Haploinsufficiency of Akt1 Prolongs the Lifespan of Mice

Aika Nojima1, Masakatsu Yamashita2, Yohko Yoshida1, Ippei Shimizu1, Harumi Ichimiya3, Naomi Kamimura3, Yoshiro Kobayashi1, Shigeo Ohta3, Naoaki Ishii4, Tohru Minamino5,6,*

1 Department of Cardiovascular Medicine, Chiba University Graduate School of Medicine, Chiba, Japan, 2Kazusa DNA Research Institute, Kisarazu, Chiba, Japan, 3Department of Biochemistry and Cell Biology, Institute of Development and Aging Sciences, Graduate School of Medicine, Nippon Medical School, Nakahara-ku, Kawasaki, Kanagawa, Japan, 4Department of Molecular Life Science, Basic Medical Science and Molecular Medicine, Tokai University School of Medicine, Isehara, Japan, 5Department of Cardiovascular Biology and Medicine, Niigata University Graduate School of Medical and Dental Sciences, Niigata, Japan, 6PRESTO, Japan Science and Technology Agency, Kawaguchi, Saitama, Japan

Abstract

There is increasing evidence that nutrient-sensing machinery is critically involved in the regulation of aging. The insulin/insulin-like growth factor-1 signaling pathway is the best-characterized pathway with an influence on longevity in a variety of organisms, ranging from yeast to rodents. Reduced expression of the receptor for this pathway has been reported to prolong the lifespan; however, the underlying mechanisms are largely unknown. Here we show that haploinsufficiency of Akt1 leads to an increase of the lifespan in mice. Akt1+/− mice had a lower body weight than their littermates with less fat mass and normal glucose metabolism. Ribosomal biogenesis and the mitochondrial DNA content were significantly reduced in these mice, along with a decrease of oxidative stress. Consistent with the results obtained in mice, inhibition of Akt-1 promoted longevity in nematodes (Caenorhabditis elegans), whereas activation of Akt-1 shortened the lifespan. Inhibition of Akt-1 led to a decrease of ribosomal gene expression and the mitochondrial DNA content in both human cells and nematodes. Moreover, deletion of ribosomal gene expression resulted in a decrease of the mitochondrial DNA content and normalized the lifespan shortened by Akt-1 activation in nematodes. These results suggest that an increase of mitochondrial amount and energy expenditure associated with enhanced protein synthesis accelerates both aging and the onset of age-associated diseases.

Introduction

Endocrine signaling was first linked to longevity when it was shown that mutations of daf-2, a homologue of the mammalian insulin/insulin-like growth factor-1 (IGF-1) receptor [1], dramatically prolonged the lifespan of the nematode Caenorhabditis elegans [2]. Genetic analysis subsequently demonstrated that reduction-of-function mutations affecting various genes in the insulin/IGF-1/phosphatidylinositol-3 kinase (PI3K)/Akt signaling pathway prolonged the lifespan of C. elegans [2–9]. Inhibiting this pathway confers longevity through changes in the expression of genes regulated by transcription factors such as the forkhead transcription factor DAF-16, the heat-shock transcription factor HSF-1, and the xenobiotic factor SKN-1 [10]. It has also been reported that the genes regulating longevity are conserved in a wide range of organisms ranging from yeast to mice. Mutation of S6K, which is homologous with Akt, extends the lifespan of yeast [11], while mutations that decrease the activity of insulin/IGF-1 pathway improve the longevity of fruit flies [12] and mice [13,14].

Target of rapamycin (TOR) is an evolutionarily conserved nutrient-sensing protein kinase that regulates growth and metabolism in all eukaryotic cells [15]. Studies performed in worms, flies, yeast, and mice support the notion that the TOR signaling network modulates aging [15–19]. Like inhibition of the insulin/IGF-1 pathway, inhibition of TOR increases resistance to environmental stress and requires transcriptional changes in order to extend the lifespan of yeast and worms [10,20,21]. In response to the intake of nutrients, TOR up-regulates translation activity, partly by activating the ribosomal subunit S6 kinase and inhibiting 4E-BP, which is a translation inhibitor. When nutrient levels and TOR activity are decreased, translation activity also declines. This appears to have a positive impact on the lifespan, since inhibition of S6 kinase improves longevity in yeast, nematodes, flies, and mice [10,15,22]. In C. elegans, DAF-16 plays an essential role in longevity related to inhibition of the insulin/IGF-1 pathway [23,24], while inhibition of TOR extends the lifespan independently of DAF-16 [17,25]. In mammals, however, the role of TOR in longevity related to inhibition of the insulin/IGF-1 pathway is largely unknown.

Here we studied Akt1+/− mice and found that their lifespan was significantly longer than that of littermates controls. We then sought to elucidate the mechanisms related to the increased longevity of these mice. Akt1+/− mice showed a decrease of TOR signaling, but phosphorylation of the forkhead transcription factors (FOXO) was not down-regulated. Gene ontology analysis suggested a crucial role of the suppression of translation and
mitochondrial activity in promoting the longevity of Akt1+/– mice, suggesting that the TOR pathway is critically involved in prolonging the lifespan of mammals by inhibiting the insulin/IGF-1 pathway.

**Materials and Methods**

**Animal Models**

All experiments using live mice were performed in strict accordance with the guidelines of the American Association for Accreditation of Laboratory Animal Care, and the study protocol was approved by Chiba University Institutional Animal Care and Use Committee. Akt1-deficient mice (Akt1+/–) were a kind gift from Dr. Morris J. Birnbaum (University of Pennsylvania School of Medicine, Philadelphia, PA). Generation and genotyping of Akt1-deficient mice have been described previously [26]. Heterozygous mice were backcrossed with wild-type C57BL/6 mice (SLC, Japan) for 6 generations. All mice were maintained under specific-pathogen-free conditions, and their lifespan was monitored by experienced technicians at Sankyo Laboratory Service (n = 101 for wild-type male mice, n = 103 for Akt1+/– male mice, n = 79 for wild-type female mice, n = 80 for Akt1+/– female mice). Survival curves were plotted by the Kaplan–Meier method, and differences between groups were evaluated by the log-rank test. The maximum lifespan was calculated as the average for the oldest 20% of the mice within each group [27]. Data shown as the mean ± s.e.m. *P<0.05, **P<0.001. Differences of lifespan between groups were evaluated by the log-rank test.

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**Cell Culture and Retroviral Infection**

Because it was very difficult to expand primary cultures of hepatocytes for infection, we utilized human endothelial cells that are highly proliferative. Human umbilical vein endothelial cells were purchased from Lonza (Walkersville, MD), and cultured according to the manufacturer’s instructions. We created a pLNCX (Clontech, Palo Alto, CA)-based vector expressing a dominant-negative form of AKT1 (AKTDN). Retroviral stocks were generated by transient transfection of a packaging cell line (293T, Clontech) and were stored at –80°C until use. Human endothelial cells (passages 4–6) were plated at 5–6×10^5 cells in 100 mm diameter dishes at 24 hours before infection. Then the culture medium was replaced by retroviral stock supplemented with 8 μg/ml polybrene (Sigma, Tokyo, Japan) for infection. After 48 hours, infected cell populations were selected by culture in 500 mg/ml G418 for 7 days. On the 8th day post-infection, 1–3×10^5 infected cells were seeded onto 100 mm diameter dishes. Oxygen consumption rates of cell cultures were determined with a 96-well BD Oxygen Biosensor System plate (BD Biosciences, San Jose, CA).

**Physiological Analysis**

We housed mice individually to monitor their body weight and food intake. Adiposity was examined by CT scanning (LaTheta, Aloka) according to the manufacturer’s protocol. We obtained CT scans at 2 mm intervals from the diaphragm to the floor of the pelvic cavity. Oxygen consumption was measured in 8-week-old and 40-week-old mice with an O_2/CO_2 metabolic measurement system (Model MK-5000, Muromachikikai), as described previ-
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**Incidence of malignancy in Wt and Akt1^{+-} mice**

|       | Female |
|-------|--------|
|       | Wt     | Akt1^{+-} |
| malignancy | 6 (60%) | 7 (70%) |
| no malignancy | 4 (40%) | 3 (30%) |

**Downregulated genes**

| GO Term                          | Genes | P-value   |
|----------------------------------|-------|-----------|
| mitochondrion                    | 151   | 1.66E-28  |
| mitochondrial membrane            | 62    | 6.96E-20  |
| ribosome                         | 28    | 2.08E-11  |
| proton-transporting two-sector ATPase complex | 8     | 1.82E-08  |
mice at 14 weeks and 40 weeks old (n = 58). Body weight was lower in Akt1+/– female mice at 8 weeks and 40 weeks old (n = 9). Visceral fat weight was reduced in Akt1+/– female mice at 8 weeks and 40 weeks old (n = 8). Glucose tolerance did not differ between the two strains. Data are shown as the mean ± s.e.m. *P<0.05, **P<0.01, and ***P<0.001 by Student’s t-test for A–H. (I) Microarray analysis of liver samples from Akt1+/– female mice and wild-type littersmates (n = 3). Gene ontology analysis demonstrated that genes related to the mitochondria and ribosomes were significantly down-regulated in Akt1+/– mice (two-way ANOVA).

Histological Analysis
Lever tissue samples were harvested and fixed in 10% formalin overnight, followed by embedding in paraffin and sectioning. Then the sections were subjected to HE staining or to immunohistochemistry with anti-4-hydroxy-2-nonenal antibody (Abcam).

Microarray Analysis
The hepatic gene profile of wild-type and Akt1+/– mice was analyzed at 8 weeks and 40 weeks of age by using Agilent Whole Mouse 44K Arrays (n = 3 per group). The raw data were subjected to log2 transformation and normalized by using the GeneSpring GX v7.3.1 (Agilent Technologies). Differentially expressed genes (p<0.01) were determined by two-way ANOVA using two parameters, which were the genetic background (Wild-type or Akt1+/–) and the age (8 weeks or 40 weeks). Gene ontology analysis was performed based on each category of two-way ANOVA. Gene expression data were deposited in the Gene Expression Omnibus database (GSE39699).

Analysis of Mitochondria
Isolation of mitochondria was performed as described previously [32]. In brief, mice were sacrificed by decapitation and their livers were harvested immediately, washed in ice-cold isolation buffer (225 mM mannitol, 75 mM sucrose, 5 mM HEPES, 1 mM EGTA), and minced with a razor blade. Then the tissue was homogenized with a motorized Teflon/glass homogenizer, the homogenate was centrifuged for 5 minutes at 500 × g at 4°C, and the supernatant was collected and re-centrifuged for 5 minutes at 500 × g. The resulting supernatant was then centrifuged for 10 minutes at 8000 × g at 4°C, and the pellet was suspended in isolation buffer. Unless otherwise indicated, all procedures were performed on ice. Protein concentrations were determined by the BCA protein assay (Pierce). Oxygen consumption was measured with an Oxygen Meter (Model 781) and a Mitocell MT200 closed respiration chamber (Strathkelvin Instruments, North Lanarkshire, UK) at 37°C with continuous stirring in respiration buffer (125 mM KCl, 1 mM K2HPO4, 5 mM MgCl2, 25 mM HEPES, 0.2 mM EGTA, and 20 mM mannitol). Mitochondria, pyruvate, and malate (2.5 mM each), 500 nM rotenone, and 5 mM succinate were added sequentially to the buffer. Oxygen consumption by complex I was defined as the rotenone-sensitive component of oxygen consumption in the presence of pyruvate plus malate. Oxygen consumption by complex II was defined as consumption after the addition of succinate minus consumption.

Isolation of Hepatocytes
Hepatocytes were isolated as described previously [33,34]. In brief, 40 week-old mice were anesthetized and the abdominal cavity was opened. A 23G needle was introduced into the portal vein, and perfusion was started with Hepatocyte Liver Perfusion Medium (1×) (Gibco) after proximal ligation of the inferior vena cava.
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A

B

C

D

E

F

G

H

I

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Figure 3. Ribosomal biogenesis and mitochondrial function in young and middle-aged female Akt1+/– mice. (A) Western blot analysis of phosphorylated mTOR, phosphorylated p70 S6 kinase, and phosphorylated FoxO3a expression in the livers of wild-type and Akt1+/– mice at 40 weeks old. (B) Pre-rRNA level in the livers of wild-type and Akt1+/– mice at 8 weeks and 40 weeks old were examined by real-time PCR (n = 10). (C) Mitochondrial DNA content of the livers prepared as Figure 2B. (D) Real-time PCR analysis of the expression of COX1 (encoding cytochrome c oxidase subunit I), ATP6 (encoding ATP synthase Fo subunit 6), and ND6 (encoding NADH dehydrogenase, subunit 6 (complex I)) in livers prepared as in Fig. 2B (n = 4). (E) Immunohistochemistry for 4-hydroxy-2-nonenal (4-HNE) in the livers of wild-type and Akt1+/– mice at 40 weeks old. Scale bar = 100 μm. The right graph displays the quantitative data on 4-HNE-positive cells (n = 6). (F) Urinary 8-isoprostane level in wild-type and Akt1+/– mice at 40 weeks old (n = 6). (G, H) Number of TMRM-positive (G) and DCF-positive (H) hepatocytes isolated from wild-type and Akt1+/– mice at 40 weeks old, as shown by FACS analysis (n = 4). (I) Oxygen consumption by complex I (left) and complex II (right) in mitochondria isolated from the livers of wild-type and Akt1+/– mice at 40 weeks old (n = 6). Data are shown as the mean ± s.e.m. *P<0.05, **P<0.01 by Student’s t-test. doi:10.1371/journal.pone.0069178.g003

Pathophysiological Features of Akt1+/– Mice

We next investigated whether there were any differences of pathophysiological features between Akt1+/– mice and their littermate controls. Consistent with the previous report of Akt1+/– mice [26], Akt1+/– mice had a lower body weight (Fig. 2A). However, the difference of body weight was not significant when corrected by femoral length (Wild-type mice, 1.52±0.04 g/mm; Akt1+/– mice, 1.48±0.04 g/mm; n = 10, p = 0.45). Although CT scanning showed that Akt1+/– mice had less fat mass than their wild-type littermates (Fig. 2B), there were no significant differences of glucose tolerance or insulin tolerance between the two groups (Fig. 2C, D). There were no significant differences of food intake, body temperature, and activity levels between the two groups (Fig. 2E–G). However, the oxygen consumption of Akt1+/– mice was slightly, but significantly, lower than that of their littermates (Fig. 2H). Although Akt1+/– mice lived significantly longer than their littermates, we did not find any differences of age-associated cardiovascular phenotypes in terms of arterial pressure and cardiac function (Fig. S2A, B). Histological examination of the aorta, bone, and skeletal muscle in aged mice also detected no age-related differences (Fig. S2C).

Since Akt1 signaling has been reported to contribute to tumorigenesis [42,43], we also investigated the effect of haploinsufficiency of Akt1 on the development of malignancy. Histological examination of 2-year-old mice demonstrated that the incidence of malignancy was not altered by Akt1 haploinsufficiency (Fig. 2I). We often observed malignant cells (chiefly lymphomas) infiltrating the tissues (such as liver, skeletal muscle, and visceral fat) of mice over 100 weeks old. Therefore, we used tissue samples from young (8-week-old) and middle-aged mice (40-week-old) for further analyses.

Ribosomal Biogenesis and Mitochondrial Function in Akt1+/– Mice

To gain some insight into the potential mechanisms leading to extension of the lifespan in Akt1+/– mice, we performed microarray analysis of liver, skeletal muscle, and visceral fat obtained from these mice and their wild-type littermates. Gene ontology (GO) analysis revealed that mitochondrial and ribosome were among the most significant GO terms (Fig. 2J and Fig. S3). Consistent with these findings, the mTOR pathway, which has a crucial role in regulating ribosomal biogenesis, protein synthesis, and mitochondrial activity [15,44], was down-regulated in Akt1+/– mice, although phosphorylation of FoxO was unaltered (Fig. 3A and Fig. S4). Indeed, ribosomal biogenesis was markedly reduced in Akt1+/– mice (Fig. 3B), along with a decrease of the mitochondrial DNA content and reduced expression of genes for mitochondrial components and transcription factors involved in mitochondrial biogenesis, when compared with their wild-type littermates (Fig. 3C, D and Fig. S5). These changes were associated with...
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A. Oxygen consumption

B. Fold increase

C. Relative N2 expression

D. Mitochondrial DNA content (ratio)

E. Survival (%)

F. Mitochondrial DNA content (ratio)

G. Relative expression

H. Survival (%)

I. Survival (%)

J. Mitochondrial DNA content (ratio)

K. Survival (%)

L. Survival (%)

P<0.001

P<0.05

P=0.001

P<0.001
Reduced oxidative stress, as demonstrated by a decrease of 4-hydroxynonenal immunoreactivity in the liver and a decline of 8-isoprostane excretion in the urine (Fig. 3E, F), suggesting that partial inhibition of Akt1 activityameliorated the age-associated increase of oxidative stress.

To further investigate the effects of Akt1 haploinsufficiency on mitochondrial function and oxidative stress, we isolated hepatocytes from Akt1+/− mice and their littermate controls. These cells were subjected to FACS analysis to detect tetramethylrhodamine, methyl ester (TMRM), and dichlorodihydrofluorescein (DCF) fluorescence. This revealed that hepatocytes from Akt1+/− mice showed a significant decrease of the mitochondrial membrane potential and levels of reactive oxygen species (ROS) compared with hepatocytes from their wild-type littermates (Fig. 3G, H). We next examined the activity of mitochondria isolated from the hepatocytes of Akt1+/− mice and their littermate controls, and found that the maximum respiration rate of isolated mitochondria did not differ between the two groups (Fig. 3I). These results indicated that the amounts of mitochondria were decreased in Akt1+/− mice compared with their littermate controls, but the activity of per mg mitochondrion was not impaired by haploinsufficiency of Akt1. We noted that the expression of FoxO-regulated antioxidant genes, such as catalase and superoxide dismutase, did not differ between Akt1+/− mice and their littermate controls (Fig. S6). Since ROS are a byproduct of normal mitochondrial respiration, a decrease in the number of mitochondria could account for a decrease of oxidative stress in Akt1+/− mice.

Ribosomal Biogenesis and Mitochondrial Function in Human Cells and C. elegans

To investigate the role of AKT1 in ribosomal biogenesis and in total mitochondrial activity, we infected human endothelial cells with a retroviral vector encoding a dominant-negative form of AKT1 (AKTDN) or an empty vector (Mock). Data represent the fold increase relative to the initial value (n = 3). P < 0.05 by two-way ANOVA. (B) Pre-RNAi level in human endothelial cells infected with AKTDN or Mock (n = 6). (C) Real-time PCR analysis of the expression of ND6 (encoding NADH dehydrogenase, subunit 6 (complex I)) in human endothelial cells infected with AKTDN or Mock (n = 6). (D) Mitochondrial DNA content of human endothelial cells infected with AKTDN or Mock assessed by real-time PCR (n = 3). Data are shown as the mean ± s.e.m. *P < 0.05 by Student’s t-test. (E) Survival of C. elegans fed with bacteria containing a control vector or the dsRNA construct targeting akt-1. Knockdown of akt-1 significantly prolonged the lifespan. P < 0.001 by the log-rank test. (F) Mitochondrial DNA content in C. elegans prepared as in Fig. 4G (n = 8). (G) Expression of ife-2 (encoding translation initiation factor 4F, cap-binding subunit (eIF4E)) and rps-11 (encoding a small ribosomal subunit 51 protein) in C. elegans prepared as in Fig. 3e (n = 8). Data are shown as the mean ± s.e.m. *P < 0.05, **P < 0.001 by Student’s t-test. (H) Activation of AKT-1 by knockdown of daf-18 (a homologue of PTEN) led to significant shortening of the lifespan of wild-type worms. P < 0.001 by the log-rank test. (I) Knockdown of rps-11 by RNAi led to significant prolongation of the lifespan of wild-type worms. P < 0.001 by the log-rank test. (J) Mitochondrial DNA content in C. elegans prepared as in Fig. 4I (n = 5). Data are shown as the mean ± s.e.m. *P < 0.05 by Student’s t-test. (K) Knockdown of atp-2 (encoding a subunit of ATP synthase, mitochondrial complex V) by RNAi significantly prolonged the lifespan of wild-type worms. P < 0.001 by the log-rank test. (L) Knockdown of rps-11 by RNAi improved shortening of the lifespan of worms with daf-18 RNAi treatment.

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Discussion

The present study demonstrated that haploinsufficiency of Akt1 significantly prolongs the lifespan of mice. Akt1+/− mice showed a decrease of ribosomal biogenesis and the amounts of mitochondrial function in human endothelial cells and the liver of Akt1+/− mice than in their littermates, an increase of mitochondria and energy expenditure associated with enhanced protein synthesis may accelerate aging and promote the onset of age-associated diseases.

An intriguing aspect of longevity pathways is that they function independently of the cells affected, since mutations of one cell type can affect the phenotype of whole organism. Neuronal cells and adipose tissue have been suggested to have a critical role in regulation of the lifespan by the insulin/IGF1 signaling pathway and the TOR pathway both promote ribosomal biogenesis and protein synthesis, as well as regulating mitochondrial function [15,44]. We found that the phosho-Akt1 level increased with age in wild-type mice, presumably due to constitutive activation of growth signals by various stresses. Given the large amount of energy consumed by ribosomal biogenesis and protein synthesis, Akt1+/− mice may utilize fewer mitochondria by reducing ribosomal biogenesis and thus minimize the enhancement of oxidative stress with aging, which could account for their longer lifespan. Decreased protein synthesis could improve the fidelity of biogenesis and prevent the accumulation of mis-folded proteins, which might also contribute to longevity associated with Akt1 deficiency. Since oxidative stress and oxygen consumption were both lower in Akt1+/− mice than in their littermates, an increase of mitochondria and energy expenditure associated with enhanced protein synthesis may accelerate aging and promote the onset of age-associated diseases.

In a recent study, the influence of inhibiting AKT-1 on ribosomal biogenesis, the mitochondrial DNA content, and the lifespan of C. elegans was investigated [14]. In agreement with the results obtained in Akt1+/− mice, inactivation of AKT-1 by RNAi resulted in a longer lifespan compared with that of wild-type (N2) C. elegans (Fig. 4E), and this change was associated with a decrease of ribosomal gene expression and reduction of the mitochondrial DNA content (Fig. 4F, G). Conversely, activation of AKT-1 by RNAi targeting daf-18 led to a shorter lifespan (Fig. 4H). Inhibition of ribosomal biogenesis by RNAi decreased the mitochondrial DNA content and extended the lifespan of wild-type animals (Fig. 4I, J), while inhibition of mitochondrial function by RNAi also increased the lifespan of wild-type animals (Fig. 4K). Moreover, inhibition of ribosomal biogenesis normalized the shortened lifespan of nematodes treated with daf-18 RNAi (Fig. 4L), suggesting that the decrease of ribosomal biogenesis and mitochondrial function were critical for improving the longevity of Akt1+/− mice.
Thus, it would be interesting to test the effects of tissue-specific deletion of Akt1 on the lifespan in the future.

Consistent with our findings, modest inhibition of respiration has been reported to prolong the lifespan of a variety of species, such as yeast, nematodes, flies, and mice [49–52]. This increase of longevity could be partly attributable to reduction of the metabolic rate in these animals. In contrast, increasing respiration was reported to promote longevity in animals with caloric restriction [53,54], so it is possible that increasing or reducing respiration can influence the lifespan in various ways.

Genetic inhibition of autophagy induces degenerative changes in mammalian tissues that resemble those associated with aging, while normal and pathological aging are often associated with a reduced autophagic potential [15,55]. Genetic manipulations that prolong the lifespan in various models often stimulate autophagy, and inhibition of autophagy compromises the longevity-promoting effect of caloric restriction or suppression of insulin/insulin growth factor signaling [15,55]. Since mTOR is a primordial negative regulator of autophagy, an increase of autophagic activity may also contribute to extending the lifespan of Akt1+/– mice. In this context, it would be interesting to examine the effect of inhibiting the TOR/autophagy pathway on the lifespan of C. elegans with akt-1 or daf-18 knockdown.

Telomeres are specialized DNA-protein structures found at the ends of eukaryotic chromosomes that serve as markers of biological aging [56]. Telomeres also play a critical role in maintaining genomic integrity and are involved in age-related diseases [28,57]. Shortening of telomeres is hazardous to healthy cells, as it is a known mechanism of premature cellular senescence and reduction of longevity. Telomerase is an enzyme that adds telomeres to the ends of chromosomes. Although the insulin/Akt pathway has been reported to positively regulate telomerase activity [58], mice have high telomerase activity and long telomeres [59,60]. Therefore, it is unlikely that Akt1 signaling regulates longevity by modulating telomerase activity in mice.

In conclusion, our results suggest that haploinsufficiency of Akt1 significantly promotes longevity in mice by mechanisms that involve reduction of both energy expenditure and oxidative stress. Further studies on improvement of longevity related to inhibition of the insulin/IGF-1 pathway should provide useful insights into the treatment of diseases associated with aging.

Supporting Information

Figure S1 Age-associated increase of phospho-Akt1 expression. Western blot analysis of phosphorylated Akt1 expression in the livers of wild-type (Wt) and Akt1+/– female mice at 8 and 40 weeks old.

(DOCX)

Figure S2 Examination of age-related phenotypes. (A) Arterial pressure of wild-type (Wt) and Akt1+/– female mice at 100 weeks old. Data are shown as the means ± s.e.m. (B) Echocardiographic analysis of wild-type (Wt) and Akt1+/– female mice at 100 weeks old. FS, fractional shortening; LVDS, left ventricular diastolic dimension. Data are shown as the means ± s.e.m. (C) Hematoxylin-eosin staining of the aorta, bone, and skeletal muscle of wild-type (Wt) and Akt1+/– female mice at 100 weeks old. Scale bar: 20 μm.

(DOCX)

Figure S3 Microarray analysis. Microarray analysis of fat and skeletal muscle samples from Akt1+/– female mice and wild-type littermates (n = 3).

(DOCX)

Figure S4 Expression of phospho-FoxO. Western blot analysis of phosphorylated FoxO3a expression in various tissues of wild-type (Wt) and Akt1+/– female mice at 100 weeks old.

(DOCX)

Figure S5 Expression of transcription factors involved in mitochondrial biogenesis. The expression of Pgc-1α (also known as Ppargc1a) and its regulating molecules related to mitochondrial biogenesis, such as nuclear respiratory factor 1 (Nrf1) and mitochondrial transcription factor A (Tfam) was examined by real-time PCR in livers of wild-type (Wt) and Akt1+/– female mice at 40 weeks old. Data are shown as the mean ± s.e.m (n = 5–8). *P<0.05.

(DOCX)

Figure S6 Expression of antioxidant genes. The expression of catalase (Cat) and superoxide dismutase 2 (Sod2) was examined by real-time PCR in livers of wild-type (Wt) and Akt1+/– female mice at 100 weeks old. Data are shown as the mean ± s.e.m (n = 4). *P<0.05.

(DOCX)

Author Contributions

Conceived and designed the experiments: AN TM. Performed the experiments: AN YY IS HI NK SO. Analyzed the data: MY YK. Contributed reagents/materials/analysis tools: NI. Wrote the paper: AN TM.
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