The Adipose Tissue Microenvironment Regulates Depot-Specific Adipogenesis in Obesity

Highlights

- High-fat diet feeding induces sex-specific patterns of adipogenesis
- The amount of obesogenic adipogenesis positively correlates with fat pad weight
- Estrogen potentiates obesogenic adipogenesis in the inguinal subcutaneous depot
- Depot-specific adipogenesis is regulated by the adipose tissue microenvironment

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In Brief

Jeffery et al. look at differences in fat depot expansion in male and female mice in response to a high-fat diet. Subcutaneous white adipose tissue (SWAT) obesogenic adipogenesis occurs only in females. Furthermore depot-specific activation and differentiation of white adipocyte precursors are determined by the microenvironment rather than cell-intrinsic differences.
The Adipose Tissue Microenvironment Regulates Depot-Specific Adipogenesis in Obesity

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SUMMARY

The sexually dimorphic distribution of adipose tissue influences the development of obesity-associated pathologies. The accumulation of visceral white adipose tissue (VWAT) that occurs in males is detrimental to metabolic health, while accumulation of subcutaneous adipose tissue (SWAT) seen in females may be protective. Here, we show that adipocyte hyperplasia contributes directly to the differential fat distribution between the sexes. In male mice, high-fat diet (HFD) induces adipogenesis specifically in VWAT, while in females HFD induces adipogenesis in both VWAT and SWAT in a sex hormone-dependent manner. We also show that the activation of adipocyte precursors (APs), which drives adipocyte hyperplasia in obesity, is regulated by the adipose depot microenvironment and not by cell-intrinsic mechanisms. These findings indicate that APs are plastic cells, which respond to both local and systemic signals that influence their differentiation potential independent of depot origin. Therefore, depot-specific AP niches coordinate adipose tissue growth and distribution.

INTRODUCTION

The increasing rates of obesity and obesity-associated pathologies around the world highlight the need for an understanding of the mechanisms that underlie white adipose tissue (WAT) growth in vivo. The distribution of WAT into subcutaneous (SWAT) and visceral (VWAT) depots in obesity has links to metabolic disease. Increased VWAT mass is strongly associated with cardiometabolic risk and mortality across ethnicities (Coutinho et al., 2011; Lee et al., 2014; Nazare et al., 2012; Phillips and Prins, 2008; Pischon et al., 2008; Wormser et al., 2011). In contrast, elevated SWAT mass has been associated with improvements in plasma lipid profile, insulin sensitivity, blood pressure, and atherosclerosis (Appleton et al., 2013; Heitmann and Lissner, 2011; Manolopoulos et al., 2010; Snijder et al., 2003). These data suggest that the mechanisms underlying depot-specific WAT growth are directly relevant to metabolic health and disease, yet the regulation of this process in vivo is not well understood.

Subcutaneous and visceral adipose depots display an array of differences including gene expression (Cohen et al., 2014; Grove et al., 2010; Karastergiou et al., 2013), developmental lineage (Chau et al., 2014; Krueger et al., 2014; Sanchez-Gurachines and Guertin, 2014), metabolic characteristics (Cohen et al., 2014; Karastergiou and Fried, 2013; Tran et al., 2008), and adipokine secretion profiles (Shi et al., 2009). Interestingly, males preferentially accumulate visceral adipose tissue, while premenopausal females accumulate more subcutaneous adipose tissue (Palmer and Clegg, 2015). In humans, these patterns of adipose distribution result in characteristically female “pear-shaped” obesity and male “apple-shaped” obesity (Gesta et al., 2007; Karastergiou et al., 2012). In postmenopausal women, adipose distribution shifts toward a male-like pattern, implicating sex hormones in maintaining a balance between subcutaneous and visceral depot mass (Palmer and Clegg, 2015). Importantly, however, the cellular and molecular mechanisms underlying differential adipose distribution in males and females are unclear.

Adipose tissue growth can occur through two distinct mechanisms: the increase in size of existing adipocytes (hypertrophy) or the increase in number of adipocytes (hyperplasia). Since mature adipocytes are post-mitotic, new adipocytes are derived from adipocyte precursor (AP) cells within the stroma of all white adipose depots (Berry et al., 2014). It was recently shown that the expansion of visceral adipose tissue in males occurs through adipocyte hyperplasia in both mice (Jeffery et al., 2015; Wang et al., 2013) and humans (Armer et al., 2013), while male subcutaneous adipose tissue in mice does not exhibit significant hyperplasia in response to obesogenic stimuli (Jeffery et al., 2015; Wang et al., 2013). Furthermore, studies of the dynamics of human adipose tissue indicate that weight loss is mediated primarily by a reduction in adipocyte size with no significant reduction in adipocyte number, even after bariatric surgery (Spalding et al., 2008). These data suggest that an increase in adipocyte number in obesity may have lasting effects on energy homeostasis and weight maintenance. Taken together, these findings indicate that depot-specific mechanisms are important in the control of WAT growth in obesity; however, the mechanisms driving these processes are not known.

We recently found that in male mice, obesogenic signals such as high-fat feeding or hyperphagia lead to the proliferation and
Figure 1. Adipocyte Hyperplasia Contributes to Depot- and Sex-Specific WAT Growth in Obesity

(A) Body fat as a percentage of total body weight after 8 weeks of SD or HFD feeding in male and female C57Bl/6J Adiponectin-creER; mTmG mice as measured by magnetic resonance imaging (n = 9–15).

(B) Weight of individual fat pads from male or female C57Bl/6J Adiponectin-creER; mTmG mice after 8 weeks of SD or HFD feeding (n = 9–15).

(C) Experimental design for Adiponectin-creER; mTmG adipogenesis experiments.

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obesity, we measured whole body fat mass (Figure 1A) and fat display similar phenotypes in the high-fat diet (HFD) model of males accumulate more fat mass than females (Figure 1A), and contribution to metabolic disease (Baglioni et al., 2012; Fried et al., 2015; Loh et al., 2015; Macotela et al., 2012). However, the mechanisms that control depot-specific AP activation in vivo have not been investigated. Here, we show that obese-genic adipogenesis in SWAT is differentially regulated in males and females, explaining the differences in WAT distribution between the sexes. Furthermore, through AP transplant studies, we find that depot-specific proliferation and differentiation of APs is determined by the depot microenvironment and not by cell-intrinsic differences in APs.

RESULTS

Adipocyte Hyperplasia Contributes to Depot- and Sex-Specific WAT Growth in Obesity

Sex-specific patterns of adipose distribution have been well documented in humans (Gesta et al., 2007; Karastergiou et al., 2012; Palmer and Clegg, 2015). To determine if C57Bl6/J mice display similar phenotypes in the high-fat diet (HFD) model of obesity, we measured whole body fat mass (Figure 1A) and fat pad weight of the perigonadal VWAT and the inguinal SWAT depots (Figure 1B) after 8 weeks of HFD or standard diet (SD) feeding. Although the precise anatomic location of these depots in mice differ from humans, these mouse depots impact metabolism similarly to human visceral adipose tissues (Barzilai et al., 1999; Foster et al., 2010, 2011; Gabriely and Barzilai, 2003) and gluteofemoral adipose tissue (Foster et al., 2013; Tran et al., 2008), respectively. We find that upon HFD feeding, males accumulate more fat mass than females (Figure 1A), consistent with previous reports (Grove et al., 2010; Medrikova et al., 2012). Interestingly, in both SD- and HFD-fed conditions, male VWAT is significantly larger than male SWAT, whereas these two depots are maintained at similar sizes in females (Figure 1B). These data indicate that, similar to humans, mice are sexually dimorphic with regard to fat mass distribution in obesity.

We recently showed that HFD feeding induces adipocyte hyperplasia in male VWAT, but not male SWAT, after 8 weeks of HFD feeding (Jeffery et al., 2015), contributing to the preferential growth of VWAT in males. To determine whether hyperplastic mechanisms influence sex-specific WAT growth, we used the Adiponectin-creER; mTmG mouse model to track the formation of new mature adipocytes in vivo in response to SD or HFD feeding. In this model, new adipocytes that form after tamoxifen injection are mTomato+ (mT), while pre-existing adipocytes will remain mGFP+ (mG) (Jeffery et al., 2014, 2015). Notably, our data indicate that the low dose of 50 mg/kg tamoxifen does not significantly affect body weight (Figure S1C). Consistent with previous findings, we observe VWAT-specific adipogenesis in males after 8 weeks of HFD feeding (Figure 1D). Interestingly, in females we find that HFD significantly increases adipogenesis in both VWAT and SWAT depots (Figure 1E), mirroring the similar expansion of these depots in response to HFD (Figure 1B). We also compared the average size of the newly formed mTomato+ adipocytes to the existing mGFP+ adipocytes within each depot after 8 weeks of diet and found no significant differences (Figures 1F and 1G), which suggests the newly formed cells contribute significantly to fat mass.

We next directly assessed the relationship between adipocyte formation and WAT depot mass. Indeed, comparison of depot size to adipocyte hyperplasia in individual animals shows that in male VWAT, female VWAT, and female SWAT depots, adipocyte formation is significantly correlated with fat pad size after 8 weeks of diet (Figures 2A–2D). Furthermore, whole-body fat mass is significantly correlated with adipogenesis within these depots (Figure S2). Conversely, there is no correlation between adipocyte formation and either fat pad mass or body fat mass in the non-adipogenic male SWAT depot (Figures 2B and S2B). These data indicate that adipocyte hyperplasia significantly contributes to depot- and sex-specific WAT growth in obesity.

The production of new adipocytes in vivo occurs through the proliferation and differentiation of APs within the adipose tissue stroma (Jeffery et al., 2015; Rodeheffer et al., 2008). In males, HFD feeding induces the transient proliferation of APs specifically in VWAT during the first week of HFD feeding, followed by adipogenesis within this depot (Jeffery et al., 2015). To determine whether the same mechanism of transient AP activation also occurs in female mice upon HFD feeding, we assessed the incorporation of bromodeoxyuridine (BrdU) into APs via flow cytometry (Figure 2E). We find that in both the female SWAT and VWAT depots, HFD feeding induces rapid and transient proliferation of APs (Figures 2F, S2E, and S2F), similar to male VWAT (Jeffery et al., 2015). Furthermore, analysis of mature adipocyte nuclei after a 1-week BrdU pulse and 7-week chase indicates that HFD induces significant adipocyte formation from proliferative precursors in both the SWAT and VWAT depots of female mice (Figure 2G), whereas males on HFD display increased BrdU-positive adipocyte formation in VWAT (Jeffery et al., 2015), but not SWAT (Figure S2G).

Interestingly, after ovariectomy females display a more male-like pattern of AP activation in response to HFD, with significantly reduced AP proliferation in SWAT (Figure 2H), which suggests that sex hormones play a role in the activation of SWAT APs. To test whether estrogen influences depot-specific AP activation, we administered estrogen to male mice via mini osmotic pumps and assessed their AP proliferation response to HFD feeding. We find that estrogen treatment of males results in

(D and E) Representative confocal images (left) and quantification (right) of adipocyte labeling in the indicated depots of Adiponectin-creER; mTmG male (D) or female (E) mice (males: n = 4–6; females: n = 12–15).
(F and G) Average adipocyte diameter of tdTomato+ (new) and eGFP+ (old) adipocytes from male (F) and female (G) mice (n = 4–6). Note: all data from (A)–(E) are from tamoxifen-treated animals. Significance was determined by comparing the indicated groups using an unpaired two-tailed Student’s t test. Scale bar is 100 µM. Error bars represent mean ± SEM. See also Figure S1.
Figure 2. Fat Pad Weight and Adipocyte Precursor Activation Mirror Depot-Specific Patterns of Adipogenesis

(A–D) Correlation between adipogenesis and fat pad weight in the male VWAT (A) and SWAT (B) depots and the female VWAT (C) and SWAT (D) depots. Each point represents one mouse (n = 21–26).

(E) Representative flow cytometry plots to measure BrdU incorporation into adipocyte precursors.

(F) BrdU incorporation into APs from the indicated depot of female mice following 1 week of SD or HFD and BrdU treatment (n = 5 per group).

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significantly increased AP proliferation in SWAT upon HFD feeding (Figure 2I), suggesting that estrogen influences the cellular response of the SWAT depot to diet. Taken together, these data show that adipocyte hyperplasia, mediated by the proliferation and differentiation of APs, is an important component of depot- and sex-specific WAT growth in obesity.

The Adipose Microenvironment Determines Adipocyte Precursor Activation

We next investigated the mechanisms regulating depot-specific AP activation. Previous in vitro studies suggest that APs from SWAT and WAT exhibit cell-intrinsic differences that control their response to environmental cues and differentiation potential (Baglioni et al., 2012; Loh et al., 2015; Macotela et al., 2012). To determine whether cell-intrinsic phenomena explain depot-specific AP activation in vivo, we performed a series of AP transplant experiments (Figure 3A). In these experiments, APs were isolated from the WAT or SWAT of tdTomato-expressing mice by fluorescence-activated cell sorting (FACS) and then transplanted into either the WAT or SWAT of congenic wild-type recipient mice. After recovery, the recipient mice were placed on either HFD or SD for 1 week, and the proliferation of tdTomato+ donor APs and tdTomato−endogenous APs was determined by BrdU incorporation. Flow cytometry analysis of APs demonstrates engraftment of tdTomato+ donor APs in recipient depots and permits the clear distinction between donor and recipient AP populations (Figures 3B, 3C, and S3A, and S3B). In recipient mice fed SD, neither donor nor endogenous APs are activated in either depot, indicating that transplantation alone does not induce AP proliferation (Figures 3D and 3E). In recipient mice fed HFD, APs derived from WAT depots display high levels of proliferation when injected back into the WAT depot, similar to endogenous cells and APs in the uninjected contralateral depot, supporting the ability of the transplanted cells to be activated by obesogenic stimuli (Figure 3D). Surprisingly, SWAT-derived donor AP cells are also induced to proliferate after HFD feeding when cells are placed in the WAT microenvironment (Figure 3D), but neither WAT- nor SWAT-derived APs proliferate significantly when injected back into SWAT (Figure 3E).

Furthermore, when male SWAT APs are injected into female SWAT, their proliferation is induced by HFD (Figure S3C), despite lower levels of engraftment (35%, 11/31) and increased background proliferation, which may be caused by sex-mismatched transplants (Volk et al., 2016; Weiss et al., 2009). These data indicate that despite their separate developmental lineages (Chau et al., 2014; Krueger et al., 2014), the depot-specific response of APs to HFD feeding is determined by the WAT depot microenvironment, not by cell-intrinsic properties of APs within distinct depots.

The depot-specific regulation of both AP proliferation (Figures 2 and 3) (Jeffery et al., 2015) and adipogenesis (Figure 1) suggests that AP differentiation is also controlled by the adipose depot microenvironment. To test this possibility, we performed transplant experiments with APs derived from the SWAT of male mice expressing luciferase under the control of the mature adipocyte-specific leptin promoter (Figure S3A) (Rodeheffer et al., 2008), as well as the mT/mG cassette (leptin-luciferase; mT/mG mice). These mice allow both non-invasive assessment of adipocyte formation via luciferase activity and direct observation of transplanted cells via tdTomato fluorescence. Importantly, male mice with this genetic background (FVB/NJ) still display a WAT-specific AP proliferation pattern in response to HFD feeding (Figure S4B). Leptin-luciferase; mT/mG SWAT APs were transplanted into the WAT or SWAT of congenic wild-type male recipient mice, and after recovery, mice were placed on HFD for 8 weeks. Luciferase signal was monitored after transplant of APs (baseline) and after 8 weeks of HFD feeding (Figure 4A). Excitingly, we find that upon HFD feeding, SWAT-derived APs display significantly increased adipogenesis when transplanted into the WAT or SWAT of congenic wild-type male recipient mice, and after recovery, mice were placed on HFD for 8 weeks. These data show that adipocyte hyperplasia, mediated by the proliferative and differentiative effects on APs (baseline) and after 8 weeks of HFD feeding (Figure 4A).

DISCUSSION

Taken together, these data show that AP proliferation in response to HFD is regulated by cell-extrinsic factors in the depot microenvironment. In the inguinal SWAT depot, systemic levels of estrogen influence this response. The finding that APs from different WAT depots are functionally interchangeable in vivo suggests that developmental lineage does not irreversibly determine depot-specific AP behavior. Rather, APs are functionally plastic cells that are influenced by their physiologic context.

We employed the Adiponectin-creER model to study WAT dynamics, which utilizes tamoxifen treatment for cre activation and cell tracing. While higher doses of tamoxifen have detrimental effects on WAT (Ye et al., 2015), reports are conflicting about how the lower tamoxifen dose used here affects WAT (Hesselbarth et al., 2015; Liu et al., 2015). To control for the potential effects of tamoxifen treatment in the Adiponectin-creER experiments, both SD and HFD groups were treated with tamoxifen. Furthermore, if the lower doses of tamoxifen used here had long-term effects on creER activation, as has been shown for higher doses (Ye et al., 2015), we would not detect any adipocyte formation in our lineage-tracing experiments. We also show similar patterns of depot-specific adipocyte formation using the tamoxifen-independent BrdU tracing assay, demonstrating that the trends in adipogenesis that we report are not an artifact of tamoxifen treatment.

(G) BrdU incorporation into adipocyte nuclei as measured via immunofluorescence in paraffin sections following 1 week of BrdU treatment (pulse) and 7 more weeks of the indicated diet (chase) (n = 8–10).

(H and I) BrdU incorporation into APs from the indicated depot of ovariecotomized female (H) or estrogen-treated male (I) mice following 1 week of SD or HFD and BrdU treatment (n = 5–10).

Significance in (A)–(D) was determined using Spearman’s non-parametric two-tailed correlation analysis. Significance in (F)–(I) was determined by comparing the indicated groups using an unpaired two-tailed Student’s t test. Error bars represent mean ± SEM. See also Figure S2.
Similar to our previous studies of male mice (Jeffery et al., 2015), the BrdU- and cre-based techniques detect similar depot-specific trends in adipogenesis but result in different quantitative measurements of adipocyte formation, with the BrdU method consistently displaying a much lower percentage of new cells formed. For example, in the VWAT of HFD-fed females, the Adiponectin-creER method identifies 35.8% ± 1.8% of adipocytes as newly formed, while the Brdu method only labels 4% ± 0.81% of mature adipocytes. This discrepancy is explained by WAT biology, as the cre-based lineage tracing method assesses the contribution of both post-mitotic preadipocytes and proliferative adipocyte progenitors, while the BrdU technique only assesses the contribution of proliferative cells. We have previously shown that within the WAT stromal vascular fraction, the abundant CD24− preadipocytes are post-mitotic, while the comparatively rare CD24+ adipocyte progenitors proliferate in response to obesogenic stimuli, but rapidly lose CD24 expression upon activation to generate and replenish the CD24−/CD24− preadipocytes (Berry and Rodeheffer, 2013; Jeffery et al., 2015). This large pool of post-mitotic preadipocytes can thus contribute to adipocyte formation without re-entering the cell cycle and incorporating BrdU. As a result, the cre-based method assesses the adipogenic capacity of both CD24+ and CD24− adipocyte precursor populations combined, while the BrdU assay only quantifies the contribution of the CD24+ adipocyte progenitors within the limited time window of the experiment. Therefore, the elevated assessment of adipogenesis in the cre model compared to the BrdU assay is accounted for by
the differentiation of the large pool of committed, post-mitotic preadipocytes that were formed from the proliferative CD24+ adipocyte progenitors prior to the initiation of the experiment. The clear links between depot- and sex-specific adipogenesis, AP activation, and fat pad weight demonstrate the important role that AP regulation and adipogenesis play in obesogenic...
WAT growth. Future studies aimed to define these regulatory signals and compare their function in adipogenic and non-adipogenic adipose tissue microenvironments may lead to novel approaches to control adipose tissue distribution in a manner that favors metabolically beneficial SWAT growth over detrimental VWAT growth.

EXPERIMENTAL PROCEDURES

Animals
All animal studies followed guidelines issued by Yale University’s Institutional Animal Care and Use Committee (IACUC). See Supplemental Experimental Procedures for details on mouse strains. VWAT refers to the perigonadal WAT and SWAT refers to inguinal WAT in mice. For BrdU experiments, mice were given 0.4–0.8 mg/ml BrdU in drinking water, refreshed every 2 days. High-fat diet is from Research Diets (D12492). Standard diet is from Harlan Laboratories (2018S). Analysis of body composition was performed by Echo MRI (Echo Medical System). For estrogen treatment, 8-week-old male mice were implanted with mini osmotic pumps (Alzet 1004) delivering cyclodextran-coated estrogen (Sigma E4389) at a dose of 2 μg/kg/day in PBS. Mice were allowed to recover for 2 weeks after pump implantation prior to experiment initiation.

Transplant Assay
0.5–1 million APs were isolated from the indicated depot of male mice and injected into the indicated pad of 3- to 6-week-old congenic wild-type mice. After recovery, mice were placed on HFD or SD and treated with BrdU for 1 week and then sacrificed or maintained on diet for 8 weeks prior to analysis. Left and right adipose tissue depots were excised and analyzed separately via flow cytometry, microscopy, or luminescence (see Supplemental Experimental Procedures).

Flow Cytometry
Flow cytometry preparations were performed as described in Jeffery et al. (2015) for BrdU analysis and Berry and Rodeheffer (2013) for cell sorting.

Confocal Microscopy
The mT/mG quantification experiments were performed as described in Jeffery et al. (2015), starting the 50 mg/kg tamoxifen treatments at 8 weeks of age. See the Supplemental Experimental Procedures for further details. For analysis of leptin-luciferase; tdTomato+ adipocytes following transplantation, tissue from the luminescent region of the VWAT was dissected and stained with HCS LipidTOX Green Neutral Lipid Stain (Invitrogen, H34475, used at 1-100) for at least 30 min before being washed in PBS and mounted onto slides in Fluormount-G (Southern Biotech; 0100-01).

Statistical Analysis
Statistical tests used are indicated in the figure legends. All statistical tests were performed using GraphPad Prism v6.02 for Windows (GraphPad Software). Data are expressed as mean ± SEM, and p < 0.05 was considered statistically significant. Outliers were identified with the ROUT method test using GraphPad Prism software, setting Q to 10%. Sample sizes for main figures are listed in Table S1 and indicate individual animals (biological replicates).

SUPPLEMENTAL INFORMATION
Supplemental Information includes Supplemental Experimental Procedures, four figures, and one table and can be found with this article online at http://dx.doi.org/10.1016/j.cmet.2016.05.012.

AUTHOR CONTRIBUTIONS
E.J. and M.S.R. designed experiments. R.B. generated the FVB/NJ; Leptin-luciferase; mT/mG mice. E.J., A.W., L.C., and C.D.C. performed experiments. E.J., A.W., B.H., J.L.K., R.S.-P., and Z.S. analyzed data. E.J., A.W., and M.S.R. interpreted data and wrote the manuscript.

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REFERENCES


Supplemental Information

The Adipose Tissue Microenvironment Regulates
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Figure S1 (Related to Figure 1) *Adiponectin-creER* targets nearly all adipocytes in female subcutaneous and visceral depots.

(A) Representative confocal images of the indicated depots from *Adiponectin-creER; mTmG* mice before (left) and after (right) tamoxifen treatment. (B) Quantification of GFP+ (targeted) adipocytes in the indicated depots following tamoxifen treatment. (n = 3 in each group). (C) Body weight measurements from female mice after treatment with tamoxifen or vehicle as performed in the lineage tracing experiments in Figure 1. (n = 5 in each group). Error bars represent mean ± S.E.M. mTmG: membrane-Tomato, membrane-GFP.
Figure S2 (Related to Figure 2) Adipogenesis in high-fat diet-responsive depots correlates with whole body fat mass.

(A-D) Correlation between whole body fat mass and adipogenesis in the indicated depots of male (A-B) or female (C-D) mice. Each point represents one mouse. (n = 21-26 each). (E-F) BrdU incorporation into adipocyte precursors at the indicated time points of diet treatment, with BrdU pulse during only the last week before sacrifice. (n = 5 in each group) (G) BrdU incorporation into adipocyte nuclei as measured by immunofluorescence in paraffin sections following one week of BrdU treatment (pulse) and 7 more weeks of the indicated diet (chase). (n = 10 in each group). Significance in A-D was determined using Spearman’s non-parametric two-tailed correlation analysis. Significance in E-F was determined using a two-way ANOVA with Bonferroni’s test for multiple comparisons. Significance in G was determined using an unpaired two-tailed student’s t-test. Error bars represent mean ± S.E.M. SD: standard diet, HFD: high-fat diet, AP: adipocyte precursor, BrdU: bromodeoxyuridine.
Figure S3. (Related to Figure 3) Characterization of adipocyte precursors after transplantation.

(A-B) Representative flow cytometry plots of cells from VWAT (A) or SWAT (B) in blue, with overlayed plots of transplanted APs in red recovered at the experimental end point. Percentages indicate the percentage of transplanted APs that fall into the indicated gate. Characteristic marker profile of APs is CD45-, CD31-, CD34+, CD29+, Sca1+.

(C) BrdU incorporation into donor APs derived from male SWAT and transplanted into female SWAT following one week of SD or HFD and BrdU. (SD n = 7, HFD n = 4). **p<0.01. Significance was determined using an unpaired two-tailed student’s t-test. Error bars represent mean ± S.E.M. SD: standard diet, HFD: high-fat diet, AP: adipocyte precursor, BrdU: bromodeoxyuridine.
Figure S4. (Related to Figure 4) The male subcutaneous adipose depot does not induce adipogenesis of transplanted adipocyte precursors upon high-fat feeding.

(A) Experimental model explaining the mature adipocyte-specific leptin-luciferase mouse model. (B) BrdU incorporation into APs from the indicated depots of adult male FVB/NJ mice following one week of SD or HFD and BrdU. (n = 5 per group) (C) Quantification of luminescent signal at the indicated experimental time points following transplant of APs from the SWAT of leptin-luciferase; mTmG mice. (n = 5) (D) Confocal images of LipidTox+ adipocytes derived from transplanted tdTomato+ APs in SWAT at baseline before starting HFD feeding, demonstrating engraftment of transplanted cells. (E) Representative flow cytometry plot of APs from SWAT following transplant and 8 weeks of HFD feeding, demonstrating that tdTomato+ donor cells are still present within the depot. Significance was determined using an unpaired two-tailed student’s t-test. Error bars represent mean ± S.E.M. Scale bar is 100μM. HFD: high-fat diet, AP: adipocyte precursor, ns: not significant.
Table S1 (Related to Experimental Procedures)

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Table S1: (Related to Experimental Methods) Experiment sample sizes.
Sample sizes are listed for individual groups in the main figure panels.
Supplemental Experimental Procedures

Animals

All mice used for these studies were on the C57BL/6J genetic background, except leptin-luciferase; tdTomato mice, and corresponding wild type recipient mice, which are on the FVB/NJ background. Adiponectin-creER mice were a generous gift from Dr. Evan Rosen (Beth Israel Deaconess Medical Center, Boston, MA) and can now be purchased at Jackson Laboratories (stock #024671). The mTmG mice were purchased from Jackson laboratories (stock #007676). Adult wild type mice were purchased from Jackson Laboratories and experiments were performed beginning at 6-8 weeks of age unless otherwise noted. Leptin-luciferase BAC transgenic mice were originally generated on the C57Bl/6 background (Birsoy et al., 2008) were reproduced by cryoinjection into FVB/NJ strain (Rodeheffer et al., 2008). C57Bl/6 mT/mG mice were backcrossed into the FVB/NJ background for >9 generations, then crossed to leptin-luciferase mice to generate leptin-luciferase; mTmG (also referred to herein as leptin-luciferase; tdTomato) mice isogenic with the FVB/NJ strain. Ovariectomized female mice were purchased from Jackson Laboratories.

Transplant assay

Isolation of APs was performed as described (Berry and Rodeheffer, 2013) from the indicated depot of male C57Bl6/J mT/mG (Figure 3) or FVB/NJ leptin-luciferase; tdTomato (Figure 4 and S2) mice and pooled from multiple animals to isolate sufficient numbers of APs. Mice were anesthetized with isoflurane and surgeries were performed using sterile technique. 0.5-1 million APs from the indicated depot were re-suspended in 10 µl of sterile PBS and injected into the left visceral or subcutaneous pad of 3-4 week old (Figure 3) or 5-6 week old (Figure 4) congenic wild type mice. Mice were allowed to recover for 2-3 weeks, then placed on HFD or SD and treated with BrdU for 1 week prior to sacrifice (Figure 3) or maintained on HFD for 8 weeks (Figure 4). Left and right adipose tissue depots were excised and analyzed separately. Preparation of stromal cells for BrdU analysis was performed as described (Jeffery et al., 2015), and transplanted cells were identified by Tomato fluorescence. For AP proliferation experiments, results were counted only for transplants in which more than 200 individual Tomato-positive donor AP cells were recovered in the recipient depot stromal vascular fraction. For transplant of male cells into male mice, this resulted in a 61% (11/18) success rate for injections of VWAT-derived cells into VWAT, a 67% (4/6) success rate for SWAT-derived cells into SWAT, a 78% (7/9) success rate for SWAT-derived cells into SWAT, and 29% (4/14) success rate for VWAT-derived cells into SWAT. For transplant of male cells into female SWAT, the success rate was 35% (11/31). To visualize luminescence in leptin-luciferase; tdTomato transplants, mice were intraperitoneally injected with 100µl of 15 mg/mL luciferin (GoldBio, Cat# LUCK-100), the signal was allowed to develop for 15 min, the mice were anesthetized with isoflurane, and the signal was detected with an IVIS Spectrum imaging system (Caliper Lifesciences).

Confocal microscopy

For mT/mG quantification experiments, starting at 8 weeks of age, mice were treated with 50mg/kg tamoxifen (Sigma) dissolved in vegetable oil by intraperitoneal injection for 5 consecutive days, and then allowed to recover for 1 week. Mice were then sacrificed for baseline analysis, or placed on HFD or remained on SD for the 8-week chase period. Ingual subcutaneous and perigonadal visceral tissue were taken from several regions throughout the depot, and analyzed by whole mount confocal microscopy for tdTomato and eGFP expression. For each data point, at least 500 adipocytes were counted from multiple images from each depot of each animal.

For adipocyte diameter measurements, the area of each adipocyte (in square pixels) was measured using ImageJ. The diameter of each adipocyte was calculated using the measured area, assuming each adipocyte is a perfect circle. At least 300 adipocytes were measured for each data point. Analysis of adipocyte nuclei was performed as described previously (Jeffery et al., 2015). 110-230 adipocyte nuclei were scored for each data point. Images were taken on a Leica TCS SP5 confocal microscope.

Supplemental References