In addition to its role in reproduction, estradiol-17β is critical to the regulation of energy balance and body weight. Estrogen receptor α-null (Erα−/−) mutant mice develop an obese state characterized by decreased energy expenditure, decreased locomotion, increased adiposity, altered glucose homeostasis, and hyperleptinemia. Such features are reminiscent of the propensity of postmenopausal women to develop obesity and type 2 diabetes. The mechanisms by which ERα signaling maintains normal energy balance, however, have remained unclear. Here we used knockin mice that express mutant ERα that can only signal through the noncanonical pathway to assess the role of nonclassical ERα signaling in energy homeostasis. In these mice, we found that nonclassical ERα signaling restored metabolic parameters dysregulated in Erα−/− mutant mice to normal or near-normal values. The rescue of body weight and metabolic function by nonclassical ERα signaling was mediated by normalization of energy expenditure, including voluntary locomotor activity. These findings indicate that nonclassical ERα signaling mediates major effects of estradiol-17β on energy balance, raising the possibility that selective ERα agonists may be developed to reduce the risks of obesity and metabolic disturbances in postmenopausal women.
Genetic rescue of nonclassical ERα signaling normalizes energy balance in obese Erα-null mutant mice

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In addition to its role in reproduction, estradiol-17β is critical to the regulation of energy balance and body weight. Estrogen receptor α-null (Erα<sup>−/−</sup>) mutant mice develop an obese state characterized by decreased energy expenditure, decreased locomotion, increased adiposity, altered glucose homeostasis, and hyperleptinemia. Such features are reminiscent of the propensity of postmenopausal women to develop obesity and type 2 diabetes. The mechanisms by which ERα signaling maintains normal energy balance, however, have remained unclear. Here we used knockin mice that express mutant ERα that can only signal through the noncanonical pathway to assess the role of nonclassical ERα signaling in energy homeostasis. In these mice, we found that nonclassical ERα signaling restored metabolic parameters dysregulated in Erα<sup>−/−</sup> mutant mice to normal or near-normal values. The rescue of body weight and metabolic function by nonclassical ERα signaling was mediated by normalization of energy expenditure, including voluntary locomotor activity. These findings indicate that nonclassical ERα signaling mediates major effects of estradiol-17β on energy balance, raising the possibility that selective ERα agonists may be developed to reduce the risks of obesity and metabolic disturbances in postmenopausal women.

Introduction

In addition to its critical functions as a reproductive hormone, estradiol-17β (E<sub>2</sub>) plays a vital role in the regulation of energy balance and body weight (1). Estrogen deficiency at menopause is associated with an increased probability of obesity as well as increased risk for the development of type 2 diabetes (2). In experimental animals, reduction of circulating estrogen levels by ovariectomy leads to the development of obesity, which can be reversed or prevented by E<sub>2</sub> treatment (1). The effects of E<sub>2</sub> on energy balance bear many similarities to the actions of leptin and insulin, key molecules involved in energy homeostasis (3, 4). Genetic and pharmacological studies have demonstrated that leptin and insulin act directly on neural networks to modulate energy homeostasis, where the net effect is to decrease food intake and increase energy expenditure (5–9). Both can activate STAT3 in various tissues, and hypothalamic leptin and insulin signaling are known to converge on the PI3K pathway (10–13). Similarly, E<sub>2</sub> is now also known to activate STAT3 as well as PI3K signaling cascades, suggestive of possible cross-talk among these molecules and possibly representing a common neuronal signaling mechanism that may help explain the similarities in their central effects on energy homeostasis (14–17).

That these metabolic actions of E<sub>2</sub> are mediated by estrogen receptor α (ERα) has been demonstrated in Erα<sup>−/−</sup> (Erα<sup>−/−</sup>) mutant mice, in which the ablation of ERα signaling results in a metabolic syndrome characterized by increased body weight, adiposity, altered glucose homeostasis, decreased energy expenditure, hyperinsulinemia, and hyperleptinemia (18–20). However, these metabolic characteristics of Erα<sup>−/−</sup> mutant mice are not evident in Erβ<sup>−/−</sup> mutant mice, which confirms that the Erβ isozyme mediates most of the effects of E<sub>2</sub> on energy balance. Moreover, in vivo ERα gene silencing in the ventromedial nucleus (VMN) of the hypothalamus has revealed this region to be a critical locus of E<sub>2</sub> regulation of energy balance (21, 22). In sum, the effects of E<sub>2</sub> on energy homeostasis appear to require activation of the ERα isoform in hypothalamic neurons.

Although the role of ERα in E<sub>2</sub> regulation of energy homeostasis and protection against high-fat diet–induced (HFD-induced) obesity are well documented, much less is known about the cellular signaling pathways through which ERs mediate their metabolic effects. In the classical ERα signaling pathway, E<sub>2</sub> binds to ERα, inducing conformational changes that permit recruitment of coactivator or corepressor molecules and binding to estrogen response element (ERE) to regulate transcription of target genes. Whereas E<sub>2</sub> acts predominantly through this pathway, emerging evidence supports the importance of nonclassical ERα signaling pathways in mediating estrogen actions. These operate independently of receptor binding directly to ERs and culminate in both genotropic and nongenotropic effects in target cells. Nonclassical genotropic ERα signaling includes protein-protein interactions with other transcription factors, such as activator protein-1 (AP1), specificity protein-1 (SP1), and NF-κB (23–26), which in turn mediate transcriptional regulation at their cognate response elements. Nonclassical ERα signaling also includes mechanisms in which cytoplasmic or membrane-associated receptor activation is coupled to stimulation of cytoplasmic signaling pathways, including the PI3K, MAPK, and cAMP-response element binding protein (CREB) signaling cascades (27–29). The extent to which E<sub>2</sub> regulates energy homeostasis via these nonclassical ERα signaling pathways remains unclear.
In the present study, we used our recently generated gene knock-in mouse, which we believe to be novel, to distinguish the relative contributions of classical versus nonclassical ERα mechanisms in a physiological context. In these mice, an AA mutant allele (E207A/G208A; referred to herein as AA), a mutant form of Erα with disrupted classical (ERE-dependent) but intact nonclassical (ERE-independent) ERα signaling capacities, was introduced onto the Erα−/− mutant background. Initial characterization of the reproductive phenotype in these nonclassical Erα-knockin (Erα−/AA) mutant mice revealed hypoplastic mammary glands, anovulation, and altered steroidogenesis (30). Certain reproductive parameters dysregulated in the Erα−/− mice, however, were restored to normal or near-normal values in the Erα−/AA mice, including E2 feedback inhibition of luteinizing hormone (31).

In this study, we examined the metabolic characteristics of female Erα−/AA mice to test the hypothesis that nonclassical ERα signaling pathways mediate regulation of energy balance by E2. We reasoned that the rescue of normal energy balance, or at least the normalization of any individual metabolic parameters that are dysregulated in Erα−/− mice, would provide in vivo evidence that nonclassical ERα signaling pathways mediate the effects of E2 on energy homeostasis. We additionally assessed whether alterations in energy balance in Erα−/− mice and any rescue of normal energy homeostasis in Erα−/AA mice are associated with corresponding changes in the E2-sensitive signaling pathways.

Results

Body weight, adiposity, and food intake. We measured the body weights of wild-type (Erα+/+), Erα−/−, and Erα−/AA female mice over a 25-week period (Figure 1, A and B). The body weights of Erα−/− females were significantly higher compared with Erα+/+ and Erα−/AA females beginning at week 8 (P < 0.05) and remained so until the end of the study. Using whole-body MRI scan, we measured percent body fat and percent lean mass in adult females and found significantly higher percent body fat in Erα−/− compared with Erα+/+ and Erα−/AA mice (Erα+/+, 35.4% ± 2.2%; Erα−/−, 18.5% ± 2.2%, P < 0.01; Erα−/AA, 24.4% ± 2.7%, P < 0.01; Figure 1C). Percent lean mass was lowest for Erα−/− compared with Erα+/+ and Erα−/AA animals (Erα+/+, 53.3% ± 2.0%; Erα−/−, 68.9% ± 1.9%, P < 0.001; Erα−/AA, 61.5% ± 2.4%, P < 0.05; Figure 1D). The percentages of body fat and lean mass were not significantly different across the 3 genotypes in mice that were 3–6 weeks old (data not shown).

Despite the significantly higher body weight and adiposity in Erα−/− compared with Erα+/+ and Erα−/AA females beginning at week 8, there was no effect of genotype on food intake among the 3 groups (Erα+/+, 11.56 ± 0.38 kcal/d; Erα−/−, 11.23 ± 0.68 kcal/d; Erα−/AA, 10.91 ± 0.51 kcal/d; Figure 1E). To test the possibility that differences in food intake may have been present prior to the observed differences in body weight, daily food intake was measured in mice beginning at weaning and lasting for a period of 3 weeks. As in the older mice, there was no difference in daily food intake among Erα+/+, Erα−/−, and Erα−/AA females between 3 and 6 weeks of age (data not shown). Placing the mice on a 45% HFD also did not result in differences in food intake across the 3 genotypes (Erα+/+, 10.78 ± 0.52 kcal/d; Erα−/−, 11.36 ± 0.64 kcal/d; Erα−/AA, 10.89 ± 0.22 kcal/d; Figure 1F); however, Erα−/− mice gained more weight on HFD than on regular chow diet (RCD) during the same period (P < 0.05; Figure 1G).

Glucose and insulin tolerance tests. Fasting blood glucose concentrations did not differ significantly in Erα+/+, Erα−/−, and Erα−/AA mice. However, blood glucose concentrations in response to glucose challenges in both unanesthetized and anesthetized mice were significantly elevated in Erα−/− mice compared with their Erα−/+ littermates, and this difference persisted at 120 minutes after glucose injection (P < 0.001; Figure 1H, data shown for anesthetized mice). The blood glucose concentrations of Erα−/AA mice were indistinguishable from those of Erα−/− mice at all time points. Blood glucose concentration in response to insulin administration was significantly elevated in Erα−/− mice compared with their Erα−/+ and Erα−/AA littermates at 15 minutes after insulin injection (P < 0.01; Figure 1I).

Akt phosphorylation in response to insulin administration. The effects of insulin on phosphorylated Akt (pAkt) expression in the soleus muscle of Erα+/+, Erα−/−, and Erα−/AA mice were assessed by Western blot analyses. A significant decrease in the pAkt/Akt ratio was observed in Erα−/+ compared with Erα−/− mice within 15 minutes of acute insulin treatment (Erα−/−, 18.4% ± 26%; Erα−/+, 25.9% ± 24%, P < 0.05; Figure 1J), indicating a significant decrease in insulin activation on Erα−/− mice. This effect was partially restored to normal levels in Erα−/AA mice (230% ± 10%). There were no significant differences in the pAkt/Akt ratio in the baseline saline-treated animals or in the total Akt level in any group.

Serum leptin levels. We examined another metabolic phenotype commonly associated with the obese state: hyperleptinemia (32, 33). Our measurements of basal serum leptin concentration after an overnight fast revealed significantly higher serum leptin concentrations in Erα−/− mice compared with Erα+/+ and Erα−/AA mice (Erα−/−, 29.13 ± 1.87 ng/ml; Erα−/+, 8.26 ± 0.06 ng/ml, P < 0.01; Erα−/AA, 14.90 ± 2.37 ng/ml, P < 0.01; Figure 1K).

Locomotor activity and energy expenditure. To determine the level of locomotor activity, running-wheel behavior was measured. Figure 2A shows representative actograms and total activity counts for each genotype. During the light/inactive phase, total activity counts were not significantly different across the 3 genotypes. During the dark/active phase, however, the mean number of revolutions recorded for Erα−/− mice was 55% lower than that for Erα+/+ animals (Erα−/−, 12.3 × 10^6 revolutions/12 hours; Erα+/+, 27.4 × 10^6 revolutions/12 hours, P < 0.05; Figure 2B). In contrast, the Erα−/AA females’ mean wheel-running activity was restored to 90% of wild-type levels (24.1 × 10^6 revolutions/12 hours). Additional measurements of wheel-running activity in young mice (4–6 weeks old), made prior to the onset of significant differences in body weight, revealed similar results: total activity counts were substantially lower in Erα−/− than in Erα+/+ and Erα−/AA mice (data not shown). To assess energy expenditure, the animals were monitored in open-circuit indirect calorimetry cages. Female Erα−/− mice exhibited significantly decreased volume of oxygen consumption (VO2) compared with Erα−/− mice (Erα−/−, 69.76 ± 0.51 ml/min/kg) compared with Erα+/+ mice (Erα+/+, 78.88 ± 0.49 ml/min/kg, P = 0.01; Figure 2C). In contrast, VO2 in Erα−/AA mice did not differ from wild-type levels (76.41 ± 0.83 ml/min/kg). Consistent with results obtained from wheel-running activity in young mice, VO2 was lower in young Erα−/− mice than in young Erα+/+ mice (P < 0.05; Figure 2D).

STAT3 phosphorylation in response to E2 and leptin administration. STAT3 phosphorylation levels were measured in the medial basal hypothalamus (MBH) after estradiol benzoate (EB) or leptin treatment in ovarioctomized Erα−/− mice by Western blot analysis. Whereas leptin rapidly and strongly induced STAT3 phosphorylation, EB treatment was without effect (Figure 3A). The absence of STAT3 activation by EB was also observed in a peroxidase immunocytochemical study like the one described below for leptin (data not shown).
Nonclassical ERα signaling restores body weight and adiposity in Erα−/− mice to wild-type levels. (A) Body weight measurements of Erα+/, Erα−/−, and Erα−AA female mice over a 25-week period beginning at weaning revealed a significant increase in Erα−/− body weight by week 8 (*P < 0.05, **P < 0.01, ***P < 0.001 versus Erα+/, n = 5–10). (B) Appearance of Erα+/, Erα−/−, and Erα−AA mice. (C) NMR imaging revealed higher percent body fat in Erα−/− mice (**P < 0.01, ***P < 0.001; n = 7–11). (D) Percent lean mass was significantly lower in Erα−/− mice (*P < 0.05, ***P < 0.001; n = 7–11). (E) Daily food intake on RCD (P = 0.69; n = 8–10). (F) Daily food intake on HFD (P = 0.67; n = 8–10). (G) Body weight gain during HFD was significantly greater in Erα−/− mice than in Erα+/− and Erα−AA mice (*P < 0.05; n = 8–10). (H) Blood glucose concentrations after glucose challenge in anesthetized mice (**P < 0.001 versus Erα+/, ***P < 0.001 versus Erα−AA; n = 6–10). (I) Blood glucose concentrations after insulin injection (**P < 0.01; n = 6–10). (J) Effect of insulin on pAkt expression in soleus muscle (*P < 0.05; n = 6–8). (K) Serum leptin concentration (**P < 0.01 versus Erα+/, n = 6–10).
STAT3 phosphorylation in response to leptin administration. The effects of leptin on pSTAT3 expression in the arcuate nucleus (ARC) of Erα+, Erα−, and Erα−AA mice were assessed by peroxidase immunohistochemical analyses. Representative photomicrographs of pSTAT3-immunoreactive (pSTAT3-ir) cells in the ARC are provided in Figure 3B. In Erα+ mice, leptin induced a robust increase in STAT3 phosphorylation (84 ± 3 pSTAT3-ir cells/section; Figure 3C). In contrast, induction of STAT3 phosphorylation by leptin decreased to less than 20% of wild-type levels in Erα− mice (16 ± 4 pSTAT3-ir cells/section, P < 0.001). In Erα−AA mice, however, it was restored to near-normal levels (64 ± 9 pSTAT3-ir cells/section). There were no differences in the number of pSTAT3-ir cells in the ARC of the vehicle-treated groups (Erα+, 2.1 ± 1.4 pSTAT3-ir cells/section; Erα−, 3.7 ± 2.2 pSTAT3-ir cells/section; Erα−AA, 4.3 ± 1.7 pSTAT3-ir cells/section).

Additional measurements were made in young lean mice (4–6 weeks old), prior to the onset of body weight differences. There were no differences in the number of pSTAT3 cells following leptin treatments across the 3 genotypes in these younger animals (Erα+, 111 ± 5 pSTAT3-ir cells/section; Erα−, 98 ± 6 pSTAT3-ir cells/section; Erα−AA, 107 ± 12 pSTAT3-ir cells/section; Figure 3D).

Akt phosphorylation in response to E2 administration. The effects of E2 on pAkt expression in the ARC and ventrolateral VMN (vlVMN) of Erα+, Erα−, and Erα−AA mice were assessed by peroxidase immunohistochemical analyses. EB had no effect on pAkt immunoreactivity in the ARC in any group (Figure 4, A and B). In contrast, acute EB treatment induced a significant increase in the number of pAkt-ir cells in the vlVMN of Erα+ mice within 1 hour compared with vehicle treatment (EB, 103.3 ± 5 pAkt-ir cells/section; vehicle, 77 ± 7 pAkt-ir cells/section, P < 0.01; Figure 4, C and D). This effect was completely absent in Erα− mice (EB, 76 ± 5 pAkt-ir cells/section; vehicle, 70 ± 6 pAkt-ir cells/section) and was restored to near-normal levels in Erα−AA mice (EB, 94 ± 5 pAkt-ir cells/section; vehicle, 63 ± 6 pAkt-ir cells/section, P < 0.05).

Discussion
Disruption of ERα signaling in Erα− mutant mice produces an obese metabolic phenotype characterized by decreased energy...
expended, but not necessarily increased energy intake (19, 20). Other features of metabolic syndrome are also evident in the absence of Ertα expression, including increased visceral adiposity, hyperinsulinemia, insulin resistance, and impaired glucose tolerance (19). These metabolic characteristics of ertα–/– mutant mice are not evident in Ebfα–/– mutant mice, and they are largely phenocopied in ovariectomized animals. It is therefore generally held that the Ertα isoform mediates most of the effects of E2 on energy balance. The mechanisms by which ERα signaling normally modulates energy balance, however, have remained unclear. Here, we present evidence for a major role for nonclassical ERα signaling in regulating energy homeostasis. We analyzed the metabolic features of ertα–/AA mice, which express mutant receptors that cannot signal through EREs, but retain the capability to signal through nonclassical ERα signaling pathways. We report that the metabolic parameters dysregulated in female ertα–/– mice — body weight, adiposity, locomotor activity, energy expenditure, blood glucose levels, and serum leptin concentration — were normalized or restored to near-normal values in these ertα–/AA mice. These observations in ertα–/AA mice demonstrate that although classical ERα signaling may play an important role, nonclassical ERα signaling is sufficient to maintain the E2-sensitive physiological regulatory mechanisms that govern energy balance.

Previous studies have suggested that the major effects of ERα signaling on body weight occur primarily as a consequence of increases in energy expenditure, and are less dependent upon modulation of energy intake. A classic finding of Wade et al., for example, demonstrated that ovariectomy induces increases in body weight and adiposity in the absence of long-term alterations in food intake (34). Furthermore, the initial phenotypic analyses of ertα–/– mice revealed that body weights are significantly greater than in ertα+/+ mice beginning at approximately 3 months of age, despite similar food intake levels in the 2 groups (19). In the present study, we similarly found that ertα–/– mice weighed significantly more than their ertα+/+ littermates and exhibited greater adiposity, despite the fact that their energy intake remained similar to that of ertα+/+ mice. Conversely, introduction of the knockin mutation was sufficient to prevent the increase in body weight to levels observed in ertα+/+ mice. Even when placed on a HFD, the body weight increases in ertα+/AA mice remained similar to those of ertα+/+ mice, suggesting the involvement of nonclassical ERα signaling in mediating the protective effects of estrogen against HFD-induced obesity. These findings confirmed the importance of ERα in the regulation of body weight and revealed in vivo that nonclassical ERα signaling is sufficient to mediate these effects through the regulation of energy expenditure.

Although we found no evidence for altered food intake among the genotypes, previous studies have established that E2 can exert some anorectic effects in female rodents (35). Our observations suggest that any such effects may be species or paradigm specific, or, more likely, that they are mediated by signaling pathways activated via ERα-independent mechanisms. As an example, a putative

Figure 3
Leptin, but not E2, activates STAT3. (A) Representative Western blot analysis of pSTAT3 and STAT3 in MBH of ertα+/+ mice 30 and 60 minutes after injection with leptin and EB, respectively. (B) Representative images of leptin-induced pSTAT3 immunoreactivity in the ARC of ertα+/+, ertα–/–, and ertα–/AA mice 30 minutes after vehicle or leptin injection. Scale bar: 100 μm. (C) The number of leptin-induced pSTAT3-ir cells was significantly lower in ertα–/– mice (**P < 0.001 versus ertα+/+ and ertα–/AA; n = 5–8). (D) In 4- to 6-week-old ertα+/+, ertα–/–, and ertα–/AA mice, there were no differences in the number of leptin-induced pSTAT3-ir cells in the ARC (n = 4–6).
membrane estrogen receptor — a Gq-coupled, membrane-initiated pathway — has been implicated in the anorectic effects of E2 in hypoestrogenic female guinea pigs (36–38).

Erα−/– mice also exhibited a metabolic syndrome, including alterations in glucose homeostasis that were normalized in Erα−/AA mice. Upon glucose challenge, Erα−/– mice exhibited a significant impairment of glucose tolerance compared with Erα+/+ animals, but nonclassical ERα signaling restored normal glucose sensitivity in Erα−/AA mice. These glucose tolerance test results were obtained in both anesthetized and awake animals, which indicates that the observed differences are not due to differences in ambient levels of physical activity.

Insulin tolerance tests similarly demonstrated decreased response to insulin in Erα−/– compared with Erα+/+ and Erα−/AA mice, indicative of insulin resistance in the obese mice that was normalized in Erα−/AA mice. In the soleus muscle, insulin’s ability to induce Akt phosphorylation was significantly lower in Erα−/– than in Erα+/+ mice, and partially rescued in Erα−/AA mice. Serum leptin concentrations were also significantly higher in Erα−/– than in Erα+/+ mice and were restored to normal levels in Erα−/AA mice. These findings clearly demonstrated that nonclassical signaling can prevent the development of a metabolic syndrome that occurs in the absence of ERα signaling.

We also sought to determine whether alterations in physical activity and energy expenditure precede — and may therefore be determinants of — the development of obesity and metabolic syndrome in Erα−/– mice. The stimulatory effects of E2 on wheel-running activity have been well documented in mice (20). Our results similarly demonstrate that the null mutation of Erα produced a significant decrease in wheel-running activity, below levels observed in Erα+/+ animals. In contrast, Erα−/AA mice exhibited activity levels comparable to those of Erα+/+ animals, demonstrating that nonclassical ERα signaling is sufficient to restore activity to normal values. Since the reduction in activity in Erα−/– mice and restoration to wild-type

Figure 4
E2 does not induce pAkt immunoreactivity in the ARC, but activates it in the vVMN. (A) Representative image of EB-induced pAkt immunoreactivity in the ARC of Erα+/+, Erα−/–, and Erα−/AA mice 1 hour after vehicle or EB injection. Scale bar: 100 μm. (B) There was no difference in the number of E2-induced pAkt-ir cells in the ARC of Erα+/+, Erα−/–, and Erα−/AA mice. (C) Representative image of E2-induced pAkt immunoreactivity in the vVMN of Erα+/+, Erα−/–, and Erα−/AA mice 1 hour after vehicle or EB injection. Scale bar: 100 μm. (D) E2 significantly increased the number of pAkt-ir cells in the vVMN of Erα+/+ and Erα−/AA mice, but not Erα−/– mice (*P < 0.05, **P < 0.01 versus vehicle; n = 5–8).
levels in Erα–/–A mice could have had secondary to differences in body weight, wheel-running activity was measured in mice prior to the onset of significant body weight differences. The wheel-running activity in young Erα–/– mice was again significantly lower than that of young Erα–/– or Erα–/–A mice, which suggests that alterations in voluntary locomotor activity precede the development of obesity. Furthermore, measurements of energy expenditure revealed differences among the genotypes that paralleled those of locomotor activity. VO2 was significantly lower in Erα–/– than in Erα–/–A mice, whereas Erα–/–A mice exhibited VO2 levels comparable to those of Erα–/–A animals in both adult mice and young lean mice. Therefore, reduced energy expenditure in Erα–/– mice compared with Erα–/– and Erα–/–A mice, despite equivalent energy intake, appears to be a critical mechanism by which obesity develops in the Erα–/– animals.

The normalization of energy balance in Erα–/–A mice may be caused by restoration of one or more nonclassical ERα signaling pathways, many of which may operate in mediobasal hypothalamic neuronal circuits known to govern energy homeostasis. Nonclassical, genotypic responses may be elicited by tethering of ERα to other DNA-bound transcription factors, such as API, SPI, and NF-kB, to regulate transcription at their cognate response elements (23, 25, 26). An alternative possibility is that the actions of E2 may initially be associated with common signaling pathways via the nonclassical, nongenotropic mechanism in the hypothalamus; one recent example is activation of STAT3 by E2 (15). In our study, however, E2 did not phosphorylate STAT3 in the MBH of Erα–/– mice, whereas leptin rapidly induced STAT3 phosphorylation. Further examination of leptin-induced STAT3 signaling in Erα–/–, Erα–/–A, and Erα–/–A mice demonstrated activation of STAT3 by leptin in the ARC of Erα–/– and Erα–/–A mice, but not Erα–/–A mice. Our findings suggest that nonclassical ERα signaling may confer STAT3 responsiveness to leptin in ARC neurons, but the mechanisms mediating this effect do not seem to involve direct activation of STAT3 by E2. Rather, leptin’s ability to activate STAT3 may be secondarily influenced by ERα-mediated alterations in metabolic status. This was confirmed by examining STAT3 phosphorylation in lean young mice, which revealed no differences in STAT3 responsiveness to leptin across genotypes.

Nonclassical, nongenotropic responses may also be initially mediated by ERα that is coupled to stimulation of MAPK, CREB-, and PI3K-mediated pathways, the latter being a particularly intriguing possibility (27, 39) because PI3K functions as a common intracellular signaling node that mediates the neural actions of metabolic regulators such as insulin and leptin (40–42). Additionally, E2 can activate PI3K through protein-protein interactions between ERα and p85 (a regulatory subunit of PI3K), which in turn activate Akt in a variety of cell types, including hypothalamic neurons, a process that is completely inhibited by the PI3K inhibitors wortmannin and LY294002 (43, 44). We therefore tested whether E2 activates Akt via ERE-independent ERα signaling in Erα–/–, Erα–/–A, and Erα–/–A mice by immunocytochemistry for pAkt in the ARC and vVMN of the hypothalamus. The ARC was a region of particular interest because of its extensive association with leptin and insulin signaling. More specifically, both leptin and insulin activated Akt in the ARC, and PI3K activity in the ARC has been linked to the regulation of energy homeostasis (10). If the effects of E2 on energy homeostasis were achieved in part by activating Akt in the ARC, we would expect increased pAkt immunoreactivity by EB in the ARC. However, our findings did not support this hypothesis: EB did not significantly increase induction of pAkt in the ARC of Erα–/–, Erα–/–A, or Erα–/–A mice.

The VMN was another region of interest for the activation of Akt, as it is a critical center of energy homeostasis. Animals with lesions to the VMN exhibit hyperphagia, increased body weight, increased adiposity, and reduced energy expenditure (45, 46). Furthermore, when VMN-lesioned animals are pair-fed with unlesioned animals, the lesioned animals exhibit significantly reduced energy expenditure prior to the onset of body weight differences, which suggests that the VMN is an important regulator of energy use. The VMN (more specifically, the vVMN) is also a nucleus with very high density of ERα (47, 48), and focused silencing of ERα in this nucleus generates a metabolic state similar to that in Erα–/– mutant mice, which suggests that ERα in the VMN plays an essential role in control of energy balance and maintenance of normal body weight (22). We report here that E2 significantly induced activation of Akt in Erα–/– mice, but not Erα–/–A mice, in the vVMN, confirming the requirement of ERα for E2 induction of pAkt. Furthermore, Akt activation by E2 was restored in Erα–/–A mice, which suggests that ERE-independent ERα signaling is sufficient to evoke rapid activation of Akt. This finding demonstrates that E2 can evoke nonclassical ERα signaling in VMN neurons and suggests at least one potential route through which nonclassical ERα signaling may convey some of E2 effects on energy homeostasis in the VMN.

In conclusion, our findings in the Erα–/–A genetic model established that nonclassical ERα signaling mechanisms are sufficient to transduce virtually all major effects of E2 on energy homeostasis in female mice. These findings raise the possibility that nonclassical ERα signaling pathways may be identifiable as new drug targets for the treatment of metabolic disorders in postmenopausal women. Different selective estrogen receptor modulators can possess agonist, antagonist, or mixed pharmacological properties that are specific to tissues, cells, target genes, and signaling pathways (49–51). Thus, treatments that selectively activate nonclassical ERα signaling pathways might have reduced oncogenic potential associated with classical signaling mechanisms (52, 53), yet retain agonistic properties that could lead to reduced risk of obesity and metabolic disturbances in women.

**Methods**

**Animals.** All animal care and use procedures were in accordance with guidelines of the Animal Care and Use Committee of Northwestern University. Mice were maintained on a 12-hour light/12-hour dark cycle and given RCD or HFD (45% kcal from fat; Research Diets Inc.) with water available ad libitum. In the HFD study, animals were maintained on a HFD for 4 weeks. Erα–/– and Erα–/–A mutant mice were generated as previously described (30, 54). All breeders were derived from previous backcrosses for 8–13 generations onto the C57BL/6 background. Compound heterozygotes (Erα–/–A) were generated by mating heterozygote Erα–/–A males with heterozygote Erα–/– females. Erα–/– mice were generated by mating Erα–/–A males with Erα–/– females. All mice were genotyped at weaning. DNA was isolated by digestion of tail tissue and amplified in 2 separate PCRs to determine the presence or absence of the wild-type Erα and the presence or absence of the knockin mutation.

**Body weight and food intake.** Measurements of body weights were taken at 5 time points: 3–4, 6–7, 8–10, 15–18, and 22–25 weeks after birth. Food intake on either RCD or HFD was measured daily at specified ages.

**Body fat and lean mass determination.** Body fat and lean mass were analyzed with a mouse-specific NMR apparatus (Echo MRI Whole Body Composition Analyzer; Echo Medical Systems).

**Locomotor activity.** Wheel-running activity was recorded for 21 consecutive days. Mice were individually housed in cages equipped with a 12.5-cm-diameter exercise wheel. Each wheel revolution triggered a microswitch mounted
on the outside of the cage near the axle of the wheel. Data were collected using Chronobiology Kit (Stanford Software Systems), and Fast Fourier Transform Analysis was used to analyze activity data (Clocklab Actimetrix).

Energy expenditure. Indirect calorimetry was performed on individually housed mice in polycarbonate calorimeter cages. Animals were placed in cages for a 24-hour acclimation period prior to calorimetric measurements. Indirect calorimetry was performed using a computer-controlled Physioscan open-circuit calorimetry system (Accuscan Instruments). VO₂ was determined as milliliters per minute per kilogram body weight (ml/min/kg⁰.⁷⁵). Calorimetric measurements were measured for each mouse at 10-minute intervals over a 24-hour period, beginning 4 hours prior to the onset of the dark phase.

Glucose and insulin tolerance tests. Animals were fasted for 16 and 4 hours before i.p. injection of 2 g/kg glucose and 0.5 U/kg insulin, respectively. Blood samples were collected from the tail vein at 0, 30, 60, and 120 minutes after glucose or insulin administration. Blood glucose concentration was measured using a blood glucose monitoring system (OneTouch; Johnson & Johnson). Glucose tolerance test was administered in both awake animals and in animals anesthetized with sodium pentobarbital (50 mg/kg fast).

Effects of insulin-Akt signaling in soleus muscle. Animals were given i.p. injection of human insulin (0.15 U/g; Eli Lilly) after a 4-hour fast. 15 minutes after insulin injection, soleus muscles were removed and flash frozen. Muscle samples were processed as previously described (55), with minor modifications. Protein content was determined by DC protein assay (Bio-Rad Laboratories). Soleus muscle lysates (50 μg protein) were resolved using SDS-PAGE on 7.5% gels and immunoblotted with antibodies specific to protein kinase B (Akt) and pAkt (ser473) (Cell Signaling Technology) and appropriate secondary antibody (goat anti-rabbit horseradish peroxidase conjugates; Cell Signaling). Ertc⁺/−, Ertc−/−, and Ertc−/−/− samples were run together on the same gels, and signals were detected by chemiluminescence (ECL Western Blotting Detection Reagent; Amersham; refs. 55, 56). Immunoblots were quantified using a high-resolution scanner, and density was calculated using NIH ImageJ software. The same internal standard was loaded on all immunoblots, and results were expressed as percent of internal standard. Phosphorylation was expressed as the ratio of phosphorylated to total protein (55, 56).

Hormone assay. Serum leptin levels were determined using RIA reagents obtained from Linco Research. The leptin assay had a lower limit of detection of 0.25 ng/ml, and the intraassay and interassay coefficients of variation were 3.93% and 8.29%, respectively.

Effects of EB and leptin on STAT3 phosphorylation. On the morning of day 7 after ovariectomy, animals were given a subcutaneous injection of EB (1 μg/mouse). 1 hour after injection, animals were anesthetized with i.p. injection of 75 mg/kg ketamine (Fort Dodge Laboratories) and 5 mg/kg xylazine (Burns Veterinary Supply). As control for STAT3 activity, leptin (5 mg/kg; R&D Systems) was administered on day 7 after an overnight fast. The MBH was dissected and homogenized in lysis buffer (20 mM Tris-HCl, pH 7.4; 150 mM NaCl; 1% Triton X-100; 0.5 mM EDTA; and proteinase inhibitors). Western blot analyses were performed according to standard protocols using antibodies against pSTAT3 and STAT3 (Cell Signaling Technology).

Effect of leptin on STAT3 phosphorylation. On day 0, mice were anesthetized and bilaterally ovariectomized. On day 6, food was removed before the start of the dark phase, and on the morning of day 7, mice were given i.p. injection of vehicle or leptin (5 mg/kg; R&D Systems). 30 minutes after injection, animals were anesthetized with 75 mg/kg i.p. ketamine and 5 mg/kg i.p. xylazine and transcardially perfused with 4% paraformaldehyde (Sigma-Aldrich). Brains were removed, postfixed in the perfusion solution for 12 hours, and then placed in 30% sucrose in 0.1 M phosphate buffer until they sank. Coronal brain sections (30 μm) were cut in a CM1900 cryostat (Leica) for subsequent immunostaining. Immunocytochemical procedures were performed as previously described (57). Briefly, sections were rinsed in 0.1 M PBS for 30 minutes and pretreated with 3% H₂O₂ in methanol for 40 minutes. After a minimum of 3 rinses in PBS, the sections were immersed in PBS with 0.3% glycine for 25 minutes. After additional rinses, sections were placed in 0.03% sodium dodecyl sulfate for 25 minutes. All sections were rinsed again and placed in 4% normal serum, 0.4% Triton X-100, and 1% BSA for 30 minutes before incubation with pSTAT3 mouse mAb (Cell Signaling) for 72 hours at 4 °C in 1% normal serum, 0.4% Triton X-100, and 1% BSA. Sections were then rinsed in PBS and incubated in biotinylated goat anti-mouse IgG diluted 1:500 for 1 hour at room temperature, followed by incubation in avidin-biotin complex (Vectastain ABC; Vector Laboratories) in PBS plus 0.4% Triton X-100 for 1 hour at room temperature. Finally, sections were incubated in 3,3′-diaminobenzidine (DAB; Vector Laboratories) for brown precipitate development at room temperature. Sections were visualized on a Nikon Eclipse TE2000-S microscope (Nikon). To define the area of interest, the boundary of the ARC was drawn using a template derived from the Paxinos and Franklin atlas of the mouse (58), creating an enclosed area. Matched sections corresponding to anterior-posterior coordinates were selected from each brain for pSTAT3 immunocytochemical analyses. The number of pSTAT3-ir cells in the ARC was counted manually using NIH ImageJ software (version 1.63). 3 sections from ARC were selected per mouse. For each mouse, the average number of cells counted in the ARC was used for statistical comparisons.

EB treatment and immunocytochemistry for pAkt. 7 days after ovariectomy, mice were given subcutaneous injection of vehicle or EB (1 μg/mouse). After 1 hour, animals were anesthetized and perfused, and their brains were fixed. Sliced sections were pretreated with 3% H₂O₂ in PBS for 20 minutes before immersion in 0.03% sodium dodecyl sulfate in PBS for 10 minutes. After incubation in 4% formaldehyde serum and 0.4% Triton X-100 in PBS for 2 hours, sections were incubated in mouse anti-pAkt (ser473; 1:500 dilution, Cell Signaling) in 4% formaldehyde serum and 0.4% Triton X-100 for 72 hours at 4 °C. After 72 hours, sections were processed as described previously (59). Incubation without the primary antibody resulted in the absence of detectable signal. Immunocytochemical analyses were performed as described above for the pSTAT3 procedure, except that the protein of interest was pAkt and the regions of interest were the ARC and vVMN.

Statistics. Statistical significance among groups was tested with 1-way ANOVA followed by Bonferroni post-hoc test. 2-way ANOVA followed by Bonferroni post-hoc test was used to test for significance in immunocytochemical studies. Repeated-measures ANOVA was used to analyze body weights and glucose levels of the 3 genotypes across stated time points. Data are presented as mean ± SEM. In all analyses, a P value less than 0.05 was considered statistically significant.

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