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Sex difference in the mouse BAT transcriptome reveals a role of progesterone

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Abstract

Brown adipose tissue (BAT) is a metabolically active organ that exhibits sex-differential features, that is, being generally more abundant and active in females than in males. Although sex steroids, particularly estrogens, have been shown to regulate BAT thermogenic function, the underlying molecular mechanisms contributing to sexual dimorphism in basal BAT activity have not been elucidated. Therefore, we assessed the transcriptome of interscapular BAT of male and female C57BL/6J mice by RNA sequencing and identified 295 genes showing ≥ 2 -fold differential expression (adjusted $P < 0.05$). *In silico* functional annotation clustering suggested an enrichment of genes encoding proteins involved in cell–cell contact, interaction, and adhesion. Ovariectomy reduced the expression of these genes in female BAT toward a male pattern whereas orchiectomy had marginal effects on the transcriptional pattern, indicating a prominent role of female gonadal hormones in this sex-differential expression pattern. Progesterone was identified as a possible upstream regulator of the sex-differentially expressed genes. Studying the direct effects of progesterone *in vitro* in primary adipocytes showed that progesterone significantly altered the transcription of several of the identified genes, possibly via the glucocorticoid receptor. In conclusion, this study reveals a sexually dimorphic transcription profile in murine BAT at general housing conditions and demonstrates a role for progesterone in the regulation of the interscapular BAT transcriptome.

Key Words

- ▶ brown adipose tissue
- ▶ gonadal steroid hormones
- ▶ primary cell culture
- ▶ RNA sequencing
- ▶ sex characteristics

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Introduction

A growing number of studies suggest that the prevalence of obesity and obesity-related diseases differs between women and men. Although the global prevalence of obesity is higher in women than in men (GBD Obesity Collaborators *et al.* 2017), obese men are more prone to develop obesity-related conditions, such as type 2 diabetes mellitus, than obese women (Logue *et al.* 2011). This sex

difference diminishes when women enter menopause, suggesting a prominent role for sex steroids in controlling adipose tissue function, since disturbances in adipose tissue function lead to obesity and associated metabolic diseases (Palmer & Clegg 2015, Longo *et al.* 2019).

In general, two types of adipose tissue with distinct physiological functions can be recognized.

While white adipose tissue (WAT) serves as an energy reservoir, brown adipose tissue (BAT) utilizes nutrient substrates for non-shivering thermogenesis, crucial for maintaining body temperature in small mammals and human infants (Townsend & Tseng 2012). In BAT, thermogenesis is activated by the sympathetic nervous system (SNS) that secretes norepinephrine (NE) upon stimuli such as cold exposure. BAT thermogenesis depends on uncoupling proton transport facilitated by the mitochondrial uncoupling protein 1 (UCP1) (Cannon & Nedergaard 2004). BAT plays a role in maintaining optimal energy balance, as illustrated by experiments in which transplantation of BAT into high-fat diet (HFD)-induced obese mice improved their body weight, fat mass, and glucose tolerance (Liu *et al.* 2013). However, conflicting results of the role of UCP1 in these beneficial metabolic effects exist. Feldmann *et al.* (2009) showed that UCP1-ablated mice housed at thermoneutral conditions become obese, whereas Maurer *et al.* (2020) demonstrated that UCP1-ablated mice did not gain weight upon HFD feeding but did develop impaired glucose tolerance. Upon confirmation of active BAT being present in adults, BAT has been studied extensively because BAT activation is considered a potential therapeutic option for obesity and metabolic syndrome (Moonen *et al.* 2019).

While sex differences in WAT and the role of sex steroids herein have been addressed by numerous studies (Newell-Fugate 2017), studies on sex differences in BAT functioning and its role in sex-dependent protection against metabolic diseases is largely lacking, although some studies suggest that sex differences may exist in the presence and activity of BAT (Mauvais-Jarvis 2015). It was observed that the prevalence of active BAT was about two-fold higher in women than in men (Cypess *et al.* 2009). Also in rodents, female rats have relatively higher BAT mass and UCP1 protein levels, and larger mitochondria in BAT than males (Rodriguez-Cuenca *et al.* 2002). Circulating sex steroid concentrations are one of the major differences between males and females of reproductive age and therefore are obvious candidates regulating sex differences in BAT activity. Indeed, the roles of sex steroids in sex-differential energy homeostasis have been demonstrated at many levels (Geer & Shen 2009, Mauvais-Jarvis 2015). Especially, the effects of estrogens on systemic energy balance and BAT activity have been studied in great detail (Mauvais-Jarvis *et al.* 2013, Gonzalez-Garcia *et al.* 2017). For example, ovariectomy in rodents led to an increase in food intake, weight gain, and lipid accumulation in

WATs, as well as reduced thermogenic activity in BAT (Yoshioka *et al.* 1988, Rogers *et al.* 2009). In addition, central or peripheral injection of 17 β -estradiol (E2) in ovariectomized mice induced BAT thermogenesis, demonstrating a role for estrogens in stimulating BAT activity via SNS activation (Martinez de Morentin *et al.* 2014). In contrast, orchietomy induced *Ucp1* mRNA expression and UCP1-positive staining in BAT coinciding with elevated body temperature, revealing an inhibitory effect of male gonadal hormones on BAT activity (Hashimoto *et al.* 2016). Treatment of cultured brown adipocytes with sex steroids confirmed that E2 facilitates while testosterone inhibits NE-induced lipolysis, an initial step of BAT thermogenesis (Monjo *et al.* 2003).

While sex steroids may control BAT thermogenic activity, the underlying molecular mechanisms contributing to the sex dimorphism in BAT function are still not fully elucidated. Therefore, we have performed RNA sequencing (RNA-seq) on the interscapular BAT of male and female mice to identify sex differences in the BAT transcriptome. Identified sex-differentially expressed genes were functionally clustered using biological databases. Gonadectomy (GDX) was performed to study whether sex steroids affected the expression of the identified genes. Progesterone was identified as one of the upstream regulators and we, therefore, studied the effects of progesterone on primary cultures of brown adipocytes isolated from male and female mice and on a brown adipocyte cell line.

Materials and methods

Animals and housing conditions

All animal experiments were approved by the Animal Ethics Committee at Erasmus MC, Rotterdam, the Netherlands. For the RNA-seq experiment, 8-week-old male and female C57BL/6J mice were purchased from Charles River Laboratories (Maastricht, Netherlands). Upon arrival, mice were housed at ~22°C on a 12 h darkness:12 h light cycle with chow diets and water available *ad libitum*. After 1-week acclimation, GDX or sham operation was performed as described previously (Kaikaew *et al.* 2017), followed by housing at standard conditions for 45 days. Next, mice were fasted for 4 h and euthanized by cardiac puncture under isoflurane anesthesia. BAT was snap-frozen and stored at –80°C until RNA isolation.

For adipose-derived stromal vascular fraction (SVF) cell culture, male and female C57BL/6J mice were bred in the Erasmus MC animal facility and housed under the

same conditions as described previously. At 20 weeks old, mice were euthanized and BAT was dissected and briefly stored in ice-cold PBS supplemented with Antibiotic-Antimycotic (Gibco, Life Technologies Europe) until SVF isolation.

RNA sequencing

Total RNA was isolated from BAT using the TriPure isolation reagent (Roche Diagnostics), contaminating genomic DNA was removed by the RQ1 RNase-free DNase (Promega Benelux), and the RNA library was prepared by the TruSeq RNA Library Prep Kit v2 (Illumina, Eindhoven, Netherlands). Paired-end sequencing with 50 base pairs in length was performed using a HiSeq 2000 sequencer (Illumina). Adapter sequences were trimmed with Cutadapt (version 1.16 with Python 3.6.4) (Martin 2011) and aligned to the mouse reference genome mm10 by RNA STAR (Galaxy version 2.6.0b-2) (Dobin *et al.* 2013). The number of reads was counted with featureCounts (Galaxy version 1.6.4) (Liao *et al.* 2014) using the genome annotation GRCm38 (release 97). The differential expression profile was analyzed in R with DESeq2 (version 1.25.11) (Love *et al.* 2014). Genes with very low expression (average counts <3) were removed and genes with the Benjamini–Hochberg (BH) adjusted *P* value <0.05 were considered sex-differentially expressed. Sequencing data can be found at ArrayExpress database of the EMBL's European Bioinformatics Institute (E-MTAB-8717).

Enrichment and pathway analysis

Genes showing ≥ 2 -fold difference in sex-differential expression were uploaded in DAVID (Bioinformatics Resources, version 6.8; <https://david.ncifcrf.gov/>) (Huang *da et al.* 2009). A default functional annotation (threshold count 2 and EASE 0.1) was performed for Gene Ontology (GO). Subsequently, default functional annotation tools were selected for functional annotation clustering with a medium classification stringency (kappa similarity term overlap 3 and threshold 0.5; multiple linkage threshold 0.5; and EASE 0.1). To identify functional upstream regulators, these differentially expressed genes with their $\log_2$ -transformed fold differences (female relative to male) were uploaded in Ingenuity Pathway Analysis (IPA, version 49309495; QIAGEN Inc., <https://www.qiagenbioinformatics.com/products/ingenuity-pathway-analysis>) (Kramer *et al.* 2014) and analyzed using Ingenuity Knowledge Base

with a default setting (direct and indirect relationships; only experimentally observed confidence).

Isolation of SVF cells, cell culture, and *in vitro* hormonal treatment

BAT depots from five to six mice were pooled (as a biological replicate) and BAT SVF cells were isolated and differentiated for 13 days as described previously (Grefhorst *et al.* 2015). However, a significant number of mice are needed to obtain enough cells for primary brown adipocyte cultures, and this limits the amount of experimental conditions that we could test in primary differentiated adipocytes. To reduce the number of mice, we, therefore, decided to use T37i cells, the female murine brown adipocyte cell line (a gift provided by Dr M. Lombès, Inserm U1185, France) (Zennaro *et al.* 1998), for progesterone stimulations. T37i cells were maintained and differentiated for 9 days as described previously (Kaikaew *et al.* 2019).

Fully differentiated BAT SVF or T37i cells were steroid-starved for 3 h in the starvation medium (medium containing 0.2% dextran-coated charcoal-treated fetal bovine serum (DCC-FBS), prepared as previously described (Kaikaew *et al.* 2019)), followed by pretreatment with 5 or 500 nM of RU486 (Sigma-Aldrich) or EtOH vehicle for 60 min. Subsequently, cells were stimulated with progesterone (Steraloids, Newport, RI, USA) or EtOH vehicle for 24 h by adding the indicated concentrations to the RU486- or EtOH-pretreated cultured media.

To study the effect of progesterone on T37i cell differentiation, cells were treated with the indicated concentrations of progesterone or EtOH vehicle supplemented to steroid-deprived differentiation medium (similar to the normal T37i differentiation medium, except replacing 10% FBS with 9% DCC-FBS and 1% FBS). For NE stimulation of differentiated T37i cells, cells were starved for 3 h in 0.2% DCC-FCS starvation medium and subsequently stimulated for 24 h with 1 μ M NE (Sigma-Aldrich) or HCl vehicle.

Adiponectin and glycerol measurement

Cultured media of progesterone and/or RU486-treated SVF-differentiated brown adipocytes and NE-stimulated T37i cells were collected, centrifuged for 10 min to remove debris, and stored at -20°C until measurement. Adiponectin concentrations were determined using the mouse HMW & total adiponectin ELISA kit (Alpco Diagnostics, Salem, NH, USA) and glycerol concentrations

were determined using the glycerol reagent set (INstruchemie, Delfzijl, Netherlands), according to the manufacturers' protocols.

Quantitative PCR

Reverse transcription and quantitative PCR (qPCR) were performed using the Transcriptor high-fidelity cDNA synthesis kit and the FastStart Universal SYBR Green Master (Rox) (both from Roche Diagnostics) with a QuantStudio 7 flex real-time PCR system (Applied Biosystems, Life Technologies). Samples that yielded too little RNA were

not included in the qPCR analysis. Gene expression was normalized to the indicated housekeeping genes using the $2^{-\Delta\Delta CT}$ method. Primer sequences are listed in Table 1.

Statistical analysis

Data were analyzed in GraphPad Prism (GraphPad Software, Inc.). Differences in gene expression levels, considered at $P < 0.05$, were analyzed with one-way or two-way ANOVA with appropriate *post hoc* tests, indicated in each figure legend. Unless otherwise indicated, data are presented as mean $\pm$ S.E.M.

Table 1 Primer sequences.

Gene	Forward (5' → 3')	Reverse (5' → 3')
<i>Adipoq</i>	GCACTGGCAAGTTCTACTGCAA	GTAGGTGAAGAGAACGGCCTTGT
<i>Ar</i>	TCCAAGACCTATCGAGGAGCG	GTGGGCTTGAGGAGAACCAT
<i>Bax</i>	TGAAGACAGGGGCCTTTTG	AATTCGCCGGAGACACTCG
<i>Bcl2</i>	GGACTTGAAGTGCCATTGGT	CATCACGATCTCCCGTTAT
<i>Bmp8b</i>	CAACCACGCCACTATGCAG	CACTCAGCTCAGTAGGCACA
<i>C7</i>	CAACTGCAAGTGGGACTCCTA	CAGCAACTGAACGCCTTCG
<i>Cdh1</i>	AGACTTTGGTGTGGGTCAGG	ATCTGTGGCGATGATGAGAG
<i>Cfd</i>	CTACAAGCGATGGTATGATGTGC	GGACCCAACGAGGCATTCT
<i>Cldn3</i>	AGTGCTTTTCCTGTTGGCGGCTCT	ATCGCGGCGCAGAATAGAGGATCT
<i>Cldn4</i>	CAGTGCAAGATGTACGACTCGAT	TACCACTGAGAGAAGCATCCCC
<i>Cldn7</i>	AGGGTCTGCTCTGGTCCT	GTACGCAGCTTTGCTTTCA
<i>Cxcl13</i>	CATAGATCGGATTCAAGTTACGCC	TCTTGGTCCAGATCACAACTTCA
<i>Epcam</i>	GCGGCTCAGAGAGACTGTG	CCAAGCATTTAGACGCCAGTTT
<i>Esr1</i>	TGGGTCATGAGAGTCCTTTGAA	CCGGGATGGAAACTGAACCTT
<i>Fabp4</i>	GCGAGTAGAATGACAGCTCCTT	CTGTCGTCTGCGGTGATT
<i>Fkbp5</i>	ATTTGATTGCCGAGATGTG	TCTTACCAGGGCTTTGTC
<i>Irf4</i>	TGCAAGCTCTTTGACACACA	CAAAGCACAGAGTCACCTGG
<i>Krt5</i>	TCTGCCATCACCCCATCTGT	CCTCCGCCAGAACTGTAGGA
<i>Krt8</i>	CAAGGTGGAAGTAGAGTCCCG	CTCGTACTGGGCACGAACTTC
<i>Krt14</i>	CCACCTTTCATCTTCCCAATTCTC	GTGCGGATCTGGCGGTTG
<i>Krt18</i>	CTGGAGGATGGAGAAGATTT	CTTTTATTGGTCCCTCAGTT
<i>Lcn2</i>	ACTTCCGGAGCGATCAGTT	CAGCTCCTTGTTCTTCCAT
<i>Lep</i>	ACCCCATCTGAGTTTGTC	TCCAGGTCATTGGCTATCTG
<i>Mup^a</i>	CAAAACAGAAAAGGCTGGTGA	TTGTGCAACCTTTCTTTGA
<i>Nr3c1</i>	CCGGGTCCCCAGGTAAAGA	TGTCCGGTAAAATAAGAGGCTTG
<i>Nr3c2</i>	GAAGAGCCCCCTCTGTTTGAG	TCCTTGAGTGATGGGACTGTG
<i>Pgr</i>	GGGGTGGAGGTCGTACAAG	GCGAGTAGAATGACAGCTCCTT
<i>Pparg</i>	GAAAGACAACGGACAATACACC	GGGGGTGATATGTTGAACCTG
<i>Ppargc1a</i>	CCCTGCCATTGTTAAGACC	TGCTGCTGTTCTGTTTTT
<i>Ptprf</i>	TGCTCTCGTGATGCTTGGTTT	ATCCACGTAATTCGAGGCTTG
<i>Scnn1b</i>	ACCCGGTGGTTCTCAATTTGT	AAGTTCGCAAGGTACACACA
<i>Sdc1</i>	TGGAGAACAAGACTTCACCTTTG	CTCCAGCACTTCCTTCTT
<i>St14</i>	TCATCGCCTACTACTGGTCAGAGT	TGGCGCGATCAACCTCTT
<i>Tsc22d3</i>	CAGCAGCCACTCAAACCAGC	ACCACATCCCCTCCAAGCAG
<i>Ucp1</i>	GGCCTCTACGACTCAGTCCA	TAAGCCGGCTGAGATCTTGT
<i>Wfdc2</i>	AACCAATTACGACTGTGTGT	TCGCTCGGTCCATTAGGCT
<i>Actb[*]</i>	AAGGCCAACCGTGAAAAGAT	GTGGTACGACCAGAGGCATAC
<i>B2m[*]</i>	ATCCAAATGCTGAAGAACGG	CAGTCTCAGTGGGGGTGAAT
<i>Hprt[*]</i>	GCAGTACAGCCCCAAAATGG	AACAAAGTCTGGCCTGTATCCAA
<i>Rn18s[*]</i>	GTAACCCGTTGAACCCCAT	CCATCCAATCGGTAGTAGCG

^aThe primers detect multiple genes of the MUP family due to its high similarity in mRNA sequences. ^{*}Housekeeping genes.

Results

BAT transcriptional profile of male and female mice

RNA-seq analysis identified 17,798 transcripts in the interscapular BAT of mice. Expression levels of these identified transcripts are visualized by the relative female-to-male fold differences in Fig. 1A. A total of 2,038 genes were identified being sex-differentially expressed (unadjusted $P < 0.05$), but when accounting for a BH correction at $P < 0.05$ only 596 transcripts remained, of which 347 genes (58.2%) showed a female-biased expression and 249 genes (41.8%) showed a male-biased expression (Fig. 1A and B). Further trimming by applying a cut-off of two-fold difference resulted in 295 significantly sex-differentially expressed genes, of which 242 genes (82.0%) showed higher expression and 53 genes (18.0%) showed lower expression in female BAT than in male BAT (Fig. 1B). Hierarchical clustering analysis of the 295 genes illustrates the differences in the transcriptional profile of BAT between male and female mice (Fig. 1C).

Analysis of biological processes of the sex-differentially expressed genes

Of the 295 sex-differentially expressed genes, 291 genes were matched in the DAVID mouse data resources.

Top GO annotation terms from enrichment analyses are presented in Table 2 and top significant functional annotation clusters are presented in Table 3. In general, among the leading significant categories and clusters are genes that encode proteins involved in cellular structure and cell-cell contact, interaction, and adhesion.

Validation of the sex-differentially expressed genes by qPCR and influences of GDX

We selected a subset of genes with a high sex-differential expression level, corresponding to the first five significant functional annotation clusters. The summary of their known biological function is reported in Supplementary Table 1 (see section [supplementary materials](#) given at the end of the article). Since sex steroids have been implicated to contribute to sex differences in BAT function (Gonzalez-Garcia *et al.* 2017), expression of these selected genes was validated in BAT of both sham-operated and gonadectomized mice by qPCR (Fig. 2). This analysis confirmed that sham-operated female mice had higher BAT mRNA expression of cell adhesion and structural molecules, for example, epithelial cadherin (E-cadherin), claudins, epithelial cell adhesion molecule (EpCAM),

Figure 1

Sex difference in BAT RNA-seq profile. (A) Scatter plot demonstrating expression levels of all identified genes. Log-transformed average normalized counts of genes in BAT are presented on the X axis and log-transformed fold changes are presented on the Y axis, with positive values indicating female-biased expression and negative values indicating male-biased expression. Blue and pink dots represent known differentially expressed genes and serve as internal controls: blue indicates Y-chromosomal genes, whereas pink indicates X gametologs of the Y-chromosomal genes and the major X inactivation gene, *Xist*. Green dots indicate the significant sex-differentially expressed genes (BH-adjusted $P < 0.05$). (B) Pie and bar chart demonstrating the number and percentage of identified genes. A two-fold cut-off was applied to select genes with a substantial sex-dependent expression. (C) Heat map and dendrogram presenting the expression of the 295 sex-differentially expressed genes (BH-adjusted $P < 0.05$ and differential expression ≥ 2 folds) in BAT of four male (M1–M4) and four female (F1–F4) mice. The heat map was visualized with the TreeScape (OmniViz, version 6.1.13.0). Red indicates upregulation and blue indicates downregulation, compared to the geometric mean of the *rlog*-normalized expression levels of each gene, and the color intensity represents the magnitude of the difference.

Table 2 Top GO terms in each annotation category by enrichment analysis in DAVID.

GO term	Count	%	P value
Biological process			
Cell adhesion	23	7.90	7.00E-7
Mammary gland alveolus development	5	1.72	1.05E-4
Cellular response to platelet-derived growth factor stimulus	5	1.72	2.30E-4
Sodium ion transport	9	3.09	2.73E-4
Positive regulation of fat cell differentiation	6	2.06	4.95E-4
Canonical Wnt signaling pathway	7	2.41	0.001
Cellular component			
Extracellular exosome	87	29.90	2.89E-16
Extracellular region	60	20.62	8.36E-12
Proteinaceous extracellular matrix	23	7.90	2.02E-10
Extracellular space	48	16.49	1.79E-8
Apicolateral plasma membrane	7	2.41	2.17E-7
Cell junction	29	9.97	2.68E-7
Molecular function			
Structural molecule activity	16	5.50	5.19E-7
Symporter activity	10	3.44	1.63E-5
Cell adhesion molecule binding	8	2.75	3.88E-5
Protein binding	78	26.80	2.43E-4
Scaffold protein binding	6	2.06	6.28E-4
Transcription activator activity, RNA polymerase II core promoter proximal region sequence-specific binding	12	4.12	8.79E-4

Count indicates the number of sex-differentially regulated genes in the GO term and % was calculated based on the total of 291 sex-differentially expressed genes uploaded in DAVID. P value indicates a modified Fisher exact test, the DAVID enrichment (EASE score) analysis.

keratins, and syndecan-1 than sham-operated male mice (Fig. 2). Ovariectomy significantly reduced the expression of this selected subset of genes, but orchiectomy only had marginal effects. Hence, female gonadal factors are likely important regulators of the expression of cell adhesion and structural molecules in interscapular BAT of mice.

Analysis of possible upstream regulators and its regulated genes

Upstream regulator analysis in IPA was used to identify possible regulators of the sex-differentially expressed genes. Top significant upstream regulators are presented in Table 4. In the gene, RNA, and protein category, erb-b2 receptor tyrosine kinase 2 (ERBB2, also known as human EGF receptor 2 (HER2)) and transforming growth factor beta-1 (TGFβ1) were identified as the top upstream regulators. The predicted state for ERBB2 on sex-differential (female-over-male) BAT transcriptome was an activation ($Z = 3.58$). This was also predicted for TGFβ1 albeit with lower confidence ($Z = 1.48$). Interestingly, another top identified upstream regulator was the estrogen receptor which is in line with E2 being the top upstream regulator in the endogenous chemical category (Table 4). The activation state for E2 on sex-differential (female-over-male) BAT transcriptome was predicted as an activation ($Z = 3.88$). Another female sex steroid, progesterone,

was predicted as the second upstream regulator in the endogenous chemical category and its activation state was possibly an activation ($Z = 1.35$). Dihydrotestosterone (DHT) was also identified as an upstream regulator, but its activation state was uncertain ($Z < 1$).

The identification of E2 and estrogen receptor as upstream regulators is in line with previous studies implicating a role for E2 in the activation of BAT (Gonzalez-Garcia *et al.* 2017). However, the involvement of progesterone in BAT function is less clear. Therefore, we further investigated the effects of progesterone on BAT.

Effects of progesterone on sex-differentially expressed genes in SVF-differentiated brown adipocytes

Direct effects of progesterone on the transcription of the validated sex-differentially expressed genes reported in Fig. 2 were assessed in SVF-differentiated primary brown adipocytes. Ingenuity identified a total of 25 genes as progesterone-regulated molecules in our dataset, of which 9 genes (*Cdh1*, *Cfd*, *Cldn3*, *Cldn4*, *Krt5*, *Lcn2*, *Pgr*, *Scnn1b*, and *Wfdc2*) were among the qPCR-validated genes in BAT. We first determined whether these 9 IPA-identified progesterone-regulated genes were detectable in SVF-differentiated brown adipocytes by qPCR. We could only detect mRNA expression of *Cfd*, *Lcn2*, and *Pgr* in

Table 3 Top functional annotation clusters by DAVID, presenting only GO terms.

Cluster 1	Enrichment score 10.351
GO terms	Extracellular region; extracellular space
Genes	Acta2, Actg2, Adamts15, Adamts18, Alcam, Ano9, Anxa2, Apln, Aqp5, Areg, Atp4b, Bglap3, Bmp8b, Bnc2, Bsn, Btn1a1, C7, Ccdc3, Ccr3, Cdcpl1, Cdh1, Cdh3, Cdh11, Cel, Cfd, Chil1, Clca3a2, Col4a6, Col9a2, Col12a1, Col17a1, Csn3, Ctse, Cxcl13, Cxcl15, Dkk1, Dmbt1, Dmkn, Dsg2, Epcam, Ephb3, Ezr, Faim2, Fam20c, Fbn2, Fcgbp, Fetub, Fgb, Flrt1, Frem1, Fxyd3, Galnt3, Gcnt4, Ggt1, Gldn, Glt1d1, Gpr156, Gpx3, Hc, Igfals, Il17b, Itgb3, Itih2, Krt8, Krt14, Krt18, Krt2ap, Lad1, Lama1, Lamc2, Large2, Lcn2, Lgals7, Ltf, Lvrn, Ly6d, Mfap2, Mfap4, Mfsd2a, Muc15, Mup1 ^a , Myc, Ncan, Nectin4, Nipal2, Npr3, Nt5e, Olfm2, Oxtr, Pdgfd, Phyhip, Piezo2, Plb1, Plet1, Plod2, Plppr3, Prlr, Proc, Prom1, Prom2, Prtn3, Ptk7, Ptn, Ptprf, Ptprn2, R3hdm1, S100a9, Scg3, Scnn1b, Sdc1, Serpinb5, Slc1a1, Slc5a1, Slc5a5, Slc5a7, Slc5a8, Slc5a9, Slc6a13, Slc12a2, Slc38a1, Slc44a4, Slco2a1, Slit2, Spint2, Spon1, St14, Sucnr1, Tacstd2, Thsd4, Tlr5, Tmem119, Tmprss2, Tmprss13, Tpbg, Tspan1, Upk3a, Vtcn1, Wfdc2, Wfdc18, Wisp2, Wnt7b
Cluster 2	Enrichment score 3.902
GO terms	Membrane; plasma membrane
Genes	Abcc8, Alcam, Ano9, Anxa2, Ap1m2, Aqp5, Areg, Atp4b, Atp6v1b1, Basp1, Bnc2, Btn1a1, Cacna1g, Cadps2, Ccr3, Cdcpl1, Cdh1, Cdh3, Cdh11, Cldn3, Cldn4, Cldn7, Cldn8, Clic6, Col17a1, Ddx3y, Dmbt1, Dsg2, Dsp, Epcam, Ephb3, Eyt3, Ezr, Faim2, Fer114, Fermt1, Flrt1, Fxyd3, Galnt3, Gcnt4, Ggt1, Gldn, Gpr156, Hpd, Irf4, Irs1, Itgb3, Kcng4, Krt5, Krt19, Lamc2, Large2, Lvrn, Ly6d, Marveld3, Mfsd2a, Mmd2, Mrgprg, Mtmr7, Muc15, Ncan, Nectin4, Nipal2, Npr3, Nt5e, Olfm2, Oxtr, Pdgfd, Perp, Pgr, Piezo2, Pkp1, Plb1, Plch2, Plet1, Plod2, Plppr3, Pmepa1, Prlr, Prom1, Prom2, Prtn3, Ptk7, Ptn, Ptprf, Ptprn2, S100a14, S100a9, Scg3, Scnn1b, Sdc1, Slc1a1, Slc5a1, Slc5a5, Slc5a7, Slc5a8, Slc5a9, Slc6a13, Slc12a2, Slc35f3, Slc38a1, Slc44a4, Slco2a1, Slit2, Smim3, Spint2, St14, Sucnr1, Syt14, Tacstd2, Tlr5, Tmem45b, Tmem56, Tmem119, Tmprss2, Tmprss13, Tpbg, Tspan1, Tusc5, Upk3a, Vtcn1, Wisp2, Wnt7b, Wwc1, 6430571L13Rik
Cluster 3	Enrichment score 3.542
GO terms	Structural molecule activity; cell periphery; scaffold protein binding; intermediate filament
Genes	Actg2, Cacna1g, Cdh1, Cgn, Cldn3, Cldn4, Cldn7, Cldn8, Col4a6, Dsp, Ezr, Fgb, Krt5, Krt8, Krt14, Krt15, Krt17, Krt18, Krt19, Lad1, Sprr1a
Cluster 4	Enrichment score 3.295
GO terms	Symporter activity; sodium ion transport; transporter activity; ion transport
Genes	Aqp5, Atp4b, Cacna1g, Clic6, Fxyd3, Kcng4, Lcn2, Ltf, Mfsd2a, Mup1 ^a , Piezo2, Scnn1b, Slc1a1, Slc5a1, Slc5a5, Slc5a7, Slc5a8, Slc5a9, Slc6a13, Slc12a2, Slc38a1, Slco2a1
Cluster 5	Enrichment score 3.022
GO terms	Apicolateral plasma membrane; bicellular tight junction; calcium-independent cell–cell adhesion via plasma membrane cell–adhesion molecules
Genes	Alcam, Cdh1, Cdh3, Cgn, Cldn3, Cldn4, Cldn7, Cldn8, Epcam, Ezr, Krt8, Krt19, Mapk13, Marveld3, Perp, Pof1b, Ptprf, Rasgrf1, Sdc1, Vtcn1

^aMup1 was entered in DAVID as a representative for Mup22 due to its highly similar protein sequences (<https://www.uniprot.org/uniprot/Q4FZE8>).

differentiated SVFs, of which levels at basal conditions confirmed the higher expression levels in female than in male BAT, although only *Cfd* and *Pgr* expression reached statistical significance (Fig. 3A). Progesterone stimulation for 24 h revealed that progesterone dose-dependently reduced *Cfd* mRNA expression in differentiated SVFs of both sexes (Fig. 3B), while it induced *Lcn2* (Fig. 3C) but reduced *Pgr* (Fig. 3D) mRNA expression more prominently in female adipocytes than in male adipocytes.

Apart from *Pgr*, which encodes the nuclear progesterone receptor (PR) and is a known progesterone-regulated gene, we also investigated the mRNA expression of other nuclear receptors to determine whether they may underlie the progesterone effects. Interestingly, progesterone dose-dependently reduced *Nr3c1* (encoding the glucocorticoid receptor (GR)) mRNA expression in adipocytes of both sexes (Fig. 3E), but progesterone did not significantly affect mRNA expression of the mineralocorticoid receptor and the androgen receptor (data not shown). Moreover,

mRNA expression of a GR-target gene *Tsc22d3* (also known as glucocorticoid-induced leucine zipper (*Gilz*)) was dose-dependently upregulated by progesterone treatment of differentiated SVFs (Fig. 3F), implicating the involvement of GR signaling in the progesterone effects in these SVFs.

As RU486 is a PR- and GR-antagonist with a more PR-specific competitive antagonism at low concentrations (i.e. 5 nM) and a dual PR- and GR-antagonism at high concentrations (i.e. 500 nM) (Kroon *et al.* 2018), we co-administered progesterone with 5 or 500 nM RU486 to study the contribution of PR and GR in the observed progesterone effects. Noticeably, the progesterone-induced upregulation of *Tsc22d3* mRNA expression remained unaffected upon 5 nM RU486 co-administration but was blocked in the presence of 500 nM RU486 (Fig. 3F), confirming the specificity of the selected RU486 doses. Similarly, we observed that progesterone-regulated *Cfd*, *Lcn2*, and *Pgr* mRNA expression levels were marginally affected by 5 nM RU486, but were completely blunted

Genes	RNA-seq F/M ratio, mean (95% CI)	qPCR relative expression level, mean±SEM				Sig.	Relative expression
		Male sham	Female sham	Male GDX	Female GDX		
<i>Epcam</i>	520.3 (88.4–3060.7)	1.00±0.37	698.2±247.9 [‡]	0.27±0.07	19.00±11.01 [‡] #	×,S,G	2 ⁹
<i>Pgr</i> [†]	439.9 (43.7–4433.0)	1.00±0.49	1.80±0.38	0.42±0.25	0.15±0.08 [#]	G	2 ⁶
<i>Wfdc2</i> [†]	343.6 (69.3–1703.1)	1.00±0.18	155.8±15.9 [‡]	0.96±0.08	6.91±2.03 [‡] #	×,S,G	2 ³
<i>Krt8</i>	332.5 (68.3–1619.0)	1.00±0.22	2616.0±588.2 [‡]	0.94±0.12	63.07±46.06 [‡] #	×,S,G	2 ⁰
<i>Cldn4</i> [†]	315.0 (49.8–1992.2)	1.00±0.07	1486.6±387.7 [‡]	1.22±0.10	56.81±7.47 [‡] #	×,S,G	2 ^{−3}
<i>Krt14</i>	243.6 (45.4–1308.0)	1.00±0.41	172.8±48.8 [‡]	0.39±0.06	5.55±3.08 ^(S) #	×,S,G	2 ^{−6}
<i>Krt5</i> [†]	197.8 (38.5–1014.7)	1.00±0.19	6.35±1.26 [‡]	1.04±0.26	1.26±0.33 [‡]	×,S,G	2 ^{−9}
<i>Krt18</i>	129.5 (37.0–453.2)	1.00±0.28	788.6±143.9 [‡]	0.34±0.08 ^(#)	20.91±7.82 [‡] #	×,S,G	
<i>Mup22</i>	38.31 (7.08–207.4)	1.00±0.20	17.17±6.73 [‡]	0.92±0.29	1.46±0.65 [‡]	×,S,G	
<i>Scnn1b</i> [†]	25.90 (4.44–151.2)	1.00±0.19	34.53±12.61 [‡]	0.60±0.08	1.99±0.42 [‡]	S,G	
<i>Cldn3</i> [†]	19.18 (7.50–49.06)	1.00±0.21	103.6±33.1 [‡]	1.59±0.49	6.26±1.29 [‡] #	×,S,G	
<i>Bmp8b</i>	18.80 (9.08–38.92)	1.00±0.57	28.46±12.26 [‡]	0.29±0.06	1.71±0.19 [‡] #	(×),S,G	
<i>Cdh1</i> [†]	17.19 (7.98–37.02)	1.00±0.04	26.86±7.88 [‡]	0.69±0.23	2.12±0.45 [‡] #	×,S,G	
<i>Lcn2</i> [†]	10.33 (5.75–18.56)	1.00±0.62	4.98±2.14 [‡]	0.77±0.39	0.32±0.13 [‡]	(×),(G)	
<i>St14</i>	8.20 (3.88–17.35)	1.00±0.28	11.90±3.05 [‡]	0.67±0.19	2.84±0.65 [‡] #	S,G	
<i>Cldn7</i>	7.33 (2.69–19.97)	1.00±0.18	25.20±6.40 [‡]	0.60±0.11	2.95±0.84 [‡] #	×,S,G	
<i>Irf4</i>	3.08 (2.15–4.41)	1.00±0.29	3.23±0.41 [‡]	2.34±0.56 ^(#)	1.11±0.30 [‡]	×	
<i>Ptprf</i>	2.53 (1.63–3.94)	1.00±0.15	1.97±0.50	0.79±0.12	0.64±0.06 [‡]	G	
<i>Cfd</i> [†]	2.41 (1.74–3.33)	1.00±0.30	5.17±1.02 [‡]	1.78±0.25	1.05±0.33 [‡]	×,(S)	
<i>Sdc1</i>	2.36 (1.70–3.26)	1.00±0.11	1.77±0.31 [‡]	0.96±0.09	0.75±0.14 [‡]	×,G	
<i>Cxcl13</i>	0.14 (0.08–0.25)	1.00±0.29	0.39±0.09	2.95±1.32	0.22±0.10 [‡]	(×),S	
<i>C7</i>	0.07 (0.06–0.10)	1.00±0.31	0.08±0.00 [‡]	1.12±0.46	0.12±0.03 [‡]	S	

Figure 2

Validation of gene expression by qPCR and influences of sex and GDX on sex-differential expression in BAT. The relative female/male (F/M) expression ratios by RNA-seq (4 mice per group) are shown in descending order in the left column. DESeq2-estimated log₂-transformed fold changes (lfc) and the standard error of the lfc (lfcSE) were used to calculate the F/M ratio by 2^{lfc} and the 95% CI by 2^{lfc−1.96(lfcSE)} and 2^{lfc+1.96(lfcSE)}. † indicates genes that the Ingenuity database identified as progesterone-regulated molecules. The other columns present relative expression levels measured by qPCR (four to six mice per group). Expression levels were normalized to *B2m* and *Rn18s* and are shown relative to the sham-operated male mice. The Sig. column lists significant effects (*P* < 0.05) analyzed from the log-transformed expression levels by two-way ANOVA: G for gonadal status (sham or GDX), S for sex, and × for an interaction between G and S. ‡ indicates a significant sex difference between mice with the same gonadal status and # indicates a significant effect of gonadal status within the same sex, by *post hoc* test (*P* < 0.05). Symbol in parentheses indicates a tendency to significance (*P* < 0.10). Color demonstrates log₂-transformed relative expression levels. GDX, gonadectomy.

with 500 nM of RU486 cotreatment (Fig. 3B, C and D), suggesting GR-mediated responses of progesterone on these genes.

Next, we determined whether progesterone could affect the expression of the 13 qPCR-validated sex-differentially expressed genes not present in the IPA database of progesterone-regulated molecules (Fig. 2). By qPCR, only the mRNA expression of *C7*, *Ptprf*, *Scd1*,

and *Krt14* was detectable in SVF-differentiated adipocytes (Fig. 3G). Unlike the expression pattern observed in interscapular BAT, we found that mRNA expression levels of *C7* were higher in female than in male SVF-differentiated adipocytes and progesterone treatment stimulated *C7* mRNA expression in differentiated adipocytes of both sexes (Fig. 3H). This stimulatory effect was abolished by 500 nM but marginally affected by 5 nM

Table 4 Upstream regulator analysis of sex-differential expression in BAT by IPA.

Upstream regulator	Molecule type	Activation Z score	P value of overlap
Genes, RNAs, and proteins			
ERBB2	Kinase	3.58	1.16E−15
Estrogen receptor	Group	1.43	2.43E−15
TGFB1	Growth factor	1.48	7.14E−13
CAV1	Transmembrane receptor	–	1.53E−12
KLF4	Transcription regulator	3.55	1.51E−11
CEBPA	Transcription regulator	0.22	5.91E−11
TNF	Cytokine	1.15	4.68E−10
CCN5	Growth factor	1.27	6.00E−10
SOX2	Transcription regulator	0.58	9.36E−10
EGF	Growth factor	1.23	1.07E−09
Endogenous chemicals			
17β-estradiol	Endogenous chemical	3.88	2.45E−11
Progesterone	Endogenous chemical	1.35	1.78E−7
Tretinoin	Endogenous chemical	−0.10	3.02E−7
Dihydrotestosterone	Endogenous chemical	0.53	6.20E−5

Activation Z score identifies upstream regulators that can explain observed gene expression fold changes in the dataset and predicts the activation state of the possible upstream regulators based on the Ingenuity Knowledge Base. *Z* > 2 indicates a significant prediction of an activated state whereas *Z* < −2 indicates a significant prediction of an inhibited state. *P* value of overlap indicates enrichment of the database-known regulated genes in the dataset.

Figure 3

Effects of progesterone on the RNA-seq-identified sex-differentially expressed genes in SVF-differentiated brown adipocytes. SVF-derived primary cultures of brown adipocytes from male and female BAT were stimulated with progesterone at indicated concentrations: P-7 (10^{-7} M), P-6 (10^{-6} M), or P-5 (10^{-5} M), together with RU486 (5 or 500 nM) or EtOH vehicle. Expression levels were normalized to *Actb* and *Hprt*, depicted relative to the level of EtOH-treated male adipocytes, and presented as mean $\pm$ s.e.m. (four biological replicates per condition). Data were log-transformed before statistical analyses. Basal mRNA expression levels of (A) the RNA-seq-identified sex-differentially expressed and IPA-identified progesterone-regulated genes and (G) the remaining RNA-seq-identified sex-differentially expressed genes. \$ indicates a significant sex difference ($P < 0.05$) or (*) indicates a tendency to differ ($P < 0.10$) by unpaired *t* tests. (B, C, D, E, F and H, I, J) mRNA expression levels of the indicated genes. Two-way repeated measures ANOVA was performed to identify the effects of sex and progesterone in each RU486 condition. Significant effects are presented as follows: (S), $P_{\text{sex}} < 0.10$; (P), $P_{\text{sex}} < 0.05$; SS, $P_{\text{sex}} < 0.01$; (PP), $P_{\text{progesterone}} < 0.10$; PP, $P_{\text{progesterone}} < 0.01$; PPP, $P_{\text{progesterone}} < 0.001$; x, $P_{\text{sex} \times \text{progesterone}} < 0.05$; and xx, $P_{\text{sex} \times \text{progesterone}} < 0.01$. Dunnett's *post hoc* tests were applied when appropriate. * indicates a significant difference ($P < 0.05$) and (*) indicates a tendency to differ ($P < 0.10$) from the EtOH control.

RU486 cotreatment. Progesterone marginally inhibited *Ptprf* mRNA expression only in male-derived differentiated adipocytes (Fig. 3I) but did not affect *Sdc1* (Fig. 3J) and *Krt14* (data not shown) mRNA expression in cells of either sex. Of note, the finding that mRNA expression of only 7 out of 22 qPCR-validated sex-differentially expressed genes in interscapular BAT were detectable in primary brown adipocytes (Fig. 3A and G) may suggest that these other 15 genes are expressed by other cell types in the BAT depot.

Effects of progesterone on adipocyte markers in SVF-differentiated brown adipocytes

Since our results suggest that progesterone may regulate some of the sex-differentially expressed genes in BAT, we next determined whether progesterone affects adipocyte function

by analyzing the expression of common adipocyte markers in differentiated SVFs. At basal conditions, mRNA expression of *Ucp1* and its transcription activator *Ppargc1a* appeared to be higher in female than in male SVFs differentiated into adipocytes, but both failed to reach significance ($P > 0.10$; Fig. 4A). Female differentiated SVFs had higher expression levels of adipogenic differentiation and maturation genes *Pparg*, *Fabp4*, and *Adipoq* than male differentiated SVFs (Fig. 4A). In addition, female differentiated SVFs had a lower expression of the classical white adipocyte marker *Lep* than male differentiated SVFs, suggesting a more metabolically active phenotype of female differentiated brown adipocytes (Fig. 4A), although it should be noted that *Lep* mRNA expression was relatively low in these cells (C_T values of samples were only $\sim 1.36 \pm 0.01$ lower than negative controls). Progesterone treatment did not significantly affect expression levels of *Ucp1* (Fig. 4B), *Ppargc1a* (Fig. 4C), *Pparg*

Figure 4

Effects of progesterone on adipocyte markers in SVF-differentiated brown adipocytes. SVF-derived primary cultures of brown adipocytes from male and female BAT were stimulated with progesterone at indicated concentrations: P-7 (10^{-7} M), P-6 (10^{-6} M), or P-5 (10^{-5} M), together with RU486 (5 or 500 nM) or EtOH vehicle. Data are presented as mean $\pm$ s.e.m. (four biological replicates per condition). (A) Basal mRNA expression levels of common adipocyte markers. *indicates a significant sex difference ($P < 0.05$) or (\$) indicates a tendency to differ ($P < 0.10$) by unpaired *t* tests. (B, C, D, E and F) mRNA expression levels of the indicated genes. mRNA expression levels were normalized to *Actb* and *Hprt*, depicted relative to the level of EtOH-treated male adipocytes, and log-transformed before statistical analyses. (G) Adiponectin (protein) concentrations in cultured media of the cells with progesterone/RU486 stimulation. Two-way repeated measures ANOVA was performed to identify the effects of sex and progesterone in each RU486 condition. Significant effects are presented as follows: (S), $P_{\text{sex}} < 0.10$; S, $P_{\text{sex}} < 0.05$; SS, $P_{\text{sex}} < 0.01$; (P), $P_{\text{progesterone}} < 0.10$; P, $P_{\text{progesterone}} < 0.05$; PPP, $P_{\text{progesterone}} < 0.001$; (X), $P_{\text{sex} \times \text{progesterone}} < 0.10$; and X, $P_{\text{sex} \times \text{progesterone}} < 0.05$. Dunnett's *post hoc* tests were applied when appropriate. *indicates a significant difference ($P < 0.05$) and (*) indicates a tendency to differ ($P < 0.10$) from the EtOH control.

(Fig. 4D), and *Fabp4* (Fig. 4E). Intriguingly, progesterone dose-dependently reduced *Adipoq* mRNA expression in differentiated SVFs of both sexes (Fig. 4F), of which the inhibitory effect of progesterone was also likely driven by GR since the effect was marginally affected by 5 nM RU486 but was blunted by 500 nM RU486 co-treatment. In addition, these results of *Adipoq* mRNA expression were confirmed at the protein level, through measurement of adiponectin levels in cultured media of the SVF-differentiated brown adipocytes (Fig. 4G).

Effects of progesterone in a brown adipocyte cell line

Next, we analyzed the effects of progesterone on adipocyte differentiation. To limit the number of mice, we first analyzed whether the effects of progesterone on

the expression of common brown adipocyte markers were also present in T37i, a female brown adipocyte cell line (Zennaro *et al.* 1998). Differentiated T37i were stimulated with progesterone for 24 h. High-dose progesterone significantly reduced *Ucp1* mRNA expression (Fig. 5A) but did not significantly affect *Pparg1a* mRNA expression (Fig. 5B). In contrast, *Pparg* mRNA expression was induced by high doses of progesterone (Fig. 5C). Interestingly, *Fabp4* mRNA expression was induced by 10^{-6} and $10^{-5.5}$ M progesterone but reduced by 10^{-5} M progesterone (Fig. 5D). Expression levels of both adipokines *Adipoq* and *Lep* were reduced by high-dose progesterone stimulation (Fig. 5E and F).

Concerning the expression of receptors that can potentially mediate progesterone effects, we found that *Pgr* mRNA expression was not detectable in T37i cells

Figure 5

Effect of progesterone on differentiated T37i brown adipocytes. Differentiated T37i brown adipocytes were treated for 24 h with progesterone at indicated concentrations. Expression levels were normalized to *Actb* and *B2m*, depicted relative to the EtOH vehicle-treated condition, and presented as mean $\pm$ S.E.M. (three biological replicates per condition). Data were log-transformed before statistical analyses with one-way ANOVA. * indicates a significant difference ($P < 0.05$) and (*) indicates a tendency to differ from the EtOH-treated condition ($P < 0.10$) by Dunnett's *post hoc* tests.

(data not shown) while *Nr3c1* mRNA expression was downregulated by progesterone (Fig. 5G). Transcription levels of the GR-target genes *Tsc22d3* (Fig. 5H) and *Fkbp5* (data not shown) were dose-dependently upregulated.

Since progesterone affected the expression of *Pparg* (the master transcriptional activator of adipogenesis) and *Ucp1* in differentiated T37i cells, we next determined the impact of progesterone on T37i differentiation by adding progesterone to the differentiation cocktail. *Ucp1* and *Ppargc1a* (the transcriptional activator of mitochondrial biogenesis) mRNA expression levels were strongly upregulated during differentiation ($P_{\text{day}} < 0.001$) but were significantly inhibited in the presence of high-dose progesterone (Fig. 6A and B). In contrast, *Pparg* mRNA expression was induced by co-stimulation with 10⁻⁶ M progesterone (Fig. 6C), while expression of the mature adipocyte marker *Fabp4* in the mature adipocytes (day 9) was not significantly affected by progesterone (Fig. 6D). However, at the early differentiation phase (day 3), 10⁻⁷ M and 10⁻⁶ M progesterone-induced *Adipoq* mRNA expression, but this effect was absent at later stages when even a downregulation was seen at 10⁻⁵ M progesterone after 9 days of differentiation (Fig. 6E). *Lep* mRNA expression was inhibited at the highest dose of progesterone treatment throughout differentiation (Fig. 6F). Of note, mRNA expression levels of a pro-apoptotic marker *Bax* and an anti-apoptotic marker *Bcl2* were not

significantly affected by progesterone co-treatment (data not shown), suggesting that progesterone did not hamper cell survival. Subsequently, we tested the NE-induced thermogenic activity in differentiated T37i cells with/without progesterone treatment during differentiation. Co-treatment with high-dose progesterone during differentiation significantly inhibited baseline mRNA expression levels of *Ucp1* (Fig. 6G) and *Ppargc1a* (Fig. 6H). However, progesterone-treated cells remained responsive to NE since NE stimulation induced the expression of these thermogenic genes in T37i cells, although NE-induced expression levels were reduced in cells treated with high-dose progesterone (Fig. 6G and H). Furthermore, NE-induced glycerol release from T37i cells (a proxy for thermogenic activity) was dose-dependently reduced by progesterone treatment during adipocyte differentiation (Fig. 6I).

Discussion

To the best of our knowledge, this is the first study to investigate sex differences in the global transcriptional profile by RNA-seq in the interscapular BAT of mice at reproductive age and at basal housing conditions at ~22°C. Functional annotation and clustering showed enrichment of genes encoding proteins involved in cellular structure, cell-cell contact, and cell adhesion. Although female BAT

Figure 6

Effect of progesterone treatment during differentiation of T37i brown adipocytes. Progesterone (P) at indicated concentrations were treated during differentiation of T37i brown preadipocytes until full differentiation at day 9. Data are presented as mean $\pm$ S.E.M. (three biological replicates per condition). Gene expression data were log-transformed before statistical analyses with two-way ANOVA. *indicates a significant difference ($P < 0.05$) and (*) indicates a tendency to differ ($P < 0.10$) from the EtOH-treated condition by Dunnett's *post hoc* tests. (A, B, C, D, E and F) Days after differentiation, when the differentiating cells were collected, are plotted on the X axis. Expression levels were normalized with *Actb* and *B2m* and depicted relative to the levels in undifferentiated cells (day 0). (G, H and I) Nine-day differentiated cells with/without progesterone during differentiation were stimulated for 24 h with 1 μ M of norepinephrine (NE) or vehicle and analyzed for mRNA expression or glycerol release in cultured media. Gene expression levels were normalized with *Actb* and *B2m* and depicted relative to levels of the EtOH-treated/vehicle-stimulated condition. Concentrations of progesterone treatment during differentiation are indicated on the X axis as follows: P-9 (10^{-9} M), P-8 (10^{-8} M), P-7 (10^{-7} M), P-6.5 ($10^{-6.5}$ M), P-6 (10^{-6} M), P-5.5 ($10^{-5.5}$ M), and P-5 (10^{-5} M).

was isolated randomly throughout the estrous cycle, the sex-differential expression remained apparent suggesting a robust sex-dependent regulation of transcription in BAT. In support, we confirmed our previous findings by this approach, showing that *Bmp8b* is sex-differentially expressed in BAT (Grefhorst *et al.* 2015).

Cell adhesion is an essential process in maintaining tissue structure and cellular interaction with neighboring cells or with the extracellular matrix (ECM) (Gumbiner 1996). Structural and adhesion proteins, such as E-cadherin, claudins, and keratins, are well-known for their role in this process as well as in epithelial-to-mesenchymal transition, cancer progression, and tumor metastasis (Basu *et al.* 2018). Keratins were identified exclusively in omental WAT, but not in subcutaneous WAT, and have been suggested to play a role in the development of the inflammation process, but also to contribute to mechanical stress of adipocytes during obesity (Perez-Perez *et al.* 2009). However, the role of structural and

adhesion proteins in BAT is less clear, although the microenvironment is shown to be important for visceral and subcutaneous WAT morphology and function, including the formation of new vessels to promote optimal tissue expansion (Choe *et al.* 2016, Pope *et al.* 2016). Upon HFD feeding of mice, male gonadal WAT showed a lack of angiogenesis whereas female gonadal WAT had a high vascular density, together with an upregulation of the angiogenic factors, for example, *Vegfa* and *Vegfr2* (Rudnicki *et al.* 2018). In support of our findings in BAT, a study comparing obesity-prone and obesity-resistant rats upon HFD feeding showed that genes encoding proteins involved in cell structure and motility constituted almost a quarter of the differentially expressed genes, with the lowest expression in the obesity-prone rats (Joo & Yun 2011). However, no major overlap was observed between the genes detected in that study with our identified genes. Importantly, we observed that the expression of most identified structural and cell-adhesion genes was very low

in our primary brown adipocyte cultures, despite having detectable expression in BAT itself. This might suggest that *in vivo* these genes are expressed in BAT by non-adipocyte cell types, such as endothelial cells and immune cells, and that these cell types play an essential role in maintaining an optimal microenvironment for BAT (Pope *et al.* 2016, Panina *et al.* 2018).

Upstream regulator analysis of the sex-differentially expressed genes identified ERBB2 and TGF β 1 as the most significant upstream regulators. ERBB2, a transmembrane tyrosine kinase receptor protein, is present in adipose-derived stem cells and is involved in adipogenic proliferation and differentiation (Scioli *et al.* 2014). TGF β 1, a member of the TGF β family of growth and differentiation factors, functions in adipose-derived stem cells to promote angiogenesis for optimal blood supply during adipose expansion (Scioli *et al.* 2014). In mice, serum TGF β 1 levels correlate with obesity and blocking of its principal signal transducer Smad3 or treatment with a TGF β 1 neutralizing antibody resulted in browning of WAT and prevented the development of obesity (Yadav *et al.* 2011). Upstream regulator analysis also identified E2, progesterone, and DHT as endogenous hormones regulating the expression of the identified genes. Our findings that ovariectomy reduced the mRNA expression levels while orchiectomy almost had no significant effect, strongly suggest a predominant role for female sex steroids in the sex-dimorphic expression pattern in BAT.

As discussed previously, most of the identified structural and cell-adhesion genes are likely expressed by non-adipocyte cell types in BAT. Additional studies, such as single-cell approaches that address the tissue cell heterogeneity, are therefore needed to determine which cell types express these genes in order to identify their role in BAT and whether these genes are directly regulated by sex steroids. Based on the literature, progesterone might regulate the expression of our identified genes, such as *Cdh1*, *Cfd*, and *Lcn2*, because these have been reported to show progesterone-dependent regulation in the endometrium, a tissue that becomes highly vascularized during the preimplantation stage (Yao *et al.* 2003, Jeong *et al.* 2005, Jha *et al.* 2006, Liu *et al.* 2016).

The role of progesterone in regulating BAT function has been studied in much less detail than the other female sex steroid E2, despite its elevated levels during the luteal phase and pregnancy. Furthermore, available data for progesterone effects on BAT activity are conflicting. Regarding the thermogenic function of BAT, progesterone treatment was found to induce NE-stimulated *Ucp1* mRNA expression in primary cultures of mouse brown adipocytes,

but surprisingly only at the lowest concentration of 10^{-9} M and not at higher concentrations of 10^{-8} or 10^{-7} M (Rodriguez *et al.* 2002). In contrast, another study showed that the mRNA expression of *Ucp1* and other thermogenic genes in murine BAT was reduced during pregnancy as well as by *in vivo* progesterone treatment. Furthermore, treatment of primary brown adipocytes by 5×10^{-7} M progesterone reduced NE-stimulated *Ucp1* mRNA expression (McIlvride *et al.* 2017). In our study, progesterone stimulation of primary brown adipocytes at a physiological concentration for female mice (10^{-7} M) (Nilsson *et al.* 2015) or at supra-physiological concentrations (10^{-6} and 10^{-5} M) did not affect *Ucp1* mRNA expression. However, in the T37i brown adipocytes, progesterone treatment at high concentrations ($10^{-5.5}$ and 10^{-5} M) clearly inhibited *Ucp1* mRNA expression. Expression of common mature brown adipocyte markers such as *Ppargc1a*, *Pparg*, and *Fabp4* was not regulated by progesterone in primary cultures, although they showed a sex-dependent expression pattern, being higher in female than in male adipocytes. Of note, concentrations of progesterone used in our study should be adequate to mediate progesterone effects because its equilibrium dissociation constant (K_d) has been reported to be 2–3 nM in various tissues (Jacobs & Smith 1981).

The difference in progesterone effects in SVF-differentiated brown adipocytes and T37i brown adipocytes might be driven by differences in the expression level of PR and its interaction with GR (Ruiz *et al.* 2020). In addition, the recently suggested functional heterogeneity of brown adipocytes and/or the presence of other cell types (e.g. macrophages, endothelial cells, fibroblasts) with adipocytes in the primary cultures could contribute to differences compared with a clonal cell (Bora & Majumdar 2017, Song *et al.* 2020). Hence, further studies are required to investigate the *in vivo* effects of progesterone on the BAT transcriptional profile and function to draw firm conclusions about the role of progesterone in the regulation of BAT activity and to determine whether it plays a role in the observed sex differences in BAT and whether this is important in humans and might be an underlying factor in clinical conditions. A study in rats has shown that energy balance and energy intake varies during the estrous cycle (Giles *et al.* 2010). So far, these changes during the cycle have been thought to be driven by changes in circulating estrogens. However, based on our results we speculate that changes in circulating progesterone, leading to an altered estrogen/progesterone ratio, also play a role. Moreover, UCP1-independent regulation of BAT thermogenesis, such as creatine and

lipid cycling, should not be ruled out (Chouchani *et al.* 2019), although the genes involved in these pathways were not identified in our *in silico* analysis.

Apart from its role in thermogenesis, BAT also serves as an endocrine organ, secreting several proteins including adiponectin and adipsin (also known as complement factor D) (Ali Khan *et al.* 2018, Villarroya *et al.* 2019) of which the gene expressions in primary brown adipocytes were dose-dependently reduced by progesterone. Adiponectin is a metabolic-favorable adipokine, having insulin-sensitizing and anti-inflammatory effects, of which the plasma concentrations are reduced in obesity (Stern *et al.* 2016, Achari & Jain 2017). The fact that ovariectomy increased while pregnancy decreased plasma adiponectin levels suggests an inhibitory effect of progesterone (or other pregnancy-related hormones) on adiponectin secretion from adipocytes (Combs *et al.* 2003), which is in line with our *in vitro* results. However, another study in female rats found no significant effect of progesterone treatment on *Adipoq* mRNA expression in BAT but instead observed reduced *Adipoq* mRNA in inguinal WAT (Stelmanska *et al.* 2012). Adipsin is necessary for initiation of the alternative complement pathway, an innate immune response for pathogen recognition and elimination (Xu *et al.* 2001) and plays a role in maintaining pancreatic β cell function (Lo *et al.* 2014). Adipsin also indirectly promotes adipogenesis and triglyceride storage in adipocytes via the product of its enzymatic action, the acylation-stimulating protein (ASP) (Cianflone *et al.* 2003, Song *et al.* 2016). Our results show that progesterone inhibited adipsin expression in primary brown adipocytes of murine origin. It remains to be determined how these results translate to human since serum adipsin levels were reduced during pregnancy but did not differ between the follicular and luteal phase in humans (Poveda *et al.* 2016).

Interestingly, using different concentrations of RU486 revealed that the progesterone-inhibiting effects on mRNA expression of various genes in brown adipocytes are possibly driven by GR rather than PR because the inhibition was blunted when RU486 was applied at a GR- and PR-antagonistic concentration but not at a more PR-specific antagonistic concentration (Kroon *et al.* 2018). Our findings agree with previous studies, for instance, progesterone was reported to act via the GR to mediate regulatory T cell enrichment (Engler *et al.* 2017) and to repress *PTGS2* mRNA expression in myometrial cells (Lei *et al.* 2012). However, non-genomic actions of progesterone, for example, through progesterone membrane receptors and the progesterone receptor membrane components, cannot be ruled out.

In summary, this study demonstrates a sex-dependent transcriptomic profile in BAT of mice at basal conditions and suggests a role for progesterone in the regulation of some of the identified sex-differentially expressed genes. Furthermore, our results highlight a role for genes encoding proteins involved in cellular structure, cell-cell contact, and cell adhesion in the sex-differential function of BAT.

Supplementary materials

This is linked to the online version of the paper at <https://doi.org/10.1530/JME-20-0210>.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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Author contribution statement

K K, A G and J A V conceived study. K K, A G, J S and A M designed and performed experiments. S M A S was involved in RNA-seq data analysis. K K, A G and J S were involved in all other data analyses. All authors contributed to result interpretation. K K, A G and J A V involved in manuscript preparation. All authors reviewed and approved the manuscript.

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