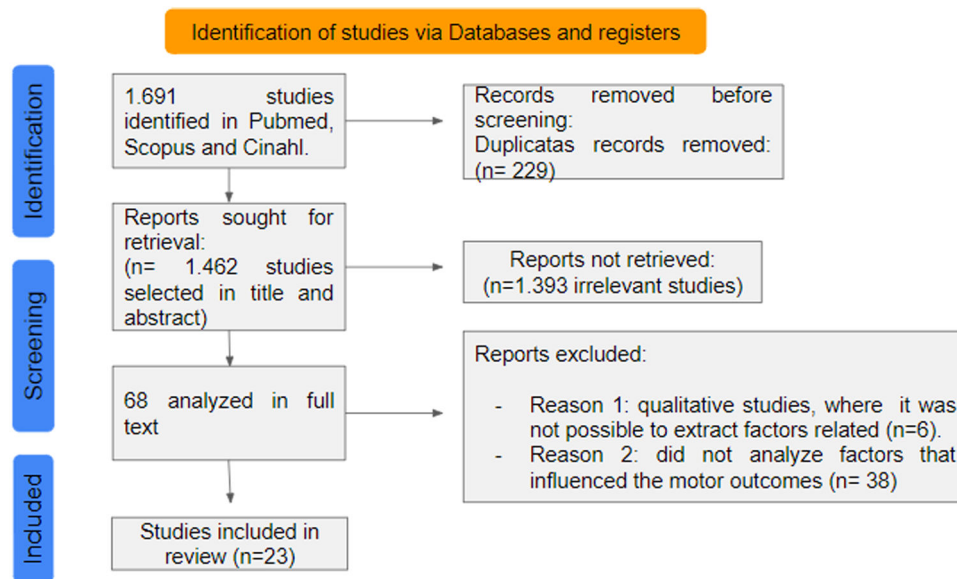


Fig. 1. PRISMA selection flow diagram.

others), altered gait parameters, and mobility performance as relevant factors. Regarding neuromusculoskeletal functions, four studies indicated factors relevant to decline such as pain, abnormal gait parameters, and increasing age (Table 3).

When assessing risk of bias in the included studies, all studies were classified as low risk (Table 4).

Additional (non-motor) outcomes demonstrating functional decline over time included self-care skills ($n = 1$), health-related quality of life ($n = 1$), and performance and satisfaction with life habits ($n = 1$), however they were infrequently reported. Other factors related to motor outcomes but not classified as significant as they did not reach the criteria established in this review included depression, emotional support,¹⁵ mental health,¹⁶ housing,¹⁷ balance,¹⁸ fatigue,¹⁵ and CP type.¹⁹

Discussion

This review aimed to identify factors relevant for motor decline in adolescents and adults with CP. Factors such as pain, mobility performance, changes in gait parameters, GMFCS level, age, history of surgery, and neuromusculoskeletal aspects were the most recurrent and relevant factors reported in the literature.

Most studies in the review were conducted in high-income countries, which has been highlighted in other reviews on individuals with CP.^{20,21} This result may also reflect larger awareness and resources directed towards research on the period of transition to adult life in these countries, and highlights the need for information on factors relevant for motor decline in low- and middle-income countries.

The most frequent methodological design was longitudinal, suitable for monitoring changes in functioning over time and obtaining knowledge about the development of these individuals.^{22,23} Of the 11 longitudinal studies, the majority included adolescents, followed by young adults. However, the total amount of studies underscores the need for more research to understand the determinants of motor changes in individuals during the transition to adulthood, as well as the impact of motor decline on broader aspects of functioning, including participation and quality of life.²²

Regarding the characterization of participants, bilateral CP was the most frequently reported subtype among the included studies, which may reflect epidemiological trends.²⁴ Most studies included individuals with CP at GMFCS levels I to III, which may be related to the fact that they are independent walkers or can walk with assistive devices, and given that gait is a common concern. Similar findings were also observed

by Furtado et al.,²⁵ reporting a higher frequency of studies on children with CP with GMFCS levels I to III. Although individuals with GMFCS levels IV and V are the most vulnerable to motor decline over the years,²⁵ literature is still scarce on motor changes after childhood. In the following sections, we will discuss the evidence on the main factors identified.

Decline in gross motor function

Sixteen studies identified factors relevant that resulted in gross motor function decline. The decline in gross motor function in individuals with CP has been consistently described through GMFM, both in overall and in standing scores over development.^{26,27} This may be a relevant tool to follow-up youth with CP across transition, however validity for adults needs to be demonstrated. A prominent relevant factor that may contribute to the decrease in gross motor scores is the presence of pain.^{15,28,29} Pain is therefore a factor that should not be neglected by healthcare professionals working with individuals with CP as it may precede relevant functional losses.

As expected, results highlighted age as a frequent factor relevant for the decline in gross motor function in adolescents and adults with CP, along with GMFCS levels.²⁹⁻³⁶ The combination of age and GMFCS predicted Pediatric Evaluation of Disability Inventory (PEDI) mobility scores, with higher development and longer period in development for individuals with mild to moderate GMFCS levels (I-III), compared to those at more severe GMFCS levels.³⁵ Similarly, these factors predicted earlier peak GMFM scores and significant declines in adolescence and adulthood in those between levels III to V.^{31,36} Several other studies supported the influence of the GMFCS level as a factor relevant for the decline in gross motor function in individuals with CP in the first decade of life and in adults, with results similar to those previously discussed.^{15,30,31,33,34,36-39} Collectively, these findings underscore the vulnerability of non-ambulant individuals to early motor decline.

Musculoskeletal conditions were also pointed out as relevant factors for gross motor change. These included muscle strength,³⁰ ROM in the upper limbs and in hip, knee, and ankle joints, spinal alignment,²⁸ and presence of stiffness and deformities.²⁹ Such findings point to the relevance of monitoring and intervening early in life on these factors because interventions addressing these issues may be strategies to prevent motor decline. Among the interventions targeting musculoskeletal aspects, single-event multilevel surgery (SEMLS) has been suggested to favor stability of GMFCS levels.³⁸

Table 2
Characteristics of the included studies.

Author/ Year	Country of origin	Study design	Participants		Outcome measures	Factors associated with motor outcomes evaluated *
			Type of CP/ number/ Levels of GMFCS	Age/Sex (F/M)		
Baird et al., 2021	UK	Cohort design or non-randomized group	CP (Spastic, Dystonic, Dyskinetic); n: 338; GMFCS: I, II, III, IV and V	8 years; (137–201)	GMFCS, FMS, Pain Questionnaire, KIDSCREEN	Pain, Neuromusculoskeletal aspects (spine and lower limbs).
Bartlett et al., 2010	Canada	Longitudinal	CP (Unilateral, Bilateral, Ataxic); n: 135; GMFCS: III, IV and V	14 years; (59–76).	SAROMM, GMFM-66, GMFCS, Pain Questionnaire, HUI-3, Self-administered physical activity questionnaire	Pain, Neuromusculoskeletal aspects (range of motion and spinal alignment); Anthropometric variables (increase in triceps skinfold, mid-arm circumference, and the relationship between arm circumference and knee height). Age, Pain, Neuromusculoskeletal aspects.
Bell et al., 2002	U.S	Longitudinal	CP (Unilateral, Bilateral); n: 48; GC: 28.	CP: 7.8 years; GC: 10.1 years; (18–10).	Gait analysis, clinical examination (passive ROM, hand muscle strength, estimated torsion measures, EMG	
Buckland et al., 2021	U.S	Retrospective review	CP; n: 84; GMFCS: IV and V	13.6 years	Clinical data (surgical and radiographic information), CPCCHILD, hip status	Neuromusculoskeletal aspects (lumbar hyperlordosis).
Church et al., 2018	U.S	Longitudinal	CP + knee surgery (Unilateral, Bilateral); n: 59; GMFCS: I, II, III and IV	20 years.	GDI, GMFM-D, GMFCS, PROMIS	GMFCS, Gait changes (kinematic variables).
Chiu et al., 2019	Taiwan	Systematic review	CP; Hemiplegic, Diplegic, Quadriplegic, n: 83; GMFCS: I, II, III, IV and V.	10.8 years (9- 12 years).	GMFCS, 6MWT, 10MWT, Sit and stand test, PowerTrackII, LEMOCOT.	GMFCS, Neuromusculoskeletal aspects (muscle contractures, spasticity, coordination and strength), Mobility performance (ability to get up from a chair and walking speed over short and long distances).
Davids et al., 2014	U.S	Prospective cohort	CP (Bilateral; n: 255; (F/M: 90–65); GMFCS: I, II, III, IV and V	13 years.	GMFCS, Portable Dynamometer (isometric muscle strength)	GMFCS, Age, Neuromusculoskeletal aspects (strength/weight ratio).
Day et al., 2007	U.S	Prospective cohort or non- randomized group	CP (Unilateral, Bilateral, Ataxic); n: 13, 271; GMFCS: I, II, III and IV	17.5 years	CDER, GMFCS	GMFCS, Age.
Gannotti et al., 2013	U.S	Longitudinal	CP (Unilateral, Bilateral); n: 26; GMFCS: I, II, III and IV	25 years; (15–11)	MIF, SF-36, COPM, GGI, GMFCS, semi-structured questionnaire (participation, personal and environmental factors), body weight, ROM.	Pain, mental health, Body weight, Participation (housing).
Gillett et al., 2018	Australia	Cross-sectional	PC; n: 33; GMFCS: I and II;	25 years; (15–18)	6MWT, LSU, TUS, Isokinetic dynamometer (isometric muscle strength), muscle ultrasound	GMFCS, Age, Isometric strength of plantar flexors.
Hanna et al., 2009	Canada	Longitudinal	CP (Unilateral, Bilateral, Ataxic); n: 657; GMFCS: I, II, III, IV and V	3.6 years (16 months- 21 years)	GMFM-66, GMFCS	GMFCS, Age.
Harvey et al., 2012	Australia	Longitudinal	CP (Bilateral)+ surgery); n: 156; GMFCS: I, II, III and IV	11.1 years; (57–99)	GMFCS; FMS	GMFCS, surgery (multilevel single event).
Himuro et al., 2018	Japan	Cross-sectional	CP; n: 386	26 years and 8 months; (172–213)	GMFCS; Self-report questionnaire and/or proxy developed for the study.	GMFCS, Age, Neuromusculoskeletal aspects.
Ho et al., 2017	Taiwan	Longitudinal	CP Unilateral, Bilateral, Ataxic, Dystonic); n: 92; GMFCS: I, II, III, IV and V	12.8 years; (35–57)	GMFM-66; GMFCS; ASKp	GMFCS, Age
Maanum et al., 2010	Norway	Cross-sectional	CP (Unilateral, Bilateral); n: 126; GMFCS: I, II and III	39 years old; (73–53)	6MWT, TUG, Borg Scale, GMFCS, FMS, Gait and Pain Questionnaire, demographic data	GMFCS, Pain, Neuromusculoskeletal aspects (popliteal angle), CP type, Mobility performance.
Morgan et al., 2013	Australia	Observational	CP (Unilateral, Bilateral); n: 25; GMFCS: I, II, III, IV and V	41 years; (16–9)	GMFCS; TC6M; TUG; FROP- community; CTSIB; Berg Balance Scale;	GMFCS, Gait changes, Balance changes.
Õunpuu et al., 2015	U.S	Longitudinal	CP (Unilateral, Bilateral) + multilevel surgery; n: 22; GMFCS: I, II, III and IV;	9.5 years; (11–11)	Physical exam, gait analysis, EMG	Age, Surgery history, Body Mass Index, Height and Weight.

(continued on next page)

Table 2 (continued)

Author/ Year	Country of origin	Study design	Participants		Outcome measures	Factors associated with motor outcomes evaluated *
Sienko 2017	U.S	Cross-sectional	CP; n: 97; GMFCS: I, II, III, IV and V	23.8 years.	GMFCS, BRFS, BPI, FAS, PHQ- 9, MHLC	Age, GMFCS, Pain, General health, Emotional support, Fatigue and Depression. GMFCS, Age.
Smits et al., 2013	Netherlands	Longitudinal	CP (Unilateral, Bilateral, Ataxic); n: 423; GMFCS: I, II, III, IV and V	9 years and 6 months; (1–22 years).	GMFM-66, GMFCS	
Smits et al., 2019	Netherlands	Longitudinal	CP; n: 551; GMFCS: I, II, III, IV and V	10 years; (1–21 years)	PEDI (mobility and self-care), GMFCS	GMFCS, Age and Self-Care.
Usuba et al., 2014	Canada	Cohort design or non-randomized group	CP; n: 54; GMFCS: I, II, III, IV and V	30.4 years; (25–29)	GMFCS, SRH, HUI3, AqoL	Age
Van Eck et al., 2008	Netherlands	Longitudinal	CP (Unilateral, Bilateral, Ataxic); n: 105; GMFCS: I, II, III, IV and V	11 years; (38–63)	VABS, GMFM-66, GMFCS	Age, motor type, CP distribution, sex and GMFCS.
Veerbeek et al., 2019	South Africa	Observational	CP (Bilateral)+SDR; n: 26; GC (without PC): 26; GMFCS: I, II and III	31.3 years; (25.4– 41.4); 35.7 years; (33.2– 44.2)	LIFE-H 3.1, FMS	Age; Decreased Mobility (FMS); Housing, Fitness, Personal Care and Recreation.

Caption: *= all factors analyzed in the study; CP, cerebral palsy; GMFCS, Gross Motor Function Classification System; GMFM, Gross Motor Function Measure; LIFE-H, Assessment of Life Habits; FMS, Functional Mobility Scale; PEDI, Pediatric Assessment of Disability Inventory; TC6, 6-minute walk test; TUG, Timed Up and Go; BBS Berg Balance Scale; LSU, lateral step-up; TSU, Timed up-stairs; BPI, Brief Pain Inventory; GGI, Gillette Gait Index; SF-36, Quality of life questionnaire; VABS, Vineland Adaptive Behavior Scales; HUI-3, Health Utilities Index; EMG, dynamic electromyography; CPCHILD, Caregiver Priorities and Child Health Index of Life with Disabilities Questionnaire; PROMIS, Global Health of PROMIS; CDER, Client Development Evaluation Report; COPM, Canadian Occupational Performance Measure; ASKp, Children's Activity Scale; FROP-community, Falls risk for older People in the community; CTSIB, Clinical Test of Sensory Interaction and Balance; BRFS, Behavioral Risk Factor Surveillance System; BPI, Scale of the Brief Pain Inventory; FAS, Fatigue Rating Scale; PHQ-9, Patient Health Questionnaire-9; MHLC, Health Locus of Control Scale; ROM, Range of motion; SAROMM, Spinal Alignment and Range of Motion Measure; F, female; M, male; NA, not applicable.

Table 3

Factors related to motor outcomes according to each category of interest.

Motor-related outcomes	Relevant factors and corresponding studies
Gross motor function	Age (Church et al., 2018; Davids et al., 2014; Gillett et al., 2018; Hanna et al., 2009; Himuro et al., 2018; Ho et al., 2017; Sienko 2017; Smits et al., 2013; Smits et al., 2019; Usuba et al., 2014; Van Eck et al., 2008; Veerbeek et al., 2019) GMFCS (Chiu et al., 2023; Church et al., 2018; Davids et al., 2014; Gillett et al., 2018; Hanna et al., 2009; Harvey, et al., 2012; Himuro et al., 2018; Ho et al., 2017; Sienko 2017; SMITS et al., 2013; Smits et al., 2019; Van Eck et al., 2008) Neuromusculoskeletal aspects (Bell et al., 2002; Bartlett et al., 2010; Chiu et al., 2023; Davids et al., 2014; Himuro et al., 2018) Pain (Bell et al., 2002; Bartlett et al., 2010; Sienko 2017; Himuro et al., 2018) Surgery history (Harvey, et al., 2012; Church et al., 2018) Altered Gait parameters (Church et al., 2018)
Gait	GMFCS (Church et al., 2018; Day et al., 2007; Gillett et al., 2018; Maanum et al., 2010; Morgan et al., 2013; Sienko 2017) Age (Bell et al., 2002; Day et al., 2007; Gillett et al., 2018; Öunpuu et al., 2015; Sienko 2017) Pain (Baird et al., 2021; Gannotti et al., 2013; Maanum et al., 2010; Sienko 2017) Neuromusculoskeletal aspects (Baird et al., 2021; Bell et al., 2002; Maanum et al., 2010) Surgery history (Veerbeek et al., 2019; Öunpuu et al., 2015; Church et al., 2018) Gait changes (Church et al., 2018; Morgan et al., 2013) Performance Mobility (Maanum et al., 2010)
Neuromusculoskeletal Functions	Neuromusculoskeletal aspects (Bell et al., 2002; Buckland et al., 2021; Chiu et al., 2023) Pain (Bell et al., 2002; Gannotti et al., 2013) Gait changes (Church et al., 2018) Age (Bell et al., 2002) Surgery History (Church et al., 2018)

Caption: GMFCS = Gross Motor Function Classification System.

In summary, the findings are quite consistent in demonstrating that during transition to adulthood there is a decline in gross motor function and mobility, with age, GMFCS level, and body structure and function aspects such as pain and neuromusculoskeletal aspects being the primary relevant factors. Thus, monitoring these factors throughout the development of children, adolescents, and young adults with CP is important so timely interventions to prevent decline and/or maintain the functional level are offered.

Decline in gait

Gait deterioration was reported as an indicator of motor decline in 11 studies. GMFCS level was pointed out as a relevant factor in several studies,^{15,16,19,26,31,39,40} often interacting with age to determine changes in gait.^{15,31,39,41,42} Other aspects of gross motor function such as the performance of the Timed Up and Go (TUG) test, the ability to walk and climb stairs, and the need for assistive devices were also relevant predictors of gait stability and decline.¹⁹ It is therefore relevant to consider the implementation of validated tests in transition programs to help predict the risk of gait deterioration.

The presence of pain was another relevant factor, predicting gait endurance,¹⁹ and gait in general,¹⁵ with reports of loss of the ability to walk in individuals who had pain.¹⁶ However, the impact of the association of pain and gait decline on functioning was not explored in the included studies, and should be addressed in the future. Another limitation was that the assessment of pain did not include detailed information such as type and location of the pain, which warrants further investigation.

The presence of musculoskeletal deformities was also reported as an important contributor to gait loss,⁴⁰ and to the performance in gait tests.^{19,41} However, there is conflicting evidence on the impact of interventions targeting such deformities on gait. Bell et al.⁴¹ showed that gait function declined over 4 years in both functional and less functional walkers without surgery, illustrated by decline walking velocity and other kinematic parameters. In contrast, a group that underwent orthopedic surgery had a more positive prognosis, suggesting that this intervention may minimize the decline over time. Possible positive effects of surgical interventions were also reported by Gannotti et al.¹⁶ and

Table 4

Risk of bias of the 24 studies included in the review.

Study	Checklist items																Total score*	Quality
	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16		
Baird et al., 2021	1	1	1	1	0	1	1	0	1	1	1	1	1	1	1	1	14	Low
Bartlett et al., 2010	1	1	0	1	0	1	0	1	1	1	1	1	1	1	1	1	13	Low
Bell et al., 2002	1	1	1	1	1	1	1	0	1	0	1	1	1	0	1	0	12	Low
Buckland et al., 2021	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	16	Low
Church et al., 2018	1	1	0	0	1	1	1	1	1	0	1	1	1	1	1	1	13	Low
Chiu et al., 2023	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	16	Low
Davids et al., 2014	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	16	Low
Day et al., 2017	1	1	1	1	1	1	1	0	1	0	1	1	1	1	1	1	14	Low
Gannotti et al., 2013	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	16	Low
Gillett et al., 2018	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	16	Low
Hanna et al., 2009	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	16	Low
Harvey et al., 2012	1	1	1	1	0	1	1	1	1	0	1	1	1	1	1	1	14	Low
Himuro et al., 2018	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	16	Low
Ho et al., 2017	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	16	Low
Maanum et al., 2010	1	1	1	1	1	1	1	1	1	0	1	1	1	1	1	0	14	Low
Morgan et al., 2013	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	16	Low
Öunpuu et al., 2015	1	1	0	1	1	1	0	0	1	0	1	1	1	1	1	1	12	Low
Sienko 2017	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	16	Low
Smits et al., 2013	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	16	Low
Smits et al., 2019	1	1	1	1	1	1	1	1	1	0	1	1	1	1	1	1	15	Low
Usaba et al., 2014	1	1	0	1	0	1	0	1	1	1	1	1	1	1	1	1	13	Low
Van Eck et al., 2008	1	1	0	1	1	1	1	1	1	1	1	1	0	1	1	1	14	Low
Veerbeek et al., 2019	1	1	1	1	0	1	1	1	1	1	1	1	1	1	1	1	15	Low

Veerbeek et al.¹⁷ These findings are however in contrast with the decline after multilevel surgery over an 11-year period reported in another study.⁴² Also, Church et al.²⁶ have suggested a mild decline (increased popliteal angle and decreased gait speed) in a 12-year follow-up after multilevel surgery. Given these findings, particularities such as type of surgery, participants' age, and other characteristics should be explored in future studies to elucidate the impact of surgery on motor prognosis.

The findings of this review highlight that gait decline is prevalent and relevant in the motor decline context. When untreated, it can result in loss of mobility and quality of life.⁴³ Thus, the identification of relevant factors for changes in gait can prevent a cascade of other potentially aggravating events for health in general. Monitoring these factors should be a priority in transition planning.

Decline in neuromusculoskeletal functions

The decline in neuromusculoskeletal functions (muscle strength, ROM, worsening of hip condition, among others) was related to factors such as GMFCS level, altered gait parameters, and age^{26,37,38,41,44} Interestingly, worsened conditions in the hips, knees, and ankles may already appear during childhood, as noted by Bell et al.⁴¹ in children followed over 4 years starting at ages as young as 5 years. Less functional walkers presented particular characteristics, such as more changes in the foot and ankle, which may relate to the increased risk of mobility decline in this subgroup and points to the importance of prevention by means of appropriate tone management and physical activity early on.⁴¹ Over a 12-year follow-up, however, changes in knee angles did not result in significant decline in the gait of young adults,¹⁶ which suggests that outcomes related to ROM must be accompanied by measures that inform about the capacity and performance of individuals, so that the proposed interventions can be functionally relevant.

One study focused more directly on hip status of non-ambulant children with CP,⁴⁴ pointing that a postoperative hyperlordosis greater than 60° was related to the increased risk of worsening the hip condition at 5 years after surgical correction of scoliosis. The implementation of adequate postural management through positioning and assistive devices is therefore a critical strategy to prevent such complications. Although not reported in the included studies, other studies indicate that in addition to spinal alignment, other factors such as age, GMFCS level, and type of gait as determinants for hip dislocation.^{26,41} Considering the

incidence of hip dislocation of up to 35 % (increasing with GMFCS) in CP, which can result in pain, reduced function, and reduced quality of life,⁴⁴ the factors relevant for this condition should be closely monitored and addressed as early as possible. The implementation of hip surveillance, which is the strategy with the highest level of evidence to minimize complications related to hip dislocation, is necessary from the first year of life of children with CP and in some cases needs to continue until skeletal maturity.^{45,46}

Given these findings, it is clear that musculoskeletal comorbidities and multimorbidities arise early and deteriorate with advancing age in adults with CP and may be more critical in the non-ambulant individuals. Thus, access to preventive services and therapeutic interventions at the right time can have a positive impact on the functioning and well-being of individuals with CP, especially those at higher risk for musculoskeletal complications, thus providing more favorable health conditions as they transition to adulthood.

Quality appraisal and future research needs

In general, the reviewed studies showed a low risk of bias, which gives more credibility and support to the conclusions made in this review. However, heterogeneity in the use of assessment instruments was found, for example, the GMFCS in some studies was used as a criterion to indicate gait ability, while in others as an indicator of general gross motor function. There was also a lack of validated assessments for older age groups of people with CP. Although the GMFM is the gold standard in CP assessment, it is not validated for adults yet. This aspect also deserves attention in future studies.

Limitations of the study include the fact that the factors were selected based on their statistical significance and the lack of measures of the strength of the recommendations. We suggest that future reviews also take into account the clinical significance of the findings and use tools to provide accurate recommendations. Interestingly, indicators of functional decline that were not directly related to motor outcomes were assessed in some studies, including decline in self-care skills, health-related quality of life, and performance and satisfaction with lifestyle habits. These indicators should not be disregarded as they illustrate areas of functioning that are essential for wellbeing. Other factors identified as relevant for motor outcomes but not classified as significant as they did not reach the criteria established in this review included

depression, emotional support,¹⁵ mental health,¹⁶ housing,⁴¹ balance,¹⁸ fatigue,¹⁵ and CP type.¹⁹ Their contribution should be investigated in future, high quality studies. Still, although the distribution of motor impairment and the presence of comorbidities have been previously pointed out as relevant factors in the literature,¹⁸ they were not identified in the results of this review. Such variability of outcomes and results is probably due to the multifactorial nature of motor decline, and indicates that individualized biopsychosocial care must be assured when planning transition services to adult life.

Regarding implications for physical therapy, the results of this review support the continuity of follow-up of individuals with CP during the transition to adulthood, especially the ones at higher risk for motor decline, while discharge due to lack of motor progress is not recommended as it may contribute to further deterioration in health status. Future studies should summarize evidence on recommended physical therapy interventions for this population. Studies that longitudinally follow adolescents and young adults with CP are also scarce, especially in low and middle-income countries, and the determinants for the motor outcomes in these countries need to be investigated in future studies.

Conclusion

This review summarized the factors that may influence the motor outcomes of adolescents and adults with CP, highlighting the role of age, GMFCS level, and presence of pain, which stood out as significant for decline in gross motor function, gait, and neuromusculoskeletal functions. Knowledge of these factors is relevant for professionals and caregivers of people with CP, as monitoring them during the transition to adolescence and adulthood may help reduce the impact of limitations caused by motor decline.

Declaration of Competing interest

The authors declare no competing interest.

Funding

This study was funded by the Coordination for the Improvement of Higher Education Personnel. Process number 131,374/2021-4 and by the National Council for Scientific and Technological Development (CNPq), grant number: 422090/2021-2. The funder had no role in the preparation of the manuscript.

This study will not need approval by the Ethics Committee, as it will not use data at the individual level of the participants.

Acknowledgements

We are grateful to the Coordination for the Improvement of Higher Education Personnel (CAPES), National Council for Scientific and Technological Development (CNPq).

Supplementary materials

Supplementary material associated with this article can be found, in the online version, at [doi:10.1016/j.bjpt.2025.101225](https://doi.org/10.1016/j.bjpt.2025.101225).

References

- Tosi LL, Maher N, Moore DW, Goldstein M, Aisen ML. Adults with cerebral palsy: a workshop to define the challenges of treating and preventing secondary musculoskeletal and neuromuscular complications in this rapidly growing population. *Dev Med Child Neurol*. 2009 Oct;51(Suppl 4):2–11. <https://doi.org/10.1111/j.1469-8749.2009.03462.x>. PMID: 19740204.
- Westbom L, Bergstrand L, Wagner P, Nordmark E. Survival at 19 years of age in a total population of children and young people with cerebral palsy. *Dev Med Child Neurol*. 2011 Sep;53(9):808–814. <https://doi.org/10.1111/j.1469-8749.2011.04027.x>. Epub PMID: 21745199.
- Bolger A, Vargus-Adams J, McMahon M. Transition of care in adolescents with cerebral palsy: a survey of current practices. *PM R*. 2017 Mar;9(3):258–264. <https://doi.org/10.1016/j.pmrj.2016.08.001>. Epub 2016 Aug 9. PMID: 27519825.
- Ryan JM, Fortune J, Walsh A, Norris M, Kerr C, Hensley O, Kroll T, Lavelle G, Owens M, Byrne M, Walsh M. Transition from child to adult health services for young people with cerebral palsy in Ireland: a mixed-methods study protocol. *BMJ Open*. 2020 Dec 22;10(12), e041425. <https://doi.org/10.1136/bmjopen-2020-041425>. PMID: 33371036; PMCID: PMC7757447.
- Myers LL, Nerminathan A, Fitzgerald DA, Chien J, Middleton A, Waugh MC, et al. Transition to adult care for young people with cerebral palsy. *Paediatr Respir Rev*. 2020;33:16–23.
- Ferreira MC, Garcia NR, Prudente COM, Ribeiro MFM. Quality of life of adolescents with cerebral palsy: agreement between self-report and caregiver's report. *Rev Lat Am Enfermagem*. 2020 Jul 1;28:e3300. <https://doi.org/10.1590/1518-8345.3928.3300>. PMID: 32609264; PMCID: PMC7332252.
- Morgan P, McGinley J. Gait function and decline in adults with cerebral palsy: a systematic review. *Disabil Rehabil*. 2014;36(1):1–9. <https://doi.org/10.3109/09638288.2013.775359>. Epub 2013 Apr 17. PMID: 23594053.
- Shearer HM, Côté P, Hogg-Johnson S, McKeever P, Fehlings DL. Pain trajectories and well-being in children and young people with cerebral palsy: a cohort study. *Dev Med Child Neurol*. 2022 Nov;64(11):1416–1424. <https://doi.org/10.1111/dmcn.15252>. Epub 2022 Apr 30. PMID: 35489074; PMCID: PMC9790202.
- Solanke F, Colver A, McConachie H.; Transition collaborative group. Are the health needs of young people with cerebral palsy met during transition from child to adult health care? *Child Care Health Dev*. 2018 May;44(3):355–363. [doi: 10.1111/cch.12549](https://doi.org/10.1111/cch.12549). Epub 2018 Jan 28. PMID: 29377236; PMCID: PMC5900977.
- Page MJ, McKenzie JE, Bossuyt PM, Boutron I, Hoffmann TC, Mulrow CD, et al. The PRISMA 2020 statement: an updated guideline for reporting systematic reviews. *BMJ*. 2021;372:n71. <https://doi.org/10.1136/bmj.n71>.
- Linsell L, Malouf R, Morris J, Kurinczuk JJ, Marlow N. Prognostic factors for cerebral palsy and motor impairment in children born very preterm or very low birthweight: a systematic review. *Dev Med Child Neurol*. 2016 Jun;58(6):554–569. <https://doi.org/10.1111/dmcn.12972>. Epub 2016 Feb 10. PMID: 26862030; PMCID: PMC5321605.
- World Health Organization; 2001. 6. [OMS] Organização Mundial da Saúde, CIF: Classificação Internacional de Funcionalidade, Incapacidade e Saúde [Centro Colaborador da Organização Mundial da Saúde para a Família de Classificações Internacionais, org.; coordenação da tradução Cassia Maria Buchalla].
- Visicato LP, da Costa CSN, Damasceno VAM, de Campos AC, Rocha NACF. Evaluation and characterization of manual reaching in children with cerebral palsy: a systematic review. *Res Dev Disabil*. 2015;36:162–174.
- Santana CAS, Dos Santos MM, de Campos AC. Interrelationships of touch and proprioception with motor impairments in individuals with cerebral palsy: a systematic review. *Percept Mot Skills*. 2022 Jun;129(3):570–590. <https://doi.org/10.1177/00315152221093904>. Epub 2022 Apr 22. PMID: 35452588.
- Sienko SE. An exploratory study investigating the multidimensional factors impacting the health and well-being of young adults with cerebral palsy. *Disabil Rehabil*. 2018 Mar;40(6):660–666. <https://doi.org/10.1080/09638288.2016.1274340>. Epub 2017 Jan 9. PMID: 28068863.
- Gannotti ME, Gorton GE IIIrd, Nahorniak MT, Masso PD. Gait and participation outcomes in adults with cerebral palsy: a series of case studies using mixed methods. *Disabil Health J*. 2013 Jul;6(3):244–252. <https://doi.org/10.1016/j.dhjo.2013.01.013>. Epub 2013 Mar 17. PMID: 23769484.
- Veerbeek BE, Lamberts RP, Fiegeen AG, Verkoijen PPJL, Langerak NG. Daily activities, participation, satisfaction, and functional mobility of adults with cerebral palsy >25 years after selective dorsal rhizotomy: a long-term follow-up during adulthood. *Disabil Rehabil*. 2021 Jul;43(15):2191–2199. <https://doi.org/10.1080/09638288.2019.1695001>. Epub 2019 Dec 9. PMID: 31815556.
- Morgan P, McGinley J. Performance of adults with cerebral palsy related to falls, balance and function: a preliminary report. *Dev Neurorehabil*. 2013;16(2):113–120. <https://doi.org/10.3109/17518423.2012.725107>. PMID: 23477464.
- Maanum G, Jahnsen R, Frøslie KF, Larsen KL, Keller A. Walking ability and predictors of performance on the 6-minute walk test in adults with spastic cerebral palsy. *Dev Med Child Neurol*. 2010 Jun;52(6):e126–e132. <https://doi.org/10.1111/j.1469-8749.2010.03614.x>. Epub 2010 Feb 12. PMID: 20163429.
- Leite HR, Jindal P, Malek SA, Rosenbaum P. Research on children with cerebral palsy in low- and middle-income countries. *Pediatr Phys Ther*. 2022 Oct 1;34(4): 551–555. <https://doi.org/10.1097/PEP.0000000000000949>. Epub 2022 Aug 12. PMID: 35960038.
- Chagas PSC, Drumond CM, Toledo AM, de Campos AC, Camargos ACR, Longo E, Leite HR, Ayupe KMA, Moreira RS, Moraes RLS, Palisano RJ, Rosenbaum P. PartiCipa Brazil Research Group. Study protocol: functioning curves and trajectories for children and adolescents with cerebral palsy in Brazil - PartiCipaBrazil. *BMC Pediatr*. 2020 Aug 20;20(1):393. <https://doi.org/10.1186/s12887-020-02279-3>. PMID: 32819318; PMCID: PMC7439543.
- Chagas PSC, Magalhães EDD, Sousa Junior RR, Romeros ACSF, Palisano RJ, Leite HR, Rosenbaum P. Development of children, adolescents, and young adults with cerebral palsy according to the ICF: a scoping review. *Dev Med Child Neurol*. 2023 Jun;65(6):745–753. <https://doi.org/10.1111/dmcn.15484>. Epub 2022 Dec 5. PMID: 36469744.
- Silva SP, Beunen G, Freitas DL, Maia JAR. Estudos longitudinais sobre o crescimento somático e desempenho motor: delineamentos, desafios, necessidades. *Rev Bras Cineantropom Desempenho Hum*. 2013;15(1):130–143. <https://doi.org/10.5007/1980-0037.2013v15n1p130>.

24. Ghali AJ, Al-Majmae KA, Nabat AA. Clinical patterns of cerebral palsy with complications and neurodisabilities associated with cerebral palsy. *ResearchGate*. 2021. <https://doi.org/10.52113/1/2410-4590/2021-54-61>.
25. Furtado MAS, Ayupe KMA, Christovão IS, Sousa Júnior RR, Rosenbaum P, Camargos ACR, Leite HR. Fisioterapia em crianças com paralisia cerebral no Brasil: uma revisão de escopo. *Dev Med Child Neurol*. 2022 May;64(5):e2–e12. <https://doi.org/10.1111/dmcn.15094>. Epub 2021 Oct 24. PMID: 34689323.
26. Church C, Ge J, Hager S, Haumont T, Lennon N, Niiler T, Hulbert R, Miller F. Flexed-knee gait in children with cerebral palsy: a long-term follow-up study. *Bone Joint J*. 2018 Apr 1;100-B(4):549–556. <https://doi.org/10.1302/0301-620X.100B4.BJJ-2017-0732.R1>. PMID: 29629589.
27. Voorman JM, Dallmeijer AJ, Knol DL, Lankhorst GJ, Becher JG. Prospective longitudinal study of gross motor function in children with cerebral palsy. *Arch Phys Med Rehabil*. 2007 Jul;88(7):871–876. <https://doi.org/10.1016/j.apmr.2007.04.002>. PMID: 17601467.
28. Bartlett DJ, Hanna SE, Avery L, Stevenson RD, Galuppi B. Correlates of decline in gross motor capacity in adolescents with cerebral palsy in gross motor function classification system levels III to V: an exploratory study. *Dev Med Child Neurol*. 2010 Jul;52(7):e155–e160. <https://doi.org/10.1111/j.1469-8749.2010.03632.x>. Epub 2010 Feb 24. PMID: 20187880.
29. Himuro N, Mishima R, Seshimo T, Morishima T, Kosaki K, Ibe S, Asagai Y, Minematsu K, Kurita K, Okayasu T, Shimura T, Hoshino K, Suzuki T, Yanagizono T. Change in mobility function and its causes in adults with cerebral palsy by Gross Motor Function Classification System level: a cross-sectional questionnaire study. *NeuroRehabilitation*. 2018;42(4):383–390. <https://doi.org/10.3233/NRE-172340>. PMID: 29660955.
30. Davids JR, Oeffinger DJ, Bagley AM, Sison-Williamson M, Gorton G. Relationship of strength, weight, age, and function in ambulatory children with cerebral palsy. *J Pediatr Orthop*. 2015 Jul-Aug;35(5):523–529. <https://doi.org/10.1097/BPO.0000000000000320>. PMID: 25264553.
31. Gillett JG, Lichtwark GA, Boyd RN, Barber LA. Functional capacity in adults with cerebral palsy: lower limb muscle strength matters. *Arch Phys Med Rehabil*. 2018 May;99(5):900–906. <https://doi.org/10.1016/j.apmr.2018.01.020>. e1.Epub 2018 Feb 10. PMID: 29438658.
32. Hanna SE, Rosenbaum PL, Bartlett DJ, Palisano RJ, Walter SD, Avery L, Russell DJ. Stability and decline in gross motor function among children and youth with cerebral palsy aged 2 to 21 years. *Dev Med Child Neurol*. 2009 Apr;51(4):295–302. <https://doi.org/10.1111/j.1469-8749.2008.03196.x>. PMID: 19391185.
33. Ho PC, Chang CH, Granlund M, Hwang AW. The relationships between capacity and performance in youths with cerebral palsy differ for GMFCS levels. *Pediatr Phys Ther*. 2017 Jan;29(1):23–29. <https://doi.org/10.1097/PEP.0000000000000332>. PMID: 27984462.
34. Smits DW, Gorter JW, Hanna SE, Dallmeijer AJ, van Eck M, Roebroek ME, Vos RC, Ketelaar M, PERRIN, Plus Study Group. Longitudinal development of gross motor function among Dutch children and young adults with cerebral palsy: an investigation of motor growth curves. *Dev Med Child Neurol*. 2013 Apr;55(4):378–384. <https://doi.org/10.1111/dmcn.12083>. Epub 2013 Jan 29. PMID: 23356539.
35. Smits DW, Gorter JW, Riddell CA, Voorman JM, Rosenbaum PL, Palisano RJ, Walter SD, Hanna SE, van Wely L, Ketelaar M. Mobility and self-care trajectories for individuals with cerebral palsy (aged 1–21 years): a joint longitudinal analysis of cohort data from The Netherlands and Canada. *Lancet Child Adolesc Health*. 2019 Aug;3(8):548–557. [https://doi.org/10.1016/S2352-4642\(19\)30122-1](https://doi.org/10.1016/S2352-4642(19)30122-1). Epub 2019 Jun 6. PMID: 31178370.
36. Usuba K, Oddson B, Gauthier A, Young NL. Changes in gross motor function and health-related quality of life in adults with cerebral palsy: an 8-year follow-up study. *Arch Phys Med Rehabil*. 2014 Nov;95(11):2071–2077. <https://doi.org/10.1016/j.apmr.2014.05.018>. e1.Epub 2014 Jun 6. PMID: 24909589.
37. Chiu HC, Ada L, Chen C. Changes in walking performance between childhood and adulthood in cerebral palsy: a systematic review. *Dev Neurorehabil*. 2020 Aug;23(6):343–348. <https://doi.org/10.1080/17518423.2019.1648579>. Epub 2019 Jul 31. PMID: 31366265.
38. Harvey A, Rosenbaum P, Hanna S, Yousefi-Nooraie R, Graham KH. Longitudinal changes in mobility following single-event multilevel surgery in ambulatory children with cerebral palsy. *J Rehabil Med*. 2012 Feb;44(2):137–143. <https://doi.org/10.2340/16501977-0916>. PMID: 22266600.
39. Day SM, Wu YW, Strauss DJ, Shavelle RM, Reynolds RJ. Change in ambulatory ability of adolescents and young adults with cerebral palsy. *Dev Med Child Neurol*. 2007 Sep;49(9):647–653. <https://doi.org/10.1111/j.1469-8749.2007.00647.x>. PMID: 17718819.
40. Baird G, Chandler S, Shortland A, Will E, Simonoff E, Scrutton D, Fairhurst C. Acquisition and loss of best walking skills in children and young people with bilateral cerebral palsy. *Dev Med Child Neurol*. 2022 Feb;64(2):235–242. <https://doi.org/10.1111/dmcn.15015>. Epub 2021 Aug 19. PMID: 34410016.
41. Bell KJ, Ounpuu S, DeLuca PA, Romness MJ. Natural progression of gait in children with cerebral palsy. *J Pediatr Orthop*. 2002 Sep-Oct;22(5):677–682. PMID: 12198474.
42. Ounpuu S, Solomito M, Bell K, DeLuca P, Pierz K. Long-term outcomes after multilevel surgery including rectus femoris, hamstring and gastrocnemius procedures in children with cerebral palsy. *Gait Posture*. 2015 Sep;42(3):365–372. <https://doi.org/10.1016/j.gaitpost.2015.07.003>. Epub 2015 Jul 23. PMID: 26260009.
43. Lennon N, Church C, Shrader MW, Robinson W, Henley J, Salazar-Torres JJ, Niiler T, Miller F. Mobility and gait in adults with cerebral palsy: evaluating change from adolescence. *Gait Posture*. 2021 Oct;90:374–379. <https://doi.org/10.1016/j.gaitpost.2021.09.177>. Epub 2021 Sep 20. PMID: 34564009.
44. Buckland AJ, Woo D, Kerr Graham H, Vasquez-Montes D, Cahill P, Errico TJ, Sponseller PD, Harms Study Group. Residual lumbar hyperlordosis is associated with worsened hip status 5 years after scoliosis correction in non-ambulant patients with cerebral palsy. *Spine Deform*. 2021 Jul;9(4):1125–1136. <https://doi.org/10.1007/s43390-020-00281-4>. Epub 2021 Feb 1. PMID: 33523455.
45. AACPD. Care pathways: hip surveillance. 2017. [internet] accessed: January 07, 2023 <https://www.aacpdm.org/UserFiles/file/hip-surveillance-care-pathway.pdf>.
46. Gibson N, Wynter M, Thomason P, Baker F, Burnett H, Graham HK, Kentish M, Love SC, Maloney E, Stannage K, Willoughby K. Australian hip surveillance guidelines at 10 years: new evidence and implementation. *J Pediatr Rehabil Med*. 2022;15(1):31–37. <https://doi.org/10.3233/PRM-220017>. PMID: 35311729.