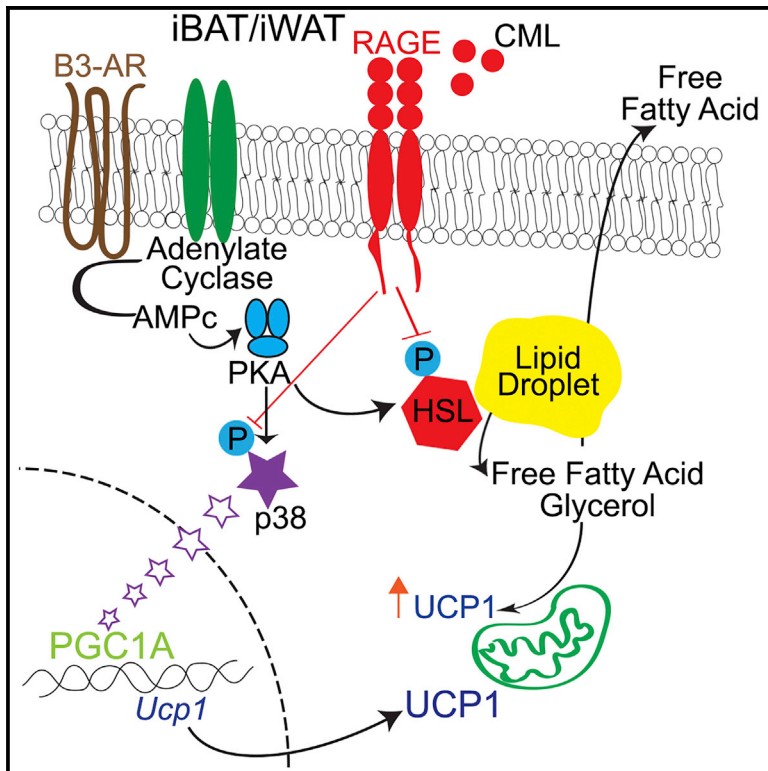


A Receptor of the Immunoglobulin Superfamily Regulates Adaptive Thermogenesis

Graphical Abstract



Authors

Carmen Hurtado del Pozo, Henry H. Ruiz, Lakshmi Arivazhagan, ..., Richard A. Friedman, Ravichandran Ramasamy, Ann Marie Schmidt

Correspondence

annmarie.schmidt@nyulangone.org

In Brief

Hurtado del Pozo et al. show that the deletion of adipocyte RAGE, whose ligands accumulate in metabolic stress, protects from obesity and cold challenges through the modulation of protein kinase A activities. This work adds RAGE to the immunometabolic networks that regulate energy expenditure in environmental and metabolic stress.

Highlights

- Ligands of RAGE accumulate in obese murine and human adipose tissues
- Adipocyte-specific RAGE deletion protects mice from obesity and cold stress
- RAGE suppresses key protein kinase A activities to restrain adaptive thermogenesis
- Transplantation of adipocyte-RAGE-deleted adipose tissue into mice prevents obesity



A Receptor of the Immunoglobulin Superfamily Regulates Adaptive Thermogenesis

Carmen Hurtado del Pozo,^{1,5} Henry H. Ruiz,^{1,5} Lakshmi Arivazhagan,^{1,6} Juan Francisco Aranda,^{1,6} Cynthia Shim,¹ Peter Daya,¹ Julia Derk,¹ Michael MacLean,¹ Meilun He,¹ Laura Frye,¹ Randall H. Friedline,² Hye Lim Noh,² Jason K. Kim,^{2,3} Richard A. Friedman,⁴ Ravichandran Ramasamy,¹ and Ann Marie Schmidt^{1,7,*}

¹Diabetes Research Program, Division of Endocrinology, Diabetes and Metabolism, Department of Medicine, NYU School of Medicine, 435 East 30th Street, New York, NY 10016, USA

²Program in Molecular Medicine, Department of Medicine, University of Massachusetts Medical School, 368 Plantation Street, Albert Sherman Center, Worcester, MA 01605, USA

³Division of Endocrinology, Metabolism and Diabetes, Department of Medicine, University of Massachusetts Medical School, 368 Plantation Street, Albert Sherman Center, Worcester, MA 01605, USA

⁴Biomedical Informatics Shared Resource, Herbert Irving Comprehensive Cancer Center, and Department of Biomedical Informatics, Columbia University Irving Medical Center, New York, NY 10032, USA

⁵These authors contributed equally

⁶These authors contributed equally

⁷Lead Contact

*Correspondence: annmarie.schmidt@nyulangone.org

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SUMMARY

Exquisite regulation of energy homeostasis protects from nutrient deprivation but causes metabolic dysfunction upon nutrient excess. In human and murine adipose tissue, the accumulation of ligands of the receptor for advanced glycation end products (RAGE) accompanies obesity, implicating this receptor in energy metabolism. Here, we demonstrate that mice bearing global- or adipocyte-specific deletion of *Ager*, the gene encoding RAGE, display superior metabolic recovery after fasting, a cold challenge, or high-fat feeding. The RAGE-dependent mechanisms were traced to suppression of protein kinase A (PKA)-mediated phosphorylation of its key targets, hormone-sensitive lipase and p38 mitogen-activated protein kinase, upon β -adrenergic receptor stimulation—processes that dampen the expression and activity of uncoupling protein 1 (UCP1) and thermogenic programs. This work identifies the innate role of RAGE as a key node in the immunometabolic networks that control responses to nutrient supply and cold challenges, and it unveils opportunities to harness energy expenditure in environmental and metabolic stress.

INTRODUCTION

The epidemic of obesity and its metabolic sequelae profoundly impact human health and longevity (Finkelstein et al., 2009; Fleegal et al., 2012). Maintaining a precise balance between energy intake and expenditure is essential for an organism's ability to store or utilize nutrients. Yet, hoarding energy is a double-edged

sword, albeit, salutary in nutrient deprivation; in over-feeding, excess energy storage confers susceptibility to obesity and type 2 diabetes.

White adipose tissue (WAT) and brown adipose tissue (BAT) in mammals store energy in the form of triglycerides and release fatty acids and glycerol in response to catecholaminergic stimulation by sympathetic nerves (Bartness et al., 2010). Whereas WAT liberates fatty acids into the circulation, BAT preferentially oxidizes fatty acids and dissipates energy through uncoupled respiration and the production of heat (Cannon and Nedergaard, 2001; Collins, 2012). The recent identification of BAT in adult humans has rekindled an interest in harnessing brown fat to spur cellular energy expenditure (Nedergaard et al., 2007; van Marken Lichtenbelt et al., 2009; Cypess et al., 2009; Virtanen et al., 2009; Zingaretti et al., 2009). The physiologic process in which energy is dissipated in the form of heat in response to stressors such as cold challenges or over-feeding is termed “adaptive thermogenesis” and is dependent on the expression and activity of uncoupling protein 1 (UCP1) in brown adipocytes (Kopecky et al., 1995). Catecholamines stimulate β -adrenergic receptors (ARs), culminating in increased activity of protein kinase A (PKA) and leading to phosphorylation of adipocyte lipases, such as hormone-sensitive lipase (HSL) and lipolysis, and of p38 mitogen-activated protein kinase (MAPK) and regulation of genes linked to lipid oxidation in brown adipocytes, including *Ucp1* and *Ppargc1a* (peroxisome proliferator-activated receptor γ [PPAR γ]-coactivator 1 α) (Collins, 2012; Yehuda-Shnaidman et al., 2010). Though a process classically ascribed to BAT, during prolonged starvation or a cold challenge, white adipocytes undergo “beiging” or “browning,” in which WAT assumes increased UCP1-expressing adipocytes bearing thermogenic capacity (Kajimura et al., 2015; Lee et al., 2015b). Despite the profound importance of such metabolic plasticity, the natural “brakes” in adipose tissues—the best described of which is insulin (Burns et al., 1979; Jungas and Ball, 1963, 1964)—remain incompletely understood.



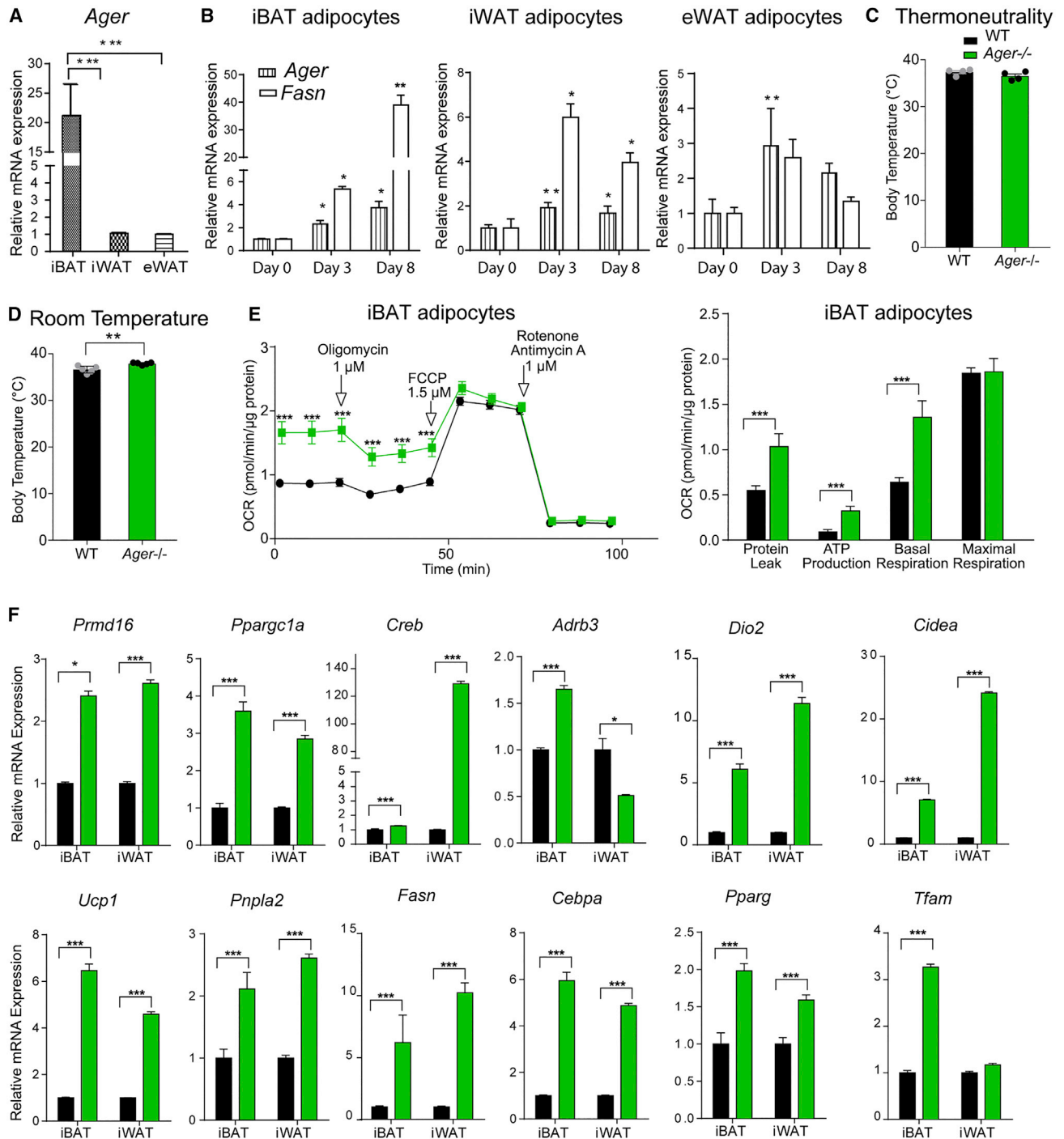


Figure 1. RAGE Is Expressed in Adipose Tissue and Affects Thermogenic Gene Programs

(A) qRT-PCR for relative *Ager* mRNA expression was performed in iBAT, iWAT, and eWAT retrieved from 12-week-old male wild-type (WT) C57BL/6 mice fed standard chow.

(B) Primary adipocytes from iBAT, iWAT, and eWAT of WT mice fed standard chow were subjected to qRT-PCR for detection of relative *Ager* and *Fasn* mRNA expression on day 0, 3, and 8 of differentiation.

(C and D) Core body temperature of 8-week-old male WT or *Ager* null mice determined in mice housed at (C) thermoneutrality (30°C) for 24 h and (D) at room temperature (~23°C).

(E) Oxygen consumption rates (OCRs) normalized to total protein were determined in primary adipocytes differentiated from iBAT from WT and *Ager* null mice. The means ± SEM are reported from five technical replicates with three biological replicates per group.

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The immunoglobulin superfamily molecule, the receptor for advanced glycation end (RAGE) products, binds to a distinct repertoire of molecules, such as the carboxymethyllysine (CML)-advanced glycation end products (AGEs), high-mobility group box 1 (HMGB1), and S100/calgranulins, which accumulate in metabolic stress (López-Díez et al., 2016; Ramasamy et al., 2012). Although these ligands are classically linked to diabetes and inflammation, recent evidence places these molecules and RAGE itself in human and murine obese adipose tissue (Gaens et al., 2014; Song et al., 2014). RAGE localization in human obese adipose tissue aligns with our recent finding that mice globally devoid of *Ager* (the gene encoding RAGE) are protected from obesity and insulin resistance when fed a high-fat diet (HFD) compared to wild-type (WT) mice (Song et al., 2014). Yet, the mediating mechanisms have remained elusive.

Here, we show that mice bearing adipocyte-specific deletion of *Ager* display significant protection from HFD-induced obesity and insulin resistance and exhibit a superior ability to thermoregulate during a cold challenge, compared to mice in which adipocytes express *Ager*. WT mice that underwent a surgical transplantation of either interscapular BAT (iBAT) or subcutaneous inguinal WAT (iWAT) from mice with adipocyte-specific deletion of *Ager* demonstrate significant protection from HFD-induced obesity and insulin resistance, compared to WT mice transplanted with iBAT or iWAT in which adipocytes expressed *Ager*. We traced these adipocyte-intrinsic, RAGE-dependent mechanisms to RAGE ligand-mediated suppression of PKA-dependent phosphorylation of HSL and p38 MAPK—processes that, collectively, dampen UCP1 and thermogenic programs.

RESULTS

RAGE Is Expressed in Mature Adipocytes

We determined the expression patterns of RAGE in WAT and BAT in WT mice fed standard chow and found that *Ager* is expressed in BAT and WAT. Compared to epididymal adipose tissue (eWAT) or iWAT, a significantly higher expression of *Ager* mRNA was observed in iBAT (Figure 1A). In the iWAT and eWAT depots, a significantly higher expression of *Ager* mRNA transcripts was observed in the floating adipocytes than in the stromal vascular fraction (SVF) (Figure S1A). When preadipocytes from the SVF of iBAT, iWAT, and eWAT were differentiated into adipocytes, in each depot, a significantly higher expression of *Ager* mRNA was noted on days 3 or 8 of differentiation versus day 0, which paralleled time-dependent increases in *Fasn*, a marker for adipocyte differentiation (Figure 1B). There were no discernible differences in the morphology of iBAT-, iWAT-, or eWAT-derived primary adipocytes or in the neutral lipid content on day 7 of differentiation from eWAT-derived adipocytes (Figures S1B and S1C). Thus, increasing *Ager* expression accom-

panies, but is not required for, the differentiation of primary adipocytes from iBAT, iWAT, and eWAT.

Deletion of *Ager* Increases Adipocyte Thermogenic Gene Programs and Mitochondrial Activity

Although core body temperature at thermoneutrality (30°C) did not differ between the WT and *Ager* null mice (Figure 1C), at room temperature, mice devoid of *Ager* displayed a significantly higher core body temperature than the WT mice (Figure 1D). Body weight, adiposity, and plasma norepinephrine levels did not differ between the *Ager* null and the WT mice fed standard chow (Figures S1D–S1H).

As the highest expression of *Ager* in the adipose depots was in iBAT, we probed the effect of *Ager* deletion on mitochondrial function in iBAT-derived primary adipocytes from mice fed standard chow. Basal respiration rates and ATP production were significantly higher in adipocytes derived from *Ager* null than from WT mice (Figure 1E). The adipocytes derived from *Ager* null iBAT exhibited pronounced mitochondrial activity, as assessed by MitoTracker Red CMXRos (Koh et al., 2009) (Figure S1I). These findings suggest that RAGE contributes to the regulation of thermogenic programs in iBAT and to browning or beigeing in iWAT.

To address this point, we retrieved adipose tissues from WT mice and mice globally devoid of *Ager* fed standard chow at room temperature. *Ager* deletion resulted in a significantly higher expression of thermogenic genes (*Prdm16*, *Ppargc1a*, *Creb*, *Dio2*, *Cidea*, and *Ucp1*) in iBAT and iWAT. Transcript levels of *Adrb3* were significantly higher in iBAT, but not iWAT, of *Ager* null than of WT mice (Figure 1F). Regarding lipolysis (*Pnpla2*) and lipogenesis (*Fasn*, *Cebpa*, and *Pparg*), mRNA levels of these genes were significantly higher in *Ager* null than in WT iBAT and iWAT. In relation to mitochondrial properties, levels of *Tfam* in iBAT (Lidell et al., 2011), but not iWAT, were significantly higher in *Ager* null than in WT mice (Figure 1F), thus suggesting that mitochondrial biogenesis was higher in iBAT of *Ager* null than of WT mice.

Although eWAT is not primarily a thermogenic adipose tissue depot, we nevertheless examined the above markers. In eWAT, the levels of *Creb*, *Dio2*, *Fasn*, *Cebpa*, and *Tfam* were significantly higher in *Ager* null than in WT; there were no differences in *Cidea*, *Ucp1*, *Pnpla2*, and *Pparg*; and the levels of *Prdm16*, *Ppargc1a*, and *Adrb3* were significantly lower in the eWAT of *Ager* null than of WT mice (Figure S1J).

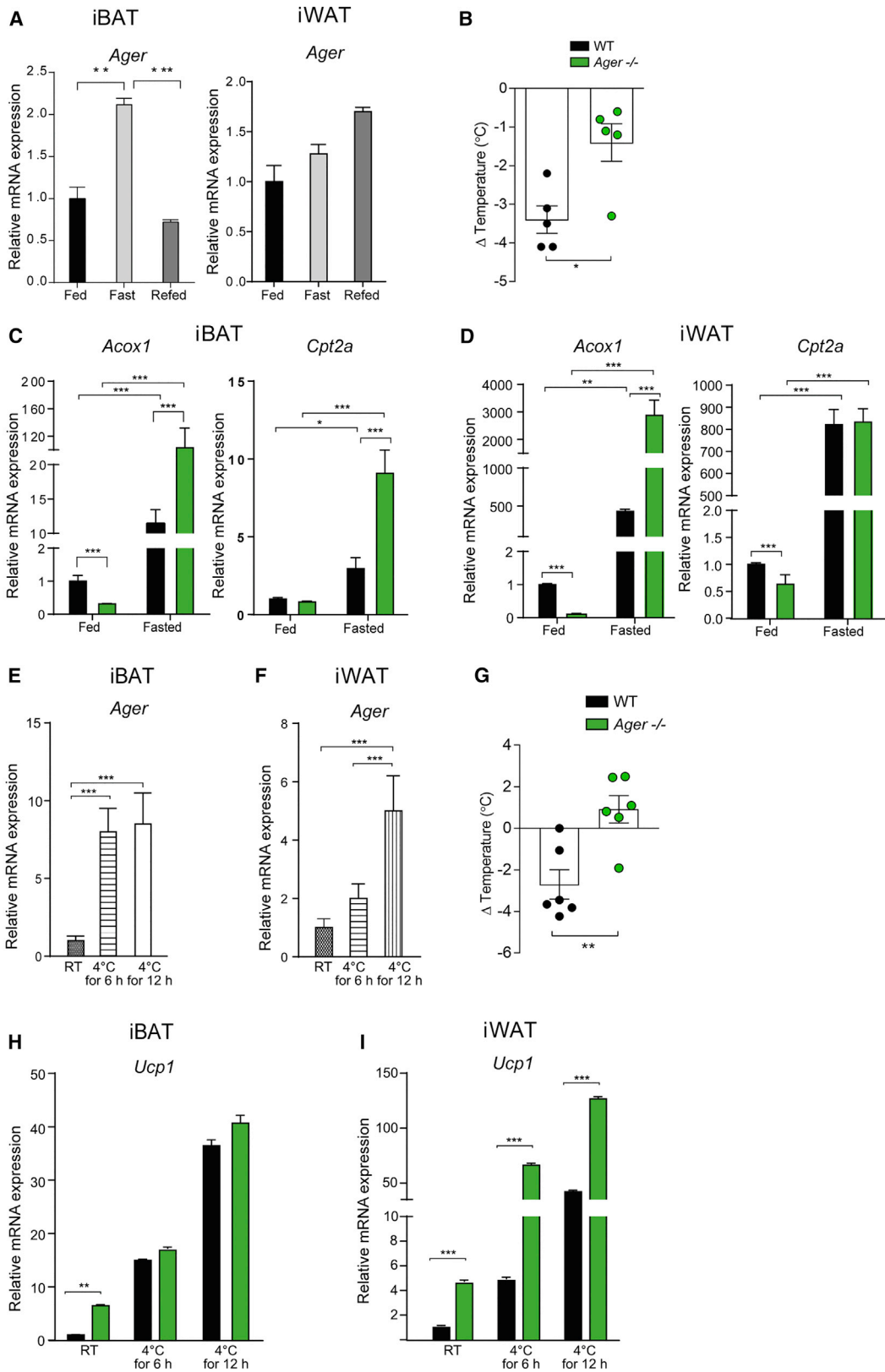
RAGE Dampens the Adaptive Response to Fasting and Cold Challenges

Our findings thus far indicate that the deletion of *Ager* upregulates adipocyte thermogenic gene programs in iBAT and iWAT. In adipocytes, β -adrenergic signals upregulate *Ucp1* transcription, and, in part through lipolysis, the released fatty acids are

(F) iBAT and iWAT retrieved from 8-week-old WT or *Ager* null mice housed at room temperature were analyzed via qRT-PCR for detection of relative mRNA expression of *Prdm16*, *Ppargc1a*, *Creb*, *Adrb3*, *Dio2*, *Cidea*, *Ucp1*, *Pnpla2*, *Fasn*, *Cebpa*, *Pparg*, and *Tfam*.

Unless otherwise stated, the data are presented as mean \pm SEM in $N = 4-5$ mice/group. Data analysis: (A and B) independent samples one-way ANOVA followed by a post hoc Tukey's HSD test; (C and D) two-tailed Student's t test; and (E, repeated measures, and F, independent samples) two-way ANOVA followed by post hoc Sidak test; * $p < 0.05$; ** $p < 0.01$, and *** $p < 0.001$.

See also Figure S1.



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proposed to enhance UCP1 activity in the mitochondria (Himms-Hagen et al., 2000; Rial et al., 1983). We tested these concepts in settings in which adaptive thermogenic programs are engaged.

First, we used a fasting-refeeding paradigm. After 24 h of fasting, WT mice displayed a significantly higher expression of *Ager* in iBAT, but not in iWAT. In iBAT, after 24 h of refeeding, the expression of *Ager* was significantly lower after fasting and returned to baseline (fed) levels (Figure 2A). Similar expression patterns were observed in eWAT under these conditions (Figure S2A). After 24 h of fasting, mice globally devoid of *Ager* displayed significantly less loss of body temperature than the WT mice did (Figure 2B). We examined the expression of *Acox1* and *Cpt2a*, genes that are associated with fatty acid beta oxidation. In iBAT, levels of *Acox1* and *Cpt2a* mRNA were significantly higher in the fasted *Ager* null than in the WT mice (Figure 2C), and in iWAT, significantly higher levels of *Acox1* mRNA, but not *Cpt2a* mRNA, were observed in fasted mice devoid of *Ager* than in the WT mice (Figure 2D). In eWAT, there were no genotype-dependent differences in the fed or the fasted states (Figure S2B).

As an alternative paradigm to induce lipolysis and thermogenic programs, we performed a cold stress (Duncan et al., 2007). Compared to room temperature, the cold challenge resulted in a significantly increased expression of *Ager* in iBAT after 6 or 12 h at 4°C (Figure 2E). In iWAT and eWAT, *Ager* expression was significantly higher after 12 h of exposure to 4°C, compared to room temperature (Figures 2F and S2C). To assess whether upregulation of *Ager* in iBAT and iWAT in mice subjected to the 4°C cold challenge affected thermoregulation *in vivo*, we assessed their core body temperature after 48 h of exposure to 4°C. Mice devoid of *Ager* exhibited significantly less loss of body temperature than did WT mice over 48 h (Figure 2G). In iBAT, at 6 or 12 h of exposure to 4°C, levels of *Ucp1* mRNA did not differ between *Ager* null and WT mice; at room temperature, levels of *Ucp1* mRNA transcripts were significantly higher in *Ager* null than in WT iBAT (Figure 2H). In iWAT, levels of *Ucp1* mRNA were significantly higher in *Ager* null mice at room temperature and after 6 or 12 h in 4°C (Figure 2I). In eWAT, levels of *Ucp1* were significantly lower in *Ager* null than in WT mice at room temperature and after 6 or 12 h at the 4°C cold challenge (Figure S2D). Thus, deletion of *Ager* protects against the loss of body temperature in mice subjected to the 4°C environment, which was accompanied by RAGE-dependent regulation of *Ucp1* in iWAT.

Adipocyte-Specific Deletion of *Ager* Protects from Diet-Induced Obesity and Cold Challenge

In addition to adipocytes, immune and vascular cells and sympathetic fibers populate adipose tissue and might contribute to the impact of RAGE expression. To definitively probe for roles for RAGE in adipocyte-intrinsic responses to the HFD and cold challenge, we generated *Ager* floxed mice (Figures S3A and S3B). *Ager*^{fllox/fllox} mice were bred with adiponectin (*Adipoq*) Cre-recombinase mice to yield *Ager*^{fllox/fllox} *Adipoq* Cre (+) (adipocyte-specific *Ager*-deleted mice) and littermate *Ager*^{fllox/fllox} *Adipoq* Cre (-) (*Ager* WT) mice. This resulted in the deletion of *Ager* from iWAT, eWAT, and iBAT, but not from the pancreas, muscle, and liver (Figure S3C). Hereafter, these animals will be referred to as *Ager*^{fllox/fllox} Cre (+) and *Ager*^{fllox/fllox} Cre (-) mice.

We phenotyped male littermate *Ager*^{fllox/fllox} Cre (+) and *Ager*^{fllox/fllox} Cre (-) mice at baseline while they were fed a standard chow diet and then after 6 weeks of HFD feeding (60% kcal/fat). At baseline, body weight and composition, average food intake, physical activity, and energy expenditure rates were indistinguishable between *Ager*^{fllox/fllox} Cre (+) (green lines) and *Ager*^{fllox/fllox} Cre (-) (red lines) mice (Figures 3A–3E). However, when fed a HFD, *Ager*^{fllox/fllox} Cre (+) mice weighed significantly less than Cre (-) mice (Figure 3A). Whole body fat mass, measured using ¹H-MRS, was significantly lower in the *Ager*^{fllox/fllox} Cre (+) than in Cre (-) mice after 3 and 6 weeks of the HFD (Figure 3B). Average food consumption did not differ by genotype, except after 6 weeks of the HFD, in which lower food consumption was noted in *Ager*^{fllox/fllox} Cre (+) than in Cre (-) mice (Figure 3C). No genotype-dependent differences in physical activity were noted at baseline or after 3 and 6 weeks of the HFD (Figure 3D). *Ager*^{fllox/fllox} Cre (+) mice displayed significantly higher energy expenditure rates than Cre (-) mice after 3 weeks of the HFD, with trends in the same direction at 6 weeks of the HFD (Figure 3E). No differences in glucose-tolerance testing (GTT) were observed after 12 weeks of the HFD (Figure 3F). *Ager*^{fllox/fllox} Cre (+) mice exhibited significantly greater insulin sensitivity than Cre (-) mice during an insulin-tolerance test (ITT) after 12 weeks of the HFD (Figures 3G and S3D).

We determined if a modulation of adipose inflammation accompanied the metabolic protection observed in *Ager*^{fllox/fllox} Cre (+) mice after the HFD. In eWAT, but not iWAT, mRNA levels of *Emr1* (F4/80) and *Ccl2* were significantly lower in *Ager*^{fllox/fllox} Cre (+) than in Cre (-) mice (Figure S3E). No differences in *Tnfa*

Figure 2. Effects of RAGE on Thermogenic Gene Expression during Fasting and Cold Challenge

(A–D) Fasting and refeeding in 11-week-old male WT and *Ager* null mice.

(A) *Ager* expression in iBAT and iWAT from WT mice in a fed, fasted (24 h) or 24-h refed state.

(B) Change in core body temperature, [(Δ) temperature], between the fasted and the refed states was measured in WT or *Ager* null male mice housed at room temperature (RT).

(C and D) Relative mRNA expression of *Acox1* and *Cpt2a* was measured by qRT-PCR in (C) iBAT and (D) iWAT from fed and fasted (24 h) WT and *Ager* null mice.

(E–I) Eight-week-old male WT or *Ager* null mice were exposed to a cold challenge (4°C).

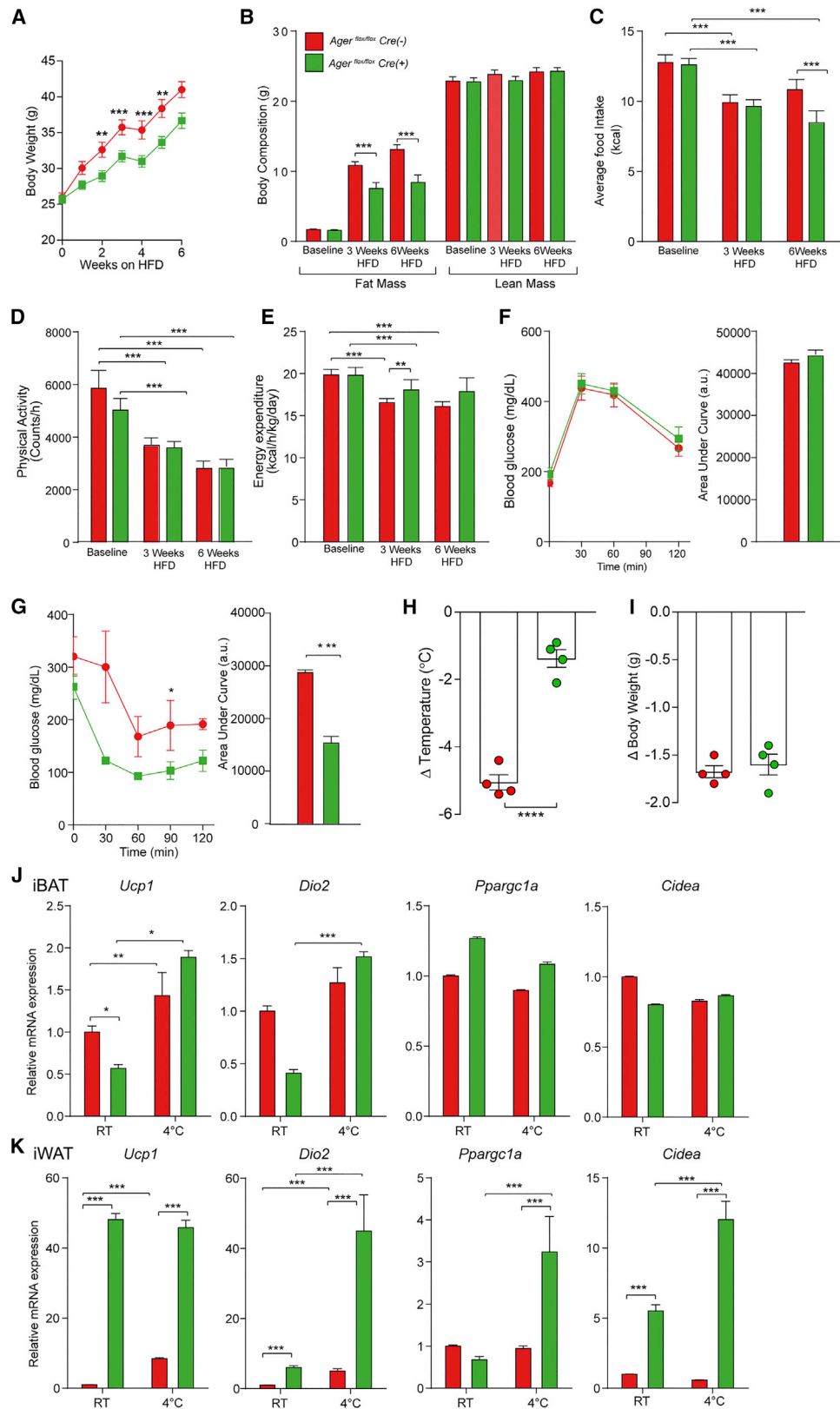
(E and F) iBAT (E) and iWAT (F) from WT mice were subjected to qRT-PCR for relative *Ager* mRNA expression.

(G) (Δ) Temperature from baseline (room temperature) and after 48 h at 4°C is shown.

(H and I) WT or *Ager* null mice were housed at room temperature (~23°C) or exposed to a 4°C cold challenge for 6 or 12 h. iBAT (H) and iWAT (I) were dissected after and subjected to qRT-PCR for detection of relative *Ucp1* mRNA expression.

In (B)–(D) and (G)–(I): WT (black bars) and *Ager* null (green bars) mice. Data are presented as mean ± SEM in N = 4–6 mice/group. Data analysis: (A, E, and F) one-way ANOVA with post hoc Tukey's HSD test; (B and G) two-tailed Student's *t* test; (C, D, H, and I) two-way ANOVA followed by a post hoc Sidak test or Tukey's test, as appropriate; **p* < 0.05; ***p* < 0.01, and ****p* < 0.001.

See also Figure S2.



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or *I110* expression were observed in eWAT or iWAT between the genotypes (Figure S3E). Transcript levels for *Irf4*, which increases adipocyte lipolysis (Eguchi et al., 2011), were significantly higher in the eWAT, iWAT, and iBAT of *Ager^{flox/flox} Cre (+)* mice than *Cre (-)* mice (Figure S3E). *Tlr4*, which has been linked to endotoxin-mediated stimulation of lipolysis in adipocytes (Zu et al., 2009), was significantly higher in eWAT, iWAT, and iBAT of *Ager^{flox/flox} Cre (+)* than *Cre (-)* mice (Figure S3E), and levels of *Tlr2* mRNA were significantly higher in eWAT and iWAT, not iBAT, of *Ager^{flox/flox} Cre (+)* than of *Cre (-)* mice (Figure S3E). Thus, decreased adipose inflammation accompanied the observed protection in *Ager^{flox/flox} Cre (+)* mice from diet-induced obesity and insulin intolerance.

We next exposed littermate *Ager^{flox/flox} Cre (+)* and *Cre (-)* mice to a cold challenge. Compared to *Ager^{flox/flox} Cre (-)* mice, *Cre (+)* mice exhibited significant protection from a loss of body temperature after 24 h of exposure to 4°C (Figure 3H), with no genotype-dependent differences observed in body weight changes (Figure 3I). We examined the expression of genes associated with thermogenic responses in iBAT and iWAT after cold exposure. In iBAT, there were no genotype-dependent differences in *Ucp1*, *Dio2*, *Ppargc1a*, or *Cidea* during the cold challenge, and only significantly lower *Ucp1* was observed in *Ager^{flox/flox} Cre (+)* at baseline (Figure 3J). In iWAT, after 24 h of 4°C, levels of *Ucp1*, *Dio2*, *Ppargc1a*, and *Cidea* mRNA were significantly higher in *Ager^{flox/flox} Cre (+)* than in *Cre (-)* mice (Figure 3K). In eWAT, levels of *Ucp1* were significantly higher in *Ager^{flox/flox} Cre (+)* than in *Cre (-)* mice at 4°C; no genotype-dependent differences were observed in *Ppargc1a* mRNA at 4°C but the levels of *Ucp1* and *Ppargc1a* were significantly higher at room temperature in the *Ager^{flox/flox} Cre (+)* than the *Cre (-)* mice (Figure S3F). These data support adipocyte-intrinsic roles for RAGE in adaptive thermogenic gene program responses after the HFD and 4°C cold challenge.

Transplantation of iBAT or iWAT from *Ager^{flox/flox} Cre (+)* Mice into WT Mice Reduces Body Weight Gain and Upregulates Thermogenic Gene Programs in High-Fat Feeding

As the *Adipoq* Cre-recombinase strategy employed above affected levels of *Ager* in both brown and white adipocytes, the relative contribution of each depot could not be discerned. To address this point, we performed an adipose tissue surgical

transplantation of iBAT bearing adipocyte-specific deleted *Ager* or adipocyte-expressing *Ager* into lean, young, WT mice. Figure S4A is a representative image demonstrating a well-vascularized, non-necrotic iBAT graft. After 4–5 days' recovery from surgery, mice were switched to a HFD and were monitored over 20 weeks. Beginning at 4 weeks post-HFD, body weights in WT mice receiving *Ager^{flox/flox} Cre (+)* iBAT were significantly lower than those of mice receiving *Ager^{flox/flox} Cre (-)* iBAT, an observation that continued and was magnified throughout the HFD feeding period (Figures 4A and 4B). Fat mass was significantly lower in the WT mice receiving *Ager^{flox/flox} Cre (+)* versus *Cre (-)* iBAT after 14 weeks of the HFD (Figure 4C). Although there were no differences in food intake or physical activity between the *Ager^{flox/flox} Cre (+)* and *Cre (-)* mice iBAT donors (Figures 4D and 4E), energy expenditure was significantly higher in the mice receiving *Ager^{flox/flox} Cre (+)* than those receiving *Cre (-)* iBAT after 16 weeks of the HFD (Figure 4F). After 12 weeks of the HFD, WT mice receiving *Ager^{flox/flox} Cre (+)* iBAT were more glucose tolerant than mice receiving *Cre (-)* iBAT (Figure 4G). Insulin sensitivity was improved in mice receiving the *Ager^{flox/flox} Cre (+)* than the *Cre (-)* iBAT (Figures 4H and S4B).

We retrieved native iBAT and iWAT, transplanted iBAT (tiBAT) and eWAT, and assessed the expression of thermogenic genes after 20 weeks of a HFD. In native iBAT, the levels of *Dio2*, *Cpt2a*, and *Pnpla2* mRNA levels were significantly higher in WT mice receiving *Ager^{flox/flox} Cre (+)* than those receiving *Cre (-)* iBAT, while levels of *Ucp1* and *Ppargc1a* mRNA were generally, but not significantly, higher (Figure 4I). Immunohistochemistry revealed that the native iBAT of mice receiving *Ager^{flox/flox} Cre (+)* versus *Cre (-)* iBAT transplants displayed a significantly higher UCP1-positive area (Figure 4J). In native iWAT, there were no significant differences in levels of *Ucp1*, *Dio2*, *Ppargc1a*, *Cpt2a*, or *Pnpla2* mRNA in WT mice receiving *Ager^{flox/flox} Cre (+)* versus *Cre (-)* iBAT transplants (Figure 4K). However, immunostaining revealed that the native iWAT of mice that were transplanted with *Ager^{flox/flox} Cre (+)* iBAT displayed significantly higher UCP1-positive areas than mice that had been transplanted with *Cre (-)* iBAT (Figure 4L). In the transplanted iBAT, levels of *Ucp1*, *Dio2*, *Ppargc1a*, *Cpt2a*, and *Pnpla2* mRNA were significantly higher in transplanted iBAT originating from *Ager^{flox/flox} Cre (+)* than from *Cre (-)* mice (Figure 4M). In native eWAT, levels of *Ucp1*, *Ppargc1a*, *Cpt2a*, and *Pnpla2* mRNA transcripts were significantly higher in the mice receiving *Ager^{flox/flox}*

Figure 3. Adipocyte-Specific Deletion of *Ager* Affects HFD-Induced Obesity and Thermoregulatory Responses to a Cold Challenge

(A–G) Eight-week-old male *Ager^{flox/flox} Adipoq Cre (-)* (red) (*Ager^{flox/flox} Cre (-)*) and *Ager^{flox/flox} Adipoq Cre (+)* (green) (*Ager^{flox/flox} Cre (+)*) mice were fed a HFD (60% kcal/fat).

(A) Body weight gain after up to 6 weeks of high-fat feeding was recorded.

(B–E) Body composition (B), average food intake (C), physical activity (D), and energy expenditure (E) were assessed in mice fed standard chow (baseline) or after 3 or 6 weeks of a HFD. In (A)–(E), the mean ± SEM is reported in N = 8 mice/group.

(F and G) Glucose (F) and insulin tolerance (G) tests were performed after 12 weeks of a HFD. The mean ± SEM is reported in N = 4–5 mice/group.

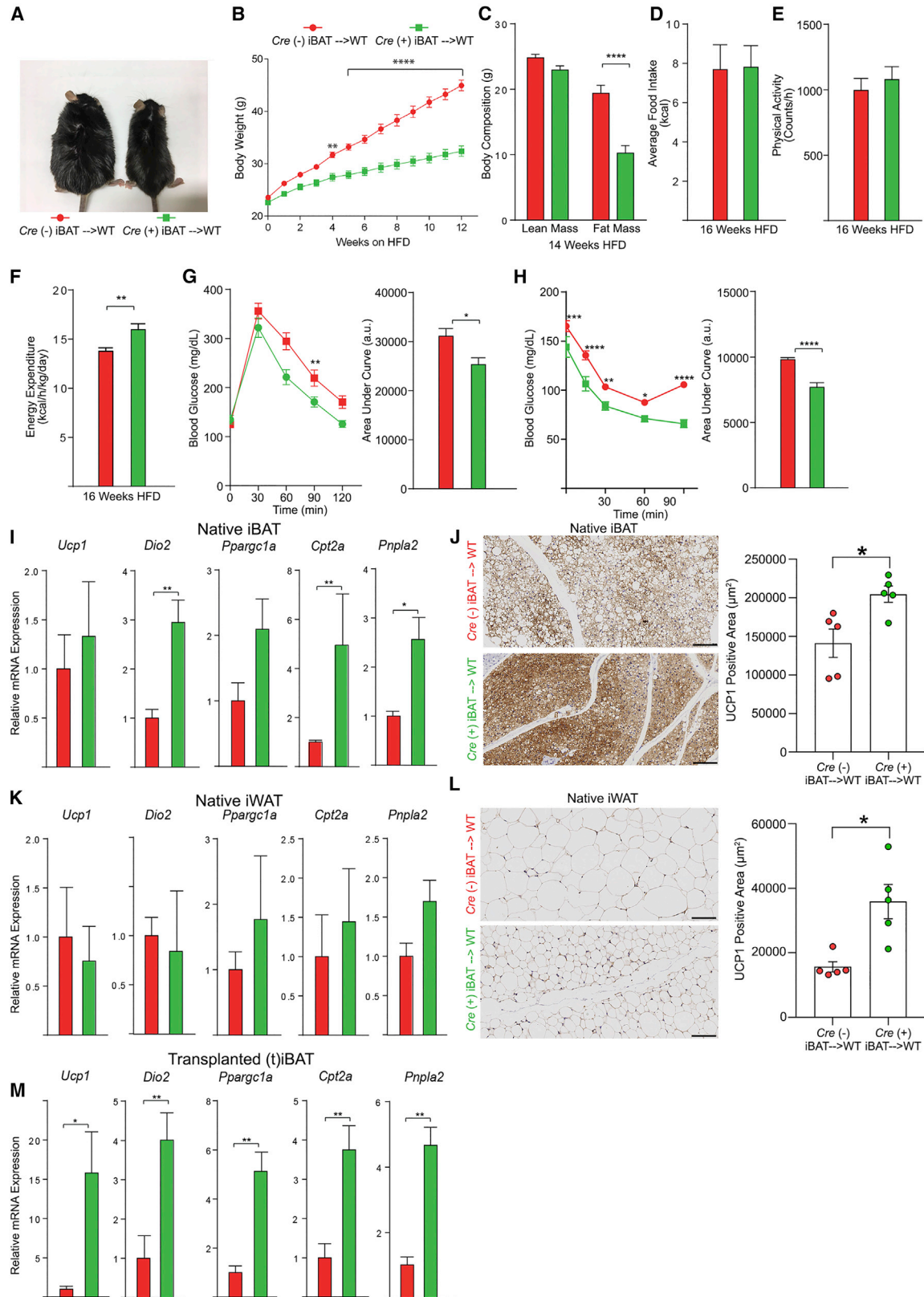
(H–K) Eight-week-old male *Ager^{flox/flox} Cre (-)* and *Ager^{flox/flox} Cre (+)* mice were housed for 24 h at 4°C.

(H and I) (Δ) Temperature (body temperature at room temperature versus 24 h of 4°C) (H) and change in body weight (I) are reported.

(J and K) Relative mRNA expression of *Ucp1*, *Dio2*, *Ppargc1a*, and *Cidea* mRNA was assessed in (J) iBAT and (K) iWAT retrieved from mice housed at room temperature or after termination of a 24-h cold challenge (4°C).

In (F)–(K), the mean ± SEM is reported in N = 4–5 mice/group. In (A)–(K), *Ager^{flox/flox} Cre (-)* shown as red and *Ager^{flox/flox} Cre (+)* shown as green. Data analysis: (A–G), repeated measures, and J and K, independent samples) two-way ANOVA followed by a post hoc Bonferroni test; (H and I) two-tailed Student's t test; *p < 0.05; **p < 0.01, ***p < 0.001, and ****p < 0.0001.

See also Figure S3.



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Cre (+) than those receiving *Cre* (–) iBAT, while levels of *Dio2* mRNA transcripts were significantly lower (Figure S4C). Figure S4D is a representative image of the secondary antibody-alone control for the UCP1 immunostaining.

We next transplanted iWAT expressing or devoid of adipocyte *Ager* into young, lean, WT mice and established the presence of vascularized non-necrotic grafts (representative image; Figure S5A). Four days after the transplantation, mice were switched to a HFD for an additional 20 weeks. Compared to recipients of *Ager^{fllox/fllox} Cre* (–) iWAT, WT mice receiving *Ager^{fllox/fllox} Cre* (+) iWAT displayed significantly less weight gain after 12 weeks of the HFD (Figures 5A and 5B). The WT mice recipients of *Ager^{fllox/fllox} Cre* (+) iWAT displayed significantly less fat mass than recipients of *Cre* (–) iWAT (Figure 5C). No differences in food intake or physical activity were observed by iWAT donor genotype (Figures 5D and 5E). As with iBAT transplants, significantly higher energy expenditure was observed in WT recipients of *Ager^{fllox/fllox} Cre* (+) than *Cre* (–) iWAT after 16 weeks of the HFD (Figure 5F). The WT mice receiving *Ager^{fllox/fllox} Cre* (+) versus *Cre* (–) iWAT displayed superior glucose tolerance (Figure 5G) and insulin sensitivity while on the HFD (Figures 5H and S5B).

After 20 weeks of the HFD, in native iBAT, WT mice receiving *Ager^{fllox/fllox} Cre* (+) versus *Cre* (–) iWAT displayed a significantly higher expression of *Ppargc1a* and *Pnpla2* mRNA transcripts, with no statistically significant differences in levels of *Ucp1*, *Dio2*, or *Cpt2a* mRNA transcripts (Figure 5I). Immunohistochemistry revealed that the native iBAT of mice that were transplanted with *Ager^{fllox/fllox} Cre* (+) versus *Cre* (–) iWAT displayed a significantly higher UCP1-positive area (Figure 5J). In native iWAT, WT mice receiving *Ager^{fllox/fllox} Cre* (+) versus *Cre* (–) iWAT displayed significantly higher levels of *Ucp1*, *Dio2*, and *Pnpla2* mRNA transcripts; significantly lower levels of *Ppargc1a* mRNA; and no differences in *Cpt2a* mRNA transcripts (Figure 5K). Immunohistochemistry revealed that there were no significant differences in the UCP1-positive area in the iWAT of mice that had been transplanted with *Ager^{fllox/fllox} Cre* (+) or *Cre* (–) iWAT (Figure 5L). In the tiWAT, the WT mice recipients of *Ager^{fllox/fllox} Cre* (+) versus *Cre* (–) iWAT displayed significantly higher levels of *Ucp1* and *Pnpla2* mRNA transcripts, a trend to

higher levels of *Dio2* expression, lower levels of *Ppargc1a*, and no significant differences in *Cpt2a* mRNA transcripts (Figure 5M). In native eWAT, significantly higher levels of *Ucp1*, *Ppargc1a*, and *Pnpla2* mRNA transcripts were observed in the recipients of *Ager^{fllox/fllox} Cre* (+) than of *Cre* (–) iWAT, with no statistically significant differences in *Dio2* or *Cpt2a* mRNA transcripts (Figure S5C). Thus, the deletion of *Ager* in iBAT or iWAT adipocytes transmits protection from diet-induced obesity and insulin and glucose intolerance and enhances expression of thermogenic genes.

To assess the effects of RAGE ligand sequestration, we previously treated WT mice fed a HFD for 18 weeks with soluble RAGE (sRAGE), the extracellular ligand-binding domains of RAGE, which act as a decoy receptor and thus suppress RAGE signaling. When treatment was given concurrently with a HFD, compared to vehicle-treated mice, mice treated with sRAGE displayed less body weight gain (Song et al., 2014). Compared to vehicle-treated mice, in the iWAT of sRAGE-treated mice, we noted the following: expression of *Ucp1* mRNA was significantly higher; expression of *Cidea* was significantly higher in the eWAT, iWAT, and iBAT; expression of *Ppargc1a* mRNA was significantly higher in the eWAT and iBAT; and expression of *Dio2* mRNA was significantly higher in eWAT (Figures S6A–S6D). Immunohistochemistry revealed more pronounced UCP1 immunoreactive epitopes in the eWAT, iWAT, and iBAT of sRAGE-treated mice (Figure S6E). Thus, both pharmacological and genetic approaches indicate that RAGE suppresses adipose tissue thermogenic programming.

RAGE Suppresses β 3-Adrenergic Stimulation of Lipolysis and *Ucp1* mRNA in Adipocytes

Compared to thermoneutrality, exposure to 4°C results in significant increases in sympathetic nerve innervation to the eWAT and iWAT (Castro et al., 2017). As β -adrenergic stimulation of adipocytes is a primary mechanism for lipolysis and the generation of fatty acid products that regulate *Ucp1* expression and activity (Cannon and Nedergaard, 2004; Collins and Surwit, 2001; Rial et al., 1983), we tested whether RAGE plays a role in this pathway in primary and cultured adipocytes.

Figure 4. Surgical Transplantation of Adipocyte *Ager*-Deficient iBAT Affects HFD-Induced Metabolic Impairment in WT Mice

(A) Representative photograph of mice surgically transplanted with iBAT from either *Ager^{fllox/fllox} Cre* (–) (red, left) or *Ager^{fllox/fllox} Cre* (+) iBAT (green, right) after 20 weeks on a HFD.

(B) Body weight curves through 12 weeks on a HFD.

(C) Body composition after 14 weeks of a HFD.

(D–F) Average daily food intake (D), physical activity (E), and energy expenditure (F) were assessed after 16 weeks on a HFD. For (B)–(F), the mean \pm SEM is reported in N = 15 mice/group.

(G and H) Glucose (G) and insulin tolerance (H) tests were performed after 12 or 13 weeks on a HFD, respectively. The mean \pm SEM is reported in N = 10–11 mice/group.

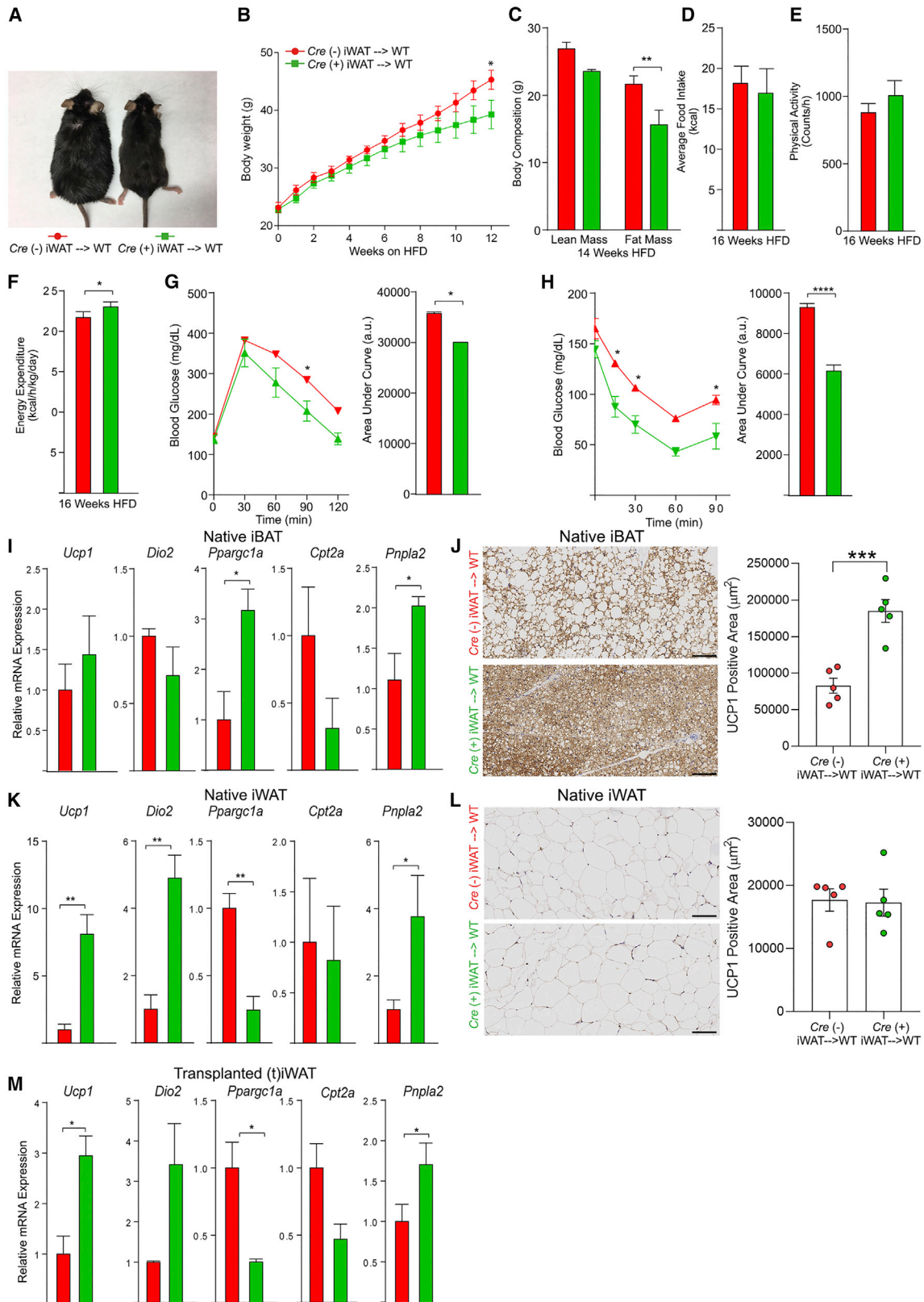
(I–M) Thermogenic programs.

(I, K, and M) The relative mRNA expression by qRT-PCR for *Ucp1*, *Dio2*, *Ppargc1a*, *Cpt2a*, and *Pnpla2* for (I) native iBAT, (K) native iWAT, and (M) transplanted iBAT (tiBAT) retrieved from the WT recipient mice after 20 weeks on a HFD is reported in N = 4–6 mice/group.

(J and L) In (J, native iBAT) and (L, native iWAT), immunohistochemistry for the detection of UCP1 was performed and image analysis employed to quantify UCP1-positive area; representative images are shown at 20 \times magnification. The mean \pm SEM is reported in N = 5 mice/group. Scale bar: 100 μ m.

(A–M) Source of donor iBAT: *Ager^{fllox/fllox} Cre* (–) shown as red and *Ager^{fllox/fllox} Cre* (+) shown as green. Data analysis: (B, G, and H, repeated measures, and C, independent samples) two-way ANOVA followed by a post hoc Bonferroni test; (D and F–H, area under the curve, and I–M) two-tailed Student's t test. Where group mean variances were statistically different ($p < 0.05$), data were analyzed post hoc using the non-parametric Mann-Whitney U test; * $p < 0.05$; ** $p < 0.01$, *** $p < 0.001$, and **** $p < 0.0001$.

See also Figure S4.



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We differentiated iBAT- and iWAT-derived adipocytes from WT and *Ager* null mice and stimulated them with CL316,243 (CL) (ARB3-specific agonist) to test the effects on lipolysis. In iBAT adipocytes, significantly higher CL316,243-stimulated levels of glycerol, and both basal and CL316,243-stimulated levels of non-esterified fatty acids (NEFAs), were observed in *Ager* null than in WT mice (Figure 6A). In iWAT adipocytes, basal and CL316,243-stimulated glycerol levels were significantly higher in adipocytes derived from *Ager* null mice, as was the CL316,243-mediated stimulation of the NEFA release (Figure 6B). Significantly higher oxygen consumption rates (OCRs) were observed in primary iBAT adipocytes from *Ager* null than from WT mice after treatment with isoproterenol (non-specific ARB agonist), which was minimally suppressed by oligomycin in the *Ager* null adipocytes, suggestive of maximum mitochondria uncoupling (Figure 6C). When primary iBAT adipocytes were treated with vehicle or norepinephrine (NE) + T3 (thyroid hormone), significantly higher *Ucp1* mRNA transcripts were observed in adipocytes devoid of *Ager* versus WT at baseline and after NE/T3 (Figure 6D). Primary iBAT-derived adipocytes from *Ager* null versus WT mice displayed significantly higher levels of UCP1 (Figure 6E). These data point to adipocyte-intrinsic roles for RAGE in β -adrenergic stimulation of lipolysis and upregulation of *Ucp1* expression and activity.

RAGE and its ligand, carboxymethyllysine (CML)-AGE, are highly expressed in obese human adipose tissue (Gaens et al., 2014). We previously showed that CML-AGEs were specific ligands of RAGE (Kislinger et al., 1999). WT primary adipocytes were treated with vehicle or NE, alone or with CML-AGE. In the basal state and upon treatment with NE, incubation with CML-AGE resulted in a significant suppression of *Ucp1* and *Pparg1a* mRNA expression in primary adipocytes from iBAT and iWAT (Figures 6F and 6G). In eWAT-derived adipocytes, CML-AGE treatment decreased *Ucp1* and *Pparg1a* at baseline and significantly suppressed the NE-stimulated induction of *Ucp1* with a similar trend noted for *Pparg1a* (Figure S6F). In iBAT-derived adipocytes, treatment with CML-AGE significantly decreased the expression of UCP1 (Figure 6H).

To determine if the effects of RAGE ligands were via direct action on *Ucp1* transcription, we employed *Ucp1* promoter

luciferase constructs (Figure 6I) transfected into C3H10T1/2 cells. Compared to the vehicle, treatment with NE resulted in a significant increase in *Ucp1* promoter luciferase activity, which was suppressed by CML-AGE (Figure 6I). We prepared mutants of the *Ucp1* promoter in which only the proximal promoter, which possesses functional CRE-binding elements, or the enhancer elements, were expressed (Figure 6J). Treatment with CML-AGE exerted only modest effects on the suppression of basal but not NE-mediated promoter luciferase activities, indicating that the entire *Ucp1* promoter was required to detect the effects of CML on transcriptional activity. On account of central roles for PKA in the β -adrenergic stimulation of thermogenic gene programs in adipocytes, we tested the effects of CML-AGE on PKA activity. In primary iBAT-derived adipocytes treated with CL316,243 and CML-AGE, a significant attenuation in PKA activity was observed (Figure 6K).

RAGE Ligands Suppress Phosphorylation of PKA Pathway Targets, Phosphorylated Hormone Sensitive Lipase (pHSL) Ser563 and p38 MAP Kinase, upon β 3-Adrenergic Stimulation

To trace the fate of lipolysis and signaling pathways downstream of PKA that regulate adaptive thermogenesis, we focused on two PKA targets: HSL and p38 MAP kinase (MAPK) (Collins, 2012). Based on the discovery that the cytoplasmic domain of RAGE is essential for RAGE signaling, in part through its binding to DIAPH1 (Hudson et al., 2008; Touré et al., 2012), we recently described the small molecule antagonist, 2-[4-(acetylamino)phenyl]-4-Quinolinecarboxylic acid, methyl ester ("RAGE inhibitor"), which binds to the cytoplasmic domain of RAGE, blocks its interaction with DIAPH1, and suppresses RAGE ligand-stimulated signal transduction in cellular and murine models (Manigrasso et al., 2016). Significantly higher levels of glycerol release were observed in C3H10T1/2 cells treated with CL316,243 and the RAGE inhibitor (RI) versus CL316,243 alone (Figure 7A). The RI had no independent effect on the numbers of lipid droplets, as assessed by staining with BODIPY (Figure 7B), nor did it affect the relative mitochondrial DNA content (Figure S7A).

Figure 5. Surgical Transplantation of Adipocyte *Ager*-Deficient iWAT Affects HFD-Induced Metabolic Impairment in WT Mice

(A) Representative photographs from mice receiving surgical transplantation of either *Ager*^{flox/flox} Cre (-) (red, left) or *Ager*^{flox/flox} Cre (+) iWAT (green, right) are shown after 20 weeks on a HFD.

(B) Body weights through 12 weeks of a HFD.

(C) Body composition after 14 weeks of a HFD.

(D–F) Average daily food intake (D), physical activity (E), and energy expenditure (F) were assessed after 16 weeks of a HFD. In (B)–(F), the mean \pm SEM is reported in N = 7–9 mice/group.

(G and H) Glucose (G) and insulin tolerance (H) tests were performed after 12 and 13 weeks of a HFD, respectively. The mean \pm SEM is reported in N = 5–6 mice/group.

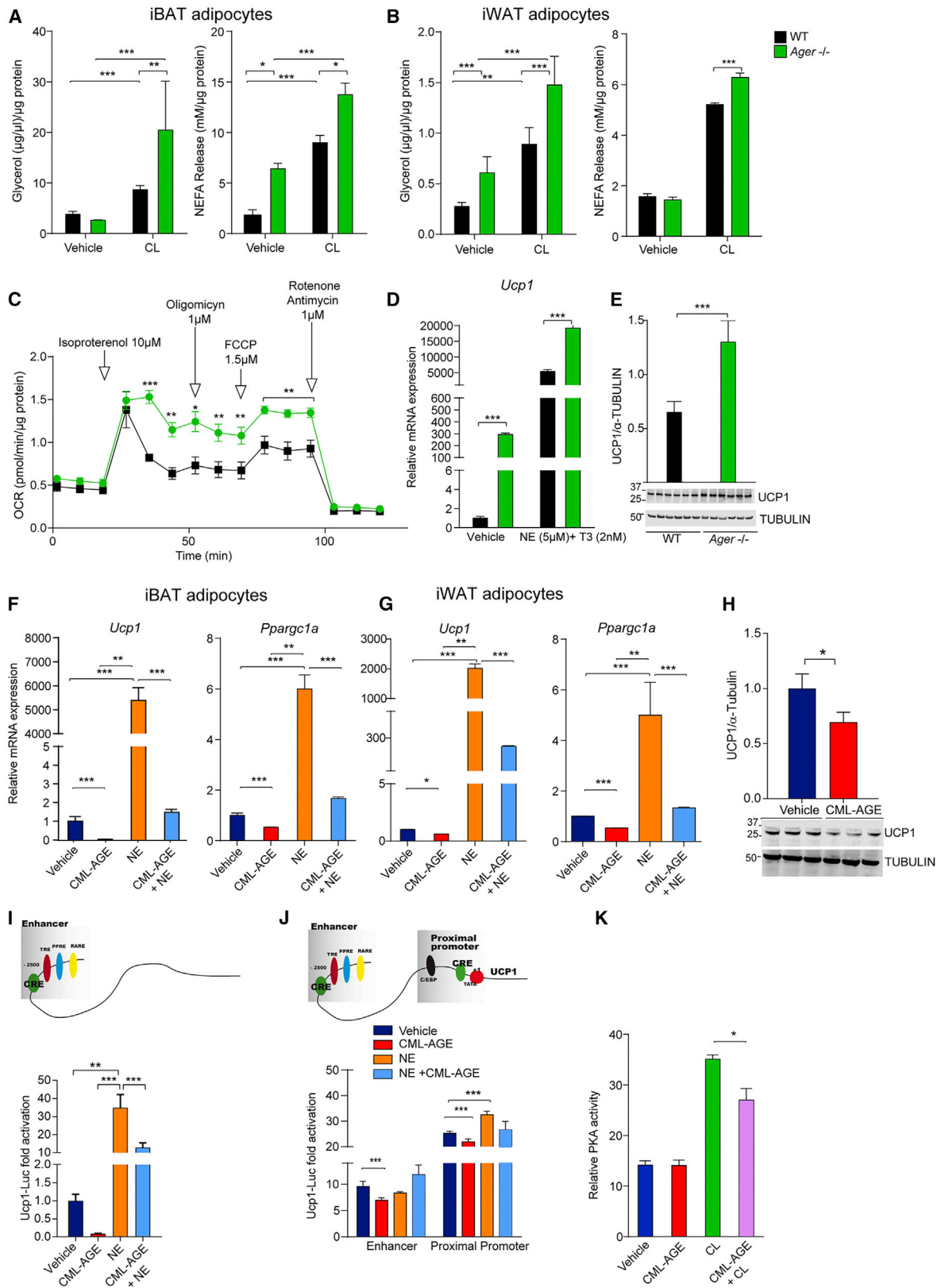
(I–M) Thermogenic programs.

(I, K, and M) The relative mRNA expression by qRT-PCR for *Ucp1*, *Dio2*, *Pparg1a*, *Cpt2a*, and *Pnpla2* mRNA transcripts in (I) native iBAT, (K) native iWAT, and (M) transplanted iWAT (tiWAT) was examined, and the mean \pm SEM is reported in N = 3–4 mice/group.

(J and L) In (J, native iBAT) and (L, native iWAT), immunohistochemistry for detection of UCP1 protein was performed and image analysis employed to quantify UCP1-positive area; representative images at 20 \times magnification are shown. The mean \pm SEM is reported in N = 5 mice/group. Scale bar: 100 μ m.

(A–M) Source of donor iWAT: *Ager*^{flox/flox} Cre (-) shown as red and *Ager*^{flox/flox} Cre (+) shown as green. Data analysis: (B, G, and H, repeated measures, and C, independent samples) two-way ANOVA followed by a post hoc Bonferroni test; (D–H, area under the curve, and I–M) two-tailed Student's t test. Where group mean variances were statistically different ($p < 0.05$), data were analyzed post hoc using the non-parametric Mann-Whitney U test; * $p < 0.05$; ** $p < 0.01$, *** $p < 0.001$, and **** $p < 0.0001$.

See also Figure S5.



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We assessed the effects of RAGE on phosphorylation of HSL Serine563 and p38 MAPK. Treatment of C3HT101/2 cells with CL316,243 resulted in significantly higher phospho/total pHSL Serine563 and phospho/total p38 MAPK compared to vehicle, which were both significantly reduced by treatment with CML-AGE. However, in cells treated with CL316,243 and CML-AGE, treatment with the RI resulted in significantly higher phospho-p38/total p38 MAPK and trended to higher phospho-HSL Serine563/total HSL, compared to treatment with CL316,243 and CML-AGE alone (Figure 7C). We hypothesized that if these effects of RAGE were through a blockade of phosphorylation of PKA targets, then direct catalytic inhibition of PKA should blunt the effects of the RI. First, we tested the efficacy of H89, a commonly used compound to inhibit PKA activity, on the suppression of the effects of CL316,243 (Chijiwa et al., 1990). In C3HT101/2 cells, treatment with H89 blocked the increased phosphorylated HSL Serine563/total HSL ratio and the increased phosphorylated p38/total p38 MAPK induced by CL316,243 (Figure S7B). We then tested if H89 blocked the effects of the RI. Compared to cells treated with CL316,243, CML-AGE, and the RI, the addition of H89 resulted in a significant reduction in phosphorylation of both HSL Serine563 and p38 MAPK (Figure 7D). Thus, direct catalytic inhibition of PKA blunted the effects of the RI on rescuing the suppressive effects of CML-AGE on the phosphorylation of pHSL Serine563 and p38 MAPK.

We next tested primary adipocytes from iBAT and iWAT. In WT, but not in *Ager* null primary adipocytes derived from iBAT (Figure 7E) and iWAT (Figure 7F), treatment with CML-AGE in CL316,243-treated cells resulted in significant suppression of phosphorylation of pHSL Serine563/total HSL and p38/total p38 MAPK. In primary iBAT- and iWAT-derived adipocytes, treatment with CL316,243 resulted in significantly

increased phosphorylated HSL Serine563/total HSL and increased phosphorylated p38/total p38 MAPK, which was significantly attenuated by H89 (Figure S7C). Treatment of *Ager* null adipocytes from iBAT or iWAT with CL316,243, CML-AGE, and H89 resulted in significant decreases in phosphorylated HSL Serine563/total HSL and phosphorylated p38/total p38 MAP kinase, compared to cells treated with CL316,243 and CML-AGE alone (Figures 7G and 7H). Thus, direct catalytic inhibition of PKA significantly reduced the effects of deletion of *Ager* on rescuing the suppression of CML-AGE on phosphorylation of these two PKA targets. *In vivo*, expression of *Lipe* (gene encoding HSL, a target of activated PKA (Holm, 2003), was significantly higher in the floating adipocyte fraction retrieved from the iWAT of *Ager* null than from the WT mice (Figure S7D).

On account of the demonstrated roles for a PKA-independent, β 3-adrenergic-mediated stimulation of extracellular signal-related kinase (ERK) MAPK on lipolysis in Robidoux et al. (2006), we tested the effects of the RAGE ligand, CML-AGE. In C3H10T1/2 cells, treatment with CL316,243 resulted in significantly increased phospho/total ERK MAPK, which was significantly suppressed by CML-AGE; the suppressive effects of CML-AGE were prevented by treatment with the RI (Figure S7E). However, in primary WT or *Ager* null adipocytes retrieved from iBAT or iWAT, no significant effects of CML-AGE on phospho/total ERK MAPK in CL316,243-treated cells were observed (Figure S7F).

There were no *Ager*-dependent differences in the protein levels of two other lipolytic molecules, adipose triglyceride lipase or monoacylglycerol lipase, upon treatment with CL316,243 (Figures S7G and S7H), and there were no genotype-dependent differences in the levels of cyclic AMP (cAMP) between WT and *Ager* null CL-316,243-treated primary adipocytes from iBAT, iWAT, or eWAT (Figure S7I).

Figure 6. RAGE Affects β -Adrenergic-Stimulated Adipocyte Lipolysis and Thermogenic Programming

Primary adipocytes were prepared from the SVF of 8-week-old WT and *Ager* null mice fed standard chow.

(A and B) iBAT- (A) and iWAT- (B) derived cells were incubated for 3 h in Krebs-Ringer bicarbonate buffer (KRB) and subjected to vehicle or CL316,243 (CL) (10 μ M) for 3 h; glycerol and non-esterified fatty acids (NEFAs) in the media were measured. The means normalized to total protein per well \pm SEM are reported from triplicate, independent experiments from cells derived from N = 5 mice/group.

(C) A representative OCR curve for iBAT-derived adipocytes during a mitochondrial stress test is shown under basal condition and following isoproterenol (10 μ M). This experiment was performed three times using cells derived from N = 3 mice/group per experiment.

(D) Primary iBAT-derived adipocytes from iBAT of WT or *Ager* null mice were stimulated with norepinephrine (NE; 5 μ M) and T3 (thyroid hormone, 2 nM) for 6 h, and relative *Ucp1* mRNA expression was determined by qRT-PCR. The mean \pm SEM is reported in N = 4 mice/group.

(E) Western blotting was performed for detection of UCP1/TUBULIN from primary iBAT adipocytes of 8-week-old WT and *Ager* null mice. The mean \pm SEM is reported in N = 6 independent cell lysates per group along with representative blots.

(F and G) Primary adipocytes from (F) iBAT and (G) iWAT of WT mice were incubated with vehicle or the RAGE ligand CML-AGE (300 μ g/ml) for 16 h alone or with NE (5 μ M) for the final 6 h. qRT-PCR for detection of relative *Ucp1* or *Ppargc1a* mRNA expression was performed. The mean \pm SEM is reported from N = 5 mice/group.

(H) Cell lysates from primary adipocytes from iBAT of WT mice treated as in (F) and (G) were used for the detection of UCP1/TUBULIN. The mean \pm SEM is reported in cell lysates from N = 3 mice/group.

(I) Schematic of *Ucp1* promoter with enhancer elements. Undifferentiated C3H10T1/2 cells were transfected with mouse 3.1 kB *Ucp1* promoter luciferase construct (*Ucp1*-luc) and stimulated for the last 6 h with NE (5 μ M) and/or CML-AGE (300 μ g/ml) for 16 h. Normalized luciferase activities are shown as fold-change compared to vehicle control.

(J) *Ucp1* promoter luciferase constructs were generated and transfected into undifferentiated C3H10T1/2 cells and the cells cultured in medium containing NE (5 μ M) for the last 6 h or CML-AGE (300 μ g/ml) for 16 h. Normalized luciferase activities are shown relative to vehicle control. The mean \pm SEM of three independent experiments is presented.

(K) PKA activity was determined in primary iBAT adipocytes from WT mice treated with CL316,243 (10 μ M) for 15 min alone or after pre-incubation with the RAGE ligand CML-AGE (300 μ g/ml) for 16 h. The mean \pm SEM of N = 8 mice/group is presented.

In (A)–(E), WT (black) and *Ager* null (green). Data analysis: (A–D, repeated measures) two-way ANOVA followed by post hoc Bonferroni test; (E and H) two-tailed Student's t test; (F, G, and I–K) one-way ANOVA followed by a post hoc Tukey's HSD test; *p < 0.05, **p < 0.01, and ***p < 0.001.

See also Figure S6.

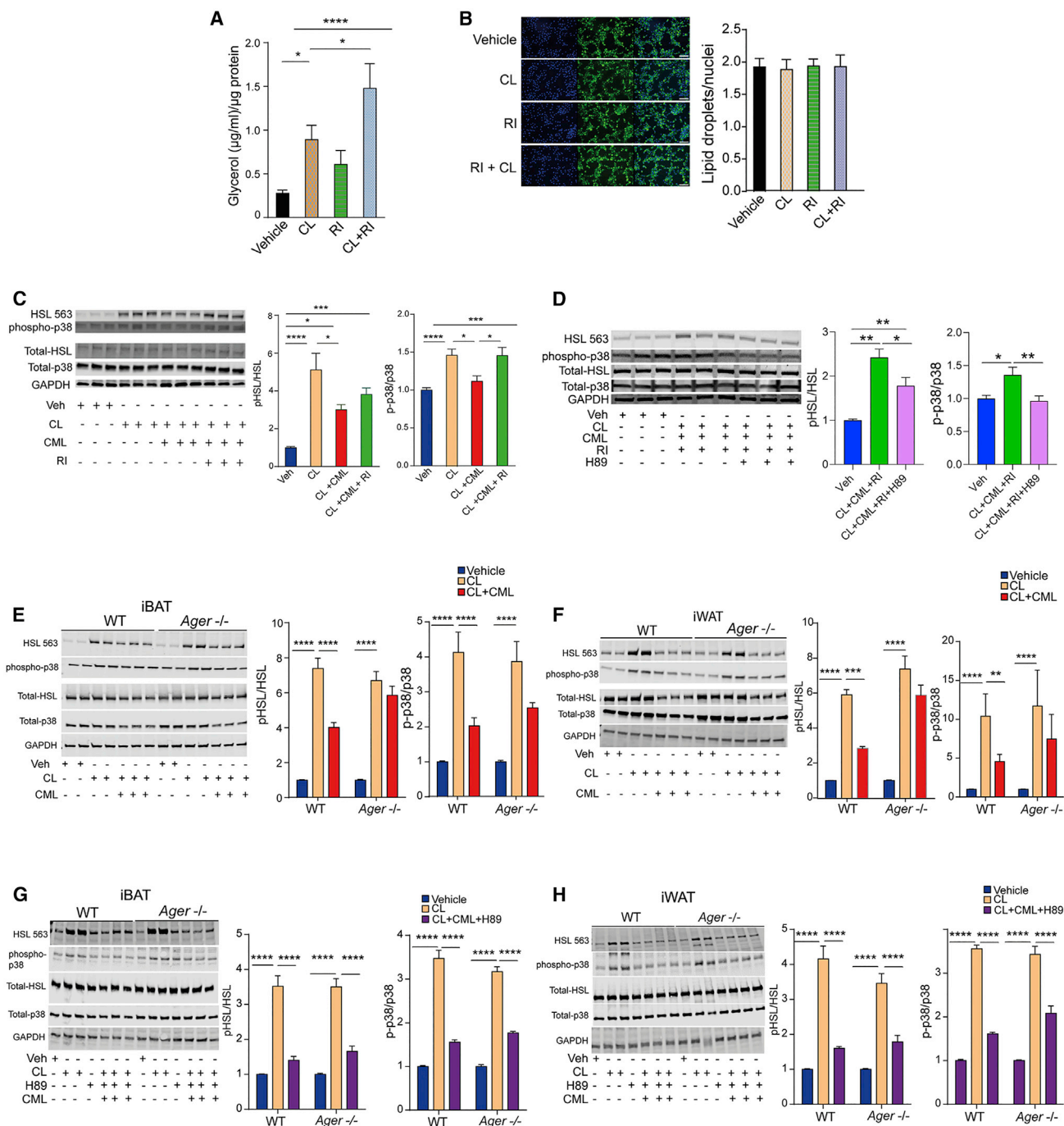


Figure 7. Effects of RAGE Signal Transduction on the PKA Signaling Pathway, Adipocyte Lipolysis, and Phosphorylation of p38 MAP Kinase (A) Differentiated adipocytes from C3H10T1/2 cells were serum starved for 3 h and pretreated with the RAGE inhibitor (2-[4-(acetylamino) phenyl] -4- Quinolincarboxylic acid, methyl ester) (1 µM) for 1 h prior to CL316,243 (10 µM) for 15 min. The mean glycerol content normalized to total protein per well ± SEM is reported from three independent experiments.

(B) C3H10T1/2 cells were differentiated to adipocytes and pre-incubated with a vehicle or the RAGE inhibitor (1 µM) for 90 min prior to CL316,243 (10 µM). Lipid droplets were stained using BODIPY and nuclei stained with DAPI. A representative fluorescence micrograph and mean ± SEM is shown. Scale bar: 100 µm.

(C and D) C3H10T1/2 cells were differentiated and treated with a vehicle or CML-AGE (300 µg/ml) for 75 min alone or after pre-treatment with the RAGE inhibitor (1 µM) for 30 min followed by CL316,243 (10 µM) treatment for 15 min.

(C) Cells were lysed and western blotting performed for phosphorylated HSL Serine563, total HSL, phosphorylated p38 MAPK, total p38 MAPK, and GAPDH. (D) Differentiated adipocytes from C3H10T1/2 cells were treated as in (C) with or without the addition of H89 (20 µM) for 30 min prior to CL316,243 (10 µM). Cells were lysed and western blotting performed for phosphorylated HSL Serine563, total HSL, phosphorylated p38 MAPK, total p38 MAPK, and GAPDH. Band

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DISCUSSION

The tethering of the biology of molecules implicated in inflammation to the regulation of metabolism underscores the significance of these two primal processes to drive and ensure organismal survival. Here, we demonstrate that the immunoglobulin superfamily molecule, RAGE, suppresses adaptive thermogenesis in both BAT and WAT, at least in part by reducing the effects of β -adrenergic signaling in adipocytes, through suppression of the phosphorylation of PKA targets. In contrast to the suppressive effects of RAGE on metabolic responses, distinct genes implicated in the inflammatory response positively regulate adipocyte lipolysis and expression of *Ucp1*. For example, IRF4 is induced in fasting, and experiments testing adipocyte-specific deletion of *Irf4* (Lee et al., 2013) uncovered its roles in the positive regulation of adipocyte lipolysis (Eguchi et al., 2011; Kumari et al., 2016). Further, a role for toll-like receptor 4 (TLR4) in endotoxin-stimulated regulation of lipolysis in adipocytes was established (Zu et al., 2009) and shown to be independent of cAMP-PKA but dependent on the ERK1/2 signal transduction pathway. The observation that levels of *Irf4* and *Tlr4* were significantly higher in adipose tissues of HFD-fed mice bearing adipocyte-specific deletion of *Ager* versus animals expressing *Ager* in adipocytes suggests that RAGE may contribute, directly or indirectly, to the regulation of these genes in adipocytes.

What are the signals that trigger these events in the RAGE pathway? Many of the families of the RAGE ligands have “double lives,” in which their intracellular regulatory functions contrast with unique roles in the extracellular zones as “damage-associated molecular pattern (DAMP)” molecules (Bertheloot and Latz, 2017; Matzinger, 1994). In the intracellular space, the RAGE ligands S100/calgranulins and HMGB1 (Taguchi et al., 2000; Hofmann et al., 1999) modulate calcium homeostasis and are nonhistone DNA binding proteins, respectively. In the extracellular space, these RAGE ligands assume response-to-stress functions, such as driving cellular inflammation through activation of nuclear factor- κ B and p38 MAPK (Gunasekaran et al., 2016; Fujiya et al., 2014). Direct roles for the RAGE ligands in obesity have been demonstrated, as blocking antibodies to HMGB1 reduced obesity and inflammation in a mouse model (Montes et al., 2015). As endogenous RAGE expression in adipocytes may be sufficient to quell thermogenic processes, it is possible that adipocytes generate and release RAGE ligands, thereby stringently regulating thermogenesis and energy expenditure. Indeed, our data in primary adipocytes from iBAT, iWAT, and eWAT revealed a time-dependent increase in *Ager* expression. In earlier work, ligand expression was linked to transcrip-

tional upregulation of *Ager* in cultured cells (Li and Schmidt, 1997).

Our data indicate that in states of RAGE ligand enrichment, such as in high-fat feeding, the activity of the RAGE axis is heightened. Although iBAT from adipocyte *Ager*-deleted mice displays lower levels of *Ucp1* and *Dio2* at room temperature and in low-fat feeding (i.e., LOW RAGE LIGAND environment) when compared to iBAT from adipocyte *Ager*-expressing mice (Figure 3J), when these iBAT tissues are transplanted into environmentally-stressed WT mice fed a HFD (i.e., HIGH RAGE LIGAND environment), the absence of RAGE in adipocytes unrestrains thermogenic programming, as evidenced by the increased expression of *Ucp1*, *Dio2*, *Ppargc1a*, *Cpt2a*, and *Pnpla2* mRNA in the transplanted iBAT of the adipocyte *Ager*-deleted versus *Ager*-expressing iBAT (Figure 4M). These considerations are supported by the work presented in Figures 6 and 7, in which the RAGE ligand CML-AGE suppresses lipolysis, phosphorylation of p38 MAPK, and thermogenic gene programs.

The activation of the β 3-adrenergic receptor, which is highly expressed in adipocytes, results in the generation of cAMP and consequent PKA-mediated phosphorylation of HSL and perilipin (Soeder et al., 1999). The present studies illustrate that recently identified inhibitors of RAGE signal transduction (Manigrasso et al., 2016) block the RAGE ligand-mediated reduction of phosphorylation of PKA targets in adipocytes. *Ager* deletion, unlike the metabolic braking effects of KCNK3 (Chen et al., 2017), exerted no effects on cAMP levels in adipocytes at baseline or after CL316,243 stimulation. The observation that catalytic inhibition of PKA (H89) effectively blocked the effects of *Ager* deletion or the effects of the RI on adaptive thermogenesis in primary and cultured adipocytes supports a notion that the RAGE effects are mediated, at least in part, through the suppression of PKA-dependent phosphorylation of its downstream targets in adipocytes. In line with recent work that demonstrated that lipolysis in BAT is not required for cold-induced thermogenesis (Schreiber et al., 2017; Shin et al., 2017), our findings reveal broader effects for RAGE, in that beyond RAGE-dependent suppression of the β -adrenergic-mediated phosphorylation of PKA target HSL in adipocytes, fundamental signaling pathways that regulate *Ucp1* and thermogenic genes, such as phosphorylation of p38 MAPK, are also downregulated by RAGE ligands and restored by *Ager* deletion or by treatment with RIs.

The present findings trace the impact of *Ager* deletion in the regulation of adaptive thermogenesis from the induction of lipolysis, particularly through the phosphorylation of HSL, to the generation of glycerol and free fatty acids and to the phosphorylation of p38 MAPK and regulation of *Ucp1* mRNA transcription. It is

intensities were normalized to the respective total HSL or total p38 MAPK, and the relative fold change is presented as mean \pm SEM. Each experiment consisted of at least three independent studies with at least three technical replicates.

(E–H) Primary adipocytes from (E and G) iBAT or (F and H) iWAT of WT and *Ager* null mice were treated with the RAGE ligand CML-AGE (300 μ g/ml) for 1 h, CL316,243 (10 μ M) for 15 min, and with or without H89 (20 μ M) for 30 min prior to CL. Cells were lysed and western blotting performed for the detection of phosphorylated HSL Serine563, total HSL, phosphorylated p38 MAPK, total p38 MAPK, and GAPDH. Band intensities were normalized to the respective total HSL or p38 MAPK, and the relative fold change is presented as mean \pm SEM from N = 4 pooled mice/group in three independent experiments with at least 2–3 technical replicates per experiment.

Data analysis: (A–H) one-way ANOVA followed by a post hoc Tukey's HSD test or Bonferroni test, as indicated; *p < 0.05; **p < 0.01, ***p < 0.001, and ****p < 0.0001.

See also Figure S7.

established that circulating fatty acids from lipolysis serve as key cofactors facilitating the activation of UCP1 and the transport of protons across mitochondrial membranes, which is essential for mammalian regulation of heat (Garlid et al., 1996; Klingenberg and Winkler, 1985; Cannon and Nedergaard, 2004). In contrast to the suppressive effects of RAGE signaling on phosphorylation of p38 MAPK in primary adipocytes (iBAT and iWAT) and in C3H10T1/2 cells, our experiments revealed inconsistent findings regarding phosphorylation of ERK. Although the reasons for the differences in ERK MAPK phosphorylation vis-à-vis RAGE between cell lines and primary adipocytes remain to be identified, our data nevertheless point to clear roles for RAGE signaling-dependent regulation of HSL and p38 MAPK phosphorylation downstream of PKA.

In the broader context of evolution and adaptive selection, then, what forces drive this metabolic braking role for *Ager*? Selective pressures in evolution favor phenotypes that safeguard survival in the face of paucities and extreme stresses (Hotamisligil, 2017). Hence, when nutrient supplies are unlimited and selective survival pressures are lost, the urge to hoard energy, if unchecked, may become a vulnerability. The gene *AGER*, located on chromosome 6 in the major histocompatibility complex (MHC) III humans (Wu et al., 2015), first appeared in Laurasiatheria (Sessa et al., 2014), a superorder of placental mammals that is part of the larger group of mammals classified as Eutheria. Eutheria are mammalian clades, and the oldest Eutherian species is believed to be *Juramaia sinensis*, which dates to 160 million years ago (mya) (Gerkema et al., 2013). A key property of the Eutherians is the expression of UCP1 in BAT, which imbues the capacity for non-shivering thermogenesis (Hughes et al., 2009). Mammals first appeared during the Mesozoic era, about 250 mya, which followed the period known as the “Great Dying,” in which a massive extinction of plants and land species ensued consequent to a period of intense global warming (Gerkema et al., 2013; Sun et al., 2012). Perhaps *AGER* evolved as a defense against starvation or swings in ambient temperatures through its ability to suppress adaptive thermogenesis.

These premises may indeed be consistent with hypotheses that reduced energy expenditure and obesity may be the unintended consequences of evolutionary pressures against severe starvation (“thrifty genotype”) (Neel, 1962; Prentice, 2005; Wang et al., 2014) or the need to protect from extreme temperature swings throughout human migration (Sellayah et al., 2014). At thermoneutrality, mice devoid of *Ager* bear body temperatures that are indistinguishable from those of WT animals, thereby implying that RAGE-dependent forces are not triggered during homeostasis, but are evoked under stress conditions in which the ligands of the receptor accumulate. Indeed, in the evolution of *AGER*, the major adaptively selected sites within the protein product RAGE are within the extracellular ligand binding domains, the intracellular adaptor binding regions, and the surfaces facilitating oligomerization and, thereby, signal transduction (Wu et al., 2015).

Notably, distinct pathways linking metabolism and inflammation, specifically TLR, insulin receptor, and tumor necrosis factor (TNF) receptor signaling, may be traced to orthologs in *Drosophila* (Hotamisligil, 2017). This is not the case for *Ager*.

However, the RAGE ligands HMGB1 and S100/calgranulins, such as S100A8/A9, may also bind TLRs (Gunasekaran et al., 2016; Hiratsuka et al., 2008). TLRs play innate roles in host defense, but the role of *Ager* in response to infectious pathogens is assuredly more complex (van Zoelen et al., 2011). Whereas in cecal ligation and puncture (sepsis) or in *Streptococcus pneumoniae* pneumonia, a deficiency of *Ager* in mice improved survival (Liliensiek et al., 2004; van Zoelen et al., 2009), the loss of *Ager* in *Klebsiella pneumoniae* pneumonia was detrimental (Achouiti et al., 2016). Together with the marked differences in appearance of *TLR* versus *AGER* in evolution and the lack of consistent roles for *AGER* in innate host defense to pathogens, it is plausible that RAGE’s ability to conserve energy underscores its primal endogenous function, which evolved, remarkably, at the expense of unremitting and chronic inflammation and the dampening of tissue repair on account of RAGE’s ability to usurp the functions of HMGB1 and S100/calgranulins ligands. Such a scenario frames a critical paradigm in the immunometabolic nexus of genes that integrate inflammatory and metabolic responses and identifies RAGE as an innate mechanism in the suppression of adipocyte metabolism and energy expenditure.

In summary, this work demonstrates that RAGE suppresses adaptive thermogenesis in both BAT and WAT during a physiological response to fasting, a cold challenge, or high-fat feeding; adds *Ager* to the cadre of genes that tether inflammation to the regulation of energy homeostasis; defines the endogenous function of RAGE in energy metabolism; and pinpoints tractable therapeutic targets to harness energy expenditure in metabolic disorders through the blockade of RAGE signal transduction.

STAR★METHODS

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Supplemental Information can be found online at <https://doi.org/10.1016/j.celrep.2019.06.061>.

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AUTHOR CONTRIBUTIONS

C.H.d.P. and H.H.R. designed the research, performed experiments, analyzed data, prepared figures, contributed to writing the manuscript, and edited the manuscript. L.A. and J.F.A. contributed to the experimental design, performed key experiments, analyzed data, prepared figures, and edited the manuscript. C.S., P.D., J.D., M.M., M.H., L.F., R.H.F., and H.L.N. performed experiments and edited the manuscript. J.K.K., R.A.F., and R.R. contributed analytically and intellectually to the experimental design and interpretation of the data and edited the manuscript. A.M.S. conceived the studies, designed the research, supervised the design and completion of the experiments, analyzed data, and wrote and edited the final manuscript.

DECLARATION OF INTERESTS

A.M.S. and R.R. are co-inventors of patent applications related to the work in this manuscript.

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STAR★METHODS

KEY RESOURCES TABLE

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Antibodies		
Anti-UCP-1 (Western Blot)	Abcam	Cat# ab10983; RRID:AB_2241462
Anti-UCP-1 (Immunohistochemistry)	Abcam	Cat# Ab23841; RRID:AB_2213764
Anti-phospho HSL (Ser563)	Cell Signaling	Cat# 4139S; RRID:AB_2135495
Anti-total HSL	Cell Signaling	Cat# 4107S; RRID:AB_2296900
Anti-Tubulin	Cell Signaling	Cat# 3873S; RRID:AB_1904178
Anti-phospho ERK	Cell Signaling	Cat# 9101L; RRID:AB_331646
Anti-total ERK	Cell Signaling	Cat # 9102; RRID:AB_330744
Anti-ATGL	Cell Signaling	Cat # 2439S; RRID:AB_2167953
Phospho-p38 MAPK (Thr180/Tyr182)	Cell Signaling	Cat #9211; RRID:AB_331641
Anti-p38 MAPK	Cell Signaling	Cat # 9212; RRID:AB_330713
Anti-Beta Actin	Santa Cruz	Cat# sc-81178; RRID:AB_2223230
Anti-Monoacylglycerol Lipase (MGL)	Abcam	Cat# ab180016
IRDye 680RD goat anti-mouse IgG	Li-Cor	Cat# 925-68070; RRID:AB_2651128
Dylight 800-conjugated goat anti-mouse IgG	Li-Cor	Cat# 925-32211; RRID:AB_2651127
Anti-GAPDH	Santa Cruz	Cat# sc-47724; RRID:AB_627678
Chemicals, Peptides, and Recombinant Proteins		
Hoeschst 33342	Invitrogen	H3570
Seahorse XF Cell Mito Stress Test	Agilent	103015-100
Seahorse XFe24 FluxPak	Agilent	100850-001
iScript cDNA Synthesis Kit, 500 × 20 μl rxns	Biorad	1708891BUN
10x Tris/Glycine Buffer	Bio-Rad	1610771EDU
10x Tris/Glycine/SDS	Bio-Rad	1610732EDU
4–15% Mini-PROTEAN® TGX Precast Protein Gel	Bio-Rad	4561086
Cell Lysis Buffer (10X)	Cell Signaling	9803
RIPA Buffer (10X)	Cell Signaling	9806
4-20% Mini-PROTEAN TGX gels	Bio-Rad	456-1096
Trans-Blot® Turbo Midi Nitrocellulose Transfer	Bio-Rad	1704159
10x Tris/Glycine/SDS Buffer	Bio-Rad	161-0772
10x Tris/Glycine Buffer	Bio-Rad	161-0771
Sybr Green	Applied Biosystems	4385618
Odyssey® Blocking Buffer (PBS)	Li-Cor	927-40003
MitoTracker® Red CMXRos	Life Technologies	M7512
DMEM, high glucose, HEPES, no phenol red	Life Technologies	21063-029
Phusion® High-Fidelity PCR Kit	New England Biolabs, Inc	E0553L
Dual-Glo® Luciferase Assay System	Promega	E2940
Glo Lysis Buffer, 1X	Promega	E2661
QIAzol Lysis Reagent	QIAGEN	7906
RNeasy Mini Kit (250)	QIAGEN	79306
Paraformaldehyde solution 4% in PBS	Santa Cruz Biotechnology	CAS 30525-89-4
(–)-Norepinephrine	Sigma	A7257
3,3',5-Triiodo-L-thyronine sodium salt	Sigma	T6397
3-Isobutyl-1-methylxanthine	Sigma	I5879
CL 316243,	Sigma	C5976

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Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Collagenase from <i>Clostridium histolyticum</i>	Sigma	C6885-1G
Dexamethasone	Sigma	D4902
Free Glycerol Reagent	Sigma	F6428
Glucose	Sigma	G8270
Glycerol Standard	Sigma	G1394
H-89 dihydrochloride hydrate	Sigma	B1427-5MG
Insulin from bovine pancreas	Sigma	I6634
Isoproterenol	Sigma	I5627
Rosiglitazone	Sigma	R2408
DMEM	ThermoFisher Scientific	11995065
Lipofectamine® 3000 Transfection Reagent	ThermoFisher Scientific	L3000001
Optimem	ThermoFisher Scientific	31985047
Bovine Serum Albumin - Fatty Acid Free	Roche	3117057001
FBS	Corning	35-010-CV
Glucose strips - FreeStyle Lite Blood Glucose Test Strips	Abbott	70819-70
D-Glucose	Alfa Aesar	A16828
Magnesium Chloride	Sigma	M8266
Sodium Chloride	Fisher Scientific	BP318-212
Potassium Chloride	Fisher Scientific	P217-500
Sodium Phosphate Dibasic	Sigma	S3264
LipidTox	ThermoFisher Reagent	H34476
BODIPY	ThermoFisher Reagent	D3922
DAPI	Life Technologies	D1306
Sodium Phosphate Monobasic	Sigma	S8282
Sodium Bicarbonate	Fisher Scientific	S3233-500
Insulin (Human) - Humulin R	Lilly	0002-8215-01
CML-AGE	Synthesized in Laboratory per Kislinger et al., 1999	N/A
2- [4- (acetylamino) phenyl] –4- Quinolinecarboxylic acid, methyl ester	Synthesized Per the Methods/Source outlined in Manigrasso et al., 2016	N/A
Critical Commercial Assays		
2-CAT (A-N) Research ELISA	Rocky Mountain Diagnostics	BA E-5400
cAMP Direct Immunoassay Kit	Abcam	Ab65355
Glycerol/Triglyceride Determination Kit	Sigma	TR0100-1KT
NEFA HR(2) color reagent A	Wako Diagnostics	99934691
NEFA-HR(2) color reagent B	Wako Diagnostics	99134891
NEFA HR(2) Solvent A	Wako Diagnostics	99534791
NEFA HR(2) Solvent B	Wako Diagnostics	99335191
cAMP Assay Kit	Abcam	ab65355
Pierce BCA Protein Assay Kit	ThermoFisher	23225
PKA Kinase Activity Assay Kit	Abcam	ab139435
Experimental Models: Cell Lines		
C3H/10T1/2, Clone 8 (ATCC® CCL-226)	ATCC American Type Culture Collection	ATCC CCL-226
Experimental Models: Organisms/Strains		
C57BL/6J mice	Jackson Laboratory	JAX:000664
Ager Flox/ Flox mice	N/A	N/A
Adipoq Cre Recombinase mice	Jackson Laboratory	28020
Ager null mice	N/A	N/A

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Continued

REAGENT or RESOURCE	SOURCE	IDENTIFIER
Recombinant DNA		
pGL3- UCP1	Koza Laboratory	N/A
PGL3 Basic	Promega	E1741
Software and Algorithms		
Graphpad Prism 8.02	GraphPad Software	https://www.graphpad.com
Leica Digital Image Hub	Leica Biosystems	https://www.leicabiosystems.com
ImageJ	NIH	https://imagej.nih.gov
Fiji (distribution of ImageJ)	Schindelin et al., 2012	https://fiji.sc
WAVE Seahorse XFe24 Analyzer	Agilent	https://www.agilent.com
LI-COR Odyssey Classic	Li-Cor	https://www.licor.com
Trans-Blot Turbo Transfer System	Bio-Rad	http://www.bio-rad.com
Image Studio Lite	Li-Cor	https://www.licor.com

LEAD CONTACT AND MATERIALS AVAILABILITY

Further information and request for resources and reagents should be directed to the lead Contact Ann Marie Schmidt (AnnMarie.Schmidt@nyulangone.org).

EXPERIMENTAL MODEL AND SUBJECT DETAILS**Mouse Strains**

All animal procedures were approved by the New York University, University of Massachusetts Medical School and Sanford Burnham Institute (Lake Nona, Orlando FLA) Institutional Animal Care and Use Committees and performed in accordance with the National Institutes of Health Animal Care Guidelines. Homozygous *Ager* null mice ([Liliensiek et al., 2004](#)) (C57BL/6 *Ager* null mice backcrossed > 20 generations into C57BL/6J [The Jackson Laboratory, Bar Harbor, ME]) and their littermate *Ager*-expressing controls or C57BL/6J were used. In order to generate adipocyte-specific *Ager*-deleted mice, we developed a floxed mouse line, suitable for the generation of a conditional deletion model for the *Ager* gene (GenOway, Lyon, France). *Ager*^{flox/flox} mice were bred with Adiponectin (*Adipoq*) Cre-recombinase mice to yield *Ager*^{flox/flox} *Adipoq* Cre (+) and *Ager*^{flox/flox} *Adipoq* Cre (–) littermate mice, resulting in deletion of *Ager* from both white and brown adipose tissue depots. Adiponectin Cre-recombinase were obtained from the Jackson Laboratory (Stock No: 028020), which were in the C57BL/6J background. C57BL/6J (WT) mice were obtained from The Jackson Laboratory (Stock No: 00664). All mice studied were male, had free access to food and water, unless otherwise noted in specific fasting/refeeding or fasting state studies, and were subjected to 12 h light/dark cycles. Mice were bred, born and housed in a pathogen-free barrier environment, which was negative for *Helicobacter* or Norovirus pathogens. Mice were housed in standard caging up to 4 mice per cage and, where, indicated, were single housed for the indicated periods of time in order to perform specific studies such as indirect calorimetry and measurement of food intake that required the presence of only one mouse per cage. At 4–5 weeks of age, mice were anesthetized with 1% isoflurane, and tail snips were analyzed by PCR. To confirm adipose-specific deletion of *Ager*, fat depots (eWAT, iWAT, and iBAT) from offspring were isolated and purification of total genomic DNA (gDNA) was performed according to the manufacturer's instructions (DNeasy Blood and Tissue, QIAGEN). Thereafter, gDNA was subjected to PCR analysis using primers spanning the deletion site (exon 6) in the *Ager* gene (GenBank Accession number NM_007425.3) forward: 5'-GGTACCCTGCACCCAACTAC-3' and reverse: 5'-TTGATGGCCCTGGGATTGAC-3'. The following conditions with Platinum TM taq DNA polymerase from Invitrogen were used for PCR: 95°C for 30 s, 58°C for 30 s, and 72°C for 1 min (29 cycles). The resulting PCR products were separated using agarose gel electrophoresis, and the bands were visualized. The primary data illustrating the targeting vector construct and the characterization of these mice are shown in [Figures S3A–S3C](#). In all cases, within genotypes, mice were randomly assigned to study groups upon review of a database, which indicates mouse date of birth, sex, identification number and body weight. Randomization and study group assignment was performed with the assistance of technical support staff in the laboratory who were not involved in the design or protocol of the specific experiments.

Cells Used in the Study

C3H10T1/2 cells, purchased from ATCC (Manassas, VA), catalog number ATCC CCL-226, were plated at low density and cultured in DMEM containing 10% (vol/vol) FBS and 1% (vol/vol) penicillin-streptomycin. Two days post-confluence (designated day 0), cells were induced to differentiate with DMEM containing 10% (vol/vol) FBS, 1 μg/ml insulin, 2 nM T3, 0.5 mM isobutylmethylxanthine, 1 μM dexamethasone and 1 μM Rosiglitazone until day 2-5. After this induction phase, complete medium was supplemented with

insulin and T3 until day 8. Before any treatment, cells were starved for 3 h in 2% fatty acid free Krebs-Ringer bicarbonate (KRB) buffer. For treatments to stimulate RAGE, CML-AGE, 300 $\mu\text{g/ml}$, was added to the media for the indicated time periods.

Primary adipocytes were derived from eWAT, iWAT and iBAT of male mice from the lines of animals described above. Fat depots were retrieved and immediately washed with PBS, minced and digested with collagenase type II (1 mg/ml), 45 mins, 37°C) in DMEM containing 1% (wt/vol) BSA. Digested tissues were filtered through sterile 100 and 40 μm cell strainers and centrifuged (2000 rpm, 10 min). The floating fractions containing adipocytes were separated from the SVF and stored at -80°C with QIAzol (QIAGEN) until further analysis. The cells were centrifuged at 2000 rpm for 5 mins, plated on a 12-well culture dish, and grown and differentiated to mature adipocytes. Briefly, cells were grown to confluence in DMEM, 10% (vol/vol) FBS and 1% (vol/vol) penicillin-streptomycin followed by standard adipogenic induction (1 $\mu\text{g}/\mu\text{l}$ insulin, 2 nM T3, 0.5 mM isobutylmethylxanthine, 1 μM dexamethasone, and 1 μM rosiglitazone). After 2-5 days induction phase, complete medium was supplemented with insulin and T3, until day 8. Before any treatment, cells were serum-starved for 1-3 h in either DMEM or KRB buffer containing 2% fatty acid free BSA.

METHOD DETAILS

Fasting experiments

Fasting and feeding studies were performed in mice fed a standard chow diet (5053 PicoLab Rodent Diet 20; LabDiet, Brentwood, MO). To determine the effect of nutritional status, 11-week-old male *Ager* null mice and WT mice were fed *ad libitum*, fasted for 24 h, or re-fed for 24 h after the 24 h fast.

Cold-induced thermogenesis and thermoneutrality studies

Cold-induced thermogenesis over 48 h was assessed at the Cardiometabolic Phenotyping Core of the Sanford Burnham Institute (Lake Nona, Orlando FL) where *Ager* null mice and WT mice were individually housed in home cages with *ad libitum* access to food and water. The cages were placed in an environmentally-controlled chamber (Powers Scientific, Inc.) with the temperature set at 23°C (standard housing temperature). The temperature was then gradually lowered to 4°C over a span of ~ 6 h. Mice remained in the chamber at 4°C for 48 h. During this 48 h period, core body temperature was measured three times per day (10:00, 14:00 and 17:00) using a rectal probe attached to a digital thermometer. Short-term cold exposure (4°C for 24 h) thermogenesis studies were carried out at NYU Medical Center. *Ager* null, WT, *Ager*^{*fllox/fllox*} *Adipoq* Cre (+) and *Ager*^{*fllox/fllox*} *Adipoq* Cre (-) male mice 8 weeks old were exposed to 4°C for 24 h. Core body temperature was measured every 6 h using a rectal probe (ThermoWorks, Alpine UT). For thermoneutrality studies, male *Ager* null and WT mice, 8 weeks of age, were housed at 30°C for 24 h in individual cages. Core body temperature was measured every 6 h using a rectal probe (ThermoWorks, Alpine UT). At 24 h, mice were removed and immediately euthanized for tissue collection.

Diet-induced obesity and indirect calorimetry

For HFD studies *Ager*^{*fllox/fllox*} *Adipoq* Cre (+) and *Ager*^{*fllox/fllox*} *Adipoq* Cre (-) were fed a HFD with 60% of calories from lard (D12492; Research Diets, Inc., New Brunswick, NJ) or low-fat diet (LFD) with 13% of calories from fat (5053 PicoLab Rodent Diet 20; LabDiet, Brentwood, MO). Metabolic studies in *Ager*^{*fllox/fllox*} *Adipoq* Cre (+) and (-) mice were performed at the National Mouse Metabolic Phenotyping Center (MMPC) at UMass Medical School (Worcester, MA) and metabolic studies in mice surgically transplanted with adipose were performed at NYU Medical Center. Metabolic cages (TSE Systems, Inc., Midland, MI) were used in conscious mice to simultaneously measure energy expenditure through indirect calorimetry analyses, food/water intake, and physical activity at before (baseline at 8 weeks of age) and after the indicated periods of HFD (Dagdeviren et al., 2016; Lee et al., 2015a).

Dual Energy X-Ray (DEXA) Absorptiometry

DEXA scans were employed to determine body composition. Before each scan session, the instrument was calibrated as per the manufacturer's direction. Mice were then weighed, briefly anesthetized via isoflurane inhalation and placed on a scanning area of a Lunar PIXImus DEXA instrument (PIXImus, WI). Mice remained immobilized on this surface for the duration of the entire scan. At least 3 scans per mouse were performed and the average lean and fat mass were recorded.

Soluble (s) RAGE

Soluble RAGE (sRAGE) (human) was prepared, purified, and rendered free of endotoxin as previously described (Park et al., 1998) and 100 $\mu\text{g/day}$ by intraperitoneal (IP) route was administered to mice beginning immediately at the time of HFD feeding at age 6 weeks. Control mice were treated with equal amounts of vehicle PBS.

Glucose and Insulin Tolerance Tests

Glucose (GTT) and insulin tolerance tests (ITT) were performed on mice after fasting for 5-6 h and acclimated to the testing room with access to water. Mice were injected intraperitoneally (IP) with either D-glucose (Sigma, 0.75-1.5 g per kg of body weight) or recombinant human regular insulin (0.5-1 U per kg of body weight); blood glucose levels were measured via tail sampling after fasting for a baseline measurement ($t = 0$) and again at $t = 15, 30, 60, 90$ and 120 mins post glucose or insulin bolus using a glucometer (Freestyle Freedom Line, Abbott Diabetes Care Inc, Alameda CA).

iBAT and iWAT transplantation

Following standard antiseptic surgical techniques, iBAT transplantation in the subcutaneous dorsal region was performed as described (Stanford et al., 2013) with the following modifications. iBAT pads were removed from 6-week old *Ager^{fllox/fllox} Adipoq Cre (+)* and *Ager^{fllox/fllox} Adipoq Cre (-)* male mice and transplanted into 6-week old WT recipient male mice. Specifically, after euthanasia of donor mice, bilateral iBAT was immediately excised, peripheral white fat was removed, and the remaining iBAT (~90 mg) washed in sterile PBS and transplanted into the subcutaneous anterior dorsal region of recipient mice as quickly as possible. With the same approach, bilateral iWAT (~250 mg) from two *Ager^{fllox/fllox} Adipoq Cre (+)* or *Ager^{fllox/fllox} Adipoq Cre (-)* donors was transplanted to a single recipient (2:1 donor:recipient) bilaterally into the posterior dorsal region of the recipient through small incisions of 2-3 mm in length beginning at the proximal end of the hind limb (Tran et al., 2008). Recipient mice were anesthetized with gas isoflurane and treated with buprenorphine during and after surgery for pain relief, mice were then monitored until recovered. In both cases, after 4 days of recovery from surgery, mice were fed a 60% HFD for the time course, as indicated in the figures. Mice were housed 4-5 per cage.

Protein extraction and western blot analysis

Protein was extracted from cells and tissue from eWAT, iWAT and iBAT or other cell or tissue lysates as indicated by using lysis buffer or RIPA (Cell Signaling) supplemented with complete protease inhibitor cocktail (Roche), and centrifuged at 13,000 g for 15 mins at 4°C. The supernatant was collected and protein concentration was measured using a bicinchoninic acid (BCA) protein quantification kit (Thermo Scientific, Waltham, MA). Samples were diluted to one constant protein concentration followed by addition of sample reducing agent and loading dye (Thermo Scientific, Waltham, MA). For western blot analyses, 20-50 µg of protein were separated in 4%-20% Mini-PROTEAN TGX gels (Bio-Rad, Hercules, CA) and blotted onto nitrocellulose membranes (Bio-Rad, Hercules, CA). Membranes were blocked at room temperature for 1 h with Odyssey LI-COR Blocking Buffer (LI-COR, Lincoln, NE) diluted 1:1 in Tris-buffered saline (TBS). Membranes were then incubated with primary antibodies (diluted 1:1000 in a 1:1 Blocking Buffer/TBS-T solution) overnight at 4°C. Primary antibodies detailed in the STAR methods key resource table were used. Membranes were washed consecutively 3 times for at least 5 mins each in TBS-T (0.1%). Blots were incubated with IRDye 680RD goat anti-mouse IgG and IRDye 800CW goat anti-rabbit IgG (LI-COR, Lincoln, NE) for 1 h at room temperature in blocking buffer containing 0.1% TBS-T and 0.1% SDS. Blots were washed 3 more times in TBS-T followed by a final wash in TBS, the blots were scanned with the LI-COR Odyssey Classic (LI-COR, Lincoln, NE) and quantified with Image Studio Lite software (LI-COR, Lincoln, NE) based on direct fluorescence measurement.

RNA analysis and Real-time quantitative PCR

Total RNA was extracted from cells and tissue samples using TRIzol and RNeasy kits (QIAGEN) followed by cDNA synthesis using iScript Reverse Transcription Supermix for RT-PCR (Bio-Rad). qPCR was performed using the TaqMan method (50°C for 2 mins, 95°C for 10 mins, and 40 cycles of 95°C for 15 s and 60°C for 1 min) with premade primer sets (Applied Biosystems). The relative abundance of transcripts was normalized according to the expression of *Actb* using the $\Delta\Delta C_t$ method. Primer sequences are listed in Table S1.

Mitochondrial DNA content

Genomic DNA was isolated from C3H10T1/2 cells using the DNeasy Blood & Tissue Kit (QIAGEN) following manufacturer's instructions. Quantitative real-time PCR assessed genomic expression of mitochondrial NADH dehydrogenase subunit 1 (*Nd1*) Fwd 5' GGCCACATGGCTACCATTTG 3' and Rev 5' CCGACCACTCACATTCACC 3' normalized to *Hbb* (β -globin) Fwd 5' ACACAAGGGACAAGAGGAATCC 3' and Rev 5' GGGTCAGGTCTTTGCTTCCA 3' in genomic extracts from adipocytes differentiated from C3H10T1/2 cells treated with vehicle or CL316243 (Sigma) with or without the small molecule RAGE inhibitor (RI); 2-[4-(acetylamino) phenyl] -4- Quinolinecarboxylic acid, methyl ester. MitoTracker® Red CMXRos (ThermoFisher) (75nM) was used to check the function of mitochondria in live iBAT-derived primary adipocytes from WT and *Ager* null mice at 37°C for 30 min to stain functional mitochondria, according to manufacturer's protocol.

XF24 Extracellular Flux Analysis (Seahorse)

SVF cells isolated from the indicated adipose tissue depots from WT and *Ager* null mice were seeded in 24-well Seahorse plates and differentiated, as described above. The oxygen consumption rate (OCR) of 8 days differentiated cells was determined with an XF24 Extracellular Flux Analyzer (Seahorse Bioscience, Proteogene, Saint Marcel, France). Isoproterenol (10 µM) was used to characterize β -adrenergic inducible respiration and injected before oligomycin was indicated. Uncoupled and maximum OCR were determined with oligomycin (1 µM) and FCCP (1.5 µM). Rotenone (1 µM) and antimycin A (1 µM) were used to inhibit complex I- and III-dependent respiration, respectively. OCR was determined by plotting the oxygen tension and acidification of the medium in the chamber as a function of time and normalized by protein concentration (picomoles per min per milligram), respectively, as per the manufacturer's instructions.

Assessment of Lipolysis

Differentiated adipocytes from iWAT or iBAT or differentiated C3H10T1/2 cells were starved for 3 h. Fresh medium (DMEM without Red Phenol) with 1% fatty acid free-BSA was added and the cells were immediately subjected to the indicated treatment (CL316243)

for the indicated times. Sampled medium was used to measure glycerol release with a free glycerol reagent (Sigma Aldrich), according to the manufacturer's instructions. Results were normalized to total sample protein content. For free fatty acid measurement, fully differentiated iWAT and iBAT-derived primary adipocytes were starved for the indicated times in KRB buffer containing 2% fatty acid free BSA (Roche) followed by treatment with either vehicle or CL316243 (Sigma Aldrich). Media and cell samples were collected at the indicated times and stored at -80°C until analyzed. Non-esterified free fatty acids (NEFA) from media were measured using a colorimetric NEFA assay (WAKO) per the manufacturer's protocol and normalized to total sample protein content.

Cell transfection and reporter assay

Undifferentiated C3H10T1/2 cells, clone 8 (ATCC® CCL-226) were transfected with pGL3 (Addgene) or complete *Ucp1* promoter luciferase reporter constructs (generously shared by Dr. R. Koza) and 220-bp BAT-specific enhancer and proximal promoter of the mouse *Ucp1* gene obtained by PCR amplification as described (Rim and Kozak, 2002) using lipofectamine 3000 (ThermoFisher). At 24 h after transfection, cells were stimulated with NE (5 μM) and CML-AGE (300 $\mu\text{g}/\text{ml}$) as indicated. Cell lysates were harvested 48 h after transfection, and luciferase activity was quantified by standard luminometer assay and normalized to Renilla signal (Renilla Luciferase Assay System, PROMEGA).

Measurement of cAMP levels

Differentiated adipocytes from iBAT iWAT and eWAT from WT and *Ager* null mice were grown with complete medium supplemented with insulin and T3. At day 7 of differentiation, complete medium was replaced with DMEM + 0.2% BSA for 16 h. The cells were then treated with serum free DMEM \pm 10 μM CL316,243 for 10 mins. Cells were collected, lysed, and intracellular cAMP levels were measured with cAMP Assay Kit (Competitive ELISA) (ab65355). cAMP levels were normalized to protein content.

Protein Kinase A Activity Assay

Differentiated primary brown adipocytes from WT mice were treated overnight with vehicle or RAGE ligand, CML-AGE (300 $\mu\text{g}/\text{ml}$) followed by stimulation with CL316,243 for 15 mins. Cells were washed twice with RIPA buffer. Fifteen μg of cell lysate protein were used for the kinase reaction according to the manufacturer's instructions (Enzo Life Science, NY). For the negative control, only the RIPA buffer was used in the reaction mixture. Data were normalized to total protein content.

Histology and Immunohistochemistry

For staining of adipose tissue for UCP1 antigen, iBAT, iWAT and eWAT samples were dissected and fixed in 4% paraformaldehyde (PFA) overnight or up to 48 h at 4°C . Samples were then washed at least three times in PBS and processed for paraffin embedding. Embedded blocks were sectioned at 5 μm thickness and placed onto histology slides. Slides were deparaffinized (Leica AR9222 dewax) and stained on a Leica BondRX® autostainer. Prior to the incubation with the primary antibody, sections underwent heat retrieval with Bond Epitope Retrieval Buffer 2 (Leica ER2, AR9640), and incubation with hydrogen peroxidase to eliminate endogenous peroxidase activity. To achieve an appropriate range of staining with different sample subsets, the primary antibody, UCP1 (Abcam ab23841) was diluted ((Leica diluent, ARD1001EA) 1:100 and 1:500 for iWAT and iBAT, respectively) and stained for 12 h. Sections were subsequently incubated with Leica anti-rabbit polymer conjugated to horse radish peroxidase (DAB KIT, Leica DS9800) and developed with diaminobenzidine. After counterstaining with hematoxylin, the sections were imaged on a Leica SCN400 whole slide scanner up to 40x magnification. Using the Slidepath Digital Imaging Hub software (Leica Biosystems), 3 randomly-selected section images per animal were captured at 20x magnification and used for quantification of UCP1 by an experimenter naive to the identification of the experimental groups.

Immunofluorescence microscopy

C3H/10T1/2 cells or SVF-derived adipocytes isolated either from WT or *Ager* null mice were grown on coverslips to obtain fully differentiated primary adipocytes. Cells were fixed with 4% PFA for 20 mins. After 5 mins of Triton X-100 0.2% permeabilization and 15 mins of blocking (PBS-BSA 3%), lipid droplets were stained using BODIPY or LipidTox for the indicated times. After washing 10 times with 1x PBS, cells were incubated with DAPI (Life Technologies) for 5 mins at room temperature. Cells were then washed extensively with 1x PBS and mounted on glass slides with Prolong-Gold (Life Technologies). All gains for the acquisition of comparable images were maintained constant and the Leica 5500B microscope was used to capture the images. Analysis of the images was performed using ImageJ (NIH) and Adobe Photoshop CC 2015.

Image analysis and quantification

MacBiophotonics ImageJ was used for image quantification and analysis of LipidTOX/BODIPY staining. To quantify the staining of lipid droplets with LipidTOX/BODIPY, the immunofluorescence of this dye was recorded as a digital, 8-bit gray-scale, 1024x1024 resolution images. The lipid droplet labeling image threshold was set at 90 on a 0–255 black to white scale (all pixels with a value under 90 were excluded from the quantification) to remove background pixels from measurement. The number of individual cells in every image was counted by DAPI staining. Quantification of LipidTOX/BODIPY staining in each condition is represented as mean intensity value divided by number of nuclei.

The Fiji distribution of ImageJ was used for image quantification and analysis of UCP1 immunostaining. To quantify the area of UCP1 staining, the images were first de-convoluted to separate the DAB (3,3'-diaminobenzidine) stain from the hematoxylin counter-stain using the Fiji plug-in Color De-convolution 1.7 (vector-H DAB). The resulting image was then subjected to automated thresholding using the Otsu method (Otsu, 1979; Ruifrok and Johnston, 2001; Schindelin et al., 2012). Quantification of UCP1-positive area was calculated as μm^2 and the average of three images per animal was used for statistical analysis.

QUANTIFICATION AND STATISTICAL ANALYSES

The Figure Legends indicate all of the subject numbers/replicates per study; the mean \pm standard error of the mean (SEM) is reported. The of details the specific statistical analyses based on the conditions and comparisons are listed in each figure legend and employed methods such as Student's t test or ANOVA followed by the appropriate and indicated post hoc tests. In the event that variances were statistically different and thus not meeting the requirement of normal distribution, a non-parametric Mann-Whitney U test was employed. P values of < 0.05 were determined *a priori* as the threshold for statistical significance. Data analysis was performed using GraphPad Prism Software Version 8.02 (GraphPad, San Diego, CA).

DATA AND CODE AVAILABILITY

We have not generated software and datasets in this manuscript.

ADDITIONAL RESOURCES

We have not generated a new website or forum.

Cell Reports, Volume 28

Supplemental Information

A Receptor of the Immunoglobulin Superfamily

Regulates Adaptive Thermogenesis

Carmen Hurtado del Pozo, Henry H. Ruiz, Lakshmi Arivazhagan, Juan Francisco Aranda, Cynthia Shim, Peter Daya, Julia Derk, Michael MacLean, Meilun He, Laura Frye, Randall H. Friedline, Hye Lim Noh, Jason K. Kim, Richard A. Friedman, Ravichandran Ramasamy, and Ann Marie Schmidt

Supplemental Information

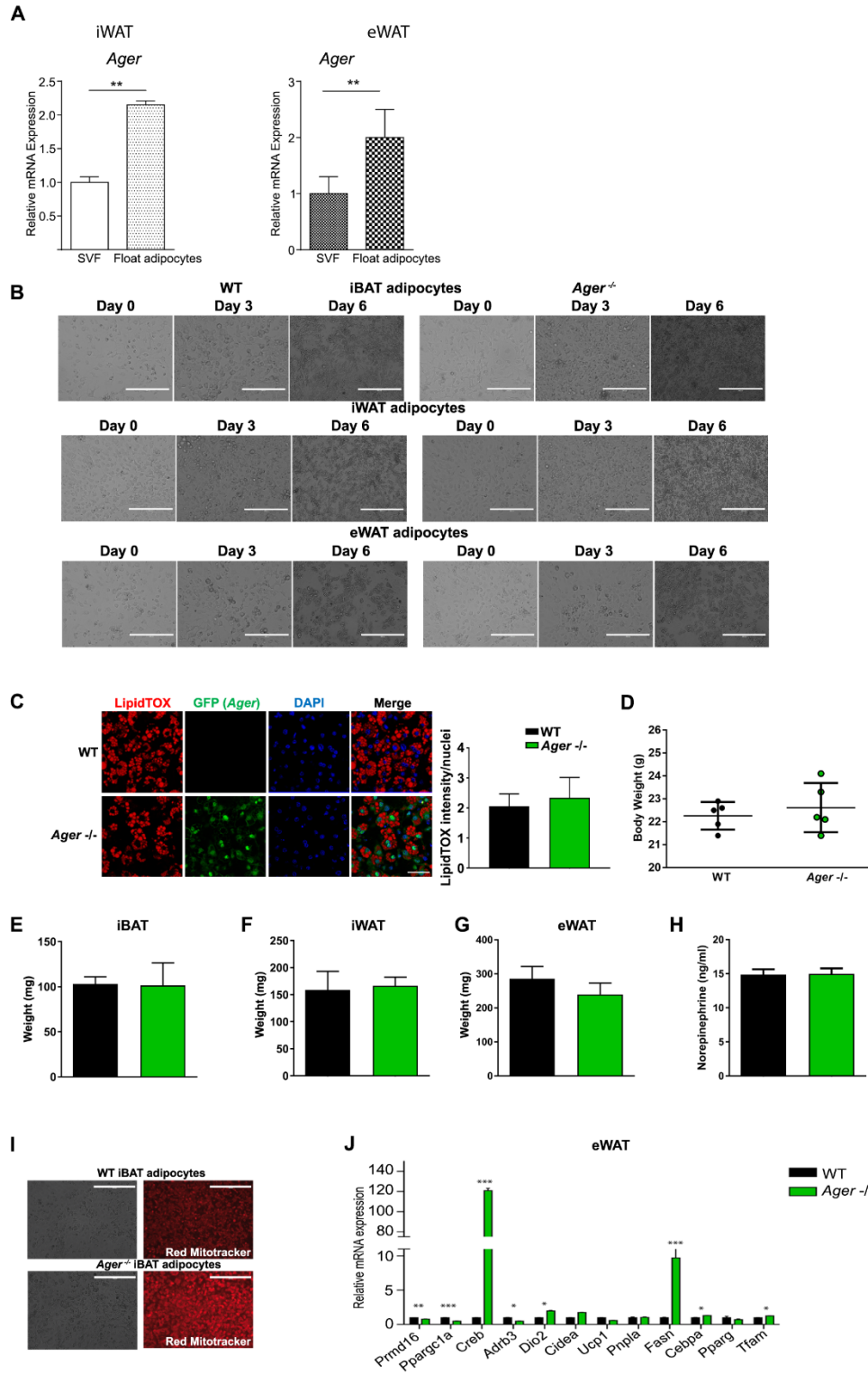


Figure S1. RAGE, primary adipocyte differentiation and adiposity. Related to Figure 1. (A) iWAT (left) and eWAT (right) were retrieved from WT mice and subjected to centrifugation for separation of floating adipocytes and

SVF. Real-time qPCR was performed for detection of relative *Ager* mRNA transcripts. The mean relative mRNA expression \pm SEM is reported in adipocytes derived from N=5 mice/group. Data analysis was performed using independent samples two-tailed Student t-test. **(B)** Representative micrographs of primary adipocytes differentiated from WT or *Ager* null iBAT (top), iWAT (middle) and eWAT (bottom) SVF of N=4 mice/group. Differential interference contrast micrographs of primary cells throughout differentiation (on day 0, 3 and 6). Scale bar: 400 μ m. **(C)** Primary adipocytes of eWAT retrieved from WT and *Ager* null mice were differentiated over 7 days and stained with LipidTOX and DAPI. GFP fluorescence denotes *Ager* null cells as *Ager* is replaced by *Gfp* in these mice. Scale bar: 20 μ m. Data are plotted as LipidTOX mean intensity/nuclei. The mean \pm SEM is reported. Data were analyzed using independent samples two-tailed Student's t-test. There were no statistically significant differences between the groups. **(D)** Body weight was recorded in 8-week old male WT or *Ager* null mice fed standard chow at room temperature. The mean \pm SEM is reported in N=5 mice/group and analyzed using independent samples two-tailed Student's t-test. **(E)** iBAT, **(F)** iWAT and **(G)** eWAT were retrieved from 8-week old standard chow fed WT and *Ager* null mice and weighed. The mean tissue weight \pm SEM is reported in N=5 mice/group and analyzed using independent samples two-tailed Student's t-test. **(H)** Plasma norepinephrine (NE) levels were measured in 8-week old WT or *Ager* null mice fed standard chow at room temperature. The mean NE levels \pm SEM are reported in N=5 mice/group and analyzed using independent samples two-tailed Student's t-test. **(I)** MitoTracker staining was performed in WT and *Ager* null primary iBAT adipocytes at day 8 of differentiation; representative micrograph is shown. Scale bar: 400 μ m. **(J)** eWAT was retrieved from 8-week old WT or *Ager* null mice and subjected to RT-qPCR to assess the relative mRNA expression of *Prdm16*, *Ppargc1a*, *Creb*, *Adrb3*, *Dio2*, *Cidea*, *Ucp1*, *Pnpla2*, *Fasn*, *Cebpa*, *Pparg* and *Tfam*. The mean relative mRNA expression \pm SEM is reported for adipocytes derived from N=5 mice/group. Data for each gene were separately analyzed using independent samples two-tailed Student's t-test. In this figure, * p <0.05, ** p <0.01 and *** p <0.001. For results presented in C-H & J, WT (black bars/circles) and *Ager* null (green bars/circles).

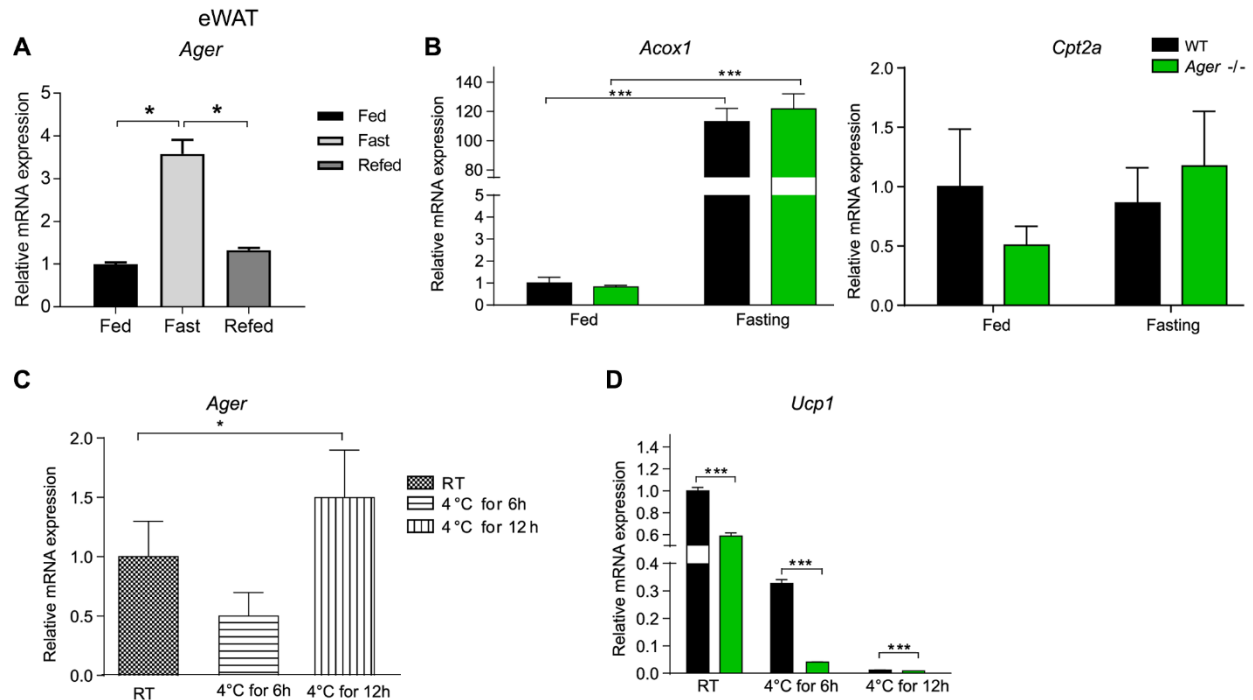


Figure S2. RAGE and effects on fasting and cold challenge. Related to Figure 2. (A) eWAT was retrieved from 11-week old WT mice in a fed, fasted (24 hours) or 24 hours refed state and subjected to RT-qPCR for detection of relative *Ager* mRNA expression. The mean relative mRNA expression \pm SEM is reported in N=5 mice/group. Data were analyzed using an independent samples one-way ANOVA followed by a *post hoc* Tukey's HSD test. (B) The relative expression of fatty acid oxidation-related genes *Acox1* and *Cpt2a* mRNA was measured by RT-qPCR in eWAT of fed and fasted (24 hours) 11-week old WT and *Ager* null mice. The mean relative mRNA expression \pm SEM is reported in N=4 mice/group. Data were analyzed using independent samples two-way ANOVA followed by *post hoc* Bonferroni test where appropriate. (C) eWAT was retrieved from WT mice immediately after exposure to 6 or 12 hours 4°C cold challenge and subjected to RT-qPCR for detection of relative *Ager* mRNA expression. The mean relative mRNA expression \pm SEM is reported in N=5 mice/group. Data were analyzed using independent samples one-way ANOVA followed by *post hoc* Tukey's HSD test. (D) Cold challenge was performed as in (C) for WT and *Ager* null male mice and immediately after sacrifice, eWAT was retrieved and subjected to RT-qPCR for detection of relative *Ucp1* mRNA expression. The mean \pm SEM is reported in N=5 mice/group. Data were analyzed using independent samples two-way ANOVA followed by *post hoc* Sidak HSD test; *** p <0.001. In this figure, * p <0.05 & *** p <0.001. For results presented in B and D, WT (black bars) and *Ager* null (green bars).

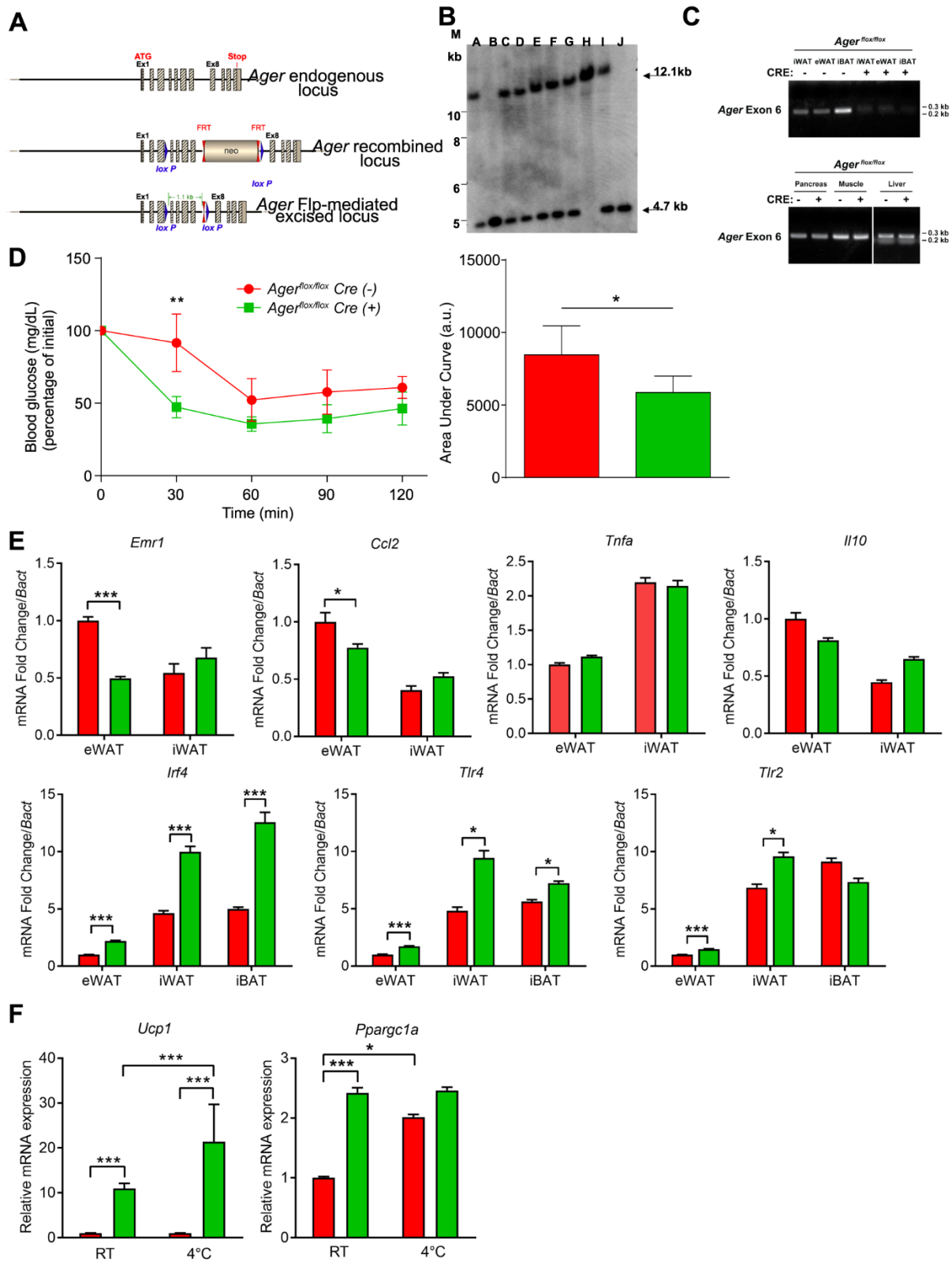


Figure S3. Generation of *Ager* floxed mice and characterization of adipose tissue inflammation and thermogenic programming. Related to Figure 3. (A) Bioinformatics analysis of *Ager* cDNA sequence

(NM_007425) indicated that the exon / intron organization of the gene revealed 11 exons separated by 10 introns. Schematic illustrating *Ager* targeting strategy leading to the insertion of a *loxP* site together with an FRT flanked neomycin selection cassette within the intron 7 and a single distal *loxP* within the intron 3 of *Ager*. This was followed by homologous recombination in embryonic stem (ES) cells (directly into C57BL/6), blastocyst injection, generation and identification of chimeric mice and breeding with *Flp* deleter mice to remove the neomycin cassette. Breeding with Cre recombinase mice thus excises the *loxP* flanked *Ager* genomic region (exons 4 to 7), thereby deleting functional RAGE. **(B)** Southern blot validation of *Ager* heterozygous and homozygous null mice. Southern blotting was performed on KpnI digested genomic DNA from WT or homozygous or heterozygous *Ager* deleted mice. In this figure, the hybridization figure corresponding to the *Ager* deleted allele is 12.1kb and signal corresponding to the *Ager* WT allele is 4.7 kb. Thus, in this Southern blot, lane H indicates homozygous *Ager* null mouse; Lanes A, C-G and I indicate heterozygous *Ager* deleted mice, and Lanes B and J are WT mice. **(C)** *Ager* floxed mice were bred with Adiponectin (*Adipoq*) Cre recombinase mice. Genomic DNA was prepared from eWAT, iWAT, iBAT, pancreas, skeletal muscle and liver of *Ager^{fllox/fllox} Adipoq Cre (-)* or *Cre (+)* mice and subjected to PCR to assess specific detection of *Ager* exon 6. The primers were designed between exon 5 and exon 6 (as the flox sequences are designed between exons 3 and 7) as follows: Primer Pair: Forward Primer: GGTACCCTGCACCCAACTAC and Reverse Primer: TTGATGGCCCTGGGATTGAC (Product length: 302 base pairs). Note that only in the *Cre (+)* mice was *Ager* selectively deleted in eWAT, iWAT and iBAT, but not in pancreas, skeletal muscle or liver whereas in *Cre (-)* mice. **(D)** Insulin tolerance test was performed in the indicated *Ager^{fllox/fllox} Cre (-)* or *Cre (+)* fed HFD and reported as % change in level of glucose from baseline and the area under the curve. The mean percent change \pm SEM is reported in N=4-5 mice/group. Data were analyzed using a repeated measures two-way ANOVA followed by a *post hoc* Bonferroni test or an independent samples Student t-test (for AUC). **(E)** *Ager^{fllox/fllox} Cre (-)* or *Cre (+)* mice were fed a HFD for 12 weeks and eWAT, iWAT or iBAT were subjected to RT-qPCR for detection of mRNA transcripts expression for *Emr1*, *Ccl2*, *Tnfa*, *Il10*, *Irf4*, *Tlr4* and *Tlr2*. The mean relative mRNA expression \pm SEM is reported in N=4 mice/group. Data analysis was performed using an independent samples Student t-test. **(F)** Mice were housed at room temperature or 4°C for 24 hours. eWAT was immediately retrieved at the end of the study and subjected to RT-qPCR for detection of relative *Ucp1* and *Pparg1a* mRNA expression. The mean relative mRNA expression \pm SEM is reported in N=4 mice/group. Data were analyzed using independent samples two-way ANOVA followed by a post-hoc *Sidak* test. In this figure, *p<0.05,

p<0.01 and *p<0.001. For results presented in D-F, *Ager^{flax/flax} Cre (-)* (red bars) and *Ager^{flax/flax} Cre (+)* (green bars).

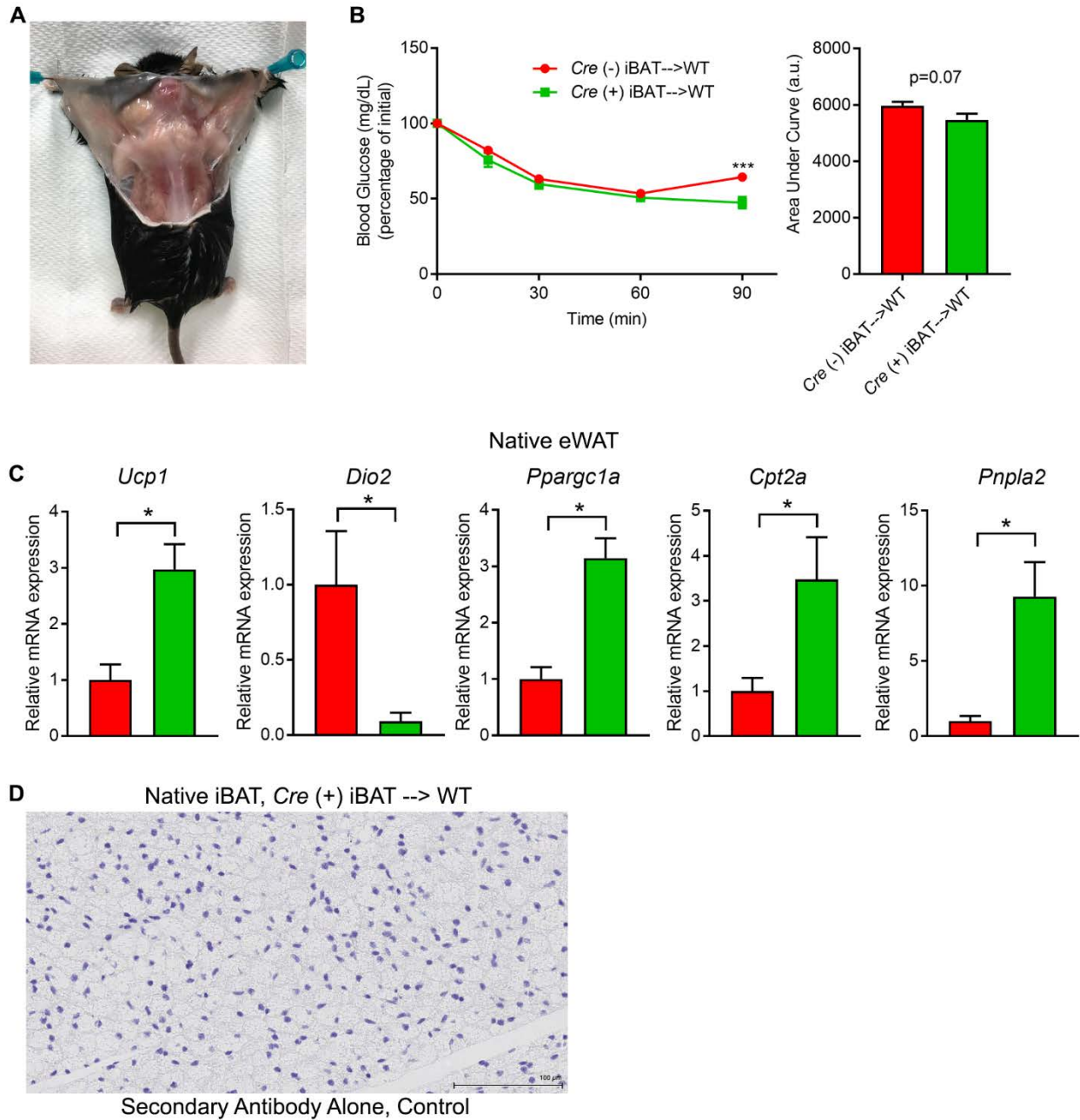


Figure S4. iBAT transplantation: effect of iBAT transplant devoid of adipocyte *Ager* to lean, male WT mice.

Related to Figure 4. At age 6 weeks, standard chow fed WT mice were surgically transplanted with iBAT derived from either *Ager*^{fllox/fllox} *Cre* (-) vs. *Ager*^{fllox/fllox} *Cre* (+) donor mice. 4 days later, mice were switched to HFD for an additional 20 weeks. **(A)** Representative photograph of iBAT transplanted graft. **(B)** Insulin tolerance test (ITT) was performed after 13 weeks of high fat feeding and the results plotted as percent change in level of blood glucose from baseline and as the area under the curve. For the percent change in glucose excursion, the mean±SEM is reported in N=10-11 mice/group. Data were analyzed using repeated measures two-way ANOVA followed by a *post hoc*

Bonferroni test or an independent samples Student t-test (for AUC); *** $p < 0.001$ & $p = 0.07$, as indicated. (C) At sacrifice, eWAT was retrieved from the recipient mice and subjected to RT-qPCR to assess the relative mRNA expression of *Ucp1*, *Dio2*, *Pparg1a*, *Cpt2a* and *Pnpla2*. The mean relative mRNA expression \pm SEM is reported in N=4-6 mice/group. Data analysis using an independent samples Student t-test or in the event that mean variances were statistically different, a *post hoc* Mann-Whitney U test was used. For results presented in B&C, *Ager^{fllox/fllox} Cre* (-) (red bars) and *Ager^{fllox/fllox} Cre* (+) (green bars). (D) Representative image of secondary antibody alone, control, for UCP1 immunohistochemistry in native iBAT post-transplantation with *Ager^{fllox/fllox} Cre* (+) iBAT. Scale bar: 100 μ m. In this figure, * $p < 0.05$ & *** $p < 0.001$.

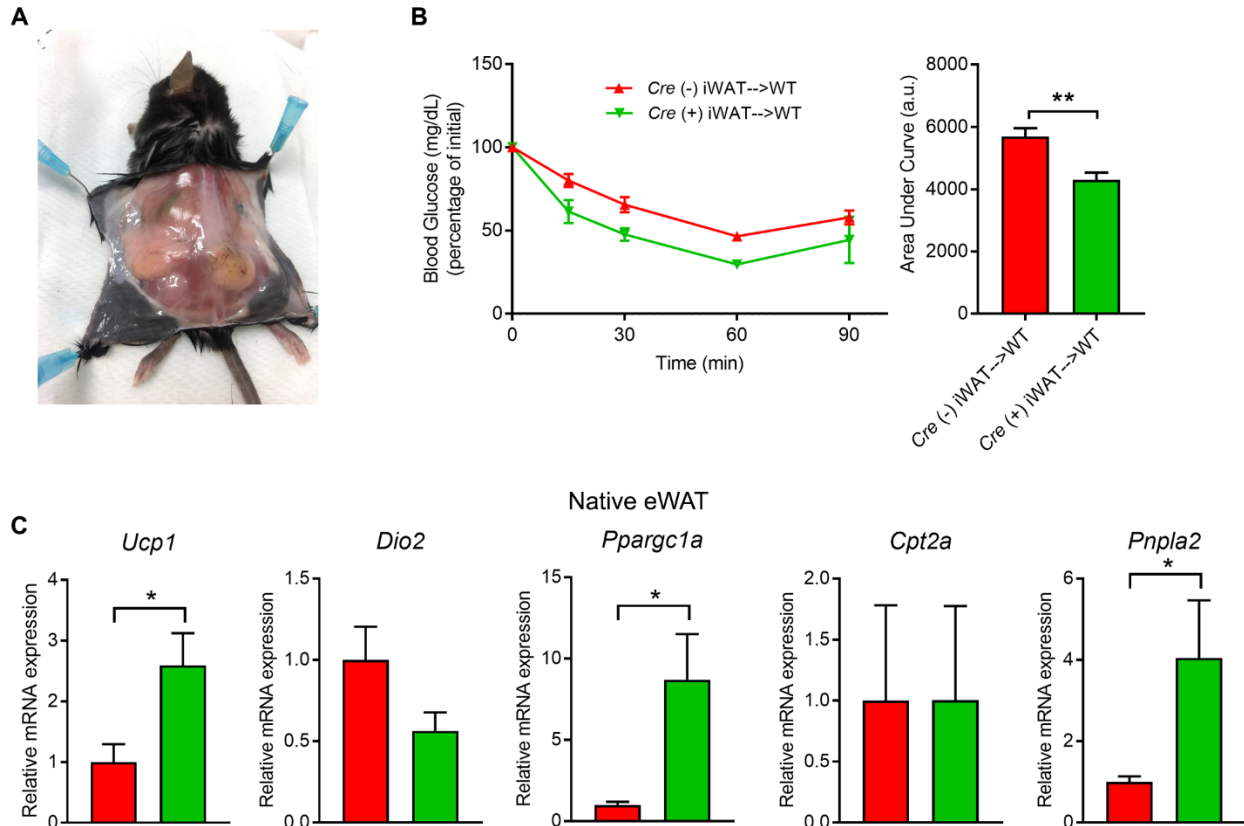


Figure S5. iWAT transplantation: effect of *Ager* devoid iWAT transplant to lean, male WT mice. Related to Figure 5. At age 6 weeks, standard chow fed WT mice were surgically transplanted with iWAT derived from either *Ager^{fllox/fllox} Cre (-)* vs. *Ager^{fllox/fllox} Cre (+)* donor mice. 4 days later, mice were switched to HFD for an additional 20 weeks. **(A)** Representative photograph of iWAT transplanted graft. **(B)** Insulin tolerance test (ITT) was performed after 13 weeks of high fat feeding and the results plotted as percent change in level of blood glucose from baseline and as the area under the curve. For the percent change in glucose excursion, the mean±SEM is reported in N=5-6 mice/group. Data were analyzed using repeated measures two-way ANOVA followed by a *post hoc* Bonferroni test or an independent samples Student t-test (for AUC). **(C)** At sacrifice, eWAT was retrieved from the recipient mice and subjected to RT-qPCR to assess the relative mRNA expression of *Ucp1*, *Dio2*, *Pparg1a*, *Cpt2a* and *Pnpla2*. The mean relative mRNA expression±SEM is reported in N=3-4 mice/group. Data analysis was performed using an independent samples Student t-test or in the event that mean variances were statistically different, a *post hoc* Mann-Whitney U test was used. For results presented in B&C, *Ager^{fllox/fllox} Cre (-)* (red bars) and *Ager^{fllox/fllox} Cre (+)* (green bars). In this figure, *p<0.05 & **p<0.01.

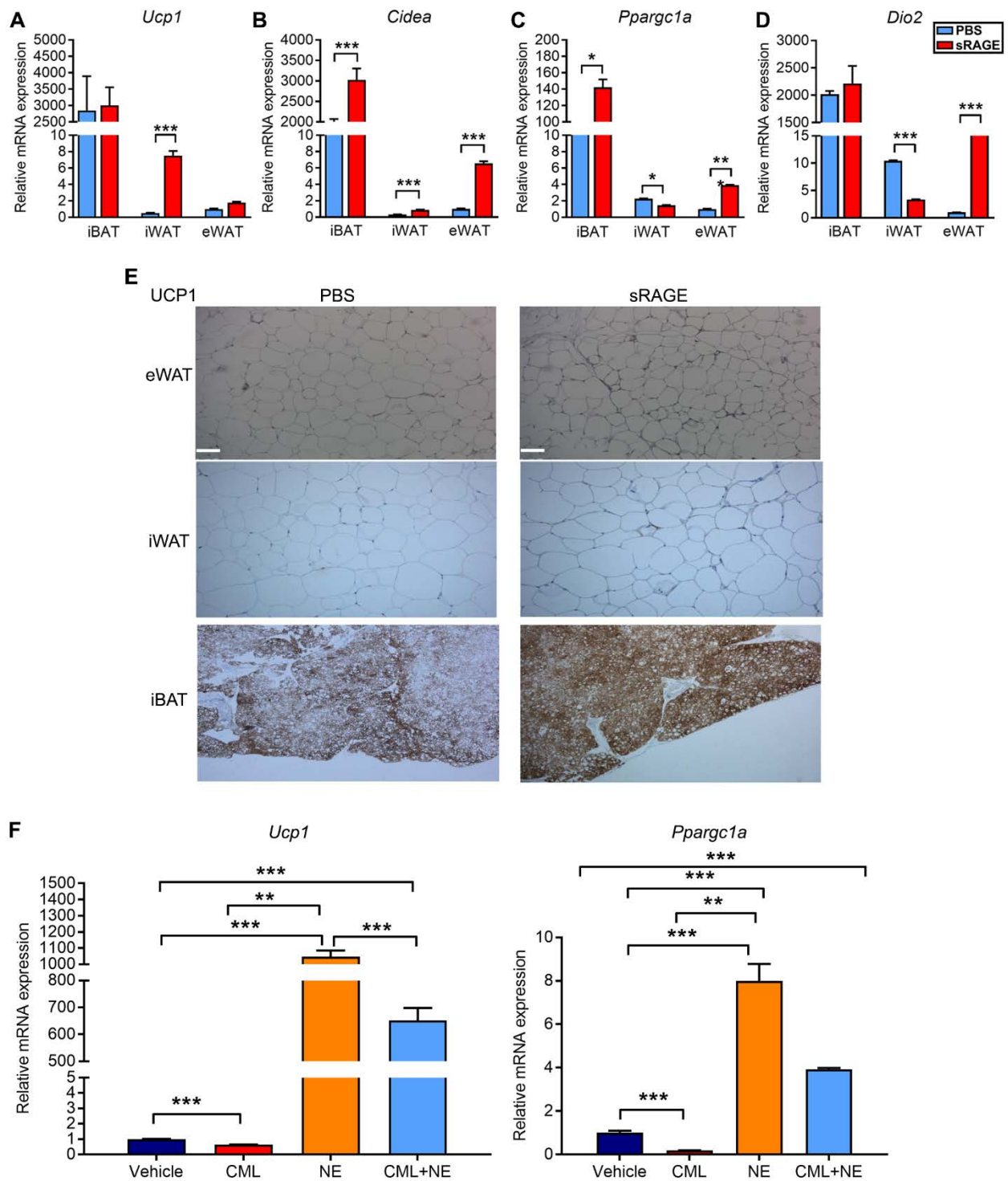


Figure S6. Effects of soluble RAGE *in vivo* and RAGE ligands on expression and induction of *Ucp1* and *Pparg1a* in eWAT-derived adipocytes. Related to Figure 6. (A-D) WT mice were fed a HFD beginning at age 6 weeks; simultaneously, vehicle (phosphate buffered saline, PBS) or recombinant sRAGE (100 μ g/day) was administered once daily for an additional 18 weeks. At sacrifice, eWAT, iWAT and iBAT were retrieved and

subjected to RT-qPCR for detection of (A) *Ucp1*, (B) *Cidea*, (C) *Ppargc1a* and (D) *Dio2* mRNA expression. The mean relative mRNA expression \pm SEM is reported in N=5 mice/group. Data were analyzed using independent samples Student t-test. (E) Representative images of eWAT, iWAT and iBAT stained for UCP1 after 18 weeks HFD and treatment with PBS vs. sRAGE. Scale bar: 100 μ m. (F) Primary adipocytes were differentiated from eWAT SVF of WT mice and incubated with vehicle or RAGE ligand CML-AGE (300 μ g/ml) for 16 hours alone or with NE (2 nM) for the final 6 hours of incubation. RT-qPCR for detection of relative *Ucp1* or *Ppargc1a* mRNA expression was performed. The mean relative mRNA expression \pm SEM is reported from cells originating from N=5 mice/group. Data were analyzed using independent samples one-way ANOVA followed by a *post hoc* Tukey's HSD test. In this figure, * p <0.05, ** p <0.01 and *** p <0.001.

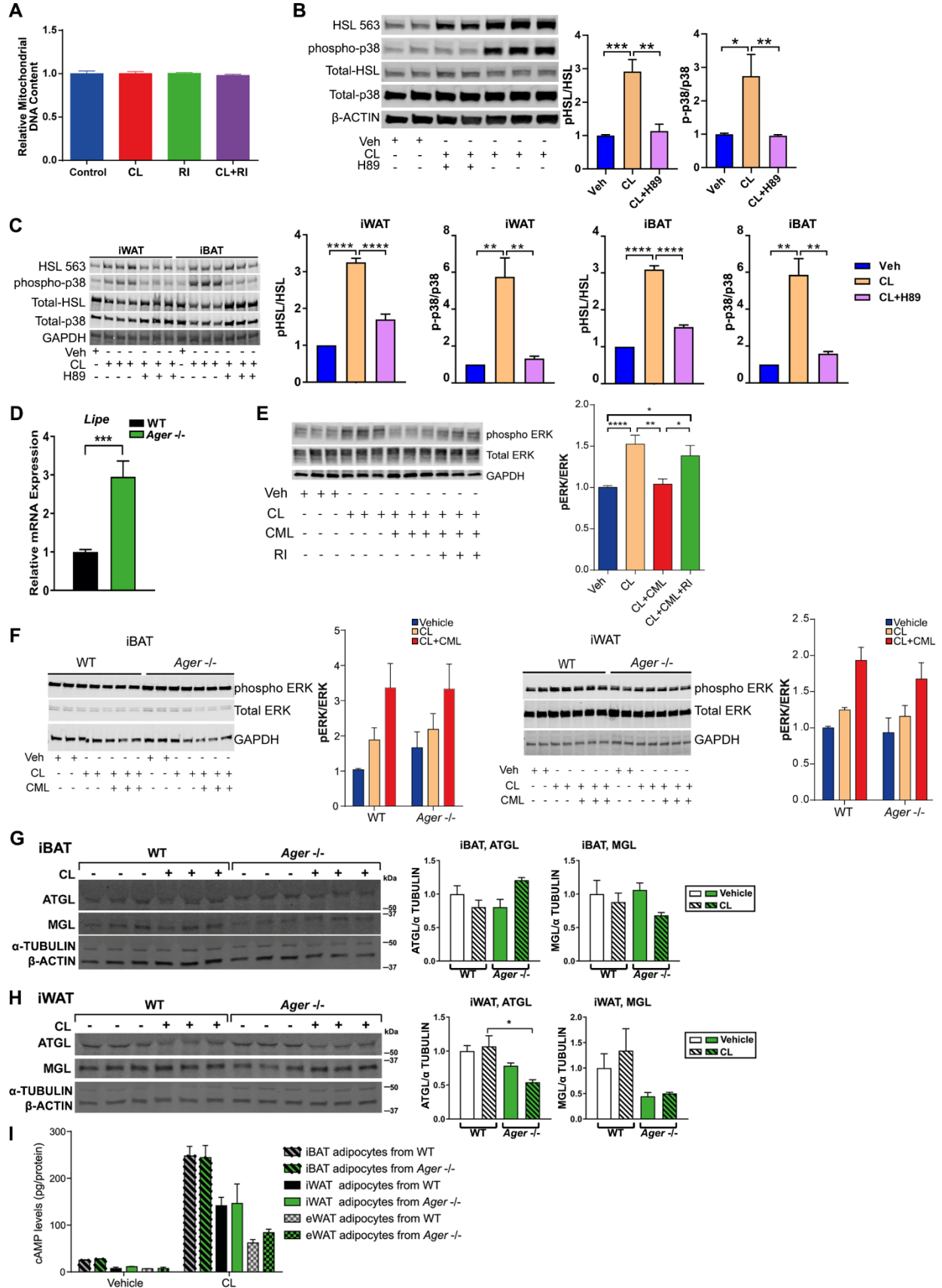


Figure S7. RAGE, intracellular levels of cAMP, expression of downstream lipolytic lipases and ERK MAPK regulation in adipocytes. Related to Figure 7. (A) Relative mitochondrial DNA content in C3H10T1/2 cells differentiated to adipocytes and treated with either CL316,243 (10 μ M) for 15 mins and/or the RAGE Inhibitor (RI) (1 μ M) for 30 mins was assessed by measurement of expression of mitochondrial gene, NADH subunit 1 (*Nd1*). Data are presented as relative mitochondrial DNA content \pm SEM and analyzed using independent samples one-way ANOVA. No statistically significant differences were observed. (B) C3H10T1/2 cells were differentiated into adipocytes and treated with vehicle or CL316,243 (10 μ M) for 15 minutes alone or after pre-treatment with the PKA inhibitor, H89 (20 μ M) for 30 mins. Cells were lysed and Western blotting performed for phosphorylated HSL Serine 563, phosphorylated p38, total HSL, total p38 and GAPDH. Band intensities were normalized to the respective total HSL or total p38 MAPK, as indicated, and the relative fold change is presented. The mean fold change \pm SEM of at least 2 independent studies with 2-3 technical replicates per experiment is shown. Data were analyzed using an independent measures one-way ANOVA followed by a *post hoc* Bonferroni test. (C) Primary adipocytes differentiated from iWAT (left) or iBAT (right) SVF of WT and *Ager* null mice were treated with vehicle or CL316,243 (10 μ M) for 15 minutes alone or after pre-treatment with the PKA inhibitor, H89 (20 μ M) for 30 minutes. Cells were lysed and Western blotting performed for phosphorylated HSL Serine 563, phosphorylated p38 MAPK, total HSL, total p38 MAPK and GAPDH. Band intensities were normalized to the respective total HSL or total p38 MAPK, as indicated, and the relative fold change is presented. The mean fold change \pm SEM is reported in cells derived from N=4 pooled mice/group in three independent experiments with at least two to three replicates per experiment. Data were analyzed using independent samples one-way ANOVA followed by a *post hoc* Tukey's HSD test. (D) Floating adipocytes were retrieved from the iWAT of WT or *Ager* null mice fed a HFD for 3 months and were subjected to RT-qPCR for detection of relative *Lipe* mRNA expression. The mean relative mRNA expression \pm SEM is reported. Data analysis was performed using independent samples Student's t-test. (E) C3H10T1/2 cells were differentiated into adipocytes and treated with vehicle, CML-AGE (300 μ g/ml) for 75 minutes alone or after pre-treatment with the RAGE inhibitor (1 μ M) for 30 minutes and with CL316,243 (10 μ M) for 15 minutes. Cells were lysed and Western blotting performed for phosphorylated ERK MAPK, total ERK MAPK and GAPDH. Band intensities were normalized to total ERK and the relative fold change is presented. The mean fold change \pm SEM of at least three independent studies with at least three technical replicates per experiment is shown. Data were analyzed

using an independent measures one-way ANOVA followed with a *post hoc* Bonferroni test. **(F)** Primary adipocytes differentiated from iBAT (left) or iWAT (right) SVF of WT and *Ager* null mice were treated with CL316,243 (10 μ M) for 15 minutes after pre-treatment with RAGE ligand CML-AGE (300 μ g/ml) for 1 hour. Cells were lysed and Western blotting performed for detection of phospho-ERK MAPK, total ERK MAPK and GAPDH. Band intensities were normalized to the respective total ERK and the relative fold change is presented as mean \pm SEM. The mean fold change \pm SEM is reported in cells derived from N=4 pooled mice/group in three independent studies with two to three technical replicates per experiment. Data were analyzed using independent samples one-way ANOVA followed by a *post hoc* Tukey's HSD test. No statistically significant differences were observed. **(G-H)** Primary adipocytes differentiated from **(G)** iBAT and **(H)** iWAT SVF retrieved from WT and *Ager* null mice were treated with CL316,243 (10 μ M) for 15 minutes and Western blotting was performed for detection of ATGL, MGL, α -TUBULIN and β -ACTIN. Detected bands were normalized to α -TUBULIN and the mean fold change from control \pm SEM is reported. Data were analyzed using independent samples two-way ANOVA followed by a *post hoc* Tukey's HSD test. **(I)** Primary adipocytes were differentiated from iBAT, iWAT and eWAT SVF of WT and *Ager* null mice and stimulated with CL316,243 (10 μ M) for 15 min. cAMP levels were measured from the cellular lysates. The mean level \pm SEM is reported in N=3 independent studies. Data analysis was performed using independent samples two-way ANOVA. Per adipose tissue depot, there were no statistically-significant genotype-dependent differences. In this figure, * p <0.05, ** p <0.01, *** p <0.001 & **** p <0.0001.

Supplemental Table S1. Related to STAR Methods.

Taqman Primers

	Assay ID	Catalog
<i>Acox1</i>	Mm00446408_m1	4448892
<i>Actb</i>	Mm00607939_s1	4352341E
<i>Adrb3</i>	Mm02601819_g1	4453320
<i>Ager</i>	Mm01134790_g1	4448892
<i>Ccl2</i>	Mm00441242_m1	4453320
<i>Cebp</i>	Mm00514283_s1	4453320
<i>Cidea</i>	Mm00432554_m1	4453320
<i>Creb</i>	Mm00457268_m1	4448892
<i>Cpt2</i>	Mm00487205_m1	4448892
<i>Dio2</i>	Mm00515664_m1	4453320
<i>Emr1</i>	Mm00802529_m1	4453320
<i>Fasn</i>	Mm00662319_m1	4453320
<i>Il10</i>	Mm01288386_m1	4453320
<i>Irf4</i>	Mm00516431_m1	4453320
<i>Pnpla2</i>	Mm00503040_m1	4453320
<i>Prdm16</i>	Mm00712556_m1	4453320
<i>Pparg</i>	Mm00440940_m1	4453320
<i>Ppargc1a</i>	Mm01208835_m1	4453320
<i>Tfam</i>	Mm00447485_m1	4453320
<i>Tlr2</i>	Mm00442346_m1	4453320
<i>Tlr4</i>	Mm00445273_m1	4453320
<i>Tnfa</i>	Mm00443258_m1	4453320
<i>Ucp1</i>	Mm01244861_m1	4453320