Peroxisome Proliferator-Activated Receptor-γ Promotes Adipogenic Changes in Growth Plate Chondrocytes In Vitro

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Chondrocytes and adipocytes are two differentiated cell types which are both derived from mesenchymal cells. The purpose of this study was to investigate whether peroxisome proliferator-activated receptor-γ (PPARγ), a transcription factor involved in lineage determination during adipogenesis, is able to induce adipogenic differentiation in growth plate chondrocytes. Isolated epiphyseal chondrocytes were infected with a PPARγ adenovirus or treated with the PPARγ agonist ciglitazone. Both of these treatments resulted in lipid droplet accumulation and expression of the adipogenic markers aP2, lipoprotein lipase, and adipsin in chondrocytes. Proteoglycan matrix synthesis was decreased in the PPARγ-infected cells, as was the expression of the chondrogenic genes Col2a1 and aggrecan. Growth plate cells transfected with a PPARγ expression plasmid under the control of the collagen α1(II) promoter also demonstrated a similar adipogenic changes. Terminal differentiation of growth plate chondrocytes induced by thyroid hormone was also inhibited by overexpression of PPARγ and ciglitazone treatment, with decreased expression of alkaline phosphatase and Runx2/Cbfa1 genes. These in vitro data suggest that PPARγ is able to promote adipogenic differentiation in growth plate chondrocytes, while negatively regulating chondrogenic differentiation and terminal differentiation.

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INTRODUCTION

Longitudinal growth of the skeleton is a result of endochondral ossification that occurs at the growth plate [1]. Endochondral ossification is a multistep process that includes differentiation of mesenchymal cells into chondrocytes, cell proliferation, hypertrophic differentiation, matrix mineralization, apoptosis, vascular invasion, and eventually the replacement of the cartilage by bone.

The first step of growth plate development is the commitment of mesenchymal stem cells to the chondrogenic lineage. Mesenchymal stem cells exhibit a high differentiation plasticity. They are capable of differentiating into chondrocytes, osteoblasts, adipocytes, and other tissues of mesenchymal origin [2]. Interconversion between mesenchymal phenotypes is thought to be under control of specific transcription factors, including the Sox family in chondrogenesis [3], Runx2/Cbfa1 in osteogenesis [4], and PPARγ (peroxisome proliferator-activated receptor-γ), and C/EBP (CCAAT/enhancer-binding protein) in adipogenesis [5].

PPARγ is a key transcriptional regulator of adipogenesis [5]. PPARγ is also expressed in preosteoblastic cells and is thought to play a role in regulation of bone metabolism. PPARγ and PPARγ activators inhibit the maturation of preosteoblastic cells to osteoblasts [6–8]. Free fatty acids activate PPARs and induce adipocyte-like differentiation of osteosarcoma cell lines [6]. Lecka-Czernik et al observed that PPARγ2 negatively regulates stromal cell plasticity by suppressing expression Osf2/Cbfa1 and osteoblast-like biosynthetic activity, while promoting differentiation to adipocytes [7]. Conversely, PPARγ insufficiency enhances osteogenesis through increased osteoblast formation from bone marrow progenitors. Homozygous PPARγ-deficient ES cells fail to differentiate into adipocytes, but increase bone mass by stimulating osteoblastogenesis from bone marrow progenitors [8].

Transdifferentiation of chondrocytes to adipocytes has been previously reported by Heermeier et al, who observed that chondrocytes of the mouse xiphoid process undergo transdifferentiation into adipocytes in the presence of 10% fetal calf serum [9].

Based on the finding that PPARγ is expressed in growth plate chondrocytes [10], as well as the evidence that PPARγ is able to compete with the thyroid hormone receptor (TR) for binding to retinoic acid receptor X to inhibit growth plate cell hypertrophy [11], the purpose of this study was to
investigate whether PPARγ and its ligands are able to promote adipogenic differentiation and suppress chondrogenic differentiation in growth plate chondrocytes.

**MATERIALS AND METHODS**

**Cell culture**

Chondrocytes were isolated from the resting zone of the distal femoral growth plate of 2-day old neonatal Sprague-Dawley rats by sequential digestion in trypsin/EDTA (Invitrogen, Carlsbad, Calif) for 1 hour at 37°C, followed by 0.3% collagenase type I (Worthington, Lakewood, NJ) for 4 hours at 37°C [12]. Cells were resuspended in DMEM/F12 medium (Invitrogen) supplemented with a defined media supplement (ITS+1, Sigma, St Louis, Mo) and plated in monolayer at a density of $5 \times 10^5$ cells/cm$^2$, or in a pellet culture of $1 \times 10^3$ cells/mL as indicated [12]. Tri-iodothyronine (T3, Sigma) at a concentration of 100 ng/mL and ciglitazone (BioMol, Plymouth Meeting, Pa) at a concentration of $5 \mu M$ were added to the medium as indicated.

**Immunoblotting**

Total cellular protein was extracted from chondrocytes treated with $5 \mu M$ of ciglitazone using RIPA buffer [9]. An equal amount of protein was subjected to SDS-PAGE, and transferred onto nitrocellulose membranes. The blots were incubated with anti-PPARγ and anti-actin (Santa Cruz Biotechnology, Santa Cruz, Calif) followed by a HRP-conjugated secondary antibody. Immunoreactive proteins were visualized by Western Blotting Chemiluminescence Luminol Reagent (Santa Cruz Biotechnology). Immunoblot bands were quantitated with Kodak 1D Image Analysis Software (Eastman Kodak Company, Rochester, NY).

**Adenovirus infection**

Recombinant adenovirus carrying PPARγ1 (Ad-PPARγ) was kindly provided by Dr J. L. Jameson (Northwestern University Medical School, Chicago, Ill). Ad-PPARγ contains mouse PPARγ1 cDNA driven by the CMV promoter/enhancer with an SV40 polyadenylation sequence [13]. Ad-Gal, which contains β-galactosidase driven by CMV promoter, was used to evaluate the efficiency of gene transduction. Eighteen hours after plating in monolayer, growth plate chondrocytes were infected with adenoviral vectors at a multiplicity of infection (MOI) of 100. Fresh media were added 24 hours after infection and incubated for 72 hours to collect the cell protein extracts. β-galactosidase expression was detected in 80% of cells after 24 hours of infection with Ad-Gal. Expression of introduced PPARγ genes was confirmed by immunoblot.

**Plasmid construction and transient transfection**

The full-length cDNA of mouse PPARγ was excised by Asp718/NheI digestion from pCMX-PPARγ (kindly provided by Dr R. Evans, Salk Institute, La Jolla, Calif). The ends of this fragment were blunted with Klenow polymerase and ligated to a blunt-ended BamHI site in the p1757 plasmid containing the rat a1(II) collagen promoter (kindly provided by Dr Y. Yamada, NIDR, Bethesda, Md) [14]. The cDNA encoding the mouse PPARγ was thus located downstream of the rat a1 (II) collagen promoter element (~977 to +110). Nucleotide sequence analysis confirmed the correct orientation of the PPARγ cDNA.

Growth plate cells were transfected with $10 \mu g$ of p1757-PPARγ or p1757 as a negative control by lipofection (Fugene 6, Roche, Indianapolis, Ind) in the presence of 4 units/mL of hyaluronidase. Sixteen hours later, the cells were trypsinized and centrifuged to pellets cultured in DMEM/F12 plus ITS+ supplements [11].

**Histochemical staining**

For the analysis of adipogenic differentiation, adipogenesis and lipid accumulation in the growth plate cells were examined by staining with Oil Red-O. After 10 days of culture, cells were washed gently with PBS followed by staining with a filtered solution of 0.5% Oil Red-O (Sigma) in 60% isopropanol for 20 minutes. After washing cells with PBS three times, cells were kept in 75% glycerol solution and observed under a phase-contrast microscope.

Alcian blue staining was used to detect chondrocyte-specific proteoglycans at 10 days of culture. Cells were stained with a 4 : 1 ratio of 0.1 M HCl/0.5% Alcian blue stock (0.5% Alcian blue in 95% ethanol) overnight at 37°C in a humidified atmosphere. Cells were then washed twice with PBS to stop reaction and once with 70% ethanol to reduce background.

For alkaline phosphatase (ALP) staining, cultured plates were rinsed with PBS at 10 days of culture, fixed in 3.7% formaldehyde at room temperature for 10 minutes, and stained in the dark for 30 minutes in a 0.1 M Tris-HCl solution (pH 8.5) containing 0.2 mg/mL of Napthol AS-MX phosphate and 0.6 mg/mL of Fast Blue BB salt (Sigma).

**Quantitative real-time RT-PCR**

The expression of chondrocyte or adipocyte-specific RNA markers was analyzed using quantitative real-time RT-PCR. Total RNA was isolated from cultured growth plate chondrocytes using the RNeasy Kit (Qiagen, Valencia, Calif) 4 days after adenovirus infection or plasmid transfection. Reverse transcription was performed using random primers and Superscript III (Invitrogen). Real-time PCR reactions were conducted in an ABI Prism 7700 Sequence Detection System using SYBR Green PCR core reagents (Applied Biosystems, Foster City, Calif). The comparative C$_T$ method ($\Delta$ΔC$_T$ method) was utilized for relative quantitation of gene levels of expression. 18S rRNA was used as an internal control for normalization of target gene expression. The forward and reverse primers for the amplifications are listed in Table 1.
**Statistical analysis**

The data for real-time PCR are represented as mean ± standard deviation. Values are assessed by one-way ANOVA with the Bonferroni post-hoc test and Student t test at a significance level of $P < .05$.

**RESULTS**

**Ciglitazone upregulates PPARγ expression in growth plate chondrocytes**

Treatment of growth plate cells with ciglitazone resulted in increases of both PPARγ mRNA and protein. PPARγ mRNA was increased 9-fold after addition of ciglitazone (5 μM) for 4 days (Figure 1(a)), while PPARγ protein levels were increased approximately 5-fold (Figure 1(b)).

**PPARγ induces adipogenic differentiation in growth plate chondrocytes**

Phase-contrast microscopy demonstrated that the Ad-PPARγ-infected growth plate chondrocytes acquired the morphology characteristic of adipocytes after culture in monolayer for 10 days. Approximately 50% of the cells had accumulated vacuoles, which were positive for Oil Red-O staining of lipid accumulation (Figure 2(a)). The control cells infected with Ad-Gal demonstrated few Oil Red-O positive vacuoles. Cells treated with 5 μM of ciglitazone alone for 10 days also showed enhanced Oil Red-O staining (20% of the total cells), while the combination of ciglitazone treatment and Ad-PPARγ infection further enhanced lipid accumulation (over 70% of cells staining positively with Oil Red-O).

**Table 1:** Primer sequences used for real-time PCR.

<table>
<thead>
<tr>
<th>Genes</th>
<th>Primers</th>
</tr>
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<tbody>
<tr>
<td>aP2</td>
<td>Forward 5′-GGCTTCGCCACCAGGAA-3′</td>
</tr>
<tr>
<td></td>
<td>Reserve 5′-CCCCCTTACGCTGATGATCAAGT-3′</td>
</tr>
<tr>
<td>LPL</td>
<td>Forward 5′-GGTGCGGCTGGTGCAAGT-3′</td>
</tr>
<tr>
<td></td>
<td>Reserve 5′-AAAGTGCCTCACATTGGGATAA-3′</td>
</tr>
<tr>
<td>Adipsin</td>
<td>Forward 5′-CCTATGTGTCGACCAACT-3′</td>
</tr>
<tr>
<td></td>
<td>Reserve 5′-CATGCTAGTGCACAGATGTG-3′</td>
</tr>
<tr>
<td>COL2A1</td>
<td>Forward 5′-GGTGGAAGCAGAAGACAA-3′</td>
</tr>
<tr>
<td></td>
<td>Reserve 5′-GTTCGCGTAAGCTGAGT-3′</td>
</tr>
<tr>
<td>Aggrecan</td>
<td>Forward 5′-CTAGCTGTAGATGAAGGATAACG-3′</td>
</tr>
<tr>
<td></td>
<td>Reserve 5′-CCGGAGAGTCCGAAAAAGC-3′</td>
</tr>
<tr>
<td>ALP</td>
<td>Forward 5′-GGCCGAAGACACACAGACT-3′</td>
</tr>
<tr>
<td></td>
<td>Reserve 5′-GGTGGAGGGTCTGAGGATG-3′</td>
</tr>
<tr>
<td>Runx2/</td>
<td>Forward 5′-TTTAGGCCCCATATC-3′</td>
</tr>
<tr>
<td>Chfα1</td>
<td>Reserve 5′-GGAGGCCGCTGTTGCTT-3′</td>
</tr>
<tr>
<td>18S</td>
<td>Forward 5′-AGTCCCTGCCCTTTGTACACA-3′</td>
</tr>
<tr>
<td></td>
<td>Reserve 5′-GATCCGAGGCTCCTACAAAC-3′</td>
</tr>
</tbody>
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Figure 1: Ciglitazone promotes PPARγ expression in the growth plate chondrocytes. Growth plate chondrocytes in pellet cultures were incubated in the presence or absence of 5 μM of ciglitazone for 4 days. (a) Total RNA was collected and real-time PCR was performed to quantitate PPARγ mRNA levels, which were normalized with respect to endogenous 18S rRNA levels. (b) Proteins were extracted for immunoblotting to detect PPARγ expression and the immunoblots quantitated using Kodak 1D image analysis software. Actin was used as an internal control.
Figure 2: Adipogenic changes in the growth plate cells in which PPARγ was overexpressed or cells treated with ciglitazone. (a) Growth plate cells were infected with Ad-PPARγ or Ad-Gal followed by a 10-day incubation in the presence or absence of ciglitazone (5 μM). Cells were observed under the phase contrast microscope with 10-fold magnification. Oil Red-O staining shows lipid accumulation within the cells. (b) Quantitative real-time PCR analysis of the adipogenic marker genes aP2, LPL, and adipsin, in growth plate cells at day 4 of the culture. Ad-Gal- infected cells were used as controls. Gene expression levels were normalized with respect to endogenous 18S rRNA. * P < .05 versus the expression in control cells.

To characterize the phenotype of the transformed cells in more detail, the cells were cultured in three-dimensional cell pellets and the expression of adipocyte differentiation marker genes examined by real-time RT-PCR at day 4 of the culture period. Compared with the control samples, the levels of expression of the adipogenic marker genes aP2, LPL, and adipsin increased 6.6-, 4.4- and 4.6-folds, respectively, on day 4 in the 5 μM ciglitazone-treated cells (Figure 2(b)). Expression of aP2, LPL, and adipsin genes increased in the Ad-PPARγ-infected cells by 75.1-, 40.2-, and 76.6-folds, respectively in the absence of ciglitazone, and 101.1-, 44.9-, and 96.3-folds, respectively, in the presence of ciglitazone (5 μM).

In order to address the possibility that PPARγ was acting on an undifferentiated mesenchymal stem cell as opposed to a differentiated chondrocyte, p1757-PPARγ expression plasmid was generated in which a PPARγ cDNA was placed under the transcriptional control of the rat COL2A1 gene promoter and enhancer sequences. Oil Red-O staining of the p1757-PPARγ-transfected cells maintained in three-dimensional pellet culture for 10 days showed markedly increased lipid accumulation (Figure 3(a)). Real-time RT-PCR demonstrated that aP2, LPL, and adipsin mRNA expressions were upregulated 21.1-, 12.9-, and 17.9-folds, respectively, compared with the cells transfected with the empty p1757 plasmid at day 4 after transfection (Figure 3(b)).

**PPARγ induces loss of chondrocytic phenotype in growth plate cells**

Alcian blue staining was used to detect the accumulation of cartilage-specific proteoglycan. At day 10, the control cultures of growth plate cells still accumulated abundant proteoglycan (Figure 4(a)). No significant difference in proteoglycan accumulation was observed in the growth plate chondrocytes that were treated with 5 μM of ciglitazone alone. Compared to the Ad-Gal-infected cells, the Ad-PPARγ-infected cultures were stained less intensely with Alcian blue, especially the surrounding of the cells that contained vacuoles. Addition of 5 μM of ciglitazone to the Ad-PPARγ-infected cultures resulted in a further decrease in proteoglycan matrix.
Quantitative RT-PCR demonstrated that the chondrocyte-specific genes COL2A1 and aggrecan were downregulated by both PPARγ and ciglitazone (Figure 4(b)). Treatment with 5 μM of ciglitazone for 4 days resulted in a 33% decrease of COL2A1 mRNA and a 17% decrease in aggrecan mRNA expression. Combination of both PPARγ adenovirus and 5 μM of ciglitazone resulted in a 50% decrease of COL2A1 mRNA and a 22% decrease in aggrecan mRNA expression.

**PPARγ inhibits T3-induced hypertrophy and mineralization in growth plate chondrocytes**

Thyroid hormone is a crucial regulator in growth plate chondrocyte hypertrophic differentiation and matrix mineralization [15–17]. Growth plate cells treated with thyroid hormone and 5 μM of ciglitazone demonstrated decreased alkaline phosphatase histochemical staining compared to cells treated with T3 alone (Figure 5(a)). Quantitative RT-PCR analysis of growth plate cells in pellet cultures treated with T3 and 5 μM of ciglitazone for 4 days confirmed a 64% decrease in ALP mRNA compared to cells treated with T3 alone (Figure 5(b)). Infection with PPARγ adenovirus in cells treated with T3 also decreased expression of ALP mRNA approximately 71% in the absence of ciglitazone, and 76% in the presence of 5 μM of ciglitazone. Runx2/Cbfa1 is expressed in chondrocytes as they initiate chondrocyte hypertrophy and maturation. Ciglitazone at a concentration of 5 μM decreased the T3-induced expression of Runx2 by 36%. Ad-PPARγ infection decreased the expression of Runx2 mRNA by 54% in the absence of ciglitazone and by 66% in the presence of 5 μM of ciglitazone.

**DISCUSSION**

Growth plate chondrocytes originate from multipotential mesenchymal stem cells that can differentiate into other cell types including adipocytes. We present evidence in this study that growth plate cells continue to display differentiation plasticity and are able to undergo adipogenic changes and a reciprocal decrease of chondrocytic markers when PPARγ is overexpressed.

It has been previously reported that chondrocytes are able to transdifferentiate into adipocytes in vitro [9]. The fatty acid content of the serum added to the culture media has been implicated as a potential cause of this transdifferentiation process [6]. We used a serum-free culture system in these experiments to avoid the possibility that fatty acids in the serum might induce the adipogenic changes observed.
Ciglitazone is one of the thiazolidinedione classes of antidiabetic compounds which can activate PPARγ [18]. Ciglitazone not only increases endogenous PPARγ transcriptional activity [11], but also upregulates PPARγ mRNA and protein expression in growth plate chondrocytes, as observed in this study.

Activation of endogenous PPARγ by ciglitazone or adenoviral overexpression of PPARγ in growth plate chondrocytes resulted in acquisition of adipogenic features in both high-density monolayer cultures and three-dimensional pellet cultures of growth plate chondrocytes, as evidenced by cell morphology, lipid accumulation, and expression of adipocyte marker genes aP2, LPL, and adipsin. Growth plate cells maintained in monolayer cultures seemed to acquire features of the adipocytic phenotype and lose features of the chondrocytic phenotype more readily than those in the pellet cultures (data not shown).

To confirm that the adipocyte-like cells were differentiated directly from chondrocytes and not from other cell types such as undifferentiated mesenchymal stem cells, growth plate cells were transfected with a PPARγ plasmid under the control of a collagen α1(II) promoter. Acquisition of the adipogenic phenotype in these transfected cells was similar to the cells infected with an adenovirus encoding PPARγ and driven by the CMV promoter/enhancer.

While PPARγ and ciglitazone converted cells of the chondrocyte lineage to an adipocytic phenotype, features of the chondrocyte phenotype were simultaneously suppressed. PPARγ inhibited the ability of chondrocytes to terminally differentiate into hypertrophic cells, and suppressed the expression of genes encoding chondrocyte-specific extracellular matrix proteins.

Slipped capital femoral epiphysis (SCFE) is an obesity-related hip disease in children characterized by weakness in the growth plate of the proximal femur, delayed skeletal maturation, and eventual mechanical failure of the physis [19–21]. We speculate that obesity may induce the expression of PPARγ isoforms in growth plate chondrocytes, resulting in phenotypic changes that interrupt normal skeletal maturation of growth plate cells and resulting decreased mineralization of the cartilage matrix would be expected to reduce the resistance of the growth plate to shear stresses. Therefore this delay in maturation at the growth plate, combined with both the increased mechanical stress resulting from increased body weight and the decreased shear stress resulting from delayed maturation, may combine to cause the proximal femoral epiphysis to slip.
ATION levels were normalized with respect to endogenous 18S rRNA. Ad-PPAR
and Runx2 genes 4 days after growth plate cells were infected with
stainings were colored in dark blue. Negative-stained background
treatment.

nonvirus carrying PPARγ
School, Chicago, Ill) for providing the recombinant ade-
We thank Dr J. L. Jameson (Northwestern University Medi-
ACKNOWLEDGMENTS

Figure 5: Inhibition of T3-induced hypertrophy and mineralization in the growth plate cells by PPARγ overexpression or cigli-
tainment in the growth plate cells by PPARγ overexpression or cigli-
(a) Growth plate cells were treated with T3
100 ng/mL) in the presence or absence of ciglitazone (5 μM) for
tained dark blue. Negative-stained background was colored in light green. (b) Quantitative PCR analysis of ALP and Runx2 genes 4 days after growth plate cells were infected with Ad-PPARγ or treated with ciglitazone. Ad-Gal-infected cells without T3 or ciglitazone treatment were used as controls. Gene expression levels were normalized with respect to endogenous 18S rRNA. * P < .05 versus the expression in the Ad-Gal-infected cells with T3-treatment.

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