Neutrophil-to-lymphocyte ratio predicts the severity of motor impairment in cerebral palsy children living at home and the rehabilitation center: A comparative study

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Abstract. The aim of the present study was to investigate the neutrophil-to-lymphocyte ratio (NLR) level in children with cerebral palsy (CP) living at home or the rehabilitation center. The correlation of NLR with different severities of motor impairment was assessed. This was a single-center, cross-sectional, observational study. A total of 80 CP children who were either living at home (n=34) or at the rehabilitation center (n=46) were included. Demographic characteristics, anthropometric parameters and complete blood counts were recorded, and the NLR values were calculated. The severity of motor impairment was evaluated and categorized according to the Gross Motor Functional Classification System (GMFCS) level. The mean age of CP participants was 8.52±1.92 years. The percentage of children with CP who were malnourished (underweight, stunted and wasted) was higher amongst those at the rehabilitation center compared with those living at home. The mean NLR of children with CP in the rehabilitation center was significantly higher compared with the patients living at home (P=0.003). Participants from the rehabilitation center had severe motor impairment (GMFCS levels IV-V) and significantly higher NLR values than those with mild motor impairments (GMFCS levels II-III; P=0.006). However, there were no differences in NLR values in relation to severity of motor impairment in CP children living at home. CP children had some degree of neuroinflammation and systemic inflammation. NLR may be a potential simple inflammatory parameter that may be used to predict the severity of the motor impairment, particularly in CP children living at a rehabilitation center.

Introduction

Cerebral palsy (CP) is a heterogeneous disorder and there are several degrees of motor impairment and permanent non-progressive neurologic deficits due to the injury of the immature brain (1). CP is the most prevalent cause of disabilities in children. The overall global prevalence of CP is 2-3 per 1,000 live births (1). The motor disorders of CP include disturbances of sensation, perception, cognition, communication and behavior; and these symptoms are caused by epilepsy and secondary musculoskeletal disorders (2). Aside from the physical and mental disabilities, children with CP are often malnourished due to disruption of the enteric neural pathways resulting in problems swallowing, and as a result exhibit gastroesophageal reflux and difficulty in breathing (3). Previous studies have reported the negative effects of malnutrition on muscle strength, impairment of motor function and weakness of the respiratory muscle (4), pre-dispositions to congestive heart failure (5), impairments in wound healing and immune function, and increased risk of postoperative complications (6).

The classification of CP is based on the motor type and distribution including ataxia, athetoid, diplegia, double hemiplegia, hemiplegia and quadriplegia (3). The Gross Motor Function Classification System (GMFCS) is used in clinical practice (3). Studies have shown that a persistent and exacerbated inflammatory response in the central nervous system can cause neuronal damage (7,8). To the best of our knowledge, there are no studies aimed at identifying inflammatory biomarkers that can indicate the severity of CP.
Peripheral blood tests are routinely used in clinical practice. Several studies have reported that the neutrophil-to-lymphocyte ratio (NLR) and lymphocyte-to-monocyte ratio (LMR) may serve as biomarkers of an inflammatory response, and may be used as prognostic factors to predict the outcome or the severity of various malignancies (9), cardiovascular disease (10), systemic lupus erythematosus (11), hip fracture and polytrauma (12). Moreover, studies have demonstrated that NLR is associated with nutritional status in geriatric and cancer patients (13,14).

The healthcare costs for CP children with co-morbidities are high. In Australia, the cost of taking care of a single CP child is $43,431 per year (15). There are two primary means of providing healthcare for children with CP: Home-based procedures and rehabilitation-center-based procedures. Home-based patients exhibit improved social acceptance, functioning, participation and physical health, emotional wellbeing and self-esteem, as well as family health; in addition, these patients do not need to spend a lot of time traveling to the doctor, and thus have lower transportation costs (15). However, the healthcare costs for these patients are higher compared to children living in a rehabilitation center.

As CP children may live in different settings, the aim of the present study was to investigate the physical growth, nutritional status and inflammatory predictors between children living at home and the rehabilitation center. It was hypothesized that CP children living at home would exhibit improved physical growth, nutritional status and lower inflammatory parameters when compared with CP children living in a rehabilitation center. The relationships between these parameters and the severity of the motor impairment in CP children were also examined.

Materials and methods

Ethics. The study protocol used in the present study conformed to the ethical standards outlined in the Declaration of Helsinki and was approved by the Institutional Review Board on Human Research of the Faculty of Medicine, Chulalongkorn University. Informed consent was obtained from one of the parents or the legal guardian of the children, as well as from the child prior to his/her participation in the study.

Study participants. A total of 86 CP children were enrolled in a cross-sectional analytical study that was performed between June and December 2013. A total of 6 children were excluded as they had a history of gastrostomy (n=2), osteotomy around the knee (n=2), or were infections during the screening period (n=2). A total of 80 children with CP (45 males and 35 females) were eligible for analysis. The mean age in this study population was 8.52±1.92 years. These children were recruited from either a home-living setting (n=34) or a rehabilitation center (n=46). The home-living participants were those who lived at home and visited the rehabilitation center during office hours. Participants who lived at the rehabilitation center for at least 9 months were enrolled into the study.

Determination of clinical parameters. For the anthropometric measurements in the present study, the body weight was obtained and the height was calculated using Stevenson's recommended formula (16). The height (cm) was measured while in a sitting position from the knee to the floor. This measurement was taken 3 times, and the mean height was used. The body mass index (BMI) was calculated. To evaluate the nutritional status of the children, the anthropometric indices, including weight-for-age, height-for-age and weight-for-height, were used to calculate the z score based on the World Health Organization reference population median (17). Z-scores >2 standard deviations lower than the mean, for weight-for-age, height-for-age and weight-for-height, were considered as underweight, stunted and wasted, respectively.

The motor function for all CP children was evaluated and assigned according to the Gross Motor Function Classification System (GMFCS) level by certified physical therapists (18). The severity levels of motor impairment were divided into two groups: Mild motor impairment (GMFCS levels I-III) and severe motor impairment (GMFCS levels IV-V).

Peripheral blood counts were obtained from all participants as part of routine analysis. An automated hematology analyzer (Advia 2120 Hematology system; Siemens Healthineers) was utilized to obtain the complete blood count (CBC). Neutrophil and lymphocyte counts were extracted from the differential blood pictures. The NLR was calculated by dividing the absolute count of neutrophils by the number of lymphocytes from the CBCs. Additionally, the LMR was determined by dividing the absolute number of lymphocytes to the number of the monocytes.

Statistical analysis. Statistical analysis was performed using SPSS version 22.0 (IBM Corp.). Continuous data are presented as the mean ± standard error of the mean. Categorical data are summarized as the frequency and percentages. Comparisons of the frequencies of various demographic variables between the two groups were performed using a χ² test. The distributions of the continuous data were examined using a Q-Q plot, Shapiro-Wilk's test and a Kolmogorov-Smirnov test. To compare the means between the two groups, an unpaired Student's t-test was used for variables that were normally distributed. A Mann-Whitney U test was used for variables that did not have a normal distribution. For comparisons between >2 groups of variables, a Kruskal-Wallis one-way ANOVA with a Dunn's post hoc test was used for all pairwise comparisons. Pearson's and Spearman's correlation analysis were used to examine the relationship between variables. P<0.05 was used to indicate a statistically significant difference.

Results

The participants were divided into two groups, those living at home and those living in the rehabilitation center. The clinical characteristics of the participants are shown in Table I. There were no significant differences in the clinical characteristics between the two groups of children. Diplegic CP was the most common type of CP amongst participants living at home and the rehabilitation center. The severity of the motor impairment was categorized by GMFCS levels. In both care settings, the children had mild and severe motor impairments.

For anthropometric parameters, the height, weight and BMI were not significantly different between the children living at
home and the rehabilitation center. However, the percentage of underweight, stunted and wasted participants was higher amongst children living at the rehabilitation center compared with those living at home (Table I). The weight-for-height percentiles were strongly correlated with BMI in CP children ($r=0.919$, $P<0.001$; Fig. 1). The distribution of weight-for-height percentiles with stacking of each GMFCS level demonstrated left shifting, particularly in children living at the rehabilitation center (Fig. 2).

The CBC analysis showed that total white blood cell counts, neutrophil counts and the NLR values were significantly greater in the CP children at the rehabilitation center compared with those living at home ($P<0.05$; Table II). There was a direct association between the total white blood cell count and the NLR ($r=0.446$, $P<0.001$). However, NLR was inversely correlated with LMR in CP children, as illustrated in Fig. 3 ($r=-0.699$, $P<0.001$).

Overall, the NLR of the children in the rehabilitation center was significantly higher than those living at home ($P=0.003$; Table II). Additionally, the NLR was significantly higher in children with severe motor impairment ($P=0.001$; Table III; Fig. 4). In the rehabilitation center, the NLR values between mild and severe motor impairments were statistically different ($P=0.006$) and the proportion of children with GMFCS level V was greater than those with levels II and III ($P=0.021$ and $P=0.04$, respectively; Table III; Fig. 4). However, the NLR values were not significantly different amongst children based on severity of motor impairment. NLR was significantly positively correlated with GMFCS level ($r=0.467$, $P=0.001$; Fig. 5).

### Table I. Clinical characteristics of CP children living at home or the rehabilitation center.

| Characteristics          | Living at home, n (%) | Living at rehabilitation centre, n (%) | P-value |
|--------------------------|-----------------------|----------------------------------------|---------|
| Total                    | 34                    | 46                                     |         |
| Sex                      |                       |                                        |         |
| Male                     | 18 (52.9)             | 27 (58.7)                              | 0.18    |
| Female                   | 16 (47.1)             | 19 (41.3)                              | 0.612   |
| CP type                  |                       |                                        |         |
| Ataxia                   | 1 (2.9)               | 0 (0)                                  | -       |
| Athetoid                 | 4 (11.8)              | 4 (8.7)                                | 1       |
| Diplegia                 | 19 (55.9)             | 33 (71.7)                              | 0.052   |
| Double hemiplegia        | 1 (2.9)               | 1 (2.2)                                | 1       |
| Hemiplegia               | 1 (2.9)               | 5 (10.9)                               | 0.102   |
| Quadriplegia             | 8 (23.5)              | 3 (6.5)                                | 0.132   |
| GMFCS level              |                       |                                        |         |
| MMI                      | 16 (47.1)             | 28 (60.9)                              | 0.07    |
| Level II                 | 3 (8.8)               | 6 (13)                                 | 0.317   |
| Level III                | 13 (38.2)             | 22 (47.9)                              | 0.128   |
| SMI                      | 18 (52.9)             | 18 (39.1)                              | 1       |
| Level IV                 | 11 (32.4)             | 15 (32.6)                              | 0.433   |
| Level V                  | 7 (20.6)              | 3 (6.5)                                | 0.206   |
| Malnutrition             |                       |                                        |         |
| Underweight              | 12 (35.3)             | 19 (41.3)                              | 0.209   |
| Stunted                  | 17 (50)               | 25 (54.3)                              | 0.217   |
| Wasted                   | 3 (8.8)               | 8 (17.4)                               | 0.132   |

CP, cerebral palsy; GMFCS, gross motor functional classification system; MMI, mild motor impairment; SMI, severe motor impairment.

Figure 1. Scatter diagram and correlation analysis of body mass index and weight-for-height percentiles in children with cerebral palsy. Body mass index was directly associated with weight-for-height; P, percentile.
The aim of the present study was to evaluate the physical growth, nutritional status, and the inflammatory parameters of NLR and LMR between CP children living at home compared with those living in a rehabilitation center. The results showed that almost half of the children had one form of malnutrition. The most common type of malnutrition was stunted growth (50% of children at home and 54.3% of children at the center), followed by being underweight (35.3% of children at home and 41.3% of children at center) and wasted (8.8% of children at home and 17.4% of children at center), in agreement with a previous study (19).

Henderson et al (20) reported that residential care children showed improved growth and nutrition scores, lower height Z scores, similar weight and muscle mass, and greater subcutaneous fat Z scores than children living at home. In contrast, the results of the present study showed higher percentages of all forms of malnutrition in CP children living at the rehabilitation center compared with those living at home.

In routine medical practice, CBC and differentiation analysis are commonly used (21,22). In the present study, the severity of...
the motor impairments was associated with higher NLR values, particularly in CP children living at the rehabilitation center. NLR values were significantly higher in CP children with severe motor impairment in the rehabilitation center.

NLR is easy to calculate and can provide useful information. Increases in NLR are indicative of systemic inflammation and physiological stress responses, including an increase in neutrophil counts, a decrease in lymphocyte counts, or both (23). Recently, NLR has been shown to be a good prognostic parameter for assessment of various diseases and was correlated with the treatment or operational outcomes of several clinical conditions, including gastric, liver and colorectal cancer (24-26). Additionally, NLR is positively correlated with several biomarkers of cancer, such as carbohydrate antigen 19-9 and carcinoembryonic antigen, and was associated with advanced stage malignancies (27,28), and overall survival and mortality for various diseases (29,30).

Persistent elevation in inflammation-related proteins during the first two postnatal weeks is associated with the diagnosis of CP, and this pattern of expression was also observed in severely cognitively impaired children (31). CP is a neurodevelopmental disorder and is attributed to injury in the developing brain. Various animal models of CP have shown that the injury to the central nervous system is due to hypoxia-ischemia and

Figure 3. Scatter diagram and correlation analysis of inflammatory parameters in children with cerebral palsy. (A) Total white blood cell counts were positively correlated with NLR. (B) NLR was inversely correlated with lymphocyte-to-monocyte ratio. NLR, neutrophil-to-lymphocyte ratio.

Figure 4. Clustered bar graph of the neutrophil-to-lymphocyte ratio for each GMFCS level between children living at home (gray) and at the rehabilitation center (black). *P<0.05. GMFCS, gross motor functional classification system.
inflammation in the brain (32). Furthermore, elevated inflammatory response-associated proteins may reflect the ongoing damage of the brain cells resulting in brain dysfunction (33). Over half of neuroanatomical abnormalities in CP patients are white matter injuries, cystic periventricular leukomalacia and periventricular hemorrhagic infarction (34). The ELGAN cohort found that systemic inflammation was associated with white matter abnormalities (35). Inflammatory cytokines and inflammation related proteins are present during the neonatal period (36,37). Additionally, levels of IL-1β, IL-6, TNF and CXCL8/IL-8 in children with CP are persistently elevated (8). Unfortunately, the markers mentioned above are rarely used in routine medical practices, and only in research. Thus, our proposed low cost, accessible inflammatory predictor, NLR, may be an improved prognostic parameter for CP; however, additional studies are required to confirm these results.

The present study has some limitations. First, this was a cross-sectional study. Therefore it was not possible to determine a cause-effect relationship. Second, a healthy control group was not available for comparison. Third, the sample size was too small, and thus made identification of significant differences between groups difficult. Finally, it is possible that

Table III. Comparison of NLR and motor impairment severity between children with CP living at home-living and those living at a rehabilitation center.

| Characteristics | Living at home, n=34 | Living at rehabilitation centre, n=46 | P-value |
|-----------------|----------------------|--------------------------------------|---------|
|                 | n (mean ± SEM)       | n (mean ± SEM)                      |         |
| MMI             |                      |                                      |         |
| GMFCS level II  | 3 (0.89±0.63)        | 6 (1.15±0.11)                       | 0.121   |
| GMFCS level III | 13 (1.41±0.15)       | 22 (1.47±0.11)                      | 0.759   |
| Total           | 16 (1.31±0.13)       | 28 (1.4±0.1)                        | 0.38    |
| SMI             |                      |                                      |         |
| GMFCS level IV  | 11 (1.3±0.21)        | 15 (2.23±0.31)                      | 0.014*  |
| GMFCS level V   | 7 (1.21±0.22)        | 3 (3.53±0.22)                       | 0.017*  |
| Total           | 18 (1.26±0.15)       | 18 (2.45±0.29)                      | 0.001b  |
| Overall         | 34 (1.29±0.1)        | 46 (1.81±0.15)                      | 0.003b  |
| P (MMI/SMI)     | 0.437                | 0.006b                              |         |
| P (GMFCS)       | 0.194                | 0.01b                               |         |

P-values for differences between the home-living and rehabilitation center. *P<0.05, **P<0.01. CP, cerebral palsy; GMFCS, Gross Motor Function Classification System; MMI, mild motor impairment; SMI, severe motor impairment; SEM, standard error of the mean.

Figure 5. Scatter diagram and correlation analysis of NLR and GMFCS of the children with cerebral palsy living at the rehabilitation center. NLR was positively correlated with GMFCS. GMFCS, gross motor functional classification system; NLR, neutrophil-to-lymphocyte ratio.
the environment, family participation and emotional stress may be different between the two groups, and this may affect the overall quality of life of CP children.

In conclusion, a high NLR value was associated with severe motor impairment in children with CP, particularly those living in rehabilitation centers. These results suggest that NLR may be used as a prognostic predictor of severe motor impairment. However, further studies on the value and relevance of NLR are required to ascertain whether it can be used to predict treatment and/or surgical outcomes, as well as mortality and survival amongst CP children.

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Availability of data and materials

The datasets used and/or analyzed during the present study are available from the corresponding author on reasonable request.

Authors' contributions

KR, CA, PO, SW, NL and SH conceived and designed the experiments. KR, CA, PO, SW, NL and SH performed the experiments and analyzed the data. CA, NL and SH provided the materials and analytical tools. CA, PO, SW and NL recruited the participants, obtained the written informed consents and collected the clinical data. KR and SH wrote and revised the manuscript. All authors have read and approved the final manuscript.

Ethics approval and consent to participate

The study protocol used in the present study conformed to the ethical standards outlined in the Declaration of Helsinki and was approved by the Institutional Review Board on Human Research of the Faculty of Medicine, Chulalongkorn University. Informed consent was obtained from one of the parents or the legal guardian, as well as from the child prior to his/her participation in the study.

Patient consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

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