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DNA Methylation differences after exposure to prenatal famine are common and timing- and sex-specific

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Abstract

Prenatal famine in humans has been associated with various later-life consequences depending on the gestational timing of the insult and the sex of the exposed individual. Epigenetic mechanisms have been proposed to underlie these associations. Indeed, animal studies and our early human data on the imprinted *IGF2* locus indicated a link between prenatal nutritional and DNA methylation. However, it remains unclear how common changes in DNA methylation are and whether they are sex- and timing-specific paralleling the later-life consequences of prenatal famine exposure. To this end we investigated the methylation of 15 loci implicated in growth and metabolic disease in individuals who were prenatally exposed to a war-time famine in 1944-45. Methylation of *INSIGF* was lower among individuals who were periconceptionally exposed to the famine (n=60) as compared with their unexposed same-sex siblings ($P=2 \times 10^{-5}$), whereas methylation of *IL10*, *LEP*, *ABCA1*, *GNASAS* and *MEG3* were higher (all $P < 10^{-3}$). A significant interaction with sex was observed for *INSIGF*, *LEP* and *GNASAS*. Next, methylation of 8 representative loci was compared between 62 individuals exposed late in gestation and their unexposed siblings. Methylation was different for *GNASAS* ($P=1.1 \times 10^{-7}$) and, in men, *LEP* ($P=0.017$). Our data indicate that persistent changes in DNA methylation may be a common consequence of prenatal famine exposure and that these changes depend on the sex of the exposed individual and the gestational timing of the exposure.

Introduction

Adverse environmental conditions during specific windows of mammalian development can have lasting effects on metabolic pathways and physiology, thereby influencing the susceptibility to chronic diseases¹. An extensive epidemiologic literature has reported associations between characteristics of early development and health outcomes later in life^{2,3}. Historical famines provide a quasi-experimental setting in which the long-term consequences of adverse conditions during development can be studied in humans. Studies of the Dutch Hunger Winter, a severe wartime famine at the end of WWII affecting the western part of The Netherlands, suggest that famine exposure *in utero* can lead to various adverse metabolic or mental phenotypes, depending on the sex of the exposed individual and the timing of the exposure during gestation⁴⁻⁷.

The period around conception may be especially sensitive to famine exposure⁸. Exposure to famine in this period is associated with diverse phenotypic outcomes such as an increased risk of adult schizophrenia⁷ and *spina bifida* at birth in men⁹. However, the effects of prenatal famine exposure are not limited to this developmental period. An example of a general, sex-specific late life effect is the increase in body mass index^{6,10} and various lipids in blood⁴ among famine-exposed women, irrespective of the precise gestational timing of the exposure. Also, increases in cerebro-cardiovascular related deaths have been reported among individuals exposed to seasonal food shortages independent of the gestational timing in a historical cohort¹¹, although preliminary results from the Dutch Famine indicated that an increased risk of coronary artery disease is specific for exposure to famine early in gestation¹². Animal experiments confirm the importance of timing and sex^{13,14}. Thus, prenatal exposure to famine can have different long-term effects that depend on the timing of the exposure and the sex of the exposed individual.

Persistent epigenetic changes induced by environmental factors are a plausible molecular mechanism underlying the relationship between early development and later life disease^{15,16}. Experiments in animal models provide

strong supporting evidence. Manipulation of the maternal diet during pregnancy lead to a persistent shift in average DNA methylation levels of specific genes in offspring resulting in permanent changes in coat color or tail shape^{17,18}. A proof-of-principle for complex diseases was reported by Bogdarina *et al.*, who showed in rats that a low protein diet during pregnancy was associated with decreased DNA methylation of the *Agtr1b* gene promoter in offspring, which may explain the increase in blood pressure among these animals¹⁹. We recently showed that similar mechanisms may be operative in humans; periconceptual exposure to famine was associated with a decrease in DNA methylation of the insulin-like growth factor 2 (*IGF2*) differentially methylated region (DMR)²⁰. However, to be a valid candidate mechanism in humans, the effect of prenatal famine on DNA methylation should be widespread, mirror the epidemiologic findings with sex- and timing-specific associations and affect genes in relevant pathways.

To further explore associations between prenatal famine and DNA methylation, including the role of the timing of exposure and the sex of the prenatally exposed individual, we assessed the methylation state of loci in 15 candidate genes involved in metabolic and cardiovascular disease and growth with diverse epigenetic features in our ongoing Hunger Winter Families Study²¹.

Results

We studied methylation of loci implicated in the transcriptional regulation of 15 candidate genes for metabolic and cardiovascular disease. The loci studied included imprinted loci (*GNASAS*, *GNAS A/B*, *MEG3*, *KCNQ1OT1*, *INSIGF* and *GRB10*), a putatively imprinted locus (*IGF2R*) and non-imprinted loci (*IL10*, *TNF*, *ABCA1*, *APOC1*, *FTO*, *LEP*, *NR3C1* and *CRH*). The selection of the loci measured was based on a combination of factors, including binding of methylation sensitive transcription factors, associations of DNA methylation with gene expression and the presence of differentially methylated regions for imprinted loci (a detailed overview is given in **Chapter 2, Table 1**). The characteristics of these loci, like the variation in DNA methylation, stability

over time and correlation with a tissue of a different embryonic lineage (buccal cells) was previously detailed (**Chapter 2**)²².

Periconceptual exposure

DNA methylation of these 15 loci was measured in 60 individuals conceived during the Dutch Famine (i.e. exposed periconceptionally) and compared with their unexposed same-sex sibling to minimize the possible confounding effects of familial environment and genetic background. For six of the fifteen loci, significant differences in DNA methylation were observed (**Table 1**). DNA methylation was increased among famine exposed individuals for the imprinted genes *GNASAS* ($P=3.1 \times 10^{-6}$) and *MEG3* ($P=8.0 \times 10^{-3}$) and the non-imprinted *IL10* ($P=1.8 \times 10^{-6}$), *ABCA1* ($P=8.2 \times 10^{-4}$) and *LEP* ($P=2.9 \times 10^{-3}$) proximal promoters. DNA methylation was decreased for the imprinted *INSIGF* promoter ($P=2.3 \times 10^{-5}$), which is part of the proximal promoter of *INS*²³. For the remaining 9 loci there was no association with periconceptual exposure to famine. All associations remained statistically significant after Bonferroni correction for multiple testing (15 loci) with the exception of *MEG3* ($P_{\text{Bonferroni}} = 0.12$). When analysed separately, individual CpG dinucleotides showed similar associations with famine exposure as the complete loci (**Figure 1A**). The loci affected did not share obvious features with respect to sequence, epigenetic features or biological function.

Late gestational exposure

Epigenetic modulation may also occur during other developmental windows^{24,25}. We therefore also studied 62 individuals who were exposed to famine late in gestation together with their unexposed, same-sex siblings. We measured DNA methylation at four loci that were associated with periconceptual famine exposure (*IL10*, *GNASAS*, *INSIGF* and *LEP*) and at four that were not (*IGF2R*, *APOC1*, *KCNQ1OT1* and *CRH*). These loci include four imprinted ones and diverse epigenetic features.

No associations were observed except for a significant reduction in methylation at the *GNASAS* locus ($P=1.1 \times 10^{-7}$, $P_{\text{Bonferroni}} = 8.8 \times 10^{-7}$) (**Table 2**). This association was consistent for the individual CpG sites within the locus (data not shown). The direction of the association was opposite to what was observed for periconceptual exposure. We then combined all periconceptual and late pregnancy exposed individuals and their controls in a single analysis to test for a statistical interaction between the famine

Table 1. DNA methylation and periconceptual exposure to famine.

Gene locus ¹	Average methylation % (SD)	Within pair difference ($\Delta\%$) ²	Effect size (SD units) ³	P-value ⁴	P-value Bonferroni corrected ⁶
<i>IL10</i>	20.8 (6.5)	2.4	0.37	1.8×10^{-6}	2.7×10^{-5}
<i>GNASAS</i>	48.8 (4.7)	1.1	0.24	3.1×10^{-6}	4.7×10^{-5}
<i>INSIGF</i>	84.8 (2.6)	-1.6	-0.61	2.3×10^{-5}	3.5×10^{-4}
<i>LEP</i>	28.6 (4.9)	1.2	0.24	2.9×10^{-3}	4.4×10^{-2}
<i>MEG3</i>	54.0 (2.4)	0.5	0.21	8.0×10^{-3}	0.12
<i>ABCA1</i>	19.9 (4.2)	0.7	0.17	0.017	0.26
<i>ABCA1</i> meth ⁵	36.9 (8.2)	1.7	0.21	8.2×10^{-4}	0.012
<i>KCNQ1OT1</i>	30.1 (1.5)	-0.2	-0.16	0.053	n.s.
<i>GRB10</i>	47.2 (4.6)	0.4	0.08	0.091	n.s.
<i>GNASAB</i>	40.3 (5.0)	0.6	0.11	0.092	n.s.
<i>APOC1</i>	16.7 (3.1)	-0.5	-0.17	0.13	n.s.
<i>IGF2R</i>	84.1 (6.9)	-0.7	-0.10	0.29	n.s.
<i>FTO</i>	97.3 (0.8)	-0.5	-0.61	0.28	n.s.
<i>CRH</i>	58.9 (6.0)	0.4	0.07	0.61	n.s.
<i>TNF</i>	9.6 (1.7)	0.1	0.06	0.63	n.s.
<i>NR3C1</i>	4.8 (1.1)	0.0	-0.01	0.79	n.s.

1. Table sorted on P-value.
2. Average absolute difference in DNA methylation between exposed and unexposed siblings.
3. Observed within pair difference divided by the standard deviation in the sibling controls.
4. Two-sided P-value resulting from a linear mixed model accounting for family relations, bisulfite batch and age at blood draw.
5. Separate analysis of methylated CpGs in *ABCA1*. In contrast to the 3' CpGs, 5' CpG dinucleotides (n=13) were methylated. The methylation of the methylated 5' CpGs was highly correlated ($R > 0.87$) amongst themselves, but not with methylation of the 3' CpG dinucleotides, which had little to no methylation (<4.5%).
6. The Bonferroni corrected P-values. Results that were already not significant before Bonferroni correction are shown as "n.s." (non significant).

associations with DNA methylation and the precise gestational timing of the exposure. The DNA methylation differences found for *IL10*, *GNASAS* and *INSIGF* were timing-specific (**Table 2**), but for *LEP* the test for interaction was not significant.

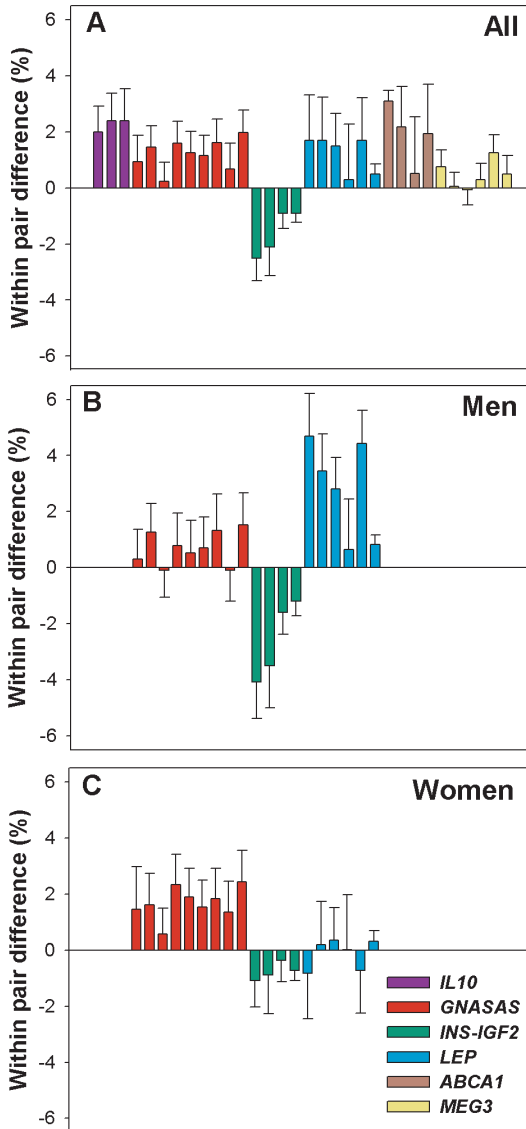


Figure 1. Difference in DNA methylation of CpG dinucleotides in siblings discordant for periconceptual exposure to famine.

Bars in the figures represent the average absolute within pair difference in DNA methylation and their standard errors for CpG dinucleotides. A positive difference indicates a higher methylation level among exposed individuals

A. The absolute within pair difference for CpG dinucleotides for which a significant overall difference of the locus in DNA methylation was observed.

B. As in panel A but for men. Only loci showing a significant interaction between sex and exposure are depicted.

C. As in panel B, but for women only.

Table 2. DNA methylation and late gestational exposure to famine.

Gene locus ¹	Average methylation % (SD) ²	Within pair difference ($\Delta\%$) ³	Effect size (SD units) ⁴	P-value ⁵	P-value Bonferroni corrected ⁶	P-value ⁷ timing specificity
<i>IL10</i>	20.7 (5.0)	-0.2	-0.04	0.76	n.s.	1.2×10^{-3}
<i>GNASAS</i>	48.8 (4.2)	-1.1	-0.26	1.1×10^{-7}	8.8×10^{-7}	3.1×10^{-12}
<i>INSIGF</i>	84.7 (2.8)	0.0	0.0	0.95	n.s.	3.2×10^{-4}
<i>LEP</i>	28.7 (4.6)	0.4	0.09	0.18	n.s.	0.13
<i>KCNQ1OT1</i>	30.2 (1.7)	0.2	0.12	0.17	n.s.	0.058
<i>APOC1</i>	16.7 (3.1)	-0.6	-0.19	0.22	n.s.	0.90
<i>IGF2R</i>	84.0 (4.4)	0.0	0.0	0.88	n.s.	0.25
<i>CRH</i>	58.9 (4.8)	0.5	0.10	0.51	n.s.	0.86

1. This table has been given the same order as table 1.
2. The batch corrected average methylation for the unexposed late gestation sibling controls and the standard deviation.
3. Average absolute difference in DNA methylation between exposed and unexposed siblings.
4. Observed within pair difference divided by the standard deviation in the sibling controls.
5. Two-sided P-value resulting from a linear mixed model accounting for family relations, bisulfite batch and age at blood draw. This test was performed on the late gestational exposed sibships (N=62)
6. The Bonferroni corrected P-values. Results that were already not significant before Bonferroni correction are shown as "n.s." (non significant).
7. Two-sided P value resulting from the test for timing specificity. The timing specificity was calculated by joining the datasets for both the periconceptional and late gestational siblings and their unexposed same-sex siblings and introducing an interaction term for gestational timing times exposure status. This test thus includes all 122 pairs.

Table 3. Sex-specific associations of DNA methylation with periconceptual exposure to famine.

Gene locus	P sex interaction ¹	Sex	Within pair difference ($\Delta\%$) ²	Effect size (SD units) ³	P-value ⁴
<i>GNASAS</i>	0.027	♂	0.7	0.15	0.013
		♀	1.5	0.33	1.1x10 ⁻⁵
<i>INSIGF</i>	8.5x10 ⁻³	♂	-2.6	-0.99	6.5x10 ⁻⁶
		♀	-0.8	-0.29	0.11
<i>LEP</i>	2.3x10 ⁻⁴	♂	2.8	0.57	3.6x10 ⁻⁷
		♀	-0.2	-0.04	0.70

1. The two sided P-value from the test for sex-specificity of the observed periconceptual effect of prenatal famine. This was tested by entering an interaction term of sex times the exposure status in the linear mixed model.
2. Average absolute difference in DNA methylation between exposed and unexposed siblings.
3. Observed within pair difference divided by the standard deviation in the sibling controls.
4. Two-sided P-value resulting from a linear mixed model accounting for family relations, bisulfitebatch and age at blood draw.

Sex-specific associations

Previous work found basal DNA methylation differences between men and women²⁶. We therefore first tested for sex differences in DNA methylation for the measured loci combining the two sibling control groups (men N=56, women N=66). DNA methylation was higher in men for *IGF2R* (+2.6%, P=0.019) and lower in men for *LEP* (-2.6%, P=3.0x10⁻³), *IL10* (-2.9%, P=0.015), and *APOC1* (-1.5, P=0.015) as compared to women. No significant differences were found for the other 4 loci (*GNASAS*, *INSIGF*, *KCNQ1OT1* and *CRH*). Next, we tested if the observed significant associations with prenatal famine were sex-specific. The interaction between sex and periconceptual famine exposure was significant for *LEP* (P=2.3x10⁻⁴), *INSIGF* (P=8.5x10⁻³) and *GNASAS* (P=0.027) (**Table 3**). For *LEP* and *INSIGF* the association of famine exposure with DNA methylation was restricted to men (P_{*LEP*, ♂}=3.6x10⁻⁷, P_{*INSIGF*, ♂}=6.5x10⁻⁶) (**Figure 1B and 1C**). For *GNASAS* the association was significant in both sexes but most pronounced in women (P_{men}=0.013; P_{women}=1.1x10⁻⁵). The association between late gestational famine exposure and *GNASAS* methylation was independent of sex.

Timing independent association

For *LEP* there was no indication for a significant interaction between the famine association with DNA methylation and the gestational timing of the exposure, even though the methylation difference was significant only following periconceptual famine exposure. Since, this association was later found to be male-specific, we tested for an interaction with sex in the late exposure group. Indeed, a significantly higher *LEP* methylation was found for men exposed late in gestation ($P=0.017$). Analysis of the whole cohort ($N=244$) including both exposure groups, revealed a significant association between prenatal exposure to famine irrespective of the precise gestational timing ($P=0.003$). Further analysis suggested that this association was male-specific ($P_{\text{interaction}} = 1.3 \times 10^{-6}$; men: +2.2% (0.83SD), $P=7.5 \times 10^{-8}$; women: $P=0.47$).

Discussion

We studied the DNA methylation levels of 15 loci for their association with prenatal exposure to the Dutch Famine at the end of WWII. For six of the loci studied, we observed significant differences in DNA methylation after famine exposure during periconception (*INSIGF*, *GNASAS*, *MEG3*, *IL10*, *LEP* and *ABCA1*). This association differed by sex for three loci (*INSIGF*, *GNASAS* and *LEP*). Of the eight loci tested, exposure to famine late in gestation was associated with methylation for *GNASAS* and for *LEP*, which was specific for men. Of interest, the differences in DNA methylation included both increases and decreases, in one case even at the same locus after exposure during different gestational periods. Together with our previous finding that the *IGF2* DMR is associated with periconceptual exposure²⁰ our data indicate that an adverse prenatal environment may trigger widespread and persistent changes in DNA methylation.

Our current and previous²⁰ observations suggest that the periconceptual period may be an especially sensitive exposure period in humans. This might be inherent to mammalian development^{27,28} and this hypothesis is also

supported by detailed animal studies^{29,30}. The association of *GNASAS* and *LEP* with late gestational exposure, however, suggests that environmentally induced DNA methylation changes may not be limited to the periconceptual period. This is in line with findings that methylation of the glucocorticoid receptor promoter depends on postnatal circumstances (e.g., maternal care in rats²⁵ and child abuse in humans²⁴). In addition to timing-specific associations, we observed sex-specific associations for three of the six loci for which DNA methylation was significantly associated with prenatal exposure to famine. In men, *LEP* methylation was associated with prenatal famine irrespective of the timing of exposure. Our observation that the methylation changes in relation to the prenatal environment may be sex-specific is in agreement with the sex-specific methylation changes found in offspring of sheep that were folate and vitamin B₁₂ restricted during periconception³⁰. How such sex-specific associations can arise is currently unknown, but interactions between sex hormones and the expression of DNA methyltransferases may be a factor³¹.

The differences in DNA methylation observed here are comparable, although slightly smaller, than we previously found for *IGF2* (absolute difference of 2.7% (5.2% relative to the mean DNA methylation level in the population)²⁰. The smaller average differences may be related to the inherent stochastic nature of epigenetic processes³² leading to a large variability in responses. The stochastic nature is strikingly illustrated by the large variation observed in the response of *agouti* gene methylation on maternal methyl donor supplementation even though the mice are inbred and the environmental conditions are highly controlled¹⁷. In human studies, genetic and environmental heterogeneity may further obscure the full impact of prenatal famine. In our study, we tried to minimize the heterogeneity caused by these factors by comparing exposed individuals with their unexposed, same-sex siblings. Another potential source of heterogeneity is the cellular diversity of whole blood, the tissue we studied. This heterogeneity is less likely to play a role for the affected imprinted loci *IGF2*²⁰, *GNASAS* and *MEG3*, since methylation is generally cell-type independent for imprinted loci³³ and the observed differences for non-imprinted loci between exposed individuals and controls

were similar. Indeed, with the exception of *IL10* we previously found that cell heterogeneity had little to no influence on the DNA methylation levels of the reported loci²².

With respect to our finding that *GNASAS* methylation was associated with exposure to famine late in gestation, it should be mentioned that a compromised late gestational development was hypothesized to lead to an immature immune system. The absolute numbers of lymphocytes were reported to be lower in children with a shorter gestation in a large, population-based study³⁴. However, since the proportions of the different cell types were not affected, it is unlikely that this has contributed to our findings.

The changes observed are comparable to those found in the liver of rats prenatally exposed to a protein-deficient diet, where promoter methylation of the *Ppara* promoter was decreased from 6.1% to 4.5% with individual CpG dinucleotides affected up to 5% and explaining up to 43% of the variance in gene expression of *Ppara*³⁵. This small absolute decrease of *Ppara* DNA methylation corresponded to a large ~26% relative change. It may be hypothesized that modest absolute changes in DNA methylation may lead to significant changes in gene expression for loci with a relatively low methylation level so that relative changes are substantial. The larger relative changes of *LEP* in men ($2.8/27.1=10.3\%$) and *IL10* ($2.4/20.8=11.5\%$) may be more likely to have functional consequences than that for *INSIGF* in men ($2.6/84.8=3.1\%$).

Similar to animal studies of methionine restriction³⁰, we observed both increases and decreases in DNA methylation, depending on the locus studied. Our results cannot be readily explained by damage due to a deficiency in dietary methyl donors due to the famine and may thus be part of an adaptive response. To definitely prove or disprove the existence of an adaptive response it will be necessary to characterize epigenetic responses of entire relevant pathways. Epigenetic mechanisms may contribute to the development of a thrifty phenotype^{1,15} and sex differences in this respect are increasingly well described¹³. It has been hypothesized that a thrifty phenotype may result from the combined effects of smaller epigenetic changes across the genome shifting metabolic networks³⁶. *INS*³⁷ and *LEP*³⁶, both affected by

prenatal famine exposure, were suggested to be particularly relevant in this respect. Our observations provide empirical evidence for these hypotheses in humans.

Our finding that the association of DNA methylation with prenatal conditions may depend on timing and sex matches the specificity in phenotypic outcomes observed for prenatal famine exposure, including neonatal outcomes⁹ and psychiatric⁷ and metabolic traits^{4,6,10}. It remains to be determined if the observed differences in DNA methylation in blood mark functional differences in relevant tissues for these traits. Previous work on *IGF2* DMR has shown that methylation in blood can mark the methylation level in a relevant tissue³⁸. Also, the CpG sites studied in the *LEP* promoter were previously reported to show similar methylation in peripheral blood and adipocytes *in vivo*³⁹ and to influence *LEP* expression *in vitro*⁴⁰. Changes in DNA methylation induced early in development, like by prenatal famine, may be propagated soma-wide contributing to the observed correlation in DNA methylation across tissues^{17,29}. A preliminary analysis of our data did not reveal significant associations between DNA methylation and plasma lipids and BMI that we previously reported to depend on prenatal exposure to famine in women in this cohort^{4,6}. But DNA methylation at most genes relevant to these particular complex phenotypes was found to be affected in men only, and not women. Similar to genetic studies, epigenetic studies linking variation in DNA methylation and complex disease will most likely require large study samples. Such studies may be particularly promising for *IL10*, which we found to be sensitive to periconceptual famine exposure. Genetic variation influencing *IL10* expression at this locus is associated with schizophrenia^{41,42}, a phenotype also particular to the periconceptual period⁷.

In summary, our study shows that exposure to famine in pregnancy may cause persistent changes in DNA methylation levels of multiple imprinted and non-imprinted genes with diverse biological functions. Our data support the hypothesis that associations between early developmental conditions and health outcomes later in life may be mediated by changes in the epigenetic information layer. Understanding how disturbances early in

human development are linked to later life disease may suggest new ways to prevent disease.

Materials and Methods

Study population

Study participants are a subset of the population from our ongoing Hunger Winter Families Study, whose design and recruitment was described previously²¹. In short, this study includes individuals exposed to famine during different periods in pregnancy and time-controls born before or after the famine. Study subjects were selected from births between 1943-1947 at three institutions in famine-exposed cities (the midwifery training schools in Amsterdam and Rotterdam and the Leiden University Medical Center). The daily rations distributed by the authorities during the famine period from November 28, 1944 to May 15, 1945 period contained an average energy equivalent of 667 kcal (SD, 151). During the famine there was little variation in the percentage of calories from proteins (12%), fat (19%) and carbohydrates (69%)⁴³. In addition to time controls from these three institutions, we recruited whenever possible a same-sex unexposed sibling of these individuals to serve as sibling control. Clinical examinations, including blood collections, were completed for 311 births in these institutions and 311 sibling controls. Ethical approval for the study was obtained from the participating institutions, and all participants provided written informed consent.

The subset selected for the present study includes two exposure groups with their unexposed sibling controls. The first exposure group comprises births conceived during the height of the famine, with periconceptual and early pregnancy exposure. The second exposure group comprises births during the height of the famine, with late pregnancy exposure. Periconceptionally exposed individuals were defined as births with a mother's last menstrual period between November 28, 1944 and May 15, 1945. This group includes 60 individuals (age 58.1y, SD 0.35), of whom 28 were male. Individuals exposed in late pregnancy were defined as births between January 28 and

May 30, 1945. This group includes 62 individuals (age 58.8y, SD 0.40), of whom 28 were male. As controls for each exposure group we used a matched same-sex sibling (age 57.1y, SD 5.50). The total study population therefore includes 244 individuals.

DNA methylation measurements

Genomic DNA was isolated from whole blood using the salting out method. One microgram of genomic DNA was bisulfite treated using the EZ 96-DNA methylation kit (Zymo Research). All samples were bisulfite treated on a total of three 96-well plates. Sibling pairs were on the same plate and periconceptionally and last trimester exposed pairs were equally distributed over the plates. DNA methylation of CpG dinucleotides were measured by a mass spectrometry based method (Epityper, Sequenom). The method determines the amount of DNA methylation by interrogating thousands of DNA copies assuming that 1 ng of genomic DNA equals ~300 copies. The quantitative nature, accuracy and reproducibility of this method has been shown extensively^{20,44,45}. All biochemical steps inherent to the methodology were performed according to the manufacturers' protocol. Bisulfite converted DNA specific PCR primers used to amplify the 15 investigated regions are summarized in **Supplement I, Table S4**. This table includes the precise genomic location of the amplified regions and an overview of the CpG sites quantized in each amplicon. All individuals exposed early in gestation and their sibling controls were measured in triplicate on a single 384 well plate for each locus and the same was true for the late gestational exposed individuals and their sibling controls. Data quality control and filtering were done as previously described⁴⁶. Data filtering consisted of the removal of CpG dinucleotides of which the measurement could be confounded by single nucleotide polymorphisms and of CpG dinucleotides of which the measurement success rate was below 80%. Common causes of a lower success rate include fragments bordering on the upper and lower limits of the mass range that can be detected and cases of fragments of which the base of the peak signal in the mass spectrum overlapped another fragment.

The success rate for the CpG containing fragments that could be measured within the limits of the methodology was 93.3%.

Statistical analysis

The analyses were performed within sib ships to minimize the possible confounding effects of differences in familial environment and genetic. We applied linear mixed models on the raw data without imputation of missing values to calculate exposure specific differences between sibling pairs. All the analyses account for family relations, age at examination, bisulfite plate and the correlation between CpG dinucleotides. Sib ship was entered as a random effect. Age at examination, bisulfite plate, exposure status and CpG dinucleotide, and where appropriate sex, were entered as fixed effects. The test for the timing specificity of the association between exposure and DNA methylation levels was done by adding a variable indicating the timing of the exposure and merging the periconceptual and last trimester datasets: timing specificity was tested by adding an interaction term of exposure status times the timing of the exposure to the linear mixed model. Testing the sex specificity of the associations between famine exposure and DNA methylation levels was done by adding a term for sex times the exposure status of the individuals in the linear mixed model.

The linear mixed model was chosen over a standard paired t-test because it allows for the analysis of multiple individual CpG dinucleotides in one test, accounts for the correlation between adjacent CpG dinucleotides, includes relevant adjustments within the model on the raw data, and uses all available data. The linear mixed model reduces to a test with identical outcome to a paired t-test if the within family difference is assessed for a single CpG dinucleotide, if no adjustments are performed and if there are no incomplete data for the sib pairs.

The basal difference in DNA methylation between men and women in the controls was calculated using ANOVA. All P-values reported are two-sided. All analyses were performed using SPSS 16.0.

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