PDE3 and PDE4 Isozyme-Selective Inhibitors Are Both Required for Synergistic Activation of Brown Adipose Tissue

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ABSTRACT

Brown adipose tissue (BAT) is a highly thermogenic organ that converts lipids and glucose into heat. Many of the metabolic and gene transcriptional hallmarks of BAT activation, namely increased lipolysis, uncoupling protein-1 (UCP1) mRNA, and glucose uptake, are regulated by the adrenergic second messenger, cAMP. Cyclic nucleotide phosphodiesterases (PDEs) catalyze the breakdown of cAMP, thereby regulating the magnitude and duration of this signaling molecule. In the absence of adrenergic stimulus, we found that it required a combination of a PDE3 and a PDE4 inhibitor to fully induce UCP1 mRNA and lipolysis in brown adipocytes, whereas neither PDE inhibitor alone had any substantial effect under basal conditions. Under submaximal β-adrenoceptor stimulation of brown adipocytes, a PDE3 inhibitor alone could potentiate induction of UCP1 mRNA, whereas a PDE4 inhibitor alone could augment lipolysis, indicating differential roles for each of these two PDEs. Neither induction of UCP1 nor lipolysis was altered by inhibition of PDE1, PDE2, or PDE8A. However, a revival of interest in brown adipose tissue types, including brown adipocytes but not in white adipocytes (Taylor et al., 1976; Marette and Bukowiecki, 1989; Virtanen et al., 2009). Previously, it was shown that animals receiving a high-fat diet have increased expression of uncoupling protein-1 (UCP1) and increased thermogenesis because of higher sympathetic tone (Rothwell and Stock, 1979). It also is known that animals lacking functional BAT, because of ablation of UCP1, gain more weight than their wild-type counterparts under thermo-neutral conditions (Feldmann et al., 2009). In rodent models of obesity, administration of β3-adrenoceptor agonists not only leads to enhanced lipolysis in white adipocytes (Murphy et al., 1993), but also enhanced insulin sensitivity, glucose use, and triglyceride clearance (Yoshida et al., 1994; Liu et al., 1998; Bartelt et al., 2011). These latter effects are partially mediated by an increase in thermogenesis, whereby circulating fatty acids mobilized from white fat stores are converted into heat by activated brown fat (Weyer et al., 1999). Recently, increased fatty acid oxidation by brown fat has been shown in adult human males exposed to cold (Ouellet et al., 2012). It is this glucose and fatty acid clearing characteristic of brown fat that makes it such an attractive potential target for drug development to treat obesity-related diseases, such as diabetes.

Differentiated brown adipocytes are activated by norepinephrine, which increases cAMP production and stimulates cAMP-dependent protein kinase (PKA) (Janssens et al., 2008). Activation of PKA in turn leads to breakdown of triglyceride stores and increased expression and activation of UCP1 to generate heat (Fedorenko et al., 2012). Increased cAMP also results in increased glucose uptake and metabolism in several tissue types, including brown adipocytes but not in white adipocytes (Taylor et al., 1976; Marette and Bukowiecki, 1989; Jensen, 2007). However, to date, an effective pharmacological strategy for combating obesity that selectively targets the cAMP-signaling pathway is lacking.

Introduction

Brown fat function and its relation to obesity and obesity-related diseases were widely studied in rodents throughout the 1980s and 1990s. However, a revival of interest in brown adipose tissue (BAT) has occurred because of the discovery that adult humans possess functionally active depots of BAT (Cypess et al., 2009; van Marken Lichtenbelt et al., 2009; Virtanen et al., 2009). Previously, it was shown that animals receiving a high-fat diet have increased expression of uncoupling protein-1 (UCP1) and increased thermogenesis because of higher sympathetic tone (Rothwell and Stock, 1979). It also is known that animals lacking functional BAT, because of ablation of UCP1, gain more weight than their wild-type counterparts under thermo-neutral conditions (Feldmann et al., 2009). In rodent models of obesity, administration of β3-adrenoceptor agonists not only leads to enhanced lipolysis in white adipocytes (Murphy et al., 1993), but also enhanced insulin sensitivity, glucose use, and triglyceride clearance (Yoshida et al., 1994; Liu et al., 1998; Bartelt et al., 2011). These latter effects are partially mediated by an increase in thermogenesis, whereby circulating fatty acids mobilized from white fat stores are converted into heat by activated brown fat (Weyer et al., 1999). Recently, increased fatty acid oxidation by brown fat has been shown in adult human males exposed to cold (Ouellet et al., 2012). It is this glucose and fatty acid clearing characteristic of brown fat that makes it such an attractive potential target for drug development to treat obesity-related diseases, such as diabetes.

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ABBREVIATIONS: BAT, brown adipose tissue; BAY 60-7550, 2-(3,4-dimethoxybenzyl)-7-[(1R)-1-[(1R)-1-hydroxyethyl]-4-phenylbutyl]-5-methyl-limidazo[5,1-f][1,2,4]triazin-4(3H)-one; BSA, bovine serum albumin; CREB, cAMP response element-binding protein; DMEM, Dulbecco’s modified Eagle’s medium; Epac, exchange protein activated by cAMP; Fdg, fluordeoxyglucose; IbmX, isobutyl-1-methylxanthine; Pcr, polymerase chain reaction; Pde, phosphodiesterase; Pka, protein kinase; Sch51866, (+)-cis-5,6a,7,8,9a-hexahydro-2-[4-[3rifluoro-methyl] phenylmethyl]-5-methyl-cyclopent[4,5]imidazo [2,1-b]purin-4(3H)one; SUV, standardized uptake value; TBSt, Tris-buffered saline containing 0.05% Tween-20; Ucp1, uncoupling protein-1.

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system, particularly via the β3-adrenoceptor, in human brown adipose tissue has not been been developed because of low expression of this receptor in humans (Barbe et al., 1996; Deng et al., 1996; Larsen et al., 2002; Ursino et al., 2009). Moreover, pharmacological strategies that avoid β-receptor activation will likely be required to avoid cardiovascular and other β-receptor–mediated adverse effects. This is of particular importance, because it has been speculated by several groups that any therapy aimed at increasing the amount of brown fat tissue by differentiation of adipocyte precursors into mature brown adipocytes may not be effective without also providing a means of increasing the activity of the new brown fat tissue (Sell et al., 2004; Gesta et al., 2007; Cannon and Nedergaard, 2009). It is therefore imperative to better understand the mechanisms that regulate the acute activation phases of heat production in vivo.

Although the cAMP dependency of brown fat metabolism has been extensively described, the field has only begun to address the roles played by the negative regulators of cyclic nucleotide signaling, the cyclic nucleotide phosphodiesterases (PDEs), that suppress or modulate each of the thermogenic processes described above. Previous studies have identified the expression of PDE2, PDE3, and PDE4 in rat BAT (Coudray et al., 1999), and at least one study suggested the presence of a PDE1 activity in differentiated brown adipocytes from NMRI mice (Bronnikov et al., 1999). However, which distinct PDE subtype regulates each of the various cAMP-dependent processes of brown fat activation has yet to be determined. Selective inhibitors to most of the PDE families have been developed, and inhibition of PDEs often leads to activation of cAMP-dependent physiologic processes, thereby making PDEs attractive pharmacological targets (Bender and Beavo, 2006).

In this study, we have determined which PDE subtypes are important regulators of three major cAMP-dependent pathways of BAT activation, namely UCPI gene expression, lipolysis, and glucose uptake. We have found that the activity of both PDE3 and PDE4 can regulate basal UCPI expression, lipolysis, and glucose uptake and that combining inhibitors to these PDEs synergistically stimulates each of these processes in BAT. We have also identified PDE8A as a potential regulator of BAT glucose uptake in vivo in conjunction with PDE3 and PDE4. Of most importance, these findings indicate that a single selective PDE inhibitor alone is not sufficient to substantially activate basal BAT function, but rather a combinatorial inhibitor approach may prove to be more effective.

**Materials and Methods**

**Animals.** Wild-type C57Bl/6 mice were purchased from Taconic (Hudson, NY) or the Jackson Laboratory (Bar Harbor, MA). A description on the generation of the PDE8A knock-out (PDE8A−/−) mouse line is provided elsewhere (Vasta et al., 2006). For the experiments involving PDE8A−/− mice reported here, wild-type littermate mice aged 10–16 weeks were used as controls. The Institutional Animal Care and Use Committee of the University of Washington, in accordance with the National Institutes of Health Guide for Care and Use of Laboratory Animals, approved all procedures.

**Real-Time Polymerase Chain Reaction.** Mouse interscapular brown adipose tissue was excised, weighed, and disrupted using a rotor-stator tissue homogenizer. RNA was isolated using the RNeasy Lipid Tissue Mini kit (Qiagen, Valencia, CA). RNA was isolated from cells in culture by using either the QIAshredder system and RNeasy Mini kit (Qiagen) or the NucleoSpin RNA II kit (Machery-Nagel, Bethlehem, PA). cDNA was then generated from 0.125–0.5 μg RNA with use of SuperScript III reverse transcriptase (Invitrogen, Grand Island, NY) and oligo-dT primers modified with a 5'-phosphate. Relative gene expression was determined by performing real-time polymerase chain reaction (PCR) on an MX3000P quantitative PCR system (Stratagene, La Jolla, CA) and analyzed using Mx-Pro software. Real-time PCR was run with iQ SYBR Supermix (Bio-Rad Laboratories, Hercules, CA) with the following thermoprofile: 95°C for 15 seconds, followed by 60°C for 45 seconds for 40 cycles. Primer (IDT, Coralville, IA) sequences for the different PDE isoforms are provided in the supplement (Supplemental Table 1). All primer sets were determined to anneal on different exons.

**Culturing and Differentiation of Immortalized Brown Adipocyte Precursors.** The immortalized brown adipocyte precursor cell line was a generous gift from Dr. Bruce Spiegelman's laboratory (Dana Farber Cancer Institute, Boston, MA). The precursor isolation, immortalization procedure, and growth conditions have been described previously (Uldry et al., 2006). In brief, cells were routinely seeded at 25,000 cells/cm2 (day 0) in culture medium, composed of high-glucose Dulbecco's modified Eagle's medium (DMEM) (HyClone, Thermo Scientific, Waltham, MA) supplemented with 10% fetal bovine serum (HyClone, Thermo Scientific) and 50 U/50 μg penicillin/streptomycin (Gibco, Life Technologies, Grand Island, NY). The following day (day 1), the media was changed to induction medium, which was culture medium supplemented with 20 nM insulin, 1 mM triiodo-thyronine (Sigma-Aldrich, St. Louis, MO), 0.5 mM dexamethasone (Sigma-Aldrich), 0.5 mM isobutyl-1-methylxanthine (IBMX, Sigma-Aldrich), and 0.125 mM indomethacin (Sigma-Aldrich). On day 3, the media was changed to culture medium supplemented with 20 nM insulin and 1 mM triiodo-thyronine and subsequently changed every day thereafter. By day 6, the cells exhibited a fully differentiated phenotype with a large accumulation of multilocular fat droplets. All experiments were conducted on day 6 unless otherwise stated.

**PDE Activity Assay.** Differentiated brown adipocytes were washed three times with phosphate-buffered saline, then lysed by briefly sonicating in ice cold 50 mM Tris, pH 7.4, 2 mM EDTA, 1 mM dithiothreitol, 1 mM sodium orthovanadate, 200 nM phenylmethylsulfonyl fluoride, and 1× protease inhibitor cocktail III (Calbiochem, EMD Millipore, Darmstadt, Germany). PDE activity was measured as described previously (Hansen and Beavo, 1982; Soderling et al., 1998). In brief, the activity assay was performed in 40 mM MOPS, pH 7.5, 15 mM magnesium acetate, 2 mM EGTA, and 0.2 mg/ml bovine serum albumin (BSA; Sigma-Aldrich) supplemented with ▼104,000 cpm ▼H-cAMP or ▼H-cGMP (PerkinElmer, Waltham, MA) in a final volume of 0.25 ml. The total substrate and inhibitor concentrations are indicated in the figure legends. The reaction time and amount of lysate were maintained so that less than 30% of the substrate was hydrolyzed.

**Preparation of Freshly Isolated Primary Brown Adipocytes.** Primary brown adipocytes were isolated using a modified Rodbell method (Rodbell, 1964; Matthias et al., 2000). Routinely, the interscapular fat pads from 3–5 C57Bl/6 mice were pooled and initially placed in a 20-ml plastic scintillation vial (Research Products International, Mount Prospect, IL) containing filter-sterilized Krebs-Ringer HEPES Isolation Buffer (20 mM HEPES, 118.5 mM NaCl, 25.3 mM NaHCO3, 1.2 mM NaH2PO4, 1.2 mM MgSO4, 10 mM D-glucose, 10 mM D-fructose, 4% bovine serum albumin, pH 7.4) warmed to 37°C and equilibrated with 95% O2–5% CO2 bubbling through a diffuser. Whole tissue was pre-incubated in 3 ml Isolation Buffer supplemented with 2.0 mg/ml Collagenase (Sigma-Aldrich; C6886) shaking in a Dubnoff shaking water bath (Thermo Scientific) for 7 minutes shaking at 90 cycles/min. Every 2 minutes, the tissue was removed and gently vortexed for 10 seconds. At the end of the 7 minutes, the tissue was removed and rinsed with Isolation Buffer. The tissue was then finely minced with surgical blades (Bard Parker, Thermo Scientific; no. 371110) in a scissors fashion and placed back into
Brown adipocyte precursors were grown and differentiated from the total. The presence of 1 M Free Glycerol Determination Kit (Sigma-Aldrich). Background lipolysis was measured using the reagents listed in the centrifuged at 16,000 g. The reaction was terminated by freezing and placed in heating blocks in a Dubnoff shaking water bath at 37°C and shaken at 90 cycles/min. The reaction was triggered by freezing the tubes in liquid nitrogen. Tubes were thawed in ice water and centrifuged at 16,000 g at 4°C. The infranatant was removed, and the glycerol content was measured using the reagents listed in the Free Glycerol Determination Kit (Sigma-Aldrich). Background lipolysis in a given experiment was defined as glycerol released in the presence of 1 μM propranolol (Sigma-Aldrich) and was subtracted from the total.

**Glycerol Release Assay.** Freshly isolated brown adipocytes were diluted to a final concentration of 75,000 cells/ml into 200 μl FA-Free Isolation Buffer containing pharmacological agents. The tubes were placed in heating blocks in a Dubnoff shaking water bath at 37°C and shaken at 90 cycles/min. The reaction was terminated by freezing the tubes in liquid nitrogen. Tubes were thawed in ice water and centrifuged at 16,000 g at 4°C. The infranatant was removed, and the glycerol content was measured using the reagents listed in the Free Glycerol Determination Kit (Sigma-Aldrich). Background lipolysis was measured using the reagents listed in the centrifuged at 16,000 g. The reaction was terminated by freezing and placed in heating blocks in a Dubnoff shaking water bath at 37°C and shaken at 90 cycles/min. The reaction was triggered by freezing the tubes in liquid nitrogen. Tubes were thawed in ice water and centrifuged at 16,000 g at 4°C. The infranatant was removed, and the glycerol content was measured using the reagents listed in the Free Glycerol Determination Kit (Sigma-Aldrich). Background lipolysis in a given experiment was defined as glycerol released in the presence of 1 μM propranolol (Sigma-Aldrich) and was subtracted from the total.

**Determination of cAMP Concentration by Enzyme Immunoassay.** Brown adipocyte precursors were grown and differentiated in 12-well plates and preincubated with PDE inhibitors in DMEM for 30 minutes, then isoproterenol in DMEM for 5 minutes. The media were removed, and the reaction was stopped with ice cold 99% ethanol-1% hydrochloric acid. The ethanol acid containing the cAMP was pipetted into Eppendorf tubes and dried in a speed vacuum (Savant, Thermo Scientific). The samples were then resuspended to 200 μl, and cAMP was measured according to the manual for the cAMP EIA Kit (American Qualex Molecular, San Clemente, CA). The cell monolayer was then resuspended in water, and protein content was assessed by BCA assay (Pierce, Rockford, IL).

**Western Blotting.** Differentiated brown adipocytes were lysed in 50 mM Tris, pH 8.0, 0.5% Triton-X100, 150 mM NaCl, 1 mM EDTA, 1 mM dithiothreitol, 1 mM sodium orthovanadate, 1 mM sodium fluoride, 200 nM phenylmethylsulfonyl fluoride, and 1× protease inhibitor cocktail III (Calbiochem, EMD Millipore) and denatured in sample buffer (125 mM Tris, 20% glycerol, 2% SDS, 2% mercaptoethanol) by boiling for 5 minutes. The samples were then separated by 10% SDS-PAGE at 0.75 mm thickness and transferred to polyvinylidene difluoride membranes for 1 hour at 60 V. Membranes were blocked overnight by shaking on an orbital shaker at 4°C in 5% BSA in Tris-buffered saline containing 0.05% Tween-20 (TBST). Membranes were probed for phospho-CREB (cAMP response element-binding protein) Ser-133 [1:1000 (v/v); 9198S; Cell Signaling Technology, Danvers, MA) or glyceraldehyde 3-phosphate dehydrogenase [1:20,000 (v/v); Fitzgerald Industries, Acton, MA) overnight with shaking on an orbital shaker at 4°C in 5% BSA in TBST. Membranes were then washed three times for 10 minutes with TBST at room temperature, rinsing with TBST between washes, then probed with secondary antibody conjugated to horseradish peroxidase [goat anti-rabbit 1:5000 (v/v), goat anti-mouse 1:20,000 (v/v); Bio-Rad Laboratories] in 5% BSA in TBST at room temperature for 1 hour. The washing steps were repeated as before, and membranes were developed with SuperSignal West Pico Chemiluminescent Substrate (Thermo Scientific). Immunoreactivity was imaged using a Bio-Rad Molecular Imager ChemiDoc XRS+ with Image Laboratory Software, and band intensity was quantified using ImageJ. For the detection of PDEs in whole mouse BAT, tissue was harvested from wild-type and PDE6A−/− mice and homogenized as described elsewhere (Nolan et al., 2004). The homogenates were centrifuged at 10,000 g for 15 minutes, and the protein content of the infranatant was determined. Samples were denatured as described above. Thirty micrograms protein was separated by SDS-PAGE and transferred to polyvinylidene difluoride for 3 hours at 60 V, and membranes were blocked in 5% milk in TBST for 1 hour at room temperature. Membranes were probed for PDE6A [1:1200 (v/v); PDE8-121AP, FabGennix International, Frisco, TX] or UCP1 [1:1000 (v/v); C-17, Santa Cruz Biotechnology] in 5% milk in TBST overnight at 4°C. Membranes were washed three times for 10 minutes with TBST and then probed with secondary antibody conjugated to horseradish peroxidase [goat anti-rabbit 1:3000 (v/v) (Bio-Rad Laboratories), or rabbit anti-goat 1:3000 (Jackson Immunoresearch Laboratories, West Grove, PA) for 1 hour at room temperature. The membranes were washed and developed as described above. Immunoreactivity was imaged using autoradiography film (Geneseel Scientific, San Diego, CA).

**18F-Flurodeoxyglucose and PET Scanning.** The mice were fasted overnight before the day of imaging. On the day of imaging, the mice were housed in portable containers that were warmed by placing them on heating pads (Deltaphase Isothermal Pad; Braintree Scientific, Braintree, MA). Thirty minutes before administration of flurodeoxyglucose (FDG), mice received an intraperitoneal injection of a single PDE inhibitor, a combination of PDE inhibitors, or vehicle control. The injected mouse was then placed in a heated chamber. The chamber was warmed using heating pads and maintained at a temperature of 30–35°C. The mouse was awake in the chamber and allowed to move freely. After 30 minutes, the mouse was administered 200–300 μCi of FDG via retro-orbital injection. The amount of FDG administered (in microcuries) was approximately 10 times the weight of the mouse in grams. The mouse was placed in a warmed induction chamber and kept lightly sedated using 1–2% isoflurane anesthesia. After a 40-minute uptake period, the mouse was placed on the small animal PET scanner (Inveon Dedicated PET system; Siemens, Munich, Germany) and imaged for 20 minutes. During the PET imaging, the mouse was kept warm by a small heating pad on the table and maintained while anesthetized with 2–3% isoflurane. A transmission scan was also collected for attenuation correction. After imaging, the mouse was euthanized and the tissue was harvested. The data were reconstructed using software provided by the vendor. The three-dimensional ordered subsets expectation maximization–maximum a posteriori (3D OSEM-MAP) method was used to reconstruct the images. A smoothing parameter to achieve 1.5 mm image resolution was used. The images were analyzed using Siemens’ ASIPRO analysis software. The system was calibrated to report image voxel values in radioactivity concentration (i.e., nCi/g). Regions of interest were drawn encompassing the two BAT regions in a coronal plane of the mouse, as shown in Supplemental Fig. 7. The maximum pixel values from the two regions of interest were averaged and used to determine the standardized uptake value (SUV) for FDG. The SUV is a semiquantitative metric used to express glucose uptake in tissue. It is calculated using the following formula:

\[
SUV = \frac{c(t)}{\text{injected activity/body weight}}
\]

where \(c(t)\) was defined as the maximum activity concentration. Injected activity was corrected to the start of scanning. Body weight of the mouse was measured in grams. When using SUV as an uptake metric, it was important that all imaging be done at the same time after injection—in this case, 40 minutes.

**Drugs.** SCH51866 [ (+)-cis-5,6a,7,8,9,9a-hexahydro-2-[(trifluoromethyl)phenylmethyl]-5-methyl-cylopen(t)4,5imidaizo [2,1-b]purin-4(3H)one] (Schering-Plough/Merck, Whitehouse Station, NJ) is a
selective PDE1 inhibitor that does not distinguish between PDE1 isoforms and was a gift from the manufacturer (structure published in Vemulapalli et al., 1996). BAY 60-7550 [2-(3,4-dimethoxybenzyl)-7-[(1R)-1-[1(R)-1-hydroxyethyl]-4-phenylbutyl]-5-methylimidazo[1,2,4]triazin-4(3H)-one] (Cayman Chemicals, Ann Arbor, MI) is a PDE2 inhibitor. Cilostamide (Tocris, Bristol, UK) is a nonselective PDE3 inhibitor. Rolipram (Enzo Life Sciences, New York, NY) is a selective PDE4 inhibitor that will inhibit PDE5A and PDE5B with similar affinities and was a gift from the manufacturer (structure published in Vang et al., 2010). IBMX (Sigma-Aldrich) is a nonselective PDE inhibitor. H-89 (Tocris) is a protein kinase inhibitor with substantial selectivity for PKA. PDE and PKA inhibitors were dissolved in dimethyl-sulfoxide (Sigma-Aldrich) and stored at −20°C. 8-Bromoadenosine-3′,5′-cyclic monophosphorothioate (Rp-8-Br-cAMPS; BioLog, Bremen, Germany) is a competitive inhibitor to the cAMP-binding site on the regulatory subunit of PKA and was dissolved in water. 8-(4-Chlorophenylthio)-2′-O-methyladenosine-3′,5′-cyclic monophosphate (8-pCPT-2-O-Me-cAMP; BioLog) is an exchange protein activated by cAMP (Epac) agonist and was dissolved in water. 8-Bromo-guanosine 3′,5′-cyclic monophosphate (8-Br-cGMP) is a protein kinase-G agonist and was dissolved in water. 10 mM isoproterenol (Sigma-Aldrich) was dissolved in 1 M ascorbic acid (8-Br-cGMP) is a protein kinase-G agonist and was dissolved in water. O-methyladenosine-3′,5′-cyclic monophosphate (8-Br-cGMP) is a protein kinase-G agonist and was dissolved in water. 10 mM isoproterenol (Sigma-Aldrich) was dissolved in 1 M ascorbic acid immediately before use in gene induction experiments. For lipolysis experiments, isoproterenol was dissolved in water immediately before use. For glucose uptake experiments, isoproterenol was dissolved in phosphate-buffered saline immediately before injection into mice.

**Results**

**PDE Expression in Mouse Brown Adipocyte Models.** We first assessed the PDE mRNA expression profile in mouse interscapular brown fat tissue, freshly isolated primary adipocytes, and brown adipocytes differentiated from a brown adipocyte precursor cell line in vitro. Quantitative real-time PCR analysis revealed that the mRNAs for PDE1A, 2A, 3B, 4B, 4D, and 8A were the predominant ones expressed in whole brown fat tissue (Fig. 1A), primary brown adipocytes isolated from BAT tissue via collagenase digestion (Fig. 1B), and immortalized brown adipocyte precursors on day 6 after differentiation (Fig. 1C). Because it was thought that PDE8A expression is limited to a small number of tissues (Soderling et al., 1998), expression of PDE8A in mouse BAT was unexpected. We further verified the presence of PDE8A protein by Western blot analysis (Supplemental Fig. 1) and also found that both PDE8A RNA and protein was absent in the BAT of PDE8A/−/− mice. (Fig. 1A; Supplemental Fig. 1). In addition, we did not detect any major compensatory changes in the expression of other PDEs in the BAT from PDE8A/−/− mice, compared with wild-type littermates (Fig. 1A). These data indicate that the pattern of PDE expression from whole BAT tissue is largely representative of that found in isolated brown adipocytes and that these patterns are similar among all models of BAT used.

Enzyme activity assays confirmed the functional presence of the major PDEs identified by the mRNA profile in immortalized brown adipocytes. The PDE1 family is stimulated by calcium-calmodulin, and PDE1A has an approximately 70-fold higher selectivity for cGMP, compared with cAMP, at low substrate levels. We therefore confirmed the presence of PDE1 activity by measuring cGMP hydrolysis in the presence and absence of calcium and calmodulin in the whole BAT extract. This hydrolytic activity was increased by 2.7-fold by calcium and was fully inhibited by 100 nM SCH51866, a relatively selective PDE1 inhibitor (IC_{50} for PDE1A, 10 nM) (Dunkern and Hatzelmann, 2007) (Fig. 2A).

PDE2 is a dual-substrate PDE, because it hydrolyzes cAMP and cGMP with approximately equal specificity. However, its cAMP-hydrolyzing activity is stimulated by cGMP. Therefore, to assay for PDE2 activity, we measured the hydrolysis of cAMP in the presence and absence of 200 nM cGMP. We included 10 µM cilostamide and 10 µM rolipram to these reactions to eliminate background PDE3 and PDE4 activity and to eliminate potential complications in the interpretation of the data, because PDE3s can be inhibited by cGMP. Under these conditions, the addition of 1 µM cGMP increased hydrolytic activity toward cAMP by 2.8-fold. This increased activity was completely inhibited by the addition of 50 nM of the PDE2 inhibitor, BAY 60-5770, a 10-fold excess over the reported IC_{50} for PDE2 (Bender and Beavo, 2006) (Fig. 2B).

PDE3 and PDE4 activities were detected by measuring the hydrolysis of 1 µM cAMP in the presence and absence of 200 nM cilostamide (PDE3 inhibitor) or 10 µM rolipram (PDE4 inhibitor), respectively. These inhibitor concentrations are 10-fold higher than their reported IC_{50} values for PDE3 and
For these studies, an immortalized brown adipocyte precursor cell line was differentiated into mature brown fat cells in vitro. The differentiated adipocytes were then pretreated with selective inhibitors to PDE1 (SCH51866), PDE2 (BAY 60-7550), PDE3 (cilostamide), PDE4 (rolipram), or PDE8 (PF-04957235), either individually or in combination. Somewhat unexpectedly, when administered individually, none of the PDE inhibitors increased the basal expression of UCP1 mRNA. However, when administered together, 10 μM cilostamide and 10 μM rolipram resulted in an approximately 40-fold increase in basal UCP1 mRNA expression, similar to that seen with a high dose of isoproterenol (10 μM) (Supplemental Fig. 2A) or IBMX (200 μM) (Fig. 3A). This latter observation strongly suggests that inhibition of PDE3 and PDE4 together is sufficient to explain the stimulatory effect of the nonselective PDE inhibitor, IBMX, on induction of UCP1. The same magnitude of potentiation was observed when the concentration of cilostamide was reduced to 300 nM (Fig. 3B). The PDE8-selective inhibitor, PF-04957325 (200 nM), did not stimulate UCP1 mRNA expression when administered either alone or in combination with IBMX, cilostamide, or rolipram (Fig. 3C; Supplemental Fig. 2B). Similarly, PDE1 or PDE2 inhibitors administered either alone or in combination with PDE3 and PDE4 inhibitors did not stimulate expression of UCP1 mRNA, nor did they augment the effect of combined PDE3 and PDE4 inhibition (Supplemental Fig. 2C). In addition, we detected a 2.7-fold increase of UCP1 mRNA in the interscapular brown fat of fasted, warmed mice that were injected with the combination of cilostamide and rolipram (P < 0.05), where an injection of either drug alone had no significant effect, compared with an injection of vehicle (Fig. 3D). A similar pattern of stimulation was also observed in immortalized brown adipocytes for basal expression of peroxisome proliferator-activated receptor-γ coactivator mRNA (Supplemental Fig. 3), a transcription factor also regulated by cAMP that is important for UCP1 gene induction during BAT activation. Finally, PDE3 and PDE4 inhibitors administered individually did not stimulate basal cAMP accumulation, but when administered together, they increased cAMP by 9.6-fold (Fig. 4). Together, these results suggest that the combined inhibition of PDE3 and PDE4 removes an important physiologic suppressive effect of these two PDEs on basal cAMP signaling and BAT activation in the absence of β-adrenoceptor agonists.

**PDE3B Controls β-Adrenoceptor Stimulation of UCP1 mRNA Expression.** Our finding that both PDE3 and PDE4 together control basal UCP1 expression prompted us to determine whether inhibition of the same combination of PDEs regulates the increase in induction of UCP1 mRNA caused by β-adrenoceptor agonists. Typically, inhibition of the PDE or PDEs that regulate a given process will shift the dose-response curve of agonists for that process to the left or up and to the left. Because β-adrenoceptor activation is known to stimulate expression of UCP1 mRNA, we hypothesized that a PDE inhibitor or combination of inhibitors would potentiate a low dose of isoproterenol based on the dose-response relationship (Supplemental Fig. 2A). For these experiments, cells were pretreated with individual inhibitors to PDE1, PDE2, PDE3, PDE4, and PDE8, followed by 1 nM isoproterenol. We found that only pretreatment with the PDE3 inhibitor, cilostamide, dose-dependently potentiated the expression of UCP1 mRNA in response to this low dose of isoproterenol (P < 0.001 at 10 μM cilostamide). In contrast, a PDE4 inhibitor, rolipram, had no effect on isoproterenol-induced UCP1 expression (Fig. 3A). Of
interest, cAMP accumulation did not follow the same pattern of potentiation (Fig. 4). At 1 nM isoproterenol, neither cilostamide nor rolipram potentiated global cAMP accumulation, even though UCPI mRNA was substantially increased under these conditions. PDE1, PDE2, and PDE8 inhibitors also did not potentiate induction of UCPI by isoproterenol (Fig. 3C; Supplemental Fig. 4).

Effects of PDE3 and PDE4 Inhibition Are PKA Dependent. Two major pathways for cAMP-mediated mechanisms for control of transcription have been described: the Epac pathway and the cAMP-dependent PKA pathway. We therefore wanted to test by which of these molecular pathway(s) PDE inhibition caused stimulation of UCPI mRNA expression. We pretreated differentiated brown adipocytes with PDE3 and PDE4 inhibitors along with increasing doses of H-89, a known PKA inhibitor. Cilostamide and rolipram alone had minor effects on PKA-substrate phosphorylation and CREB H-89, a known PKA inhibitor. Cilostamide and rolipram alone

![Graph A](image1)

**Fig. 3.** PDE3 and PDE4 inhibitors increase UCPI and Fgc-1α mRNA expression in differentiated brown adipocytes. UCPI mRNA expression was measured in differentiated brown adipocytes that were (A) pretreated with 10 μM cilostamide, 10 μM rolipram, both, or 200 μM IBMX for 30 minutes, then stimulated by vehicle or 1 nM isoproterenol for 4 hours (n = 3–13); (B) treated with 300 nM cilostamide or 10 μM cilostamide with or without 10 μM rolipram (n = 3–8); or (C) treated with either 200 nM PF-04957325, 200 μM IBMX, or both for 30 minutes, then stimulated by vehicle or 1 nM isoproterenol for 4 hours (n = 3). UCPI mRNA was quantified relative to 18S mRNA with use of RT-PCR as described in Materials and Methods. (D) UCPI mRNA expression was measured from the RNA extracted from the interscapular BAT pads harvested from mice undergoing PET scanning after the procedure was terminated (Materials and Methods). UCPI mRNA was quantified relative to GAPDH (glyceraldehyde 3-phosphate dehydrogenase) mRNA with use of real-time (RT)-PCR as described in Materials and Methods. Data are presented as mean fold over vehicle control ± S.E.M., and statistical analyses were performed using one-way analysis of variance with Dunnet post hoc test: ***P < 0.001; *P < 0.05, versus vehicle in each group.

![Graph B](image2)

![Graph C](image3)

![Graph D](image4)

A selective agonist for Epac, 8-pCPT-2-O-Me-cAMP, did not stimulate UCPI mRNA (unpublished data). Because it has been shown that H-89 can inhibit protein kinase-G (K_i = 0.5 μM) with a K_i that is only 10-fold higher than the K_i for PKA (K_i = 0.05 μM) in vitro (Hidaka and Kobayashi, 1992), we also tested whether 8-Br-cGMP, an agonist to cGMP-dependent protein kinase, could stimulate accumulation of UCPI mRNA; we observed no appreciable change after application of this inhibitor (unpublished data). Together,
these data strongly suggest that the stimulatory effect of combined PDE3 and PDE4 inhibition on UCP1 mRNA accumulation is largely through the canonical PKA-dependent signaling pathway.

**PDE3 and PDE4 in Combination Regulate Lipolysis in Primary Brown Adipocytes.** Noradrenergic regulation of BAT function also requires stimulation of lipolysis to provide both the energy for establishing the mitochondrial proton gradient and the allosteric activation of UCP1 by free fatty acids. Therefore, we were interested in determining which PDEs were important for cAMP-dependent regulation of lipolysis in brown fat. Freshly isolated primary brown adipocytes were pretreated with 10 μM cilostamide, 10 μM rolipram, or both and then treated with vehicle or isoproterenol for 1 hour. Lipolysis was measured as glycerol released into the media. We found that cilostamide alone did not significantly alter basal lipolysis or lipolysis stimulated by 10 nM isoproterenol. Conversely, rolipram alone significantly potentiated both basal and isoproterenol-stimulated glycerol release by 7.8-fold and 2.8-fold, respectively, over the vehicle control (Fig. 6A). This suggested that PDE4, but not PDE3, was a primary regulator of the pool of cAMP that regulates lipolysis in primary brown adipocytes. However, when both inhibitors were combined, there was an even larger 30-fold increase in lipolysis that is of the same magnitude as that seen with the nonselective PDE inhibitor IBMX (Fig. 6A). Of interest, despite substantial expression of PDE8A in brown fat, the PDE8 inhibitor did not augment lipolysis in primary brown adipocytes, either alone or in combination with IBMX, cilostamide, and/or rolipram (Fig. 6B; Supplemental Fig. 6, A–C).

These data indicate synergistic roles for PDE3 and PDE4, but not for PDE8A, on lipolysis in brown adipocytes.

**Inhibition of PDE3 and PDE4 Stimulate In Vivo Glucose Uptake in BAT.** Glucose uptake in brown adipocytes can be regulated by cAMP-dependent signaling (Marette and Bukowiecki, 1989). We therefore hypothesized that selective inhibition of the appropriate PDE(s) would potentiate glucose uptake in BAT. To test this, we treated fasted, warmed, wild-type, and PDE8A−/− littersmates with 3 mg/kg cilostamide, 3 mg/kg rolipram, a combination of both, or the vehicle control in each group; #P < 0.05; ###P < 0.001, versus vehicle control in each group; *P < 0.05; **P < 0.001, versus 1 nM isoproterenol plus 10 μM cilostamide without H-89.

**Fig. 5.** The potentiating effect of PDE inhibitors on Ucp1 mRNA and CREB phosphorylation is PKA-dependent. (A) UCP1 mRNA expression was measured in differentiated brown adipocytes that were pretreated with H-89 for 1 hour, then 10 μM cilostamide and 10 μM rolipram for 4.5 hours. RNA was isolated, and UCP1 mRNA was quantified relative to 18S mRNA with use of real-time (RT)-PCR as described in Materials and Methods. Values were then normalized to the cilostamide and rolipram treatment in the absence of H-89 control. (B) CREB phosphorylation after 40 minutes of stimulation by 10 μM cilostamide and 10 μM rolipram in differentiated brown adipocytes. (C) Quantification of Western blot in (B). (D) UCP1 mRNA expression in differentiated brown adipocytes that were pretreated with H-89 at the indicated doses for 1 hour, followed by PDE inhibitors added 30 minutes before addition of 1 nM isoproterenol. RNA was isolated, and UCP1 mRNA was quantified relative to 18S mRNA with use of RT-PCR as described in Materials and Methods. Values were then normalized to the cilostamide and isoproterenol treatment in the absence of H-89 control. (E) CREB phosphorylation after 10 minutes of isoproterenol stimulation after a 30-minute preincubation with 10 μM cilostamide in differentiated brown adipocytes. (F) Quantification of Western blot in (E). Data are presented as mean ± S.E.M. (n = 3–7), and statistical analyses were performed by one-way analysis of variance with Dunnett post hoc: ***P < 0.001, **P < 0.01, versus vehicle control in each group; *P < 0.05; **P < 0.001, versus 1 nM isoproterenol plus 10 μM cilostamide without H-89.
BAT glucose uptake between wild-type and PDE8A−/− mice. Taken together, these data suggest that each of these three PDEs can suppress basal cAMP-dependent glucose uptake in BAT and that all three need to be inhibited to allow full induction of UCP1 mRNA. Because we did not observe changes in global cAMP accumulation at any of the conditions described in the article were designed to test the hypothesis that inhibition of the specific PDEs that regulate BAT activation might provide a conceptual basis for the design of new PDE-related therapies for the treatment of obesity-related diseases.

In this study, we addressed the question of which subtype(s) of PDE regulate lipolysis, glucose uptake, and mRNA expression of UCP1. We identified PDE3 and PDE4 (likely PDE3B and PDE4B/D) as the major regulators of these major BAT processes both in vitro and in vivo (Figs. 2, 3, 6, and 7). Under basal conditions, the effects of the individual PDE3 and PDE4 inhibitors were synergistic in nature, because individual inhibitors to PDE3 and PDE4 had little or no effect on their own but, when combined, would produce a large, synergistic response. In fact, the same magnitude of potentiation on UCP1 expression was observed over a wide range of cilostamide doses, suggesting a truly synergistic relationship between rolipram and cilostamide (Fig. 3B). The only conditions in which a single selective PDE3 or PDE4 inhibitor was sufficient to potentiate UCP1 induction or lipolysis, respectively, were those in which an adrenoceptor agonist was also present (Figs. 3 and 6). Finally, we also observed that PDE8A may play a role in the regulation of basal BAT glucose uptake, but this global knockout effect was only observed when PDE3 and PDE4 were also inhibited (Fig. 7).

We also found that PDE1, PDE2, or PDE8 inhibitors had virtually no effect on UCP1 mRNA either alone or in any combination with either PDE3 or PDE4 inhibitors (Fig. 3C; Supplemental Fig. 2B), despite the fact that we can detect PDE1, PDE2, and PDE8 mRNAs and enzyme activities (Figs. 1 and 2, A and B) in extracts from these differentiated brown adipocytes. Therefore, inhibition of both PDE3 and PDE4 together is necessary to fully activate basal UCP1 induction and lipolysis and is sufficient to fully account for the IBMX response. We observed what appears to be a greater, though not statistically significant, effect of IBMX on cAMP levels, compared with the cilostamide-rolipram combination at 10 nM isoproterenol. This may suggest that PDE1 and PDE2 have some role under high cAMP levels, which is consistent with what has been shown previously in several other tissue types (Bronnikov et al., 1999; Beavo et al., 2007). Therefore, different PDEs can regulate different processes in BAT, but not all PDEs regulate all processes.

We next investigated which phosphodiesterase subtypes regulate adrenoceptor-mediated lipolysis and induction of UCP1 mRNA transcription. We found that PDE3 inhibition potentiated induction of UCP1 mRNA in response to 1 nM isoproterenol, whereas PDE1, PDE2, PDE4, and PDE8 inhibition did not (Fig. 3A; Supplemental Fig. 2C). This observation suggests that PDE3, but not PDE4, regulates a local pool of cAMP that is generated by the stimulation of β-adrenoceptors and leads to full induction of UCP1 mRNA. Because we did not observe changes in global cAMP accumulation at any of the various concentrations of isoproterenol and PDE inhibitors...
The pathway(s) and mechanism(s) by which cAMP stimulates glucose uptake in BAT are controversial. It has been suggested that there may be an indirect response of the cell to the increased ATP demand because of the uncoupling process stimulated by lipolysis in rat brown adipocytes (Marette and Bukowiecki, 1991). Because PDE3 and PDE4 inhibition stimulates lipolysis to the same order of magnitude as IBMX in freshly isolated primary adipocytes, our data suggest that the injection of these isozyme selective inhibitors into mice might synergistically stimulate glucose uptake possibly via free fatty acid stimulation of UCP1. Furthermore, one study indicated that there was an increase in glucose transporters to the surface of primary isolated mouse brown adipocytes stimulated by cAMP-elevating agents (including IBMX) (Omatsu-Kanbe and Kitasato, 1992), and another study indicated that the cAMP effect was largely on the affinity of GLUT-1 receptors, and not on translocation, in differentiated adipocyte precursors isolated from the stromal fraction of mouse BAT (Shimizu et al., 1998). Despite the lack of a potentiating effect of the PDE8-selective inhibitor on either UCP1 mRNA expression or lipolysis (Figs. 3 and 6), we found that glucose uptake in response to PDE3-PDE4 dual inhibition was 1.8-fold higher in BAT from PDE8A−/− mice than BAT from their wild-type littermates (Fig. 7, A and B), whereas isoproterenol had no differential effect between the two genotypes (Fig. 7C). These results suggest that there may be at least two cAMP-dependent pathways that converge on glucose uptake: one mediated by lipolysis and uncoupling via free fatty acid stimulation of UCP1. Furthermee, one study indicated that there was an increase in glucose transporters to the surface of primary isolated mouse brown adipocytes stimulated by cAMP-elevating agents (including IBMX) (Omatsu-Kanbe and Kitasato, 1992), and another study indicated that the cAMP effect was largely on the affinity of GLUT-1 receptors, and not on translocation, in differentiated adipocyte precursors isolated from the stromal fraction of mouse BAT (Shimizu et al., 1998). Despite the lack of a potentiating effect of the PDE8-selective inhibitor on either UCP1 mRNA expression or lipolysis (Figs. 3 and 6), we found that glucose uptake in response to PDE3-PDE4 dual inhibition was 1.8-fold higher in BAT from PDE8A−/− mice than BAT from their wild-type littermates (Fig. 7, A and B), whereas isoproterenol had no differential effect between the two genotypes (Fig. 7C). These results suggest that there may be at least two cAMP-dependent pathways that converge on glucose uptake: one mediated by lipolysis and uncoupling via free fatty acid stimulation of UCP1. Furthermore, one study indicated that there was an increase in glucose transporters to the surface of primary isolated mouse brown adipocytes stimulated by cAMP-elevating agents (including IBMX) (Omatsu-Kanbe and Kitasato, 1992), and another study indicated that the cAMP effect was largely on the affinity of GLUT-1 receptors, and not on translocation, in differentiated adipocyte precursors isolated from the stromal fraction of mouse BAT (Shimizu et al., 1998).
further suggested by the observation that activated human BAT can take up and metabolise circulating fatty acids and glucose (Ouellet et al., 2012) and that obese persons can exhibit reduced thermogenesis and low responsiveness to noradrenaline (Jung et al., 1979). In addition, because human fat tissues of any type express a negligible number of β3-adrenoceptors, selective activation of the tissue with β3-adrenoceptor agonist approaches have failed in clinical trials. Therefore, alternative approaches that bypass receptor-mediated activation may be useful in the future.

A synergistic relationship between PDE3 and PDE4 has been described previously in other tissues, such as rat vascular smooth muscle cells (Palmer et al., 1998) and even in the regulation of lipolysis in white adipocyte models from mice (Snyder et al., 2005). In addition, dual PDE3/PDE4 inhibitors have been tested clinically for the treatment of chronic obstructive pulmonary disease with limited success. However, these drugs were fairly well tolerated in early-phase clinical trials with low incidence of adverse effects that are traditionally associated with effective doses of PDE4 inhibitors alone (Banner and Press, 2009). To our knowledge, the effects of these dual inhibitors on BAT function have not been determined.

In summary, understanding the regulation of the basal state of CAMP/KA in BAT may prove to be critically important in the design of future therapies for disease states. In this regard, it appears that there are multiple BAT PDEs that suppress tissue activation in the absence of agonists that drive CAMP production. Removing this inhibition with the correct combination of PDE inhibitors is sufficient to fully activate these processes. It is now well known that identifying small molecules that target only subsets of PDEs is possible from a medicinal chemistry point of view, and examples of single molecules that selectively inhibit both PDE3 and PDE4 have been described (reviewed in Bender and Beavo, 2006). However, it remains to be seen whether such an approach, perhaps involving a single molecule that inhibits multiple PDEs, might be efficacious in a clinical setting.

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Authorship Contributions

Participated in research design: Kraynik, Miyaoka, Beavo. Conducted experiments: Kraynik, Miyaoka. Performed data analysis: Kraynik, Miyaoka, Beavo. Wrote or contributed to the writing of the manuscript: Kraynik, Miyaoka, Beavo.

References


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