



Repeatability and validation of Gait Deviation Index in children: Typically developing and cerebral palsy



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ABSTRACT

The Gait Deviation Index (GDI) is a dimensionless parameter that evaluates the deviation of kinematic gait from a control database. The GDI can be used to stratify gait pathology in children with cerebral palsy (CP). In this paper the repeatability and uncertainty of the GDI were evaluated. The Correlation between the GDI and the Gross Motor Function Classification System (GMFCS) was studied for different groups of children with CP (hemiplegia, diplegia, triplegia and quadriplegia). Forty-nine, typically developing children (TD) formed our database. A retrospective study was conducted on our 3D gait data and clinical exams and 134 spastic children were included. Sixteen TD children completed the gait analysis twice to evaluate the repeatability of the GDI (test–retest evaluation). Monte Carlo simulations were applied for all groups (TD and children with CP) in order to evaluate the propagation of errors stemming from kinematics. The repeatability coefficient (2SD of test–retest differences), obtained on the GDI for the 16 TD children (32 lower limbs) was ± 10 . Monte Carlo simulations showed an uncertainty ranging between 0.8 and 1.3 for TD children and all groups with CP. The Spearman Rank correlation showed a moderate correlation between the GDI and the GMFCS ($r = -0.44$, $p < 0.0001$).

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1. Introduction

Given the different gait patterns and pathologies in children with cerebral palsy (CP), a global analysis is essential in clinical practice. For this purpose, estimation of gait deviations from normative values is required and is helpful to improve therapeutic interventions.

In the CP pathology, three dimensional (3D) gait analysis for ambulant children and clinical evaluations are commonly used as assessments and are part of the international standard of CP care [1]. Even with 3D gait analysis, which is becoming an essential tool to assess ambulant children with CP [2], it is sometimes difficult to define objectively the amount of abnormalities and the degree by which an abnormal gait deviates from normal patterns. Three dimensional (3D) gait analysis provides a large amount of interdependent data and variables corresponding to different gait patterns [3]. The quantity and complexity of the data have pushed many authors to describe indices based on 3D gait

analysis, developed primarily to evaluate clinical changes after a therapeutic intervention like the hip flexor index (HFI) [4] and the Gillette Gait Index (GGI) [5]. The GGI is calculated using discrete parameters incorporating 16 kinematic and temporal distance variables, chosen arbitrarily, allowing us to describe and to quantify the amount of pathology in an individual's gait pattern, and its repeatability has been evaluated [6]. These tools ignore the relation that exists between gait variables, contrary to other indices, described later, like the Gait Profile Score (GPS) [3] and the Gait Deviation Index (GDI) [7]. The GDI is an alternative to the GGI, measuring the subject's gait deviation from a normative database. It is calculated using kinematic variables, studied point by point, during the entire gait cycle. It is a scaled distance between 9 kinematic parameters of pathological gait and the average of 9 kinematic parameters of normal gait (group of typically developing (TD) children) [8]. The GDI is represented as a single number: when the number decreases, clinical involvement increases, and when the number increases, the gait profile is closer to a normal profile (≥ 100).

Many authors were interested in studying the correlation between indices calculated from the computerized gait analysis and clinical evaluation tools [9–11]. Molloy [12] studied the correlation between the GDI and clinical functional measures like the Gross Motor Function Measure (GMFM) and Gross Motor

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Function Classification system (GMFCS) and concluded that the GDI is a valid tool to describe motor impairments in CP.

However, for clinical use, it would be interesting to evaluate the repeatability of the GDI, by the test–retest method, since there is no previous study on this subject. Moreover, the method of calculation of the GDI involves kinematic curves and their transformation into vectors, matrices, euclidian distances and singular value decomposition [7]. Kinematics are subject to errors in 3D gait analysis and these errors can lead to uncertainties on the value of the GDI. It would be interesting to evaluate the propagation of errors during the calculation of the GDI.

The aim of this study is to evaluate the repeatability of the GDI within typically developing (TD) children. The error propagation during the calculation of the GDI was evaluated by applying Monte Carlo simulations on TD children and children with CP.

2. Methods

2.1. Samples

A retrospective study was conducted on our 3D gait data and clinical assessments performed between 2006 and 2012. One hundred ninety one ($N = 191$) children were referred to our gait laboratory for an orthopedic evaluation, an orthotic intervention or for a baseline gait assessment. Forty-nine ($N = 49$) TD children formed our asymptomatic database with a mean age of 10.3 years ($SD = 3$, minimum = 5 years, maximum = 15 years) [6]. One hundred thirty four ($N = 134$) children with the diagnosis of spastic cerebral palsy were included in this study with a mean age of 10.5 years ($SD = 4.5$, minimum = 5 years, maximum = 20 years). Six Vicon MX3 cameras were used for the data acquisition (Vicon Motion Systems, Oxford, UK). Sixteen reflective markers based on the Davis protocol [2] were placed to calculate joint kinematic angles. Six to seven trials at self-selected walking speed were collected for each subject and one representative cycle for a barefoot stride for each limb was selected, for each subject, jointly by a bioengineer and a physical therapist, both experienced in gait analysis. The GMFCS score was assessed by the physical therapist during the clinical examination, which was carried out immediately before the subject underwent 3D gait analysis.

Each child was classified by the referring clinician, according to the distribution of spasticity and limb involvement during visual gait assessment and clinical examination. Children with hemiplegia ($N = 31$) have one lower limb involved with the ipsilateral upper limb. Children with diplegia ($N = 86$) have both lower limbs involved with minimal involvement of the upper limbs, while children with triplegia ($N = 6$) present spasticity in the lower limbs with major involvement of one upper limb. Quadriplegia ($N = 11$) is characterized by moderate to severe involvement of the four limbs. The GDI score was calculated in a Matlab (Mathworks®, Natick, MA, USA) function, separately for each leg, using the method provided by Schwartz [7]. Nine kinematic variables are used for this calculation: pelvis and hip motion in the 3 planes (sagittal, frontal, horizontal), knee and ankle motion in the sagittal plane and the foot progression angle in the horizontal plane [7].

The database of TD children, available in our laboratory ($N = 49$ subjects) was divided into 2 groups: the reference group ($N = 33$ who performed the gait analysis one time) and the able-bodied group ($N = 16$ subjects who performed the exam twice at one week interval). The reference group formed the database for GDI calculation. The calculation of the GDI for the able-bodied group helped in the verification of GDI implementation. The repeatability of the GDI was calculated on the 32 lower limbs from the 16 TD children who formed the able-bodied group. The correlation between the GDI and the GMFCS was evaluated on the groups with CP.

2.2. Monte Carlo simulations

In order to evaluate the sensitivity of the GDI to the variability of kinematic parameters, Monte Carlo simulations were applied on the able-bodied group and on each group of children with CP. Monte Carlo simulations are widely used in biomedical studies to assess the propagation of uncertainties in a calculation process [6,13]. The application of this simulation requires: definition of the measurand (the mathematical model), association of uncertainty on each variable entry, generation of N random realizations of entry, calculation of the N measurands and their standard deviations (95% confidence interval: mean $\pm$ 2SD), via the model. In this study, the measurand defined is the GDI. The variable entries are the kinematic curves of the subjects. Ten thousand random realizations of entries were created by adding a random noise to each curve. The means of the random noises were set to zero and the standard deviations were the same as the inter-session variability found on each kinematic parameter in our laboratory [6]. The standard deviations were those found on the overall range of motion for each joint/segment (pelvis, hip, knee, ankle, foot progression) in each plane of interest (sagittal, frontal and horizontal). Ten thousand values of the GDI were obtained and their standard deviations were calculated (95% confidence interval by calculating the 2SD).

2.3. Statistical analysis

Xlstat® (Addinsoft, New York, USA) and Matlab® (Mathworks®, Natick, MA, USA) were used for statistical calculations. The Shapiro Wilk test was applied to study the distribution of all the variables. The Fisher test was applied in order to verify the equality of variances, then, the Student paired t -test was applied to compare the average of the GDI between session 1 and session 2 for the 16 TD children who completed the exam twice. The confidence interval was evaluated by calculating 2SD of the difference between the 2 sessions, by using the method described by Bland and Altman [14,15]: there were two samples of GDI values ($x_{i,1}$ and $x_{i,2}$) from the two sessions of the 16 subjects who completed the gait analysis twice (index i referring to subject's number). To evaluate the extent of the differences between repeated tests a within-subject inter-sessions difference was calculated, $D_i = (x_{i,1} - x_{i,2})$. The coefficient of repeatability was 2SD of the 16 values of D_i , as reported by Bland and Altman.

One way Anova with Duncan's post hoc test was applied to determine the ability of the GDI to differentiate between children with cerebral palsy, defined by their topographic classifications and GMFCS levels. A Power analysis was performed. The Spearman Rank correlation was applied to study the correlation between the GMFCS and the GDI. Significance level was set to 0.05 for all tests.

3. Results

3.1. GDI for children with CP

The demographic characteristics of the children with CP and TD children are summarized in Table 1. The GDI was calculated separately for each lower limb. The GDI was normally distributed in each GMFCS level for children with CP and for TD children. The mean value of the GDI according to GMFCS levels is represented in Table 2. The mean value of the GDI decreases when the GMFCS increases. The histogram of the GDI is shifted toward 100 when the GMFCS decreases (Fig. 1 (left)). The mean values of the GDI (mean $\pm$ 1SD) for the different groups of children with CP are presented in Table 2. We noticed an overlap of GDI values for various CP sub-types most noted between hemiplegia, diplegia and triplegia (Fig. 1 (right)).

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