Identification and Characterization of Sebaceous Gland Atrophy-Sparing DGAT1 Inhibitors

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Abstract

Inhibition of Diacylglycerol O-acyltransferase 1 (DGAT1) has been a mechanism of interest for metabolic disorders. DGAT1 inhibition has been shown to be a key regulator in an array of metabolic pathways; however, based on the DGAT1 KO mouse phenotype the anticipation is that pharmacological inhibition of DGAT1 could potentially lead to skin related adverse effects. One of the aims in developing small molecule DGAT1 inhibitors that target key metabolic tissues is to avoid activity on skin-localized DGAT1 enzyme. In this report we describe a modeling-based approach to identify molecules with physical properties leading to differential exposure distribution. In addition, we demonstrate histological and RNA based biomarker approaches that can detect sebaceous gland atrophy pre-clinically that could be used as potential biomarkers in a clinical setting.

Introduction

Diacylglycerol O-acyltransferase 1 (DGAT1) is ubiquitously expressed and catalyzes the final step in triglyceride (TG) synthesis [1]. TG biosynthesis has pleiotropic roles in various tissues. TG can be taken up by the diet and resynthesized in the small intestine by DGAT1 or can be synthesized de novo by either DGAT1 or DGAT2 in the liver and/or adipose tissues [2]. Inhibition of DGAT1 in the intestine has been shown to enhance circulating levels of gut incretin levels such as Glucagon-like peptide 1 (GLP-1) and Peptide YY (PYY) post-prandially [3,4]. In addition to DGAT1’s role in these tissues, DGAT1 and DGAT2 have also been demonstrated to be expressed in the skin of mice [5,6] and human (data not shown). Mice with a deletion of the DGAT1 enzyme (DGAT1 -/-) are protected from diet induced obesity and energy expenditure [7]. However, in addition to these metabolic phenotypes, DGAT1 -/- mice develop leptin-dependent abnormal skin phenotypes, characterized by sebaceous gland atrophy and hair loss [5]. The metabolic effects and the skin phenotype were shown to be recapitulated with pharmacological inhibition of DGAT1 [6]. Skin composition between human and preclinical species varies; wax dieter is the major sebum lipid in mouse while TG is the major form in human [8]. Although the exact role of sebum in human is not fully understood, sebum production could be decreased with pharmacological inhibition of skin DGAT1 activity.

Since the identification and the characterization of DGAT1 -/- mice, multiple pharmaceutical companies have been actively pursuing the discovery of small molecule DGAT1 inhibitors to reproduce the beneficial metabolic phenotypes of these mice [9,10]. Recent early clinical data with DGAT1 inhibitors have uncovered gastrointestinal adverse effects (AEs) as a major issue with no report of adverse skin effects [10–12]. However, considering the role of DGAT1 in the skin, such inhibitors represent potential liabilities related to skin AEs as well. To that end one of our goals was to develop small molecule DGAT1 inhibitors with differential exposures at the site of action vs. skin. Low exposures in the skin would protect from skin liabilities while maintaining the beneficial metabolic benefits associated with DGAT1 inhibition in other tissues such as the small intestine. Based on molecular modeling we demonstrated the correlation between lipophilicity of several DGAT1 small molecule inhibitors, skin histological findings and systemic and skin drug exposures. In addition we proposed an RNA-based approach that could be utilized as clinical biomarkers to detect sebaceous gland atrophy driven by DGAT1 inhibitors.

Results

Skin effects of DGAT1 inhibitors

Several DGAT1 inhibitors across different structural classes were tested for their effect on skin morphology after chronic treatment in mice (Figure 1 and Table 1). Compounds were separated into structural classes and assigned to groups A to E. Representative structures from groups A, B, and C are shown in Figure 1 (structures of compounds from groups D and E will be the subject of future reports). After 14 days of oral dosing several compounds either induced sebaceous gland atrophy in the skin or showed no response. As shown in Figure 2, the sebaceous glands in
the skin of mice treated with either vehicle or Cpd1 (3 mg/kg, 14 days) appeared normal while the skin of mice treated with Cpd2 (30 mg/kg, 14 days) had moderate to marked atrophic sebaceous glands on the dorsal surface, which were characterized by an overall decreased amount and size of sebaceous gland acini. Skin of mice treated with Cpd3 (30 mg/kg, 14 days) showed minimal to mild effects. The affected sebaceous gland units had fewer acinar cells and/or cells with decreased amount of cytoplasmic vacuolation. Frequently the sebaceous gland acini had consolidated, eosinophilic cytoplasm and pyknotic nuclei. No other histomorphologic changes were observed in these skin sections. Compound plasma exposures (Plasma μM) did not correlate with skin AEs (Scoring) or with skin exposure (Skin/Plasma Ratio; Table 1). However, skin exposures as it related to IC50 did correlate with skin AEs (Skin/IC50). The effects of the compounds were similar on ventral skin (Figure S1). Of note, longer exposure of Cpd1, which did not cause sebaceous gland atrophy at 14 days, in rats at up to 300 mg/kg for 4 weeks had no adverse effects in skin (data not shown). Additional studies will be required to study the effects of these compounds in skin after longer-term exposure.

Plasma and Skin Concentrations of Compounds

Plasma and corresponding skin concentrations of the compounds are shown in Table 1. As can be seen, following dosing of the compounds at 30 mg/kg for 14 days, the relative trough levels of the compounds in plasma and skin do not have a direct correlation. Compounds that exhibit high plasma exposures do not appear to have high skin exposure and vice versa. Similar lack of correlation is also observed with unbound plasma concentrations and skin exposures.

Lipophilicity of the Compounds

To gain a better understanding of the chemical physical properties of DGAT1 inhibitors as it relates to skin AEs, we evaluated lipophilicity (Log D) of these compounds which can be either measured by HPLC (mlogD) or calculated using the ACD software (clogD). Distribution into skin can be understood as a partitioning equilibrium dependent on compound properties. As listed in Table 1, 7 out of 22 compounds tested (Cpd2, 3, 5, 6, 12, 15, 22) were found to induce either “minimal to mild” or “moderate to marked” skin sebaceous gland atrophy after 14 days of treatment. Compounds with sufficient potency against the mouse enzyme (mDGAT1 IC50 nM) only show skin toxicity as measured by histopathology at sufficient exposure levels in the skin (Table 1). Interestingly, we find that compounds with skin effects can be distinguished from compounds without observable skin effects by plotting lipophilicity (clogD) against the observed histopathology (Figure 3). Compounds which show significant or minimal adverse effects in skin generally have a predicted clogD of greater than 1, while compounds with no observable effects all have a predicted clogD of less than 0.7.

We also plotted mlogD versus observed histopathology and found considerable overlap between the compounds showing and lacking adverse skin effects (Figure 3). While the mlogD and clogD are correlated, there is a poorer correlation of the most polar compounds and the mlogD does not correctly predict histopathological findings.

To better understand why measured logD more poorly differentiated compounds than calculated logD we plotted the frequency of logD measurements for about 4,000 DGAT1 compounds by method (Figure 4). While the two logD methods were correlated (R2 0.76), we found that the dynamic range of measured logD was significantly less than the calculated range with values ranging from 0 to 5.5, while calculated logD measurements ranged from −3 to 9. The shoulder at mlogD five represents the more lipophilic series B compounds and is less pronounced, but visible, in the clogD distribution. The narrower range of HPLC measured logD measurements does not significantly distinguish compounds, while the ACD calculated logD values allow for more differentiation.

RNA profiling of skin biopsies

To identify biomarkers underlying the skin pathophysiology associated with DGAT1 inhibition we initiated global gene
Discussion

Potent DGAT1 inhibitors are currently being developed for the treatment of hyper-triglyceridemia and obesity [6]. The major adverse effect observed so far in the clinic relates to gastrointestinal effects (nausea, diarrhea, and vomiting), with no report of skin issues [10]. Nevertheless, to avoid potential skin AEs, especially after chronic treatment, effective therapeutics will selectively inhibit DGAT1 in the liver and gut with minimal activity on other tissues such as skin. We described compound lipophilicity as a predictor of skin exposure with subsequent induction of sebaceous gland atrophy. The distribution coefficient, D, is a pH dependent measure of the propensity of a molecule to differentially dissolve in two immiscible phases, taking into account all ionized and unionized forms (microspecies). It serves as a quantitative descriptor of lipophilicity.

Interestingly, many compounds which carry a carboxylic acid moiety are associated with a lack of skin AEs (Cpd1 is an example). It is likely that the carboxylic acid leads to decreased lipophilicity which prevents the compound from entering the skin. To address this hypothesis, we modified Cpd1 and replaced the carboxylic acid with a tertiary alcohol group (Cpd22). As predicted, this moiety is associated with a lack of skin AEs (Cpd1 is an example).

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Table 1. Compound characteristics. Cpd1 is the same as DGAT1i in Lin, H et al (2013) [4], while Cpd2 is the same as Compound L in Liu, J et al (2013) [3].

| Study | Compound | Series | Plasma µM | Skin µM | Skin/Plasma Ratio | Fu Plasma | mlogD | clogD | mDGAT1 IC50 nM | mDGAT1 IC50 µM | Skin/IC50 | Scoring | clogD/mIC50 |
|-------|----------|--------|-----------|---------|-------------------|-----------|-------|-------|----------------|----------------|-----------|---------|-------------|
| 1     | Cpd1     | A      | 0.194     | 0.0621  | 0.320            | 0.310     | 0.35  | 0.00  | 34.6           | 0.035          | 1.80      | not remarkable | 0          |
| 1     | Cpd2     | B      | 0.163     | 2.25    | 2.25             | 0.01      | 4.04  | 4.27  | 31.4           | 0.031          | 71.61     | moderate to marked | 136         |
| 1     | Cpd3     | C      | 1.38      | 0.135   | 0.098            | 0.063     | 1.73  | 1.83  | 11              | 0.011          | 12.27     | minimal to mild | 166         |
| 1     | Cpd4     | D      | 0.855     | 0.111   | 0.130            | 0.090     | 0.90  | 0.44  | 58.9           | 0.059          | 1.89      | not remarkable | 7          |
| 2     | Cpd5     | D      | 0.461     | 0.138   | 0.298            | 0.09      | 1.08  | 1.61  | 5.2            | 0.005          | 26.45     | minimal to mild | 310         |
| 2     | Cpd6     | D      | 2.22      | 0.430   | 0.194            | 0.09      | 1.27  | 1.60  | 18.8           | 0.019          | 22.87     | moderate to marked | 85          |
| 2     | Cpd7     | D      | 0.171     | 0.002   | 0.012            | 0.098     | 1.37  | 0.42  | 32.4           | 0.032          | 0.07      | not remarkable | 13         |
| 2     | Cpd8     | D      | 0.0072    | 0.010   | 1.345            | 0.033     | 1.11  | 0.48  | 4.9            | 0.005          | 1.96      | not remarkable | 98         |
| 2     | Cpd9     | D      | 0.0047    | 0.006   | 1.324            | 0.133     | 0.94  | 0.43  | 12.3           | 0.012          | 0.50      | not remarkable | 35         |
| 3     | Cpd10    | E      | 0.0782    | 0.0159  | 0.203            | nd        | 1.81  | 0.00  | 10.9           | 0.011          | 1.46      | not remarkable | 0          |
| 3     | Cpd11    | D      | 0.199     | 0.0478  | 0.241            | 0.0836    | 0.73  | 0.00  | 29             | 0.029          | 1.65      | not remarkable | 0          |
| 3     | Cpd12    | A      | 2.84      | 0.669   | 0.236            | 0.256     | 1.34  | 2.07  | 40.4           | 0.040          | 16.56     | moderate to marked | 51         |
| 3     | Cpd13    | D      | 0.276     | 0.0131  | 0.047            | 0.06      | 1.37  | 0.00  | 33.3           | 0.033          | 0.39      | not remarkable | 0          |
| 3     | Cpd14    | D      | 0.209     | 0.0265  | 0.127            | 0.073     | 1.05  | 0.00  | 46.9           | 0.047          | 0.57      | not remarkable | 0          |
| 3     | Cpd15    | D      | 1.21      | 0.190   | 0.157            | nd        | 1.15  | 0.77  | 10.7           | 0.011          | 17.72     | moderate to marked | 72         |
| 3     | Cpd16    | E      | 0.196     | 0.0347  | 0.177            | 0.059     | 1.81  | 0.00  | 10.4           | 0.010          | 3.34      | not remarkable | 0          |
| 3     | Cpd17    | E      | 1.42      | BLQ     | 0.1021           | 1.58      | 0.09  | 6.3   | 0.006          | not remarkable | 0         |
| 4     | Cpd18    | D      | 1.13      | BLQ     | nd                | 0.99      | 0.65  | 11.7  | 0.012          | not remarkable | 56         |
| 4     | Cpd19    | D      | 5.38      | BLQ     | nd                | 1.20      | 0.00  | 17.5  | 0.018          | not remarkable | 0          |
| 4     | Cpd20    | D      | 2.19      | BLQ     | 0.0495           | 0.99      | 0.35  | 34.2  | 0.034          | not remarkable | 10         |
| 4     | Cpd21    | D      | 8.01      | BLQ     | nd                | 1.04      | 0.11  | 17.2  | 0.017          | not remarkable | 6          |
| 4     | Cpd22    | A      | 5.14      | BLQ     | 0.112            | 2.15      | 2.87  | 25.9  | 0.026          | moderate to marked | 111         |
| 4     | Cpd6     | D      | 8.39      | BLQ     | 0.033            | 1.27      | 1.60  | 18.8  | 0.019          | moderate to marked | 85          |

Plasma = plasma concentration; Skin = skin concentration; Skin/Plasma Ratio = skin to plasma concentration ratio; Fu Plasma = unbound fraction in plasma; mlogD = measured logD; clogD = calculated logD using ACD (measure of lipophilicity); mDGAT1 IC50 = in vitro potency; Skin/IC50 = skin concentration vs in vitro potency; Scoring = histological adverse effect score; clogD/mIC50 = calculated logD vs in vitro potency; BLQ = below detection. Bold = compounds causing atrophy.

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studies were approved by the Merck Research Laboratories Institutional Animal Care and Use Committee (Rahway, NJ). Mice were acclimated to nonspecific stress for 7 days before the onset of the chronic dosing studies where they were dosed orally with vehicle (0.5% methylcellulose) or compounds at 30 mg/kg, or at doses indicated, for 14 days.

Histology

Skin samples from DIO mice treated with DGAT1 inhibitors for 14 days were collected from the dorsal and/or ventral surface and immediately fixed in 10% buffered formalin for 24 hr at room temperature and embedded in paraffin. Paraffin specimens were sectioned at 5 μm and stained with hematoxylin and eosin, and were evaluated blindly with light microscopy. In general, the severity scores were determined using the following criteria: “not remarkable” = no abnormalities; “minimal”, “mild”, “moderate”, “marked”, and “severe” = less than 10%, 10–25%, 25–50%, 50–75%, and greater than 75% of sebaceous gland units with some evidence of atrophy, respectively. One slide/location with at least 50 sebaceous glands was used for scoring purposes.

Digital images were captured using a 20× objective and the ScanScope XT slide scanner system (Aperio Technologies, Vista, CA) and managed using Spectrum (Aperio). The size of the sebaceous glands were measured using ImageScope of Aperio system by the method previously described [17].

Plasma Compound Concentration

Concentrations of compounds in plasma were determined by LC-MS/MS following protein precipitation. To 100 μL of plasma sample, 20 μL of an acetonitrile solution of the internal standard (2,500 ng/ml), was added. This step was followed by protein precipitation with 300 μL of acetonitrile. Samples expected to contain analyte concentrations exceeding the upper limits of the calibration curves were diluted with control plasma. Sample mixtures were vortex-mixed for 15 minutes. After spinning in a centrifuge for 15 minutes at 1,000 × g (at 10°C), the supernatants were transferred by a Tomtec Quadra 96 Model 320 (Hamden, CT) to Empore 96 filter plates (Fisher Scientific, Springfield, NJ), and the resulting filtrates collected. These filtrates were dried at 25°C under a stream of nitrogen gas, on a Zymark TurboVap 96.
The residues were reconstituted with 100 μL of a 8:2 v/v mixture of mobile phases A (20 mM aqueous ammonium acetate containing 0.01% formic acid), and B (80:20 v/v, acetonitrile:methanol), vortex-mixed, spun in a centrifuge at 1,000 g (at 10°C), then analyzed by LC-MS/MS.

The LC-MS/MS system comprised of a Leap Technologies HTS-PAL autosampler (Carrboro, North Carolina), two Perkin Elmer Series 200 Micro Pumps (Norwalk, CT), an in-line precolumn filter, a Type W 6-port Valco switching valve (Vici, Houston, TX), and a Sciex API 363 mass spectrometer (Toronto, Canada). Chromatographic separation of the analytes was achieved by RP-HPLC.

Data were acquired and processed by Sciex Analyst version 1.1 software. Peak area ratios of analyte to internal standard were plotted as a function of the nominal concentrations of the analyte. A line of best fit was generated from the curve points by linear regression with a weighting factor of 1/x2. The equation of this line was then used to calculate the plasma ratios in samples.

Skin Compound Concentration

The skin tissue samples from mice were weighted and placed in 48-well plates (or tubes). A volume of water equivalent to three times the skin weight was added to each tube and homogenized by a Tissue Terror (Biospec Products, Inc.; Model 985-370). An aliquot of 100 μL homogenized blank skin tissue samples were transferred to 96-well plates designated for standard curve and QCs. Standard curve and QCs were prepared by adding aliquots of 25 μL working standard solutions (1–8000 ng/mL), to generate a tissue calibration curve of 1–8000 ng/mL. An aliquot of 100 μL dosed skin tissue homogenates were transferred to the 96-well plate and a 25 μL aliquot of acetonitrile:water 1:1 was added into each well containing dosed tissue samples to make the volumes of all samples, stds, and QCs equal. Protein precipitation was used for sample extraction by the addition of 500 μL of acetonitrile crash solution, containing IS. Internal standard used was Alprazolam (Sigma Aldrich) at 100 μM in the crash solution. Samples were vortexed for 30 seconds and then centrifuged at 4000 rpm for 5 min. The supernatant was transferred to a separate plate.

A Thermo Scientific-Cohesive LX-2 system consisting of a CTC Analytics autosampler, Flux Instruments AG pumps, and a valve module, controlled by Aria software was used. A Sciex API-4000 mass spectrometer (Toronto, Canada) was the detector using Analyst 1.4.2 software. The aqueous mobile phase (A) was water.
2.1 mm samples. The equation was used to determine the concentration in the unknown linear regression line with a \(1/x^2\) weighting factor. The line’s measure the analyte response. The peak area ratio of analyte to analytes. Multiple reaction monitoring (MRM) was used to ionspray interface was used in positive ion mode for detection of compound (5\(\text{m}\)). Unbound was determined as the amount of compound in buffer (unbound) was added to the opposite side. The plate was then sealed and placed in a shaking incubator set to maintain a temperature of 37\(\text{C}\) and constant shaking rate of 60 rpm for 4 hr. At the end of the incubation period, 50 \(\mu\text{L}\) aliquots of plasma and buffer were removed and analyzed by LC-MS/MS. Percent unbound was determined as the amount of compound in buffer (post incubation) divided by amount of compound in plasma (post incubation) \(\times 100\).

**Measured logD**

LogD was measured by high-throughput (HT) HPLC at pH 7.0. The chromatographic system consisted of an Agilent 1200 HPLC system composed of a G1379B degasser, G1312B binary pump, G1367C well-plate autosampler, G1316B thermo-statted column compartment, G1315C diode-array UV-vis detector, and ChemStation software, all from Agilent Technologies, USA. The separations were carried out on a Poroshell 120 EC-C18, 30 mm \(\times\) 3.0 mm I.D., 2.7 \(\mu\text{m}\), (Agilent Technologies, USA). The mobile phase consisted of phosphate buffered saline at pH 7 (mobile phase A) and acetonitrile (mobile phase B). The column oven temperature was set to 30\(\text{C}\). The HPLC analysis began with an isocratic step of 0.2 minutes at 3\% B at 1.5 ml/min, followed by a gradient from 5\% to 98\% B in 1.0 minute at 1.5 ml/ min. A second isocratic step of 0.2 minutes at 98\% B with a changing flow rate from 1.5 to 2.0 ml/min was then followed by a gradient from 98 to 5\% B in 0.1 minutes with the flow rate changing from 2.0 to 1.5 ml/min. The equilibration time between injections is 0.2 minutes at 5\% B. The injection volume is 5 \(\mu\text{L}\) and the spectrophotometric detection was set to 215, 238 and 254 nm.

A 10 mM DMSO stock solution of compound was delivered for analysis. A 100 \(\mu\text{M}\) standard solution was generated by diluting 10 \(\mu\text{L}\) of the 10 mM stock solution with 990 \(\mu\text{L}\) of diluent (10\% DMSO/10\% MeCN/80\% MeOH, \(v/v/v\)). The chromatographic system was calibrated with a set of standards with published shake-flask logD values. Linear regression was used to determine the calibration line relating the retention time to logD for the calibration standards. This line was then used to determine the HT HPLC logD of compound from the measured retention time by the HPLC/DAD analysis of the 100 \(\mu\text{M}\) standard solution.

**Calculated logD**

LogD was calculated using the ACD Labs v10 LogD Predictor software at pH 7.4 for all compounds (ACD Labs, Toronto, Canada)[10].

**DGAT1 Enzyme Assay**

The activity of DGAT1 inhibition was measured using membrane preparations from Pichia overexpressing human DGAT1 and DG/oleoyl CoA as substrates at Km concentrations in the presence of CPM (Life Technologies, D346), which is weakly fluorescent until reacted with free thiols of CoA released from oleoyl CoA after it is incorporated into diacylglycerol to form TG. Mock membranes showed minimal activity (data not shown). IC50s were calculated using either Assay Data Analyzer or GraphPad Prism4 software [3].

**RNA Profiling**

Skin biopsies (3–5 mm\(^2\)) from dorsal and/or ventral surfaces from D10 mice treated with compounds for 14 days were obtained and placed in RNAlater solution (Ambion, Austin, TX) at 4\(\text{C}\), according to the manufacturer’s instructions. After removal from the RNAlater solution, the samples underwent a freeze/manual pulverization step prior to Trizol (Invitrogen, Carlsbad, CA) polytron chloroform extraction. RNA was isolated using the Promega SV-96 total RNA kit (Promega, Madison, WI) according to the manufacturer’s instructions. Samples were amplified and labeled using a custom automated version of the
NuGEN Ovation WB protocol. Hybridization on custom mouse Affymetrix microarrays (Santa Clara, CA), labeling and scanning using Affymetrix ovens, fluidics stations and scanners were performed according to the protocols recommended (NuGEN, San Carlos, CA). Sample amplification, labeling, and microarray processing were performed by the Covance Genomics Laboratory in Seattle, WA. The raw gene expression data has been deposited into the Gene Expression Omnibus database (http://www.ncbi.nlm.nih.gov/geo/query/acc.cgi?acc=GSE49375).

One-way ANOVA analysis was performed with Matlab (The Mathworks, Natick, MA). Probesets had to pass a pre-filter of Affymetrix MAS5 present call p-value < 0.05 in more than 50% of the samples to qualify for further analysis. Average fold changes between compound and vehicle treatment groups were generated

Figure 7. Immune-regulated genes are up-regulated, while lipid metabolism genes are down-regulated, with sebaceous gland atrophy. Box plots of probesets regulated by the skin-positive DGAT1 inhibitors (those that produce sebaceous gland atrophy) but not by the skin-negative compounds (those that do not produce sebaceous gland atrophy). Plotted are the LogIntensity values across the replicates in each group, and across the three studies. Ccl1 (A; chemokine (C-C motif) ligand 1) is involved in the recruitment of T cells in skin inflammation; and Scd3 (B; stearoyl-coenzyme A desaturase 3) is a sebaceous gland specific gene.

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Table 2. RNA biomarkers for sebaceous gland atrophy in skin. Listed are the 41 unique genes from the 42 probesets identified in the Training Set as shown in Figure 4 (Cxcl16 had 2 probesets).

| Gene Symbol | Entrez ID | Fold Change p | Fold Change | Fold Change p | Fold Change | Fold Change p | Fold Change p | Fold Change p | Fold Change p |
|-------------|-----------|---------------|-------------|---------------|-------------|---------------|---------------|---------------|---------------|
| Ccl1        | 20290     | 19.9          | 10.2        | 9.3           | 6.6         | 1.2           | 1.2           |               |               |
| Cyp2g1      | 13108     | 2.4           | 5.7         | 3.8           | 1.5         | 1.1           | 1.0           |               |               |
| Psca        | 72373     | 1.9           | 2.2         | 1.9           | 1.8         | *             | −1.0          | 1.1           |               |
| Cxcl16      | 66102     | 2.0           | 1.8         | 1.8           | 1.5         | *             | 1.1           | 1.2           |               |
| Defb6       | 116746    | 1.5           | 1.6         | 1.6           | 1.3         | *             | 1.1           | −1.1          |               |
| Cxcl16      | 66102     | 2.1           | 1.7         | 1.6           | 1.2         | *             | 1.1           | 1.2           | *             |
| Tnfsf4      | 22163     | 2.1           | 1.6         | 1.5           | 1.0         |               | 1.1           |               |               |
| Defb1       | 13214     | 1.7           | 1.7         | 1.7           | 1.3         | *             | −1.1          | −1.0          |               |
| Ano1        | 101772    | 1.5           | 1.4         | 1.6           | 1.3         |               | 1.1           | 1.1           | **            |
| Gch1        | 14528     | 1.6           | 1.3         | 1.4           | 1.2         |               | 1.1           | −1.2          |               |
| Sox7        | 20680     | 1.2           | 1.5         | 1.4           | 1.5         | **            | −1.1          | 1.2           |               |
| Pdlim4      | 30794     | 1.3           | 1.5         | 1.3           | 1.1         |               | 1.0           |               |               |
| Cldn12      | 64945     | −1.2          | −1.3        | −1.3          | 1.0         |               | 1.0           |               |               |
| Lrp6        | 16974     | −1.2          | −1.3        | −1.4          | 1.1         |               | 1.0           |               |               |
| Sgpl1       | 20397     | −1.4          | −1.3        | −1.4          | −1.4        | **            | 1.0           |               |               |
| Lamb1       | 16777     | −1.5          | −1.3        | −1.5          | −1.2        |               | 1.0           | −1.1          |               |
| Chst14      | 72136     | −1.6          | −1.4        | −1.8          | −1.6        |               | 1.1           |               |               |
| Elolv5      | 68801     | −1.7          | −1.4        | −1.6          | −1.5        | **            | 1.0           |               |               |
| Crat        | 12908     | −2.2          | −1.4        | −1.9          | −1.3        |               | 1.1           |               |               |
| Mansc1      | 67729     | −2.2          | −1.8        | −1.8          | −1.9        |               | 1.0           |               |               |
| Abhd3       | 106861    | −2.4          | −1.7        | −2.1          | −2.1        |               | −1.1          | −1.0          |               |
| Lass5       | 71949     | −2.4          | −2.1        | −2.2          | −2.2        |               | −1.1          | −1.1          |               |
| Ccdd136     | 232664    | −2.6          | −2.3        | −2.3          | −2.1        |               | −1.1          |               |               |
| Slda19      | 74338     | −3.1          | −1.9        | −2.2          | −2.1        | *             | −1.2          | −1.1          |               |
| Sec1414     | 103655    | −3.3          | −2.4        | −3.0          | −1.8        |               | −1.0          |               |               |
| Hpd1        | 242642    | −4.1          | −2.0        | −2.5          | −2.6        |               | 1.1           |               |               |
| Hsd17b2     | 15486     | −4.2          | −1.7        | −2.6          | −2.7        |               | −1.2          | −1.0          |               |
| Acx2        | 93732     | −3.4          | −2.6        | −3.4          | −3.1        |               | 1.0           | 1.2           |               |
| Dgat2l6     | 668257    | −3.3          | −2.9        | −3.3          | −3.0        |               | −1.0          |               |               |
| Adam3       | 11497     | −5.0          | −2.1        | −3.6          | −3.6        |               | −1.1          | −1.1          |               |
| Cyp1a1      | 13076     | −5.4          | −2.5        | −4.0          | −2.2        |               | 1.2           |               | −1.1          |
| A4gnt       | 333424    | −5.3          | −2.0        | −3.9          | −3.4        |               | −1.1          |               |               |
| Cyp2j2-12   | 242546    | −5.0          | −2.1        | −4.7          | −3.0        |               | −1.1          | 1.2           |               |
| Ces4a       | 234677    | −5.2          | −2.6        | −4.2          | −3.1        |               | 1.0           | 1.1           |               |
| Pnpla5      | 75772     | −5.7          | −2.5        | −3.9          | −3.2        |               | 1.0           | −1.0          |               |
| Ear5        | 54159     | −3.5          | −3.4        | −5.3          | −5.7        |               | −1.1          | 1.0           |               |
| Gabra4      | 14397     | −6.4          | −2.1        | −5.6          | −3.8        |               | −1.2          | −1.0          |               |
| Gm13177      | 435815    | −6.2          | −4.1        | −7.6          | −4.7        |               | 1.0           |               |               |
| Aox4        | 71872     | −7.9          | −4.1        | −7.6          | −6.3        |               | −1.2          | −1.0          |               |
| Scd3        | 30049     | −14.1         | −4.1        | −6.2          | −6.0        |               | 1.1           |               | 1.1           |
regulated 13 probesets as follows: 29 probesets from the averaged expression value of the up-regulated and Cpd18, skin-negative treatments). The score was calculated by the treatments in the independent Test Set (Cpd6, skin-positive negative compound treatment and its respective vehicle treatment were removed (as well as unknown RIKEN genes).

Of the 42 probesets identified in the Training Set, 26 were also significantly regulated in the Test Set by the skin-positive compound (Cpd6; 1.2 fold change and ANOVA p<0.01) and not by the skin-negative compound (Cpd18; ANOVA p>0.1).

### Supporting Information

**Figure S1** DGAT1 inhibitors with high lipophilicity induce sebaceous gland atrophy. Shown are hematoxylin and eosin stains of ventral skin biopsies from DIO mice treated with either vehicle (A), Cpd1 (B), or Cpd2 (C) for 14 days at doses indicated. Scoring refers to the histological adverse effect score as described in Table 1. Bar = 50 μm. (PDF)

**Table S1** DGAT1 inhibitors with high lipophilicity induce sebaceous gland atrophy. Shown are the sebaceous gland sizes from dorsal skin biopsies from DIO mice treated with vehicle, Cpd1, Cpd2, or Cpd3 for 14 days. The data are plotted in Figure 2E. (XLSX)

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