Decades of research in rodent models has shown that early postnatal overnutrition induces excess adiposity and other components of metabolic syndrome that persist into adulthood. The specific biologic mechanisms explaining the persistence of these effects, however, remain unknown. On postnatal day 1 (P1), mice were fostered in control (C) or small litters (SL). SL mice had increased body weight and adiposity at weaning (P21), which persisted to adulthood (P180). Detailed metabolic studies indicated that female adult SL mice have decreased physical activity and energy expenditure but not increased food intake. Genomewide DNA methylation profiling identified extensive changes in hypothalamic DNA methylation during the suckling period, suggesting that it is a critical period for developmental epigenetics in the mouse hypothalamus. Indeed, SL mice exhibited subtle and sex-specific changes in hypothalamic DNA methylation that persisted from early life to adulthood, providing a potential mechanistic basis for the sustained physiological effects. Expression profiling in adult hypothalamus likewise provided evidence of widespread sex-specific alterations in gene expression. Together, our data indicate that early postnatal overnutrition leads to a reduction in spontaneous physical activity and energy expenditure in females and suggest that early postnatal life is a critical period during which nutrition can affect hypothalamic developmental epigenetics. *Diabetes* 62:2773–2783, 2013

Environmental influences on the development of body weight regulatory mechanisms may be an important factor in the worldwide obesity epidemic (1,2). Evidence in humans indicates that overnutrition during early postnatal life can permanently alter body weight regulation, increasing susceptibility to obesity throughout life (3,4). Accordingly, various animal models have been developed to explore the effects of infant overnutrition on lifelong obesity risk. Artificial feeding of rodent pups by intragastric cannula provides clear evidence for sustained effects of early postnatal overnutrition (5) but requires raising newborn rodents in isolation, which itself has long-term consequences. Overfeeding dams during lactation, with a high-fat diet, for example, could indirectly overnourish pups. Indeed, two recent rodent studies (6,7) report that the obesogenic effect of maternal high-fat diet occurs specifically during the suckling period. Pups from high-fat-fed dams are not consistently heavier at weaning, however (8,9), indicating that maternal overnutrition does not reliably induce early postnatal overnutrition.

In the rodent small litter model of early postnatal overnutrition (10), offspring from several litters born on the same day are randomized and fostered to either normal size (control [C]) or small litters (SL). Suckling in SL is naturalistic, easy to implement, and consistently induces early postnatal overnutrition, providing an apt model in which to study potential long-term effects of infantile overnutrition by excessive formula feeding (11). The early postnatal exposure induces elevated body weight and adiposity that persists to adulthood (10–12), with concomitant increases in plasma insulin (11,13) and leptin concentrations (13) and impaired glucose tolerance (11,13,14). It remains unresolved, however, whether the sustained increase in adiposity of adult SL rodents results from increased energy intake or decreased energy expenditure (13,15). Moreover, the fundamental mechanisms by which the metabolic effects of SL exposure persist to adulthood are unknown.

Environmental influences on developmental epigenetics (16,17) provide a likely mechanism. Epigenetic mechanisms regulate mitotically heritable alterations in gene expression that are not caused by changes in DNA sequence (18) and are known to play key roles in brain development (19). DNA methylation, the most stable epigenetic modification (20), is a likely mechanism to explain effects that persist for a lifetime (2). Given its central role in regulating food intake and energy expenditure (21), the hypothalamus is an obvious tissue in which to explore a potential epigenetic basis for induced alterations in body weight regulation. We therefore set out to determine 1) whether the persistently elevated adiposity of SL mice is caused by increased food intake or decreased energy expenditure, and 2) whether early postnatal overnutrition causes persistent changes in hypothalamic epigenetic regulation that may perpetuate altered body weight regulation.

RESEARCH DESIGN AND METHODS

For the litter size studies, virgin FVB/NJ females (The Jackson Laboratory) were mated with FVB/NJ males at age 5 weeks. In each batch, 14–15 mating pairs were set up on the same day; four independent batches of mice were studied over the course of 2 years. On postnatal day 1 (P1), pups from all litters born on the same day (P0) were weighed, sexed, and pooled randomly. Only pups from a birth litter size of 6–12 were included. Foster dams received either four (SL) or nine (C) pups. There were two females and two males in each SL and four to five females and males in each C litter. Litter assignment was performed systematically to balance body weight at P1. At P21, offspring from both groups were weaned onto a fixed-formula, soy protein–free diet (2020X; Harlan Teklad); females were housed two to five per cage, and males were housed individually. Body composition, food intake, energy expenditure, and physical activity were measured at P21–P25 and approximately P180. The P21 vs. P0 methylation–specific amplification and microarray hybridization (MSAM) comparisons used female C57BL/6J mice, and the pyrosequencing validation studies were performed in C57BL/6J and FVB/NJ mice of both sexes.
power of these time-series data while recognizing the nonindependence of the 24 multiple measures within each mouse. Analysis of food intake, energy expenditure, and physical activity were performed both with and without lean mass and fat mass included as independent variables to adjust for group differences in body size and composition (30). Group differences in DNA methylation by pyrosequencing were analyzed by repeated-measures ANOVA, with CpG site as the repeated effect (Supplementary Table 2). Loci that showed significant group effects on methylation but no significant group × age interaction were considered persistently altered by SL suckling. Requiring the same group difference, in the same direction, at both ages (in independent sets of mice) afforded substantial protection against type 1 error; these analyses were therefore not otherwise adjusted for multiple testing. Akaike information criterion (AIC) model selection by adjusted R² (SAS Proc Reg) was performed based on individual average dark-period energy expenditure and physical activity data. 

RESULTS
Early postnatal overnutrition reduces adult energy expenditure in females. We used the SL mouse model (Fig. 1A) to study persistent effects of overnutrition during the suckling period. We studied four independent batches (groups of litters cross-fostered at one time) over 2 years, including offspring from 24 C and 26 SL litters total. Consistent with previous studies, SL mice were heavier at P21 and remained so into adulthood (P < 0.0001 in both females and males) (Fig. 1B). Although the increase in adult body weight was modest, effects on body composition were substantial. Both male and female SL adults had 50% higher fat mass and percent body fat compared with C mice (P < 0.005 in all comparisons) (Fig. 1C). There were no group differences in lean mass. Clearly, suckling in a small litter induces persistent changes in regulatory mechanisms that affect adult body composition.

To determine whether these changes involve alterations in food intake and/or energy expenditure, we used metabolic cages to simultaneously monitor food intake, energy expenditure, and voluntary physical activity. In an attempt to identify persistent metabolic differences that might explain the sustained group differences in adiposity, we performed the metabolic measurements shortly after weaning (P25) and in adulthood (P180). (Again, these data represent four batches of mice studied over the course of 2 years.) After appropriate least squares normalization for lean mass and fat mass (30), food intake of SL mice tended, surprisingly, to be slightly lower than that of C mice at both P25 and P180 (Fig. 2A), but these differences were not statistically significant. Energy expenditure (normalized for lean mass and fat mass [30]) was nearly identical between SL and C mice at P25 (Fig. 2B). At P180, however, energy expenditure of SL females was significantly lower than that of C females (P = 0.002); this group difference was significant during both the light and dark periods. Resting metabolic rate was estimated as the lowest average energy expenditure within 1 h for each mouse. After least squares normalization for lean mass and fat mass, female mice showed no group differences in resting metabolic rate at either age. Resting metabolic rate of SL males, however, was higher at P25 (P = 0.02) and lower at P180 (P = 0.03) relative to C males. There were no group differences in respiratory exchange ratio. Group differences in voluntary physical activity, again normalized for lean mass and fat mass, were consistent with those in energy expenditure: none were found at P25, but SL females were significantly less active than C females at P180, specifically during the dark period (group × light interaction P = 0.0003) (Fig. 2C).

It is noteworthy that adult SL females were less physically active even after including body weight and body composition in the model; hence, their lower activity
was not caused by their excess adiposity. (For comparison, unnormalized data on food intake, energy expenditure, and physical activity are shown in Supplementary Fig. 1.) Including physical activity in the model for energy expenditure of P180 females drastically reduced the significance of the SL effect (from $P = 0.002$ to $P = 0.01$), suggesting that physical activity explains much of the group difference in energy expenditure. Together, these data indicate that the persistent alterations in energy balance of female SL mice are due not to excess food intake but, rather, to reduced energy expenditure. The sex specificity of this effect may be related to the male-specific decline of physical activity with age (Supplementary Fig. 1C).

**Extensive epigenetic development occurs in the early postnatal hypothalamus.** Ontogenic periods when epigenetic mechanisms are being established or undergoing maturation constitute critical periods during which environmental influences can cause persistent changes in epigenetic regulation (31,32). To determine whether the suckling period might be a critical period for developmental epigenetics in the hypothalamus, we tested for changes in hypothalamic DNA methylation. We used MSAM, which is based on sequential digestion of genomic DNA with the methylation-sensitive and -insensitive isoschizomers SmaI and XmaI (23,33). Two independent P21 vs. P0 MSAM cohybridizations (incorporating a dye swap) were performed. Using stringent criteria validated in our previous studies (24,33), 868 SmaI/XmaI intervals changed methylation from P0 to P21. Only 31 intervals lost methylation (Fig. 3A), and the genomic distribution of associated SmaI/XmaI cut sites was not different from that on the array (Fig. 3B and Supplementary Fig. 2). Methylation increased at 837 intervals (Fig. 3A); associated cut sites were significantly underrepresented at promoters ($P < 0.0001$) and overrepresented in introns ($P < 0.0001$) (Fig. 3B). In a larger number of P0 and P21 mice, we used bisulfite pyrosequencing (23,33) to measure P0–P21 changes in hypothalamic CpG methylation at 10 intervals identified by

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**FIG. 1.** SL mice are persistently heavier and fatter than C mice. **A:** Overview of the litter size experiment. FVB mice were cross-fostered at P1 and randomly assigned to SL (green) or C (orange) litters. Quantitative magnetic resonance (QMR) and CLAMS measurements were performed after weaning (P21–P25) and at P180. Hypothalami were isolated at approximately P25 and P180. Four independent batches of SL and C mice were studied over the course of 2 years. **B:** Body weight of SL and C mice did not differ at P1 ($P > 0.7$). SL mice showed significantly higher body weight at P21 (insets), which was maintained to adulthood (P180) ($P < 0.0001$ in both females and males). Data are presented as means ± SEM of 20–94 mice in each group, sex, and age. (Error bars are smaller than symbols.) Box plots (insets) represent median (mid-line), 25th–75th percentiles (box), and 5th–95th percentiles (whiskers). **C:** Both male and female SL mice have higher adiposity at P25 (left panel) ($P < 0.01$). By P180 (right panel), group differences in body composition are much greater both in absolute terms (top panel) ($P < 0.002$) and as percent of body weight (bottom panel) ($P < 0.004$). Plots represent 20–30 mice of each group, sex, and age (**$P < 0.01$).
MSAM; all 10 validated (100%) (7 are shown in Fig. 4). Additionally, since the P21 vs. P0 MSAM studies were performed in C57BL/6J mice, we confirmed (at a subset of loci) that these methylation changes also occur in both sexes of FVB/NJ mice (the strain used for the litter size studies) (Supplementary Fig. 3).

Functional significance of the developmental changes in DNA methylation was evaluated by gene ontology analysis. Relative to all potentially informative genes on the array, no enriched ontologies were found for the few genes associated with intervals that lost methylation. Genes that gained methylation, however, were significantly enriched for 20 biological process categories (Fig. 3C); of these, 17 are explicitly related to development, including neurodevelopmental processes such as axon guidance and neuron differentiation. Postnatal development of the mouse hypothalamus clearly involves functionally important epigenetic changes. This may be a critical period during which environment can influence these processes, with long-term consequences.

Early postnatal overnutrition causes persistent and sex-specific alterations in hypothalamic DNA methylation and gene expression. We therefore examined DNA methylation differences among SL and C hypothalami.

Intrigued by the large group differences in energy expenditure and physical activity in P180 females, we used MSAM to compare hypothalamic DNA methylation of SL and C females at P180. Two independent SL vs. C cohybridizations were performed, with each hypothalamic DNA sample pooled from five females drawn from different foster litters. The results, however, provided no evidence of persistent group differences in hypothalamic DNA methylation.

Reasoning that DNA methylation changes in SL mice might be too subtle to detect by MSAM, we used bisulfite pyrosequencing to examine a panel of candidate genes. Since genomic regions undergoing methylation change from P0 to P21 are most likely to show persistent effects of overnutrition during this period (2), most of the genomic regions that we selected were those identified in our P21 vs. P0 MSAM experiments. In addition to 10 of those already validated (Fig. 4), we examined six hits near genes previously reported to change expression in hypothalamus from P0 to P21 (34) and two showing interindividual variation in DNA methylation. Additionally, promoters of a few genes critical to hypothalamic function and development (Agrp, Fto, and Pomc) were included. In total, 24 loci

FIG. 2. Adult SL females have reduced energy expenditure and physical activity. A: Hourly data on food intake are presented as least squares means, adjusting for lean mass and fat mass. Light/dark cycle is indicated by shading. Food intake of female and male SL mice did not differ from that of C mice at any age. A, B, and C; mean ± SEM of 20–30 mice in each group, sex, and age. B: Hourly data on energy expenditure are presented as least squares means, adjusting for lean mass and fat mass. No group differences were found at P25. At P180, SL females had significantly lower energy expenditure (P = 0.002), and this group difference was seen during both the light and dark cycles. SL males likewise tended to have lower energy expenditure at P180, but this difference was not statistically significant. C: Hourly data on physical activity are presented as least squares means, adjusting for lean mass and fat mass. No group differences were found at P25. At P180, SL females had significantly lower activity, specifically during the dark cycle (P = 0.0009). SL males likewise tended to have lower activity at P180, but this difference was not statistically significant (**P < 0.01; ***P < 0.001).
Of these, 15 showed no DNA methylation differences between SL and C hypothalamus at P25 (Supplementary Fig. 4 and Supplementary Table 2) and were not examined at P180. The remaining nine loci were examined at both P25 and P180 (Supplementary Fig. 5 and Supplementary Table 2). Considering this an exploratory data analysis, we set an $a$-level of 0.1 for main effects and 0.2 for interactions. The initial model including both sexes showed significant main effects for age but not group (SL vs. C); interestingly, however, four loci showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2). We therefore performed sex-specific analyses. In females, AK145544, Aqp4, and Nolz1 and in males Gadd45b showed significant group × sex interactions (Supplementary Table 2).
correction, there was an enrichment of low P value probes in the group × sex analysis (Fig. 6A), suggesting subtle sex-specific effects. We therefore performed gene ontology analysis on the 37 and 732 transcripts with at least a 20% group difference (P, 0.05) in females and males, respectively. (The reference list comprised the 14,628 transcripts significantly expressed in hypothalamus.) No enriched gene ontology terms were found in females. In males, however, for both the 381 genes upregulated and the 351 downregulated in SL hypothalamus, the foremost biological process related to formation of neuronal projections. Examination of the gene ontology terms associated with the genes comprising these enrichments (Fig. 6B and Supplementary Table 5) suggests a subtle shift in expression profile that may favor neuronal remodeling in the hypothalamus of adult SL males. Additionally, in an analysis of gene networks associated with the expression changes in male hypothalamus (Supplementary Fig. 7), two of the top three networks were related to cellular development and nervous system development. These networks are centered on Atn1 and dynein, respectively (both regulators of neurodegeneration), again supporting potential alterations of neuronal remodeling in SL males.

FIG. 4. Validation of P21 vs. P0 MSAM by bisulfite pyrosequencing. A: At AK04543, the P0-P21 methylation decrease identified by MSAM was confirmed. At all other regions analyzed, methylation increases identified by MSAM were confirmed: Amn (B), Podn (C), Tmem154 (D), Tnfrsf1a (E), Nolz1 (F), and Shank3 (G). Gray columns indicate the SmaI/XmaI cut sites. Data are represented as means ± SEM of 5–10 mice per age. CpG site locations are provided relative to transcription start site (TSS) or transcription end site (TES).
DISCUSSION

Here we showed that early postnatal overnutrition, known to permanently increase body weight and adiposity, also reduces voluntary physical activity and energy expenditure in adult females. These physiological changes were associated with persistent alterations in hypothalamic DNA methylation at specific loci. Overall, these findings provide support for the hypothesis that early postnatal overnutrition causes subtle but widespread changes in hypothalamic epigenetic regulation that persist to influence adult energy balance.

Our study addresses a key outstanding question: whether the persistently altered energy balance of SL rodents is due to increased food intake or decreased physical activity. Previous studies reported increased food intake (13,36) and energy expenditure (37) in adult SL rodents. Those conclusions, however, were based on non-normalized data, disregarding the altered weight and body composition of SL rodents. Here, we used least squares means to appropriately adjust expenditure and intake data for body weight and composition (30). Compared with C mice of the same weight and body composition, adult SL mice were not hyperphagic (Fig. 2A). Their energy expenditure, however, again compared with C mice of the same weight and body composition, was lower (Fig. 2B), significantly so in females. Hence, our data provide strong evidence that reduced energy expenditure, not increased food intake, explains the increased adiposity of female adult SL mice.

In addition to food intake and energy expenditure, however, there are other determinants of energy balance, such as nutrient absorption, which were not measured in this study. Also, it is possible that group differences in central regulation of food intake may have been unmasked if mice were provided a highly palatable diet (38). Other than physical activity, we did not measure additional determinants of energy expenditure, such as brown adipose tissue activity. These shortcomings may explain why the excess adiposity of male SL mice occurred without measurable differences in physical activity or energy expenditure. (Notably, a recent study found age-associated alterations in thermogenic capacity of brown adipose tissue.)
demonstrated by increased tissue in male SL rats [39], consistent with our finding that resting metabolic rate is increased at P21 but decreased at P180 in male SL mice.) In females, the reduced energy expenditure was largely explained by reduced physical activity (Fig. 2C). In an earlier report in rats, prenatal undernutrition likewise caused persistent reductions in locomotor activity, most prominently in females (40). Given the worldwide trends of decreasing physical activity (41), it is crucial to determine whether, in humans as in rodents, nutrition during early life modulates voluntary physical activity for a lifetime.

Despite its importance in central regulation of food intake and energy expenditure (21), our understanding of the molecular mechanisms driving functional development of the hypothalamus remains limited. Mouse hypothalamic development continues into early postnatal life, a critical period for formation of leptin-sensitive neuroanatomic projections that function in energy homeostasis (42) and major alterations of hypothalamic gene expression (34). Here, we have shown for the first time that during this same period widespread changes in DNA methylation—mostly increases—are underway. The association of these methylation increases with genes involved in neural development (Fig. 3C) suggests a process of postnatal epigenetic maturation. Because projections from the arcuate nucleus of the hypothalamus to other brain regions form prenatally in primates but postnataally in rodents (43), it is often proposed that hypothalamic development during the
suckling period in the mouse is comparable with that in a third-trimester human. It is currently unknown, however, whether and when the epigenetic maturation we have documented in the postnatal mouse occurs in humans. Moreover, our findings that postnatal overnutrition leads to a decrease in physical activity in female mice raises the question of whether the mouse is a good model for physical activity in humans. Although we currently have only a rudimentary understanding of the neurobiological regulation of spontaneous physical activity (44), the hypothalamus and other brain regions are known to be involved, as are several highly conserved neuropeptides including cholecystokinin, corticotrophin-releasing hormone, leptin, and orexins.

It was recently reported that SL rats have alterations in hypothalamic DNA methylation (45,46). Those studies, however, assessed DNA methylation only at P21. Our data therefore provide the first evidence that early postnatal overnutrition induces persistent epigenetic changes in the hypothalamus. Additionally, unlike previous studies on related models (45–48), rather than focus on single CpG sites we performed integrated analysis of all CpG sites represented in each assay because 1) regional changes in DNA methylation are more likely to affect gene expression and 2) concordant changes at multiple adjacent sites are less likely to occur by chance. Notably, contrary to the previous report of increased DNA methylation (at 2 of 20 CpG sites measured) at the Pomc promoter in the hypothalamus of P21 SL rats (45), our methylation assay spanning five nearby CpG sites found no SL vs. C differences at P25 (Supplementary Fig. 4).

We developed the strategy of examining genomic regions undergoing DNA methylation changes from P0 to P21, based on the conjecture that these changes may be susceptible to environmental influences. Indeed, 4 of 21 regions undergoing P0–P21 DNA methylation change showed evidence of persistent methylation differences between SL and C mice, supporting the utility of this approach. Hence, the ~900 loci that we report that undergo postnatal methylation changes may provide useful candidate regions for future studies of environmental influences on hypothalamic developmental epigenetics.

With the potential exception of AK145544, all four genes with persistent changes in hypothalamic DNA methylation in SL hypothalami play important roles in neural development or function (49–51). At each of these genes, the methylation change in SL mice was modest (Fig. 5); the cumulative effect of such changes at hundreds or thousands of genes, however, could be considerable. This interpretation is supported by the AIC model selection (Supplementary Fig. 6), which included as significant predictors of adult energy expenditure expression of three and methylation at two of the genes that we identified, in most cases with $P$ values comparable with that of lean body mass.

The results of transcriptional profiling in P180 hypothalamus mirrored our DNA methylation analyses in detecting subtle, widespread, and sex-specific alterations in gene expression. Analyzing the corpus of genes with potentially altered expression in male SL hypothalamus identified highly significant gene ontology enrichments pertaining to regulation of neuronal projections (Fig. 6). The adult rodent hypothalamus maintains significant synaptic plasticity (52); our data suggest that early postnatal overnutrition in males may persistently augment this capability. Adult mice that become obese owing to a high-fat diet, conversely, appear to have reduced hypothalamic neurogenesis (53).

All the potential explanatory effects we found—changes in energy expenditure, physical activity, DNA methylation, and gene expression—were sex specific. The long-term consequences of early life exposures have long been recognized to differ by sex (16). Our findings of sexual dimorphism in the epigenetic responses to early postnatal environment suggest that nutrition may interact with the epigenetic mechanisms regulating hormone-dependent sexualization of the neonatal hypothalamus (54). In fact, the sex differences found here might provide an answer as to why the lower physical activity in SL females arose only in adulthood. In male mice, physical activity declined with age in both groups, but in females this decline was seen only in SL mice (Supplementary Fig. 1C). Our results may suggest, therefore, that postnatal overnutrition is leading to masculinization of the CNS pathways that regulate age-related changes in physical activity.

Encouraged by earlier studies that gained insights into hypothalamic developmental epigenetics (55), we too studied DNA methylation in whole hypothalamus. The interpretation of our data is therefore complicated by the heterogeneity of the hypothalamus at both the regional and the cellular level. The hypothalamus is comprised of distinct regions, or “nuclei,” with specialized functions, gene expression patterns (21), and epigenetic regulation (56). Additionally, the nervous system includes diverse cell types; the simplest classification distinguishes neurons and glia, which are epigenetically distinct (57,58). To improve our understanding of how early postnatal overnutrition causes persistent changes in regulation of body weight and body composition, it will be advantageous to characterize epigenetic effects within specific hypothalamic nuclei and specific cell types. For example, based on our current data we cannot exclude the possibility that the persistent alterations in DNA methylation that we identified represent a shift in the proportion of hypothalamic cell types rather than induced alterations in epigenetic regulation within specific cell types. Moreover, since early postnatal life is a critical period for not only epigenetic but also neuroanatomic development (42), studying these processes in an integrated fashion will likely be necessary to gain a clear understanding of how early postnatal nutrition affects the establishment of hypothalamic body weight regulation.

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G.L. and J.J.K. performed experiments and wrote the manuscript. W.Z. and E.L. performed experiments. G.K.-R. performed bioinformatic analyses. M.S.B. performed experiments. M.L.F. provided critical guidance on experimental procedures and edited the manuscript. R.A.W. designed the study and wrote the manuscript. R.A.W. is the guarantor of this work and, as such, had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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