Macrophage PPARγ, a Lipid Activated Transcription Factor Controls the Growth Factor GDF3 and Skeletal Muscle Regeneration

Graphical Abstract

Highlights
- Macrophage PPARγ is required for skeletal muscle regeneration
- PPARγ regulates GDF3 in muscle infiltrating Ly6C− repair macrophages
- The Gdf3 locus has multiple PPARγ:RXR heterodimer-bound active enhancers
- GDF3 regulates muscle regeneration and enhances primary myoblast fusion

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In Brief
PPARγ is a lipid activated transcription factor essential for fat cells and linked to lipid processing in macrophages. Varga et al. (2016) now show that PPARγ is transcriptionally active and regulates the growth factor Gdf3 in repair macrophages in regenerating skeletal muscle, forming a paracrine axis connecting macrophages to muscle progenitor fusion.

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Macrophage PPARγ, a Lipid Activated Transcription Factor Controls the Growth Factor GDF3 and Skeletal Muscle Regeneration

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SUMMARY

Tissue regeneration requires inflammatory and reparatory activity of macrophages. Macrophages detect and eliminate the damaged tissue and subsequently promote regeneration. This dichotomy requires the switch of effector functions of macrophages coordinated with other cell types inside the injured tissue. The gene regulatory events supporting the sensory and effector functions of macrophages involved in tissue repair are not well understood. Here we show that the lipid activated transcription factor, PPARγ, is required for proper skeletal muscle regeneration, acting in repair macrophages. PPARγ controls the expression of the transforming growth factor-β (TGF-β) family member, GDF3, which in turn regulates the restoration of skeletal muscle integrity by promoting muscle progenitor cell fusion. This work establishes PPARγ as a required metabolic sensor and transcriptional regulator of repair macrophages. Moreover, this work also establishes GDF3 as a secreted extrinsic effector protein acting on myoblasts and serving as an exclusively macrophage-derived regeneration factor in tissue repair.

INTRODUCTION

Tissues suffer damage during an organism’s lifetime. In order to maintain the body’s integrity and homeostasis, it is critically important to achieve complete regeneration. In many cases, a straightforward paradigm can be applied whereby organ injury induces expansion and differentiation of a quiescent population of tissue-specific stem-cell-like progenitors. Impaired injury-related immune response has been shown to greatly influence regeneration in liver, central nervous system, or skeletal muscle (Chazaud, 2014; Duffield et al., 2005; Laflamme and Murry, 2011; Rapalino et al., 1998). Immune cells and in particular macrophages sense the injury, remove damaged tissues, then initiate restoration of tissue integrity via promoting repair mechanisms. During this latter phase the immune response regulates the reengagement of tissue progenitor cell populations to support cell growth and differentiation. Our knowledge is fragmented on how macrophages employ sensory and regulatory mechanisms and use effector functions to serve their reparatory roles. We sought to identify such integrated regulatory mechanisms that equip a macrophage with the capacity to contribute to a timely progression of repair.

We found that the fatty acid regulated transcription factor, peroxisome proliferator-activated receptor gamma (PPARγ) (Tontonoz et al., 1998), was required in repair macrophages during skeletal muscle regeneration. Mice with a deletion of PPARγ in their myeloid lineages showed a pronounced delay in regeneration. PPARγ regulated the expression of a secreted factor, GDF3, in repair macrophages. GDF3 deficiency impaired muscle regeneration and recombinant GDF3 enhanced repair in vivo and the fusion of primary myogenic precursor cells (MPCs) in vitro cultures. Our data reveal a PPARγ-GDF3 pathway with sensory, gene regulatory, and effector components in which PPARγ in repair macrophages responds to signals and support the timely promotion of tissue repair during skeletal muscle regeneration.
RESULTS

PPARγ Is Expressed in Macrophages of the Cardiotoxin Induced Skeletal Muscle Injury Model

Skeletal muscle possesses robust regenerative capacity; therefore, it provides us with an excellent model system to study regeneration. The best characterized experimental model of skeletal muscle injury is the toxin-induced injury and regeneration. We triggered skeletal muscle damage in the tibialis anterior (TA) muscle of mice by intramuscular injection of the snake venom, Cardiotoxin (CTX), to induce a homogenous and synchronous muscle damage that is repaired with the active contribution of infiltrating immune cells. We isolated macrophage populations from injured muscle and interrogated their gene-expression profiles by microarray analysis. When the expression profiles of inflammatory Ly6C+ and repair Ly6C− macrophages derived from injured muscle at day 2 CTX injury were compared, gene ontology (GO) annotation categories belonging to lipid and carbohydrate metabolism dominated the biological processes that were the most robustly upregulated in the Ly6C− (repair) macrophages (Figure S1A). When analyzing the expression data, we found that a master regulator of metabolism, Pparg, was highly expressed in these macrophages. Using publicly available gene-expression data within the Immunological Genome Project, we compared the expression of Pparg in muscle infiltrative macrophages to that of their direct precursors, Ly6C+ monocytes (Varga et al., 2013), and various other myeloid cells (Figure S1B). We found that Pparg in muscle macrophages was highly expressed, and that only two in vivo macrophage subtypes, alveolar macrophages and splenic red pulp macrophages expressed Pparg higher. In contrast to Pparg, Ppara was not expressed in muscle infiltrative macrophages, while the expression of Ppard showed a declining expression in the course of regeneration (Figure S1C).

On the basis of these findings, we hypothesized that macrophage PPARγ is a metabolic sensor and regulator of skeletal muscle regeneration. To test this hypothesis, we used the Pparg−/− Ly22-cre mouse strain, which is deficient in PPARγ specifically in myeloid lineages (Clausen et al., 1999). When CD45+ cells, which comprise all infiltrating hematopoietic cells, or sorted macrophages, were isolated from injured skeletal muscle, the expression of Pparg was detected in these cells by RT-qPCR (Figures S1D and S1E) in wild-type (WT) animals. Furthermore, the expression of Pparg was greatly diminished in corresponding CD45+ cells and macrophages isolated from Pparg−/− Lyz2-cre animals, validating the suitability of this genetic model for these experiments.

Macrophage PPARγ Regulates Skeletal Muscle Regeneration

WT and Pparg−/− Lyz2-cre animals were injected with CTX to induce TA muscle injury and then regeneration was analyzed by a combination of morphometric and flow cytometry analysis. We found Pparg−/− Lyz2-cre animals showed a pronounced delay in muscle regeneration (Figures 1A–1D and S2A). First, the cross-sectional area (CSA) of the regenerating muscle fibers was significantly smaller in the Pparg−/− Lyz2-cre than in WT mice at day 8 and day 21 following CTX injury (Figures 1C and S2A). Second, there were a significantly higher number of phagocytic and/or necrotic fibers present at day 8 post CTX in Pparg−/− Lyz2-cre mice (Figures 1A and 1B), indicating either a delayed clearance of dying myofibers or an altered dynamics of muscle fiber death in Pparg−/− Lyz2-cre animals. Third, increased inflammatory infiltration persisted in small regions in the regenerative areas in Pparg−/− Lyz2-cre muscles at day 8 (Figure 1A), which were resolved by day 21 (Figure S2B). Next, we wanted to ascertain whether PPARγ deficiency in the hematopoietic compartment was the major contributor to the observed phenotype. To prove this, we used a second genetic model, in which bone marrow from the epiblastic conditional ablation of Pparg (Pparg−/−, Sox2-cre+) (Nadra et al., 2010) or WT animals were used to reconstitute the hematopoietic compartment in irradiated WT animals (bone marrow transplanted or BMT animals). TA muscles of recipient BMT animals were injected with CTX 12 weeks after BMT and histological analysis of muscle regeneration was carried out 22 days post injury. When compared with animals that received WT bone marrow (WT BMT), mice that received bone marrow deficient in PPARγ (Pparg−/−, Sox2-cre+ BMT) exhibited a profound deficit in regeneration (Figures 1E and 1F). Further underlying the importance of PPARγ in muscle regeneration, full body Pparg−/− Sox2-cre+ animals displayed impairment in their skeletal muscle regeneration (Figure S2C).

PPARγ Deficiency Does Not Alter Macrophage Infiltration or Differentiation in Injured Muscle

Several possible reasons could explain why macrophage PPARγ deficiency leads to such impairment in muscle regeneration. One underlying reason behind our observations could be a decreased macrophage infiltration in Pparg−/− Lyz2-cre animals. To monitor the cellular dynamics of immune infiltration in CTX injured muscle, we treated WT and Pparg−/− Lyz2-cre animals with CTX, then isolated and analyzed immune cells from injured muscles on days 1, 2, or 4, using CD45+ magnetic bead selection. We found no major difference between the numbers and types (Ly6Cmid F4/80−, Ly6Cmid F4/80+ and Ly6C− F4/80high macrophages) of infiltrating immune cells in WT vs. Pparg−/− Lyz2-cre animals (Figure S3), with the exception of minor alterations in the ratio of neutrophils at day 1 and in the total number of CD45+ cells at day 6.

Next, we wanted to explore which macrophage functions might be relevant to muscle regeneration and regulated by PPARγ activity. To test the possible contribution of impaired phagocytosis, we used bone-marrow-derived macrophages (BMDMs) isolated from WT or Pparg−/− Lyz2-cre animals, and from bone marrow transplanted WT BMT or Pparg−/−, Sox2-cre+ BMT animals (Figures S4A for experimental setup and S4B for data). We set up a phagocytosis assay, in which fluorescently labeled necrotic C2C12 myoblasts were co-incubated with BMDMs labeled with a different fluorescent dye. Pparg−/− Lyz2-cre BMDMs showed no significant increase in the number of phagocytosing BMDMs or in the amount of phagocytosed substrate as compared with WT BMDMs (Figure S4B). Similar results were obtained using BMDMs derived from WT BMT or Pparg−/−, Sox2-cre+ BMT animals, except that Pparg−/−, Sox2-cre+ BMT BMDMs were able to phagocytose a greater load. Our results indicated that an inadequate phagocytic clearance was unlikely to be responsible for the observed delay.
Figure 1. Impaired Regeneration of Skeletal Muscle in PPARγ-Deficient Animals

(A) Representative images of HE stained skeletal muscle from WT and Pparg<sup>fl/fl</sup> Lyz2-cre animals prior (day 0) or post CTX induced injury (day 8) are shown. Asterisk labels phagocytic and/or necrotic fibers and arrow points to foci of inflammatory infiltrations. IHC detection of desmin (red), F4/80 (green), and nuclei (blue) at day 8 post CTX injury is also shown. Scale bars in the upper left represent 50 μm.

(B) The ratio of phagocytic and/or necrotic fibers relative to all regenerative fibers at day 8 of regeneration in WT and Pparg<sup>fl/fl</sup> Lyz2-cre muscle sections is shown.

(C) Fiber size repartition of regenerating muscle in WT or Pparg<sup>fl/fl</sup> Lyz2-cre animals at day 8 and day 21 post CTX injury.

(D) Average fiber cross section area (CSA) of regenerating muscle at indicated timepoints post CTX injury in WT and Pparg<sup>fl/fl</sup> Lyz2-cre animals.

(E) n = (numbers of individual muscles, derived from WT or Pparg<sup>fl/fl</sup> Lyz2-cre mice): 4 and 4 for day 0 samples, 5 and 6 for day 8 samples, 5 and 5 for day 21 samples and 5 and 5 for day 63 samples, respectively. Representative images of HE stained skeletal muscle 22 days after CTX injury from bone marrow transplanted (BMT) animals that received either WT or Pparg<sup>fl/fl</sup> Sox2-cre bone marrow.

(F) Muscle fiber CSA of BMT animals 22 days post CTX injury. n = 8 muscles for both genotypes. In all bar graphs, mean values ± SEM are shown. For Pparg expression in macrophages and CD45<sup>+</sup> cells and for additional histological analysis, see Figures S1 and S2. For the FACS analyses of infiltrating cells, see Figure S3.
Figure 2. PPARγ-Regulated Macrophage Functions and Genes

(A) Effect of BMDM derived conditioned media on the proliferation of primary myoblasts (± SEM). n = 4 or 3 for WT or Ppargfl/fl Lyz2-cre BMDM supernatant. For the complete analysis, see Figures S4C and S4D. n = 6 for both genotypes.

(B) Effect of BMDM derived conditioned media on the differentiation of primary myotubes (± SEM). For the complete analysis, see Figures S4C and S4D. n = 6 for both genotypes.

(C) Transcriptional analysis of the Ly6C+ and Ly6C- macrophage populations derived from WT and Ppargfl/fl Lyz2-cre BMDM supernatant. For the complete analysis, see Figures S4C and S4D. n = 6 for both genotypes.

(C–E) Heatmap representation of genes that show differential (p = 0.05, min. 1.5X FC) expression in the four sorted WT vs. Ppargfl/fl Lyz2-cre macrophages in day 1 Ly6C+, D2 Ly6C+ and D2 Ly6C−, and D4 Ly6C− cells (labeled as D1 Ly6C+ etc.). In each heatmap, the differentially expressed genes are highlighted within a red square and the expression pattern of these genes in the other macrophage subtypes is also shown for reference. Blue and red arrows label genes that are upregulated and downregulated, respectively.
Macrophage PPARγ Regulates Myoblast Differentiation in a Paracrine Manner In Vitro

These results led us to test whether macrophage PPARγ activity confers a yet unidentified muscle differentiation-promoting phenotype to macrophages, which could explain the observed delayed muscle regeneration in animals deficient in PPARγ in macrophages. To test this hypothesis, we used in vitro muscle precursor cell proliferation or differentiation assays that utilize primary myoblasts isolated from WT mice (Figures 2A and 2B). In the first assay, we cultured primary myoblasts with conditioned medium derived from non-treated, interferon-γ (IFN-γ), or interleukin-4 (IL-4)-treated WT and Pparg<sup>fl/fl</sup> Ly22-cre BMDMs, in conditions favoring cell proliferation and measured the proliferation index by detecting Ki67<sup>+</sup> cells by immunofluorescence (IF). As expected, conditioned medium derived from IFN-γ-treated WT BMDMs increased myoblast proliferation (Mounier et al., 2013). Conditioned medium from non-treated Pparg<sup>fl/fl</sup> Lyz2-cre BMDMs phenocopied the proliferation enhancing effect of inflammatory WT BMDMs on myoblasts (Figure 2A). These results indicated that PPARγ in macrophages modulated an unknown signaling system that could influence myoblast proliferation in a paracrine manner. Next, we tested the effect of BMDM-derived conditioned media on the differentiation of myoblasts by counting the number of cell nuclei within freshly formed desmin-positive myotubes cultured in differentiation medium (Figure 2B and Figure S4C). As expected, we observed a large increase in differentiation when myoblasts were grown in conditioned medium derived from IL-4-treated WT BMDMs. Importantly, this increased differentiation was abrogated when conditioned medium from IL-4-treated Pparg<sup>fl/fl</sup> Lyz2-cre BMDMs was added to differentiating myoblasts. This effect was seen in several independently isolated primary myoblast cell lines that were used for the experiments (Figure S4C). BMDM supernatant derived from IFN-γ-treated cells, on the other hand, did not alter myoblast differentiation (Figure S4D). Our results raised the possibility that similar PPARγ-dependent paracrine signaling events took place in situ during regeneration, where muscle infiltrative macrophages and MPCs might interact to achieve a synchronized and timely regeneration.

PPARγ Regulates Cell-Type-Specific Genes in Muscle Infiltrating Macrophages

Next, we set out to identify PPARγ-dependent regulatory circuits that connect macrophages to myotube differentiation in a paracrine manner. Because PPARγ is a transcription factor, we presumed that a relevant change in the gene expression in muscle macrophages must shed light on the regulatory circuit that is abrogated in Pparg<sup>fl/fl</sup> Lyz2-cre macrophages. We isolated populations of macrophages from regenerating muscle from WT and Pparg<sup>fl/fl</sup> Lyz2-cre animals and analyzed their gene-expression profiles by microarrays (Figures 2C–2E and S4E and S4F). We selected inflammatory Ly6C<sup>+</sup> macrophages at day 1 and 2, and repair Ly6C<sup>−</sup> macrophages at day 2 and 4 post CTX injury and compared their gene expression by two-way ANOVA tests (Table S1). We created heatmaps for all four examined macrophage subsets (Figure 2C). These heatmaps show all genes that were differentially expressed in one relevant subset and also show the expression pattern of these genes in all the other macrophage subsets. The top five genes that were most differentially regulated in WT vs. Pparg<sup>fl/fl</sup> Lyz2-cre cells are shown in Figure 2D. The number of genes that were concordantly regulated in a PPARγ-mediated manner in more than one macrophage subtype is shown in Figure 2E. We hypothesized those genes could be under regulation by PPARγ that were expressed differently in more than one subtype of muscle macrophages. Accordingly, we combined the lists of upregulated genes reported by the ANOVA analysis of WT vs. Pparg<sup>fl/fl</sup> Lyz2-cre comparisons. Although many genes were differentially regulated in a single type of muscle macrophages, only five genes (Saa3, Hebp1, Plxnd1, Apold1, Tsg101) were upregulated in all four investigated subtypes of PPARγ-deficient muscle macrophages (Figure 2E and Table S1). Next, we analyzed the gene sets that were downregulated in Pparg<sup>fl/fl</sup> Lyz2-cre macrophages. There was only one gene, namely growth differentiation factor 3 (Gdf3), that was consistently downregulated in all four investigated macrophage populations (Figures 2D and 2E). Thus, we identified several putative PPARγ target genes that showed consistent PPARγ dependency in more than one muscle macrophage subset. To ascertain the PPARγ-dependent regulation of some representative genes, we measured the mRNA expression of Gdf3, Apold1, Hebp1, and Plxnd1 by RT-qPCR in macrophage subsets sorted from injured muscle (Figure S5A). This analysis confirmed the results derived from the microarray experiments. The expression pattern of a short panel of previously described PPARγ-dependent (M2) alternative genes (Odegaard et al., 2007) indicated that the repair macrophages in CTX injured muscles were not canonical M2 macrophages, and that PPARγ exerted little, if any, influence on their expression (Figure S5B). Along the same line, while a total body deficiency in STAT6, the master regulator of IL4 signaling, caused increased presence of phagocytic and/or necrotic fibers at day 8 (Figure S5C), it did not affect the CSA of new myofibers (Figure S5D).

The genes we identified as PPARγ-dependent in muscle macrophages did not belong to the group of canonical PPARγ-regulated genes described in various myeloid cells in earlier studies (such as Plin2, Cd36, Angptl4, or Fabp4) (Szanto et al., 2010; Welch et al., 2003). One possible reason for this discrepancy could be that most in vitro studies apply synthetic or natural ligands of PPARγ to study the transcriptional activity of the receptor upon ligand activation. Therefore, we wanted to see whether synthetic PPARγ ligand activation of infiltrating macrophages gave rise to transcriptional changes that are more reminiscent of the list of previously identified PPARγ target genes. For this reason, we treated WT animals with rosiglitazone (RSG) via gavage and analyzed the ligand-dependent gene expression downregulated or upregulated in WT vs. Pparg<sup>fl/fl</sup> Lyz2-cre cells, respectively. The blue and red arrows point to the direction of increasing fold change difference. For RT-qPCR validation of mRNA expression, see Figure S5S.

(D) Top 5 up and downregulated genes in the four sorted macrophage populations in Pparg<sup>fl/fl</sup> Lyz2-cre macrophages. Table lists gene symbols and fold change differences (FC). Gdf3 and Apold1, the genes that are down- or upregulated in Pparg<sup>fl/fl</sup> Lyz2-cre in all four subtypes, are highlighted in color.

(E) Venn-diagrams show the overlap of the number of genes that are down- or upregulated in Pparg<sup>fl/fl</sup> Lyz2-cre macrophages in the four analyzed populations.
changes in macrophages (Figures S4F and S5E, and Table S1). We found that many more genes were regulated by RSG treatment in Ly6C⁺ than in Ly6C⁻ cells. Again, the genes that showed differential expression upon RSG treatment in Ly6C⁺ cells did not contain established PPARγ-regulated genes, nor the six differently regulated genes that appeared to be under PPARγ regulation in all macrophage subsets. Although RSG treatment caused the differential regulation of fewer genes in Ly6C⁻ cells, the most robustly upregulated gene was Angpt4, one of the best-characterized PPARγ target genes. This suggests that not only Ly6C⁺ macrophages at day 2 expressed PPARγ, but that the receptor was also sensitive to the activating effect of an exogenous ligand in Ly6C⁻ cells. It is important to note that Gdf3, the gene that was found to be consistently downregulated in Pparg⁻/⁻ Lyz2-cre macrophage subsets, was also regulated by RSG treatment (only) in Ly6C⁺ macrophages. Next, we took the list of 43 genes that showed ligand dependent upregulation in Ly6C⁺ macrophages upon RSG treatment and created a heatmap representation to see how these genes were regulated in the absence of RSG treatment (Fig S5E). Even without RSG treatment, most of the otherwise RSG dependent genes showed a characteristic induction as Ly6C⁺ macrophages differentiated into Ly6C⁻ cells and an even further induction by day 4. This observation raised the intriguing possibility that the underlying reason behind the limited number of PPARγ ligand regulated genes in Ly6C⁺ macrophages was that most of these genes were already induced during muscle regeneration, even in the absence of exogenous synthetic ligand treatment. Related to this hypothesis, we detected a dynamic in situ regulation of eicosanoid synthesis during regeneration. While inflammatory eicosanoids (e.g., PGE₂ and PGF₂α) were detectable in the early inflammatory stages of injury, they were later replaced by lipid mediators produced by murine 12/15-lipoxygenase (Alox15) that have been implicated in ligand activation of PPARγ such as 12-HETE and 15-HETE (Figure S5F) (Huang et al., 1999).

**GDF3 Is a Macrophage-Derived PPARγ-Dependent Member of the TGF-β Family**

To focus on putative PPARγ regulated genes whose activity could promote muscle regeneration, we interrogated the list of differentially expressed genes for genes that (1) were PPARγ-dependent in more than one macrophage subset, (2) coded a secreted factor and (3) whose activity might be linked to muscle differentiation. Of note, one gene, Gdf3 (Levine and Brivanlou, 2006; Levine et al., 2009; Shen et al., 2009), fit all these criteria. Gdf3 was statistically significantly downregulated in Pparg⁻/⁻ Lyz2-cre cells in all four investigated macrophage subsets (Figures 2D and 2E, Table S1). Gdf3 belongs to the TGF-β family, whose members are secreted factors acting in a paracrine manner. Finally, several members of the TGF-β family are known regulators of muscle regeneration, including GDF8 (also known as Myostatin) (McPherron et al., 1997). Therefore, we selected Gdf3 as the most likely PPARγ-dependent gene that contributes to muscle regeneration for further analysis.

**PPARγ Occupies a Complex Set of Active Enhancers around the Gdf3 Locus**

Next, we wanted to characterize the genomic events that are responsible for the regulation of Gdf3 by PPARγ. We elected to use BMDMs, a readily available in vitro model system that allowed us to employ high-throughput genomic and epigenomic methods to interrogate the regulatory mechanism exerted by PPARγ on the Gdf3 locus. We established that WT and Pparg⁻/⁻ Lyz2-cre BMDMs provided a platform with good correlation to study the PPARγ-dependent regulation of Gdf3, as PPARγ deficiency in BMDMs abrogated the expression of both the canonical PPARγ target gene Angpt4 and that of Gdf3 (Figure 3A). Then, we compiled epigenomic and genomic data to identify the relevant enhancers that were active and possibly under PPARγ regulation in BMDMs (Figure 3B). We included CTCF as a binding factor of insulator regions and RAD21, as a component of the cohesin complex to determine the boundaries of potential chromatin loops or topological domains, PU.1 as a key lineage determining factor in macrophages, RXR (the obligate heterodimeric partner of PPARγ), and PPARγ chromatin immunoprecipitation (ChIP)-seq data derived from thioglycolate elicited peritoneal macrophages and adipocytes. We combined these data with active epigenetic marks from H3K4me3 ChIP-seq experiments and GRO-seq data from BMDMs. Based on the common CTCF and RAD21 binding sites (Daniel et al., 2014; Merkenschlager and Odom, 2013), the transcription unit of Gdf3 appeared to be approximately between −50 Kb to +50 Kb. Our definition of putative, active enhancers included (1) binding of PU.1, (2) presence of detectable enhancer transcript (GRO-seq signal), and (3) RXR or PPARγ binding. This approach was validated by applying the same criteria to the Angpt4 locus, in which we readily identified its PPARγ-dependent enhancer (Figure S6A). On the basis of these criteria, we nominated 14 putative active enhancers at a distance from +38 Kb to −47 Kb relative to the transcription start site of Gdf3 (Figures 3B and S6B). As we show in Figure 3C, binding of PPARγ and RXR could be readily detected on five of these selected enhancers (at +7.3 Kb, −21 Kb, −25 Kb, −44 Kb, and −47 Kb) if we compared WT to Pparg⁻/⁻ Lyz2-cre BMDMs. These data strongly suggested that Gdf3 was regulated by one or several of these PPARγ:RXR binding sites.

**GDF3 Is a Regulator of Myoblast Proliferation, Differentiation and Muscle Regeneration**

Next, we analyzed the GDF3 protein expression in whole muscle lysates of CTX injured WT mice, which provided a snapshot of GDF3 protein level during regeneration. The protein expression followed the induction seen at the mRNA level in macrophages and showed a pronounced induction, which peaked at day 4 (Figure 4A), at the time when inflammation subsides and regenerative processes start to dominate within the injured muscle. Importantly, the induction of GDF3 expression was detectable in the CD45⁺ (hematopoietic) compartment and was diminished at both mRNA and protein amount in Pparg⁻/⁻ Lyz2-cre animals (Figures 4B–4D). Next, we further investigated GDF3 expression in alternative models of muscle injuries. We found that, similarly to CTX injury, GDF3 protein expression was induced during glycerol-mediated injury and regeneration in WT but diminished in PPARγ macrophage-deficient animals (Figure 4E). Furthermore, not only the mRNA expression of Gdf3, but the entire panel of genes that showed strong PPARγ dependency in the CTX model, was regulated concordantly in the two models of injury (Figure 4F). GDF3 protein expression was also induced in muscle.
Figure 3. Gdf3 Is a PPARγ Target Gene in BMDMs
(A) mRNA expression (± SD) of Angptl4, a canonical PPARγ target gene, Pparg, Gdf3, and Apobec1, a nearby, not regulated gene, are shown in BMDMs (n = 4 for WT and n = 5 for Pparg<sup>fl/fl</sup> Lyz2-cre).

(B) Identification of possible enhancers around the Gdf3 locus. The selection criteria for enhancers possibly involved in Gdf3 regulation are described in the text and in Figures S6A and S6B. Putative enhancers are labeled by vertical lines. Blue verticals highlight enhancers without PPARγ ChIP enrichment, red verticals label enhancers where enrichment in PPARγ binding in WT BMDMs was detected by PPARγ ChIP.

(C) ChIP on the putative enhancer regions reveal PPARγ binding at +7.3 Kb, -21 Kb, -25 Kb, -44 Kb, and -47 Kb enhancers around the Gdf3 locus. Representative graphs showing PPARγ, RXR, or IgG ChIPs carried out on two samples are shown. Angptl4 enhancer and Gdf3 +16 Kb enhancer are shown as positive and negative controls, respectively.
samples exposed to crush- and freeze-injuries, which are toxin-free methods (Figure 4G). Due to recent publications that reported a high tendency for false positive detection of GDF proteins in protein detection applications (Egerman et al., 2015), it is important to note that the GDF3 protein induction during CTX injury was undetectable in muscle samples from \textit{Gdf3}/\textit{C0}/\textit{C0} animals (Figure 4H). To summarize, GDF3 is a macrophage-derived protein whose expression is induced in various models of muscle regeneration in a PPAR\textsubscript{g}-dependent manner.

According to our model, the regeneration delay in macrophage PPAR\textsubscript{g}-deficient animals was, at least partly, attributable to a diminished macrophage-derived GDF3 secretion within regenerating muscles. This model posits that GDF3 deficiency in macrophages should yield impairment in regeneration comparable to what was observed in \textit{Pparg}\textsuperscript{fl/fl} \textit{Lyz2-cre} animals. Indeed, muscle regeneration after CTX injury was altered in full-body \textit{Gdf3}/\textit{C0}/\textit{C0} animals at day 8 (Figures 5A and 5B). It has been reported that the full-body deletion of \textit{Gdf3} shows incomplete penetrance (Shen et al., 2009), which suggests possible compensatory mechanisms. To limit their involvement and ascertain the hematopoietic source of GDF3 during muscle regeneration, we generated BMT animals reconstituted with \textit{Gfd3}/\textit{C0}/\textit{C0} BM. When the GDF3 chimeric animals were challenged with CTX induced muscle injury, they exhibited impairment in regeneration at day 16 and 20 (Figures 5C and 5D). When compared with WT BMT animals, \textit{Gdf3}/\textit{C0}/\textit{C0} chimeras contained more regenerating myofibers with smaller CSA and the regenerating muscle was replete with lipid accumulations, which are hallmarks of defective muscle regeneration (Figures 5C and 5D). Other cell types, such as fibro-adipogenic progenitors (FAPs) are involved in muscle regeneration (Heredia et al., 2013; Lemos et al., 2015). In line with our results from the \textit{Gdf3}/\textit{C0}/\textit{C0} BMT experiment (Figures 5C and 5D) and with the mRNA and protein expression data showing GDF3 expression in the CD45\textsuperscript{+} compartment (Figure 4), \textit{Pdgfra} expressing FAP cells isolated from D2 regenerating muscle barely expressed \textit{Gdf3} and \textit{Lyz2} mRNA (Figure 5E), rendering the involvement of FAPs unlikely in the macrophage-derived GDF3-driven effects on muscle regeneration.

To further prove the requirement for GDF3 in muscle regeneration, we injected recombinant GDF3 into CTX injured muscles of \textit{Pparg}\textsuperscript{fl/fl} \textit{Lyz2-cre} mice. We found that the exogenously added GDF3 rescued the regeneration deficit seen in these animals (Figures 6A and 6B). To characterize the function of GDF3 in detail, we cultured primary myoblasts with or without recombinant GDF3. We found GDF3 slightly decreased myoblast proliferation (Figure 6C, left panel). We detected an even more robust effect of GDF3 on myotube formation, as myoblasts...
cultures showed a pronounced increase in their fusion index in the presence of GDF3 (Figures 6C, right panel, and 6D). Myotube formation depends on cell motility, terminal differentiation and cell fusion. In a specific fusion assay, we showed that GDF3 was a potent inducer of myotube formation (Figure S7 A), while a differentiation assay indicated that GDF3 did not affect the terminal differentiation of myoblasts into myocytes (Figure S7 B).

Next, we investigated whether the SMAD2 phosphorylation pathway, which is involved in the signal transduction of several TGF-β superfamly members, is engaged during muscle regeneration. We found a detectable induction of in situ pSMAD2 signals in muscles at day 4 of regeneration (Figure 7A), at the time when GDF3 expression peaked in the injured muscle. Furthermore, SMAD2 phosphorylation was significantly increased during in vitro treatment of primary myoblasts with GDF3 (Figures 7 B and 7C).

In search for the molecular changes triggered in muscle progenitors in the presence of GDF3, we differentiated in vitro primary myoblasts with or without GDF3 and interrogated the gene-expression changes by RNA-seq. First, we compared the profile of primary myoblasts and myoblast-derived myotubes that were cultured in the presence or absence of GDF3. The expression pattern of a preselected list of genes relevant to muscle differentiation (Figure 7 D) validated our experimental system. Next, we compared the expression profile of differentiating myotubes cultured with or without GDF3. The list of the differentially regulated genes (Figure 7E and Table S2) showed that a limited set of transcripts were either induced or repressed in the presence of GDF3. Several of the differentially regulated genes, including Bex1, (Jiang et al., 2016; Koo et al., 2007), Sgca (Matsumura et al., 1992), and Camk1g, have been implicated in muscle regeneration, muscle structure and/or Ca²⁺ homeostasis, showing that macrophage-derived GDF3 could elicit biologically relevant changes during muscle regeneration.

If GDF3, a macrophage-derived secreted factor, can regulate in vitro and in situ muscle differentiation and regeneration, then we wanted to ask whether GDF3 is the only macrophage-derived TGF-β family member that is relevant in the context of CTX induced muscle injury. Therefore, we reanalyzed the transcriptomic features of muscle infiltrative macrophages to chart the expression and dynamics of the TGF-β family signaling system (Figure 7F and Figure S7C). Three ligands (Gdf3, Gdf15, and Inhba) showed notable gene expression dynamics in muscle infiltrative macrophages. GDF3 expression peaked in repair macrophages and showed definitive, consistent regulation by PPARγ. The two other family members (Figure S7C), Gdf15...
DISCUSSION

Skeletal muscle possesses excellent regenerative capacity, therefore it was striking to see that after CTX injury, full-body Ppargfl/fl Sox2-cre animals showed signs of residual inflammation and impaired regeneration. The true extent of the involvement of macrophage PPARγ in the regeneration failure in these animals is unclear for several reasons, including the uncharacterized, but presumably inflammatory state of these animals and the potential involvement of non-macrophage (e.g., muscle) PPARγ in regeneration. Therefore we used two distinct genetic models (BMT and conditional PPARγ deficiency, Ppargfl/fl Lyz2-cre), which allowed us to focus on the role of PPARγ in macrophages. The delay in regeneration in macrophage PPARγ-deficient animals was less profound than in the epiblastic Ppargfl/fl mice, yet it was detectable as long as three weeks after the initial injury, thus appearing to be among the most dramatic reported deficiencies in regeneration caused by impairments in macrophage functions (Mounier et al., 2013).

Our analysis did not reveal a gross difference in macrophage number or differentiation in Ppargfl/fl Lyz2-cre animals, unlike two other reported experimental systems where AMPK or IGF1 deficiency in muscle infiltrative macrophages led to altered macrophage differentiation (Mounier et al., 2013; Tonkin et al., 2015). Although alternatively activated macrophages have been implicated in tissue repair and PPARγ has been reported to be a regulator of alternative macrophage polarization (Odegaard et al., 2007), we have previously reported that muscle Ly6C+ and Ly6C- macrophages do not correspond to canonical alternatively polarized macrophage populations (Varga et al., 2016) in the CTX model. Therefore, it is not surprising that, in this model, PPARγ is controlling genes other than alternative macrophage-related ones, reported to be PPARγ-dependent in other tissue compartments and contexts (Odegaard et al., 2007). The fact that the regeneration impairment in Stat6−/− animals did not manifest in a decrease in CSA, also suggest that PPARγ, in this experimental context, acts through mechanisms other than modulating alternative macrophage activation. Systematic transcriptomic analyses, however, provided clues about both the sensory and the regulatory roles of PPARγ in muscle-infiltrating macrophages. It is important to stress that earlier descriptions of direct PPARγ transcriptional target genes often reported lipid metabolic genes as the main targets of PPARγ in macrophages, which could poorly explain the anti-inflammatory role of the receptor (Szanto et al., 2010; Welch et al., 2003). We report here that the transcriptional activity of PPARγ is unique in muscle macrophages, because the most robustly changing genes (such as Saa3, Hebp1) were linked to inflammation, rather than to lipid metabolism. Second, in vivo treatment with RSG identified the Ly6C+ repair macrophages as an in situ macrophage subtype that could be activated by a synthetic ligand for PPARγ. The surprising fact that RSG treatment elicited characteristically different gene-expression changes in Ly6C+ and Ly6C− macrophages isolated from the same tissue and timepoint underscores the notion that distinct macrophage subsets have differential responses to environmental cues. A possible interpretation of the available data would be the involvement of a yet unidentified endogenous ligand for PPARγ whose activity is restricted to the Ly6C− compartment, which could

and Inhba, were also regulated during muscle regeneration, and both genes exhibited partial PPARγ dependency. The PPARγ-GDF3 regulatory axis described in this study therefore identifies a sensory-regulatory-effector mechanism, by which macrophages are regulators of the tissue progenitor compartment, namely MPCs. This axis orchestrates tissue regeneration, possibly in unison with other members of the TGF-β family, leading to synchronous regeneration.
Figure 7. Effects of GDF3 on Myogenesis

(A) Increased pSmad2 phosphorylation in regenerating muscles peaking at day 4 post CTX injury.
(B and C) Increased Smad2 phosphorylation in primary myoblasts treated with rGDF3. IF images and percentage of pSMAD2-positive cells are shown. Mean values ± SEM are shown. n = 3.
(D) Heatmap representation of the expression changes of myogenic genes validating the utilized in vitro primary myoblast assay.
(E) Heatmap representation of genes that are differentially expressed (min. fold change difference of 1.2X between differentiated myoblasts ± rGDF3) in the presence of recombinant GDF3 during myoblast differentiation.
(F) Heatmap representation of members of the TGF-β superfamily signaling system that are expressed and regulated, or expressed but not regulated in muscle derived macrophages. For non-expressed members, see Figure S7C.
explain the tendency of otherwise RSG inducible genes to be upregulated in the Ly6C+ macrophages even in the absence of the synthetic ligand. Whether the dynamic regulation of in situ eicosanoid synthesis we detected during regeneration could be behind the apparent ligand activation of the receptor requires further investigation.

From the perspective of muscle regeneration, the most notable finding was the identification of GDF3, a TGF-β family member, which showed consistent regulation by PPARγ in all relevant macrophage subtypes. To ascertain that GDF3 was not only a PPARγ-dependent factor, but also a direct PPARγ target, we analyzed an extensive range of genomic and epigenomic data. Although it is clear that GDF3 is expressed in a PPARγ-dependent fashion and can be induced by ligand in muscle-derived Ly6C+ macrophages, direct regulation by PPARγ is challenging to prove, because ligand-dependent regulation appears to be macrophage subtype specific and not detectable in BMDMs. However, we have provided data that are consistent with direct regulation, even in BMDMs.

It is noteworthy that both GDF3 gene and protein expressions were much lower in the CD45+ fraction isolated from injured muscle than in the hematopoietic compartment. Considering that the separation of CD45+ cells is inherently incomplete, our results indicate that macrophages are the predominant, if not the only, source of GDF3 within the injured tissue. This exclusivity sets GDF3 apart from other macrophage-derived regenerative factors, such as IGF1 (Tonkin et al., 2015), which is also produced by muscle and in the liver upon injury. The timing and localization of GDF3 protein in the CTX and other, unrelated injury models firmly suggested that GDF3 is a general, macrophage-specific regulator of muscle regeneration.

To link macrophage biology to tissue regeneration, we analyzed the role of macrophage derived GDF3 in muscle regeneration in a combination of in vivo and in vitro approaches. Foremost, two genetic models of GDF3 deficiency reported a delay in regeneration. While the decrease in average CSA in Gdf3−/− animals was comparable to that seen in Pparg−/− Lyz2-cre animals, Gdf3−/− animals did not display persistent inflammation and delayed resolution of necrotic and/or phagocytic fibers. This suggested that PPARγ regulated several relevant pathways during regeneration. Notably, a gain of function experiment revealed that exogenous GDF3 could counteract the deleterious effect PPARγ deficiency in macrophages. Our in vivo results with BMDM supernatants and myoblasts indicated the presence of a regulatory circuit between macrophages and muscle cells and showed that GDF3 appeared to be an especially robust enhancer of myoblast fusion.

Because other cell types are also involved in the regeneration process (Heredia et al., 2013; Joe et al., 2010; Uezumi et al., 2010), it cannot be excluded that GDF3 is only one of the TGF-β family members that are active during regeneration and that it has effects on other cell types such as FAPs as well. It is remarkable, though, that the key elements of the myogenic cross talk between cell types can be modeled in vitro using macrophages and myoblasts only, arguing that these two cell types and their interactions are critical to support regeneration.

Our findings also carry potential implications for pathological circumstances in which recurrent muscle damage and asynchrony in repair due to genetic conditions leads to debilitating degenerative muscle diseases, such as Duchenne Muscular Dystrophy (DMD). It is of great importance to determine whether GDF3 is also a regulator of muscle regeneration in DMD or other types of myopathies, which are most of the time associated with the permanent presence of inflammatory cells, especially macrophages.

**EXPERIMENTAL PROCEDURES**

For more detailed descriptions of experimental procedures, please see Supplemental Experimental Procedures.

**Mice**

Ppargfl/flLy22-cre+ and wild-type C57BL/6J controls, Ppargfl/− Sox2-cre+ and littermate control Ppargfl/− Lyz2-cre− animals, and Gdf3fl/fl and littermate C57BL/6 albino controls were used in the experiments. All experimental procedures conducted on animals were carried out in accordance with institutional regulations.

**Muscle Injury**

Mice were anaesthetized with isoflurane and 50 μl of cardiotoxin (12 × 10−6 mol/l in PBS) was injected in the TA muscle. Muscles were recovered for flow cytometry analysis at day 1, 2, or 4 post-injury or for muscle histology at day 8 post-injury.

**Histological Analysis of Muscle Regeneration**

Muscles were removed and snap frozen in nitrogen-chilled isopentane (~160°C). 8 μm thick cryosections were cut and stained with hematoxylin-eosin (HE). HE stained sections were analyzed for cross sectional area (CSA) or for the presence of phagocytic fibers. Day 8 post CTX slides were also IF stained for Desmin / F4/80 / DAPI.

**Macrophage Cell Culture for Conditioned Medium Generation**

Macrophages were obtained from bone marrow (BM) precursor cells that were cultured in DMEM medium containing 20% FBS and 30% conditioned medium of L929 cell line (enriched in CSF-1) for 7 days. Macrophages were activated with IFN-γ (50 ng/ml) or IL-4 (10 ng/ml) to obtain macrophage-conditioned medium.

**Myogenic Precursor Cell Culture**

Murine MPCs were obtained from TA muscle and cultured using standard conditions in DMEM/F12 (Gibco Life Technologies) containing 20% FBS and 2% Ultroser G (Pall, Inc). For proliferation studies, MPCs were incubated for 1 day with conditioned medium + 2.5% FBS or with 2.5% FBS medium containing GDF3 mouse recombinant protein. Cells were then incubated with anti-kil67 antibodies (15580 Abcam), which were subsequently visualized using cy3-conjugated secondary antibodies (Jackson Immunoresearch, Inc). For differentiation studies, MPCs were incubated for 3 days with conditioned medium containing 2% horse serum or with 2% horse serum medium containing GDF3. Cells were then incubated with anti-desmin antibodies (32362 Abcam), in combination with a cy3-conjugated secondary antibody (Jackson Immunoresearch Inc).

**Phagocytosis Assay**

BMDM cells and C2C12 cells were stained with CellVue or PKH67 (Sigma), respectively. Heat-killed stained C2C12 were used as phagocytic substrates for stained BMDMs and fluorescent intensity was measured with a FACScalibur instrument.

**Image Capture and Analysis for Myoblast Cultures**

Fusion index (for myogenic cells) was calculated as the number of nuclei within myotubes divided by the total number of nuclei, nuclei number being estimated using the ImageJ software.

**Isolation of Macrophages from Muscle**

CD45+ cells were isolated from CTX injected muscles using magnetic sorting (Miltenyi Biotec). CD45+ cells then were labeled with fluorescently labeled
antibodies and Ly6C\(^+\) F4/80\(^{low}\) macrophages, Ly6C\(^-\) F4/80\(^{+}\) macrophages, and Ly6C\(^{md}\) F4/80\(^{+}\) neutrophils were analyzed and sorted with a BD FACSaria III sorter.

**RNA Isolation from Sorted MFs**

Macrophage subsets were sorted from day 1, 2, and 4 post-injury muscles with a FACSaria III sorter and total RNA was isolated with TRIzol reagent according to the manufacturer’s recommendation.

**Microarray Analysis of Muscle Macrophages**

Global expression pattern was analyzed on Affymetrix GeneChip Mouse Gene 1.0 ST arrays. The microarray data are publicly available (GEO: GSE71155).

**Chromatin Immunoprecipitation**

ChIP was carried out in BMDMs using antibodies against pre-immune immunoglobulin G (IgG) (Millipore, 12-370), (pan) RXR (sc-774 Santa Cruz Biotechnology), and PPARγ (Perseus #PP-A3409A).

**Bioinformatic Analysis of the Active Enhancers around the Gdf3 and Angptl4 Locus**

The list of published and/or publicly available datasets used for visualization in IGV2 to identify active enhancers can be found in the Supplemental Experimental Procedures section.

**Western Blotting**

GDF3 protein expression was measured using western blot analysis. Samples from CXT injected TA muscles or CD45\(^+\) cells were lysed in RIPA buffer. GDF3 was targeted using rabbit monoclonal Anti-GDF3 primary antibody (ab109617, Abcam) at 1:1,000 dilution in TBS-T supplemented with 5% BSA overnight at 4°C. Anti-GAPDH mouse monoclonal primary antibody (AM4300, Ambion) was used as a protein loading control at 1:10,000–1:20,000 dilution in TBS-T supplemented with 5% BSA overnight at 4°C.

**RNA Sequencing Library Preparation for Myoblast Gene-Expression Analysis**

cDNA library for RNA-seq was generated from 1 μg total RNA using TruSeq RNA Sample Preparation Kit (Illumina) according to the manufacturer’s protocol. The RNA-Seq data are publicly accessible (SRA: PRJNA290560).

**General Statistical Analyses**

All experiments were performed using at least three different samples. Student’s t-tests and two-way ANOVA analyses were performed and p < 0.05 was considered significant (\(p < 0.05\), \(p < 0.01\), \(p < 0.001\)). Mean and SD values, or mean and SEM values are shown in graphs.

**ACCESSION NUMBERS**

The microarray data are publicly available in the GEO database under accession number GSE71155 and the RNA-Seq data are publicly accessible under SRA: PRJNA290560.

**SUPPLEMENTAL INFORMATION**

Supplemental Experimental Procedures can be found with this article online at http://dx.doi.org/10.1016/j.immuni.2016.10.016.

**AUTHOR CONTRIBUTIONS**


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