RESEARCH COMMUNICATION

Neuronal SIRT1 regulates endocrine and behavioral responses to calorie restriction

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Mammalian life span can be extended by both calorie restriction (CR) and mutations that diminish somatotrophic signaling. Sirt1 is a mediator of many effects of CR in mammals, but any role in controlling somatotrophic signaling has not been shown. Since the somatotrophic axis is controlled by the brain, we created mice lacking Sirt1 specifically in the brain and examined the impacts of this manipulation on somatotrophic signaling and the CR response. These mutant mice displayed defects in somatotrophic signaling when fed ad libitum, and defects in the endocrine and behavioral responses to CR. We conclude that Sirt1 in the brain is a link between somatotrophic signaling and CR in mammals.

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Mammalian life span can be extended by both dietary and genetic interventions. Calorie restriction (CR), a reduction of food intake while maintaining adequate nutrition, dramatically extends the life span in both invertebrates and mammals (for review, see Bordone and Guarente 2005). In multiple species, the beneficial effects of CR have been linked to the function of the sirtuin family of NAD⁺-dependent protein deacetylases (Sirt1 in mouse). In yeast, fruit flies, and possibly mice, genetic ablation of Sirt1 orthologs blocks the extension of life span by CR (Lin et al. 2000; Rogina and Helfand 2004; Li et al. 2008). Because sirtuins require NAD⁺ as an essential cofactor for their deacetylase activity, they are thought to read the metabolic state of an organism and direct changes in transcription and metabolism in response (Imai et al. 2000). Sirt1 regulates multiple metabolic pathways in mammals, and is thought to play a critical role in directing protective changes in response to alterations in nutritional status (Guarente 2009).

Genetic and physiological analysis of Sirt1 in mice supports the model that this sirtuin mediates the beneficial effects of CR. Sirt1 protein levels increase in many, but not all, mammalian tissues during CR (Cohen et al. 2004; Chen et al. 2008). Furthermore, whole-body Sirt1

knockout mice show neither the increase in physical activity nor many of the physiological changes normally associated with CR (Chen et al. 2005; Boily et al. 2008; Li et al. 2008). Conversely, transgenic mice that overexpress Sirt1 show metabolic phenotypes that partially overlap those of CR (Bordone et al. 2007; Banks et al. 2008; Pfluger et al. 2008). Finally, small molecules that increase the enzymatic activity of Sirt1 induce transcriptional changes that significantly overlap changes induced by CR (Barger et al. 2008a,b; Pearson et al. 2008).

In addition to CR, mouse life span can be extended by single-gene mutations. Collectively, these mutations point to the importance of the somatotrophic axis as a major regulator of mouse life span. Long-lived Ames and Snell dwarf mice harbor mutations that disrupt pituitary development, thereby reducing levels of growth hormone (GH) as well as several other pituitary hormones (for review, see Tatar et al. 2003). These mutations also diminish levels of insulin-like growth factor 1 (IGF-1), a downstream mediator of many of the effects of GH whose role in the regulation of life span is less certain (Holzenberger et al. 2003; Ladiges et al. 2009). It has long been known that CR down-regulates somatotrophic signaling in mice, as indicated by a reduction in circulating IGF-1 (Al-Regaiey et al. 2005), suggesting that CR and mutations that block somatotrophic signaling may extend life span in part through shared mechanisms. Indeed, mice in which GH signaling is blocked by ablation of the GH receptor receive no further extension of overall longevity when placed on a CR diet (Bonkowski et al. 2006). The mechanisms by which CR might direct changes in the somatotrophic axis, however, have remained unclear.

We wished to test the idea that Sirt1 might mediate some of its effects on CR by controlling the somatotrophic axis. Secretion of GH from the pituitary is directed by the hypothalamus, a region of the brain that is anatomically adjacent to the pituitary. We therefore used Cre/loxP technology to specifically ablate Sirt1 in the brain, and examined the effects of this manipulation on somatotrophic signaling in mice under normal and CR conditions. We found that these mutant mice are viable and have grossly normal brains, but show specific defects in the somatotrophic axis. Moreover, Sirt1 activity in the brain is required to mediate changes in somatotrophic signaling and physical activity that occur in response to CR. This study establishes Sirt1 in the brain as the critical link between the GH and CR longevity pathways in mammals.

Results and Discussion

Generation of brain-specific Sirt1 knockout (BSKO) mice

We crossed a conditional allele of Sirt1 (Sirt1^{lox}) (Cheng et al. 2003) that had been backcrossed onto the C57Bl/6 genetic background (Chen et al. 2008) to a transgenic mouse strain in which Cre recombinase expression is driven by the Nestin promoter, also on the C57Bl/6 background (Tronche et al. 1999). Nestin-cre was always passed through the male germline. The nestin promoter drives Cre expression only in the cells that give rise to neurons, astrocytes, and glia. Sirt1^{lox/lox}; Nestin-Cre (BSKO) mice and Sirt1^{lox/lox} littermate controls

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(referred to for simplicity as wild-type [WT]) were born at Mendelian ratios. BSKO mice were fertile and displayed grossly normal brain morphology (Fig. 1A). For all experiments, male mice were analyzed except where otherwise noted, but similar phenotypes were observed in both genders (data not shown).

The Sirt1^{flox} allele has loxP sites flanking exon 4 of the Sirt1 gene, which encodes the catalytic domain of the protein. Previous studies of these mice demonstrated that the excised version of this allele produces a stable (but nonfunctional) protein product that can be distinguished from wild-type Sirt1 protein by size (Cheng et al. 2003). To confirm excision of the Sirt1^{flox} allele in BSKO mice, we analyzed brain homogenates from these animals and littermate controls by Western blotting. Sirt1 has two distinct protein isoforms in the brain (Fig. 1B, closed arrowheads). BSKO brains contained no wild-type Sirt1 protein, and only a small amount of the mutant protein (Fig. 1B, open arrowheads). A similar pattern was observed in lysates prepared from the hypothalamus of these animals (Supplemental Fig. S1). Importantly, in the pituitary, the majority of which arises from a non-neuronal lineage, we detected only wild-type Sirt1 in BSKO mice (Fig. 1C). Likewise, only wild-type Sirt1 protein was detected in peripheral tissues such as liver, muscle, and pancreas (Supplemental Fig. S1). Thus, Nestin-Cre expression was confined to the neuronal lineage in our mice.

BSKO mice are dwarfed and have reduced somatotrophic signaling

BSKO mice and their wild-type littermates were indistinguishable at birth and for their first month of life. However, beginning at 4–5 wk of age, we observed that BSKO mice were significantly smaller than their littermates, and that this size difference was maintained into adulthood

(Fig. 1D). This reduction in body mass was accompanied by a reduction in body length (Fig. 1E), and could not be attributed to a decrease in food consumption (Supplemental Fig. S1). Because somatotrophic signaling is critical for post-weaning growth and for directing elongation of the long bones of the animal, we hypothesized that the dwarf phenotype of BSKO mice might arise from defects in the somatotrophic axis.

To examine somatotrophic signaling directly, we first measured serum GH levels in 4-wk-old mice, where GH levels are highest (Diaz-Torga et al. 2002). Mean circulating GH levels in BSKO mice were nearly 50% lower than in wild-type littermate controls (Fig. 2A). We also measured IGF-1, which is released into the circulation predominantly by the liver in response to GH signaling. Serum IGF-1 levels, as well as IGF-1 mRNA levels in the liver, were significantly lower in adult BSKO as compared with wild-type animals (Fig. 2B,C). Finally, we examined transcript levels of genes that are expressed in a sexually dimorphic pattern in mouse liver in direct response to the different patterns of GH release in the two sexes (Waxman and O'Connor 2006). The expression pattern of these genes was significantly disrupted in BSKO mice of both sexes, providing additional evidence for alterations in GH secretion in adult BSKO mice (Supplemental Fig. S2).

Because GH is synthesized, stored, and released by the pituitary, we next examined this gland in BSKO mice. Pituitaries from BSKO animals were smaller than those of wild-type littermates, but displayed no major defects upon histological examination (Fig. 2D; Supplemental Fig. S3). The mass of the pituitaries of BSKO animals was reduced in comparison with wild-type littermates, even when the smaller body mass of BSKO mice was taken into account (Fig. 2E). Furthermore, each pituitary gland contained significantly less stored GH in BSKO mice as compared with controls, although the density of GH-positive cells was comparable between mice of both genotypes (Fig. 2F; Supplemental Fig. S3). To determine whether the defect in GH reflected a more general deficit in pituitary hormone production, we measured serum concentrations of three additional pituitary hormones—adrenocorticotropic hormone (ACTH), prolactin (PRL), and thyroid-stimulating hormone (TSH)—and two hormones produced by the thyroid in response to TSH (T3 and T4) (Fig. 2G). PRL concentrations were somewhat increased in BSKO mice as compared with wild type, while concentrations of ACTH, TSH, T3, and T4 were unchanged. Thus, the pituitary deficit of BSKO mice appears to be restricted to GH.

Together, these findings imply that Sirt1 acts in the brain to stimulate production and secretion of GH but not other pituitary hormones. Because development, proliferation, and function of the pituitary are under the control of the hypothalamus (for review, see Zhu et al. 2007), we hypothesize that the pituitary defects observed in BSKO mice are due to alterations in hypothalamic function in the absence of Sirt1. The production and release of GH is controlled by the opposing action of two hypothalamic neuropeptides: GH-releasing hormone and somatostatin (Low et al. 2001; Alba and Salvatori 2004). The expression of the mRNA encoding these neuropeptides is unaffected in BSKO mice (data not shown), and thus the mechanism by which hypothalamic Sirt1 acts to control the somatotrophic axis is not certain.

Figure 1. BSKO mice lack functional Sirt1 protein in the brain and are dwarfed. (A) Brains from wild-type (WT) and BSKO mice exhibit similar gross morphology. (B) Western blotting of Sirt1 protein in whole-brain lysates from wild-type and BSKO mice. Closed arrowheads indicate wild-type Sirt1 isoforms, and open arrowheads indicate mutant isoforms. Tubulin is shown as a loading control. (C) Western blotting of Sirt1 protein in pituitary lysates. (D) Body mass of male wild-type and BSKO animals at the indicated ages; $n = 6$ –10 per genotype and time point. Curves are significantly different by repeated-measures ANOVA; $P < 0.01$. (E) Snout–anus body length of male wild-type and BSKO animals at 10 wk of age; $n = 6$ –10 per genotype. All data are shown as mean $\pm$ SEM. (*) $P < 0.05$ by two-tailed Student's t -test.

Figure 2. The somatotrophic axis is disrupted in BSKO mice. (A) Serum GH concentrations in 4-wk-old BSKO mice and littermate controls; $n = 15$ –20 per group. (B) Serum IGF-1 concentrations in 10-wk-old BSKO mice and controls; $n = 6$ –10 per group. (C) IGF-1 mRNA levels in BSKO liver normalized to wild-type (WT) levels; $n = 5$ per group. (D) Pituitaries from BSKO and wild-type littermates. (E) Pituitary mass normalized to body mass of 10-wk-old BSKO and wild-type mice; $n = 6$ –10 per group. (F) Pituitary GH stores in 10-wk-old BSKO mice; $n = 5$ –6 per group. (G) Measurements of additional pituitary hormones in BSKO and wild-type mice. For PRL, TSH, and T4, $n = 10$ –14 females per group. For T3 (females) and ACTH (males), $n = 6$ –10 per group. All data are shown as mean $\pm$ SEM. (*) $P < 0.05$ by two-tailed Student's t -test.

Young BSKO mice display near-normal glucose metabolism

Many mouse strains with decreased GH/IGF-1 signaling display improvements in baseline insulin sensitivity. To investigate whether this might also be true of BSKO mice, we measured fasted glucose and insulin in young adult BSKO animals and littermate controls. Young BSKO animals displayed a slight reduction in fasting blood glucose levels and a trend toward a reduction in fasted insulin levels that did not achieve statistical significance (Supplemental Fig. S4). We then assessed the performance of wild-type and BSKO mice in insulin and glucose tolerance tests (ITT and GTT, respectively). At 3 mo of age, BSKO and wild-type mice performed similarly on ITT (Fig. 3A). Since BSKO mice started with lower fasting glucose levels, the levels of blood glucose in these animals remained lower than that of wild type throughout the test (curves are significantly different by two-way ANOVA, $P < 0.05$). Young wild-type and BSKO mice performed identically on the GTT (Fig. 3B). Thus, only minor differences in glucose tolerance and insulin sensitivity were detected in young BSKO mice.

We wondered whether these small differences in glucose metabolism could lead to protection against the metabolic changes brought on by a high-fat diet. To test this, we fed BSKO mice and littermate controls either a high-fat diet or matched control diet beginning at 6 wk of age and continuing for 18 wk. All groups of mice ate a similar amount of food per gram of body weight (data

not shown) and gained weight equally on the high-fat diet (Supplemental Fig. S5). By ITT, high-fat-fed BSKO mice appeared to remain somewhat more insulin-sensitive than wild-type mice fed the same diet, but similar protection was not observed by GTT (Supplemental Fig. S5). Thus, ablation of *Sirt1* in the brain did not provide significant protection against the deleterious effects of a high-fat diet.

Old BSKO mice display glucose intolerance and altered CR responses

Next, we asked whether BSKO animals would display the improvements in insulin sensitivity and glucose tolerance normally found in animals maintained on a CR diet. To test this, we placed BSKO mice and wild-type littermates on a CR diet, with CR mice being fed daily 60% by mass of the food consumed by ad libitum-fed controls. Animals of both genotypes lost weight similarly on CR (Supplemental Fig. S6). After 7 mo of CR, when all of the mice in this study were 10 mo old, we repeated the ITT

Figure 3. BSKO mice have altered glucose homeostasis with aging. (A) ITT on 3-mo-old BSKO and wild-type (WT) mice; $n = 10$ –14 per group. Curves are significantly different by two-way ANOVA, $P < 0.05$. (B) GTT on 3-mo-old BSKO and wild-type mice; $n = 10$ –14 per group. (C) ITT on 10-mo-old mice of indicated diet and genotype groups; $n = 6$ –8 per group. (AL) Ad libitum fed; (CR) calorie restricted. WT-CR and BSKO-CR curves are significantly different by two-way ANOVA, $P < 0.05$. (D) GTT on 10-mo-old mice of indicated diet and genotype groups; $n = 6$ –8 per group. All data are shown as mean $\pm$ SEM. (E) Blood glucose and insulin levels in 1-yr-old wild-type and BSKO mice 45 min after IP injection of 2 g/kg glucose in PBS; $n = 6$ –7 per group. Insulin values are significantly different; $P < 0.01$. The P -value for blood glucose comparison is 0.0727 by a two-tailed Student's t -test. (F) PEPCK, G6pase, and Pcx mRNA expression in liver of 0.5- to 1-yr-old female wild-type and BSKO mice 45 min following vehicle (PBS) or 2 g/kg glucose (GLU) injection; $n = 7$ –14 per group. (*) $P < 0.05$ by two-tailed Student's t -test versus WT-PBS for each gene.

and GTT experiments to determine the impact of CR on glucose homeostasis in BSKO animals. As in young animals, older ad libitum-fed wild-type and BSKO mice performed similarly on the ITT (Fig. 3C). As expected, CR dramatically increased the insulin sensitivity of wild-type animals. However, CR induced only a modest improvement in insulin sensitivity in BSKO animals, suggesting that neuronal Sirt1 is required to achieve the full benefit on insulin sensitivity that is normally conferred by CR. In contrast, the GTT revealed that BSKO and wild-type animals received a significant and equivalent benefit in glucose tolerance when maintained on a CR diet (Fig. 3D). This test also allowed us to compare older ad libitum animals to the young animals studied in Figure 3, A and B. Surprisingly, older BSKO animals developed severe glucose intolerance that was not observed in young mice.

We first hypothesized that the glucose intolerance of BSKO mice might arise from a failure of the pancreatic islets to produce sufficient insulin during the GTT. However, we found that BSKO mice of this age actually produce more insulin in response to glucose injection than wild type (Fig. 3E). We next reasoned that the glucose intolerance phenotype was unlikely to be due to a cell-autonomous effect on key peripheral tissues such as liver or muscle, since Sirt1 is ablated only in the neuronal lineages in BSKO animals (Supplemental Fig. S1). However, the ability of the liver to properly down-regulate gluconeogenesis in response to increasing insulin levels (as occur during a GTT) is in part under central control (for review, see Schwartz and Porte 2005). Insulin-responsive neurons in the hypothalamus direct the down-regulation of gluconeogenesis in the liver via the vagus nerve, which innervates the liver and other visceral organs. Mice in which the vagus nerve has been severed do not down-regulate gluconeogenic genes in response to insulin (Pocai et al. 2005). To determine if this brain–liver circuit was impaired in BSKO mice, we injected older adult (0.5- to 1-yr-old) wild-type and BSKO mice with a bolus of glucose, and after 45 min monitored expression of the mRNA encoding the gluconeogenic enzymes *Pepck*, *G6pase*, and *Pcx* as markers for gluconeogenesis in liver. In all cases, the genes were significantly down-regulated in wild-type mice in response to glucose in-

jection, whereas their expression was unchanged in BSKO mice (Fig. 3F). This finding supports the idea that the glucose intolerance of older BSKO mice arises from a breakdown in the brain–liver circuit that regulates glucose homeostasis.

The fact that CR rescues the glucose intolerance of older BSKO animals may be explained in two ways. First, it is possible that CR blocks the degeneration of the hypothalamic–liver circuit through a Sirt1-independent mechanism. Second, CR dramatically increases the insulin sensitivity of skeletal muscle, so it is possible that the muscles of BSKO mice on CR are able to take up the extra glucose present in the blood and thus mask the inability of the liver to halt gluconeogenesis. Breakdown of the brain–liver circuit may also help to explain our finding that BSKO mice apparently do not receive a full benefit in insulin sensitivity on CR, as measured by ITT; BSKO mice on CR may appear less insulin-sensitive because they are unable to turn off gluconeogenesis in response to an injection of insulin, and thus maintain higher blood glucose values throughout the test. It is also possible that Sirt1 plays some additional role in the central control of peripheral insulin sensitivity, since it is known from other genetic studies that the brain can, in fact, control insulin sensitivity through other pathways (Kahn et al. 2005).

Neuronal Sirt1 regulates somatotrophic signaling on CR

A reduction in somatotrophic signaling is one of the endocrine hallmarks of CR. Because our initial results demonstrated that neuronal Sirt1 regulates somatotrophic signaling, we hypothesized that Sirt1 activity in the brain might be required to direct down-regulation of this axis in response to CR. After wild-type and BSKO mice had been subjected to 10 wk of CR, we measured serum IGF-1 levels as a marker for somatotrophic signaling in ad libitum and restricted animals of both genotypes. As expected, serum IGF-1 levels were significantly lower in wild-type mice on CR than in wild-type ad libitum controls. In contrast, IGF-1 levels remained unchanged in BSKO animals on CR as compared with ad libitum (Fig. 4A). Thus, Sirt1 activity in the brain is absolutely

Figure 4. BSKO mice have altered endocrine and behavioral responses to CR. (A) Serum IGF-1 concentrations after 10 wk of CR; $n = 6$ –8 per group. Interaction between diet and genotype is significant by two-way ANOVA; $P < 0.05$. Data are shown as mean $\pm$ SEM. (B–E) Each bar represents the average number of revolutions run per 15-min period over 15 d for mice of the indicated genotype and diet; $n = 5$ per group.

required to regulate the somatotrophic axis in response to diet.

Since *Sirt1* appears to be a positive regulator of GH production and signaling, our findings also suggest that *Sirt1* activity in the hypothalamus might decrease during CR. Because no changes in *Sirt1* protein levels were detected in the hypothalamus or pituitary in mice fed a CR diet (Supplemental Fig. S7), it is likely that other factors such as changes in the NAD/NADH ratio act to decrease *Sirt1* activity in the hypothalamus during CR. This is in contrast to other tissues, such as skeletal muscle and white adipose tissue, where *Sirt1* activity has been reported to increase in response to CR. However, previous findings indicate that *Sirt1* activity in the murine liver decreases during CR (Chen et al. 2008). Thus, the liver and hypothalamus may be members of a subset of tissues in which it is a decrease, rather than an increase, in *Sirt1* function that mediates physiological changes during CR.

Neuronal Sirt1 regulates physical activity on ad libitum and CR diets

A well-established behavioral consequence of CR in mice is up-regulation of physical activity, presumably a correlate of increased foraging behavior. Previous work using animals lacking *Sirt1* throughout the body demonstrated that *Sirt1* was required for this up-regulation of activity in response to CR (Chen et al. 2005). However, this study left open the question of whether *Sirt1* activity is required in the nervous system or the periphery to generate this behavioral change. To test the requirement for *Sirt1* in the brain in the behavioral response to CR, we used running wheels to monitor physical activity in ad libitum and restricted animals of both genotypes for a 2-wk period beginning after 7 mo of CR. As has been shown previously, wild-type animals dramatically up-regulated their activity in response to CR (Fig. 4B,C). BSKO animals were somewhat more active than wild-type animals on an ad libitum diet (Fig. 4D), and their activity did not increase—and, in fact, slightly decreased—on the CR diet (Fig. 4E). We conclude that *Sirt1* activity is required specifically in the brain to regulate physical activity in ad libitum mice and to drive the changes in activity brought on by CR.

Conclusions

This study brings together three major pathways known to affect aging—CR, the somatotrophic axis, and sirtuins—in what is likely a single mechanism operating in the brain. Our findings suggest that CR triggers a reduction in *Sirt1* activity in hypothalamic neurons governing somatotrophic signaling to lower this axis, in contrast with the activation of *Sirt1* by CR in many other tissues. *Sirt1* may have evolved to positively regulate the somatotrophic axis, as it does insulin production in β cells, to control mammalian health span and life span in an overarching way. However, the fact that *Sirt1* is a positive regulator of the somatotrophic axis may complicate attempts to increase murine life span by whole-body activation of this sirtuin.

It is interesting that *Sirt1* activity in the brain governs the production of GH, but not other pituitary hormones, even though the production of all pituitary hormones is regulated at least in part by signals from the hypothalamus. It will be of interest to dissect the function of this

sirtuin in specific brain regions with more targeted knockout strategies. Likewise, it will be important to study *Sirt1* in the pituitary itself to address whether this sirtuin also controls the other major endocrine axes.

Materials and methods

Animals

All animal procedures were carried out in accordance with the Massachusetts Institute of Technology Committee on Animal Care. Mice were group-housed in a 12-h light/12-h dark cycle at controlled temperature ($25^{\circ}\text{C} \pm 1^{\circ}\text{C}$). For high-fat diet studies, mice were fed diets with 60% or 10% of calories from fat (high fat and control, respectively; Research Diets, Inc.). For CR studies, the amount of food consumed by ad libitum control mice was determined weekly, and CR mice were fed 60% of that value daily. For GTT and ITT, mice were fasted overnight prior to injection with either 2 g/kg glucose (Sigma) or 1 IU/kg porcine insulin (Sigma) delivered in sterile saline. Blood glucose was measured at the indicated time points using a LifeScan OneTouch glucometer. Activity was measured by individually housing animals with low-profile wireless running wheels (Med Associates, Inc.). For tests of the response of animals to a bolus of glucose, mice were fasted for 14 h and then injected with either 2 g/kg glucose or an equivalent volume of saline vehicle and were analyzed 45 min after injection.

Western blotting

Tissues were lysed in TNT buffer (20 mM Tris at pH 8.0, 150 mM NaCl, 1% Triton X-100) supplemented with protease inhibitors (Roche), and blotted with antibodies against *Sirt1* (Millipore) and tubulin (Sigma).

Hormone measurements

Serum for hormone measurements was collected by retro-orbital puncture under isoflurane anesthesia. ELISA kits for measuring hormone concentrations (listed in the Supplemental Material) were used according to the manufacturer's instructions.

Quantitative RT-PCR

Total RNA was isolated using Trizol reagent (Invitrogen) and was treated with DNase (Promega) prior to reverse transcription with MMLV reverse transcriptase (Invitrogen) and random hexamer primers (New England Biolabs). Real-time PCR reactions were performed on a LightCycler 480 II (Roche) using iQ SYBR Green Supermix (Bio-Rad). The relative abundance of transcripts of interest was calculated by first normalizing to a housekeeping gene (β -actin or 18S RNA) and then normalizing to wild-type controls using the $2^{-\Delta\Delta\text{CT}}$ method. Primer sequences were taken from the Harvard Primer Bank (Wang and Seed 2003; Spandidos et al. 2008). For sex-specific gene expression in the liver, primer sequences were as described (Holloway et al. 2008).

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Erratum

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Neuronal SIRT1 regulates endocrine and behavioral responses to calorie restriction

Dena E. Cohen, Andrea M. Supinski, Michael S. Bonkowski, Gizem Donmez, and Leonard P. Guarente

Due to an error during figure preparation for the above-mentioned paper, one of the graphs in Figure 4 was inadvertently duplicated such that the graphs in panels B and E appeared identical. A corrected version of the figure is shown below, such that B remains as originally presented and E is replaced by the correct, nonduplicated graph. This error in no way affects the conclusions of the paper.

The authors apologize for any confusion caused by this error.

Please note that this has been corrected in the online version of this paper.

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