Proopiomelanocortin (POMC) neurons in the arcuate nucleus of the hypothalamus are central components of systems regulating appetite and energy homeostasis. Here we report on the establishment of a mouse model in which the ribonuclease III ribonuclease Dicer-1 has been specifically deleted from POMC-expressing neurons (POMC<Hsup>−/−</sup>DCR), leading to postnatal cell death. Mice are born phenotypically normal, at the expected genetic ratio and with normal hypothalamic POMC-mRNA levels. At 6 weeks of age, no POMC neurons/cells could be detected either in the arcuate nucleus or in the pituitary of POMC<Hsup>−/−</sup>DCR mice. POMC<Hsup>−/−</sup>DCR develop progressive obesity secondary to decreased energy expenditure but unrelated to food intake, which was surprisingly lower than in control mice. Reduced expression of AgRP and ghrelin receptor in the hypothalamus and reduced uncoupling protein 1 expression in brown adipose tissue can potentially explain the decreased food intake and decreased heat production, respectively, in these mice. Fasting glucose levels were dramatically elevated in POMC<Hsup>−/−</sup>DCR mice and the glucose tolerance test revealed marked glucose intolerance in these mice. Secondary to corticotrope ablation, basal and stress-induced corticosterone levels were undetectable in POMC<Hsup>−/−</sup>DCR mice. Despite this lack of activation of the neuroendocrine stress response, POMC<Hsup>−/−</sup>DCR mice exhibited an anxiogenic phenotype, which was accompanied with elevated levels of hypothalamic corticotropin-releasing factor and arginine-vasopressin transcripts. In conclusion, postnatal ablation of POMC neurons leads to enhanced anxiety and the development of obesity despite decreased food intake and glucocorticoid deficiency. (Molecular Endocrinology 27: 1091–1102, 2013)
signals impacting on appetite and energy expenditure. The first is the neuropeptide Y (NPY) and agouti-related protein (AgRP) containing neurons that are stimulated by ghrelin and inhibited by leptin with the net effect of increasing appetite (3); and second are the proopiomelanocortin (POMC)- and cocaine and amphetamine-regulated transcript-containing neurons that are stimulated by leptin. POMC is processed to α-MSH that acts on melanocortin receptor 4 (MC4R) expressed in neurons situated in the hypothalamic paraventricular nucleus (PVN) and the nucleus of the solitary tract (NTS) in the hindbrain, thus suppressing appetite and increasing energy expenditure (4). NPY/AgRP neurons may reduce the melanocortinergic tone by activation of Y1 receptors on POMC neurons, and/or by direct AgRP antagonism of MC4R-expressing PVN neurons (2).

The clinical importance of the melanocortin system in weight regulation is evident from patients harboring mutations in the pomc gene (5), in the prohormone convertase 1 (6), and in the mc4r gene (7). These monogenic forms of obesity are characterized by early-onset hyperphagia and an extreme obese phenotype. Similar phenotypes also occur in POMC knockout mice models (8, 9). Although slightly different strategies were used to generate these POMC-deficient mice, the end point was similar, ie, a complete ablation of POMC-derived peptides. Despite that, it was striking to find significant phenotype differences between these models, mainly related to the onset of obesity, thyroid function, and glucose levels. The possible impact of compensatory changes on normal physiology and on our ability to interpret the resulting phenotypic changes is a well-recognized limitation of developmental models of genetic manipulations (10). Furthermore, knowledge obtained from isolated knockout of the pomc gene may not fully reveal the whole span of functions mediated by the POMC neuron.

To better characterize the physiological roles of the POMC neuron, we sought to establish a mouse model in which ablation of POMC-expressing neurons is achieved early in the postnatal period. To this end, we chose to genetically delete the ribonuclease III ribonuclease Dicer-1 (dcr-1) gene specifically from POMC neurons (POMCΔDcr mice), based on the knowledge that Dicer ablation has been shown to deplete cells of mature microRNAs, leading to apoptosis and cell death (11). Using this model, we show that postnatal ablation of POMC neurons leads to the development of obesity, insulin resistance, and enhanced anxiety-like behavior despite decreased food intake and glucocorticoid deficiency.

Materials and Methods

Animals

Mice were maintained in a pathogen-free, temperature-controlled (22°C ± 1°C) mouse facility on a reverse 12-hour light, 12-hour dark cycle at the Weizmann Institute of Science, according to institutional guidelines. Food and water were given ad libitum. Mice were fed regular chow diet (Harlan Biotech Israel Ltd, Rehovot, Israel). All experimental protocols were approved by the Institutional Animal Care and Use Committee of the Weizmann Institute of Science.

For the experiments described here, Pomp-Cre mice (stock number 005965; Jackson Laboratory, Bar Harbor, Maine) were initially mated with mice carrying the Dicer-1floxed allele (12). Hemizygous Pomp-Cre and heterozygous Dicer1floxed offspring were then intercrossed to produce study animals. For visualizing POMC neurons, mice were crossed with the conditional R26R YFP reporter line (12). The Pomp-Cre mice were genotyped using the following primers: forward (TGCCACGACCAAGTCACACG), reverse (CCAGGTTACGGATATAGTTCAT), with an annealing temperature of 55°C. The Dicer-1floxed mice were genotyped as previously described (13). The conditional R26R YFP reporter mice were genotyped using the following oligonucleotide primers: wild type (GCCGATAGTTGCTTCACACC), transgene (AAAGTCGCTCTGTTGGTAT), and wild type reverse (GAGCCGGAGAATGGATGTG), with an annealing temperature of 60°C.

Histological and immunohistological analysis

Mice were crossed with a Rosa26 YFP reporter mouse line for the detection of POMC neurons in the adult. Mice were transcardially perfused with 4% paraformaldehyde, and brains were sectioned on a freezing sliding microtome. Immunostaining for yellow fluorescent protein (YFP) was performed using biotinylated anti-green fluorescent protein antibody raised in goat as primary antibody (Abcam, Cambridge, United Kingdom). Streptavidin-conjugated Cy3 was used as secondary antibody (Jackson ImmunoResearch Laboratories Inc, West Grove, Pennsylvania). Fluorescence images were captured using a Nikon i80 microscope (Nikon, Tokyo, Japan).

For morphological studies, the pituitaries were sectioned horizontally and slides were stained with hematoxylin and eosin and with immunohistochemistry to localize ACTH as described previously (14). Immunolocalization was detected with the streptavidin-biotin-peroxidase complex technique and 3,3′-diaminobenzidine. Negative controls were performed with normal rabbit serum replacing the primary antiserum and after preabsorption of the primary antiserum with purified antigen.

Adrenal paraffin sections were stained by Mayer’s hematoxylin and eosin (H&E) by a standard procedure.

Interscapular brown adipose tissue (BAT) was processed for immunohistochemical staining for Ucp1. Immunohistochemistry was carried out on paraffin sections with the avidin-biotin-peroxidase (avidin biotin complex method; Vector Laboratories, Burlingame, California) using the rabbit anti-Ucp1 Ab (Abcam). Sections were lightly counterstained with hematoxylin.
Circulating corticosterone levels
For the evaluation of the endocrine response to stress, tail blood samples were collected before (basal), immediately after 25 minutes of restraint stress, and at 80 and 120 minutes after stress initiation. Restraint stress was induced using a 50-mL ventilated conical tube. Plasma samples were immediately centrifuged and stored at −80°C until assays for hormone measurement were conducted. Corticosterone concentrations were quantified using a corticosterone enzyme immunoassay kit (Cayman Chemical Co, Ann Harbor, Michigan).

Plasma hormones and lipid level measurement
After a 5-hour fasting period, mice were killed and trunk blood was collected. Plasma samples were immediately centrifuged and stored at −80°C. Free T₄ and TSH were measured by chemiluminescence (Advia Centaur; Siemens, Melvern, Pennsylvania). Cholesterol and triglycerides were measured in a chemical autoanalyzer (Advia 1650; Siemens, Melvern, Pennsylvania). The MILLIPLEX MAP mouse gut hormone panel (Millipore Corp, St Charles, Missouri) was used for the simultaneous quantification of active glucagon-like peptide 1 (GLP-1), insulin, and leptin in mouse plasma according to the manufacturer’s protocol. Briefly, MILLIPLEX MAP is based on the Luminex xMAP technology, which uses internally color-coded microspheres with fluorescent dyes, thus creating distinctly colored bead sets, each of which is coated with a specific capture antibody. After an analyte from a test sample is captured by the bead, a biotinylated detection antibody is introduced, and the reaction mixture is then incubated with the streptavidin-phycoerythrin conjugate, the reporter molecule, to complete the reaction on the surface of each microsphere. The antibody pairs in the panel are specific to the desired analyte and exhibit no or negligible cross-reactivity with other analytes in the panel. Samples are read on a Luminex 200. The intraassay and interassay variations are less than 7% and less than 23%, respectively. Assay sensitivities [minimum detectable concentrations (picograms per milliliter)] for insulin, leptin, and GLP-1 are 25, 22, and 15 pg/mL, respectively.

Homeostasis model assessment insulin resistance index (HOMA-IR) calculation
The baseline degree of insulin resistance was estimated by homeostasis model assessment according to the method described by Matthews et al (15). In particular, an insulin resistance score (HOMA-IR) was computed with the formula: fasting plasma glucose (milligrams per deciliter) × fasting serum insulin (milliunits per liter)/405. Low HOMA-IR values indicate high insulin sensitivity, whereas high HOMA-IR values indicate low insulin sensitivity (insulin resistance).

Metabolic studies
Indirect calorimetry and food and water intake as well as locomotor activity were measured using the Labmaster system (TSE-Systems, Bad Homburg, Germany). The LabMaster instrument consists of a combination of feeding and drinking sensors for automated online measurement. The calorimetry system is an open-circuit system that determines O₂ consumption, CO₂ production, and respiratory exchange ratio. A photobeam-based activity monitoring system detects and records ambulatory movements, including rearing in every cage. All the parameters are measured continuously and simultaneously. Data were collected after 48 hours of adaptation in acclimated singly housed mice. Cumulative food intake data was recorded every 30 seconds and was further used for meal microstructure analysis.

Meal microstructure analysis
We used a 2-state hidden Markov model (16) to distinguish between a state of hunger and a state of satiation, which are characterized by different distribution of food and water intake. These distributions were estimated from the eating and drinking patterns of 8 control mice during 1 day. In addition, these data were used to estimate the probabilities of transition between the states. These probabilities as well as the state distribution were estimated using the Baum-Welch algorithm (17). In the test, we found the most likely state sequence that matched the data. In this way we were able to identify epochs of eating out of hunger, which we dubbed as meals. The sequences were found using the Viterbi dynamic programming algorithm (18).

Glucose tolerance tests
Glucose tolerance tests were performed in awake animals during the dark phase of the day cycle. Following 6 hours of fasting, glucose (2 g/kg of body weight) was injected ip, and whole venous blood obtained from the tail vein at 0, 15, 30, 60, 90, and 120 minutes after the injection was measured for glucose using an automatic glucometer (Accu-Check performa; Roche Diagnostics GmbH, Mannheim, Germany).

Core body temperature
Core body temperature was measured using rectal thermometer at a constant hour during 3 consecutive days.

Body composition
Body composition was assessed using Echo-MRI (Echo Medical Systems, Houston, Texas).

Behavioral tests for anxiety-like behavior
All behavioral studies were performed during the dark phase after habituation to the test room for 2 hours before any test. For the assessment of anxiety-like behaviors, the light-dark transfer and the elevated plus maze tests were used. The set-up for the light-dark transfer test consists of 2 compartments, a dark one (14 × 27 × 26 cm) and a 950-lux illuminated lit compartment (30 × 27 × 26 cm), connected by a small passage. Mice were placed in the dark compartment to initiate a 5-minute test session. The time spent in the lit compartment, the number of entries to the lit compartment, and the latency of entering the lit zone were measured. The indices collected in these tests were quantified using an automated video tracking system (VideoMot2; TSE Systems).

The elevated plus maze apparatus is comprised of a central part (5 × 5 cm), 2 opposing open arms, and 2 opposing closed arms (30.5 × 5 × 15 cm). The apparatus was elevated to a height of 55 cm, and the open arms were illuminated with 6 lux. Mice were placed in the center, facing an open arm to initiate a 5-minute session test. The time spent in the open arms and the number of entries to the open arms was measured.
Acoustic startle response test

The acoustic startle response (ASR) test was performed as previously described (19). Briefly, the ASR session started with a 5-minute acclimatization period, with background white noise [70 dB(A)], which was maintained throughout the session. Overall, 32 startle stimuli [120 dB(A), 40 milliseconds; intertrial interval (randomly varying): 12–30 seconds] were presented; the stimulus presentations were divided into three blocks: blocks 1 and 3 each consisted of 6 startle stimuli, whereas block 2 consisted of 10 startle stimuli and 10 no stimuli [70 dB(A), 40 milliseconds; ie, equivalent to the background white noise] that were presented in a quasirandom manner. The entire ASR test session was completed in 20 minutes. Two indices were recorded: 1) mean maximal ASR (grams) (ie, the mean maximal ASR of the responses to the startle stimuli in block 2); and 2) the rate of habituation to the startle stimuli within a session [ie, the percentage change in mean maximal ASR (grams) of block 3 compared with that of block 1].

RNA preparation and real-time PCR

Total RNA was extracted from hypothalami, pituitaries, or BAT using a 5 Prime PerfectPure RNA cell and tissue kit (5 Prime GmbH, Hamburg, Germany). RNA was reverse transcribed to generate cDNA using high capacity cDNA reverse transcription kit (Applied Biosystems Inc, Foster City, California). The cDNA products were used as templates for real-time PCR analysis. Sense and antisense primers were selected to be located on different exons to avoid false-positive results caused by DNA contamination. Oligonucleotide primers for real-time PCR analysis were designed using Primer Express software (Applied Biosystems, PerkinElmer, Dublin, California). The specificity of the amplification products was checked by melting curve analysis.

Table 1. Primers Used for Real-Time PCR Analysis

<table>
<thead>
<tr>
<th>Gene</th>
<th>Forward Primer</th>
<th>Reverse Primer</th>
<th>GenBank Accession Number</th>
</tr>
</thead>
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<tr>
<td>Adrb3</td>
<td>GMACAAAATCTGCTGTATAT</td>
<td>GCCATTTAAGAGATCTCCAC</td>
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<tr>
<td>AgRP</td>
<td>AAAGCTTTGGCAGAGCTAGA</td>
<td>AACGAGGACTCTGCAAGCTACA</td>
<td>NM_007427.2</td>
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<tr>
<td>AVP</td>
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<td>NM_009732.1</td>
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<td>CRFR1</td>
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<td>AAAAGCGATAGAGGTCTCAG</td>
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<tr>
<td>DIO2</td>
<td>TCTCCTAGATCATCAAAAGAG</td>
<td>ATTCAGAGATGAGGTCTCAG</td>
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<tr>
<td>GHSR</td>
<td>AAGGACAGATGAGAGTCTGCGT</td>
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<tr>
<td>HPRT1</td>
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<td>GGTCTTTTTTACCAACAGCT</td>
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<td>Leptin R</td>
<td>ACATCGAGGAGGAGACATG</td>
<td>GGTTCTTTAGTAAATGCTCCT</td>
<td>NM_146104</td>
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<tr>
<td>MC3R</td>
<td>CTCAGACCAAGGGTCTGCGT</td>
<td>ATCAGCTGCTGCTCACAC</td>
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<td>MC4R</td>
<td>CAGAAGATCTGCACTCCACCA</td>
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<td>NPY</td>
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<td>Pomc</td>
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<td>SF-1</td>
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<td>NM_139051.3</td>
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<td>Trh</td>
<td>AACTTTGCTGCTTTAGATTC</td>
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Abbreviations: HPRT1, hypoxanthine guanine phosphoribosyl transferase 1; Leptin R, leptin receptor; SF-1, steroidogenic factor 1.

Statistical analysis

Results are expressed as means ± SEM. Statistical analysis was performed using ANOVA or repeated-measurements 2-way ANOVA with post hoc Student’s t tests or paired Student’s t tests, as appropriate using JMP software (SAS Institute, Cary, North Carolina) (*, P < .05; **, P < .01).

Results

POMC<sub>ADCR</sub> mice lose their POMC-expressing neurons and pituitary corticotropes

To establish a mouse line in which the POMC neurons are ablated postnatally, we cross-bred mice carrying conditional Dicer alleles (13), DCR<sup>lox/lox</sup>, with mice expressing Cre recombinase specifically in POMC-expressing neurons/cells (21) (Figure 1A). The R26R-YFP conditional reporter alleles (12) in the genetic background of the DCR<sup>lox/lox</sup> mouse line enabled the detection of Cre-mediated recombination by the induction of YFP expression (Figure 1, A and B).

POMC<sub>ADCR</sub> mice are born phenotypically normal and at the expected genetic ratio. In contrast with the strong YFP immunostaining obtained in control animals, no POMC neurons could be detected in the arcuate nucleus of the hypothalamus in POMC<sub>ADCR</sub> mice at 6 weeks of age, indicating that microRNAs are essential for survival of these neurons (Figure 1B). Accordingly, POMC-mRNA expression in the hypothalamus of 2-month-old mice was reduced by 96% compared with control animals (P < .001, Figure 1C). To establish the time course of this neuronal loss, POMC-mRNA expression was examined in hypothalami excised at sequential time points. On postnatal day (PND) 1, POMC-mRNA expression is similar in POMC<sub>ADCR</sub> and their control littermates, whereas at PND7 and PND21, there was a reduction of 35% and
83% in POMC-mRNA expression levels, respectively (Figure 1C). POMC-mRNA expression was similar in mice heterozygous for the Dicer allele (POMC<sup>ΔDCR</sup>/H<sub>11001</sub>/H<sub>11002</sub>) and control mice in all the tested time points (data not shown), indicating that loss of only 1 dicer allele does not lead to POMC neuronal loss. In contrast to the hypothalamus, POMC expression in the NTS was not affected, as demonstrated by immunostaining for YFP (Figure 1D).

Loss of POMC-expressing cells was evident in the pituitaries of POMC<sup>ΔDCR</sup> mice. ACTH immunostaining was negative in both the anterior and the intermediate lobes of the pituitary gland (Figure 1E). Furthermore, there was marked atrophy of the pituitary intermediate lobe, indicating loss of melanotrope cells, which are the predominant cell type in this region (Figure 1E). POMC and corticotropin-releasing factor receptor 1 (CRFR1) mRNA were undetectable in the pituitary gland of POMC<sup>ΔDCR</sup>, consistent with loss of corticotropes (Figure 1F). In contrast, steroidogenic factor 1, a gonadotrope marker, was expressed at similar levels in POMC<sup>ΔDCR</sup> and control animals. As expected from the loss of the entire population of pituitary corticotrope cells, both basal and stress-induced corticosterone levels were undetectable in these mice (Figure 1G). Loss of ACTH tone led
to changes in both adrenal weight and morphology. Adrenal weight of POMCΔDCR mice was markedly reduced compared with that of control animals (Figure 1H), which could be mainly attributed to atrophy of the zona fasciculata layer (Figure 1I).

POMCΔDCR mice develop obesity and diabetes at an early age

At 4 weeks of age, it was already evident that both male and female POMCΔDCR mice were heavier than POMCΔDCR+/− and control littermates (males, 19.28 ± 0.75, 17.07 ± 0.57, and 14.2 ± 0.96 g, respectively, P < .001; females, 19.35 ± 0.62, 14.58 ± 0.32, and 14.6 ± 0.91 g, respectively, P < .001; Figure 2, A and B). Although at this age it was apparent that POMCΔDCR+/− males' weight was higher than control mice (P < .05), this difference was subsequently lost as POMCΔDCR weight curves rapidly diverged upward in comparison with POMCΔDCR+/− and control curves that progressed in parallel (2 way ANOVA P < .0001 for both genders, Figure 2, A and B). We next explored the possibility that POMCΔDCR+/− may have an intermediate obese phenotype when challenged with a high-fat diet, as has been described in heterozygous pomc mice (8). Weight gain was similar to that observed in controls (data not shown), reflecting the similar hypothalamic POMC-mRNA levels found in these genotypes. In addition to the prominent weight gain, the POMCΔDCR mice showed a significant increase in percentage fat mass and a decrease in lean mass compared with control or POMCΔDCR+/− littermates (Figure 2C). Interestingly, despite the higher percentage of fat mass, total cholesterol, high-density lipoprotein cholesterol, and low-density lipoprotein cholesterol were significantly lower in POMCΔDCR mice, whereas triglyceride levels were similar (Figure 2, D–G). Fasting glucose levels were dramatically elevated in POMCΔDCR mice (141.5 ± 5.0 vs 94.3 ± 13.2 in control mice, P < .05) and a glucose tolerance test revealed marked glucose intolerance in POMCΔDCR mice (ANOVA P < .01), with peak glucose levels reaching 357 ± 34 mg/dL vs 247 ± 13 mg/dL at 30 minutes (P < .001) (Figure 2H).

Insulin levels and HOMA-IR were already significantly higher in POMCΔDCR mice at the age of 7 weeks (Table 2). Interestingly, at 7 weeks of age, basal GLP-1 levels were slightly higher in the POMCΔDCR mice (93 ± 1.4 pg/mL) in comparison with control mice (85 ± 0.7 pg/mL, P = .005), but at 4 months, this difference was lost (Table 2).

Food intake and metabolic rate are altered in POMCΔDCR mice

At the age of 6 weeks, food intake was significantly higher in POMCΔDCR mice (Figure 3A), but when corrected to metabolic body weight (BW0.75), the food intake was significantly lower (Figure 3B). In adulthood, a nonsignificant tendency toward lower absolute food intake of the POMCΔDCR mice was observed (Figure 3A). However, when corrected to metabolic body weight, the food intake was significantly lower relative to control littermates (Figure 3, A and B).

![Figure 2](image_url). Increased body weight and adiposity in POMCΔDCR mice is accompanied by alterations in lipids and glucose homeostasis. A and B, Growth curves of males (A) and females (B) POMCΔDCR, POMCΔDCR+/−, and control littermates (n = 7–12), demonstrating a significant increase in body weight in the POMCΔDCR mice. C, Body composition demonstrating increased adiposity and reduced lean mass in POMCΔDCR mice (n = 7). D–G, Total cholesterol (D), high-density lipoprotein (HDL) cholesterol (E), low-density lipoprotein (LDL) cholesterol (F), and triglycerides (G), demonstrating altered lipid profile of POMCΔDCR mice. H, Glucose tolerance test showing impaired glucose tolerance in POMCΔDCR mice (n = 6). *, P < .05; **, P < .01. Values are expressed as the mean ± SEM.
Microstructural analysis of eating behavior was performed for the dark (active) phase of the day cycle. As seen in Table 3, the number of meals consumed as well as the food and water amount per meal was similar between the groups. Interestingly, POMC<sup>ΔDCR</sup> mice meal duration was significantly longer due to slower eating rate.

Indirect calorimetry revealed that POMC<sup>ΔDCR</sup> mice had a decreased metabolic rate, which was reflected by the reduction of both light and dark phase oxygen consumption and carbon dioxide production as well as decreased heat production (Figure 3, C–E). These changes were not reflected in the respiratory exchange ratio, which remained similar in both groups (not shown). The decrease in metabolic rate and energy expenditure could not be attributed to alteration in locomotor activity that was similar in both groups (Figure 3F).

POMC<sup>ΔDCR</sup> mice had reduced body core temperature (Figure 4A) and reduced expression of UCP1, peroxisome proliferator-activated receptor γ coactivator 1α, β-3 adrenergic receptor (Adrb3), and type II iodothyronine de-iodinase (DIO2), genes associated with thermogenesis in BAT (Figure 4, B and C), suggesting that the reduced heat production in these mice is a consequence of reduced sympathetic tone. The reduced expression of DIO2 as well as the reduced metabolic rate were not secondary to decreased thyroid function because free T<sub>4</sub> levels was similar in POMC<sup>ΔDCR</sup> and control mice (Figure 4D). However, despite the equivalent free T<sub>4</sub> levels, hypothalamic TRH mRNA levels were significantly lower in POMC<sup>ΔDCR</sup> mice (Figure 4E), probably reflecting the low melanocortinergic tone in these mice. To explore additional mechanistic changes that may contribute to the metabolic phenotype of POMC<sup>ΔDCR</sup> mice, we have examined the hypothalamic mRNA expression levels of neuropeptides and receptors involved in the regulation of energy homeostasis as well as plasma levels of leptin. AgRP mRNA levels were robustly reduced by 87% (<i>P < .01</i>) in POMC<sup>ΔDCR</sup> mice, potentially explaining the decreased food intake in these mice (Figure 4E). In accordance to the lepintin resistance state associated with obesity, plasma leptin levels were already significantly elevated in 7-week-old POMC<sup>ΔDCR</sup> mice, and a further increase was observed at the age of 4 months compared with the control mice (Table 2). The expression of hypothalamic leptin receptor, ghrelin receptor, and MC4Rs were decreased in the POMC<sup>ΔDCR</sup> mice, whereas MC3R and NPY expression was similar to that found in the control mice (Figure 4E).

**POMC neuron ablation induced a basal state of enhanced anxiety**

Because loss of corticotropes in POMC<sup>ΔDCR</sup> mice resulted in inactivation of the hypothalamus-pituitary-adrenal (HPA) axis, we explored whether this affected the mice performance in anxiety-like be-

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**Table 2. Cytokine and HOMA-IR in Control and POMC<sup>ΔDCR</sup> Mice**

<table>
<thead>
<tr>
<th></th>
<th>Seven Weeks</th>
<th>Four Months</th>
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<tr>
<td></td>
<td>Control</td>
<td>POMC&lt;sup&gt;ΔDCR&lt;/sup&gt;</td>
</tr>
<tr>
<td>Insulin, ng/mL</td>
<td>1.12 ± 0.18</td>
<td>2.68 ± 0.63&lt;sup&gt;a&lt;/sup&gt;</td>
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<tr>
<td>HOMA-IR</td>
<td>11.98 ± 1.99</td>
<td>35.73 ± 9.78&lt;sup&gt;a&lt;/sup&gt;</td>
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<tr>
<td>Leptin, ng/mL</td>
<td>1.42 ± 0.21</td>
<td>6.60 ± 1.86&lt;sup&gt;a&lt;/sup&gt;</td>
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<tr>
<td>GLP-1, pg/mL</td>
<td>84.95 ± 0.75</td>
<td>93.06 ± 1.43&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

<sup>a</sup> <i>P < .05</i> compared with control mice at the same age.

<sup>b</sup> <i>P < .01</i> compared with control mice at the same age.

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(*Figure 4A*) and reduced expression of UCP1, peroxisome similar among experimental groups. Values are expressed as the mean ± SEM, *, <i>P < .05</i>; **, <i>P < .01</i> vs control (n = 7–8).

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**Figure 3.** Food intake and metabolic rate are altered in POMC<sup>ΔDCR</sup> mice. A and B, Daily food intake in young and adult POMC<sup>ΔDCR</sup> and control littermates in absolute values (A) and normalized to metabolic body weight (BW<sup>0.75</sup>) (B). Demonstrated a significant decrease in food intake in the POMC<sup>ΔDCR</sup> mice. C–E, Heat production (C), oxygen consumption (VO<sub>2</sub>) (D), and carbon dioxide production (VCO<sub>2</sub>) (E) are lower in adult POMC<sup>ΔDCR</sup> mice during both the dark and light phases of the day cycle, indicative of lower metabolic rate. F, Locomotor activity is similar among experimental groups. Values are expressed as the mean ± SEM, *, <i>P < .05</i>; **, <i>P < .01</i> vs control (n = 7–8).
havioral tests. POMCΔDCR mice exhibit a robust anxiogenic phenotype as assessed by the dark-light transfer test, the elevated plus maze, and the acoustic startle response. In the dark-light transfer test, POMCΔDCR mice spent less time in the light compartment (Figure 5A), showed longer latency to enter the light compartment (Figure 5B), and traveled a shorter distance during the test (Figure 5C). In the elevated plus maze, the POMCΔDCR mice spent less time in the open arms (Figure 5D). The acoustic startle response test is known to be corticosterone independent because it is not modified by adrenalectomy (22). Therefore, we performed this test with the intent of analyzing the anxiety response while accounting for the absent stress induced corticosterone response in the POMCΔDCR mice. The POMCΔDCR and the control mice had similar reaction times (POMCΔDCR: 9.64 ± 0.72 seconds; control: 8.46 ± 0.37 seconds), but the maximum response (Figure 5E) and the within-session rate of habituation (Figure 5F) were markedly higher and lower, respectively, in the POMCΔDCR mice. The anxiogenic phenotype of POMCΔDCR mice was accompanied by elevated levels of both corticotropin-releasing factor (CRF) and arginine-vasopressin (AVP), both known to have an anxiogenic effect (Figure 5G), and probably reflecting a compensatory response to the decreased POMC levels and to the absence of the negative feedback provided by corticosterone.

### Discussion

By deleting the ribonuclease III ribonuclease Dicer-1 gene specifically from POMC neurons and corticotropes, we succeeded in establishing a mouse model in which the POMC cells are ablated postnatally and mice are born with hypothalamic POMC-mRNA levels equivalent to that of the control littermates. By avoiding the partial embryonic lethality observed in previous models (8, 9) and by delaying POMC neuron ablation to the postnatal period, we may have reduced possible compensatory mechanisms that could confound the phenotype of these mice.

One of the main findings of the current study is that the obese phenotype generated by POMC neuron ablation is maintained by a decrease in energy expenditure without the expected increase in food intake, which one would anticipate from loss of the melanocortinergic tone. In fact, corrected food intake was decreased in young animals and even more so in adult POMCΔDCR mice, in striking contrast to the hyperphagia described in mice models of pomc gene deletion (8, 9, 23). One could speculate that ablation of POMC neurons abrogated feeding regulatory pathways that are not mediated through activation of the pomc gene. Possible explanations for the decrease in food intake are the significantly reduced hypothalamic mRNA levels of the orexigenic peptide AgRP as well as that of the ghrelin receptor (GHSR). Reduced AgRP expression has also been described in the pomc-null mouse model generated by the group of O’Rahilly and colleagues (8, 24).
The absence of peripheral glucocorticoids is probably one of the major factors contributing to the reduced expression of AgRP because it has been shown that AgRP expression was dramatically reduced in adrenalectomized mice (25) and that corticosterone administration increased AgRP expression levels both on wild-type and pomc-null background (24, 25). The reduced expression of GHSR may also be partially attributed to the absence of glucocorticoids because dexamethasone was shown to stimulate the GHSR mRNA and protein expression in a hypothalamic cell line (26). The critical role played by glucocorticoids is well exemplified by the reversal of the obese phenotype and the restoration of the hypothalamic melanocortic tone in leptin-deficient ob/ob mice by adrenalectomy (25).

Of note, in the pomc-null mice characterized by Smart et al (23), a similar tendency for a decrease in food intake toward adulthood was found; although corrected food intake was higher than controls at 6 weeks, it was significantly lower at 26 weeks (23). A similar transient hyperphagia has been described in mice lacking leptin receptors from both POMC and AgRP neurons (27), suggesting a differential weight regulation at young age. A parallel could be made to humans harboring heterogeneous mutations in the MC4R, who present with increased appetite in childhood, with the progressive amelioration of the phenotype such that the ad libitum energy intake at the age group of 11–15 years was approximately half of that in children up to 5 years of age, as expressed by kilocalories per kilogram of lean mass (28).

Two significantly different paradigms have been previously used to study the effects of acute or gradual POMC neuron ablation on parameters of energy homeostasis. Diphtheria toxin administration to adult mice selectively expressing a diphtheria toxin receptor in POMC neurons caused acute degeneration or necrosis of more than 85% of these neurons within 72 hours of treatment initiation (29). An increase in food intake and body weight, without changes in blood glucose levels, was observed during the 10-day follow-up period. In the second model, POMC-expressing cells were gradually ablated by selective deletion of the mitochondrial transcription factor A (tfam), leading to a progressive loss of cellular respiration and consequent cell death (30). Reduction in α-MSH immunostaining was evident in pomc-specific tfam mutant mice only at 7 months of age, although excess weight gain, secondary to both increased food intake and reduced energy expenditure, was already apparent at 4 months of age and proceeded progressively compared with control animals and as such were suggested as a model for age-related obesity. Clearly, other than the resultant obese phenotype, it is extremely difficult to draw

Figure 5. Ablation of POMC-expressing cells leads to increased anxiety-like behavior. A–D, Anxiety-like behaviors of POMC<sup>ΔDCR</sup> and control littermates as measured by the dark-light transfer (DLT) (A–C) and by the elevated plus maze (EPM) tests (D). The reduced percentage of time spent in the light compartment (A), longer latency entering the light compartment (B), shorter distance traveled in the light compartment of the DLT (C), and the reduced time spent in the open arms of the EPM (D) are indicative of anxiogenic behavior. E and F, Anxiety-like behaviors as measured by the ASR. The increased maximal ASR (E) and reduced within-session ASR rate of habituation (F) points toward an increase in anxiety-like behavior. Hypothalamic expression of the anxiogenic peptides, CRF and AVP (G). *, P < .05; **, P < .01 vs control. Values are expressed as the mean ± SEM (n = 8–10). Max g, maximum response.
other parallels between our model, in which POMC-cell ablation is generated in the neonatal period, and those in which neuronal loss occurs acutely at adulthood or gradually toward senescence.

Another major point to be taken from our study is that the main factor leading to the maintenance of obesity in adulthood was the decreased metabolic rate and not increased food consumption. The decreased metabolic rate of POMC\textsuperscript{ADCR} mice could be explained by alteration in BAT activity. Energy expenditure in BAT is highly regulated by sympathetic stimulation. Norepinephrine released from sympathetic nerve terminals activates the BAT via binding to \( \beta \)-adrenergic receptors. Adrenergic innervations regulate both Ucp1 expression and activity and increases DIO2 activity, leading to increased intracellular concentrations of T\(_3\) (31). The reduced BAT Adbrb3 expression in POMC\textsuperscript{ADCR} mice suggests a reduction in sympathetic tone and may explain, at least in part, the reduction in Ucp1 and DIO2 expression level. Because NTS POMC neurons are intact in POMC\textsuperscript{ADCR} mice, the reduced sympathetic tone could be due to changes in sympathetic activation arising from the hypothalamus. This can involve both first-order neurons because POMC neurons were demonstrated, using retrograde tracing to project polysynaptically to BAT (32) or via second-order neurons in the PVN, a key source of autonomonic outflow (33).

Maintenance of obesity due to diminished metabolic rate cannot be overemphasized in view of its clinical implications for the care of patients suffering from hypothalamic obesity. These subjects, frequently children, develop obesity secondary to hypothalamic damage by tumors such as craniopharyngiomas or secondary to their treatment (34). In general, parents or family members report an acute increase in food intake at diagnosis or shortly after surgical tumor excision, although, to the best of our knowledge, this hyperphagia has not been formally quantified. Nevertheless, at a later stage of the disease, these patients were found to have normal or even decreased food intake, coupled with a significant reduction in basal metabolic rate (35, 36), which may be attributed to the abnormal sympathetic tone secondary to hypothalamic damage. Hence, clinical interventions based on caloric restriction are doomed to fail in achieving sustained weight loss, thus leading to frustration and further psychological distress (34). Interestingly, obese subjects heterozygous for \( mc4r \) were shown to have decreased urinary norepinephrine excretion (37), and their body mass index was inversely correlated with vasoconstrictive muscle sympathetic nerve activity, a direct measure of central sympathetic nervous outflow (38). Thus, in addition to their reported hyperphagia, reduced metabolic rate emerges as another important factor to be addressed in patients harboring \( mc4r \) mutations, which accounts for approximately 5% of cases with body mass index greater than 30 kg/m\(^2\) (28).

The phenotypic characterization of our model highlighted another significant difference consequent of POMC neuron ablation compared with knockout of the \( pome \) gene: POMC\textsuperscript{ADCR} mice were diabetic and insulin resistant from an early age, in contrast to the model created by Hochgeschwender and colleagues (9, 39) in which mice with a deletion of the coding region for all POMC peptides had normal fasting and insulin levels as well as normal glucose tolerance, despite their marked obesity. In the mouse model generated by the group of O’Rahilly and colleagues (8, 24), POMC-deficient mice have higher fed blood glucose but equivalent insulin levels compared with their wild-type littermates and become diabetic only upon corticosterone administration from weaning but not in adulthood. Similarly, normoinsulinemic POMC-null mice developed insulin resistance only upon restoration of HPA axis function by introducing a POMC transgene selectively in the pituitary gland.

Our results confirm and strengthen the published data indicating the central role of POMC neurons in the regulation of whole-body glucose homeostasis, including glucose sensing (40) and modulation of pancreatic glucagon secretion and hepatic insulin sensitivity (41). Activation of leptin, insulin (42), and serotonin 2C receptors (43) on POMC neurons has been shown to regulate different aspects of glucose homeostasis. The observation that isolated \( pome \) gene ablation causes no or minimal alteration in glucose levels may suggest that the melanocortinergic pathways may not be involved or essential in mediating these effects. Alternatively, it is possible that \( pome \) gene-deleted mice developed in utero compensatory mechanisms to circumvent the striking disruption of glucose homeostasis that was observed with POMC neuron ablation.

In view of the expanding interest in the relationship between stress-regulatory systems and eating behavior, we explored the effect of POMC neuron ablation on anxiety parameters. We hypothesized that POMC\textsuperscript{ADCR} mice would exhibit decreased anxiety behavior in view of the known anxiogenic effects of \( \alpha \)-MSH through the activation of MC4R (44). Surprisingly, POMC\textsuperscript{ADCR} mice exhibit a robust anxiogenic phenotype as assessed by several well-established behavioral tests for assessing anxiety-like behavior. The elevated CRF and AVP levels, both anxiogenic peptides, may reflect a compensatory response to the decreased POMC levels and to the absence of the negative feedback provided by corticosterone (44). Additional detailed pharmacological or genetic studies are needed to fully elucidate the molecular and cellular
mechanisms that are mediating the behavioral changes observed in these mice. In this sense, this is a unique model to study the specific effects of CRF on anxiety, independent of the effects secondary to the increased HPA axis activation characteristic of models of CRF overexpression or central CRF administration (45). The observed decrease in ghrelin receptors, possibly leading to reduced ghrelin action may be an additional contributory factor because this peptide has been shown to have anxiolytic and antidepressant effects (46, 47).

In summary, this model emphasizes that the main mechanism leading to obesity secondary to POMC neuron ablation is a reduction in metabolic rate and not appetite control dysregulation. Furthermore, unlike isolated deletion of the pomc gene, loss of POMC neurons leads to pronounced insulin resistance and diabetes. Finally, the increased anxiety exhibited by these mice possibly indicates that up-regulation of CRF and/or AVP tone mediate the stress response despite the loss of melanocortinergic tone and the lack of HPA axis activation.

Acknowledgments

We thank Mr. S. Ovadia for his devoted assistance with animal care. We also thank Dr. R. Eilam for her assistance with the BAT immunostaining.

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This work was supported by FP7 Grant 260463 from the European Research Council; a research grant from the Israel Science Foundation; a research grant from Roberto and Renata Ruhman; a research grant from the Legacy Heritage Biomedical Science Partnership; a grant from Mr. and Mrs. Mike Kahn; a research grant from Mr. Jorge David Ashkenazi; a research grant from Roberto and Renata Ezzat; a research grant from Mr. and Mrs. Mike Kahn; a research grant from Mrs. and Mr. Barry Wolfe; and a research grant from Nella and Leon Benoziyo Center for Neurosciences.

Disclosure Summary: The authors have nothing to disclose.

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