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Differential effects of myeloid cell PPAR δ and IL10 in regulating macrophage recruitment, phenotype and regeneration following acute muscle injury.

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Abstract

Changes in macrophage phenotype in injured muscle profoundly influence regeneration. In particular, the shift of macrophages from a pro-inflammatory (M1-biased) phenotype to a pro-regenerative (M2-biased) phenotype characterized by expression of CD206 and CD163 is essential for normal repair. According to the current canonical mechanism regulating for M1/M2 phenotype transition, signaling through PPAR δ is necessary for obtaining the M2-biased phenotype. Our findings confirm that the murine myeloid cell targeted deletion of *Ppard* reduces expression *in vitro* of genes that are activated in M2-biased macrophages; however, the mutation in mice *in vivo* increased numbers of CD206+ M2-biased macrophages and did not reduce the expression of phenotypic markers of M2-biased macrophages in regenerating muscle. Nevertheless, the mutation impaired CCL2-mediated chemotaxis of macrophages and slowed revascularization of injured muscle. In contrast, null mutation of IL10 diminished M2-biased macrophages but produced no defects in muscle revascularization. Our results provide two significant findings. First, they illustrate that mechanisms that regulate macrophage phenotype transitions *in vitro* are not always predictive of mechanisms that are most important *in vivo*. Second, they show that mechanisms that regulate macrophage phenotype transitions differ in different *in vivo* environments.

Keywords

skeletal muscle; inflammation; acute muscle injury; macrophage; PPAR δ ; interleukin-10

Introduction

The inflammatory response to skeletal muscle injury or disease has important influences on the rate and extent of muscle repair. Although the initial population of myeloid cells that enter muscle following injury is dominated by neutrophils that can amplify muscle damage, their numbers rapidly decline and they are replaced by phenotypically-complex populations of macrophages that can remain at elevated concentrations in the regenerating muscle for weeks (1, 2, 3). Over that prolonged period of muscle regeneration, the predominant macrophage phenotypes shift from a population that expresses pro-inflammatory cytokines and increases proliferation of myogenic cells to a population that produces anti-inflammatory cytokines, promotes muscle fiber growth and increases connective tissue production (4, 5, 6, 7, 8, 9). The importance of the shifts in macrophage phenotypes on the course of muscle repair and regeneration has been well-established by investigations that have shown that disrupting the normal recruitment or phenotype specification of macrophages amplifies muscle damage and slows regeneration (10, 11, 12, 13, 14, 15).

The initial characterization of macrophage phenotypes as M1 (pro-inflammatory) or M2 (anti-inflammatory) was based upon changes in gene expression that occurred in macrophages *in vitro* following stimulation with Th1 cytokines (interferon-gamma [IFN γ]; tumor necrosis factor [TNF]) or Th2 cytokines (interleukin-4 [IL4]; IL13) (16, 17, 18). However, in the complex inflammatory environment *in vivo*, numerous other immunomodulatory molecules can influence macrophage activation to produce a broad and continuous spectrum of phenotypes that range between the M1 and M2 designations (19). Whether macrophage activation is biased along the spectrum toward the M1 or M2 phenotype is assessed by determining their expression of numerous phenotypic markers. Within injured and diseased muscles of mice, macrophages that are biased toward the M1 phenotype express inducible nitric oxide synthase (iNOS), CD68 and high levels of CD11b and Ly6C. Intramuscular macrophages biased toward the M2 phenotype express CD163, CD206, arginase-1 (Arg1) and low levels of CD11b and Ly6C (8, 10, 20, 21, 22).

Multiple signaling pathways contribute to activating macrophages to an M2-biased, pro-regenerative phenotype. For example, macrophage activation to a CD163+/CD206+, M2-biased phenotype in inflamed muscle is strongly influenced by IL10, transforming growth factor- β (TGF β) and insulin-like growth factor-1 (IGF1) that are expressed by macrophages at the site of muscle damage (7, 8, 21, 23). Furthermore, diminishing expression of these regulators of M2-biased macrophage phenotype in injured or diseased muscle causes impaired muscle repair and retarded muscle growth (8, 21, 23), which emphasizes the functional importance of intramuscular macrophages biased to an M2-biased phenotype. However, recent discoveries also demonstrated that changes in the relative prevalence of lipid mediators that influence macrophage phenotype change over the course of muscle repair following injury (24). For example, lipid mediators that reduce inflammation (specialized, proresolving lipid mediators; SPMs) are more prevalent in Ly6Clo macrophages and intramuscular injection of one of the SPMs, resolvin D2, produced increases in Ly6Clo cell numbers in injured muscles and improved muscle function (24). Those findings suggest that SPM-induction of an M2-biased phenotype may also be able to improve muscle regeneration.

Although untested, the collective findings of numerous investigations support the hypothesis that lipid mediators may influence the shift of macrophages to an M2-biased phenotype in injured muscle by activating peroxisome proliferator-activated receptor- δ (PPAR δ). PPAR δ is a ligand activated nuclear receptor that is activated by polyunsaturated fatty acids and other lipid mediators that can be produced during breakdown of apoptotic cells that are engulfed by macrophages (25). Activated PPAR δ then dimerizes with retinoic acid receptors (RXR), translocates to the nucleus and activates expression of M2-associated transcripts including *Mrc1* (which encodes CD206), *Arg1*, *Chi3l3*, *Retnla* and *Ppard* (26, 27). The importance of PPAR δ in activating macrophages to an M2-biased phenotype was demonstrated clearly in Kupffer cells in the liver and in adipose tissue macrophages (28, 29) in which the deletion of *Ppard* in LysM expressing cells or in bone marrow derived cells prevented activation of macrophages to an M2-biased phenotype. However, fatty acid ligation of PPAR δ in macrophages can also cause an increase in IL10 expression that requires PPAR δ (25) and induction of IL10 expression is increased by PPAR δ in endothelial cells (30). Those findings show that IL10-mediated signaling and PPAR δ -mediated signaling pathways can interact, suggesting the possibility that IL10 and lipid mediators may affect intramuscular macrophage phenotype through overlapping mechanisms.

In this investigation, we generated a mouse line in which there is a targeted deletion of *Ppard* in LysM expressing myeloid cells to test the hypothesis that activation of macrophages to an M2-biased phenotype in acutely-injured muscle is mediated by PPAR δ . We also assayed whether *Ppard* mutation in macrophages influences macrophage function or affects muscle repair, growth and revascularization following injury. Finally, we assayed whether the same parameters of macrophage activation and muscle repair are similarly influenced by ablation of *Il10* expression to address the question of whether IL10-mediated and PPAR δ -mediated regulation of macrophage phenotype have similar effects on muscle inflammation, repair and regeneration, which would indicate a potential interaction between the two regulatory pathways in acutely injured muscle. The direct comparison of the effects of the two mutations in the same injury model enabled us to demonstrate the distinct roles of myeloid cell PPAR δ and IL10 in regulating macrophage recruitment, phenotype and regeneration following acute muscle injury.

MATERIALS AND METHODS

Ethical approval and animals

All animals were handled according to guidelines provided by the Chancellor's Animal Research Committee at the University of California, Los Angeles. C57BL/6, *Ppard*flox/flox (#005897), LysM-cre (#004781) and *Il10*^{-/-} (#002250) mice were purchased from The Jackson Laboratory (Bar Harbor, ME) and bred in specific pathogen-free vivaria at the University of California, Los Angeles. *Ppar δ* fl/fl and LysM-cre mice were crossed to generate myeloid cell-specific *Ppar δ* mutant mice. Knockdown of *Ppar δ* was confirmed by quantitative real time polymerase chain reaction (QPCR) assays on bone marrow-derived macrophages (BMDMs) from *Ppar δ* fl/fl LysM-cre (*Ppar δ* mutants) and *Ppar δ* fl/fl (*Ppar δ* WT) mice. *Il10* mutation was confirmed by QPCR of skeletal muscle from C57BL6 and *Il10*^{-/-} mice. Mice were housed on a 12/12 hour light/dark cycle and were provided food

and water ad libitum. Following euthanasia by inhalation of isoflurane, muscles were collected, weighed and flash-frozen for histological or biochemical analyses.

Acute muscle injury

Mice were anesthetized with isoflurane inhalation in a chamber (4-5% isoflurane with 100% oxygen) then moved to a nose cone (1-2% isoflurane). Anesthesia was checked by testing mice for a positive reflex response to a hind foot pinch and by monitoring respiration. The lower limb was wiped with 70% ethanol before intramuscular injection. Sterile muscle injury was induced by the intramuscular injection of 50 μ l of a 1.2% barium chloride (BaCl₂) solution into the midbelly of tibialis anterior (TA) muscles of healthy, 4-6 months old male mice. Animals were monitored daily until they recovered.

RNA isolation and QPCR

RNA was isolated from cells and muscle homogenates and electrophoresed on agarose gels and its quality assessed by 28S and 18S ribosomal RNA integrity (31). RNA samples (2 μ g) were reverse transcribed with Super Script Reverse Transcriptase II using oligo dTs to prime extension (Invitrogen, Carlsbad, CA, USA) to produce cDNA. Expression of selected transcripts was assayed using iTaq SYBR green qPCR Supermix according to the manufacturer's protocol (BioRad, Hercules, CA, USA) and a QuantStudio 5 Real-time PCR System (Applied Biosystems, Foster City, CA, USA). Established guidelines for experimental design, data normalization and data analysis for QPCR were used to maximize the rigor of quantifying the relative levels of mRNA (32, 33). We empirically tested reference genes and identified those with the least variability between experimental groups. Based on that analysis, the relative expression of transcripts of interest were normalized to the reference genes *Srp14* and *Rnps1* for muscle regeneration, *Srp14* and *Tpt1* for *Ppar δ* BMDMs, and *Tbp* and *Rplo* for *Il10*^{-/-} BMDMs QPCR assays. The normalization factor for each sample was calculated by geometric averaging of the Ct values of reference genes. Expression for each gene in control samples was set to 1 and the other values were scaled to the control. Primers used for QPCR are listed in Table 1.

Muscle fiber cross-sectional area

Frozen, cross-sections of TA muscles were sectioned at the midbelly and used for fiber cross-sectional area measurements. Sections were stained with hematoxylin and the cross-sectional areas of fibers were measured using a digital imaging system (Bioquant, Nashville, TN, USA). In muscle that had undergone acute injury, only centrally-nucleated, regenerating muscle fibers (34) were measured from the area of the central lesion, identified as the region that was least regenerated, as previously described (35). The classification of small and large fibers was determined by setting three standard deviations from the mean cross-sectional area for the *Ppar δ* WT group at 15-dpi time point and quantifying the proportion of myofibers measured that fell within these ranges (adapted from reference 36). The threshold for small and large fibers at 15-dpi was determined to be less than 719 μ m² or more than 2304 μ m², respectively.

Production of Pax7 antibody

Pax7 hybridoma cells were purchased from Developmental Studies Hybridoma Bank anti-Pax7 (Developmental Studies Hybridoma, Iowa City, Iowa, USA; RRID: 2299243). Cells were cultured and antibody was isolated from the supernatant as previously described (37).

Immunohistochemistry

Muscles were dissected from euthanized mice and then rapidly frozen in liquid nitrogen-cooled isopentane. For identification of myogenic progenitor cells, cross-sections, 10 μm thick were taken from the mid-belly of muscles, air-dried for 30 minutes and then fixed in 4% paraformaldehyde for 10 minutes. Sections were then immersed in antigen retrieval buffer (10 mM sodium citrate, 0.05% Tween 20, pH 6.0) at 95–100° C for 40 minutes. Endogenous peroxidase activity in the tissue was quenched by immersion in 0.3% H_2O_2 . Sections were then treated with blocking buffer from a mouse-on-mouse immunohistochemistry kit (M.O.M kit; Vector Labs, Burlingame, CA, USA) for 1 hour and immunolabeled with affinity purified mouse anti-Pax7 (1:500) antibody overnight at 4° C. Sections were washed with phosphate-buffered saline solution (PBS) and then incubated with biotin-conjugated anti-mouse IgG (1:250) for 30 minutes. Sections were subsequently washed with PBS and then incubated for 30 minutes with ABC reagents from the M.O.M kit. Staining was visualized with the peroxidase substrate 3-amino-9-ethylcarbazole (AEC; Vector), yielding a red reaction product.

For identification of macrophages and endothelial cells, cross-sections 10- μm thick were taken from the mid-belly of muscles and fixed in ice-cold acetone. Endogenous peroxidase activity in the sections was quenched by immersion in 0.3% H_2O_2 . Sections were blocked for 1 hour with blocking buffer (3% bovine serum albumin (BSA), 2% gelatin and 0.05% Tween-20 in 50 mM Tris-HCl pH 7.6 containing 150 mM NaCl). Sections were then incubated with: rat anti-mouse F4/80 (1:100, overnight at 4° C, eBioscience #14-4801, San Diego, CA, USA; RRID: AB2314387), rat anti-mouse CD68 (1:100, 3 hours at room temperature (RT), Biorad #MCA1957, Hercules, CA, USA; RRID: AB322219), rabbit anti-mouse CD163 (1:100, 3 hours at RT, Santa Cruz Biotech #33560, Dallas, TX, USA; RRID: AB2074556), rat anti-mouse CD206 (1:50, 3 hours at RT, Biorad #MCA2235, Hercules, CA, USA; RRID: AB324622), and rat anti-CD31 (BD Pharmingen #01951D, San Diego, CA, USA). The sections were washed with PBS and probed with rabbit anti-rat IgG and goat anti-rabbit IgG (Vector Labs #BA-4001 and #BA-1000, Burlingame, CA, USA). Sections were then washed with PBS and incubated with avidin D-conjugated HRP (1:1000, 30 minutes at RT, Vector Labs, Burlingame, CA, USA). Staining was visualized with 3-amino-9-ethylcarbazole.

Stereology

The number of cells per volume of muscle was determined by measuring the total volume of each section using a stereological, point-counting technique to determine section area and then multiplying that value by the section thickness (10 μm) (38). The numbers of immunolabeled cells in each section were counted and expressed as the number of cells per unit volume of each section. In inflammatory lesions with high concentrations of

macrophages, boundaries of individual cells were identified by using Nomarski optics (resolution ~400 nm) and changing plane of focus in the Z-axis.

Assay for ischemic muscle area

Ischemic areas of muscles were quantified by imaging the entire muscle section and merging the images using Image Composite Editor (Microsoft, Redmond, CA, USA). Bioquant digital imaging software (Nashville, TN, USA) was used to calculate the entire area of the muscle followed by measuring and subtracting areas of the tissue containing artifacts or imperfections (tears or folds). Areas of ischemia that were sparsely occupied by CD31+ endothelial cells were quantified by an investigator blinded to the identity of tissue sections. Data were expressed as the proportion of the ischemic area divided by the total muscle area.

Immunofluorescence

The immunofluorescence double-labeling protocol was similar to macrophage immunohistochemistry described above. Sections from the mid-belly of TA muscles were fixed with 4% PFA for 10 minutes and then blocked with PBS containing 10% horse serum and 0.1% Tween-20 for 60 minutes. Muscle sections were incubated with rat anti-mouse F4/80 (1:100) and goat anti-Ki67 (1:200; Santa Cruz Biotech #sc-7846, Dallas, TX, USA; RRID: AB2142374) or rabbit anti-CCR2 (1:50; Abcam #32144, Cambridge, UK; RRID:AB1603737) overnight at 4° C. Sections were subsequently incubated with a combination of donkey anti-rat IgG Alexa 488 (Abcam #ab102260, Cambridge, UK) and horse anti-goat IgG Dylight 594 (Vector #DI-3094, Burlingame, CA, USA) or horse anti-rabbit IgG Dylight 594 (Vector #DI-1094, Burlingame, CA, USA) and cover-slipped with Prolong Gold anti-fade reagent with DAPI (Invitrogen, Carlsbad, CA, USA).

Assay for muscle fiber damage

Damaged muscle fibers were identified in sections that were blocked in a 1% gelatin solution in PBS for 30 minutes and then labeled with horse anti-mouse IgG Dylight 488 (1:100; Vector Labs #DI-2488, Burlingame, CA, USA) and cover slipped with Fluorogel (Electron Microscopy Sciences, Hatfield, PA, USA). Intracellular labeling of muscle fibers denoted the presence of the extracellular mouse IgG protein in the muscle fiber indicating damage to the muscle fiber membrane. The proportion of a muscle cross sections that contained damaged fibers was determined by using a sampling grid that was superimposed over the tissue after which the number of grid intercepts that overlaid IgG-positive fibers was counted and expressed as a percentage of the total number of intercepts (39).

Arginase activity assay

Arginase activity was measured in TA muscles at 7-dpi and in contralateral, noninjured TA muscles that were homogenized in 5 volumes of 25 mM Tris-HCl (pH 7.4) containing 5 mM MnCl₂, 0.2% Triton X-100 and protease inhibitor cocktail (Sigma Aldrich, St. Louis, MO, USA). Lysates were centrifuged and the supernatant was heated to 56° C for 10 minutes to activate arginase. Substrate hydrolysis was performed by adding 0.5 M arginine, pH 9.7 to the tissue supernatant followed by 1-hour incubation at 37° C. The reaction was stopped by adding H₂SO₄ and H₃PO₄ in water (1:3:6 parts volume/volume). Samples were heated to

100° C for 45 minutes after adding 9% α -isonitrosopropriophenone. Urea content was then measured spectrophotometrically at 540 nm. Values were normalized to protein content of the tissue lysate. Six tissues were assayed at each per condition (adapted from 40).

Preparation of bone marrow-derived macrophages

For preparation of BMDMs, bone marrow cells were aseptically flushed from femurs and tibiae and differentiated using RPMI-1640, 20% heat-inactivated fetal bovine serum (FBS; Omega Scientific, Inc, Tarzana, CA, USA), 100 U/ml penicillin and 100 μ g/ml streptomycin (1% P/S), and 10 ng/ml macrophage colony stimulating factor (M-CSF; Cell Applications Inc., #RP2008, San Diego, CA, USA) *in vitro* to BMDMs (31). BMDMs were stimulated for 24-hours with activation media consisting of Dulbecco's Modified Eagle Medium (DMEM) with 0.25% heat-inactivated FBS, 1% P/S, 10 ng/ml M-CSF and recombinant IL4 (25 ng/ml) with IL13 (10 ng/ml) (BD Pharmingen #550067 and #554599, San Diego, CA, USA), recombinant TNF α (10 ng/ml) with IFN γ (10 ng/ml) (BD Pharmingen #554589 and #554587, San Diego, CA, USA) or recombinant IL10 (10 ng/ml) (BD Pharmingen #550070, San Diego, CA, USA).

Macrophage mobility and chemotaxis

Cell migration and chemotaxis was evaluated using a modified Boyden chemotaxis chamber according to the manufacturer's protocol (Neuro Probe, Gaithersburg, MD, USA). Briefly, the lower chambers were filled with a volume of DMEM with 1% BSA or DMEM with 1% BSA and 100 ng/ml recombinant CCL2 (R&D Systems #479-JE-010, Minneapolis, MN, USA) resulting in a slight positive meniscus over the well. Next, a 5 μ m porous membrane, silicone gasket and the upper plastic chambers were placed over top of the lower chambers. BMDMs were prepared as described above. Cells were dissociated from the plate and resuspended at a concentration of 2×10^6 cells/ml after which 105 cells were loaded into the upper chambers. The filled chambers were incubated for 2 hours in an incubator at 37° C in humidified air with 5% CO₂. After 2 hours, the remaining cells were aspirated from the upper chamber, the chamber was disassembled and the cells on the non-migrated side of the membrane were cleared. The membrane was stained with hematoxylin, mounted to a slide and the number of cells that migrated across the membrane were quantified.

Results

Transcriptional regulation of macrophage activation by myeloid PPAR δ in vitro

We generated myeloid specific *Ppard* mutant mice by crossing *Ppard*^{fl/fl} mice (*Ppard* WT) with lysozyme Cre mice to assess the regulatory roles of PPAR δ in muscle inflammation and repair following acute injury. First, we assayed the influence of *Ppard* mutation on myeloid cell phenotype and function *in vitro* by assaying changes in gene expression in BMDMs. Quantitative PCR (QPCR) analysis confirmed > 90% knockdown of *Ppard* expression in *Ppard* mutant macrophages (Figure 1A). The mutation also resulted in a 93% increase in the expression of the M1-biased transcript *Nos2* while reducing the expression of M2-biased transcripts *Arg1*, *Mrc1* and *Chil3* by 61%, 30%, and 31%, respectively (Figure 1B). These findings generally agree with previous investigations showing that PPAR δ signaling promotes the M2 phenotype in macrophages (28,29).

Because the activation state of macrophages *in vitro* along the M1/M2 spectrum of phenotypes is largely determined by the presence of Th1 or Th2 cytokines (17, 19), we assayed the influence of *Ppard* mutation on changes in gene expression that were induced by cytokines that may be modulated in muscle following acute injury or disease. Stimulation of wild-type (WT) BMDMs with the Th2 cytokines IL4 and IL13 showed the anticipated increases in expression of M2-biased genes (*Arg1*, *Arg2*, *Mrc1*, *Retnla* and *Chil3*), and *Ppard* mutation diminished the increase for each of those transcripts, except for *Arg2* (Figure 1C). These observations are consistent with the previously-reported, *Ppard*-dependent increase in the expression of *Mrc1*, *Retnla* and *Chil3* in BMDMs stimulated with IL4 only (29). Stimulation of WT BMDMs with the Th1 cytokines TNF and IFN γ produced expected increases in some M1-biased genes (*Nos2* and *Tnf*) while reducing expression of some M2-biased genes (*Mrc1*, *Chil3*). Unexpectedly, TNF/IFN γ stimulation of WT BMDMs also increased expression of *Arg1* and *Arg2*. *Ppard* mutation also caused a small increase in *Nos2* expression in TNF/IFN γ stimulated BMDMs and diminished the increased expression of *Arg1* that was caused by stimulation with the Th1 cytokines (Figure 1D). Those observations show that *Ppard* mutation somewhat amplifies the M1-biased phenotype in TNF/IFN γ activated cells. In addition, the findings show that the influence of *Ppard* mutation on the expression of some genes, such as the increase in *Nos2* and reduction in *Arg1*, occurs regardless of the Th1 vs. Th2 cytokine environment. However, reductions in *Mrc1* and *Chil3* expression that were caused by *Ppard* mutation in unstimulated BMDMs or IL4/13 stimulated BMDMs, did not occur in TNF/IFN γ stimulated BMDMs. This suggests that the regulatory role of PPAR δ signaling may vary with the stage of inflammation in injured tissue, as the cytokine environment changes.

Modulation of Th1 and Th2 cytokine gene expression following acute muscle injury

Because *in vitro* observations showed that the response of BMDMs to stimulation with either Th1 or Th2 cytokines *in vitro* was influenced by changes in *Ppard* expression, we assayed whether the expression of those cytokines changed over the time course of muscle repair following acute injury caused by BaCl₂ injection. In addition, IFN γ , TNF and IL4 have all been shown previously to influence the muscle inflammation and repair following acute injury or disease (5, 8, 41, 42). Although changes in the expression of IL4 and IFN γ have been observed following muscle injury by snake toxin (5; 41), the inflammatory responses to muscle injury by snake toxin and BaCl₂ differ (43). QPCR data showed that *Tnf* expression was greatly elevated at 3-days post-injury (dpi), remained elevated at 7-dpi and then declined to basal levels by 15-dpi (Figure 2A), which was consistent with a previous report that BaCl₂-induced injury increases TNF expression in muscle (44). However, we also observed that *Ifng*, *Il4* and *Il13* expression were elevated at 7-dpi, with *Il4* and *Il13* returning to levels that did not differ from non-injured muscle by 15-dpi (Figures 2B-D). Notably, the increase in *Il4* and *Il13* expression at 7-dpi coincided to the time post-injury at which CD206⁺ macrophage populations are elevated (23, 45, 46). Because IL4 and IL13 can increase expression of *Mrc1* (which encodes CD206) through a PPAR δ -mediated mechanism (29), the observations support the expectation that elevations in IL4 and IL13 expression in muscle at 3- to 7-dpi could lead to increased activation of macrophages to the CD206⁺ M2-biased phenotype through a PPAR δ -mediated process.

Myeloid PPAR δ affects resident macrophage phenotype in healthy muscle without affecting muscle mass

We tested whether *Ppard* mutation in myeloid cells affected their numbers or phenotype in healthy muscle *in vivo*. Using antibodies to the pan-macrophage marker F4/80, we found that deletion of *Ppard* in myeloid cells did not affect total numbers of macrophages in healthy, adult muscle (Figures 3A, B, I). However, the mutation yielded significant reductions of CD163+ (38%; Figures 3E, F, I) and CD206+ (55%; Figures 3G, H, I) M2-biased macrophages and we observed a strong trend ($P = 0.06$) for an increase in CD68+ macrophages per volume of muscle (Figure 3C, D, I). The reduction in CD206+ macrophages in *Ppard* mutant mice is consistent with our *in vitro* observations showing that the mutation reduced expression of *Mrc1* (Figure 1). Despite the established role of M2-biased macrophages in promoting muscle growth following injury (10, 14, 22), we did not observe any influence of the reduction of intramuscular CD206+ M2-biased macrophages on the mass in healthy, adult TA muscles (Figures 3J-L).

The influence of myeloid cell deletion of *Ppard* differs between healthy and injured muscles

We tested whether PPAR δ served similar regulatory roles for regulating macrophage phenotype in healthy and injured muscle by assaying numbers of macrophages in muscle that expressed CD68, CD163 or CD206 over the course of muscle inflammation following injury. Contrary to our observations on resident macrophages in healthy muscle, numbers of CD68+ macrophages were significantly reduced at 3-dpi, returning to levels present in non-mutant mice at 7-dpi (Figures 4A, B, C). Also contrary to the effect of the mutation in healthy muscle, numbers of CD163+ macrophages were not affected at any stage of inflammation following muscle injury (Figures 4D, E, F). However, in most distinct contrast to healthy muscle in which the mutation caused large reductions in CD206+ M2-biased macrophages, numbers of CD206+ cells were significantly elevated in mutant mice TAs at 7-dpi (Figures 4G, H, I).

We also observed that the influence of *Ppard* deletion on the expression of transcripts related to the M2-biased phenotype differed greatly in macrophages *in vitro* compared to myeloid cells in injured muscles. For example, *Ppard* mutation in macrophages *in vitro* reduced *Arg1* expression, regardless of the cytokine environment (Figure 1), but the mutation increased *Arg1* expression in inflamed muscle at 7- and 15-dpi (Figure 4J) which was reflected by elevated Arg activity in the injured muscle (Figure 4K). Similarly, *Retnla*, a stereotypical indicator of the M2-biased phenotype, was significantly elevated at 15-dpi (Figure 4L). Thus, ablation of *Ppard* in macrophages within acutely injured muscle increased their expression of transcripts associated with activation of the M2-biased phenotype.

Ppard mutation in myeloid cells impairs macrophage accumulation in injured muscle and reduces chemotaxis

We assayed whether *Ppard* mutation in myeloid cells affected the kinetics of macrophage accumulation in injured muscle by assaying changes in total macrophage numbers, using antibodies to the pan macrophage marker, F4/80. The concentration of F4/80+ cells was substantially reduced in *Ppard* mutant mice at 3-dpi (Figures 5A-C). However, the number of

F4/80+ cells at 7-dpi was significantly greater in injured muscles of mutant mice than in WT muscles and a strong trend ($P = 0.08$) for sustained elevations of F4/80+ cells in muscle persisted through 15-dpi (Figure 5C).

Because previous investigators have shown that *Ppard*-deficient macrophages have a markedly reduced mitogenic response to IL4 stimulation *in vitro* (29), we tested whether the delayed accumulation of macrophages in injured muscles of *Ppard* mutants was attributable to impaired macrophage proliferation *in vivo*. However, the proportion of F4/80+ cells that co-expressed Ki67, a marker of cell proliferation, did not differ between injured muscles of control and mutant mice (Figures 5D, E), indicating that suppression of proliferation did not underlie the slowed kinetics of macrophage accumulation in injured muscles.

Alternatively, the delayed accumulation of macrophages in injured muscles of mutant mice could result from impaired recruitment because several previous investigations have shown that perturbations of CCL2/CCR2 signaling are particularly important in disrupting macrophage recruitment to injured muscle (6, 47-49). We observed that the proportion of F4/80+ cells that expressed detectable levels of CCR2 was significantly reduced in *Ppard* mutant muscles at 3-dpi (Figures 3F-H), suggesting that reductions in recruitment via CCR2 can contribute to the reduced numbers of total macrophages at this stage. In addition, we confirmed that *Ppard* mutation caused large reductions (41%) in *Ccr2* expression by BMDMs *in vitro* (Figure 5I), consistent with previous reports (50). Furthermore, *Ppard* mutant BMDMs exhibited impaired chemotaxis towards CCL2, but did not affect cell mobility independent of the CCR2 ligand (Figure 5J). Finally, we also tested whether expression of either CCL2 or CCR2 differed in mutant and WT muscles at 7-dpi, when numbers of F4/80+ cells in mutant muscles increased, suggesting a recovery in a defect in CCR2-mediated chemotaxis. Although CCL2 expression did not differ between WT and mutants at 7-dpi (Figure 5K), CCR2 expression was significantly elevated and greater in mutant muscles at this stage (Figure 5L). Thus, our data indicate that myeloid cell PPAR δ is important for influencing macrophage recruitment into muscle after acute injury, possibly by regulating CCR2 expression and reducing chemotaxis at early stages post-injury.

Ppard mutation in myeloid cells slows muscle repair and revascularization following acute injury. Intramuscular macrophages can contribute to amplifying muscle damage or promoting muscle repair following injury or disease (3) which led us to examine the influence of *Ppard* mutation in myeloid cells on muscle injury and repair. We assayed muscle injury by measuring the proportion of total muscle cross-sections that contained injured fibers with large membrane lesions that allowed the unregulated influx of IgG (Figures 6A, B), which does not cross the intact cell membrane of healthy muscle fibers. The proportion of the muscle that contained injured fibers exceeded 73% at 3-dpi, and did not differ between WT and mutant mice at this stage (Figure 6C), indicating that PPAR δ signaling is not an essential mediator of macrophage cytotoxicity in this injury model. Furthermore, because the numbers of macrophages in muscles at 3-dpi were greatly reduced (Figure 5C) although muscle injury did not differ between WT and mutant muscles at that stage (Figure 6C), the observations show that the magnitude of the inflammatory response is not an important determinant of the extent of muscle damage at this stage in this injury model. In addition, we observed that the regions of the most extensive muscle damage at 3-

dpi contained few macrophages and regions of muscle fiber repair and regeneration were relatively enriched in macrophages (Figures 6D, E). Collectively, these observations show that neither the magnitude of macrophage accumulation, the localization of macrophages or their expression of PPAR δ is an important determinant of the initial magnitude of muscle damage in this injury model. However, at 7-dpi, the area of injury in WT muscles returned almost to levels of non-injured muscle, although injury in muscles of mutant mice remained greatly elevated (Figure 6C), coinciding with the large elevation of macrophages in the muscle at this stage (Figure 5C). Repair of injury in muscles of mutant mice was complete by 15-dpi (Figure 6C), indicating that *Ppard* mutation slowed the rate of muscle repair.

Previous investigators demonstrated that treatments that reduce macrophage numbers in injured muscles can disrupt revascularization, which was associated with impaired muscle regeneration (6, 48, 51). We tested whether the myeloid cell targeted deletion of *Ppard* impaired revascularization, using immunohistochemistry for vascular endothelial cells that expressed CD31. We found that areas of muscle in which there was extensive muscle fiber damage contained few CD31+ cells, although areas in which regenerative muscle fiber displayed intact membranes were richly-vascularized (Figures 6F, G). We tested whether *Ppard* mutation affected the extent of ischemia in muscle during the repair phase following acute injury by assaying for the proportion of the muscle cross-section that was devoid of CD31+ vascular endothelial cells and found that mutant muscles showed over four-fold greater area of ischemia than WT at 7-dpi (Figures 6H-J). In addition, we assayed levels of expression of the hypoxia-inducible gene *Hif1a* as an independent index of ischemia and oxidative stress and found significantly elevated levels of *Hif1a* at each stage of muscle repair in *Ppard* mutants (Figure 6K).

Ppard mutation in myeloid cells slows myogenesis following acute injury

Macrophage-derived factors can influence the expansion of myogenic satellite cell populations and affect the growth of regenerative muscle fibers following muscle injury or disease (7, 23, 31). We tested whether PPAR δ in myeloid cells played a significant role in these pro-myogenic roles of macrophages by assaying the effect of *Ppard* mutation on the numbers of Pax7+ myogenic cells and the size of regenerative muscle fibers over the course of muscle repair. *Ppard* mutants showed large reductions in the number of Pax7+ cells at 3-dpi which persisted in mutants at 7-dpi (Figures 7A-C). However, by 15-dpi, the numbers of Pax7+ cells in WT and mutant muscles were greatly reduced and did not differ between the two genotypes (Figure 7C). The negative influence of the *Ppard* mutation on myogenesis also manifests as a reduction in the cross-sectional area of central-nucleated, regenerative fibers (Figures 7D-F), which was 23% less in mutant mice at 15-dpi. In addition, the mutation produced a significant reduction in the proportion of large diameter muscle fibers (> 2304 μm^2) and a nearly significant increase ($p = 0.06$) in small diameter fibers (Figure 7G).

IL10 is a greater effector of macrophage activation to a CD163+/CD206+ M2-biased phenotype in injured muscle than PPAR δ

Our unexpected finding that *Ppard* mutation did not diminish macrophage activation to a CD163+/CD206+ M2-biased phenotype in injured muscle led us to assay for other effector

molecules that activate the CD163⁺/CD206⁺ phenotype in injured muscle. Our previous observation that ablation of *Il10* in mice experiencing increased muscle use greatly reduced CD163⁺ M2-biased macrophages while increasing CD68⁺ macrophages (21) implicated *IL10* as a significant modulator of intramuscular macrophage phenotype. In the present investigation, we showed that IL10 expression is greatly elevated in muscles following acute injury, reaching expression levels at 3-dpi that are more than 150-times the expression in uninjured muscle (Figure 8A). We also confirmed that IL10 is a strong activator of the M2-biased phenotype in BMDMs *in vitro*, causing large increases in the expression of *Arg1*, *Arg2* and *Mrc1* although *Cd163* expression was not affected (Figure 8B).

In contrast to the effects of *Ppard* gene ablation in myeloid cells, *Il10* mutation greatly influenced macrophage phenotype without affecting macrophage accumulation during repair of injured muscle. Assays for changes in F4/80⁺ total macrophages showed that *Il10* mutation did not affect the numbers of macrophages or the time course of their accumulation in injured muscle (Figures 9A-C), which indicates that expansion and recruitment of macrophage populations were unaffected by the mutation. However, the mutation caused significant increases in the numbers of CD68⁺ macrophages at 15-dpi (Figures 9D-F) and caused large reductions in numbers of CD163⁺ cells at earlier stages of repair (Figures 9G-I). The most prominent treatment effect that we observed was the greatly reduced numbers of CD206⁺ macrophages that occurred at every stage of repair (Figure 9J-L), showing that IL10-mediated signaling rather than PPAR δ signaling is an important regulator of the CD206⁺ M2-biased phenotype in injured muscle. Preventing expansion of CD206⁺ macrophages was not accompanied by a reduction in either *Arg1* or *Arg2* in injured muscles of *Il10* mutant mice (Figures 9M, N), in contrast to the tremendous induction of *Arg* expression in IL10-stimulated BMDMs (Figure 8B).

Because our observations of the effects of *Ppard* mutation on muscle repair showed PPAR δ played a significant role in regulating the expansion of satellite cell numbers, reducing the extent of muscle damage and revascularization of muscle, we also assayed whether IL10 contributed to these features of muscle repair. Unlike the effects of *Ppard* mutation, *Il10* mutation had no effect on the time course or magnitude of expansion of satellite cell populations following injury (Figures 9O-Q). Also unlike *Ppard* mutants in which the initial magnitude of muscle damage was unaffected by the mutation, the *Il10* mutants experienced a small decrease in the proportion of the muscle that showed damage at 3-dpi (Figures 9R-T). However, at 15-dpi, *Il10* mutants had more muscle damage than WT mice (Figure 9T). Finally, in further contrast to *Ppard* mutants, neither the extent of revascularization of muscle at 7-dpi (Figures 9U-W) or the levels of *Hif1* expression, was affected by *Il10* mutation (Figure 9X).

Discussion

Landmark discoveries by previous investigators demonstrated the importance of *Ppard* expression and activation in regulating the phenotype of resident macrophages in liver and adipose tissue *in vivo* and in BMDMs *in vitro* (28, 29). In particular, the reduced expression of markers of M2-biased macrophage activation (e.g., *Arg1*, *Mrc1* (or CD206), *Retnla*, *Chi3l3*, *Il10*) in myeloid targeted *Ppard* mutants in these models (28, 29) served as the

foundation for the broad view that PPAR δ -mediated signaling provides a crucial control for the switch of macrophages from a pro-inflammatory, M1-biased phenotype to an anti-inflammatory, M2-biased phenotype (52). The present investigation shows that this regulatory function for PPAR δ in myeloid cells does not exist in inflamed, regenerating muscle following acute injury. Conversely, we observed that the myeloid-cell-specific ablation of *Ppard* caused either increases or no change in the numbers of CD206+ macrophages over the course of muscle inflammation and caused either an increase or no change in the expression of phenotypic markers of M2-biased macrophages. These findings show that regulation of macrophage phenotype *in vivo* differs with the tissue environment in which phenotype switching occurs.

Our findings also confirm that regulatory controls of macrophage phenotype that are apparent *in vitro* are not necessarily predictive of controls that are most important *in vivo*, as emphasized previously by luminaries in the field of macrophage biology (19). Although our *in vivo* observations showed that myeloid cell targeted ablation of *Ppard* enhanced expression of markers of the M2-biased phenotype in injured muscle, our *in vitro* observations showed that the same mutation in BMDMs *in vitro* diminished their expression of M2-biased phenotypic markers. In addition, the mutation diminished the elevated expression of M2-biased phenotypic markers in BMDMs activated with the Th2 cytokines IL4 and IL13, as reported previously (29).

The distinct differences between the roles of PPAR δ in controlling macrophage phenotype *in vivo* may reflect differences in the tissue in which activation occurred (i.e. fat or liver vs. skeletal muscle); however, our observations indicate that tissue type is not the most important variable for determining the influence of PPAR δ on macrophage phenotype. Contrary to our findings in injured muscle, in which the myeloid-targeted mutation of *Ppard* increased CD206+ M2-biased macrophage number, the mutation reduced CD206+ resident macrophages in healthy, non-injured muscle. This effect of *Ppard* mutation on resident macrophages in healthy muscle resembles the influence of *Ppard* ablation in adipose tissue macrophages in healthy mice (29). Together, the observations suggest that differences between the regulatory roles of *Ppard* on macrophages in injured muscle vs. liver or adipose tissue may be attributable, in part, to differences in phenotype regulation in injured vs. non-injured tissue, rather than differences in tissue type. In contrast to the evidence that *Ppard* mutation does not prevent macrophage activation to an M2-biased phenotype in injured muscle, we observed that *Il10* mutation produced large reductions in the numbers of CD206+ and CD163+ macrophages over the entire course of muscle regeneration that we studied. This is consistent with previous observations that IL10 plays a significant role in macrophage phenotype transitions in muscles of dystrophic mice and mice experiencing modified muscle loading (8, 21) and that treatment of infected muscles with exogenous IL10 increases the proportion of macrophages that express high levels of CD206 (53). We also observed that the rate of repair of the damaged muscle was impaired in *Il10* mutant mice, which is consistent with previous findings that showed that diminishing macrophage phenotype transitions during muscle regeneration increased damage or slowed repair (7, 8, 21, 23). Thus, our findings collectively show that IL10 contributes significantly to the shift of macrophages to a CD206+, CD163+, M2-biased phenotype that is independent of PPAR δ signaling and accelerates muscle repair.

Although our findings show that IL10 expression is mandatory for normal acquisition of a CD206+ M2-biased phenotype in acutely injured muscle and PPAR δ is not required, it is feasible that PPAR δ can contribute to acquisition of CD163+/CD206+ M2-biased phenotype in injured muscle, but that function is assumed by other molecules in mutant mice. Many other molecules could potentially serve that function, including other members of the PPAR family. For example, PPAR γ has a well-established role in regulating macrophage function (54), although it is not essential for development of the macrophage lineage (55). Although numerous findings indicate that PPAR γ -mediated signaling can promote an M2-biased phenotype in macrophages (56-60), other findings indicate that inhibition or loss of PPAR γ can bias macrophages to a CD163+/CD206+ macrophage phenotype (61). These different outcomes may reflect differences in the environment in which macrophage activation occurred, and suggests that the role of PPAR γ signaling in regulating macrophage phenotype may also vary *in vivo* according to the tissue and inflammatory context. However, there is no current evidence to show a role for PPAR γ in regulating macrophage phenotype in injured muscle. Specifically, in skeletal muscle injured by snake toxin, ablation of PPAR γ in myeloid cells did not affect the proportion of Ly6c+ macrophages (associated with an M1-biased phenotype) relative to Ly6c-macrophages (associated with an M2-biased phenotype) (62), in skeletal muscle injured by snake toxin.

Although mutation of *Ppard* in myeloid lineage cells did not prevent activation of macrophages to a CD206+ M2-biased phenotype in injured muscle, our data indicate that the mutation affected the kinetics of the inflammatory process. In particular, the amplification of F4/80+ macrophage numbers in injured muscles of mutant mice was delayed and was accompanied by reductions in the expression of CCR2 by intramuscular macrophages. Further, we observed an impaired chemotactic response of mutant macrophages to CCL2 *in vitro*. Because discoveries by other investigators showed that signaling via CCL2/CCR2 is tremendously important in promoting the traffic of macrophages into diseased or injured tissue (6, 47, 48, 63, 64), disrupting that signaling pathway by *Ppard* mutation would affect the time course and magnitude of inflammation. However, elevations in PPAR δ signaling can also decrease CCL2/CCR2-mediated chemotaxis, at least in other tissue types and states of tissue damage. For example, activation of PPAR δ by agonists suppresses atherogenic inflammation and atherosclerosis, which was attributed to the ability of PPAR δ to repress the expression of CCL2 (65, 66) and thereby reduce transendothelial migration of monocytes, at least *in vitro* (65). These unpredictable relationships between changes in PPAR δ signaling and signaling via CCL2/CCR2 and chemotaxis can be explained in part by opposing influences of PPAR δ on CCL2 expression. Both pharmacological activation and genetic deletion of PPAR δ can inhibit CCL2 promoter activity (66, 67), which indicates that any perturbation of *Ppard* expression or activity has the potential to reduce CCL2/CCR2-driven chemotaxis.

Notably, CCL2/CCR2 signaling does not only influence macrophage chemotaxis; it can also influence macrophage polarization. For example, disruption of CCL2/CCR2 signaling in human macrophages *in vitro* impairs their activation to a CD163+ M2-biased phenotype (68) and direct activation of human monocytes with CCL2 shifts them to a CD206+ M2-biased phenotype (69). However, in contrast with those observations, other investigators report that CCL2/CCR2 signaling in macrophages suppresses the M2 phenotype and promotes the M1

phenotype. For example, ablation of CCR2 in mice on a high-fat diet increased expression of M2 phenotypic markers in macrophages (70). Those observations suggest that the elevated numbers of CD206+ M2-biased macrophages that we observed in *Ppard* muscles at 7-dpi could reflect reductions of CCR2-mediated regulation of macrophage phenotype.

Because we found that *Ppard* mutation in myeloid cells produced either an increase or no change in numbers of macrophages that expressed markers of the M2-biased, pro-regenerative phenotype, we were surprised that the mutation greatly slowed muscle repair between 3 and 15-dpi. However, we observed that the mutation affected the distribution of macrophages in the injured muscles and that macrophages were more broadly distributed in injured WT muscles than in injured mutant muscles. In addition, macrophages were located in areas of regeneration and absent in areas of necrosis in both genotypes. These observations are consistent with a defect in macrophage recruitment to sites of muscle injury in the mutant mice, as a consequence of reduced expression of CCR2. Associated with the reduced dispersal of macrophages in the injured muscle of mutant mice, we found significantly larger areas of muscle lacking vascularization. Although we cannot definitively address the cause of this defect, the failure to recruit pro-regenerative macrophages to the sites of muscle damage may contribute. Previous investigators have established that macrophages play a significant role in affecting muscle revascularization following injury (71) and that reduction in numbers of macrophages in regenerative muscle is associated with impaired revascularization and slowed repair of injured muscle (6, 48, 51). This is similar to the outcomes that we observed. Interestingly, the systemic delivery of PPAR δ agonists to experimental animals increases angiogenesis in synthetic implants *in vivo* (72) and in skeletal muscle (73). Because endothelial cells express PPAR δ and signaling through PPAR δ increases proliferation of endothelial cells (72, 74), the pro-vascular effects of PPAR δ signaling were partially attributed to more rapid expansion of endothelial cell populations. However, our observations suggest the possibility that part of the increased angiogenesis in experimental animals treated with PPAR δ agonists may result from an increase in PPAR δ signaling in myeloid cells, in addition to endothelial cells.

Collectively, our findings indicate that PPAR δ signaling in myeloid cells and IL10-mediated signaling both influence muscle inflammation and improve muscle repair following acute injury. However, their effects appear to be mediated by distinct influences on the inflammatory process, in which PPAR δ signaling affects the chemotactic response of monocytes and macrophages to the injured muscle and IL10 signaling influences the shift in macrophages to an M2-biased phenotype, reflected by the numbers of cells that expressed the phenotypic markers CD206 and CD163. In addition to contributing to our understanding of processes that regulate muscle inflammation and repair following acute injury, these findings illustrate that the transcriptional control of macrophage phenotype switching *in vivo* varies with tissue type and is not always reliably predicted by *in vitro* findings.

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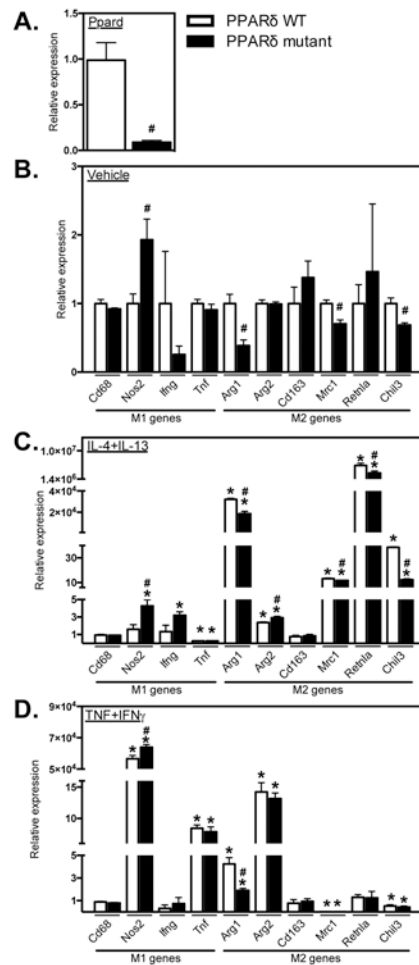
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Key points:

1. Macrophage deletion of PPAR δ reduces inflammation/revascularization of injured muscle
2. PPAR δ expression is not required for M2 macrophage activation in injured muscle
3. IL10 is required for M2 macrophage activation after sterile muscle injury

**Figure 1:**

Ppard mutation reduced expression of M2-biased transcripts and promoted an M1-biased phenotype. A: QPCR data showing reduced *Ppard* expression in BMDMs from *Ppard* mutant mice compared to *Ppard* WT mice. B: Expression of the M1-biased transcript *Nos2* was increased and M2-biased transcripts *Arg1*, *Mrc1* and *Chil3* were reduced in mutant BMDMs compared to WT in control media. C: With M2 activation (IL4 + IL13), *Tnf* expression was reduced and *Arg1*, *Arg2*, *Mrc1*, *Retnla* and *Chil3* expression were increased compared to control media in *Ppard* WT BMDMs. Induction of M2-associated transcripts *Arg1*, *Mrc1*, *Retnla* and *Chil3* was attenuated in mutant BMDMs. *Nos2* and *Arg2* expression were increased in M2-activated mutant BMDMs compared to WT. D: M1 activation (TNF α + IFN γ) increased the expression of *Nos2*, *Tnf* and had differential effects on M2-biased transcripts increasing *Arg1*, *Arg2* and reducing *Mrc1* and *Chil3* expression in WT BMDMs. With M1 activation, *Nos2* expression was increased and *Arg1* expression was attenuated in mutant BMDMs compared to WT. Values in each data set were normalized to *Ppard* WT BMDMs in vehicle control media, set at 1. * indicates significant difference ($P < 0.05$) from control BMDMs within same genotype. # indicates significant difference ($P < 0.05$) versus treatment-matched *Ppard* WT BMDMs. P-values based on two-tailed t-test. $N = 3-4$ for each data set. Data are presented as mean \pm SEM.

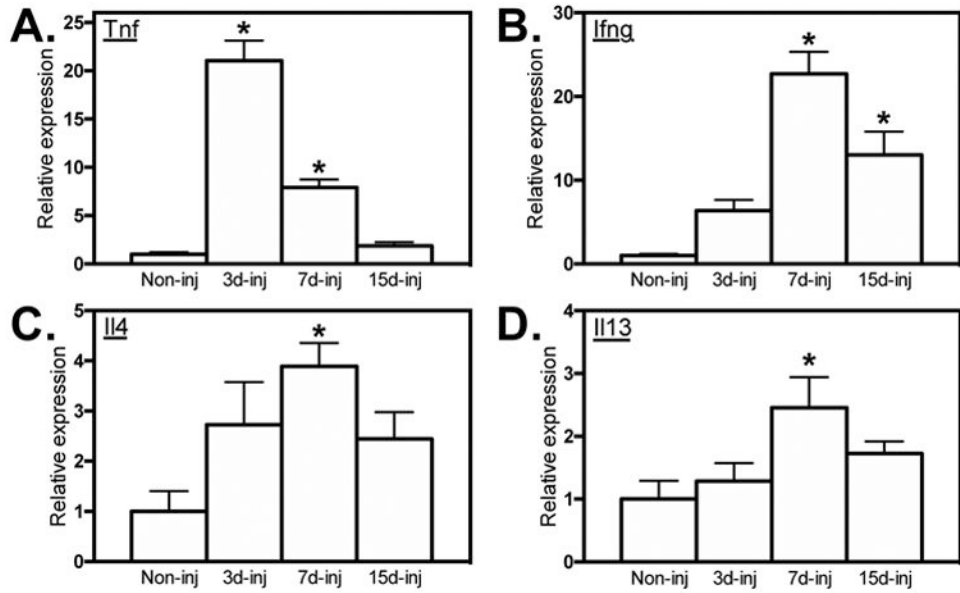


Figure 2. Elevated cytokine expression in skeletal muscle following acute injury. QPCR analysis showed elevated expression of *Tnf*, (A), *Ifng* (B), *Il4* (C) and *Il13* (D) in the TA muscles 3-, 7- and 15-dpi relative to non-injured muscle. All values were normalized to non-injured WT TA muscle and set at 1. * indicates significant difference ($P < 0.05$) from non-injured muscle. P-values based ANOVA with Tukey’s multiple comparison test. $N = 4-5$ for each data set. Data are presented as mean \pm SEM.

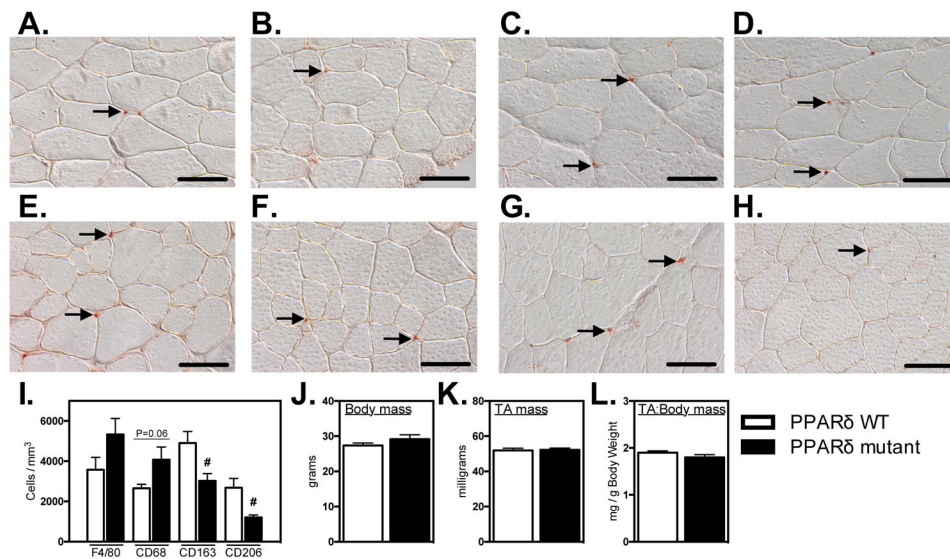
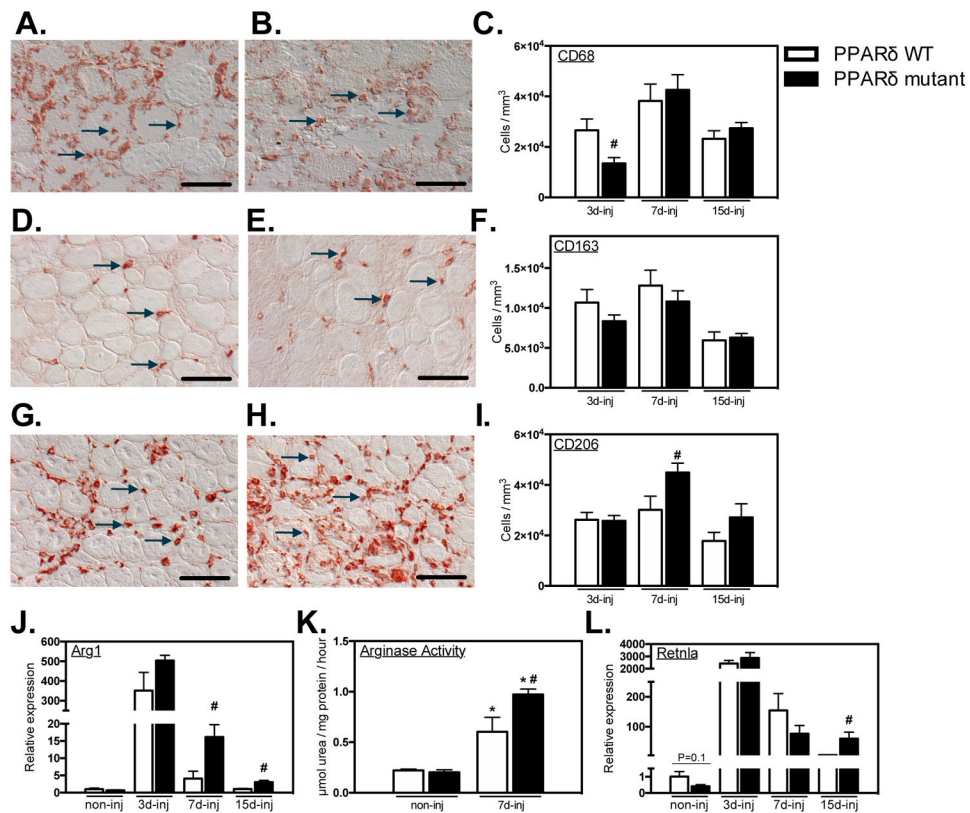


Figure 3.

Ablation of *Ppard* in myeloid cells reduces M2-biased macrophages in healthy muscle without affecting muscle mass. A-H: Cross-sections of non-injured *Ppard* WT (A, C, E, G) and mutant (B, D, F, H) TA muscles were immunolabeled with anti-F4/80 (A, B), anti-CD68 (C, D), anti-CD163 (E, F) or anti-CD206 (G, H). Representative, positively labeled cells are indicated with arrows. Scale bars = 50 μm. I: Quantification of numbers of immunolabeled cells per sectioned muscle volume show that CD163+ and CD206+ cells were reduced in muscles of mutants. J-L: Myeloid *Ppard* deficiency did not affect body mass (J), TA muscle mass (K) or TA muscle mass-to-body mass ratio (L). # indicates significant difference ($P < 0.05$) from *Ppard* WT. P-values based on two-tailed t-test. N = 4-5 for each data set. Data are presented as mean ± SEM.

**Figure 4.**

Ablation of *Ppard* in myeloid cells shifts macrophages to an M2-biased phenotype in injured muscle. A- I: Representative images and quantification of intramuscular macrophages after injury. Bars = 50 μ m. A, B: CD68⁺ macrophages in PPAR δ WT (A) and PPAR δ mutant (B) muscles at 3-dpi. Representative CD68⁺ cells are indicated by arrows. C: Numbers of CD68⁺ cells per volume of sectioned tissue at 3-, 7- and 15-dpi. D, E: CD163⁺ macrophages in PPAR δ WT (D) and PPAR δ mutant (E) muscles at 7-dpi. Representative CD163⁺ cells are indicated by arrows. F: Quantification of numbers of CD163⁺ cells per volume of section tissue at 3-, 7- and 15-dpi. G, H: CD206⁺ macrophages in PPAR δ WT (G) and PPAR δ mutant (H) muscles at 7-dpi. Representative CD206⁺ cells are indicated by arrows. I: Numbers of CD206⁺ cells per volume of sectioned tissue at 3-, 7- and 15-dpi. J: QPCR analysis revealed increased expression of M2-biased transcript *Arg1* in muscle of mutant mice after injury. K: Elevated Arg enzymatic activity was detected in *Ppard* WT and mutant TA muscles 7-dpi compared to non-injured contralateral control muscle. Arg activity was greater in mutant injured TA muscles compared to WT. L: QPCR analysis for expression of the M2-biased transcript *Retnla* in muscle of mutant mice after injury. All values for QPCR assays were normalized to non-injured *Ppard* WT TA muscle and set at 1. # indicates significant difference ($P < 0.05$) from WT within a time point. P-values based on two-tailed t-test. $N = 4-6$ for each data set. For Arg activity assay, * indicates significant difference ($P < 0.05$) from non-injured muscle of the same genotype. # indicates significant difference ($P < 0.05$) from PPAR δ WT within a time point. P-values based ANOVA with Tukey's multiple comparison test. $N = 4-5$ for each data set. Data are presented as mean \pm SEM.

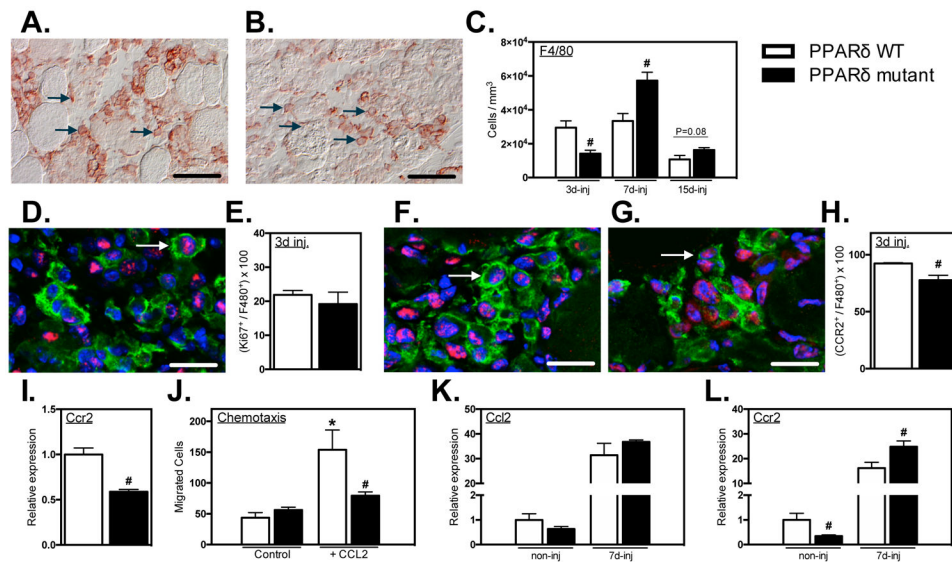


Figure 5.

Myeloid PPAR δ mutation delayed macrophage accumulation in muscle after injury, regulated CCR2 expression and affected CCL2-mediated chemotaxis. A-C: Cross-sections of TA muscles 3-, 7- and 15-dpi were immunolabeled with anti-F4/80. Representative images of F4/80+ macrophages in PPAR δ WT (A) and PPAR δ mutant (B) muscles at 3-dpi. Representative F4/80+ cells are indicated by arrows. C: Numbers of F4/80+ cells per volume of sectioned tissue at 3-, 7- and 15-dpi. D: Cross-section of TA muscles 3-dpi from *Ppard* WT and mutant mice were immunolabeled with anti-F4/80 (green) and anti-Ki67 (red). Nuclei stained blue with DAPI. Arrow indicates an example of an F4/80+ cell with nuclear Ki67. E: Proportion of F4/80+ cells co-expressing nuclear Ki67 in PPAR δ WT and PPAR δ mutant muscles. F, G: Muscle sections were co-labeled with anti-F4/80 (green) and anti-CCR2 (red). Arrows indicate examples of F4/80+ cells that are CCR2+ at 3-dpi in PPAR δ WT and PPAR δ mutant muscles. Bars = 10 μ m. H: Numbers of F4/80+ cells that were CCR2+ per volume of sectioned tissue at 3-dpi in PPAR δ WT (F) and PPAR δ mutant (G) muscles. H: Proportion of F4/80+ cells that were CCR2+ at 3-dpi in PPAR δ WT and PPAR δ mutant muscles. I: QPCR analysis of BMDMs demonstrated that *Ppard* mutation reduced *Ccr2* expression. Values in each data set were normalized to *Ppard* WT BMDMs and set at 1. # indicates significant difference ($P < 0.05$) from *Ppard* WT within a time point. P-values based on two-tailed t-test. N = 4-5 for each data set. J: A chemotaxis assay was used to assess whether PPAR δ affects macrophage mobility or chemotaxis. CCL2 increased chemotaxis but the *Ppard* mutation completely abrogated the chemotactic effect. Macrophage migration was unaffected by PPAR δ in the absence of CCL2. * indicates significant difference ($P < 0.05$) from nontreated BMDMs within same genotype. # indicates significant difference ($P < 0.05$) from *Ppard* WT within a treatment group. P-values based ANOVA with Tukey's multiple comparison test. N = 6 for each data set. Data are presented as mean \pm SEM. K, L: QPCR analysis of non-injured or injured muscle at 7-dpi showed that *Ppard* mutation did not affect *Ccl2* expression (K) but decreased *Ccr2* expression in non-injured muscle and increased *Ccr2* expression at 7-dpi (L). Values in each data set were normalized to *Ppard* WT BMDMs and set at 1. # indicates significant difference ($P < 0.05$)

from *Ppard*^{WT} with same treatment conditions. P-values based on two-tailed t-test. N = 5 for each data set.

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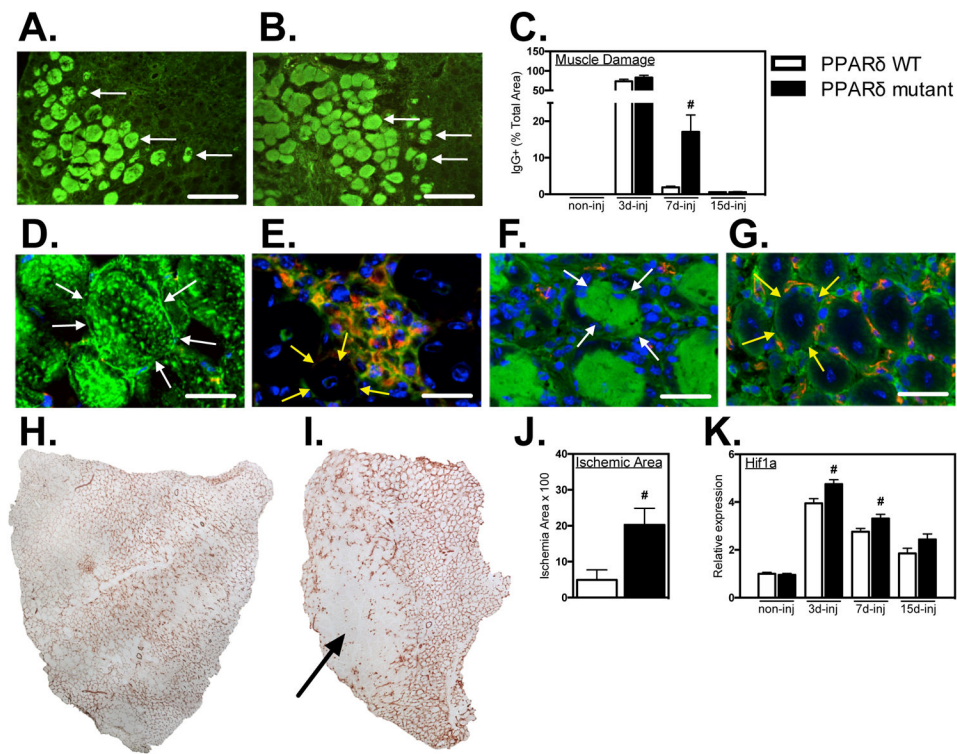


Figure 6. Myeloid *Ppard* mutation impairs muscle repair and revascularization following injury. A, B: Representative cross-sections of TA muscles were immunolabeled with anti-mouse IgG (green) to assay for sarcolemma damage at 7-dpi in PPAR δ WT and PPAR δ mutant muscles. Arrows indicate examples of IgG⁺ muscle fibers. Bars = 100 μ m. C: The volume fractions of TA muscles occupied by IgG⁺ fibers were quantified at 3-, 7- and 15-dpi. # indicates significant difference ($P < 0.05$) from WT within a time point. P-values based on two-tailed t-test. $N = 4-6$ for each data set. D-G: Muscle cross-sections at 7-dpi were co-labeled with anti-mouse IgG (green) and anti-CD68 (red) (D, E) or anti-mouse IgG (green) and anti-CD31 (red) (F, G). Bars = 15 μ m. Both CD68⁺ and CD31⁺ cells were present in low concentrations in areas marked by muscle fiber damage (D, F) and in high concentrations in areas of regeneration containing centrally-nucleated muscle fibers (E, G). Arrows outline the surface of representative, IgG⁺, injured fibers (white arrows; D, F) or IgG⁻, central-nucleated, regenerative fibers (yellow arrows; E, G). H, I: Representative images of whole TA muscle cross-sections immunolabeled for CD31 in *Ppard* WT (H) and mutants (I) at 7-dpi. Arrow indicates ischemic region of muscle sparsely populated by CD31⁺ cells. J: Quantification of the volume fraction of the muscle sparsely occupied by CD31⁺ cells at 7-dpi. # indicates significant difference ($P < 0.05$) from *Ppard* WT. P-values based on two-tailed t-test. $N = 5$ for each data set. K: QPCR assays showed elevated expression of the hypoxia inducible transcript *Hif1a* in mutant muscle 3- and 7-dpi compared to WT. Values were normalized to non-injured *Ppard* WT TA muscle and set at 1. # indicates significant difference ($P < 0.05$) from *Ppard* WT within a time point. P-values based on two-tailed t-test. $N = 4-6$ for each data set. Data are presented as mean \pm SEM.

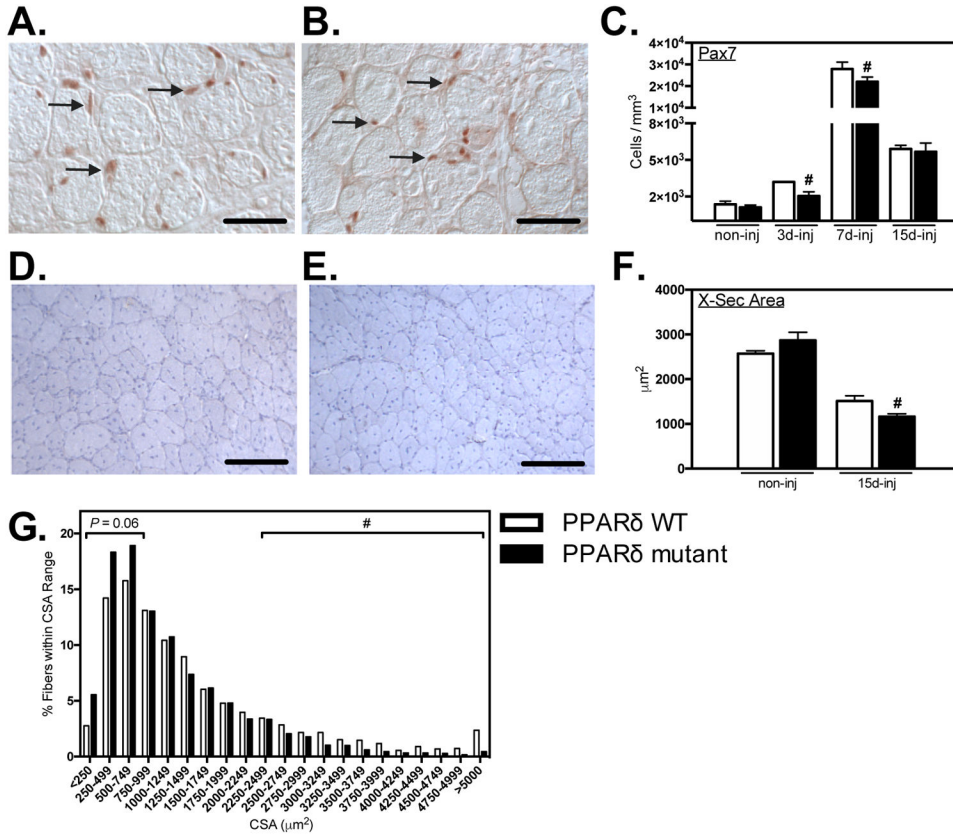
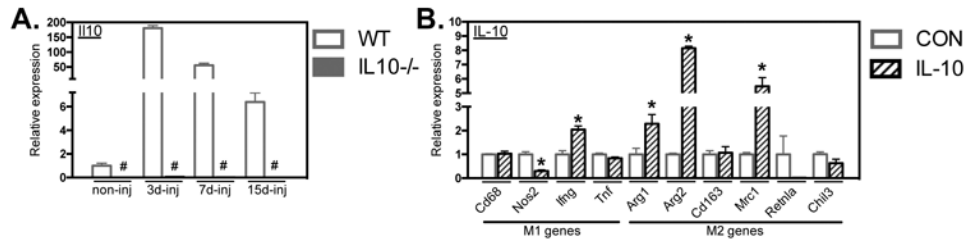


Figure 7. Myeloid *Ppard* mutation delayed expansion of satellite cell populations and reduced the size of regenerative fibers in injured muscle. A, B: Representative images of muscle cross-sections from PPARδ WT (A) and PPARδ mutant (B) muscles at 7-dpi immunolabeled with anti-Pax7. Arrows indicate examples of Pax7+ cells. Bars = 50 µm. C: Numbers of Pax7+ cells per volume of sectioned tissue at 3-, 7- and 15-dpi. # indicates significant difference ($P < 0.05$) from *Ppard* WT at a time point. P-values based on two-tailed t-test. $N = 4-6$ for each data set. Representative images of cross-sections of TA muscles from *Ppard* WT (D) and mutants (E) 15-dpi stained with hematoxylin to assay cross-sectional area of non-injured muscles and muscles 15-dpi. Bars = 100 µm. F: Cross-sectional areas of centrally-nucleated regenerating muscle fibers at 3-, 7- and 15-dpi. # indicates significant difference ($P < 0.05$) from *Ppard* WT within a time point. P-values based on two-tailed t-test. $N = 5-6$ for each data set. G: Frequency distribution of fiber cross-sectional areas for TA muscles from *Ppard* WT and mutants 15-dpi. $N = 5-6$ for each data set. # indicates a significant ($P < 0.05$) reduction in the proportion of large muscle fibers from mutant muscle 15-dpi compared to WT. Small ($< 719 \mu\text{m}^2$) and large fibers ($> 2304 \mu\text{m}^2$) were 3 standard deviations below and above the average cross-sectional area of *Ppard* WT mice 15-dpi. Data are presented as mean \pm SEM.

**Figure 8.**

IL10 stimulation shifted BMDMs to an M2-biased phenotype *in vitro*. A: QPCR data showed induction of *I10* gene expression in WT muscle after injury. *I10* expression was ablated in non-injured and injured muscle 3-, 7- and 15-dpi from *I10*^{-/-} mice. Values were normalized to non-injured WT TA muscle. # indicates significant difference ($P < 0.05$) from WT at a time point. P-values based on two-tailed t-test. N = 4-5 for each data set. B: QPCR data showing that WT BMDMs stimulated with IL10 reduced iNOS and increased *Irfg*, *Arg1*, *Arg2* and *Mrc1* expression. All values were normalized to non-stimulated (CON) WT BMDMs and set at 1. # indicates significant difference ($P < 0.05$) from CON BMDMs. P-values based on two-tailed t-test. N = 3-4 for each data set. Data are presented as mean \pm SEM.

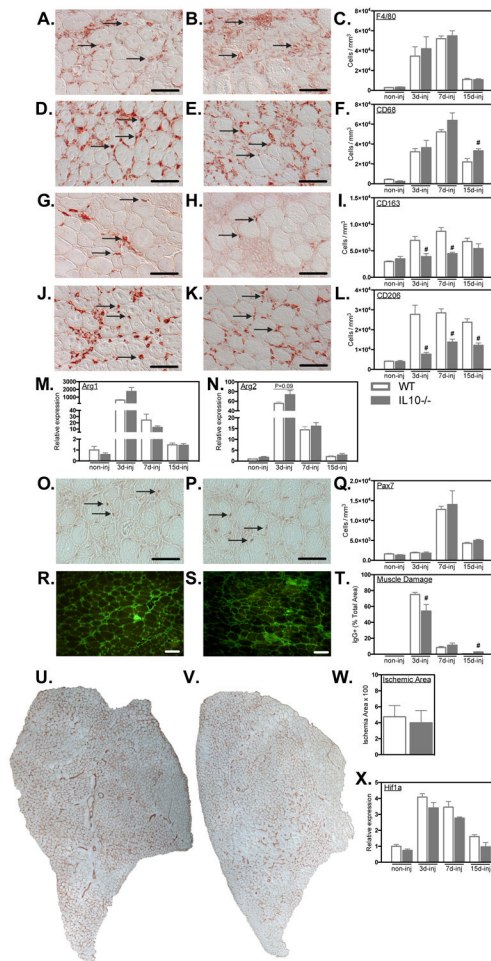


Figure 9.

Il10 mutation attenuated the expansion of M2-biased macrophages in injured muscle affecting muscle injury and repair. A-L: Cross-sections of WT and *Il10*^{-/-} TA muscles were immunolabeled with anti- F4/80 (A, B), anti-CD68 (D, E), anti-CD163 (G, H) and anti-CD206 (J, K). Representative images from WT (A, D, G, J) and mutant (B, E, H, K) muscles at either 7-dpi. Arrows indicate examples of antibody-labeled leukocytes. Bars = 50 μ m. Numbers of F4/80+ (C), CD68+ (F), CD163+ (I) or CD206+ (L) cells per volume of sectioned tissue at 3-, 7- and 15-dpi are summarized in the histograms. M, N: QPCR data showed no change in the induction of *Arg1* and *Arg2* expression in WT and *Il10*^{-/-} TA muscles after injury. O, P: Representative images of muscle cross-sections from WT (O) and *Il10* mutant (P) muscles at 7-dpi immunolabeled with anti-Pax7. Arrows indicate examples of Pax7+ cells. Bars = 50 μ m. Q: Numbers of Pax7+ cells per volume of sectioned tissue at 3-, 7- and 15-dpi. R, S: Representative cross-sections of TA muscles were immunolabeled with anti-mouse IgG (green) to assay for sarcolemma damage at 15-dpi in WT and *Il10* mutant muscles. Bars = 100 μ m. T: The volume fractions of TA muscles occupied by IgG+ fibers were quantified at 3-, 7- and 15-dpi. # indicates significant difference ($P < 0.05$) from WT within a time point. P-values based on two-tailed t-test. N = 4-6 for each data set. U, V: Representative images of whole TA muscle cross-sections at 7-dpi immunolabeled for CD31 in WT (U) and *Il10* mutants (V). W: Quantification of the volume fraction of the muscle

sparsely occupied by CD31+ cells at 7-dpi. X: QPCR analysis of *Hif1a* transcript expression in WT and *I10* mutant muscles at 3-, 7-, and 15-dpi. # indicates significant difference ($P < 0.05$) from WT within a time point. P-values based on two-tailed t-test. N = 4-5 for each data set. Data are presented as mean \pm SEM.

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Table 1:

Primer sequences used for QPCR

Gene	Forward	Reverse
Arg1	CAATGAAGAGCTGGCTGGTGT	GTGTGAGCATCCACCCAAATG
Arg2	GAAGTGGTTAGTAGAGCTGTGTC	GGTGAGAGGTGTATTAATGTCCG
Ccr2	CCTGTAAATGCCATGCAAGTTC	GTATGCCGTGGATGAACTGAG
Cd68	CAAAGCTTCTGCTGTGGAAAT	GACTGGTCACGGTTGCAAG
Cd163	GCAAAAAGTGGCAGTGGG	GTCAAAATCACAGACGGAG
Chil3	GGGCATACCTTTATCCTGAG	CCACTGAAGTCATCCATGTC
Hif1a	GCTTACACACAGAAATGGCCC	CCTCCACGTGCTGACTTG
Ifng	GACAATCAGGCCATCAGCAAC	CGGATGAGCTCATTGAATGCTT
Il4	GGATGTGCCAAACGTCCTC	GAGTCTTCTTCAAGCATGGAG
Il10	CAAGGAGCATTTGAATTCCC	GGCCTTGTAGACACCTTGGTC
Il13	GTCTGGCTCTTGCTTGC	CACTCCATACCATGCTGCC
Mrc1	GGATTGTGGAGCAGATGGAAG	CTTGAATGGAAATGCACAGAC
Nos2	CAGCACAGGAAATGTTTCAGC	TAGCCAGCGTACCGGATGA
Ppard	GAACACACGCTTCCTCCAGC	CACCCGACATTCCATGTTGAG
Retnla	TCGTGGAGAATAAGGTCAAGG	GGAGGCCCATCTGTTTCATAG
Rnsp1	AGGCTCACCAGGAATGTGAC	CTTGGCCATCAATTGTCCT
Rplp0	GGACCCGAGAAGACCTCCTT	GCTGCCGTTGTCAAACACC
Srp14	AGAGCGAGCAGTTCCTGAC	CGGTGCTGATCTTCCTTTTC
Tbp	TCCCCTCTGCACTGAAATC	AGTGCCGCCCAAGTAGCA
Tpt1	GGAGGGCAAGATGGTCAGTAG	CGGTGACTACTGTGCTTTTCG
Tnf	CTTCTGTCTACTGAACTTCGGG	CACTGGTGGTTTGCTACGAC