Lessons on Conditional Gene Targeting in Mouse Adipose Tissue

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Lessons on Conditional Gene Targeting in Mouse Adipose Tissue

Kevin Y. Lee,1 Steven J. Russell,1 Siegfried Ussar,1 Jeremie Boucher,1 Cecile Vernochet,1 Marcelo A. Mori,1 Graham Smyth,1 Michael Rourk,1 Carly Cederquist,1 Evan D. Rosen,2 Barbara B. Kahn,2 and C. Ronald Kahn1

Conditional gene targeting has been extensively used for in vivo analysis of gene function in adipocyte cell biology but often with debate over the tissue specificity and the efficacy of inactivation. To directly compare the specificity and efficacy of different Cre lines in mediating adipocyte specific recombination, transgenic Cre lines driven by the adipocyte protein 2 (aP2) and adiponectin (Adipoq) gene promoters, as well as a tamoxifen-inducible Cre driven by the aP2 gene promoter (aP2-CreERT2), were bred to the Rosa26R (R26R) reporter. All three Cre lines demonstrated recombination in the brown and white fat pads. Using different floxed loci, the individual Cre lines displayed a range of efficacy to Cre-mediated recombination that ranged from no observable recombination to complete recombination within the fat. The Adipoq-Cre exhibited no observable recombination in any other tissues examined, whereas both aP2-Cre lines resulted in recombination in endothelial cells of the heart and nonendothelial, nonmyocyte cells in the skeletal muscle. In addition, the aP2-Cre line can lead to germline recombination of floxed alleles in ~2% of spermatozoa. Thus, different “adipocyte-specific” Cre lines display different degrees of efficiency and specificity, illustrating important differences that must be taken into account in their use for studying adipocyte biology. Diabetes 62:864–874, 2013

A

diocyte function in vivo has benefited from the development of mouse lines that use the Cre/LoxP site-specific recombination system to inactivate specific genes in fat (3). The use of such targeting systems has allowed researchers to clarify the relative contribution of the adipose tissue in many metabolic phenotypes and circumscribe lethality that might be associated with inactivation of genes at the whole-body level. Several different Cre transgenes have been used for this purpose. The most common use the promoter of the mouse adipocyte protein-2 (aP2) gene, which encodes fatty acid-binding protein-4 (Fabp4). A 5.4-kb piece of the aP2 promoter/enhancer has been shown to be sufficient to direct expression in adipocytes (4,5). At least three independent laboratories have developed aP2-Cre transgenic mice. The first aP2-Cre line was created by Kleanthis Xanthopoulos (6); subsequently, the aP2-CreBI line was created by Barbara Kahn (Beth Israel, Boston, MA) (7), and the aP2-CreBI was created by Ronald Evans (Salk Institute, San Diego, CA) (8). In addition, the aP2 promoter has been used by the Chambon laboratory (Institut de Génétique et Biologie Moléculaire et Cellulaire, Paris, France) to drive the expression of a tamoxifen-inducible Cre transgene (aP2-CreERT2), which is only able to recombine floxed alleles in the presence of 4-hydroxytamoxifen (4-OHT) (9,10).

Although aP2/Fabp4 was originally identified as an adipocyte-specific protein, recent studies have shown that Fabp4 is also expressed in other cell types (11), including macrophages (12–14), the lymphatic system (15), and during embryogenesis (16). To circumvent the possible side effects of gene deletion of the aP2-Cre in tissues other than adipocytes, two laboratories have developed adiponectin-Cre transgenic mice (Adipoq-Cre), with expression of a Cre recombinase driven by the promoter/regulatory regions of the mouse adiponectin locus using a bacterial artificial chromosome (BAC) transgene (17) or by a 5.4-kb promoter fragment (18).

In the current study, we have directly compared the specificity and efficacy of three mouse transgenic Cre lines—the aP2-CreBI, aP2-CreERT2, and Adipoq-Cre BAC transgenic mouse lines—in mediating adipocyte-specific recombination using a number of different floxed alleles as well as by breeding these mice to the LacZ-Gt(ROSA)26Sor-1Bgr (termed R26R-lacZ) reporter mouse, in which Cre-mediated recombination irreversibly activates a lacZ reporter gene (19). We find that all of the Cre lines induce recombination in the adipose tissue. In addition, the aP2-CreBI and aP2-CreERT2 lines both induce recombination in the capillary endothelium in the heart and in intermyofibrillar cells in the skeletal muscle, but not in macrophages in adipose tissue. Interestingly, we find that different floxed gene loci display differential sensitivity to Cre-mediated recombination and that different adipose depots recombine to different extents. The aP2-CreBI can also lead to germline recombination of floxed alleles. These results illustrate the differences between “adipose-specific” Cre lines and caveats in their use that are critical for interpretation of research using these models.

RESEARCH DESIGN AND METHODS

Animals and diets. aP2-CreBI and aP2-CreERT2 mice were maintained on a C57BL/6 background. Adipoq-Cre mice had also been backcrossed to...
C57BL/6; however, single nucleotide polymorphism panel analysis revealed that these mice, although largely C57BL/6, still have markers of a mixed genetic background [http://jaxmice.jax.org/strain/010803.html].

The Cre mice were bred to Gr(Rosa)26Sortm1Wtsc (B6;129S8-Tg(CAG-EGFP)35Ccd/129S8-Tm1Ley), obtained from Jackson Laboratories on the C57BL/6 background. Mice with floxed alleles of insulin receptor (IR), Hif1α, Igf1r, Dicer, Sho2, and Tfam have previously been described (20–25), as have the generation of fat-specific knockouts of GLUT4, peroxisome proliferator-activated receptor-γ (Pparg), IR, Hif1α, Tfam, and Irbg/f1r with the aP2-Cre mouse (7,26–30). The generation of fat-specific knockouts of PTP1B, Pparg-coactivator-1α (Pgc1α), and SH2-containing protein tyrosine phosphatase-2 (Shp2) with the Adipoq-Cre have also been described (31–33). All mice were housed in a mouse facility on a 12-h light/dark cycle in a temperature-controlled room with ad libitum access to water and food.

For tamoxifen treatment of aP2-CreERT2, tamoxifen was dissolved in a solution of 10% ethanol in sunflower seed oil at a concentration of 50 mg/mL. Mice were gavaged with this solution (100 µL) daily for 5 days and killed 2 weeks later. Daily gavage with 4-hydroxytamoxifen (4-OHT) led to a slight decrease (~3 g) in the body weight of mice. Similar recombination efficiencies, without effects on body weight, were observed after three cycles of gavage every other day and killing the animals 2 weeks after the last dose. Longer treatment protocols did not increase deletion efficacy. Other treatment methods led to undesirable complications. Intraperitoneal injection 4-OHT led to excess oil unabsorbed in the abdomen and mild peritonitis, and subcutaneous injection led to the formation of granulomas at the site of injection (data not shown).

Animal care and study protocols were approved by the Joslin Diabetes Center Animal Care Committee and were in accordance with the National Institutes of Health guidelines.

Detection of β-galactosidase activity. Fat pads were whole-mount stained. Other tissues were cut into 10-µm cryosections and stained for β-galactosidase activity, as previously described (34).

Macrophage isolation. Peritoneal macrophages were collected by injecting 5 mL ice-cold PBS supplemented with 1% BSA into the peritoneal cavity and collecting the resultant cell suspension. Fat resident and recruited macrophages were collected by excising and mincing epididymal white adipose tissue into 5 mL with Liberase Thermolysin Medium (21.5 µg/mL, Dicer Roche) and incubated at 37°C for 45 min with shaking. Larger particles were removed using 250-µm nylon sieves and the filtrates were centrifuged at 300g for 5 min to separate floating adipocytes from the stromal-vascular fraction (SVF) pellets. SVF pellets containing macrophages and cell suspensions containing peritoneal macrophages were treated with 100 µL ACK lysis buffer (Lanza) for 5 min at room temperature, and washed with PBS. Macrophages were stained in 100 µL staining buffer (PBS containing 1% BSA with F4/80-APC; E-Bioscience). Cells were again washed, filtered through a 40-µm mesh, and F4/80-positive cells were sorted by FACSAria (BD Biosciences).

Immunofluorescence and immunohistochemistry. Localization of Fabp4 and CD31 was assessed in the fat by whole-mount immunofluorescence using hamster anti-mouse CD31 monoclonal antibodies (1:500; BD Biosciences) and rabbit anti-Fabp4 polyclonal antibodies (1:200; Cell Signaling), antihamster-FTTC (1:1,000; BD Biosciences), and goat anti-rabbit Alexa 564 (1:1,000; Invitrogen). Photographs were generated using an inverted confocal microscope. Colocalization in heart and muscle was assessed using cryosections with the same antibodies. F4/80 localization was assessed using rat anti-F4/80 (1:100; Abcam) and goat anti-rat Alexa 488 (1:1000; Invitrogen).

Quantification of β-galactosidase activity. Total protein lysates were made by homogenizing tissue in M-PER Reagent (Pierce). Five hundred micrograms of concentrated protein lysate was added to 100 µL β-galactosidase

FIG. 1. Analysis of Cre-induced recombination in the aP2-CreR26R-lacZ Mice. A: Whole-mount X-gal staining of fat pads from the aP2-CreR26R-lacZ mouse. Brown, subcutaneous, and perigonal adipose tissue from 3-month-old chow-fed aP2-CreR26R-lacZ (n = 4) after overnight incubation in X-gal staining solution (scale bar = 1 cm). B: Representative sections from aP2-CreR26R-lacZ X-gal stained fat pads. Fat pads from panel A were fixed in Bouin’s fixative and embedded in paraffin; 5-µm sections were hematoxylin stained. Representative images are shown. C: Representative sections from aP2-CreR26R-lacZ tissues stained with X-gal. Lung, heart, testis, and muscle from 3-month-old chow-fed aP2-CreR26R-lacZ (n = 4) were embedded in optimal cutting temperature medium; 10-µm sections were X-gal stained, and counterstained with nuclear fast red. Representative images are shown.
staining solution (34) and incubated overnight. Absorbance was measured at 635 nmol/L.

Assessment of germline recombination. DNA from tail biopsies was extracted and subjected to PCR. In addition to primers that discriminate between floxed and wild-type alleles, an additional primer to assess the presence of recombined floxed alleles (Δ allele) was added. For the IR locus: forward primer 5′-ctgaatagctgagaccacag-3′; reverse primer 5′-gttggaacccctagtcg-3′; delta primer 5′-ctataactgtgctag-3′. For the Tfam locus: forward primer 5′-tgctctctcttgccggcgg-3′; reverse primer 5′-gaaccgcagaactgtg-3′; delta primer 5′-ctctgaagcatggtcata-3′.

RESULTS

aP2-CreBI recombination in adipose tissue, heart, skeletal muscle, and testis. aP2-CreBI mice were bred with R26R-lacZ reporter mice to visualize the tissues in which Cre-mediated recombination occurred. X-gal staining of fat pads of 3-month-old mice revealed strong recombination in the brown adipose tissue (BAT), subcutaneous fat, and perigonadal fat (Fig. 1A). Sections from these fat pads show extensive lacZ staining in adipocytes but no staining in large blood vessels in the subcutaneous fat, as indicated by the arrows (Fig. 1B). This staining was relatively fat-specific, because no staining, or only scattered rare blue nuclei, were observed in other tissues, including the liver, kidney, brain, thymus, adrenal, pancreas, spleen, uterus, ovary, skin, and salivary gland (Supplementary Fig. 1). However, X-gal staining was observed in the heart, lung, and selected nuclei in skeletal muscle (Fig. 1C). Strikingly, strong recombination was also observed in the developing spermatogonium of ~2% of the seminiferous tubules but was completely absent in the other 98% (Fig. 1C, lower left), suggesting clonal germ line recombination occurs during spermatogenesis.

Localization of endogenous Fabp4 expression. When crossed with R26R-lacZ reporter mice, aP2-CreBI mice show recombination limited mostly to the fat pads, but also in the heart and muscle. Quantitative (q)PCR analysis of Fabp4 and Adiponectin mRNA across a wide panel of tissues revealed that both are primarily expressed in fat, with very low expression levels in other tissues (Supplementary Fig. 2). To better define these additional cell types exhibiting recombination in the aP2-CreBI mice, we localized endogenous aP2/Fabp4 in mouse adipose tissue,

FIG. 2. Endogenous expression of Fabp4 in wild-type mice. A: Double immunofluorescence for the endothelial cell markers CD31 and Fabp4 in the perigonadal fat, heart, and muscle of 2-month-old C57BL/6 male mice. Perigonadal fat pads were whole-mount stained and representative composite (original magnification ×10) pictures created from Z-stacks of several images taken by confocal microscopy are shown (scale bar = 50 μm). Images for double immunofluorescence for CD31 and Fabp4 in heart and muscle are representative original magnification ×20 images (scale bar = 200 μm). B: Double immunofluorescence for the macrophage marker, F4/80 and Fabp4 in the perigonadal fat of wild-type mice. Representative images (original magnification ×40) of double immunofluorescence for F4/80 and Fabp4 in the perigonadal fat of 12-month-old wild-type C57BL/6 mice (scale bar = 50 μm). C: qPCR analysis of Fabp4 from sorted F4/80-positive cells from the peritoneal cavity and adipose tissue and whole adipose tissue of male mice (n = 6) after 12 weeks of a high-fat diet (started at age 6 weeks). Data are normalized to the expression of TATA binding protein (Tbp) and are shown as mean ± SEM.
myocardium, and skeletal muscle by double immunofluorescence with anti-Fabp4 and anti-CD31, which stains endothelial cells, or an anti-F4/80 antibody, which marks macrophages (Fig. 2A). Confocal images of the whole-mount stained perigonadal adipose tissue demonstrated only very slight colocalization of CD31 and Fabp4 (Fig. 2A). No colocalization was observed between anti-F4/80 and anti-Fabp4 antibodies in perigonadal fat (Fig. 2B), demonstrating that Fabp4 is not highly expressed in macrophages or endothelial cells. The immunofluorescence demonstrates that Fabp4 expression is limited almost exclusively to adipocytes. Furthermore, qPCR analysis of sorted F4/80 positive macrophages from the peritoneal fluid and perigonadal adipose tissue shows that levels of Fabp4 in macrophages are several thousandfold lower than those found in whole adipose tissue (Fig. 2C).

Colocalization of CD31 and Fabp4 was not observed in skeletal muscle; however, in the heart, Fabp4 did colocalize with CD31 staining in small capillaries. Interestingly, large CD31-positive blood vessels in the heart did not stain positively for Fabp4, indicating that endogenous Fabp4 is expressed only in a specific subset of endothelial cells (Fig. 2A).

**aP2-CreERT2 recombination in adipose tissue, heart, muscle, and salivary gland.** The 4-OHT-inducible Cre driven by the aP2 promoter (aP2-CreERT2) allows for temporal control of recombination (9). In the absence of 4-OHT, no X-gal staining was observed in any tissue in male or female mice (Fig. 3A and C, Supplementary Fig. 3). Upon oral 4-OHT treatment, X-gal staining was observed in the subcutaneous and perigonadal fat pads (Fig. 3A). Sections through these fat pads show limited X-gal staining (Fig. 3B). In agreement with observations from the aP2-CreERT2 mice, recombination was observed in capillaries of the myocardium and in interfibrillar cells in skeletal muscle (Fig. 3C). Unlike the aP2-CreERT2 mice, recombination was also observed in the epithelium of the salivary gland after 4-OHT exposure (Fig. 3C). No staining was observed in any other tissue after 4-OHT treatment (Supplementary Fig. 4).

**Adipoq-Cre recombination in adipose tissue.** Whole-mount X-gal staining of fat pads of 6-week-old Adipoq-Cre mice crossed to R26R-lacZ reporter mice revealed strong recombination in the brown, subcutaneous, and perigonadal fat (Fig. 4A). Upon sectioning, robust lacZ staining can be seen in adipocytes from all depots, with no staining in the blood vessels of the subcutaneous fat (Fig. 4B, arrows). The recombination from the Adipoq-Cre mice was fat-specific, because absolutely no staining was observed in other tissues, including the liver, kidney, brain, thymus, adrenal, pancreas, spleen, lung, testis, salivary gland, ovary, or uterus (Supplementary Fig. 5). In contrast to the aP2-CreERT2 lines, no expression was observed in the myocardium or skeletal muscle; however, some recombination was observed in dermal adipocytes of the skin (Fig. 4C).

**Quantification of Cre expression and recombination of the aP2-CreERT2, aP2-CreERT2, and Adipoq-Cre lines.** Cre mRNA levels from subcutaneous, perigonadal, and brown fat, as well as muscle, heart, and liver, were analyzed in aP2-CreERT2, aP2-CreERT2, and Adipoq-Cre mice using qPCR. In all three fat pads, the aP2-CreERT2 exhibited the highest Cre expression at the mRNA level, with expression also observed in the heart and muscle (Fig. 5A). The aP2-CreERT2 expressed low levels of Cre mRNA in the fat pads and also expressed in the heart and liver. Conversely, in the Adipoq-Cre mice, Cre mRNA

![Fig. 3. Analysis of Cre-induced Recombination in the aP2-CreERT2 R26R-lacZ mice.](image-url)
expression was found exclusively in the fat pads, with no detectable expression in heart, muscle, or liver. Assessment of Cre protein level by Western blot showed that in the subcutaneous and brown fat, Cre protein levels were high both in the aP2-CreBI and Adipoq-Cre, with only minimal expression in the aP2-CreERT2 (Fig. 5B).

Whole-mount β-galactosidase staining demonstrated that the aP2-CreBI and Adipoq-Cre mouse lines effectively recombine the Rosa locus. However, to quantify this recombination, a colorimetric assay was developed using β-galactosidase activity as a readout for the degree of recombination of tissues from aP2-CreBI and Adipoq-Cre lines crossed to the R26R-lacZ reporter mice. Lysates from R26R-lacZ reporter mice alone were used to establish baseline readings. β-Galactosidase activity was found in all fat pads of aP2-CreBI and Adipoq-Cre mice. aP2-CreBI Cre showed a greater extent of recombination in BAT, whereas Adipoq-Cre showed slightly greater recombination in the subcutaneous fat and perigonadal fat. A small amount of β-galactosidase activity over background was observed in the aP2-CreBI in the heart and muscle, but not in liver (Fig. 5C).

Recombination efficiency of the aP2-CreBI is allele- and age-dependent. The aP2-CreBI mouse has been crossed to multiple lines harboring different floxed alleles to generate fat-specific knockout mouse models. In most cases, recombination was adipose-specific as determined by qPCR analysis, which showed no significant recombination in the liver, heart, or brain with any of the floxed alleles. Interestingly, in some cohorts, up to a 60% reduction of IR mRNA was observed in the skeletal muscle of fat-specific insulin receptor knockout (FIRKO) animals, but this was not observed with any of the other genes (Table 1). There was also differential recombination efficiency across different alleles and in different fat pads. This may represent differences in efficacy of recombination or differential expression of the targeted locus in different cells within the fat pad. qPCR analysis showed reductions of Hif1β RNA of 63, 68, and 89% in whole fat perigonadal, subcutaneous, and brown fat compared with floxed controls. In the white fat depots, the lower recombination rate was at least partly due to the presence of Hif1β in other cells of the fat pad. Thus, qPCR analysis of isolated adipocytes revealed 83 and 86% reduction of Hif1β mRNA in perigonadal and subcutaneous adipocytes, respectively. There was also modest reduction of Hif1β in the SVF of the perigonadal and subcutaneous fat of 27 and 40%, suggesting recombination in preadipocytes or early adipocytes present in the fraction.

Recombination of Shox2 in aP2-CreBI mice resulted in reductions of Shox2 mRNA of 48, 58, and 81% in perigonadal, subcutaneous, and brown fat compared with floxed controls, respectively. Recombination of the Ppara locus was also very efficient with the aP2-CreBI mice. Fat-specific ablation of Ppara with the aP2-CreBI led to a complete absence of Ppara mRNA in brown fat. In the white fat pads, mRNA Ppara levels were reduced more than 90% in the subcutaneous white fat and completely ablated in the perigonadal fat pad.
Conversely, using the aP2-Cre line and mice with a floxed Tfam (mitochondrial transcription factor A) allele, there was a 67% reduction at the mRNA level in the brown fat, but no significant reduction of Tfam mRNA was observed in the white fat pads. When the fat pads were digested, there was a significant reduction in Tfam protein expression of adipose tissues from aP2-CreBI and Adipoq-Cre mice. Western blot analysis of subcutaneous, perigonadal, and brown fat from 3- to 5-month-old aP2-CreERT2 (♂ = 4), Adipoq-Cre(♀ = 4), and Adipoq-Cre(♀ = 2) mice. All animals were maintained on a chow diet. Data are represented as a ratio of mRNA expression compared with control mice. mRNA levels were normalized to the expression of glyceraldehyde-3-phosphate dehydrogenase (GAPDH) and are expressed as mean ± SEM.
### Table 2: Protein expression (ratio of protein expression to floxed controls)

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<td></td>
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Recombination efficiency as assessed by Western blot of fat-specific knockout mouse models in fat, isolated adipocytes, the SVF of fat, and other tissues of different floxed alleles. All animals were maintained on chow diet, and data are represented as a ratio of protein expression compared with floxed controls. N/A, not applicable; NC, not changed; ND, not detectable; PG, perigonadal, SC, subcutaneous. *Macrophage expression. †Glut4 is only expressed in adipocytes, and not in other cell types in the adipose tissue. ‡Estimation. §No induction during chronic cold exposure.
efficient ablation with the Adipoq-Cre in whole fat pads, with reductions of 80–90%. In addition, PTP-1B protein was undetectable in isolated adipocytes, but no change was seen in the SVF of fat pads, the liver, or the muscle (Table 2) (21). Efficient ablation by the Adipoq-Cre was also observed with Dicer, with reductions of Dicer mRNA of 70, 61, and 81% in whole perigonadal, subcutaneous, and brown fat compared with floxed controls. Similar reductions of Dicer mRNA was also observed in isolated adipocytes, with 61 and 74% reduction of Dicer mRNA in perigonadal and subcutaneous adipocytes, respectively, with no significant reductions in the SVFs of perigonadal or subcutaneous fat. Efficient ablation using the Adipoq-Cre mouse has also been reported for the floxed alleles of Pgc1α, and Shp2 (31,33). qPCR analysis showed Adipoq-Cre reduced Pgc1α RNA more than 80% in the perigonadal and subcutaneous fat, and more than 90% in brown fat. A >90% reduction of Pgc1α RNA was found in perigonadal adipocytes, with no changes in expression in the liver or the muscle (Table 1). Similarly, no Pgc1α protein was detectable even after chronic cold exposure (Table 2). These data demonstrate specific and efficient recombination of multiple alleles in adipocytes using the Adipoq-Cre mice.

Incidence of germline deletion when using the aP2-Creβi. Mice harboring Cre transgenes have been shown to lead to occasional germline recombination of floxed alleles (38). Because recombination is observed in testis of aP2-Creβi mice, we tested our mice lines for evidence of germline recombination. A PCR-based assessment of recombination was developed for each floxed allele and performed with isolated tail DNA. For the IR locus, a 280-bp amplified product indicates a wild-type exon 4, and a 320-bp product represents an intact floxed exon 4, and a 220-bp product represents a floxed allele after Cre-mediated recombination. The recombinant allele is termed the delta (Δ) allele (Fig. 6A). A similar PCR strategy was developed for Tfam in which a 350-bp product indicates a wild-type allele, a 420-bp product represents an intact Tfam allele with exon 6 and 7 floxed, and a 300-bp product represents the ΔTfam allele created by Cre-mediated recombination (Fig. 6B). No evidence of germline recombination was found with the aP2-CreERT2 (mated before tamoxifen induction) or Adipoq-Cre mouse lines. However, Cre-mediated excision of floxed transgene occurs in a number of mice upon crossing with the aP2-Creβi line. The Δ allele, was detectable in the tail DNA, in the absence of any cell previously demonstrated to express aP2, and

FIG. 6. Germline deletion of floxed alleles when using the aP2-Creβi. A: Evidence of recombination of the IR locus from the aP2-Creβi in DNA from mouse tails. DNA was extracted from tails of progeny of the aP2-Creβi mouse and mice with the floxed IR. The recombination status of the IR locus (from left to right) are IR+/-, IRfl/fl, and IRfl/fl WT, wild-type. B: Evidence of recombination of the Tfam locus with the aP2-Creβi in DNA from mouse tails. DNA was extracted from tails of progeny of the aP2-Creβi mouse and mice with a floxed Tfam locus. The recombination status of the Tfam locus (from left to right) are Tfam+/-, Tfamfl/fl, and Tfamfl/fl WT, wild-type. C: Frequency of the Δ allele is illustrated in progeny of the aP2-Creβi mouse and mice with floxed alleles of IR, Tfam, Dicer, and Hif1β. D: Expression of IR was compared using qPCR in perigonadal fat of 3-month-old male IR+/- (LOX), aP2-Creβi IRfl/fl (FIRKO), IRfl/fl (DELTA), or aP2-Creβi IRfl/fl (DELTA FIRKO) mice. Data are normalized to the expression of Tbp and are shown as mean ± SEM of six to seven samples. E: Expression of IR was compared using qPCR in the liver of 3-month-old male animals as in panel D. Data are normalized to the expression of Tbp and are shown as mean ± SEM of six to seven samples.
the incidence of the Δ allele varies across floxed transgenes, from as low as 1.6% in IR-floxed animals to as high as 14.4% in Tfam-floxed animals (Fig. 6C). Furthermore, qPCR analysis of IR mRNA from tissues of animals possessing a Δ IR allele, shows that the presence of the Δ allele, independently of the presence of the aP2-Cre transgene, leads to an ~60% loss of expression in perigonadal fat, as well as liver and brain (Fig. 6D–F). These results indicate that the aP2-Cre\(^{fl}\) line can lead to germline recombination of the floxed alleles, resulting in whole-body heterozygous knockout mice.

**DISCUSSION**

During the past 2 decades, advances in the use of site-specific recombinases have added greatly to our ability to manipulate cells and gene expression. These site-specific recombinases bind to and recombine specific sequences of DNA, allowing researchers to heritably label cells, conditionally inactivate or activate genes, and even ablate cells based on their gene expression. Thus far, studies examining the in vivo role of specific genes in adipocytes by conditional ablation have largely relied on the adipose-specific expression of the aP2 promoter (4,5), with (9,10) or without (7,8) additional regulation by tamoxifen. Recently, Adipoq-Cre lines were created as a possible alternative model to conditionally ablate genes in adipose tissue (17,18). More than 70 published reports have relied upon aP2-Cre or Adipoq-Cre lines, and although several caveats in their use have been alluded to, thus far, these have not been systematically assessed. In the current study, we have performed a direct comparison and analysis of these fat-specific Cre lines.

The tissue and cell-type specificity of these different mouse transgenic Cre lines was determined by crossing them to the R26R-lacZ reporter mouse and analyzing the tissues from the resultant mice for recombination. Of the three Cre lines analyzed in this study, the Adipoq-Cre showed the most specific expression in white and brown fat. The aP2-Cre\(^{fl}\) R26R-lacZ mice and the aP2-CreERT2 R26R-lacZ mice also showed recombination both in brown and white fat; however, there was nonadipose expression with both aP2-Cre lines. This included expression in the capillaries of the myocardium and in perivascular cells in the skeletal muscle. On the one hand, that two independent aP2-Cre mouse lines gave similar patterns suggests that the recombination in these tissues is due to eutopic expression of Cre recombinase directed by the aP2 promoter. On the other hand, the differences between the aP2-CreERT2 and aP2-Cre\(^{fl}\) mice, including the staining within the testis and salivary gland, may be due to the different sites of transgene integration.

The large degree of recombination in the endothelial cells of the heart observed in the aP2-Cre\(^{fl}\) and the aP2-Cre\(^{fl}\) may explain why fat-specific Dicer-knockout animals created with aP2-Cre exhibit perinatal lethality due to Dicer’s known roles in vessel formation and maintenance (39). However, such cell-type specific loss of gene expression in a tissue of mixed cell types, such as the heart or skeletal muscle, may be difficult to detect by qPCR or Western blot because the other cells in the tissue continue normal expression. Likewise, stochastic recombination in the germ cell lines would be difficult to detect without histological analysis. It is also worth noting that the endogenous Fabp4 expression is observed only in the small capillaries and not in the large vessels of the heart. This specific expression in a distinct subset of endothelial cells suggests developmental differences between endothelial cells. Some studies of the FIRKO mouse have observed a reduction of IR gene expression in skeletal muscle. However, because this reduction is only observed in the FIRKO, this effect may be a secondary effect specific to this line and not due to aP2-Cre deletion.

Although previous reports have shown Fabp4 expression in many other tissues, including the kidney, liver, and skin (11), little or no recombination is observed in the aP2-Cre lines in any of these tissues. The levels of Fabp4 are very low in nonadipose tissue, and the lack of recombination in these tissues may be due to low levels of Cre expression that are below the threshold needed for recombination. Alternatively, the enhancer regions needed for expression in these tissues may be missing from the aP2 promoter fragment used to generate these Cre mice. Previous reports have also shown that Fabp4 is expressed in macrophages (40) within the heart. Here, we demonstrate that macrophages within the adipose tissue do not express significant levels of Fabp4. These findings are confirmed by the absence of recombination that we observe in the macrophages of the aP2-Cre\(^{fl}\).

In addition to differences among the Cre mice, we demonstrate that the different floxed gene loci display a range of sensitivity to recombination when using these different Cre lines. With the aP2-Cre\(^{fl}\), very efficient recombination was observed in the Hif1β, Shox2, Glut4, and IR loci, with far less recombination found in the combination of the IR/IGF1R double knockout loci and the PTP1B and Tfam locus. Several factors may contribute to this range of sensitivity to Cre-mediated recombination. First, different floxed alleles are known to have differential sensitivity to Cre-mediated recombination, with some loci inherently more difficult to recombine (41). Another potential factor that may confound these results is that some of the genes being inactivated may be necessary for adipogenesis or survival. Work from in vitro models demonstrates that ablation of Hif1β (28), Shox2 (unpublished data), or Glut4 (7) does not affect adipogenesis or cellular survival. Similarly, differentiation of preadipocytes that lack IR can be compensated for by expression of IGF1R (30). On the other hand, Dicer appears to be indispensable for adipogenesis (42), and combined loss of IR and IGF1R results in complete failure of adipocyte differentiation (30). Although cells with a knockdown of Tfam differentiate into adipocytes (43) ablation of Tfam leads to increased apoptosis (44–46). Thus, in models that ablate genes necessary for adipogenesis or cellular survival, recombination would appear to be less efficient because the knockout adipocytes never develop. Conversely, genes that are expressed exclusively in the mature adipocytes, including Glut4 (47) and Ppard (48), would have a greater recombination efficiency than those expressed in multiple cell types.

Within the FIRKO animals, we also find that there is an age-dependent increase in recombination efficiency. Several factors may be contributing to the improved recombination. First, as animals age, the cellular composition of the fat pads may shift, including a decrease of preadipocytes (49). This would lead to a relative increase in the proportion of adipocytes compared with other cells, and because the aP2-Cre is active in adipocytes, recombination efficiency would appear to be improved. In addition, transcriptional activity and changes in splicing from the IR locus have been observed during aging (50),...
suggesting that epigenetic modifications to the IR locus occur with age. It is possible that these changes in the IR locus may change its susceptibility to Cre-mediated recombination.

Cre mRNA levels in the fat pads of aP2-Cre line mice are higher than in the aP2-CreERT2 or Adipoq-Cre mice. Because the same 5.4-kb promoter fragment was used in the creation of the aP2-Cre line and aP2-CreERT2 lines, the difference in expression between these two lines is most likely due to differential copy number or to different sites of transgene integration. Conversely, because only one to two copies of a BAC transgene integrate into the genome, the levels of Cre transgene observed in the Adipoq-Cre mice most likely reflect the high endogenous expression of Adipoq mRNA. Despite the lower Cre expression level, the Adipoq-Cre mice demonstrate slightly greater recombination in both the subcutaneous and perigonadal fat. These results suggest that the greater Cre expression in the aP2-Cre line is due to higher expression of Cre per cell and not to a greater number of cells expressing Cre. In fact, the greater recombination efficiency observed in the Rosa locus, as well as in the Tjam, Dicer, and PTP1B loci, suggests that the Adipoq-Cre may direct Cre expression to a more complete population of adipocytes than the aP2-Cre line.

Finally, we have shown that transmission of the aP2-Cre line can lead to germline deletion (Δ) of floxed alleles. The incidence of Δ alleles varies across floxed transgenes and leads to a reduction of gene expression. Strong staining and specific LacZ staining could be seen in ~2% of seminiferous tubules of aP2-Cre line R26R-LacZ testis, leading to populations of recombined spermatid precursors. Because within individual mice the Δ allele can be independent of the presence of Cre, recombination in the germ cells most likely occurs early in spermatogenesis, thus allowing Cre and Δ alleles to segregate during meiosis. Although we did not observe oocytes with LacZ staining, the Δ allele can also arise when the female is the carrier of the Cre transgene, suggesting that germline recombination may also occur in the oocyte. In our studies of using the aP2-Cre, we excluded all animals carrying the Δ allele from further analysis or breeding. However, if such mice are not excluded from the breeding strategy, the Δ allele leads to 100% of Δ-Lox progeny, which is expected to maintain the Δ allele in the progeny. In conclusion, in this study we have analyzed the efficacy and specificity of transgenic Cre lines driven by the aP2 and adiponectin gene promoters. All three lines we examined demonstrated recombination in fat with minimal staining in most other tissues. The aP2-Cre line demonstrated both an allele- and age-dependent sensitivity to Cre-mediated recombination within the fat. Finally, we show that the aP2-Cre line can lead to germline recombination of floxed alleles. These results are the first systematic analysis of these Cre lines that have been widely used to study adipocyte biology and highlight important considerations and implications not only in their use but also in the need for careful characterization of Cre lines in general.

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