

University of Groningen

Altered neurodevelopmental DNA methylation status after fetal growth restriction with brain-sparing

Richter, Anne E; Bekkering-Bauer, Iris; Verkaik-Schakel, Rikst Nynke; Leeuwerke, Mariëtte; Tanis, Joziën C; Bilardo, Caterina M; Kooi, Elisabeth M W; Scherjon, Sicco A; Bos, Arend F; Plösch, Torsten

Published in:

Journal of developmental origins of health and disease

DOI:

[10.1017/S2040174421000374](https://doi.org/10.1017/S2040174421000374)

IMPORTANT NOTE: You are advised to consult the publisher's version (publisher's PDF) if you wish to cite from it. Please check the document version below.

Document Version

Publisher's PDF, also known as Version of record

Publication date:

2022

[Link to publication in University of Groningen/UMCG research database](#)

Citation for published version (APA):

Richter, A. E., Bekkering-Bauer, I., Verkaik-Schakel, R. N., Leeuwerke, M., Tanis, J. C., Bilardo, C. M., Kooi, E. M. W., Scherjon, S. A., Bos, A. F., & Plösch, T. (2022). Altered neurodevelopmental DNA methylation status after fetal growth restriction with brain-sparing. *Journal of developmental origins of health and disease*, 13(3), 378-389. <https://doi.org/10.1017/S2040174421000374>

Copyright

Other than for strictly personal use, it is not permitted to download or to forward/distribute the text or part of it without the consent of the author(s) and/or copyright holder(s), unless the work is under an open content license (like Creative Commons).

The publication may also be distributed here under the terms of Article 25fa of the Dutch Copyright Act, indicated by the "Taverne" license. More information can be found on the University of Groningen website: <https://www.rug.nl/library/open-access/self-archiving-pure/taverne-amendment>.

Take-down policy

If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.

Altered neurodevelopmental DNA methylation status after fetal growth restriction with brain-sparing

Original Article

Cite this article: Richter AE, Bekkering-Bauer I, Verkaik-Schakel RN, Leeuwerke M, Tanis JC, Bilardo CM, Kooi EMW, Scherjon SA, Bos AF, and Plösch T. Altered neurodevelopmental DNA methylation status after fetal growth restriction with brain-sparing. *Journal of Developmental Origins of Health and Disease* doi: [10.1017/S2040174421000374](https://doi.org/10.1017/S2040174421000374)

Received: 13 October 2020

Revised: 25 April 2021


Accepted: 7 June 2021

Keywords:

Fetal growth restriction (FGR); brain-sparing; neurodevelopment; DNA methylation

Address for correspondence:

Anne E. Richter, Division of Neonatology, Beatrix Children's Hospital, University Medical Center Groningen, Hanzeplein 1, 9713 GZ Groningen, The Netherlands.
Email: a.e.richter@umcg.nl

Anne E. Richter¹ , Iris Bekkering-Bauer^{1,2}, Rikst Nynke Verkaik-Schakel², Mariëtte Leeuwerke², Jozien C. Tanis², Caterina M. Bilardo^{2,3}, Elisabeth M. W. Kooi¹, Siccio A. Scherjon², Arend F. Bos¹ and Torsten Plösch²

¹Division of Neonatology, Beatrix Children's Hospital, University Medical Center Groningen, University of Groningen, Groningen, The Netherlands; ²Department of Obstetrics and Gynecology, University Medical Center Groningen, University of Groningen, Groningen, The Netherlands and ³Department of Obstetrics and Gynecology, Amsterdam University Medical Center, VU University Medical Center, Amsterdam, The Netherlands

Abstract

It is under debate how preferential perfusion of the brain (brain-sparing) in fetal growth restriction (FGR) relates to long-term neurodevelopmental outcome. Epigenetic modification of neurotrophic genes by altered fetal oxygenation may be involved. To explore this theory, we performed a follow-up study of 21 FGR children, in whom we prospectively measured the prenatal cerebroplacental ratio (CPR) with Doppler sonography. At 4 years of age, we tested their neurodevelopmental outcome using the Wechsler Preschool and Primary Scale of Intelligence, the Child Behavior Checklist, and the Behavior Rating Inventory of Executive Function. In addition, we collected their buccal DNA to determine the methylation status at predefined genetic regions within the genes hypoxia-inducible factor-1 alpha (*HIF1A*), vascular endothelial growth factor A (*VEGFA*), erythropoietin (*EPO*), EPO-receptor (*EPOR*), brain-derived neurotrophic factor (*BDNF*), and neurotrophic tyrosine kinase, receptor, type 2 (*NTRK2*) by pyrosequencing. We found that FGR children with fetal brain-sparing (CPR <1, $n = 8$) demonstrated a trend ($0.05 < p < 0.1$) toward hypermethylation of *HIF1A* and *VEGFA* at their hypoxia-response element (HRE) compared with FGR children without fetal brain-sparing. Moreover, in cases with fetal brain-sparing, we observed statistically significant hypermethylation at a binding site for cyclic adenosine monophosphate response element binding protein (CREB) of *BDNF* promoter exon 4 and hypomethylation at an HRE located within the *NTRK2* promoter (both $p < 0.05$). Hypermethylation of *VEGFA* was associated with a poorer Performance Intelligence Quotient, while hypermethylation of *BDNF* was associated with better inhibitory self-control (both $p < 0.05$). These results led us to formulate the hypothesis that early oxygen-dependent epigenetic alterations due to hemodynamic alterations in FGR may be associated with altered neurodevelopmental outcome in later life. We recommend further studies to test this hypothesis.

Introduction

Fetal growth restriction (FGR) is a serious complication of pregnancy, often followed by altered brain structure, cognitive deficits, motor disability, and neuropsychological disorders.¹ Several genetic and environmental factors may induce FGR, the most common being placental dysfunction, which leads to abnormal maternal–fetal exchange and fetal hypoxia.² Compensatory fetal hemodynamic redistribution with cerebral vasodilation attempts to spare the fetal brain from hypoxic damage, which is known as fetal brain-sparing.³ It is still under debate whether and how this may benefit the developing brain.

In the same cohort of FGR children as used in this study, we have previously observed that the sonographic presence of fetal brain-sparing was associated with higher neonatal cerebral oxygen saturations.⁴ At follow-up, fetal brain-sparing and higher cerebral oxygen saturations were related to better behavior and executive functioning at 4 years of age.⁵ However, high cerebral oxygen saturation levels have also been associated with poorer cognition, in particular, a poorer Performance Intelligence Quotient (IQ).^{5,6}

There is evidence that the early-life environment influences long-term neurodevelopment through epigenetic mechanisms.^{7,8} Among these mechanisms, methylation of cytosine-phosphate-guanine (CpG) dinucleotides within promoter regions contributes to maintaining long-lasting states of gene repression.⁹ Oxygen levels have shown to alter DNA methylation and directly affect the activity of transcription factor hypoxia-inducible factor-1 α (HIF1 α), which has popular target genes exerting neurotrophic functions.^{10–13}

© The Author(s), 2021. Published by Cambridge University Press in association with International Society for Developmental Origins of Health and Disease. This is an Open Access article, distributed under the terms of the Creative Commons Attribution-NonCommercial-NoDerivatives licence (<http://creativecommons.org/licenses/by-nc-nd/4.0/>), which permits non-commercial re-use, distribution, and reproduction in any medium, provided the original work is unaltered and is properly cited. The written permission of Cambridge University Press must be obtained for commercial re-use or in order to create a derivative work.

The aim of this study was therefore to analyze whether in our FGR cohort, fetal brain-sparing was associated with altered methylation at predefined loci of the genes encoding HIF1 α , its neurodevelopmentally important target genes (vascular endothelial growth factor A [VEGFA], erythropoietin [EPO], brain-derived neurotrophic factor [BDNF], and neurotrophic tyrosine kinase, receptor, type 2 [NTRK2]) and the EPO-receptor gene (EPOR) at 4 years of age. We used buccal DNA as it is convenient to sample at this age and has been demonstrated to show methylation patterns comparable with neuronal DNA.¹⁴ We further explored whether differential methylation of these genetic loci was related to neurodevelopmental outcome (intelligence, behavior, and executive functions) at this age, which may explain our previous findings.

Method

Study design and population

This was a prospective observational cohort study including 21 children born following FGR between June 2012 and May 2014 at the University Medical Center Groningen (UMCG), The Netherlands.

Antenatal inclusion was based on FGR diagnosed by a fetal abdominal circumference or estimated fetal weight below the 10th percentile or by flattening of the fetal growth curve by more than 30 percentiles compared with the preceding examination. Exclusion criteria were structural or chromosomal abnormalities, multiple pregnancy, or evidence of intrauterine infection. At the age of 4 years, surviving children with complete perinatal hemodynamic assessment (antenatal cerebroplacental Doppler and postnatal cerebral oxygenation measurements), consent for follow-up, and sufficient knowledge of the Dutch language received neurodevelopmental testing. Buccal DNA was sampled during this visit. Participants declining DNA sampling were excluded for the purpose of this study.

We chose to assess the children at the age of 4 years, as children enter school at this age in The Netherlands and are able to participate in the Wechsler Preschool and Primary Scale of Intelligence (WPPSI) IQ test. Written informed parental consent for participation was obtained and the study was approved by the Medical Ethical Committee of the UMCG.

Perinatal data

We prospectively collected gestational data (parental ethnicity, maternal age, body mass index [BMI], smoking habits, antidepressants, pregnancy complications, antenatal steroids, and magnesium sulfate), fetoplacental data (Doppler measurements and placental histology as examined by a perinatal pathologist according to criteria applicable at the time of examination^{15–26}), and neonatal data (gestational age, delivery mode, sex, birth weight, head circumference, Apgar score, admission to the neonatal intensive care unit, postnatal complications, the need for steroids, and cerebral oxygen saturation as measured with near-infrared spectroscopy during first three days after birth).

Fetal brain-sparing

Antenatal Doppler sonography was performed to assess the pulsatility index (PI) of the umbilical and middle cerebral artery. The cerebroplacental ratio (CPR) was calculated by dividing the latter by the first.²⁷ A CPR below 1 was considered as evidence for fetal brain-sparing. The measurements were performed at least once a

week upon diagnosis of FGR. The last measurement before birth was included for analysis.

Methylation of neurodevelopmental genes at 4 years of age

Gene selection

Based on our hypothesis, we chose to analyze oxygen-dependent regulatory genomic regions encoding HIF1 α and well-known neurotrophic factors, including EPO, VEGFA, and BDNF, which play important roles during brain development and contain binding sites for HIF1 α .^{28–30} The selected DNA sequences with its CpG positions and relevant transcription factor binding sites are presented in Fig. 1. We selected the promoter region of *HIF1A*, which encodes HIF1 α . This region contains a hypoxia-response element (HRE), to which HIF1 α is able to bind and increase expression of HIF1 α under hypoxia in an autoregulatory fashion.³¹ Next to the HRE lies a binding site for the transcription factor Kaiso (also known as ZBTB33), which has been suggested to repress HIF1 α expression.³² We further selected the promoter and the distal enhancer region of *EPO*, which contain binding sites for HIF1 β and HIF1 α , respectively, and together are responsible for the expression of EPO under hypoxic conditions.^{33,34} In addition, the promoter region of *EPOR* was analyzed. *EPOR* does not contain any known HREs, but the selected region has been implicated in developmental downregulation of EPOR expression in the brain.³⁵ We further selected an HRE locus in the promoter region of the gene encoding VEGFA.^{36,37} Likewise, an HRE locus within the promoter region of *NTRK2* was selected, which encodes Tropomyosin receptor kinase B (TrkB), a receptor for BDNF and other neurotrophins.¹¹ In addition, we selected a region within the promoter of *BDNF* exon 4. This region contains two binding sites for the transcription factor cyclic adenosine monophosphate response element binding protein (CREB), which has shown to mediate neuroprotective effects of EPO in cerebral ischemia.^{38–41}

DNA sampling and isolation

Buccal cells were collected during follow-up using Isohelix Buccal Swabs (Cell Projects Ltd, Kent, UK). All samples were stored at 4°C until DNA isolation. Once all samples were collected, isolation of DNA was performed according to the protocol of the BuccalFix Plus DNA Isolation Kit (Cell projects Ltd, Kent, UK). Quality and concentration of the isolated DNA were checked by gel electrophoresis and the NanoDrop 2000c spectrophotometer (Thermo Fisher Scientific, Waltham, MA). Isolated DNA was subsequently stored at –80°C until bisulfite treatment.

Analysis of DNA methylation by pyrosequencing

Before pyrosequencing, 250 ng of isolated DNA was treated with bisulfite to convert unmethylated cytosine residues into uracil leaving only methylated cytosine residues. This was done using the EZ DNA Methylation-Gold Kit (Zymo Research, Irvine, CA) according to the manufacturer's protocol with a discard of the flow-through and an extra round of 30 s centrifuging at full speed after step 8. For polymerase chain reaction (PCR), we prepared a mastermix containing 12.5 μ l HotStarTaq DNA Polymerase, 10.5 μ l of sterile water, and a 1 μ l of forward and reverse primer mix (each 10 μ M) per 1 μ l bisulfite template. A negative control without template was included to check for contamination. We used the T100 Thermal Cycler (Bio-Rad, Hercules, CA) and the following conditions for PCR amplification: 95°C for 15 min, 45 cycles of 94°C for 30 s, assay specific temperature for 30 s,

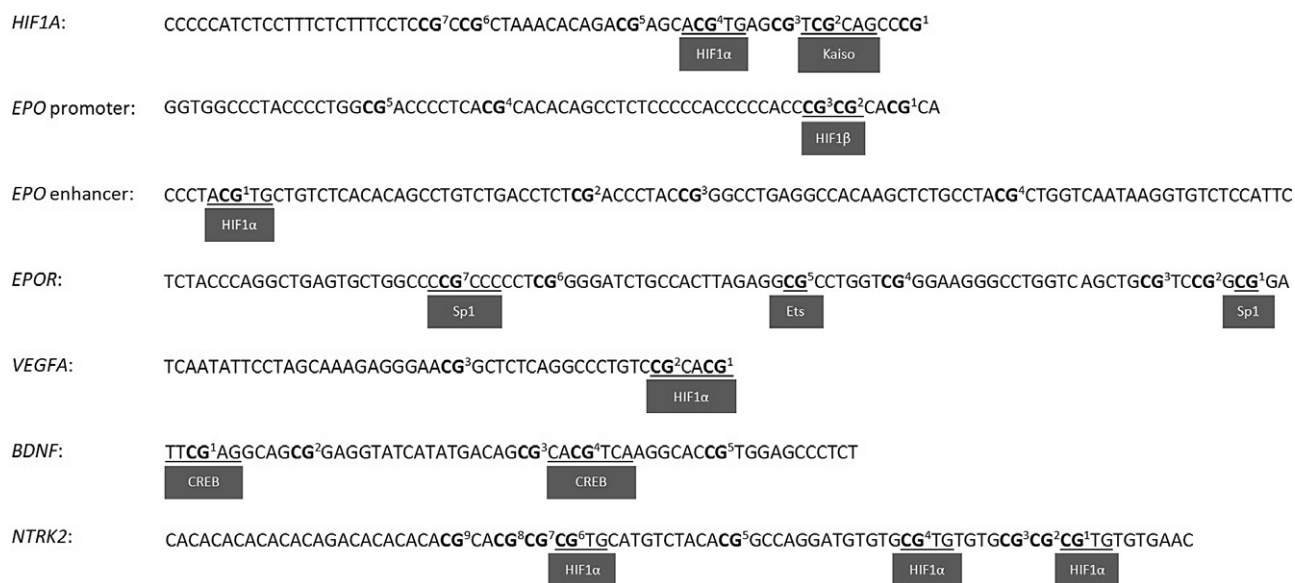


Fig. 1. DNA sequence to analyze before bisulfite treatment depicting CpG positions (bolded and numbered in the direction of sequencing) and important transcription factor binding sites (underlined with transcription factors in grey boxes). *BDNF*, brain-derived neurotrophic factor; CpG, 5'-cytosine-phosphate-guanine-3' dinucleotide; CREB, cyclic adenosine monophosphate response element binding protein; *EPO*, erythropoietin; *EPOR*, erythropoietin receptor; Ets, E26 transformation-specific; *HIF1A/HIF1α*, hypoxia-inducible factor-1 alpha; *HIF1β*, hypoxia-inducible factor-1 beta; *NTRK2*, neurotrophic tyrosine kinase, receptor, type 2; Sp1, specificity protein 1; *VEGFA*, vascular endothelial growth factor A.

72°C for 30 s, followed by a final step of 72°C for 7 min. Assay specific temperatures: 56°C for *HIF1A* promoter and *NTRK2*; 58°C for *EPO* promoter, *EPOR*, and *VEGFA*; 62°C for *EPO* enhancer and *BDNF* exon 4. The modified DNA was then stored at -20°C.

PCR and pyrosequencing primers (Table 1) were designed using the PyroMark Assay Design software (Qiagen, Hilden, Germany). Primers used for *HIF1A*, *VEGFA*, and *EPO* were previously designed by Bekkering *et al.*⁴² After PCR amplification of the DNA region of interest, pyrosequencing was performed using PyroMark Q24 and PyroMark Q48 Autoprep (Qiagen). Methylation levels of each CpG position (given as percentages) and the bisulfite conversion rate were analyzed using the PyroMark Q24 Software and PyroMark Q48 Advanced Software (Qiagen). The maximum allowed percentage of non-conversion was set at 5% in the analysis software. The actual non-conversion rate in the samples was between 0.5% and 3%, wherefore we considered the bisulfite conversion successful. Samples which failed quality control were replicated up to three times until quality control was passed. The sample was excluded if low quality persisted (Supplementary Table S1).

Neurodevelopmental outcome

Neurodevelopmental outcome was assessed at the age of 4 years based on cognition, behavior, and executive functions. Cognition was tested using the WPPSI for children aged 4–7 years (3rd edition, Dutch version), yielding a normed Full Scale, Verbal, and Performance IQ. Total, internalizing, and externalizing behavior (normed *T*-scores) were assessed using the Child Behavior Checklist for ages 1.5–5 years. Executive functions were examined using the Behavior Rating Inventory of Executive Function–Preschool Version for children aged 2–5 years. More specifically, inhibitory self-control, (emotional) flexibility, and emergent metacognition (problem solving using working memory

and planning) were tested, which produced normed *T*-scores for the respective indices and total executive functioning.

Statistical analysis

The statistical software package SPSS 23.0 (IBM Corporation, Armonk, NY, USA) was used for analyses. A (two-sided) *p*-value below 0.05 was considered significant and a (two-sided) *p*-value between 0.05 and 0.1 a possible trend or tendency toward a significant association. As this was an explorative, hypothesis-forming study, we refrained from correcting our *p*-values to reduce the chance of type 2 error.

First, each variable was tested for normal distribution using the Shapiro–Wilk test. Second, differences in population characteristics and the percentage of methylation per CpG location between FGR children with and without evidence of fetal brain-sparing were tested using a Student's *t*-test or Mann–Whitney U test, depending on normality of the data. To test for differences between two binary variables a chi-square test was used. In addition, to better understand how methylation of individual CpG sites within one region relate to each other and potential transcription factor binding sites, we performed correlation analyses between CpGs using Pearson or Spearman's rank correlation analysis, depending on normality of the data. Third, cohort characteristics, which are likely to have a separate effect on DNA methylation, such as perinatal steroid use, gestational age, sex, child age and BMI (*z*-score) at DNA sampling, maternal smoking, medication, age, BMI, and parental ethnicity, were regarded as potential confounders.^{43–45} If in our population, these variables were significantly associated with both brain-sparing and CpG methylation, their confounding effect was adjusted for in a multivariate linear regression model. Finally, to secondarily assess whether differential CpG methylation was associated with neurodevelopmental outcome, correlation analyses were applied using Pearson or Spearman's rank correlation.

Table 1. PCR and sequencing primers, sequence to analyze after bisulfite treatment, and the respective genomic region per gene as based on the *Homo Sapiens* GRCh38.p 13 primary assembly

Gene	Primers	Sequence to analyze	Genomic region
<i>HIF1A</i>	PCR Forward: 5'-AGGAGGTTAGTTGAGGTA TAGTTGG-3' PCR Reverse: 5'-Biotin-CACCCCATCTCCT TTCT-3' Sequencing: 5'-GTTGAGGTATAGTTGGGA-3'	YGGGTTGYGAYGTTTAYGTGTTTGTGTTTAYG GYGGAGAAAGAGAAAGGAGATGGGG	Chromosome 14, 61695200-61695263
<i>EPO</i> promoter	PCR Forward: 5'-GGGGTAGGGGTGTT ATTTGTATG-3' PCR Reverse: 5'-Biotin-CCCAAACCTCCTACC CCTACTCTAAC-3' Sequencing: 5'-GGGTTGTTATTTGTATGTG-3'	TGYGTGYGGGGTGGGGTGGGGAGAGGTTGTG TGYGTGAGGGGTGTTAGGGGTAGGGGTTATT	Chromosome 7,100720640- 100720705
<i>EPO</i> enhancer	PCR Forward: 5'-GGGAAAAGAGGGGTGGAGG-3' PCR Reverse: 5'-Biotin-CTCCCTCCTTAAT AACAACTCAAC-3' Sequencing: 5'-GTGGAGGGGGTTGGG-3'	TTTTAYGTGTTTATATAGTTTGTGTTGATTTTTY ATTTTATYGGGTTGAGGTTATAAGTTTTGTTAYGTTGG TTAATAAGGTGTTTTTATT	Chromosome 7, 100723816-100723913
<i>EPOR</i>	PCR Forward: 5'-GGAGTAGATTTGGGGTTAG AGGG-3' PCR Reverse: 5'-Biotin-AAAAACCCCTACC TCCT-3' Sequencing: 5'-GTTGGGTTAGTAGTTGTT-3'	TTYGTGYGAYGTAGTTGATTAGGTTTTTGYATTAGGY GTTTTAAGTGTTAGATTTTTYAGGGGGYGGGGTTAGT ATTTAGTTGGGTAGA	Chromosome 19, 11384387-11384295
<i>VEGFA</i>	PCR Forward: 5'-GGGAGTAGGAAAGTGAGGT-3' PCR Reverse: 5' Biotin-TTCCCTACCCCTT CAATAT-3' Sequencing: 5'-AGTAGGAAAGTGAGGTTA-3'	YGTGYGGATAGGGTTTGGAGTYGTTTTT TTTTGTAGGAATATGA	Chromosome 6, 43769854-43769901
<i>BDNF</i>	PCR Forward: 5'-GGGGTTGGAAGTAAAA TATTTGTA-3' PCR Reverse: 5' Biotin-CCCCATCAACAAAA ACTCCATTAATCTC-3' Sequencing: 5'-GTAATTAGTGTATTAGAGTGT TTAT-3'	TTYGAGGTAGYGGAGGTATTATATGATAGY TAYGTTAAGGTATYGTGGAGTTTTT	Chromosome 11, 27701701-27701645
<i>NTRK2</i>	PCR Forward: 5'-Biotin-GTTTATTTAGAGGT ATTTGGATGTAAATG-3' PCR Reverse: 5'-TAACCAAAAAACAACAAC AACATA-3' Sequencing: 5'-ACAACAACAACAACATAT AAAA-3'	ATTCACACACRCRCACACACACATCCTAACRT ATAAACATACACRCRCRTACRTATATATCTATATATATATA	Chromosome 9, 84669441-84669524

BDNF, brain-derived neurotrophic factor; *EPO*, erythropoietin; *EPOR*, erythropoietin receptor; *HIF1A*, hypoxia-inducible factor-1 alpha; *NTRK2*, neurotrophic tyrosine kinase, receptor, type 2; PCR, polymerase chain reaction; *VEGFA*, vascular endothelial growth factor A.

Results

From an FGR cohort originally including 51 fetuses, three infants died in the neonatal period, six had incomplete perinatal hemodynamic measurements, another two declined follow-up at perinatal inclusion, and one withdrew consent for the whole study. At the age of 4 years, three infants were lost to follow-up and the parents of 12 infants withdrew from follow-up after initial agreement. In three children, buccal DNA sampling was denied or not feasible due to severe developmental problems. This resulted in the inclusion of 21 children with complete information on the CPR, DNA methylation levels, and neurodevelopmental test results. Children lost to or declining follow-up and DNA sampling ($n = 20$) less often required intensive care, but did not significantly differ in gestational age, birth weight

(z -score), head circumference (z -score), PI of the umbilical artery (not more often above the 95th percentile), presence or absence of fetal brain-sparing, postnatal cerebral oxygenation, or gestational and neonatal complications from the included study population (Supplementary Table S2).

Characteristics of the study population

Eight children had evidence of fetal brain-sparing, while 13 children did not. Impaired flow in the umbilical artery (i.e., decreased, absent, or reversed end-diastolic flow, implying greater placental resistance) at last prenatal ultrasound was significantly more often observed in children showing fetal brain-sparing (Table 2). Moreover, in the same children, placental weight tended to be more often below the 10th percentile. Placental signs of infection

Table 2. Cohort characteristics

	Fetal brain-sparing, <i>n</i> = 8	No fetal brain-sparing, <i>n</i> = 13	<i>p</i> -value
Gestational characteristics			
Caucasian mother	8 (100)	11 (85)	0.243
Caucasian father	7 (88)	10 (83)	0.798
Maternal age (years)	29.3 [26.5; 35.2]	32.4 [24.9; 40.3]	0.089 [#]
Maternal BMI	24.4 [19.1; 34.7]	25.3 [17.7; 36.0]	0.718
Maternal smoking	3 (38)	3 (23)	0.477
Diabetes mellitus	1 (13)	2 (15)	0.854
Preeclampsia	2 (25)	2 (15)	0.586
HELLP	1 (13)	1 (8)	0.243
PPROM	1 (13)	1 (8)	0.716
Maternal antidepressants	0 (0)	1 (8)	0.421
Antenatal steroids	6 (75)	5 (39)	0.104
Antenatal MgSO ₄	1 (13)	5 (39)	0.201
Cesarean section	8 (100)	7 (54)	0.023*
Placental histology			
Maternal vascular underperfusion	5 (71), <i>n</i> = 7	5 (46), <i>n</i> = 11	0.280
Fetal thrombotic vasculopathy	4 (57), <i>n</i> = 7	3 (27), <i>n</i> = 11	0.205
Ascending intrauterine infection	1 (14), <i>n</i> = 7	2 (18), <i>n</i> = 11	0.829
Chronic deciduitis	2 (29), <i>n</i> = 7	4 (36), <i>n</i> = 11	0.732
Villitis of unknown etiology	0 (0), <i>n</i> = 7	1 (9), <i>n</i> = 11	0.412
Increase in nucleated RBCs	2 (29), <i>n</i> = 7	2 (18), <i>n</i> = 11	0.605
Placental weight < 10th percentile	7 (100), <i>n</i> = 7	7 (64), <i>n</i> = 11	0.070 [#]
Doppler characteristics at last prenatal ultrasound			
Abnormal flow UA (PI > 95th percentile or absent/reversed flow)	8 (100)	6 (46)	0.011*
Abnormal flow MCA (PI < 5th percentile)	5 (63)	1 (8)	0.007*
Abnormal flow DV (PI > 95th percentile/absent or reversed a-wave)	5 (63)	5 (46)	0.463
Neonatal characteristics			
Female	5 (63)	7 (54)	0.697
Gestational age (weeks)†	31.4 [29.1; 37.3]	34.4 [28.0; 39.9]	0.045*
Birth weight (z-score)	-3.28 [-4.66; -1.79]	-2.68 [-5.87; -0.29]	0.278
Head circumference (z-score)	-1.78 [-3.62; -0.43]	-2.09 [-2.69; -1.34]	0.451
Apgar score at 5 min†	6 [4; 10]	9 [5; 10]	0.104
Admission to NICU	7 (88)	8 (62)	0.201
Mechanical ventilation	3 (38)	6 (46)	0.697
Bronchopulmonary dysplasia	1 (13)	1 (8)	0.716
Hemodynamically significant PDA	0 (0)	2 (15)	0.243
Necrotizing enterocolitis	0 (0)	0 (0)	-
Sepsis	1 (13)	0 (0)	0.191
IVH/PVL	0 (0)	0 (0)	-
Postnatal steroids	0 (0)	1 (8)	0.421
r _c SO ₂ on postnatal day 1†	85 [73; 90], <i>n</i> = 8	81 [56; 92], <i>n</i> = 13	0.500
r _c SO ₂ on postnatal day 2	87 [80; 94], <i>n</i> = 8	79 [64; 92], <i>n</i> = 10	0.035*
r _c SO ₂ on postnatal day 3	85 [74; 93], <i>n</i> = 8	77 [65; 91], <i>n</i> = 9	0.083 [#]

(Continued)

Table 2. (Continued)

	Fetal brain-sparing, <i>n</i> = 8	No fetal brain-sparing, <i>n</i> = 13	<i>p</i> -value
Childhood characteristics at follow-up			
Age (years)†	4.3 [4.3; 4.5]	4.2 [4.0; 4.7]	0.140
BMI (z-score)†	-0.86 [-1.38; 0.29]	-0.33 [-1.93; 3.26]	0.571
Head circumference (z-score)	-0.59 [-1.81; 2.09]	-0.42 [-1.78; 1.08]	0.727
Reported ASD	0 (0)	2 (15)	0.243
Cognitive outcome (IQ)			
Full Scale	92 [63; 115], <i>n</i> = 8	94 [79; 107], <i>n</i> = 8	0.851
Verbal	99 [71; 120], <i>n</i> = 8	93 [76; 108], <i>n</i> = 8	0.405
Performance	91 [72; 107], <i>n</i> = 8	99 [85; 110], <i>n</i> = 9	0.143
VIQ > PIQ discrepancy	5 (63), <i>n</i> = 8	2 (25), <i>n</i> = 8	0.131
Behavioral outcome (<i>T</i> -score)			
Total behavior	49 [34; 58], <i>n</i> = 8	58 [45; 72], <i>n</i> = 12	0.011*
Internalizing behavior	54 [43; 63], <i>n</i> = 8	57 [45; 73], <i>n</i> = 12	0.299
Externalizing behavior	46 [28; 59], <i>n</i> = 8	56 [44; 66], <i>n</i> = 12	0.022*
Executive function (<i>T</i> -score)			
Total executive function	51 [41; 65], <i>n</i> = 8	61 [46; 76], <i>n</i> = 12	0.047*
Inhibitory Self-Control Index	50 [41; 66], <i>n</i> = 8	60 [42; 76], <i>n</i> = 13	0.042*
Flexibility Index†	58 [41; 60], <i>n</i> = 8	62 [39; 95], <i>n</i> = 13	0.336
Metacognition Index	51 [42; 65], <i>n</i> = 8	59 [46; 71], <i>n</i> = 12	0.064 [#]

For good comparability, data are presented as mean/median [total range] or absolute number (percentage). †indicates, where a Mann-Whitney U test was used and a median is presented due to non-normality of the data. *T*-scores are to be interpreted as the lower, the better. [#] and * present differences between both groups at $p < 0.1$ and $p < 0.05$, respectively. ASD, autism spectrum disorder; BMI, body mass index; DV, ductus venosus; HELLP, syndrome of hemolysis, elevated liver enzymes, and low platelets; IQ, intelligence quotient; IVH, intraventricular hemorrhage; MCA, middle cerebral artery; MgSO₄, magnesium sulfate; NICU, neonatal intensive care unit; PDA, patent ductus arteriosus; PI, pulsatility index; PIQ, performance intelligence quotient; PPRM, prolonged premature rupture of membranes (>12 h); PVL, periventricular leukomalacia; RBC, red blood cell; r_cSO₂, regional cerebral oxygen saturation (measured with near-infrared spectroscopy); UA, umbilical artery; VIQ, verbal intelligence quotient.

were infrequent and equally distributed among groups. Maternal and gestational characteristics were largely similar, but delivery mode and duration of pregnancy were significantly different. Children with fetal brain-sparing had to be delivered by cesarean section more frequently and were born at a younger gestational age. Before birth, infants delivered by Cesarean section more often experienced heart rate decelerations as detected by cardiotocography (8 vs. 0, $p = 0.015$) and impaired flow in the umbilical artery (12 vs. 2, $p = 0.020$). Spontaneous or induced labor before Cesarean delivery was only seen in four cases, of which one was preterm and another one belonged to the group with fetal brain-sparing. There were no differences in sex. After birth, infants with fetal brain-sparing had significantly higher regional cerebral oxygen saturations on day 2, which was also highly related to an abnormally increased blood flow in the middle cerebral artery ($p = 0.037$), but not impaired umbilical flow, gestational age, Cesarean delivery, or birth weight z-score ($p > 0.1$). At 4 years of age, infants with fetal brain-sparing had better total behavior and executive functions, in particular, better externalizing behavior and inhibitory self-control (i.e., lower *T*-scores). Two infants (15%) without fetal brain-sparing were reported to be diagnosed with or highly suspected of autism spectrum disorder. Neither of the two groups showed intraventricular hemorrhage nor periventricular ischemic lesions as assessed by postnatal cranial ultrasound.

Differences in methylation between FGR children with and without brain-sparing

The methylation patterns for FGR children with and without brain-sparing are presented per analyzed genomic region in Fig. 2. In children with fetal brain-sparing, there was a trend toward a significantly higher percentage of methylation at CpG site 4 of the analyzed *HIF1A* locus than in FGR children without fetal brain-sparing (mean [μ] \pm standard deviation [SD]) of 1.16 ± 0.28 vs. 0.086 ± 0.40 , $p = 0.093$), while all other CpGs were not methylated differently (Table 3). CpG 4 lies exactly within a binding site for HIF1 α , as demonstrated in Table 1.

The analyzed *EPO* promoter and enhancer region were not methylated differently, but CpG position 4 of the *EPOR* gene tended to be significantly less methylated in FGR children with fetal brain-sparing than without ($\mu \pm SD = 3.20 \pm 1.53$ vs. 4.97 ± 1.85 , $p = 0.052$; Table 3). Methylation levels of CpGs 1–3 of *EPOR* were not different between the groups, and unfortunately, CpG 5–7 had to be excluded from analyses due to insufficient quality of measurements.

In children with fetal brain-sparing, we found a trend toward significantly higher methylation levels at CpG 1 of the selected *VEGFA* locus ($\mu \pm SD = 1.31 \pm 0.43$ vs. 0.93 ± 0.40 , $p = 0.054$; Table 3), which lies within an HRE (Fig. 2). Furthermore, FGR children with fetal brain-sparing had significantly higher

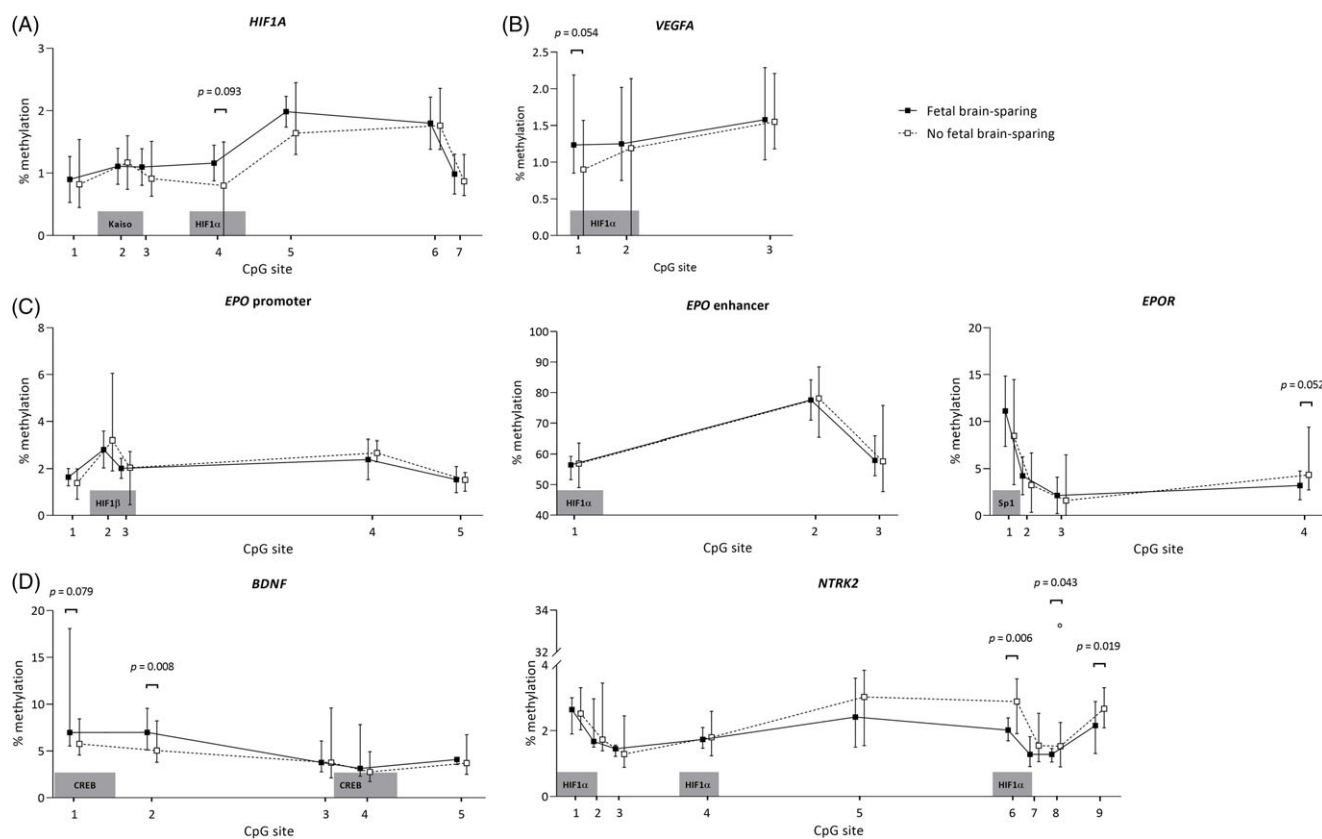


Fig. 2. Buccal DNA methylation patterns at selected regions of (A) *HIF1A*, (B) *VEGFA*, (C) *EPO* and *EPOR*, and (D) *BDNF* and its receptor gene *NTRK2* in 4-year-old children born following fetal growth restriction with ($n=8$) and without fetal brain-sparing ($n=13$). Data are presented as medians (total range); the circle represents an extreme outlier (33.21%). The p -values given were gained using Student's t -test or Mann-Whitney U test, involved numbers (n) per analyzed CpG, and group are given in Table 3. † $p > 0.1$ when adjusted for gestational age in multivariate regression analysis. *BDNF*, brain-derived neurotrophic factor; CpG, 5'-cytosine-phosphate-guanine-3' dinucleotide; CREB, cyclic adenosine monophosphate response element binding protein; *EPO*, erythropoietin; *EPOR*, erythropoietin receptor; *HIF1A/α*, hypoxia-inducible factor-1 alpha; *HIF1β*, hypoxia-inducible factor-1 beta; *NTRK2*, neurotrophic tyrosine kinase, receptor, type 2; *Sp1*, specificity protein 1; *VEGFA*, vascular endothelial growth factor A.

methylation levels at CpG 2 ($\mu \pm SD = 6.91 \pm 1.41$ vs. 5.24 ± 1.17 , $p = 0.008$) and a tendency toward higher methylation of CpG 1 ($\mu \pm SD = 8.25 \pm 4.04$ vs. 6.06 ± 1.15 , $p = 0.079$) of the selected *BDNF* locus (Table 3). Only the latter lies within a transcriptionally important CREB binding site (Figs. 1 and 2), but methylation levels of both CpG sites highly correlated (Supplementary Table S3). The *BDNF* receptor gene *NTRK2* demonstrated significantly lower methylation levels at CpG 6 (which lies within an HRE, $\mu \pm SD = 2.08 \pm 0.30$ vs. 2.80 ± 0.54 , $p = 0.006$), CpG 8 (median [interquartile range] = 1.28 [1.09; 1.44] vs. 1.63 [1.37; 2.13], $p = 0.043$), and CpG 9 ($\mu \pm SD = 2.10 \pm 0.51$ vs. 2.64 ± 0.34 , $p = 0.019$) in FGR children with fetal brain-sparing (Table 3; Fig. 2). Correlation coefficients between these CpG sites were high (Supplementary Table S2). CpG 8 of *NTRK2* contained an extreme outlier within the group of FGR children without fetal brain-sparing (33.21% methylation). Excluding the outlier did not relevantly change our results ($\mu \pm SD = 1.28 \pm 1.58$ vs. 1.62 ± 0.43 , $t(11) = 2.23$, $p = 0.049$). As quality of methylation analysis was high, the presented data in Table 3 and Fig. 2 include this outlier.

Multiple regression analysis

Gestational age, which was significantly lower in children with fetal brain-sparing, also significantly correlated with methylation of

EPOR CpG 4 (Pearson's correlation coefficient = 0.479, $p = 0.038$). As gestational age is known to have a separate effect on DNA methylation, we included it as a confounder in a linear regression model. After correcting for gestational age, the association between brain-sparing and methylation of *EPOR* at CpG 4 lost its significance (B [95% confidence interval] = -1.31 [-3.19 ; 0.58], $t = -1.48$, $p = 0.160$).

Association between differentially methylated CpGs and neurodevelopmental outcome

To secondarily assess whether differential CpG methylation among children with and without brain-sparing may explain differences in neurodevelopmental outcome between the two groups, correlation coefficients between CpG methylation levels and outcome were calculated (Table 4). Increased methylation of CpG 4 of *HIF1A* was associated with a trend toward higher Full Scale IQ (Pearson coefficient = 0.449, $p = 0.081$) and higher Verbal IQ (Pearson coefficient = 0.429, $p = 0.098$). Moreover, methylation of CpG 1 of the *VEGFA* locus significantly and inversely correlated with Performance IQ (Pearson coefficient = -0.660 , $p = 0.003$), while hypermethylation of CpG 2 (and to a minor extent CpG 1) of *BDNF* correlated with better executive function (i.e., lower T -scores), in particular, better inhibitory self-control (Pearson coefficient = -0.493 , $p = 0.020$).

Table 3. Differences in percentage methylation per CpG of selected gene locations at 4 years of age between children born following fetal growth restriction with or without brain-sparing, as assessed with Student's *t*-test or Mann-Whitney U test

Gene	CpG	Mean/median (n)		<i>t</i> (df)/U	<i>p</i> -value
		Brain-sparing	No brain-sparing		
<i>HIF1A</i>	1	0.90 (8)	0.84 (13)	−0.40 (19)	0.697
	2	1.11 (8)	1.16 (13)	0.37 (18)	0.717
	3	1.10 (7)	0.97 (13)	−1.01 (18)	0.328
	4	1.16 (7)	0.86 (13)	−1.77 (18)	0.093 [#]
	5	1.98 (7)	1.77 (13)	−1.39 (18)	0.181
	6	1.80 (7)	1.80 (13)	−0.03 (18)	0.980
	7	0.99 (7)	0.94 (13)	−0.36 (18)	0.727
<i>EPO promoter</i>	1	1.64 (8)	1.33 (13)	−1.71 (19)	0.104
	2†	2.63 (8)	3.20 (13)	35.0	0.238
	3	2.01 (8)	1.83 (13)	−0.69 (19)	0.497
	4†	2.70 (8)	2.67 (13)	47.0	0.750
	5	1.53 (8)	1.49 (13)	−0.22 (8.6)	0.859
<i>EPO enhancer</i>	1	56.00 (8)	57.22 (13)	0.77 (19)	0.449
	2	77.85 (8)	77.75 (13)	−0.03 (19)	0.973
	3†	58.55 (8)	58.91 (13)	0.13 (19)	0.903
<i>EPOR</i>	1	11.11 (8)	9.02 (13)	−1.42 (19)	0.171
	2	4.32 (8)	3.57 (13)	−0.77 (19)	0.449
	3†	1.51 (8)	1.60 (13)	49.5	0.860
	4	3.20 (7)	4.97 (11)	2.10 (16)	0.052 [#]
<i>VEGFA</i>	1	1.31 (8)	0.93 (13)	2.06 (19)	0.054 [#]
	2	1.29 (8)	1.25 (13)	0.19 (19)	0.854
	3	1.64 (8)	1.70 (13)	0.31 (19)	0.761
<i>BDNF</i>	1	8.25 (8)	6.06 (13)	−1.49 (19)	0.079 [#]
	2	6.91 (8)	5.24 (13)	−2.94 (19)	0.008 [*]
	3	3.82 (7)	4.29 (13)	0.56 (18)	0.583
	4†	3.13 (7)	2.77 (13)	32.0	0.311
	5	3.96 (7)	3.99 (13)	0.10 (15)	0.925
<i>NTRK2</i>	1	2.59 (8)	2.53 (13)	0.41 (19)	0.689
	2†	1.68 (8)	1.73 (13)	42.5	0.500
	3†	1.45 (8)	1.29 (13)	51.0	0.972
	4†	1.74 (8)	1.80 (12)	39.5	0.521
	5	2.50 (8)	2.86 (11)	1.22 (17)	0.240
	6	2.08 (7)	2.80 (10)	3.54 (15)	0.006 [*]
	7	1.35 (7)	1.59 (10)	1.24 (15)	0.235
	8†	1.28 (7)	1.63 (10)	14.0	0.043 [*]
	9	2.10 (7)	2.64 (10)	2.64 (15)	0.019 [*]

†Mann-Whitney U test was used due to non-normality of the data, for which median and U are given. [#] and ^{*} mark differences between means or medians at (an uncorrected) $p < 0.1$ and $p < 0.05$, respectively. *BDNF*, brain-derived neurotrophic factor; CpG, 5'-cytosine-phosphate-guanine-3' dinucleotide; df, degrees of freedom; *EPO*, erythropoietin; *EPOR*, erythropoietin receptor; *HIF1A*, hypoxia-inducible factor-1 alpha; *NTRK2*, neurotrophic receptor tyrosine kinase 2; *VEGFA*, vascular endothelial growth factor A.

Discussion

In this prospective follow-up study we compared the buccal DNA methylation status of predefined genomic regions of *HIF1A* and some of its neurotrophic target genes between FGR

children with and without fetal brain-sparing, relating it to their neurodevelopmental outcome. The regions analyzed were selected based on their proven or theoretical potential to alter activity of the respective gene in an oxygen-dependent manner. We found that

Table 4. Correlation coefficients between the percentage methylation of brain-sparing associated CpGs and neurodevelopmental outcome at 4 years following fetal growth restriction

	Gene, CpG site						
	<i>HIF1A</i> , 4	<i>VEGFA</i> , 1	<i>BDNF</i> , 1	<i>BDNF</i> , 2	<i>NTRK2</i> , 6	<i>NTRK2</i> , 8†	<i>NTRK2</i> , 9
Cognition (IQ)							
Full Scale	0.449 (0.081) [#]	-0.325 (0.203)	0.045 (0.865)	0.103 (0.694)	0.175 (0.568)	0.326 (0.278)	0.029 (0.926)
Verbal	0.429 (0.098) [#]	0.055 (0.834)	0.248 (0.336)	0.284 (0.270)	0.080 (0.794)	0.269 (0.374)	0.105 (0.732)
Performance	0.412 (0.100)	-0.660 (0.003) [*]	-0.119 (0.637)	0.021 (0.934)	0.287 (0.320)	0.454 (0.103)	-0.001 (0.997)
Behavior (T-score)							
Total	-0.016 (0.945)	-0.130 (0.575)	-0.205 (0.372)	-0.229 (0.318)	0.238 (0.358)	0.189 (0.468)	0.270 (0.294)
Internalizing	0.093 (0.696)	-0.172 (0.456)	-0.091 (0.696)	0.036 (0.877)	-0.068 (0.796)	0.052 (0.844)	0.048 (0.856)
Externalizing	-0.098 (0.680)	0.029 (0.901)	-0.133 (0.567)	-0.271 (0.235)	0.328 (0.199)	0.157 (0.547)	0.237 (0.361)
EF (T-score)							
Total	-0.133 (0.578)	0.088 (0.706)	-0.287 (0.207)	-0.408 (0.066) [#]	0.015 (0.954)	0.088 (0.736)	0.102 (0.696)
ISCI	-0.245 (0.284)	0.152 (0.500)	-0.344 (0.117)	-0.493 (0.020) [*]	0.102 (0.687)	0.226 (0.366)	0.105 (0.679)
FI†	0.036 (0.875)	0.152 (0.498)	0.014 (0.952)	-0.260 (0.243)	-0.116 (0.646)	0.146 (0.564)	-0.035 (0.890)
EMI	-0.094 (0.694)	0.008 (0.972)	-0.385 (0.085) [#]	-0.398 (0.074) [#]	0.134 (0.608)	0.393 (0.118)	0.205 (0.430)

†Spearman's rank correlation analysis was used due to non-normality of the data. *P*-values are given in brackets and were not corrected for multiple testing. [#] and ^{*} mark associations at *p* < 0.1 and *p* < 0.05, respectively. *T*-scores are to be interpreted as the lower, the better. *BDNF*, brain-derived neurotrophic factor; CpG, 5'-cytosine-phosphate-guanine-3' dinucleotide; EF, executive function; EMI, Emergent Metacognition Index; *EPOR*, erythropoietin receptor; FI, Flexibility Index; *HIF1A*, hypoxia-inducible factor-1 alpha; ISCI, Inhibitory Self-Control Index; IQ, Intelligence Quotient; *NTRK2*, neurotrophic tyrosine kinase, receptor, type 2; *VEGFA*, vascular endothelial growth factor A.

FGR children with prenatal evidence of brain-sparing showed a trend toward hypermethylation at the HRE of *HIF1A* and *VEGFA*. Moreover, we found hypermethylation at a CREB binding site within the promoter region of *BDNF* exon 4 and hypomethylation at an HRE located within the promoter region of its receptor *NTRK2*.

HIF1α regulates the cellular response to hypoxic conditions and among its targets are many neurotrophic factors. In a previous study, we found that preferential perfusion of the fetal brain in FGR was associated with better neurodevelopmental outcome at 4 years of age than FGR without fetal brain-sparing.⁵ This seemed to be mediated through higher cerebral oxygen saturations as suggested by postnatal tissue oxygenation monitoring with near-infrared spectroscopy. In the same cohort, we now find a trend toward hypermethylation at the autoregulatory HRE of the *HIF1A* promoter in the buccal DNA of these children. Since buccal methylation patterns closely correlate with those of neuronal DNA due to their common ectodermal origin, this may reflect suppression of the hypoxic response in brain tissue.¹⁴ This could be caused by higher cerebral oxygen saturations, which are evident in FGR neonates with fetal brain-sparing.⁴ Moreover, hypermethylation

of this locus was associated with a trend toward better Full Scale and Verbal IQ. This may be mediated through its effects on neurotrophic factors, since postnatal cranial ultrasounds did not demonstrate a difference in ischemic lesions between FGR infants with and without fetal brain-sparing. However, it needs to be emphasized at this point that these trend may also merely represent accidental findings and require confirmation.

Several genetic factors play critical neurotrophic roles during early human brain development, whose expression has shown to be affected by hypoxia. *VEGFA* expression is increased by *HIF1α* with important pro-angiogenic effects under hypoxic conditions but also stimulatory effects on axonal outgrowth and the proliferation, migration, and survival of neurons and neuroglia.⁴⁶ Oosthuysen *et al.* demonstrated impaired hypoxic upregulation of neural VEGF and motor neuron degeneration in knockout mice lacking another HRE in the *VEGFA* promoter.⁴⁷ Since hyperoxia downregulates VEGF expression, significantly higher postnatal cerebral oxygenation levels in FGR children with fetal brain-sparing than in those without brain-sparing may possibly explain the observed trend toward hypermethylation at the HRE of *VEGFA* in FGR children with fetal brain-sparing.

Although this tendency toward increased methylation at this locus may also be an accidental finding, if found to be true and related to reduced expression of *VEGFA*, it could explain the inverse correlation between *VEGFA* methylation and Performance IQ at 4 years, since Performance IQ is related to motor function.⁴⁸ That there may be a link between early cerebral oxygenation levels, altered *VEGFA* levels and neurodevelopmental functioning, was already suggested by us in a previous study, which demonstrated a lower Performance IQ in children with comparably high oxygen saturations during the first days after birth.⁵

While above theory is based on a small cohort and studies testing the methylation of the analyzed HRE in relation to expression of *VEGFA* are not yet available, it further remains debatable whether increased buccal *VEGFA* methylation at later age could indeed result from fetal brain-sparing. We recently demonstrated an association between fetal brain-sparing and hypermethylation of the same CpG of *VEGFA* in the placental tissue of this FGR cohort.⁴² However, one may rather expect placental hypomethylation at this locus, since fetal brain-sparing is generally accepted to be a compensatory fetal response to placental hypoxia. Instead, both placental and buccal hypermethylation may therefore reflect an anti-angiogenic state during early fetoplacental development, which leads to placental insufficiency and subsequently fetal brain-sparing.⁴⁹ This angiogenic dysbalance seems to persist in the neonate.⁵⁰ More severe placental insufficiency in children with fetal brain-sparing, which may be suggested by a decreased flow in the umbilical artery (indicating a higher placental resistance) and a tendency towards lower placental weight, could therefore also explain our findings. Although an angiogenic dysbalance (lower circulating VEGF levels) has also been evidenced in patients with ASD, we have reported in our previous paper that both children also presented with impaired umbilical flow even if no signs of fetal brain-sparing were present.^{5,51} Regardless of its origin, *VEGFA* hypermethylation does not seem to benefit neurodevelopmental outcome.

Although the *BDNF* gene does not contain any HREs, transcriptional activation upon hypoxia has shown to occur through interaction of the transcription factor CREB with the promoter region of exon 4, mediated through EPO-enhanced phosphorylation of CREB.³⁹ Accordingly, we found significant hypermethylation close to the designated binding sites of CREB at this promoter in FGR children with fetal brain-sparing. However, we did not observe any significant differences in methylation of the selected HRE regions in *EPO*, suggesting that this may not fully explain altered *BDNF* methylation. Moreover, hypermethylation of *BDNF* seemed associated with better executive functioning, in particular significantly better inhibitory self-control, although we expected hypermethylation to cause poorer executive functioning by reducing expression of *BDNF*. However, the analyzed CREB binding sites have also been implicated in calcium-mediated, activity-dependent upregulation of *BDNF* through N-methyl-D-aspartate (NMDA)-receptor activation by glutamate.^{38,41} *BDNF* is known to sustain NMDA activation through TrkB signaling, creating a positive feedback loop, which promotes neuronal sprouting and synaptogenesis but may also lead to hyperexcitability.⁵² Our findings may therefore reflect a reduction of perinatal hypoxia-induced glutamate excitotoxicity through hypermethylation of *BDNF* in the presence of fetal brain-sparing.⁵³ This may also contribute to increased *BDNF* levels in autism spectrum disorder, which has been related to an excitatory/inhibitory imbalance and, in our cohort, was reported in 15% of FGR children without fetal brain-sparing.^{54,55}

Interestingly, regarding the receptors of neurotrophic factors, brain-sparing was also significantly associated with hypomethylation of the *BDNF* receptor gene *NTRK2* at CpG position 6, which corresponds to HRE1 reported by Martens *et al.*, and the closely located CpG positions 8 and 9.¹¹ Likewise, we found a trend toward significant hypomethylation of *EPOR*, although this seemed to be confounded by lower gestational age, supporting the hypothesis that methylation of this region is unaffected by oxygenation status but involved in developmental downregulation of *EPOR*.³⁵ Hypomethylation of the HRE of *NTRK2*, however, was unexpected and may be related to placental hypoxia rather than brain-sparing, since it also did not correlate with neurodevelopmental outcome. This is in line with studies demonstrating elevated *NTRK2* levels in the placental tissue of FGR pregnancies to stimulate endothelial cell survival and angiogenesis.⁵⁶ Although low *BDNF* levels have also been demonstrated in early preeclampsia and may contribute to an anti-angiogenic placental environment, the analyzed promoter region of exon 4 is highly tissue-specific for the brain, which may also explain paradoxical differences in methylation between *BDNF* and its receptor gene.^{57,58}

To our knowledge, this is the first study analyzing the association between fetal brain-sparing and buccal DNA methylation patterns of neurodevelopmentally important genes in primary school children born following FGR. The strength of our study lies within the specific analyses of binding sites implicated in the hypoxic upregulation of these genes and their relationship to outcome established by prospective neurodevelopmental testing. However, we acknowledge some limitations. First, the sample size of this study was small, which limited our power to detect significant associations. In addition, it limited our study to a candidate gene approach, while an epigenome-wide association study would yield far more information. Second, we performed multiple testing without controlling for it, as we considered this a hypothesis-generating study. However, while this reduced type II error, it may also have increased our false discovery rate. Reported trends ($p < 0.1$), in particular, but also significant findings ($p < 0.05$) are therefore to be interpreted with great caution and require validation by larger studies. Third, due to the small sample size, we may have neglected some important confounders of the association between brain-sparing and methylation as well as the association between methylation and neurodevelopmental outcome, such as sex or ethnicity. Moreover, although we tried to study causation between brain-sparing and altered methylation patterns by analyzing oxygen-dependent regulatory genomic regions, some of our findings may also be explained by underlying anti-angiogenic pathology of placental insufficiency or resulting hypoxia, which could also explain differences in DNA methylation.⁹ Finally, differences in methylation were small, and it remains to be investigated whether they would be similar in neuronal DNA and sufficient to alter gene expression. Therefore, we encourage replication of our findings by larger trials with additional gene expression analysis in both buccal and neuronal DNA.

In conclusion, this explorative study, limited in sample size, showed that fetal brain-sparing in FGR is associated with a trend toward buccal hypermethylation of *HIF1A* and *VEGFA*, significant hypermethylation of *BDNF*, and significant hypomethylation of *NTRK2* at regions implicated in oxygen-mediated regulation of these genes. Moreover, increased methylation of these regions within *BDNF* and *VEGFA* was significantly associated with better inhibitory self-control and poorer Performance IQ, respectively. However, before we can draw conclusions about possible causation and mechanisms behind these findings, they need to be confirmed

by larger hypothesis-testing studies, which also need to involve gene expression analysis.

Supplementary Material. To view supplementary material for this article, please visit <https://doi.org/10.1017/S2040174421000374>

Acknowledgements. We thank A. Britt Foreman and Sahar Salavati for their help and assistance during follow-up and Anne den Heijer for her expertise, advice, and training during neurodevelopmental testing. We thank Mirthe H. Schoots for analyzing the placentas of this FGR cohort, thereby providing valuable information to this work.

Financial Support. This study was part of the research program of the Research Institute of Behavioral and Cognitive Neurosciences (BCN), Graduate School of Medical Sciences, University of Groningen, participation in which is financially supported by the Junior Scientific Master Class of the University Medical Center Groningen, University of Groningen, The Netherlands.

Conflicts of Interest. None.

Ethical standards. The authors assert that all procedures contributing to this work comply with the ethical standards of the relevant national guidelines on human experimentation (Wet medisch-wetenschappelijk onderzoek met mensen) and with the Helsinki Declaration of 1975, as revised in 2008, and has been approved by the institutional committees (Medisch Ethische Toetsingscommissie van het UMCG).

References

1. Miller SL, Huppi PS, Mallard C. The consequences of fetal growth restriction on brain structure and neurodevelopmental outcome. *J Physiol (Lond)*. 2016; 594(4), 807–823.
2. Nardoza LMM, Caetano ACR, Zamarian ACP, *et al*. Fetal growth restriction: current knowledge. *Arch Gynecol Obstet*. 2017; 295(5), 1061–1077.
3. Giussani DA. The fetal brain sparing response to hypoxia: physiological mechanisms. *J Physiol*. 2016; 594(5), 1215–1230.
4. Tanis JC, Boelen MR, Schmitz DM, *et al*. Correlation between doppler flow patterns in growth-restricted fetuses and neonatal circulation. *Ultrasound Obstet Gynecol*. 2016; 48(2), 210–216.
5. Richter AE, Salavati S, Kooi EMW, *et al*. Fetal brain-sparing, postnatal cerebral oxygenation, and neurodevelopment at 4 years of age following fetal growth restriction. *Front Pediatr*. 2020; 8, 225.
6. Verhagen EA, Van Braeckel KN, van der Veere CN, *et al*. Cerebral oxygenation is associated with neurodevelopmental outcome of preterm children at age 2 to 3 years. *Dev Med Child Neurol*. 2015; 57(5), 449–455.
7. Stroud H, Su SC, Hrvatin S, *et al*. Early-life gene expression in neurons modulates lasting epigenetic states. *Cell*. 2017; 171(5), 1151–1164. e16.
8. Schachtschneider KM, Welge ME, Auvil LS, *et al*. Altered hippocampal epigenetic regulation underlying reduced cognitive development in response to early life environmental insults. *Genes (Basel)*. 2020; 11(2), 162. doi: [10.3390/genes11020162](https://doi.org/10.3390/genes11020162).
9. Weber M, Hellmann I, Stadler MB, *et al*. Distribution, silencing potential and evolutionary impact of promoter DNA methylation in the human genome. *Nat Genet*. 2007; 39(4), 457.
10. Bernaudin M, Nedelec A, Divoux D, MacKenzie ET, Petit E, Schumann-Bard P. Normobaric hypoxia induces tolerance to focal permanent cerebral ischemia in association with an increased expression of hypoxia-inducible factor-1 and its target genes, erythropoietin and VEGF, in the adult mouse brain. *J Cereb Blood Flow Metab*. 2002; 22(4), 393–403.
11. Martens LK, Kirschner KM, Warnecke C, Scholz H. Hypoxia-inducible factor-1 (HIF-1) is a transcriptional activator of the TrkB neurotrophin receptor gene. *J Biol Chem*. 2007; 282(19), 14379–14388.
12. Thienpont B, Steinbacher J, Zhao H, *et al*. Tumour hypoxia causes DNA hypermethylation by reducing TET activity. *Nature*. 2016; 537(7618), 63–68.
13. Mutoh T, Sanosaka T, Ito K, Nakashima K. Oxygen levels epigenetically regulate fate switching of neural precursor cells via hypoxia-inducible factor 1 α -notch signal interaction in the developing brain. *Stem Cells*. 2012; 30(3), 561–569.
14. Smith AK, Kilaru V, Klengel T, *et al*. DNA extracted from saliva for methylation studies of psychiatric traits: evidence tissue specificity and relatedness to brain. *Am J Med Genet B Neuropsychiatr Genet*. 2015; 168B(1), 36–44.
15. Lewis SH, Perrin EV. *Pathology of the Placenta* (2nd ed.), 1999. Churchill Livingstone, London.
16. Khong TY, Bendon RW, Qureshi F, *et al*. Chronic deciduitis in the placental basal plate: definition and interobserver reliability. *Hum Pathol*. 2000; 31(3), 292–295.
17. Redline RW, Boyd T, Campbell V, *et al*. Maternal vascular underperfusion: nosology and reproducibility of placental reaction patterns. *Pediatr Dev Pathol*. 2004; 7(3), 237–249.
18. Redline RW, Faye-Petersen O, Heller D, *et al*. Amniotic infection syndrome: nosology and reproducibility of placental reaction patterns. *Pediatr Dev Pathol*. 2003; 6(5), 435–448.
19. Redline RW. Elevated circulating fetal nucleated red blood cells and placental pathology in term infants who develop cerebral palsy. *Hum Pathol*. 2008; 39(9), 1378–1384.
20. Redline RW, Ariel I, Baergen RN, *et al*. Fetal vascular obstructive lesions: nosology and reproducibility of placental reaction patterns. *Pediatr Dev Pathol*. 2004; 7(5), 443–452.
21. Redline RW. Inflammatory responses in the placenta and umbilical cord. *Semin Fetal Neonatal Med*. 2006; 11(5), 296–301.
22. Redline RW. Villitis of unknown etiology: noninfectious chronic villitis in the placenta. *Hum Pathol*. 2007; 38(10), 1439–1446.
23. Roberts DJ, Post MD. The placenta in pre-eclampsia and intrauterine growth restriction. *J Clin Pathol*. 2008; 61(12), 1254–1260.
24. Pathak S, Lees CC, Hackett G, Jessop F, Sebire NJ. Frequency and clinical significance of placental histological lesions in an unselected population at or near term. *Virchows Archiv*. 2011; 459(6), 565–572.
25. Stevens D, Al-Nasiry S, Bulten J, Spaanderman M. Decidual vasculopathy in preeclampsia: lesion characteristics relate to disease severity and perinatal outcome. *Placenta*. 2013; 34(9), 805–809.
26. Bendon RW. Review of autopsies of stillborn infants with retroplacental hematoma or hemorrhage. *Pediatr Dev Pathol*. 2011; 14(1), 10–15.
27. Gramellini D, Folli MC, Raboni S, Vadora E, Merialdi A. Cerebral-umbilical doppler ratio as a predictor of adverse perinatal outcome. *Am J Obstet Gynecol*. 1992; 79(3), 416–420.
28. Buemi M, Cavallaro E, Floccari F, *et al*. Erythropoietin and the brain: from neurodevelopment to neuroprotection. *Clin Sci (Lond)*. 2002; 103(3), 275–282.
29. Carmeliet P, Storkebaum E. Vascular and neuronal effects of VEGF in the nervous system: implications for neurological disorders. *Semin Cell Dev Biol*. 2002; 13(1), 39–53.
30. Gottmann K, Mittmann T, Lessmann V. BDNF signaling in the formation, maturation and plasticity of glutamatergic and GABAergic synapses. *Exp Brain Res*. 2009; 199(3–4), 203–234.
31. Koslowski M, Luxemburger U, Türeci Ö, Sahin U. Tumor-associated CpG demethylation augments hypoxia-induced effects by positive autoregulation of HIF-1 α . *Oncogene*. 2011; 30(7), 876–882.
32. Pierre CC, Longo J, Basseby-Archibong BI, *et al*. Methylation-dependent regulation of hypoxia inducible factor-1 alpha gene expression by the transcription factor kaiso. *Biochim Biophys Acta*. 2015; 1849(12), 1432–1441.
33. Steinmann K, Richter AM, Dammann RH. Epigenetic silencing of erythropoietin in human cancers. *Genes Cancer*. 2011; 2(1), 65–73.
34. Dewi FR, Fatchiyah F. Methylation impact analysis of erythropoietin (EPO) gene to hypoxia inducible factor-1alpha (HIF-1alpha) activity. *Bioinformation*. 2013; 9(15), 782–787.
35. Wallach I, Zhang J, Hartmann A, *et al*. Erythropoietin-receptor gene regulation in neuronal cells. *Pediatr Res*. 2009; 65(6), 619–624.
36. Sundrani DP, Reddy US, Joshi AA, *et al*. Differential placental methylation and expression of VEGF, FLT-1 and KDR genes in human term and preterm preeclampsia. *Clin Epigenetics*. 2013; 5(1). doi: [10.1186/1868-7083-5-6](https://doi.org/10.1186/1868-7083-5-6).

37. Pisani F, Cammalleri M, Dal Monte M, *et al.* Potential role of the methylation of VEGF gene promoter in response to hypoxia in oxygen-induced retinopathy: beneficial effect of the absence of AQP4. *J Cell Mol Med.* 2018; 22(1), 613–627.
38. Fang H, Chartier J, Sodja C, *et al.* Transcriptional activation of the human brain-derived neurotrophic factor gene promoter III by dopamine signaling in NT2/N neurons. *J Biol Chem.* 2003; 278(29), 26401–26409.
39. Viviani B, Bartesaghi S, Corsini E, *et al.* Erythropoietin protects primary hippocampal neurons increasing the expression of brain-derived neurotrophic factor. *J Neurochem.* 2005; 93(2), 412–421.
40. Aid T, Kazantseva A, Piirsoo M, Palm K, Timmusk T. Mouse and rat BDNF gene structure and expression revisited. *J Neurosci Res.* 2007; 85(3), 525–535.
41. Kundakovic M, Gudsnuk K, Herbstman JB, Tang D, Perera FP, Champagne FA. DNA methylation of BDNF as a biomarker of early-life adversity. *Proc Natl Acad Sci U S A.* 2015; 112(22), 6807–6813.
42. Bekkering I, Leeuwerke M, Tanis JC, *et al.* Differential placental DNA methylation of VEGFA and LEP in small-for-gestational age fetuses with an abnormal cerebroplacental ratio. *PLoS One.* 2019; 14(8). doi: [10.1371/journal.pone.0221972](https://doi.org/10.1371/journal.pone.0221972).
43. Hogg K, Price E, Hanna C, Robinson W. Prenatal and perinatal environmental influences on the human fetal and placental epigenome. *Clin Pharmacol Ther.* 2012; 92(6), 716–726.
44. McKennan C, Naughton K, Stanhope C, *et al.* Longitudinal data reveal strong genetic and weak non-genetic components of ethnicity-dependent blood DNA methylation levels. *Epigenetics.* 2020, 1–15.
45. Virani S, Dolinoy DC, Halubai S, *et al.* Delivery type not associated with global methylation at birth. *Clin Epigenetics.* 2012; 4(1), 8-7083-4-8.
46. Carmeliet P, de Almodovar CR. VEGF ligands and receptors: implications in neurodevelopment and neurodegeneration. *Cell Mol Life Sci.* 2013; 70(10), 1763–1778.
47. Oosthuysen B, Moons L, Storkebaum E, *et al.* Deletion of the hypoxia-response element in the vascular endothelial growth factor promoter causes motor neuron degeneration. *Nat Genet.* 2001; 28(2), 131–138.
48. Kopp S, Beckung E, Gillberg C. Developmental coordination disorder and other motor control problems in girls with autism spectrum disorder and/or attention-deficit/hyperactivity disorder. *Res Dev Disabil.* 2010; 31(2), 350–361.
49. Helmo FR, Lopes AMM, Carneiro ACDM, *et al.* Angiogenic and antiangiogenic factors in preeclampsia. *Pathol Res Pract.* 2018; 214(1), 7–14.
50. Hentges CR, Silveira RC, Prociyanoy RS. Angiogenic and antiangiogenic factors in preterm neonates born to mothers with and without preeclampsia. *Am J Perinatol.* 2015; 32(12), 1185–1190.
51. Emanuele E, Orsi P, Barale F, di Nemi SU, Bertona M, Politi P. Serum levels of vascular endothelial growth factor and its receptors in patients with severe autism. *Clin Biochem.* 2010; 43(3), 317–319.
52. Murray PS, Holmes PV. An overview of brain-derived neurotrophic factor and implications for excitotoxic vulnerability in the hippocampus. *Int J Pept.* 2011; 2011. doi: [10.1155/2011/654085](https://doi.org/10.1155/2011/654085).
53. Burd I, Welling J, Kannan G, Johnston MV. Excitotoxicity as a common mechanism for fetal neuronal injury with hypoxia and intrauterine inflammation. *Adv Pharmacol.* 2016; 76, 85–101.
54. Qin X, Feng J, Cao C, Wu H, Loh YP, Cheng Y. Association of peripheral blood levels of brain-derived neurotrophic factor with autism spectrum disorder in children: a systematic review and meta-analysis. *JAMA Pediatr.* 2016; 170(11), 1079–1086.
55. Gao R, Penzes P. Common mechanisms of excitatory and inhibitory imbalance in schizophrenia and autism spectrum disorders. *Curr Mol Med.* 2015; 15(2), 146–167.
56. Dunk CE, Roggensack AM, Cox B, *et al.* A distinct microvascular endothelial gene expression profile in severe IUGR placentas. *Placenta.* 2012; 33(4), 285–293.
57. Boule F, Van Den Hove D, Jakob S, *et al.* Epigenetic regulation of the BDNF gene: implications for psychiatric disorders. *Mol Psychiatry.* 2012; 17(6), 584–596.
58. D'Souza VA, Kilari AS, Joshi AA, Mehendale SS, Pisal HM, Joshi SR. Differential regulation of brain-derived neurotrophic factor in term and preterm preeclampsia. *Reprod Sci.* 2014; 21(2), 230–235.