Asymmetry of cerebral glucose metabolism in very low-birth-weight infants without structural abnormalities

Jae Hyun Park¹, Chun Soo Kim¹, Kyoung Sook Won², Jungsu S. Oh³, Jae Seung Kim³, Hae Won Kim²*

¹ Department of Pediatrics, Keimyung University Dongsan Medical Center, Daegu, Republic of Korea, ² Department of Nuclear Medicine, Keimyung University Dongsan Medical Center, Daegu, Republic of Korea, ³ Department of Nuclear Medicine, Asan Medical Center, University of Ulsan College of Medicine, Seoul, Korea

* hwkim.nlm@gmail.com

Abstract

Even when structural abnormalities are not observed on the brain magnetic resonance images (MRI) of very low-birth-weight (VLBW) infants, such infants are at increased risk for poor neurodevelopment. The aim of the present study was to evaluate cerebral glucose metabolism in VLBW infants without apparent structural abnormalities on MRI.

Methods

Thirty-six VLBW infants who underwent F-18 fluorodeoxyglucose (F-18 FDG) brain PET and MRI were prospectively enrolled, while infants with evidence of parenchymal brain injury on MRI were excluded. The regional glucose metabolic ratio and asymmetry index were calculated. The asymmetry index more than 10% (right > left asymmetry) or less than -10% (left > right asymmetry) were defined as abnormal. Regional cerebral glucose metabolism were compared between right and left cerebral hemispheres, and between the following subgroups: multiple gestations, premature rupture of membrane, bronchopulmonary dysplasia, and low-grade intraventricular hemorrhage.

Results

In the individual analysis, 21 (58.3%) of 36 VLBW infants exhibited asymmetric cerebral glucose metabolism. Fifteen infants (41.7%) exhibited right > left asymmetry, while six (16.7%) exhibited left > right asymmetry. In the regional analysis, right > left asymmetry was more extensive than left > right asymmetry. The metabolic ratio in the right frontal, temporal, and occipital cortices and right thalamus were significantly higher than those in the corresponding left regions. In the subgroup analyses, the cerebral glucose metabolism in infants with multiple gestations, premature rupture of membrane, bronchopulmonary dysplasia, or low-grade intraventricular hemorrhage were significantly lower than those in infants without these.
Conclusion
VLBW infants without structural abnormalities have asymmetry of cerebral glucose metabolism. Decreased cerebral glucose metabolism are noted in infants with neurodevelopmental risk factors. F-18 FDG PET could show microstructural abnormalities not detected by MRI in VLBW infants.

Introduction
Over the last decade, improvements in perinatal care have resulted in increased rates of survival for very low-birth-weight (VLBW) infants [1, 2]. However, an increase in preterm births and remarkable improvements in survival for infants have outpaced any concomitant decrease in rates of long-term neurodevelopmental disability [3]. As many as 42–47% of cerebral palsy cases in children can be attributed to preterm births [4, 5]. In addition, subtle motor and cognitive dysfunction due to high-grade intraventricular hemorrhage (IVH) or white matter injury has been associated with extreme prematurity, most often resulting in regulatory, attentional, or adaptive impairments [6]. Furthermore, VLBW infants without severe brain injury are at risk for cognitive dysfunction and may require more support than do full-term infants.

Brain magnetic resonance imaging (MRI) is considered the most sensitive and specific modality for diagnosing brain injury and predicting neurodevelopmental outcomes in VLBW infants. The most extensively studied MRI abnormalities shown to predict neurodevelopmental outcomes are white matter lesions (e.g., periventricular leukomalacia [PVL], punctate white matter lesions, and diffuse excessive high signal intensities) [7]. However, although conventional MRI can be used to detect macrostructural brain abnormalities in VLBW infants, such images are unable to detect micropathology in VLBW infants [8]. Indeed, previous studies have demonstrated that some children with cerebral palsy or cognitive dysfunction exhibit no apparent abnormalities on conventional MRI [9, 10].

However, F-18 fluorodeoxyglucose (F-18 FDG) positron emission tomography (PET) imaging may be helpful in detecting cerebral glucose metabolic abnormalities at a much earlier stage, prior to the development of structural and morphologic abnormalities [11]. Previous studies have demonstrated that patterns of cerebral glucose metabolism are correlated with the later development of different types of cerebral palsy in children [12, 13]. Although there have been no reports regarding cerebral glucose metabolism in VLBW infants, it is generally accepted that many neurodegenerative diseases are associated with significantly decreased glucose metabolism on F-18 FDG PET images, even though conventional MRI images reveal no specific abnormalities [14, 15]. Thus, the aim of the present study was to evaluate cerebral glucose metabolism in VLBW infants without structural abnormalities on brain MRI.

Material and methods
Subjects
VLBW infants (<1500 g) born at less than 30 weeks gestation admitted to the neonatal intensive care unit of Keimyung University Dongsan Medical Center between December 2014 and October 2015 were included in the present prospective study. All infants underwent MRI scans at term-equivalent age, and infants who exhibited evidence of parenchymal brain injury on MRI (cystic PVL, congenital cerebral anomaly, or grade 3 or 4 IVH) were excluded [16]. In all infants, F-18 FDG PET was performed at term-equivalent age. The institutional review
board of Dongsan Medical Center approved the present study, and written informed consent was obtained from the parents/guardians of all participants.

**Protocol for \(^{18}\)F-FDG PET acquisition**

A PET/CT system was used to acquire F-18 FDG PET images (Biograph mCT-64, Siemens Healthcare, Knoxville, TN). Infants received intravenous administration of 3.7 MBq/kg. Each infant was sedated using thiopental 40 minutes after F-18 FDG injection, and images were acquired 50 minutes after injection. A non-enhanced low-dose CT scan was obtained for attenuation correction and localization. A light, foam-rubber holder was used for fixation of the head. The PET images were reconstructed iteratively using ordered subset expectation maximization. Attenuation correction of PET images was performed using attenuation data from CT images. All fusion images were viewed using dedicated workstations for each PET/CT system.

**Quantitative analyses**

Image processing was performed using SPM12 (Wellcome Department of Imaging Neuroscience, Institute of Neurology, University College London) within MATLAB 2013a (MathWorks Inc., MA, USA) and MRicro version 1.37 (Chris Rorden, Columbia, SC, USA, [www.mricro.com](http://www.mricro.com)). Quantitative analyses were conducted on volumes of interest (VOIs) using the software program PMOD (PMOD Technologies Ltd, Zurich, Switzerland), as previously described [17]. All reconstructed PET images were co-registered to the corresponding MR images and spatially normalized to a neonate brain atlas using a neonate brain MRI template [18]. For analysis of cerebral glucose metabolism, 24 regional VOIs were identified via the neonate automated anatomic labeling (nAAL) template, as previously described [18, 19]. We investigated the following regional VOIs: the central region; lateral, medial, and orbitofrontal lobes; lateral and medial temporal lobe; lateral parietal lobe; lateral and medial occipital lobe; caudate nucleus; putamen; and thalamus. The VOI template was automatically applied directly to the spatially normalized individual PET images for the analysis of cerebral glucose metabolism.

The activity concentration was calculated for each VOI of the F-18 FDG PET images. The activity concentration in the whole brain was also calculated as a reference. The regional cerebral glucose metabolic ratio (CMRgl) for each VOI was calculated as follows: CMRgl = count of regional VOI / count of whole brain. The asymmetry index (AI\(_{\text{voi}}\)) for each VOI was calculated as follows: AI\(_{\text{voi}}\) = (count of right VOI—count of left VOI) / (count of right VOI + count of left VOI) × 200. In the regional analysis, AI\(_{\text{voi}}\) values more than 10% (right > left asymmetry) or less than -10% (left > right asymmetry) were defined as abnormal [20]. The asymmetry of cerebral glucose metabolism was also evaluated using voxel based morphometry (VBM): We created mirror images of each PET image, following which we spatially normalized both the original PET and mirror images. Overlay masks were generated from the averaged spatially normalized PET image using a region-growing algorithm, as previously described [21]. The voxelwise asymmetry index (AI\(_{\text{vbm}}\)) was calculated using the following formula: AI\(_{\text{vbm}}\) = (\(\sum X_{\text{orig}} - (\sum X_{\text{mirror}})\) / (\(\sum X_{\text{orig}} + \sum X_{\text{mirror}}\)) × 200, where \(\text{orig}\) and \(\text{mirror}\) represent the original PET and mirror images, respectively, and \(X\) denotes voxel intensity of the PET image. The mean values of AI\(_{\text{vbm}}\) were overlaid onto a standardized neonate brain MRI.

In the individual analysis, infants with abnormal AI\(_{\text{voi}}\) values in any regional VOI were classified into the abnormal AI\(_{\text{voi}}\) group. We also compared regional CMRgl between VLBW infants with and without risk factors for poor neurodevelopmental outcome, including premature rupture of membrane (PROM), bronchopulmonary dysplasia (BPD), multiple gestations (MG), grade 1 or 2 IVH.
Statistical analyses

The regional CMRgl values were compared between the left and right cerebral hemispheres using paired samples t-tests. The regional CMRgl values were compared between each subgroup (MG, PROM, BPD, and IVH) using two independent t-tests. The P values were corrected for multiple comparisons via false discovery rate correction. One-sample t-tests were used to determine whether the AI\textsubscript{ROI} of each region was higher than 0. A P value of less than 0.05 was considered to indicate statistical significance. Data for all study variables are expressed as the mean ± SD.

Results

Characteristics

Of the 96 live-born VLBW infants eligible for participation, 80 (83.3%) survived to term-equivalent age. Among these, the parents of 29 infants refused to participate in the study, and 15 further infants were excluded due to evidence of parenchymal brain lesions on MRI: cystic PVL (n = 12) and congenital cerebral anomaly (n = 3). Therefore, the final sample consisted of 36 VLBW infants who underwent F-18 FDG brain PET (Fig 1). The mean GA was 27.2 ± 1.3 weeks, and mean birth weight was 925.0 g. The mean chronological age at evaluation of F-18 FDG PET was 10.0 weeks. No significant differences in gestational age, birth weight, or sex were observed among the PROM, BPD, MG, and IVH subgroups. Table 1 summarizes the clinical characteristics of the infants.

Cerebral glucose metabolism

The CMRgl of the right medial and orbitofrontal, right lateral temporal, right lateral and medial occipital lobes, and right thalamus were significantly higher than those of the corresponding regions in the left hemisphere (P < 0.001, P < 0.001, P = 0.002, P = 0.002, P = 0.008, and P < 0.001, respectively). The CMRgl of the right putamen was significantly lower than that of the left putamen (P = 0.019). Table 2 shows the mean AI\textsubscript{ROI} of each VOI in VLBW infants. The VBM analysis also revealed right dominant asymmetry of cerebral glucose metabolism.

Fig 1. Flow diagram of the study population. PVL: periventricular leukomalacia.

https://doi.org/10.1371/journal.pone.0186976.g001
The mean AI_vbm values in the right cerebral hemisphere were positive, while the mean AI_vbm values in the left cerebral hemisphere were negative (Fig 2).

In the individual analysis, 21 (58.3%) of 36 VLBW infants exhibited abnormal AI_voi values. Fifteen infants (41.7%) exhibited right > left asymmetry for cerebral glucose metabolism (Table 3). These asymmetries were observed in the central region, frontal lobes, temporal lobes, occipital lobe, caudate nucleus, putamen, and thalamus. Fig 3 includes a representative F-18 FDG brain PET image of a VLBW infant with right > left asymmetry. Six infants (16.7%) exhibited left > right asymmetry. These asymmetries were observed in the central region, frontal lobe, temporal lobe, parietal lobe, caudate nucleus, putamen, and thalamus. In the regional analysis, right > left asymmetry was observed more frequently and was more extensive than left > right asymmetry. The AI_voi values of all brain regions were significantly higher than 0 (P < 0.001).

In the subgroup analyses, the PROM group had a significantly lower CMRgl in the left lateral parietal lobe than did the non-PROM group (P = 0.030) (Fig 4). The BPD group had a significantly lower CMRgl in the right central region and right lateral temporal lobe than did

| Characteristic                              | Infant data |
|--------------------------------------------|-------------|
| Mean gestational age at birth (wk)         | 27.2 ± 1.3  |
| Mean birth weight (g)                      | 925.0 ± 207.2 |
| Chronological age at F-18 FDG PET (wk)     | 10.0 ± 2.2  |
| Male sex, no. (%)                          | 14 (38.9)   |
| Premature rupture of membrane, no. (%)     | 14 (38.9)   |
| Bronchopulmonary dysplasia, no. (%)        | 8 (22.2)    |
| Multiple gestation, no. (%)                | 13 (36.1)   |
| Apgar score                                |             |
| at 1 min, mean (SD)                        | 5.0 ± 1.4   |
| at 5 min, mean (SD)                        | 7.3 ± 1.3   |

Table 2. Cerebral metabolic ratio and asymmetry index in very low-birth-weight infants.

| VOI*                      | Cerebral metabolic ratio | Asymmetry index (%) | P value† |
|---------------------------|--------------------------|---------------------|---------|
|                           | Left                     | Right               |         |
| Central region            | 1.06 ± 0.05              | 1.07 ± 0.04         | 0.6 ± 5.0 | 0.500 |
| Lateral frontal lobe      | 0.90 ± 0.04              | 0.90 ± 0.05         | -0.6 ± 4.2 | 0.477 |
| Medial frontal lobe       | 0.89 ± 0.05              | 0.92 ± 0.06         | 3.8 ± 4.9 | < 0.001† |
| Orbital frontal lobe      | 0.89 ± 0.04              | 0.91 ± 0.04         | 2.0 ± 2.1 | < 0.001† |
| Lateral temporal lobe     | 1.03 ± 0.05              | 1.06 ± 0.03         | 2.9 ± 4.5 | 0.002‡ |
| Medial temporal lobe      | 1.19 ± 0.12              | 1.20 ± 0.12         | 2.3 ± 3.1 | 0.231 |
| Lateral parietal lobe     | 0.87 ± 0.04              | 0.86 ± 0.04         | -1.60 ± 3.2 | 0.06 |
| Lateral occipital lobe    | 0.87 ± 0.05              | 0.89 ± 0.04         | 3.0 ± 4.7 | 0.002‡ |
| Medial occipital lobe     | 0.96 ± 0.04              | 0.98 ± 0.03         | 1.4 ± 2.8 | 0.008‡ |
| Caudate nucleus           | 0.89 ± 0.09              | 0.90 ± 0.09         | 1.0 ± 7.2 | 0.477 |
| Putamen                   | 1.35 ± 0.12              | 1.32 ± 0.11         | -2.6 ± 5.8 | 0.019‡ |
| Thalamus                  | 1.35 ± 0.19              | 1.42 ± 0.18         | 5.3 ± 7.5 | < 0.001† |

*VOI: volume of interest.
†Comparison of cerebral metabolic ratios between right and left hemispheres.
‡Statistically significant results.

https://doi.org/10.1371/journal.pone.0186976.t001

https://doi.org/10.1371/journal.pone.0186976.t002
the non-BPD group ($P = 0.038$ and $P = 0.035$). The MG and IVH groups had a significantly lower CMRgl in the right medial occipital lobe than did the non-MG and non-IVH groups ($P = 0.029$ and $P = 0.011$, respectively). The MG and IVH groups had a significantly higher CMRgl in the left putamen than did the non-MG and non-IVH groups ($P = 0.033$ and $P = 0.038$, respectively). There were no significant differences in AI\textsubscript{voi} between any of the subgroups. S1 and S2 Tables provide full results of the subgroup analyses.

![Voxelwise asymmetry index (AI\textsubscript{vbm}) of cerebral glucose metabolism in a very low-birth-weight infant.](https://doi.org/10.1371/journal.pone.0186976.g002)

**Fig 2.** Voxelwise asymmetry index (AI\textsubscript{vbm}) of cerebral glucose metabolism in a very low-birth-weight infant. Voxel-based morphometry revealed right dominant asymmetry of cerebral glucose metabolism. Red indicates a region of relatively high glucose metabolism (where AI\textsubscript{vbm} is positive) in comparison to corresponding regions. Blue indicates a region of relatively low glucose metabolism (where AI\textsubscript{vbm} is negative) in comparison to corresponding regions.

| Region                  | Left > Right | Normal | Right > Left |
|-------------------------|--------------|--------|--------------|
| VLBW infants*           | 6            | 15     | 15           |
| Central region          | 2            | 33     | 1            |
| Lateral frontal lobe    | 0            | 35     | 1            |
| Medial frontal lobe     | 1            | 31     | 4            |
| Orbital frontal lobe    | 0            | 36     | 0            |
| Lateral temporal lobe   | 0            | 35     | 1            |
| Medial temporal lobe    | 1            | 33     | 2            |
| Lateral parietal lobe   | 1            | 35     | 0            |
| Lateral occipital lobe  | 0            | 34     | 2            |
| Medial occipital lobe   | 0            | 36     | 0            |
| Caudate nucleus         | 1            | 32     | 3            |
| Putamen                 | 2            | 33     | 1            |
| Thalamus                | 2            | 25     | 9            |

*VLBW infant: vere-low-birth-weight infant.

https://doi.org/10.1371/journal.pone.0186976.t003
Discussion

In the present study, we demonstrated that more than half of the included VLBW infants exhibited asymmetric cerebral glucose metabolism, although there was no apparent evidence of parenchymal brain injury on brain MRI. Furthermore, the present study revealed that VLBW infants with neurodevelopmental risk factors had a significantly lower CMRgl in the left lateral parietal lobe than those without risk factors. Abnormalities in glucose metabolism that extend beyond the site of a structural lesion may help clinicians to determine the extent of injury in VLBW infants [11]. However, few studies have evaluated cerebral glucose metabolism in preterm infants, and these studies primarily included infants with severe brain injury [22] or a small number of low birth weight infants [13, 23]. The present study is the first to utilize F-18 FDG PET for the evaluation of cerebral glucose metabolism in a sufficient number of VLBW infants without structural abnormalities on brain MRI.

Cerebral glucose metabolism is usually homogeneous and symmetrical in adults with typical cognitive function/development. Generally, metabolic asymmetry is considered to be abnormal if it exceeds 10% [24]. Slight asymmetries of cerebral glucose metabolism have been observed in the Wernicke area, frontal eye fields, and angular gyrus, with a prevalence of generally less than 10% [20]. In accordance with the findings of previous studies, we observed that VLBW infants exhibited relatively high glucose metabolism in the thalamus, putamen, medial temporal lobe, and central region [13, 18]. However, in present study, 58.3% of VLBW infants exhibited abnormal AI vo values (>10%), although there was no evidence of parenchymal brain injury on brain MRI. Thus, our findings suggest that microstructural white matter injury—which cannot be detected using MRI—disrupts corticostriatal connectivity and results
in decreased activity of the cerebral cortex. This hypothesis is consistent with the known pathogenesis of macrostructural white matter injury in preterm infants, which can impair myelination of projection and association fibers or interfere with cortical neuronal development [25, 26].

Cerebral palsy develops in approximately 5 to 15% of surviving VLBW infants [5], while cognitive delay has been reported to occur in as many as 40% of these children upon reaching school age [3]. The characteristic symptoms of cerebral palsy are decreased or increased motor activity and abnormal posture, which can be symmetric or asymmetric [27]. The clinical phenotype of cerebral palsy can be predicted in VLBW infants based on patterns of cerebral metabolism.
glucose metabolism on F-18 FDG PET images. Kerrigan et al. reported that bilateral thalamic hypometabolism predicts the spastic diplegic type of cerebral palsy, and that the topography of focal cortical areas exhibiting hypometabolism is correlated with specific cognitive deficits [28]. Similar to the findings of previous studies, our results demonstrated that 41.7% of VLBW infants exhibited right dominant asymmetry, while 16.7% exhibited left dominant asymmetry. In addition, right dominant asymmetry was observed more frequently and was more extensive than left dominant asymmetry. Such findings may be explained by factors unique to the circulation of each fetus, as both prenatal factors and perinatal hypoxia-ischemia are known to play a major role in most cases of cerebral palsy [29]. In fetal circulation, the right common carotid artery is located closer to the left ventricle than the left common carotid artery and has two sources of blood flow from the left ventricle and ductus arteriosus [30]. A relatively weak blood supply from the left common carotid artery can easily cause microstructural white matter injuries in prenatal and perinatal hypoxic states [31]. The unique cerebrovascular anatomy and physiology of premature infants underlies the exquisite sensitivity of the white matter to the abnormal milieu of preterm extrauterine life, particularly with regard to ischemia and inflammation [32].

Even in the absence of structural brain abnormalities, VLBW infants remain at high risk for cognitive dysfunction, impaired language development, and neurobehavioral impairment [5, 10]. Previously identified risk factors for poor cognitive development in VLBW infants include chorioamnionitis, MG, PROM, BPD, and IVH [5, 33, 34]. Many studies have revealed that clinical risk factors can reliably predict neurodevelopmental impairment in VLBW infants [33–35]. Lodha et al. [35] reported that the Clinical Risk Index for Babies score predicted major neurodevelopmental impairment (odds ratio, 1.57; 95% CI, 1.26–3.01) and poor outcome (odds ratio, 1.46; 95% CI, 1.31–1.71) at 36 months’ corrected age. In agreement with previous studies, our results revealed that glucose metabolism in the cerebral cortices of VLBW infants with PROM, BPD, MG or IVH were significantly lower than those in infants without risk factors. The decreased glucose metabolism was shown in the lateral parietal and temporal lobes, the medial occipital lobes and central region depending on the risk factors. This result provides indirect evidence that F-18 FDG PET can detect microstructural abnormalities, which are related with poor neurodevelopmental outcome, in VLBW infants.

The present study possesses some limitation of note. First, we did not perform VOI analysis based on partial volume correction. The partial volume effect can influence the quantitative analysis of PET images, resulting in a lower CMRgl than the actual value. However, we believe that the partial volume effect on the AI_{voi} was minimal in the present study: Bilateral cerebral hemispheres without structural abnormalities on MRI were analyzed using the same method, and the partial volume effect on CMRgl was similar in both cerebral hemispheres. Second, we did not directly investigate the association between long-term neurodevelopmental outcomes and cerebral glucose metabolism in VLBW infants. Although the present study revealed asymmetry of cerebral glucose metabolism, further longitudinal studies are required to relate F-18 FDG PET images to long-term neurodevelopmental outcomes. Finally, the radiation exposure to the infants during F-18 FDG PET may raise concerns if proposed for clinical use in the prediction of neurodevelopmental outcomes. However, most PET scans in neonates can be accomplished using effective doses approximately equal to the yearly background radiation exposure, or less than the exposure during a clinical CT scan of the head [36, 37].

**Conclusion**

Our findings demonstrated that VLBW infants without structural abnormalities have asymmetry of cerebral glucose metabolism. Decreased cerebral glucose metabolism are noted in infants
with neurodevelopmental risk factors. F-18 FDG PET could show microstructural abnormalities not detected by MRI in VLBW infants.

Supporting information

S1 Table. Comparison of metabolic ratios between PROM and non-PROM groups and between BPD and non-BPD groups.
(DOCX)

S2 Table. Comparison of metabolic ratios between MG and non-MG groups and between IVH and non-IVH groups.
(DOCX)

S1 Data. Data underlying the findings.
(XLSX)

Acknowledgments

Ethical approval: All procedures involving human participants were conducted in accordance with the ethical standards of the institutional research committee and with the 1964 Declaration of Helsinki.

Informed consent: Written informed consent was obtained from the parents/guardians of all participants.

Author Contributions

Conceptualization: Jae Hyun Park, Hae Won Kim.

Data curation: Jae Hyun Park.

Formal analysis: Chun Soo Kim, Hae Won Kim.

Funding acquisition: Jungsu S. Oh, Hae Won Kim.

Investigation: Jae Seung Kim, Hae Won Kim.

Methodology: Kyoung Sook Won, Hae Won Kim.

Project administration: Jae Hyun Park, Hae Won Kim.

Resources: Jae Hyun Park.

Software: Jungsu S. Oh, Hae Won Kim.

Supervision: Jae Seung Kim.

Validation: Kyoung Sook Won.

Visualization: Jungsu S. Oh.

Writing – original draft: Jae Hyun Park, Hae Won Kim.

Writing – review & editing: Chun Soo Kim, Kyoung Sook Won, Hae Won Kim.

References

1. Goldenberg RL, Culhane JF, Iams JD, Romero R. Epidemiology and causes of preterm birth. Lancet. 2008; 371(9608):75–84. https://doi.org/10.1016/S0140-6736(08)60074-4 PMID: 18177778.

2. Beck S, Wojdyla D, Say L, Betran AP, Merialdi M, Requejo JH, et al. The worldwide incidence of preterm birth: a systematic review of maternal mortality and morbidity. Bull World Health Organ. 2010; 88 (1):31–8. https://doi.org/10.2471/BLT.08.062954 PMID: 20426351
3. Marlow N, Wolke D, Bracewell MA, Samara M, Group EPS. Neurologic and developmental disability at six years of age after extremely preterm birth. N Engl J Med. 2005; 352(1):9–19. https://doi.org/10.1056/NEJMoa041367 PMID: 15635108.

4. Behrman RE, Stith Butler A. Institute of Medicine Committee on understanding premature birth and assuring healthy outcomes board on health sciences outcomes: preterm birth: causes, consequences, and prevention. Preterm birth: causes, consequences, and prevention, National Academies Press, Washington, DC. 2007.

5. Linsell L, Malouf R, Morris J, Kurinczuk JJ, Marlow N. Prognostic Factors for Poor Cognitive Development in Children Born Very Preterm or With Very Low Birth Weight: A Systematic Review. JAMA Pediatr. 2015; 169(12):1162–72. https://doi.org/10.1001/jamapediatrics.2015.2175 PMID: 26457641.

6. Msall ME, Park JJ. The spectrum of behavioral outcomes after extreme prematurity: regulatory, attention, social, and adaptive dimensions. Semin Perinatol. 2008; 32(1):42–50. https://doi.org/10.1053/j.semperi.2007.12.006 PMID: 18249239.

7. Woodward LJ, Anderson PJ, Austin NC, Howard K, Inder TE. Neonatal MRI to predict neurodevelopmental outcomes in preterm infants. New England Journal of Medicine. 2006; 355(7):685–94. https://doi.org/10.1056/NEJMoa053792 PMID: 16914704.

8. Son SM, Ahn YH, Sakong J, Moon HK, Ahn SH, Lee H, et al. Diffusion tensor imaging demonstrates focal lesions of the corticospinal tract in hemiparetic patients with cerebral palsy. Neurosci Lett. 2007; 420(1):34–8. https://doi.org/10.1016/j.neulet.2007.04.054 PMID: 17512661.

9. Leitner Y, Weinstein M, Myers V, Ullel S, Geva K, Berger I, et al. Diffuse excessive high signal intensity in low-risk preterm infants at term-equivalent age does not predict outcome at 1 year: a prospective study. Neuroradiology. 2014; 56(8):669–78. https://doi.org/10.1007/s00234-014-1373-8 PMID: 24823447.

10. Korzeniewski SJ, Birbeck G, DeLano MC, Potchen MJ, Paneth N. A systematic review of neuroimaging for cerebral palsy. J Child Neurol. 2008; 23(2):216–27. https://doi.org/10.1177/0883073807307983 PMID: 18263759.

11. Chugani HT. Positron emission tomography scanning: applications in newborns. Clinics in perinatology. 1993; 20(2):395–409. PMID: 8358958.

12. Altman DI, Volpe JJ. Positron emission tomography in newborn infants. Clin Perinatol. 1991; 18(3):549–62. PMID: 1934855.

13. Shi Y, Jin RB, Zhao JN, Tang SF, Li HQ, Li TY. Brain positron emission tomography in preterm and term newborn infants. Early Hum Dev. 2009; 85(7):697–72. PMID: 19269116.

14. Silverman DH. Brain 18F-FDG PET in the diagnosis of neurodegenerative dementias: comparison with perfusion SPECT and with clinical evaluations lacking nuclear imaging. J Nucl Med. 2004; 45(5):940–607. Epub 2004/04/10. PMID: 15073255.

15. Chetelat G, Desgranges B, de la Sayette V, Viader F, Eustache F, Baron JC. Mild cognitive impairment: Can FDG-PET predict who is to rapidly convert to Alzheimer’s disease? Neurology. 2003; 60(8):1374–7. PMID: 12707450.

16. Bax M, Tydeman C, Flodmark O. Clinical and MRI correlates of cerebral palsy: the European Cerebral Palsy Study. JAMA. 2006; 296(13):1602–8. https://doi.org/10.1001/jama.296.13.1602 PMID: 17018805.

17. Mikolajczyk K, Szabatin M, Rudnicki P, Grodzki M, Burger C. A JAVA environment for medical image data analysis: initial application for brain PET quantitation. Med Inform (Lond). 1998; 23(3):207–14. PMID: 9785322.

18. Shi F, Yap PT, Wu G, Jia H, Gilmore JH, Lin W, et al. Infant brain atlases from neonates to 1- and 2-year-olds. PLoS One. 2011; 6(4):e18746. https://doi.org/10.1371/journal.pone.0018746 PMID: 21533194.

19. Tzourio-Mazoyer N, Landeau B, Papathanassiou D, Crivello F, Etard O, Delcroix N, et al. Automated anatomical labeling of activations in SPM using a macroscopic anatomical parcellation of the MNI MRI single-subject brain. Neuroimage. 2002; 15(1):273–89. https://doi.org/10.1006/nimg.2001.0978 PMID: 11771995.

20. Ivancic V, Alavi A, Souder E, Mozley PD, Gur RE, Benard F, et al. Regional cerebral glucose metabolism in healthy volunteers determined by fluorodeoxyglucose positron emission tomography: appearance and variance in the transaxial, coronal, and sagittal planes. Clin Nucl Med. 2000; 25(8):596–602. PMID: 10944013.

21. Wilke M, Lidzba K. LI-tool: a new toolbox to assess lateralization in functional MR-data. J Neurosci Methods. 2007; 163(1):128–36. https://doi.org/10.1016/j.jneumeth.2007.01.026 PMID: 17386945.
22. Higuchi Y, Maihara T, Hattori H, Furusho K, Okazawa H, Ishizu K, et al. [18F]-fluorodeoxyglucose-positron emission tomography findings in protein infants with severe periventricular leukomalacia and hypsarrhythmia. Eur J Pediatr. 1997; 156(3):236–8. PMID: 9083768.

23. Chugani HT. Positron emission tomography scanning: applications in newborns. Clin Perinatol. 1993; 20(2):395–409. PMID: 8358958.

24. Shen X, Liu H, Hu Z, Hu H, Shi P. The relationship between cerebral glucose metabolism and age: report of a large brain PET data set. PloS one. 2012; 7(12):e51517. https://doi.org/10.1371/journal.pone.0051517 PMID: 23284706

25. Volpe JJ. Encephalopathy of prematurity includes neuronal abnormalities. Pediatrics. 2005; 116(1):221–5. https://doi.org/10.1542/peds.2005-0191 PMID: 15995055.

26. Schmahmann JD, Smith EE, Eichler FS, Filley CM. Cerebral white matter. Annals of the New York Academy of Sciences. 2008; 1142(1):266–309.

27. Ahnsjo S. [Cerebral palsy. VI. Psychiatric aspects]. Nord Med. 1955; 54(44):1657–8. PMID: 13280139.

28. Kerrigan JF, Chugani HT, Phelps ME. Regional cerebral glucose metabolism in clinical subtypes of cerebral palsy. Pediatr Neurol. 1991; 7(6):415–25. PMID: 1797007.

29. Nelson KB, Blair E. Prenatal Factors in Singletons with Cerebral Palsy Born at or near Term. N Engl J Med. 2015; 373(10):946–53. https://doi.org/10.1056/NEJMra1505261 PMID: 26332549.

30. Degani S. Fetal cerebrovascular circulation: a review of prenatal ultrasound assessment. Gynecol Obstet Invest. 2008; 66(3):184–96. https://doi.org/10.1159/000143157 PMID: 18607112.

31. McClure MM, Riddle A, Manese M, Luo NL, Rovik DA, Kelly KA, et al. Cerebral blood flow heterogeneity in preterm sheep: lack of physiologic support for vascular boundary zones in fetal cerebral white matter. J Cereb Blood Flow Metab. 2008; 28(5):995–1008. https://doi.org/10.1038/sj.jcbfm.9600597 PMID: 18091757

32. Khwaja O, Volpe JJ. Pathogenesis of cerebral white matter injury of prematurity. Arch Dis Child Fetal Neonatal Ed. 2008; 93(2):F153–61. https://doi.org/10.1136/adc.2006.108837 PMID: 18296574

33. Hayakawa M, Ito Y, Saito S, Mitsuda N, Hosono S, Yoda H, et al. Incidence and prediction of outcome in hypoxic-ischemic encephalopathy in Japan. Pediatr Int. 2014; 56(2):215–21. https://doi.org/10.1111/ped.12233 PMID: 24127879

34. Xiong T, Gonzalez F, Mu DZ. An overview of risk factors for poor neurodevelopmental outcome associated with prematurity. World J Pediatr. 2012; 8(4):293–300. https://doi.org/10.1007/s12519-012-0372-2 PMID: 23151855.

35. Lodha A, Sauve R, Chen S, Tang S, Christianson H. Clinical Risk Index for Babies score for the prediction of neurodevelopmental outcomes at 3 years of age in infants of very low birthweight. Dev Med Child Neurol. 2009; 51(11):895–900. https://doi.org/10.1111/j.1469-8749.2009.03284.x PMID: 19416333.

36. Ruotsalainen U, Suohon-Polvi H, Eronen E, Kinnala A, Bergman J, Haaparanta M, et al. Estimated radiation dose to the newborn in FDG-PET studies. J Nucl Med. 1996; 37(2):87–93. PMID: 8667081.

37. Xia T, Alessio AM, De Man B, Manjeshwar R, Asma E, Kinahan PE. Ultra-low dose CT attenuation correction for PET/CT. Phys Med Biol. 2012; 57(2):309–28. https://doi.org/10.1088/0031-9155/57/2/309 PMID: 22156174