Inhibition of adipogenesis and development of glucose intolerance by soluble preadipocyte factor–1 (Pref-1)

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Preadipocyte factor-1 (Pref-1) is a transmembrane protein highly expressed in preadipocytes. Pref-1 expression is, however, completely abolished in adipocytes. The extracellular domain of Pref-1 undergoes two proteolytic cleavage events that generate 50 and 25 kDa soluble products. To understand the function of Pref-1, we generated transgenic mice that express the full ectodomain corresponding to the large cleavage product of Pref-1 fused to human immunoglobulin-γ constant region. Mice expressing the Pref-1/hFc transgene in adipose tissue, driven by the adipocyte fatty acid–binding protein (aP2, also known as aFABP) promoter, showed a substantial decrease in total fat pad weight. Moreover, adipose tissue from transgenic mice showed reduced expression of adipocyte markers and adipocyte-secreted factors, including leptin and adiponectin, whereas the preadipocyte marker Pref-1 was increased. Pref-1 transgenic mice with a substantial, but not complete, loss of adipose tissue exhibited hypertriglyceridemia, impaired glucose tolerance, and decreased insulin sensitivity. Mice expressing the Pref-1/hFc transgene exclusively in liver under the control of the albumin promoter also showed a decrease in adipose mass and adipocyte marker expression, suggesting an endocrine mode of action of Pref-1. These findings demonstrate the inhibition of adipogenesis by Pref-1 in vivo and the resulting impairment of adipocyte function that leads to the development of metabolic abnormalities.

differentiation (10–12, 15). In this regard, glucocorticoids enhance adipose differentiation partly by downregulation of Pref-1 expression (15). Recently, we found that the transmembrane form of Pref-1 is cleaved at the juxtamembrane region to generate a 50-kDa large soluble form that contains the full ectodomain of Pref-1 (11). In addition, smaller fragments are also produced due to processing that occurs at a more N-terminal region. Together, these observations suggest that Pref-1 may function either in a juxtransin/paracrine or in an endocrine manner to inhibit adipocyte differentiation.

Apart from preadipocytes, Pref-1 is also expressed in various tissues during embryogenesis, and its expression is abolished after birth (10, 16, 17). In this regard, the gene encoding for Pref-1 has been reported recently to be an imprinted gene that is paternally expressed due to differential methylation (18–20). Given the role of imprinted genes in fetal growth and development, in general, and the expression of Pref-1 in embryonic tissues, Pref-1 may have functions beyond the regulation of adipogenesis (18–22).

Here we show the inhibitory role of the large soluble form of Pref-1 on adipogenesis by generating transgenic mice expressing the full ectodomain of Pref-1 as a human immunoglobulin-γ constant (hFc) region fusion protein. The large soluble form of Pref-1 specifically expressed in adipose tissue caused a decrease in fat pad weight, with a decrease in average adipocyte cell size, as well as in the expression of adipocyte marker genes, indicating an impaired adipogenesis. Hepatic expression of the same Pref-1/hFc transgene also produced a decrease in adipose tissue mass, suggesting the endocrine mode of Pref-1 action. Transgenic mice with a substantial reduction in adipose tissue mass showed hypertriglyceridemia, impaired glucose tolerance, and reduced insulin sensitivity, demonstrating the significance of adipose tissue mass to the development of the diabetes-like syndrome.

Methods

Transgene constructs and generation of transgenic mice. The transgene constructs were designed for either adipose- or liver-specific expression of the soluble Pref-1/hFc fusion protein under the control of the adipocyte fatty acid–binding protein (aP2) promoter or the albumin promoter, respectively. To generate a soluble Pref-1/hFc fusion protein, a PCR product encoding the large soluble form that contains the full ectodomain of Pref-1 (5′-CAC GAG CGT CCT GTT CAG CAG CC-3′) and the 5′ end of the human Fc cDNA sequence (5′-CTT GAC CTC AGG GTC TTC GTG-3′) were used to amplify a 254-bp fragment. Transgenic mice and their wild-type control littermates were maintained under standard temperature and lighting.

Measurements for body weight, food intake, and total body fat. All mice were weighed at 5-day intervals from 3 to 10 weeks of age. For food intake measurement, 10- to 12-week-old transgenic mice and their wild-type littermates were individually housed. Body weight and daily food consumption were measured for 8 days. Body fat and fecal fat content were determined by the modified method of Bligh and Dyer (25).

Western blot analysis. Serum (3 μl) from 10-week-old wild-type and transgenic mice were subjected to 10% SDS-PAGE and transferred onto nitrocellulose membranes. Endogenous Pref-1 or Pref-1/hFc fusion protein was probed with a polyclonal rabbit anti-Pref-1 Ab (1:5,000) and a goat anti-rabbit IgG Ab conjugated with peroxidase (1:5,000) as a secondary Ab. Pref-1/hFc fusion protein was also probed with an anti-human IgG Ab conjugated with peroxidase (1:3,000). Protein (20 μg) isolated from extracts of 12-day-old whole embryos were used for Western blotting analysis.

Tissue collection and histology. Mice were anesthetized in a CO2 chamber and euthanized by cervical dislocation. Inguinal, renal, and epididymal or parametrial depots of white adipose tissue and interscapular brown adipose tissue were collected from 10-week-old mice. For histology, renal fat tissue was excised and fixed in Bouin’s fluid, embedded in paraffin, and sliced into 8-μm sections. Sections were stained with hematoxylin and eosin. Images of adipose tissue sections were captured and adipocyte volumes were measured from at least 300 cells by NIH image software.

Northern blot analysis and RT-PCR. Total RNA from tissues or whole embryos was prepared using TRIzol reagent (Invitrogen Corp., San Diego, California, USA). Total RNA (15 μg) was subjected to electrophoresis through a 1.2% formaldehyde-agarose gel in 2.2 M formaldehyde, 20 mM Mops, 1 mM EDTA, and transferred to a nylon membrane (Hybond N; Amersham Pharmacia Biotech, Piscataway, New Jersey, USA). After cross-linking by UV, the membranes were hybridized with 32P-labeled cDNA probes for aP2, FAS, SCD-1, UCP-1, C/EBPδ, ADSF/resistin, adiponectin, and leptin in ExpressHyb solution (CLONTECH Laboratories Inc., Palo Alto, California, USA). For RT-PCR, total RNA from epididymal or parametrial fat pads from 10-week-old mice was reverse transcribed with SuperScript II (Invitrogen Corp.). The endogenous Pref-1 cDNA was amplified using primers (forward: 5′-GAC GAG CCT CGT TTT CTC AAC AAG TG-3′; reverse: 5′-GTA AGC ATA GGC TTC ACT CGA TTC-3′) corresponding to the 3′ noncoding region, enabling detection of all forms of endogenous Pref-1 mRNA. β-actin primers (forward, 5′-TCC TAT GTG GGT GAC GAG GC-3′; reverse, 5′-CAT GG CGT GGG TGT TGA AGG-3′) were used as internal control.
Glucose and insulin tolerance tests. Glucose and insulin tolerance tests were performed on 10- to 12-week-old transgenic mice, and their wild-type control littersmates. For glucose-tolerance test, D-glucose (2 mg/g of body weight) was intraperitoneally injected into overnight fasted mice, and glucose levels were monitored at 0, 30, 60, and 120 min after injection using an Accu-Chek glucometer (Roche Diagnostics Corp., Indianapolis, Indiana, USA). Insulin-tolerance tests were performed on mice following a 5-hour fast. Animals were injected intraperitoneally with 0.5 U/kg body weight of insulin (Eli Lilly Co., Indianapolis, Indiana, USA). Tail-blood samples were taken at time 0, 30, 60, 90, and 120 min after injection for measurement of blood glucose levels.

Measurements of serum triglyceride and insulin levels. Serum triglycerides were analyzed with triglyceride INT 10 kit (Sigma-Aldrich, St. Louis, Missouri, USA). Insulin levels after overnight fasting were measured using a Linco Rat RIA kit. Leptin and adiponectin levels in serum were determined by RIA (Linco Research Inc., St. Charles, Missouri, USA).

Skeletal preparations. Embryos or adult mice were skinned, eviscerated, and fixed in 90% ethanol for at least 1 week. For cartilage staining, carcasses were incubated with 0.01% alcian blue in 80% ethanol and 20% glacial acetic acid for 3 days. Then samples were rehydrated with series of ethanol, 70, 40, and 15% for 2 hours. Subsequently, mice were incubated with alizarin red in 1% KOH for 5 days for bone staining. After rinsing with 1% KOH, samples were stored in 100% glycerol.

Statistical analysis. Data are expressed as the mean ± SEM. The statistical significance of differences in mean values between transgenic and wild-type littersmates was assessed by Student t test.

Results and Discussion
We employed the full Pref-1 ectodomain corresponding to the large soluble form of Pref-1 (10, 11, 26) as an Fc fusion protein to generate transgenic mice expressing the fusion gene in adipose tissue under the control of the aP2 promoter. Since Pref-1 has been known to function as a dimer, addition of human Fc to generate a Pref-1 fusion protein would enhance its dimerization and bioactivity (27, 28). The large soluble form of Pref-1 expressed in adipocytes in this manner may affect preadipocyte differentiation in a paracrine fashion as well as in a long-range endocrine action via the circulation. Five transgenic founder lines expressing the transgene were generated. Northern blot analysis of various tissues from adult transgenic mice demonstrated that the aP2-Pref-1/hFc transgene transcript was expressed exclusively in adipose tissues, including inguinal, reproductive, and renal white adipose tissue, as well as in interscapular brown fat. The expression of the aP2-Pref-1/hFc transgene was not detected in any

Figure 1
aP2-Pref-1/hFc transgene expression. (a) Analysis of transgene expression by Northern blot analysis in 3AS line of aP2-Pref-1/hFc mice. Total mRNA was extracted from the tissues of 10-week-old transgenic mice (TG) and wild-type littersmates and probed with radiolabeled Pref-1 cDNA probe. S, spleen; Th, thymus; Li, liver; Lu, lung; K, kidney; SM, skeletal muscle; BAT, brown adipose tissue; Rep, reproductive fat pad; Ing, inguinal fat pad; Ren, renal fat pad. (b) Western blot analysis for Pref-1/hFc fusion protein in serum of three lines (1AS, 2AS, and 3AS) of aP2-Pref-1/hFc transgenic mice. The serum proteins were separated by SDS-PAGE and probed with human Fc antibody. The antibody detected the 75-kDa of Pref-1/hFc fusion protein in three founder lines of transgenic but not in wild-type mice.

Figure 2
Growth curve and organ weight of 3AS aP2-Pref-1/hFc transgenic mice. (a) Growth curves for male and female wild-type (open circles) and transgenic (filled circles) (TG) mice fed a chow diet. Body weight of mice measured at 5-day intervals is shown; each point represents mean ± SEM from 6 to 13 mice. Body weight of Pref-1/hFc transgenic mice was significantly lower (P < 0.01) than those of wild-type mice at all ages. (b) Organ weights of 10-week-old mice are presented as percentage of body weight (n = 8–10 per group). Statistically significant differences between the groups are indicated as *P < 0.05 and **P < 0.01.
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Reduced fat pad weight in 3AS aP2-Pref-1/hFc transgenic mice. Fat depot weights from 10-week-old mice are presented (n = 6 per group). BAT, brown adipose tissue; Renal, renal fat pad; Ing, inguinal fat pad; Para, parametrial fat pad; Epi, epididymal fat pad. Statistically significant differences between the groups are indicated as *P < 0.05 and **P < 0.01.

Figure 3

Histological analysis of adipose tissue and cell size distribution. (a) Paraffin-embedded sections of renal white adipose tissue from 10-week-old male mice were stained with hematoxylin and eosin. Scale bar, 50 µm. (b) Distribution of the adipocyte volume in wild-type and aP2-Pref-1/hFc transgenic mice. The volume of at least 300 cells per sample (mean of four mice per group) was determined with the NIH image software.

other tissue examined, including spleen, thymus, liver, lung, kidney, and muscle (Figure 1a). High levels of circulating Pref-1/hFc fusion protein were detected in three aP2-Pref-1/hFc transgenic lines (1AS, 2AS, and 3AS) (Figure 1b), whereas two other lines had barely detectable levels of plasma Pref-1/hFc. Wild-type mice, as expected, did not show any Pref-1/hFc in circulation as assessed by Western blot, using a specific Ab against hFc (Figure 1b).

At 3 weeks of age, 3AS aP2-Pref-1/hFc transgenic mice, expressing the highest levels of circulating Pref-1/hFc, had lower body weight than their wild-type littermates (Figure 2a). The body weight of the 3AS aP2-Pref-1/hFc transgenic mice was lower than that of wild-type mice throughout life up to 10 weeks of age (P < 0.01) (Figure 2a). We observed no significant changes in food intake (relative to body weight) or in fat absorption as monitored by daily food consumption and fecal lipid content (data not shown). Despite the lower body weight, the mass of various organs, expressed as a percentage of body weight, was larger in the transgenic mice (Figure 2b). We then investigated whether the lower body weight of the transgenic mice was due to a decrease in fat pad mass. The 3AS transgenic mice had smaller inguinal, renal, and epididymal/parametrial fat pad weight at 10 weeks of age compared with wild-type littermates (in males, 42, 75, and 34%, respectively; in females 38, 39, and 50%, respectively; P < 0.01; Figure 3). In the midlevel expresser line (2AS), inguinal, renal, and reproductive fat pad weight was reduced to a lesser extent (in males, 26, 57, 25%, respectively; n = 5–6, P < 0.05; in females, 18, 27, and 38%, respectively; n = 5–8, P < 0.05). This indicates that the magnitude of decrease in white fat pad weight was positively correlated with the levels of transgene expression. We also observed a significant 36% reduction in interscapular brown adipose tissue in 3AS male transgenic mice, although the decrease was not obvious in the midlevel expresser line 2AS. Total body lipid content was also measured to examine whether the reduced proportion of fat pad mass was reflected in the total lipid content of the animals. The 3AS transgenic mice had less total body lipid (1.82 ± 0.08 g) than their wild-type littermates (2.86 ± 0.16 g, P < 0.01). The 36% reduction of total body lipid content in transgenic mice was similar to the decrease in fat pad weight described above, indicating that fat content in other tissues probably did not change significantly, and the decrease in total body fat can be attributed mainly to a reduction of adipose tissue mass.

A decrease in adipose tissue mass can result from a reduction in adipocyte cell size and/or a decrease in adipocyte cell number due to an impairment of preadipocytes’ differentiation into adipocytes. We first measured the total amount of DNA in fat pads to establish whether decreased fat pad weight of transgenic mice was accompanied by a decreased number of cells. The total DNA content of adipose tissue did not differ between wild-type and transgenic mice (data not shown). Histological analysis of renal fat pads from the 3AS transgenic mice and wild-type littermates at 10 weeks of age revealed that adipocytes from transgenic mice were markedly smaller than those from wild-type mice (Figure 4a). Regarding the distribution of renal adipocyte volume, transgenic mice showed a shift of peaks down to smaller sizes, around 10^4 µm^3 and 1 to 4 × 10^5 µm^3 (Figure 4b). Adipose cell differentiation involves changes in expression of genes that result in the acquisition of fat cell phenotype and adipocyte function. Not only specialized in triglyceride storage, mature adipocytes are characterized by the ability to express and secrete a wide range of molecules involved in many physiological processes, including energy homeostasis. Northern blot analysis showed that aP2-Pref-1/hFc transgenic mice had decreased expression of transcription factors known to be involved in adipogenesis, such as C/EBPα, as well as other adipocyte markers, such as aP2, FAS (fatty acid synthase), and SCD (stearoyl-CoA desaturase1), in the three different fat pads examined (Figure 5a). The expression of factors secreted by adipocytes, including leptin, adiponectin, and ADSF/ resistin was also decreased. In contrast, expression of...
endogenous Pref-1, a preadipocyte marker (12, 13), was increased in Pref-1/hFc transgenic mice as detected by RT-PCR (Figure 5b). Moreover, interscapular brown adipose tissue also exhibited a decrease in expression of aP2, FAS, and SCD, as well as the brown adipocyte marker UCP-1 (uncoupling protein 1) (Figure 5c). Taken together, these observations suggest that the decrease in fat mass in aP2-Pref-1/hFc transgenic mice reflects an impairment of adipocyte differentiation by Pref-1/hFc in all adipose tissue pads, including brown fat. The inhibition of adipogenesis and the consequent decrease in fat mass in these transgenic mice agree with our previous in vitro studies on Pref-1 in which constitutive expression of Pref-1 leads to inhibition of 3T3-L1 adipocyte differentiation, whereas antisense Pref-1 enhances adipocyte differentiation (10, 15). Furthermore, the effectiveness of the large extracellular domain of Pref-1 in transgenic mice led us to conclude that the large soluble Pref-1 is sufficient to inhibit adipocyte differentiation. In agreement with these results, we have reported recently that only the large, soluble form of Pref-1 inhibits 3T3-L1 differentiation in vitro (29).

Since the aP2 promoter was employed to express the transgene in adipose tissue, the high levels of Pref-1/hFc fusion protein secreted in adipose tissue and the amounts present in the circulation of transgenic mice may have affected differentiation of preadipocytes to adipocytes in a paracrine or in an endocrine manner, respectively. We therefore generated transgenic mice expressing the Pref-1/hFc fusion gene under the control of the albumin promoter. Three different Alb-Pref-1/hFc transgenic lines (4Alb, 21Alb, and 23Alb) expressing the transgene at different levels were obtained (Figure 6a, upper panel). Northern blot analysis revealed that the transgene was ex-pressed exclusively in the liver (Figure 6a, lower panel). As shown by Western blot analysis, hepatic expression of the Pref-1/hFc transgene produced a soluble form of the Pref-1/hFc fusion protein in circulation (Figure 6a, upper panel). Transgenic lines 4Alb and 21Alb, which express the highest levels of the transgene, were analyzed to investigate whether the soluble Pref-1/hFc secreted by the liver was able to affect adipose tissue development. The circulating levels of Pref-1/hFc fusion protein in Alb-Pref-1/hFc mice was similar to that of the 2AS aP2-Pref-1/hFc mice and was approximately 50% of that seen in the 3AS aP2-Pref-1/hFc line (Figures 6a and 1b). Compared with wild-type littermates, 21Alb transgenic mice did not show any change in body weight at birth, but did show a slight tendency to gain less weight during the study period, although the differences were not statistically significant (data not shown). The 21Alb Alb-Pref-1/hFc transgenic mice had smaller fat pad weight when compared with wild-type littermates, with 10- to 14-week-old mice showing significant 33.1, 21.2, and 24.1% reductions in renal, inguinal, and reproductive fat pad weight, respectively (Figure 6b). Moreover, a 12% reduction in brown adipose tissue was detected, although due to the variability it was not statistically significant. The lower fat mass was accompanied by decreased expression of adipocyte markers, including C/EBPα, FAS, SCD, ADSF/resistin, leptin, and adiponectin, indicating an impairment of adipogenesis in Alb-Pref-1/hFc transgenic mice (Figure 6c). The 4Alb Pref-1/hFc transgenic line, which expressed comparable levels of transgene to 21Alb line, was unable to transmit the transgene to the F1 generation; therefore the fat pad weight of the founder was compared with the wild-type littermate. Similar to the 21Alb transgenic line, the 4Alb Pref-1/hFc founder showed a 21% reduction in total fat pad weight as compared with wild-type littermates. Northern blot analysis also revealed a reduction in the expression of adipocyte markers (data not shown). A strong correlation between the

Figure 5
Gene expression in adipose tissue from aP2-Pref-1/hFc transgenic mice. (a) Northern blot analysis of adipocyte marker expression in white adipose tissue of wild-type and 3AS aP2-Pref-1/hFc transgenic mice. Total RNA from three different fat pads (Ing, inguinal; Rep, reproductive; Ren, renal) was probed with cDNA probes for different adipocyte markers. (b) Endogenous Pref-1 expression analyzed by RT-PCR. A primer set was designed to amplify only the endogenous Pref-1 message (see Methods). β-actin was used as an internal control. (c) Northern blot analysis for adipocyte markers in interscapular brown adipose tissue.
expression level of Pref-1/hFc and the degree of decrease in fat pad weight was observed in the various transgenic lines of aP2-Pref-1/hFc and Alb-Pref-1/hFc, regardless of whether the transgene was expressed in adipose tissue or liver (Figure 7). The total fat pad weight showed a 45% reduction in adipose tissue mass in 3AS aP2-Pref-1/hFc transgenic line, the highest expresser of the transgene in adipose tissue. The decrease in total fat pad weight in 2AS aP2-Pref-1/hFc, 4Alb, and 21 Alb Alb-Pref-1/hFc lines was 27, 21, and 25%, respectively. These three lines of transgenic mice, whether the transgene is expressed in adipose tissue or in liver, had similar levels of circulating Pref-1/hFc protein corresponding to approximately 50% of that observed in the 3AS aP2-Pref-1/hFc transgenic mice. The ability of hepatocellular expression of Pref-1/hFc to affect fat pad weight to a similar degree suggests that the large soluble form of Pref-1 may act in an endocrine manner to inhibit adipocyte differentiation. It is also interesting to note that Delta, which was originally reported to function as a membrane form of a ligand, has been demonstrated to release a soluble extracellular domain capable of binding to and acting through Notch (30, 31). However, Pref-1 lacks the DSL domain that is conserved in all Notch ligands and mediates Notch receptor-ligand interaction (10). Thus, it is unlikely that Pref-1 acts as a Notch ligand. Identification of Pref-1–interacting proteins or its receptor will make possible to understand the molecular mechanisms underlying Pref-1 action.

Changes in adipose tissue mass are frequently associated with alterations in glucose homeostasis. We therefore carried out insulin- and glucose-tolerance tests to determine if insulin and glucose homeostasis would be affected by ectopic expression of Pref-1. When glucose-tolerance tests were performed (Figure 8a) on the 3AS aP2-Pref-1/hFc transgenic line, mice
expressing Pref-1/hFc showed higher levels of blood glucose (Table 1), and the glucose clearance rate was slower than in wild-type mice. Insulin-tolerance tests showed that the transgenic mice had overt insulin insensitivity (Figure 8b). Levels of insulin after overnight fasting were higher in transgenic mice than in their wild-type littermates (Table 1). We also measured serum triglyceride levels, since insulin resistance, hyperglycemia, and hyperinsulinemia have all been associated with dyslipidemia (32, 33). Plasma triglyceride levels were higher in transgenic mice than in wild-type mice (Table 1). These data suggest that expression of the Pref-1/hFc transgene and the resulting decrease in adipose tissue mass has promoted development of abnormalities in glucose homeostasis and insulin sensitivity. Increased fasting insulin levels in Pref-1 transgenic mice suggest that the capacity for insulin secretion by islet β cells is probably not impaired. Severe or complete loss of fat mass in the transgenic mice overexpressing dominant negative A-ZIP/F-1 or SREBP-1c in adipose tissue have been reported to accompany hypertriglyceridemia and syndromes similar to type II diabetes (34, 35). In addition to these animal models, studies in human patients with lipodystrophy also have revealed a strong correlation between decrease in adipose tissue mass and insulin resistance (36, 37). Insulin insensitivity in 3AS Pref-1/hFc transgenic mice accompanies the substantial loss of adipose tissue, although the loss is less complete than in lipoatrophic models. It is plausible that the reduced capacity for lipid storage in adipose tissue of these mice may contribute to these metabolic syndromes. Transgenic lines showing lower expression of the Pref-1/hFc fusion protein, 2AS aP2-Pref-1/hFc and 21Alb, 4Alb Alb-Pref-1/hFc, did not show significant differences in glucose clearance rates and insulin resistance (data not shown), indicating that the reduction in adipose tissue mass in those mice was not sufficient to bring about the insulin resistance and glucose intolerance associated with lipoatrophic models. The mechanisms, however, by which changes in adipose tissue mass cause insulin resistance still remain controversial (38). An increasing number of studies suggests an important role of fat-derived hormones in the etiology of insulin resistance associated with altered adipose tissue mass. Indeed, leptin and adiponectin have been reported to reverse insulin resistance associated with lipodystrophy (39, 40). On the other hand, ADSF/resistin, a recently discovered factor secreted by adipocytes (41, 42) has been suggested, although controversially, to contribute to insulin resistance (42). In this regard, it should be noted that the 3AS aP2-Pref-1/hFc transgenic mice exhibited a marked reduction in the expression and secretion of the insulin-sensitizing hormones leptin and adiponectin (Figure 5a and Table 1). The reduced levels of leptin and adiponectin in serum of 3AS aP2-Pref-1/hFc transgenic mice, which correlate with the expression levels in adipose tissue, may have contributed to the observed insulin resistance. The smaller reduction in leptin and adiponectin expression shown in adipose tissue from Alb-Pref-1/hFc transgenic mice may not have been sufficient to bring about the metabolic abnormalities observed in 3AS aP2-Pref-1/hFc transgenic mice. Further studies are needed to clarify the implications of the reduced expression of leptin and adiponectin in the insulin resistance and glucose intolerance observed in our Pref-1 transgenic mice.

Since we observed lower body weight in transgenic mice at weaning, we examined embryonic expression of the Pref-1/hFc transgene driven by the aP2

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### Table 1

<table>
<thead>
<tr>
<th>Serum parameters of 3AS aP2-Pref-1/hFc male transgenic mice and wild-type littermates</th>
<th>Wild-type</th>
<th>3AS aP2-Pref-1/hFc</th>
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<tbody>
<tr>
<td>Fasted glycemia (mg/dl)</td>
<td>66.4 ± 2.7</td>
<td>81.5 ± 7.9^a</td>
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<tr>
<td>Fasted insulin (ng/ml)</td>
<td>0.22 ± 0.04</td>
<td>0.36 ± 0.02^a</td>
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<tr>
<td>Triglycerides (mg/dl)</td>
<td>98.3 ± 3.2</td>
<td>125.9 ± 8.1^a</td>
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<td>Leptin (ng/ml)</td>
<td>2.3 ± 0.2</td>
<td>1.7 ± 0.4</td>
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<tr>
<td>Adiponectin (µg/ml)</td>
<td>12.1 ± 0.6</td>
<td>4.9 ± 2.7^a</td>
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Data are expressed as mean ± SEM (n = 4–5 animals in each group). ^aP < 0.05.
promoter in two transgenic lines. Pref-1/hFc transgene expression was detected at embryo day 12 and was further increased at later stages (Figure 9a). Interestingly, the 3AS aP2-Pref-1/hFc transgenic embryos showed growth retardation and skeletal abnormalities, primarily in the distal vertebra. The two other aP2-Pref-1/hFc transgenic lines also showed similar skeletal abnormalities, with the severity correlated with the levels of circulating Pref-1/hFc (Figure 9, b–d). Recently, pref-1/dlk1 has been reported to be a paternally expressed gene located in a chromosomal region containing six imprinted genes (18, 19, 43, 44). Paternal monoallelic expression of pref-1/dlk1 has been observed in the syntenic chromosomes 12, 18, and 14 in mice, sheep, and humans, respectively (18–20, 43, 44). Although the gene(s) responsible for the phenotype is not known, perturbation of imprinting gene expression in human paternal uniparental disomy (pUPD) 14 and mouse (pUPD) 12 causes growth retardation and bone malformation (45–55). Given our findings of decreased adipose tissue mass in Pref-1/hFc transgenic mice, it is possible that the lean phenotype of clpg sheep is due to the antiadipogenic action of Pref-1.

In conclusion, we provide evidence that forced expression of Pref-1/hFc results in decrease in fat mass, fat cell size, and inhibition of adipogenesis. Moreover, we show that the large cleavage product of Pref-1/hFc transgenic embryos had a smaller thoracic cavity with short ribs. Vertebrae were fused and disorganized, resulting in scoliosis. 

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