Retinoic Acid Mediates Visceral-Specific Adipogenic Defects of Human Adipose-Derived Stem Cells

Increased visceral fat, rather than subcutaneous fat, during the onset of obesity is associated with higher risk of developing metabolic diseases. The inherent adipogenic properties of human adipose-derived stem cells (ASCs) from visceral depots are compromised compared with those of ASCs from subcutaneous depots, but little is known about the underlying mechanisms. Using ontological analysis of global gene expression studies, we demonstrate that many genes involved in retinoic acid (RA) synthesis or regulated by RA are differentially expressed in human tissues and ASCs from subcutaneous and visceral fat. The endogenous level of RA is higher in visceral ASCs; this is associated with upregulation of the RA synthesis gene through the visceral-specific developmental factor WT1. Excessive RA-mediated activity impedes the adipogenic capability of ASCs at early but not late stages of adipogenesis, which can be reversed by antagonism of RA receptors or knockdown of WT1. Our results reveal the developmental origin of adipocytic properties and the pathophysiological contributions of visceral fat depots.

Obesity is defined as excess fat mass in the body and is generally associated with increased risk of developing metabolic diseases, such as cardiovascular diseases and type 2 diabetes (1). At least two main types of white adipose tissue (WAT) are present in human and animals—namely, subcutaneous (SC) fat and visceral (VS) fat. Body fat distribution is increasingly recognized as one of the key factors explaining the metabolic heterogeneity of obesity. Increased visceral adiposity is particularly associated with the risk of developing metabolic complications, whereas increased SC fat presents no or little risk and, rather, is considered to be protective (2–4). These two types of fat differ in their pathophysiological properties, including insulin sensitivity, adipokine secretion, lipolysis, and development of inflammation (5).

Adipose tissue expands not only through increased lipid storage in existing adipocytes (leading to hypertrophy) but also by the differentiation of new adipocytes from progenitor/stem cells (leading to hyperplasia). There are intrinsic differences in the properties of cells from different depots of WAT in vivo and in vitro. It is generally believed that when excess lipids systemically accumulate in the overnutrition state, cells from SC fat mainly undergo hyperplasia, whereas cells from VS fat tend to expand by hypertrophy in vivo (6). Although regulation of adipocyte differentiation has been extensively characterized (7,8), little is known about the molecular
basis of regional differences in adipogenic differentiation capacities. Adipose-derived stem cells (ASCs) and adipose progenitor cells from SC and VS depots have intrinsic differences in vitro, such as proliferation and differentiation potentials (9–12). ASCs derived from SC fat differentiate easily into mature adipocytes, whereas VS-derived ASCs differentiate poorly in response to a standard induction cocktail (9). This explains the different expression levels of key adipogenic factors such as peroxisome proliferator–activated receptor (PPAR)–γ and C/EBPα in mature adipocytes and adipose tissue (13,14). As another example of inherent molecular differences, we recently demonstrated that distinct, selective cell surface markers are expressed in SC ASCs versus VS ASCs and reflect their adipogenic properties (15). In addition, previous reports showed that adipose tissue cells from different depots have distinct patterns of gene expression, especially in the category of developmental genes (e.g., the Hox family), in humans and rodents (16–18). However, how these differences in developmental gene expression lead to functional differences of ASCs is not clear. We hypothesize that intrinsic differences in certain signaling pathways at the progenitor or stem cell level may account for depot-specific differences, with consequences in adipose cell properties and body fat distribution.

In this study, we found WT1-mediated upregulation of the retinoic acid (RA) signaling pathway in ASCs from VS fat, which leads to early, but not late-stage, inhibition of adipogenesis. Our data suggest a contribution of RA to controlling the depot-specific gene program during the functional development of adipocytes in human WAT.

**RESEARCH DESIGN AND METHODS**

**Isolation and Culture of ASCs**

WAT was isolated from the SC depot of the abdominal region and the VS depot of the omental region of 10 human volunteers (S1–7 and S11–13) undergoing bariatric surgery, with approval by the National Healthcare Group Domain Specific Review Board, Singapore. The subjects S1–7, S11, and S12 have been described previously (15). S13 is a 47-year-old Chinese man. ASCs were isolated from WAT and cultured, as previously described (19). Only cells with a doubling time shorter than 36 h were used (up to p9), and cell samples with similar passage numbers were used for any comparative studies. Mesenchymal stem cell surface markers and the multipotency of ASCs used in this study were confirmed by flow cytometry and differentiation assays, respectively (15).

**Adipogenesis and AdipoRed Staining**

On day (D) 0, 2 days after reaching the confluent state, cells were induced with an adipogenic differentiation cocktail containing 1 μmol/L dexamethasone, 0.5 mmol/L isobutylymethykanthine, and 167 nmol/L insulin plus 100 μmol/L indomethacin, 8 mg/L biotin, and 4 mg/L pantothenate. On D6, cells were switched to a medium with 1 μmol/L dexamethasone and 167 nmol/L insulin plus 100 μmol/L indomethacin and then maintained until at least D12. The cells then were washed with PBS and stained with AdipoRed (Lonza) according to the manufacturer’s protocol. After 30 min, fluorescence readings were recorded at 485-nm excitation and 572-nm emission. Fluorescence images were captured using ImageXpress Micro (Molecular Devices).

**Real-Time PCR**

Total RNA was extracted using the TRizol reagent (Invitrogen) and purified with the Column RNeasy Kit (Qiagen), according to the manufacturer’s instructions. cDNA was converted using the RevertAid H Minus First Strand cDNA Synthesis Kit (Fermentas). Quantitative PCR (qPCR) was performed using SYBR Green PCR Master Mix on a StepOnePlus Real-Time PCR System (Applied Biosystems) using the primer pairs shown in Supplementary Table 4. Relative mRNA were calculated and normalized to the level of GAPDH.

**RA-Responsive Luciferase Assay**

To determine relative intracellular levels of RA activity, a firefly luciferase under the control of an RA-responsive element (RARE) in the pGL3 vector and control plasmid pRL-Renilla luciferase under the control of the cytomegalovirus promoter were transfected together into 293T cells with Lipofectamine 2000 (Invitrogen) per the manufacturer’s protocol. Then, 24 h after transfection, the 293T cells were incubated with either fresh media (DMEM + 15% FBS) or human ASC–conditioned media. After another 24 h, RA activities were assessed by measuring the dual activities of firefly and Renilla luciferases per the manufacturer’s protocol (Promega) and normalizing firefly luciferase activities relative to the control (Renilla) luciferase.

**Electrophoretic Mobility Shift Assay**

Nuclear fractions from SC and VS ASCs were isolated using a previously described method (20). The oligonucleotides containing the WT1 binding site on the human ALDH1A2 promoter (5′-AGAACTCAGAGTGGGAGATGTTCCCT-3′) were hybridized to their respective complementary strands and labeled at the 3′ end with biotin tetraethylene glycol (Sigma-Aldrich). Nuclear extracts obtained from ASCs were incubated with the biotin-labeled probe and then subjected to electrophoresis, transferred to a nylon membrane, and ultraviolet cross-linked. The membrane then was probed with stabilized Streptavidin–horseradish peroxidase conjugate and developed using the LightShift Chemiluminescent electrophoretic mobility shift assay (EMSA) kit (Thermo Scientific/Pierce Biotechnology).

**Chromatin Immunoprecipitation Assay**

Chromatin immunoprecipitation assay was performed based on a previous protocol (21), with minor modifications. Briefly, ASCs were fixed with 1% formaldehyde, stopped by adding 125 mmol/L glycine, and then washed and collected in ice-cold PBS. The cell pellets were resuspended in lysis buffer, and the resulting pellet of crude nuclear
extract was resuspended in a high-salt lysis buffer. The nuclear extracts were sonicated on ice using a Misonix 3000 to yield DNA fragments 200–500 base pairs in size. Immunoprecipitation was conducted overnight at 4°C with 2 μg of anti-WT1 antibody (sc-192x; Santa Cruz Biotechnology) or anti-IgG antibody. PureProteome pre-blocked protein A magnetic beads (100 μL; Millipore) were incubated with the samples at 4°C for 2 h with rotation and eluted for 2 h at 65°C. After the cross-linking was reversed overnight at 65°C, DNA was extracted and PCR performed using the following set of primers: WT1 forward: 5’-ATCCCATTTCTATCTACTCC-3'; WT1 reverse: 5’-CCTGCACTGCTTTGTCTATCT-3’.

**Gene Knockdown Through Lentivirus**

Short hairpin RNAs (shRNAs) targeting WT1 were designed using publicly available algorithms from GenScript. Nine different shRNAs were synthesized, and the one that provided the best knockdown efficiency was selected for further experiments. Stable knockdown of WT1 was achieved using lentivirus expressing the shRNA sequence GCTCCAATCCTATCAAAACTCAGAGTGTGAGGAGGAGGAGGGTTT. Sense and antisense sequences are denoted in italics, whereas the loop sequence is underlined. A nontargeting shRNA was used as scrambled control. Lentiviral particles were produced by transient transfection of 293T cells with a transfer vector and pVSV-G, pMDL/prRE, and pRSV-Rev packaging vectors. The viral supernatant was collected, filtered, and concentrated by ultracentrifugation. Stable cell lines were generated by transducing human ASCs at a multiplicity of infection of 4 in complete media containing polybrene.

**Liquid Chromatography/Mass Spectrometry Analyses of Retinoids**

To measure cellular retinol levels, highly sensitive ultraperformance liquid chromatography coupled with mass spectrometry (LC-MS) was used (22). For all-trans RA (atRA) extraction, samples were subjected to a two-step acid-base extraction as described before (23), with minor modifications. The details are described in the Supplementary Data.

**Western Blot Analysis**

Protein from ASCs was extracted in radioimmunoprecipitation assay buffer and subjected to Western blot analysis. The concentration of protein in lysates was determined using Bradford assay. Protein (20 μg) was separated on 4–20% Mini-PROTEAN gels (Bio-Rad) and then transferred to nitrocellulose membranes. The membranes were probed with primary antibodies, WT1 (1:1,000 dilution; Santa Cruz Biotechnology) and α-tubulin (1:5,000 dilution; Sigma). After subsequent washes and incubation with respective secondary antibodies, the horseradish peroxidase activity was detected using chemiluminescent reagents.

**Data Analysis**

All results are presented as means ± SEMs. Comparisons between groups were analyzed using two-sided, paired t tests. Differences with a P value <0.05 were considered significant.

**RESULTS**

**RNA-Seq and Microarray Analysis of SC and VS Fat Depots Highlight the Differential Regulation of Genes Involved in Retinoid Metabolism**

We performed next-generation sequencing analysis of eight paired SC and VS WAT depots from subjects without diabetes using the Illumina HiSeq 2000 system. Through the RNA-seq analysis, a total of 1,185 annotated genes exhibited significant differences between VS and SC fat depots (significance was defined as fold change >2.0 and a false discovery rate <0.05) (Supplementary Table 1). Of these genes, 792 had higher expression in VS fat, whereas 393 genes were predominant in SC fat. Ontological analysis of these differentially regulated genes highlighted just three categories that reached significance as assessed by P value <0.05 after Bonferroni correction (Fig. 1A). One of these was found to be “retinol metabolism,” which includes VS-enriched genes (set red in Supplementary Table 1) and SC-dominant genes (set blue in Supplementary Table 1).

Because different WAT regions contain a number of distinct cell types, especially infiltrating immune cells, it is plausible that many of the differentially regulated genes mentioned above are influenced by cell type differences in SC and VS depots. To determine what portion of these genes are influenced by intrinsic differences in the stem cell population, we performed global gene expression analysis of human ASCs from six paired SC and VS depots using an Illumina HumanHT-12 array. Of these, 171 genes showed a fold change >2.0 and a positive false discovery rate <0.05 (Supplementary Table 2). Among these genes, 85 were expressed predominantly in SC ASCs (Supplementary Table 2A) and 86 were expressed more in VS ASCs (Supplementary Table 2B). In our analysis, several HOX genes were differentially expressed in ASCs from different depots, as previously reported by others (16–18): HOXA2, HOXA4, and HOXA5 were upregulated in VS ASCs, whereas HOXC6, HOXC8, HOXC9, HOXA9, and HOXA10 were upregulated in SC ASCs. Consistent with our previous report (15), the cell surface protein MME/CD10 was selective for both SC ASCs and SC fat, whereas CD200 was higher in both VS ASCs and VS tissue at the gene expression level (Supplementary Tables 1 and 2).

Similar to the RNA-seq analysis of tissues, ontological analysis of these differentially expressed genes in ASCs from different depots also revealed enrichment of genes categorized in the “retinoid metabolic process.” It was previously estimated that RA can modulate the expression of a few more than 530 genes (24). Of these, 15 RA-responsive genes among 86 of the upregulated genes in VS ASCs and 6 RA-responsive genes among 85 of the upregulated genes in SC ASCs were identified. We found that genes participating in RA synthesis (RDH10, CRBP1, and DHRS3) and reportedly modulated by RA (CRBP1, MGP, RARRES1, BAPX1, RARRES2, NR2F1, PTGS1, CDKN2B, PLAT, HOXA4, HOXA5, F3, HSD17B1, MEIS1, and CDH6) were upregulated in VS ASCs (Supplementary Table 3A). Also, six putative RA-responsive genes (MME, PITX2, TNC,
ENPP2, CTSK, and CRYAB) were upregulated in SC ASCs (Supplementary Table 3B). The acquisition of RA-responsive genes in a depot-specific manner is highly significant: 15 of 86 VS-enriched genes (17.4%) and 6 of 85 SC-enriched genes (7.1%) versus 532 of the 31,000 total RA-responsive genes/all annotated genes from our microarray data (1.7%). These results indicate that a significant number of gene expression differences are present, even in stem cell populations between distinct fat depots, and that the retinoid/RA metabolism pathway may represent one of the most important categories underlying pathophysiological differences in VS versus SC fat.
Validation of Depot-Specific Expression of RA-Related Genes by qPCR

Our microarray results showed that the expression of genes involved in the first retinoid synthesis—RDH10, CRBP1, and DHR53—was increased in VS ASCs when compared with that in SC ASCs. This observation led us to question whether other RA-related metabolic genes were differentially expressed in SC and VS ASCs. Retinol, a vitamin A species, is metabolized to hydroxy-RA by cytochrome P450 proteins CYP26A1, CYP26B1, and CYP26C1. RDH10, which converts retinaldehyde to retinal, is expressed in VS ASCs (Fig. 2A). CYP26A1 expression was significantly upregulated in VS ASCs in almost all 10 subjects. In the second step, retinaldehyde dehydrogenases, which include ALDH1A1, ALDH1A2, and ALDH1A3, irreversibly oxidize retinol to RA. ALDH1A1 (Fig. 2D) and ALDH1A3 (Fig. 2F) did not show depot specificity, whereas ALDH1A2 expression (Fig. 2E) was significantly higher in VS ASCs than in SC ASCs. Finally, RA is metabolized to hydroxy-RA by cytochrome P450 proteins CYP26A1, CYP26B1, and CYP26C1. CYP26A1 (Fig. 2G) and CYP26C1 (Fig. 2I) did not show depot specificity, whereas CYP26B1 expression (Fig. 2H) was significantly lower in VS ASCs when compared with SC ASCs. In the downstream pathway, RA activates the RA receptor (RAR) family (RARs α, β, and γ) or retinoid X receptors [RXRs] α, β, and γ), which contains ligand-dependent transcription factors of the nuclear receptor family, to regulate the expression of many RA target genes (Fig. 1B). Of the six genes measured, only RARB expression was upregulated in VS ASCs (Fig. 2K), whereas no change was observed in expression of the other five genes (Fig. 2J, L, M–O). Uregulation of RARB coincides with the results of an earlier report, which showed that gene expression of RARB, but not other RA receptors, is inducible by RA itself (26).

In General, RA Induces VS ASC Genes and Suppresses SC ASC Genes

Using qPCR, we confirmed the depot specificity of 14 of the 15 genes that showed significantly higher expression in VS ASCs (Figs. 2B and 3A–M); only changes in HSD17B1 (Fig. 3N) did not reach statistical significance. All of the six genes were validated to exhibit significantly higher expression in SC ASCs (Fig. 3O–T).

To investigate the RA-mediated responses of these genes, we treated SC ASCs with different concentrations of exogenous RA for 48 h. Eleven of 14 VS ASC genes (CRBP1, MGP, RARRES1, RARRES2, NR2F1, CDKN2B, PLAT, HOXA4, HOXA5, F3, CDH6) showed increased expression upon RA treatment, generally in a concentration-dependent manner (Supplementary Fig. 1A–F). In contrast to the VS ASC genes listed above, four of six SC ASC genes (MME, TNC, ENPP2, CRYAB) showed decreased expression upon RA treatment in a dose-dependent manner (Supplementary Fig. 1B and I). These data indicate that VS ASC-enriched genes are mostly induced by exogenous RA, whereas SC ASC genes are generally downregulated by RA treatment.

VS ASCs Exhibit Higher Endogenous RA Levels

Since higher RA-mediated activity was observed in VS ASCs than in SC ASCs, we assessed endogenous RA levels using three different methods. First, a Luciferase reporter assay using the RARE indicated that VS ASCs rendered a significantly higher RA-responsive activity than SC ASCs (Fig. 4A). Second, endogenous levels of RA were assessed by ultrasensitive surface-enhanced Raman spectroscopy using the scheme and spectra shown in Supplementary Fig. 2A–C. The surface-enhanced Raman spectroscopy measurement indicated significant upregulation of RA in VS ASCs compared with SC ASCs (Supplementary Fig. 2D). A similar increase of endogenous RA in VS ASCs was observed when the conventional method using LC-MS was used (Fig. 4B). The level of the RA precursor retinol, as measured by LC-MS, also indicated a significant increase in VS ASCs compared with SC ASCs (Fig. 4C).

RA Inhibits Adipocyte Differentiation of Human ASCs at Early Stages

It was previously reported that RA is a potent inhibitor of adipocyte differentiation in mice (27, 28). To determine whether RA affects the adipocyte differentiation of human ASCs and, if so, at which adipogenic stage, we treated ASCs with two different concentrations of RA (1 and 10 μmol/L) at different times during adipocyte differentiation. The results showed that the early addition of RA drastically inhibited adipocyte differentiation during the predifferentiation period and during the first 6 days after differentiation (D−2 to D0, D0 to D3, or D3 to D6) and did so to a greater extent in SC ASCs (Fig. 5A and B1–v) than VS ASCs (Fig. 5A and Supplementary Fig. 3i–v). A higher concentration (10 μmol/L) of RA corresponded with a greater reduction in adipocyte formation, especially that of SC ASCs, when compared with a lower concentration (1 μmol/L) of RA. Importantly, the adipogenic level of SC ASCs treated with 10 μmol/L RA during D0 to D3 or D3 to D6 was similar to that of nontreated VS ASCs. Also, just 2 days of pretreatment with RA itself was sufficient to significantly reduce adipocyte formation in ASCs. RA treatment at later stages of differentiation—that is, D6 to D9 and D9 to D12—either minimally inhibited or failed to block the adipocyte formation in both SC and VS ASCs (Fig. 5A and Bvi–vii; Supplementary Fig. 3vi–viii). Furthermore, as shown in Supplementary Fig. 4, treatment of SC and VS ASCs with retinol (Supplementary Fig. 4A and ii) or retinal (Supplementary Fig. 4B and ii) intermediates in the RA synthesis pathway, significantly inhibited adipocyte differentiation at various time points during adipogenic induction, as observed by AdipoRed staining.
Figure 2—Depot-specific expression profiles of genes involved in RA metabolism. qPCR was performed on 10 pairs of patient-derived SC and VS ASCs (S1–7 and S11–13). The graphs show differential mRNA expression of RDH10 (A), CRBP1 (B), DHRS3 (C), ALDH1A1 (D), ALDH1A2 (E), ALDH1A3 (F), CYP26A1 (G), CYP26B1 (H), CYP26C1 (I), RARA (J), RARB (K), RARG (L), RXRA (M), RXRB (N), and RXRG (O) in S1–7 and S11–13. The values are relative arbitrary units and are normalized to the housekeeping gene GAPDH. **P < 0.01 denotes significant differences in pairs of ASCs (n = 10).
Figure 3—Differential expression of RA-regulated genes in SC and VS ASCs. The graphs show differential mRNA expression of genes upregulated in VS ASCs, including MGP (A), RARRES1 (B), RARRES2 (C), BAPX1 (D), NR2F1 (E), PTGS1 (F), CDKN2B (G), PLAT (H), HOXA4 (I), HOXA5 (J), F3 (K), MEIS1 (L), CDH6 (M), and HSD17B1 (N), and genes upregulated in SC ASCs, such as MME (O), PITX2 (P), TNC (Q), ENPP2 (R), CTSK (S), and CRYAB (T), in S1–7 and S11–13. The values are relative arbitrary units and are normalized to the housekeeping gene GAPDH. *P < 0.05 and **P < 0.01 denote significant differences in pairs of ASCs (n = 10). See also Supplementary Fig. 1.
the expression of early and late adipogenic markers in RA-treated SC ASCs (Fig. 6E–H) were very similar to those in VS ASCs (Fig. 6A–D). The results imply that the adipogenic defect of VS ASCs may be at least partially influenced by higher RA-mediated activities.

Visceral-Specific Developmental Factor May Act to Facilitate the RA Signaling Pathway by Modulating the Expression of the RA Synthesis Enzyme

It was recently reported that the Wt1 gene, a mesothelial developmental marker, is specifically expressed in VS fat, but not in SC fat, in mice (30). Consistent with this result, we found that WT1 is selectively expressed 24-fold more, on average, in VS ASCs compared with SC ASCs, as assessed using qPCR; this reflects the presence of WT1 in the stem cell population of human VS fat (Fig. 7A). In the search for a developmental factor linking to the RA synthesis pathway, we found a WT1 binding site within the promoter region of the human ALDH1A2 gene (Fig. 7B) that is upregulated in VS ASCs and produces the dehydrogenase enzyme that oxidizes retinol into RA (Fig. 1B). Using EMSA, the antibody against WT1 was found to specifically bind the DNA probe spanning the ALDH1A2 promoter with higher affinity in VS ASCs than in SC ASCs (Fig. 7C). The specificity of WT1 was confirmed when a diminished band intensity or the disappearance of a shifted band is observed by (pre)incubating the VS ASC nuclear extracts with 100× the concentration of competitor oligos or WT1 antibodies. In addition, chromatin immunoprecipitation experiments showed that the anti-WT1 antibody, but not the control IgG antibody, pulled down the ALDH1A2 promoter region at a higher level in VS ASCs than SC ASCs (Fig. 7D). Finally, the WT1 gene was knocked down through the lentiviral construct in SC ASCs and VS ASCs. qPCR and Western blotting showed effective knockdown of WT1 in both gene and protein expression (Fig. 7Ei and ii). It was found that high expression of the ALDH1A2 gene in VS ASCs was dramatically suppressed by knockdown of WT1 compared with the scrambled control (Fig. 7F). Importantly, the knockdown of WT1 in VS ASCs significantly increased adipogenesis, as shown by the AdipoRed staining in Fig. 7Gi and ii. These results suggest that the developmental WT1 protein can upregulate the RA pathway directly through the induction of the ALDH1A2 gene in VS ASCs, and genetic ablation of WT1 improves their adipogenic capacity.

RA-Mediated Adipogenic Defects Are Reversed by Antagonism of the Downstream Target RAR

Finally, to address whether adipogenic defects caused by excessive RA can be reversed by modulating the RA pathway in VS ASCs, we used BMS493, an antagonist of RARs (α, β, γ), at different points during adipocyte differentiation. The result of AdipoRed staining and quantification demonstrated that treatment of BMS493 during D−2 and D12 significantly improved the adipogenic capacity of VS ASCs (Fig. 8A and B). BMS493 also reversed the adipogenic defect caused by excessive RA at all the
Figure 5—RA inhibits the early stage of adipocyte differentiation of ASCs. Using a standard adipogenic cocktail, adipocyte differentiation was induced in ASCs from S11, with or without pretreatment/treatment with RA at various time points. A: A representative graph showing relative fluorescence units (RFUs) of AdipoRed staining on ASCs from S11. The ASCs were treated with 1 or 10 μmol/L of RA at different time points, as indicated by D0 (D−2 to D0), D3 (D0 to D3), D6 (D3 to D6), D9 (D6 to D9), and D12 (D9 to D12). *P < 0.05 denotes significant fold change (in RFUs) against respective “No RA” control samples, corresponding to the quantitation of lipid accumulation during adipocyte differentiation. B: Representative images (original magnification ×10) showing the lipid accumulation (AdipoRed, green) and the nuclei (Hoechst 33342, blue) in SC ASCs from S11 treated with 1 or 10 μmol/L of RA at different time points: no induction (i), without RA (ii), with RA treatment during D−2 to D0 (iii), D0 to D3 (iv), D3 to D6 (v), D6 to D9 (vi), or D9 to D12 (vii). The field in ii is magnified and merged with the bright-field image to show the overlap of AdipoRed staining and lipid droplet structures (top right). Similar results were obtained from experiments using cells from S12. For the purpose of presentation, the fluorescent intensities were enhanced to the same degree for all original images. Scale bar = 100 μm. See Supplementary Fig. 3 for representative images in VS ASCs.
Figure 6—RA-mediated differential expression of adipogenic genes during adipocyte differentiation of ASCs. Adipocyte differentiation was induced in SC and VS ASCs from S11 using a standard adipogenic cocktail. RNA was collected at various time points during differentiation (D0, D2, D4, and D6), and the expression of adipogenic genes was analyzed by qPCR. The representative graphs show gene expression of CEBPB (A), CEBPD (B), CEBPA (C), and PPARG (D) in SC and VS ASCs from S11 at the various time points indicated (n = 2). Adipocyte differentiation was induced in SC ASCs from S11 using a standard adipogenic cocktail, with or without RA (10 μmol/L) treatment. RNA was collected at various time points (D0, D2, D4, and D6). The representative graphs show gene expression of CEBPB (E), CEBPD (F), CEBPA (G), and PPARG (H) in SC ASCs treated with or without RA during differentiation at various time points (n = 2). *P < 0.05 denotes significance. Similar results were obtained in cells from S12.
Figure 7—WT1 mediates the upregulation of the RA synthesis enzyme in VS ASCs. A: A qPCR graph showing substantially higher WT1 expression in VS ASCs from S1–S7 and S11–13. **P < 0.01 denotes significant change in pairs of ASCs (n = 10). B: Consensus WT1 binding site was found within the promoter region of the human ALDH1A2 gene. C: EMSA was performed using nuclear extracts from SC and VS ASCs from S11. The representative gel shows increased WT1 binding in VS ASCs, as indicated by the shifted band in lane 3, when compared with that of SC ASCs (lane 2) (lane 1 represents oligonucleotides [oligo] only). Diminished band intensity or the disappearance of a shifted band is observed when VS ASC nuclear extracts were incubated/preincubated with 100× the concentration of competitor oligonucleotides (lane 4) or with diluted 1:1 (dil.; lane 5) or concentrated (conc.; lane 6) WT1 antibody (n = 2). D: A representative agarose gel image showing the increased binding of WT1 to the ALDH1A2 promoter in VS ASCs from S11 (lane 9) compared with SC ASCs (lane 5).
times tested (Fig. 8A and B). We also treated VS ASCs with LE135, a RARβ-specific antagonist. AdipoRed staining revealed that treatment with LE135 significantly improved the adipogenic potential of VS ASCs, as shown in Fig. 8C and D. Together, these results propose a potential therapeutic strategy of RAR or RARβ antagonism that can relieve adipogenic dysfunction mediated by excessive RA signaling in VS fat.

**DISCUSSION**

In this study, through the ontological analysis of global gene expression, we found that a pathway regulated by RA is one of the major metabolic responses to developmental differences of stem cells from human fat depots. RA is a vitamin A metabolite that regulates many genes through specific binding to nuclear transcription factors, especially RARs (24,31). The endogenous concentration of RA is tightly regulated during embryonic and postnatal development and is essential for the proper functionality of adult tissues and cells. Several recent reports have indicated potential roles of retinoids (RA, retinol, retinal, and other vitamin A derivatives) in adipose tissue (32–34). Although the liver is the primary storage site of retinoids, adipose tissue is the second largest site and actively mediates retinoid metabolism (35,36). However, little is known about its fat depot–specific mechanisms and contributions, especially in the context of human stem/progenitor cells.

Our microarray and qPCR analysis indicated that genes that lead to RA synthesis, such as retinoid oxidoreductases (RDH10, DHRS3, and ALDH1A2), had higher expression in VS ASCs, whereas CYP26B1, a gene involved in the breakdown of RA, had higher expression in SC ASCs (Fig. 1B). RDH10 is a retinol dehydrogenase that has been shown to be essential for atRA biosynthesis in embryonic development (37), and DHRS3/retSDR1 seems to be a major retinal reductase that is also important for development in human and mice (38,39). ALDH1A2 is the primary isoform of retinal dehydrogenases that are critical for embryogenesis (40), whereas 1A1 and 1A3 isoforms are more important for postnatal and adult development and endocrine functions. In addition, the 1A2 isoform has a much lower Michaelis constant, Km (0.66 μmol/L), than that of 1A1 (11.6 μmol/L) and 1A3 (3.9 μmol/L) (25), indicating that, when all three isoforms are present, this isoform may be predominant in producing RA from retinal. Finally, for the catabolic enzymes, both CYP26A1 and CYP26B1 are important for embryonic development, but CYP26B1 is a major isoform in adult nonbrain tissues, at least in mice (25).

We found 14 genes that had higher expression in VS ASCs and were induced by RA. Of these, CDKN2B and CRBP1 are known to have an anti-adipogenic function (41,42). NR2F1 is expressed more in VS ASCs, and its paralog, NR2F2, has anti-adipogenic capacity as well (43). One VS ASC–specific and RA-inducible gene that is previously well studied is CRBP1. It was reported that overexpression of CRBP1 inhibits adipogenesis and knockdown of CRBP1 enhances adipogenesis in 3T3-L1 cells (41). Interestingly, CRBP1 knockout mice had increased adiposity, especially of epididymal fat (VS fat), but remained more glucose tolerant and insulin sensitive when fed a high-fat diet. Expression of the master adipogenic regulator PPAR-γ was significantly increased in WAT of CRBP1 knockout mice (41). Whether these VS-dominant genes are involved in the adipogenic defect or other VS-specific phenotypes in humans warrants further investigation.

Consistent with earlier studies showing a general inhibitory effect of RA in adipogenesis in vivo (27,44) or in vitro (28,45) in mice, we demonstrated that exogenous RA impaired adipocyte differentiation of human ASCs from different depots at the early stage of adipogenesis, although this occurred to a greater extent in SC ASCs. Interestingly, RA is known to enhance the commitment of embryonic stem cells into an adipocyte lineage when they were pretreated during embryoid body formation (46,47). It is possible that RA exerts distinct effects during different developmental stages or different stages of differentiation. Alternatively, only a higher concentration of RA, as used in our study (1 and 10 μmol/L) and observed in VS ASCs, may inhibit adipocyte differentiation by “overactivating” RAR-mediated signaling; the minimum concentration required for activating RAR is in the nanomolar range. However, the RA dose used for embryonic stem cells’ commitment into adipocytes is 0.1 to 1 μmol/L; we found that as little as 0.1 μmol/L RA is sufficient to inhibit ASC adipogenesis (data not shown). Further studies are necessary to delineate the mechanisms regulated by RA in different developmental/differentiation stages of cells. Together, our results suggest that RA, more of which is produced in VS ASCs than SC ASCs, modulates gene expression differences in ASCs from different depots, influences...
Figure 8—RA-mediated adipogenic defects are reversed by the antagonism of the downstream target RAR. A: A representative graph showing the relative fluorescence units (RFUs) of AdipoRed staining on VS ASCs from S11. Adipocyte differentiation was induced in ASCs using a standard adipogenic cocktail with or without pretreatment/treatment with BMS493 and/or RA at various time points. The ASCs were treated with 1 μmol/L of BMS493 and/or 10 μmol/L of RA at the time points indicated: D0 (D−2 to D0), D3 (D−2 to D3), and D12 (D−2 to D12). *P < 0.05 and ^P < 0.05 denote significant fold change in RFUs corresponding to the quantitation of lipid accumulation during adipocyte differentiation (n = 2). B: Representative images (original magnification ×10) showing lipid accumulation (AdipoRed, green) in VS ASCs from S11 treated with 1 μmol/L of BMS493 and/or 10 μmol/L of RA at the different time points indicated: D0 (D−2 to D0), D3 (D−2 to D3), and D12 (D−2 to D12). For the purpose of presentation, the fluorescent intensities were enhanced to the same degree for all images. Scale bar = 100 μm (n = 2). C: A representative graph showing the RFUs of AdipoRed staining on VS ASCs from S11. ASCs were treated with LE135 (10 or 25 nmol/L) for 3 days upon the induction of adipogenic differentiation with a standard adipogenic cocktail. *P < 0.05 and **P < 0.01 denote significant fold change in RFUs (n = 2). D: Representative images (original magnification ×10) showing lipid accumulation (AdipoRed, green) in VS ASCs from S11 treated with LE135 (10 or 25 nmol/L) for 3 days upon the induction of adipogenic differentiation. For the purpose of presentation, the fluorescent intensities were enhanced to the same degree for all images. Scale bar = 100 μm (n = 2). Similar results were obtained from experiments using cells from S12.
adipocyte differentiation, and may cause defects in the properties of adipose cells from visceral depots and in body fat distribution.

In addition, we found substantially higher expression of WT1, a developmental marker that marks all visceral organs within the abdominal and thoracic cavities, in human VS ASCs. We found that WT1 may control the expression of the VS-specific ALDH1A2 gene by directly binding to its promoter region in human VS ASCs. WT1 knockdown in VS ASCs led to the reduced expression of ALDH1A2 and improved adipogenesis. This result points to the developmental origin of accelerated RA production and signaling in the human VS depot.

Our finding of differential RA signaling may help to understand how cells from VS fat are compromised and lead to pathophysiologica phenotypes of this depot, such as immune cell infiltration, inflammation, limited lipogenesis, altered adipokine secretion, and insulin resistance. It was recently reported that peritoneal macrophages receive the RA signal from the omental region and migrate to the peritoneal cavity in a manner dependent on RA-induced genes (48). Interestingly, RA synthesis enzymes, especially Raldh2 (the mouse ortholog of ALDH1A2), are abundantly expressed in peritoneum-associated tissues with mesothelial origins. It would be of interest to explore further how the higher level of RA in VS fat affects the functional polarization and migration of macrophages and other immune cells, compared with the SC fat depot, which is relatively free from proinflammatory immune infiltration and reactions. Taken together, our investigations pave the way for understanding the developmental basis of visceral adiposity and the potentially immune infiltrative milieu of this depot.

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