

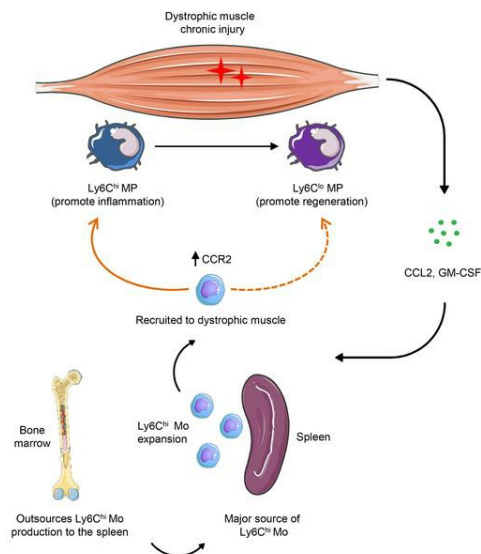
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Splenic Ly6C^{hi} monocytes are critical players in dystrophic muscle injury and repair

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Conflict of interest

The authors declare no conflicts of interest.

Abstract

Dystrophic muscle is characterised by chronic injury, and a steady recruitment of inflammatory Ly6C^{hi} monocytes. Recent studies have identified the spleen as the dominant reservoir of these cells during chronic inflammation. Here we investigated the hitherto unexplored contribution of splenic Ly6C^{hi} monocytes to dystrophic muscle pathology. Using the *mdx* mouse model of muscular dystrophy, we show that Ly6C^{hi} monocytes accumulate in great numbers in the spleen over the course of the disease. The chemokine receptor CCR2 was upregulated on Ly6C^{hi} monocytes in *mdx* spleen before disease onset, thereby enabling their recruitment to dystrophic muscle. Splenectomy performed before disease onset significantly reduced the number of Ly6C^{hi} monocytes infiltrating dystrophic limb muscle. Moreover, in the absence of splenic Ly6C^{hi} monocytes there was a significant reduction in dystrophic muscle inflammation and necrosis, along with improved regeneration during early disease. However, during late disease, lack of splenic Ly6C^{hi} monocytes adversely affected muscle fiber repair, due to a delay in the phenotypic shift of pro-inflammatory F4/80⁺/Ly6C^{hi}/CD206^{lo} to anti-inflammatory F4/80⁺/Ly6C^{lo}/CD206⁺ macrophages. Overall, we show that the spleen is an indispensable source of Ly6C^{hi} monocytes in muscular dystrophy, and that splenic monocytes are critical players in both muscle fiber injury and repair.

Introduction

Duchenne muscular dystrophy (DMD) is a genetic disease in which lack of dystrophin leads to chronic muscle injury and inflammation (1). The immune response plays a critical role in the pathogenesis of DMD, and both the innate and the adaptive immune response contribute to the formation of the dystrophic muscle infiltrate (2–5). Monocytes and macrophages comprise the major part of the inflammatory cell infiltrate in dystrophic muscle (2,6). There are two types of monocytes in circulation, classical Ly6C^{hi} monocytes characterised by the expression of the chemokine receptor CCR2, and Ly6C^{lo} or patrolling monocytes, which lack expression of CCR2 but express the fractalkine receptor CX₃CR1 (7). The classical Ly6C^{hi} monocytes also known as inflammatory monocytes are among the first innate immune cells to be mobilised from the bone marrow into the circulation and recruited to the site of tissue injury (8). Once inside the tissue, and depending on the cytokine milieu, they give rise to both inflammatory and anti-inflammatory macrophages (7,8). On the other hand, Ly6C^{lo} monocytes patrol the vasculature under steady state, and participate in the resolution of inflammation (7–9). We recently described the kinetics of the early inflammatory cell infiltrate in lower limb muscle in the *mdx* mouse model of muscular dystrophy and showed that Ly6C^{hi} monocytes begin to infiltrate muscle early during the pre-necrotic stage of the disease, at 2 weeks of age (6). They reach a peak during the necrotic phase at 4 weeks and gradually decline by 12 weeks of age when anti-inflammatory macrophages begin to predominate. In spite of their important role in dystrophic muscle pathology, little is known about the origin of Ly6C^{hi} monocytes in dystrophic muscle. Although the bone marrow has traditionally been considered as the major source of monocytes, there is now compelling

evidence suggesting that the spleen is also an important reservoir of Ly6C^{hi} monocytes, and that the bone marrow likely outsources their production to the spleen in diverse disease settings (10). Thus, splenic monocytes were found to contribute to myocardial healing following ischemia, the formation of atherosclerotic lesions and to disease progression in ALS (10–12). However, the contribution of splenic monocytes to dystrophic muscle pathology remains to be elucidated.

In this study we used the *mdx* mouse model of muscular dystrophy to investigate whether splenic Ly6C^{hi} monocytes participate in muscle injury, inflammation and regeneration. We show for the first time that Ly6C^{hi} monocytes accumulate in significant numbers in the spleen of *mdx* mice and are recruited to dystrophic muscle where they contribute to both muscle fiber injury and repair.

Results

Ly6C^{hi} monocytes accumulate in the spleen of mdx mice over the course of the disease.

We previously showed that Ly6C^{hi} monocytes begin to infiltrate *mdx* limb muscle before the onset of overt disease at 2 weeks of age, and reach a peak at 4 weeks of age when necrosis becomes evident (6). We therefore hypothesized that if the spleen represents an important source of Ly6C^{hi} monocytes in the context of dystrophic muscle injury, then we should be able to detect changes in their number and or phenotype. To examine this hypothesis we analysed the number of Ly6C^{hi} monocytes in the spleen of *mdx* and WT mice at 1, 2, 4 and 12 weeks of age. Ly6C^{hi} monocytes were identified by flow cytometry as CD11b⁺/F4/80^{lo}/Ly6g^{lo}/CCR2⁺/Ly6C^{hi} cells (Figure 1A). Interestingly, while there were no differences at 1 week of age, as one might expect, we observed a striking increase in the number of Ly6C^{hi} monocytes at 2 weeks of age in *mdx* spleen compared to WT, and this significant increase was sustained at 4 weeks of age, gradually declining by 12 weeks of age when their numbers were comparable to WT (Figure 1B). We also noticed an increase in Ly6g^{hi} neutrophils in the spleen of *mdx* mice, and this will be the subject of future investigations. In support of these data, we found a significant increase in the frequency and number of Ly6C^{hi} monocytes positive for the proliferation marker Ki-67 (Figure 1C-E), and an increase in the expression level of myeloid cell cytokines IL-1 β and GM-CSF mRNA in *mdx* spleen at 2 weeks of age (Figure S1B and C). Of note, we observed increased frequency and number of myeloid progenitor cells in *mdx* spleen (Figure S2A-C), which likely give rise to Ly6C^{hi} monocytes as previously described (12,13). Furthermore, a flow cytometric analysis of splenic Ly6C^{hi} monocytes in *mdx* mice at 2 weeks of age, revealed an upregulated expression of the chemokine

receptor CCR2 (Figure 1F, G), suggesting an activated status, and a preparedness to be recruited and extravasate from circulation into the site of tissue injury.

These results show that there is a marked expansion of the splenic reservoir of Ly6C^{hi} monocytes in *mdx* mice before the onset of disease, suggesting a possible role in dystrophic muscle inflammation.

Splenectomy reduces the number of Ly6C^{hi} monocytes/macrophages infiltrating dystrophic muscle. In order to investigate the contribution of the splenic reservoir of Ly6C^{hi} monocytes to dystrophic muscle inflammation we splenectomised *mdx* mice at 2 weeks of age, which is the time when Ly6C^{hi} monocytes begin to infiltrate muscle (6), and performed analyses 2 weeks later, at 4 weeks of age, when typically *mdx* muscle inflammation and damage peak, 8 weeks of age when inflammation begins to subside, and 12 weeks of age when inflammation is minimal (6) (Figure 2A). Upon recruitment, Ly6C^{hi} monocytes give rise to pro-inflammatory F4/80⁺/CCR2⁺/Ly6C^{hi} macrophages (or recently recruited monocytes/macrophages (Mo/MP)), which over time lose the expression of Ly6C and CCR2 and differentiate into anti-inflammatory or pro-regenerative F4/80⁺/CCR2⁻/Ly6C^{lo} macrophages (Figure 2B) (6,7,14). Splenectomy significantly reduced the number of Ly6C^{hi} Mo/MP in *mdx* limb muscle at 4 (Figure 2C, F) and 8 weeks of age (Figure 2D, F). No significant differences were observed at 12 weeks of age (Figure 2E, F). While there was no difference in the kinetics of Ly6C^{hi} Mo/MP in dystrophic muscle at 4, 8 and 12 weeks, the number of Ly6C^{hi} monocytes was consistently lower in splenectomised *mdx* mice compared to control (Figure 2F). The expression of CCR2 by the recently recruited F4/80⁺Ly6C^{hi} Mo/MP in muscle of *mdx*

control and splenectomised mice at 4 weeks of age was similar (Figure 2G). Concomitant with the reduction in the number of inflammatory Mo/MP, the ligand for the CCR2 chemokine receptor, CCL2, which drives the recruitment of monocytes into the site of tissue injury (8), and the GM-CSF cytokine, which drives the expansion and activation of monocytes (15) were significantly reduced in limb muscle of splenectomised *mdx* mice compared to control (Figure H, I).

Overall, these data suggest that the spleen serves as the dominant reservoir of Ly6C^{hi} monocytes during the acute phase of dystrophic muscle inflammation.

In the absence of the spleen the bone marrow increases the production of Ly6C^{hi} monocytes. Our and previous findings suggest that the bone marrow outsources the supply of inflammatory monocytes to the spleen during chronic inflammation (12,13). Therefore, we reasoned that in the absence of the spleen, the bone marrow (BM) might compensate by increasing the production of Ly6C^{hi} monocytes. Indeed, 2 weeks after splenectomy, at 4 weeks of age, the number of Ly6C^{hi} monocytes in the BM of splenectomised *mdx* mice was significantly increased compared to control (Figure 3A). Intriguingly, the number of Ly6C^{hi} monocytes was also increased in the BM of splenectomised WT mice compared to control, suggesting that the spleen might be a source of monocytes also in the steady state (Figure 3B). The expression of CCR2 on bone marrow Ly6C^{hi} monocytes was slightly increased in both splenectomised *mdx* and WT mice compared to control (Figure 3C). On the other hand there was a modest increase in the number of circulating Ly6C^{hi} monocytes in splenectomised compared to control *mdx* mice (Figure 3D) and a significant increase in splenectomised compared to

control WT mice (Figure 3E). Similar to BM, there was a small increase in the expression of CCR2 by Ly6C^{hi} monocytes in splenectomised compared to control *mdx* mice (Figure 3F). These results show that the bone marrow increases the production of Ly6C^{hi} monocytes following splenectomy, but as shown in Figure 2, fails to fully compensate for the absence of splenic Ly6C^{hi} monocytes.

Lack of splenic Ly6C^{hi} monocytes is associated with a reduction in inflammation and early muscle fiber necrosis. Inflammatory Mo/MP contribute to early inflammation-mediated muscle damage and facilitate the recruitment of other inflammatory cells (2). Thus, removal of the splenic reservoir of Ly6C^{hi} monocytes was associated with a significant reduction of the inflammatory infiltrate in muscle of splenectomised *mdx* mice compared to control (Figure 4A, B). The total number of mononuclear cells isolated from limb muscle was significantly reduced in splenectomised *mdx* mice compared to control at all ages analysed (Figure S3A-C). In addition, the number of CD45⁺ cells was significantly reduced at 4 and 8 weeks of age (Figure 4C, D). There was no significant difference in the number of infiltrating CD45⁺ cells at 12 weeks of age, when muscle inflammation is minimal (Figure 4E). Although the kinetics of CD45⁺ cell infiltration of dystrophic muscle did not change, the number was consistently lower in splenectomised versus control *mdx* mice (Figure 4F). In addition, the number of neutrophils was also significantly reduced at 4 and 8 weeks of age in splenectomised *mdx* muscle (Figure 4G and H), most likely because of the reduced local production of GM-CSF as shown in Figure 2I. There was no significant difference in the number of T cells recruited to dystrophic muscle at 4 and 8 weeks of age in splenectomised compared to control *mdx*

mice (Figure 4I and J), suggesting that in the absence of the spleen T cells are recruited from regional lymph nodes. Finally, lack of Ly6C^{hi} monocytes ameliorated early dystrophic muscle necrosis at 4 weeks (Figure 4K, L).

These findings suggest that splenic Ly6C^{hi} monocytes contribute to dystrophic muscle inflammation and early muscle fiber damage.

Removal of the splenic reservoir of Ly6C^{hi} monocytes improves early regeneration but hinders later stage muscle fiber repair. Monocytes recruited to sites of tissue injury perform diverse effector functions, ranging from pro-inflammatory cytokine production and amplification of the immune response, to clearance of dead cells and stimulating angiogenesis and regeneration (7,14). We therefore wondered whether the absence of splenic Ly6C^{hi} monocytes might affect the ability of dystrophic muscle to regenerate. In the *mdx* mouse model, the acute inflammatory phase in limb muscle begins at 2 weeks of age and is characterised by a peak in inflammation at 4 weeks, after which inflammation subsides and returns to almost basal level by 12 weeks of age (4,6). Throughout this time, regeneration is ongoing and reaches a peak as inflammation begins to wane. We found that the frequency of embryonic myosin heavy chain (eMHC) positive regenerating fibers in limb muscle of splenectomised *mdx* mice was significantly increased compared to control at 4 weeks of age (Figure 5A, B). At this stage, there was no difference in the size (Figure 5C) or mean cross sectional area (CSA) of regenerating fibers (FigureS5E). In support of these data, the cytokine typically associated with regeneration, IL10 was significantly increased in limb muscle from splenectomised *mdx* mice compared to control at 4 weeks of age (Figure S4E). These results suggest that in the absence of

splenic Ly6C^{hi} monocytes regeneration is augmented in dystrophic muscle during the early phase of the pathology, at 4 weeks. This increase in regeneration was not due to increased damage since muscle necrosis and inflammation were reduced in splenectomised *mdx* mice (Figure 4K and L). Another possible reason for the boost in early regeneration following splenectomy could be increased activation of satellite cells. However, we found no significant difference in the level of myogenic markers including Pax7, MyoD and myogenin mRNA in muscle from control and splenectomised *mdx* mice (S6A). While the regenerating area remained slightly increased at 8 weeks compared to control (Figure S5C), the number of small regenerating fibers (CSA < 500 μ m) remained high in limb muscle of splenectomised *mdx* mice, whereas the number of bigger fibers (CSA 1000-2000 μ m) was reduced compared to control (Figure 5D). There was no significant difference in the average CSA of regenerating fibers between splenectomised and control *mdx* mice at 8 weeks (Figure S5F). During the later phase of the disease at 12 weeks of age, regeneration was reduced (Figure S5D), and the size of regenerating fibers remained smaller in muscle of splenectomised *mdx* mice compared to control (Figure 5E). Interestingly, the impaired muscle fiber growth and regeneration at 8 and 12 weeks of age was accompanied by increased necrosis (Figure S5A and B). These findings suggest that splenic Ly6C^{hi} monocytes are necessary for efficient muscle regeneration in muscular dystrophy. Impaired muscle regeneration has been associated with a reduction in angiogenesis and increased fibrosis (16). Indeed, the number of CD31⁺ capillaries per fiber was reduced in the absence of splenic Ly6C^{hi} monocytes (Figure 5F and G). Furthermore, although a histological analysis of collagen deposition in muscle following splenectomy revealed a modest increase in fibrosis in splenectomised *mdx* mice at 12

weeks of age, the expression level of the fibrosis marker Collagen 1A was significantly increased in muscle of splenectomised *mdx* mice compared to control (Figure S7 A-C). Finally, muscle function in splenectomised *mdx* mice was compromised as evidenced by a significant reduction in muscle strength (Figure 5H).

A delay in pro-inflammatory to anti-inflammatory macrophage shift contributes to impaired regeneration following splenectomy. Soon after reaching dystrophic muscle, Ly6C^{hi} monocytes differentiate into F4/80⁺/Ly6C^{hi} pro-inflammatory macrophages. As inflammation progresses, a phenotypic transition begins to take place whereby pro-inflammatory F4/80⁺/Ly6C^{hi} macrophages gradually lose the expression of Ly6C and up-regulate the expression of CD206, eventually giving rise to pro-regenerative F4/80⁺/Ly6C^{lo}/CD206⁺ macrophages (6,7,14). In the absence of splenic Ly6C^{hi} monocytes we observed a delay in this shift in *mdx* muscle at 12 weeks of age (Figure 6A-C). A comparison of the forward (size) and side scatter (granularity) characteristics of F4/80⁺/Ly6C^{hi}/CD206^{lo} and F4/80⁺/Ly6C^{lo}/CD206⁺ macrophages, revealed that they are of a similar size but were more granular in muscle from control compared to splenectomised *mdx* mice, suggesting differences in cytoplasmic complexity (Figure 6D). Finally, as mentioned above, as Ly6C^{hi} macrophages lose the expression of Ly6C, they upregulate the expression of CD206, a marker of anti-inflammatory macrophages. As illustrated in Figure 6E, while Ly6C^{lo} macrophages in control *mdx* muscle gradually increased the expression of CD206 over the course of the disease, this increase was not observed in Ly6C^{lo} macrophages in splenectomised *mdx* muscle (Figure 6E). The expression level of IL-1 β , a pro-inflammatory cytokine produced by activated Mo/MP,

was significantly increased in muscle of splenectomised *mdx* mice compared to control (Figure S4C), reflecting the delay in the shift from pro-inflammatory to anti-inflammatory macrophages. These findings suggest that one of the ways by which lack of splenic Ly6C^{hi} monocytes impedes dystrophic muscle regeneration is by causing a delay in the phenotypic shift from pro-inflammatory to pro-regenerative macrophages, thereby altering the tissue cytokine environment in ways that do not support adequate muscle fiber growth.

Discussion

Inflammatory Ly6C^{hi} monocytes are the principal effector cells in dystrophic muscle pathology (2,6,14). The origin of these cells in dystrophic muscle has not been directly studied before, and the general assumption has been that they originate from the bone marrow. However, following the identification of the spleen as an alternative source of Ly6C^{hi} monocytes by Swirski et al (10), a number of studies have emphasized the critical role of the spleen as an emergency reservoir from which these cells are deployed to sites of inflammation, as shown during ischemic myocardial injury and atherosclerosis (12,13,17). Here we provide further evidence of the spleen as a dominant reservoir of Ly6C^{hi} monocytes in the context of chronic muscle injury in muscular dystrophy. Reminiscent of the kinetics of the inflammatory response in dystrophic muscle (6), Ly6C^{hi} monocytes in the spleen showed the first signs of expansion early, before the onset of overt disease, and remained elevated until the end of the acute inflammatory phase. This early expansion was accompanied by an increase in the expression of CCR2, a chemokine receptor known to be necessary for monocyte egress. Such an early response

in the spleen has also been observed in a mouse model of ALS, where similar to our findings, Ly6C^{hi} monocytes in the spleen upregulated CCR2 expression prior to disease onset (11). Interestingly, these early events in the spleen of *mdx* mice coincided with an increase in limb muscle of CCL2, the ligand for CCR2, which mediates monocyte recruitment, and of GM-CSF, a cytokine, which in Ly6C^{hi} monocytes stimulates chemotaxis, phagocytosis and induction of genes linked to inflammasome function (15). It is possible that GM-CSF produced by activated macrophages and fibroblasts in dystrophic muscle stimulate myelopoiesis in distal sites such as the spleen and the bone marrow, as recently shown in acute myocardial ischemia (18). Nevertheless, unlike in the spleen, we found no significant increase in the number of Ly6C^{hi} monocytes in the bone marrow of *mdx* mice, and no upregulation of CCR2, suggesting that the spleen represents the dominant supplier of these cells during the early stages of dystrophic muscle injury, as previously demonstrated during ischemic myocardial injury and atherosclerosis (12,13,17). Thus, splenectomy performed before disease onset, reduced the number of Ly6C^{hi} monocytes infiltrating dystrophic limb muscle by 70%, and ameliorated inflammation and necrosis during the early stages of the disease. Although the number of Ly6C^{hi} monocytes increased in the bone marrow of splenectomised *mdx* mice compared to control, the bone marrow failed to fully compensate for the absence of the spleen, most probably due to the reduced level of the monocyte mobilising chemokine CCL2 and the GM-CSF cytokine released from dystrophic muscle following splenectomy. The fact that these cytokines were reduced in dystrophic muscle after splenectomy, suggests that splenic Ly6C^{hi} monocytes contribute to the early muscle fiber damage and amplification of the inflammatory response.

Previous studies have shown that targeting Ly6C^{hi} monocytes through CCR2, or treatment with anti-Ly6C mAb reduces dystrophic muscle damage. Mojumdar et al. demonstrated that CCR2 deficiency in *mdx* mice reduces the number of Ly6C^{hi} monocytes recruited to the diaphragm, and ameliorates diaphragm muscle degeneration during the early stages of the disease (19). Interestingly, the authors found that splenectomy had no effect on the number of Ly6C^{hi} monocytes recruited to the diaphragm. The reason for the different outcomes between this and our study could be that the focus was on different tissues (diaphragm versus limb muscle) with different inflammation kinetics. For example, in our study we performed splenectomy at 2 weeks of age because our kinetics data showed that to be the earliest time when monocytes begin to infiltrate limb muscle. We sacrificed the animals 2 weeks later in order to examine the effect on the peak of inflammation, and 6 and 10 weeks later in order to study regeneration. Although much less striking, the difference in the number of Ly6C^{hi} Mo/MP observed at 4 weeks, was still present at 8 weeks, but absent at 12 weeks of age. Mojumdar et al. splenectomised the animals at 3 weeks of age and examined the diaphragm 6 weeks later when inflammation likely begins to subside and any differences in numbers that might have existed become less obvious (19).

A more recent study, using *mdx*^{5cv}-CCR2^{-/-} mice also showed a beneficial effect on muscle pathology by reducing the number of infiltrating Ly6C^{hi} monocytes. However, the improvement in diaphragm and limb muscle damage and fibrosis observed at 14 weeks was not sustained at 6 months (20). Similar to this study, we found that the beneficial effect of splenectomy observed during the early phase was not sustained during the later stages of the disease. While lack of splenic monocytes reduced inflammation and

necrosis and ameliorated regeneration early in the acute disease phase, a defect in muscle fiber repair became evident by 8 weeks and was still apparent at 12 weeks. Smaller regenerating fibers predominated in splenectomised *mdx* mice, and there was a reduced appearance of bigger fibers, suggesting impaired muscle fiber growth or maturation. This compromised repair of muscle fibers, together with the reduced number of CD31⁺ capillaries per fiber, led to a reduction in muscle strength, suggesting that in the absence of splenic Ly6C^{hi} monocytes the dystrophic phenotype worsens. Removal of the splenic reservoir of Ly6C^{hi} monocytes did not affect the ability of satellite cells to undergo activation, and the impairment in muscle fiber growth is likely due to the delay in the appearance of the pro-regenerative F4/80⁺/Ly6C^{lo}/CD206⁺ macrophage subset and the consequent changes in the tissue cytokine environment that do not adequately support muscle fiber growth, as previously proposed in various contexts of muscle injury (14,21,22). On the other hand, the balance of pro-inflammatory Ly6C^{hi} to pro-regenerative Ly6C^{lo} macrophages in splenectomised *mdx* muscle was not affected during the early phase of the disease, which, together with the higher level of pro-regenerative cytokine IL10 and the decrease in inflammation might have contributed towards the increase in early regeneration we observed at 4 weeks of age.

Given that Mo/MP perform diverse effector functions both in the context of tissue injury and regeneration, these findings were not surprising. The optimal response to tissue injury depends on a fine balance between inflammatory monocyte numbers and their phenotype. Recruitment of too few or too many is likely to create an unfavourable environment for the resolution of tissue inflammation, and compromise the re-establishment of tissue homeostasis. Thus, it seems probable that the 70% reduction in

the number of Ly6C^{hi} monocytes in dystrophic muscle following splenectomy, precluded the effective clearance of necrotic cells and debris, eventually hindering repair and worsening fibrosis. It is also possible, that the spleen provides a distinct monocyte subtype or subtypes that differentially participate in the inflammatory and regenerative phase of dystrophic muscle injury and are recruited as part of separate waves. Future studies are needed to test these possibilities and broaden our understanding of the origin and function of monocytes and macrophages in muscular dystrophy.

Finally, we cannot exclude the possibility that absence of cells other than monocytes, recruited from the spleen might have contributed to these outcomes. Nevertheless, we found no difference in the number of CD3⁺ T cells recruited to muscle of splenectomised animals, and although we did not examine regulatory T cells, we found similar levels of the immunoregulatory cytokines IL-10 and TGF- β in muscle of splenectomised versus control *mdx* mice.

In conclusion, we show that the spleen is an indispensable source of Ly6C^{hi} monocytes in muscular dystrophy, and that splenic monocytes contribute to dystrophic muscle injury early in the course of the disease, but are also necessary for effective regeneration during the later stages of the pathology.

An important question to be addressed in the future is whether the spleen is just an extramedullary site to which the bone marrow outsources the production of Ly6C^{hi} monocytes during chronic inflammation, or whether it also represents an important monocyte training ground.

The present study underscores the complexity of monocyte responses in tissue injury and highlights the need for a greater understanding of the origin, time and context dependent

effector functions of these cells. Only then, will we be able to design novel pharmaceuticals to target monocytes in muscular dystrophy in a way that will reduce damage without compromising regeneration.

Methods

Mice. C57BL/10ScSn-Dmd^{mdx} and C57BL/10ScSn mice were purchased from The Jackson laboratory. Only males were used.

Cell isolation and processing. Muscle single cell suspensions were obtained as previously described (6). Briefly, mice were perfused with PBS, and hind limb musculature (tibialis anterior, gastrocnemius and quadriceps) was excised, weighed, cut into 1-mm pieces and digested with 1mg/ml Collagenase type 4 (Worthington) for 1.5h in a pre-warmed (37C) water bath with agitation. Hind limb muscle from each mouse was processed individually. The digested muscle preparation was first passed through a 70µm cell strainer (BD Falcon), and then through a 40µm cell strainer, washed in DMEM/10% FBS and resuspended in FACS buffer (PBS 1%FBS). Single cells were counted with a haemocytometer and analysed by flow cytometry.

Splenocytes were obtained by grinding the spleen between the rough edges of two microscopy slides. Blood was collected by cardiac puncture. Following the lysis of RBCs, cells were counted, stained with the relevant antibodies and analysed by FACS.

Splenectomy. Splenectomy was performed on zoletil (50/50 mg/ml) and rompun (20 mg/ml) anesthetized 2-week-old *mdx* or WT mice. After opening of the abdominal cavity, the spleen vessels were cauterized and the spleen was carefully excised. Control animals

underwent anesthesia and opening of the abdominal cavity but without removal of the spleen. Animals were sacrificed 2, 6 or 10 weeks after splenectomy for analysis.

Flow Cytometry. Prior to staining, single cell suspensions were incubated with anti-CD16/32 (Biolegend, clone 93, 1 μ l/test) to block nonspecific binding to Fc receptors. Samples were stained with the relevant antibodies listed in Supplementary Table 1. Cell viability was assessed with 4',6-diamidino-2-phenylindole, dilactate (DAPI) (Biolegend). Compensation was set using the ABC compensation beads kit (Thermo Fisher). Samples were acquired with a CyAn ADP (DAKO) and acquired data were analysed using FlowJo software version 10.

Histology. Tibialis muscles were embedded in Tissue freezing medium (Leica) and snap-frozen in liquid nitrogen-cooled isopentane. Frozen muscles were cut into 8 μ m sections and stored at -20 C until use. For morphological assessment, frozen sections were stained with H&E. The extent of necrosis was determined by staining with fluorescently labeled IgG (Sigma, 1:100). Regeneration was assessed by staining for embryonic myosin heavy chain (eMHC) (Hybridoma Bank, clone F1.652, 1:10). The cross sectional area (CSA) of regenerating fibers was examined using anti-laminin antibody (Sigma, 1:200) to visualise the fibers and Image J software for calculating the area of regeneration (23). Angiogenesis was examined by staining of muscle sections with CD31 antibody (Biolegend, 1:100) to visualise the capillaries as previously described (23). All analyses were done in blinded fashion and on 3 muscle sections per mouse. Fibrosis was assessed by staining of muscle sections with sirius red to visualise areas of collagen deposition and quantified using Image J software.

Hanging-wire test. Each mouse was placed on top of a wire lid from a typical housing cage. The lid was then turned upside down with the mouse hanging by its fore limbs and the time until fall was recorded. Each mouse was given three attempts to reach the 180 s limit before releasing its grip and falling, with 30 s recovery period prior to the next attempt. The longest suspension time was recorded.

Quantitative real-time PCR. RNA was isolated from muscle single cells using Trizol (Sigma) according to the manufacturer's instructions. cDNA was synthesised using the High-Capacity cDNA Reverse Transcription kit (Applied Biosystems). PCR amplification was performed using the SensiMix™ SYBR® Low-ROX Kit, following the manufacturer's protocol. Data analysis was performed 7500 Software v2.0.6 provided by Applied Biosystems. Data are expressed as fold-change in expression levels. Relative expression levels were normalised to *GAPDH* mRNA. The primer pairs used for amplification are listed in Supplementary materials.

Statistical analysis. All statistical analyses were performed using GraphPad Prism software version 6. Data are presented as mean \pm SEM. Statistical significance was determined using unpaired 2-tailed Student's *t* test with Welch's correction for unequal variances, or 1-way ANOVA followed by Sidak's correction for multiple comparisons as appropriate. A *P* value of ≤ 0.05 was considered statistically significant.

Study approval. All mice were bred and housed in our pathogen-free Histology Department-accredited animal facility. All procedures involving mice were approved by the Italian Ministry for Health and were conducted according to the U.S. National Institutes of Health (NIH) guidelines and Directive 2010/63/EU.

Author contributions

B.L.O conceived the study, designed the experiments, analysed and interpreted data. G.R and R.D.M performed experiments and analysed data. A.B and J.M helped with sample collection and histological analysis. MB contributed to experimental design and data interpretation, and provided resources and financial support. BLO sourced funding for the study, and wrote the manuscript. All authors critically read, edited and approved the final manuscript.

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References

1. Emery AE. The muscular dystrophies. *Lancet*. 2002;359(9307):687–95.
2. Rosenberg AS, Puig M, Nagaraju K, Hoffman EP, Villalta SA, Rao VA, et al. Immune-mediated pathology in Duchenne muscular dystrophy. *Sci Transl Med*. 2015;7(299):299rv4.
3. Evans NP, Misyak SA, Robertson JL, Bassaganya-Riera J, Grange RW. Immune-mediated mechanisms potentially regulate the disease time-course of duchenne muscular dystrophy and provide targets for therapeutic intervention. *PM R*. 2009;1(8):755–68.

4. Spencer MJ, Tidball JG. Do immune cells promote the pathology of dystrophin-deficient myopathies? *Neuromuscul Disord*. 2001;11(6-7):556-64.
5. Villalta SA, Rosenberg AS, Bluestone JA. The immune system in Duchenne muscular dystrophy: Friend or foe. *Rare Dis*. 2015;3(1):e1010966.
6. Lozanoska-Ochser B, Benedetti A, Rizzo G, Marrocco V, Di Maggio R, Fiore P, et al. Targeting early PKC θ -dependent T-cell infiltration of dystrophic muscle reduces disease severity in a mouse model of muscular dystrophy. *J Pathol*. 2017;
7. Guilliams M, Mildner A, Yona S. Developmental and Functional Heterogeneity of Monocytes. *Immunity*. 2018;49(4):595-613.
8. Ingersoll MA, Platt AM, Potteaux S, Randolph GJ. Monocyte trafficking in acute and chronic inflammation. *Trends Immunol*. 2011;32(10):470-7.
9. Guilliams M, Scott CL. Does niche competition determine the origin of tissue-resident macrophages? *Nat Rev Immunol*. 2017;17(7):451-60.
10. Swirski FK, Nahrendorf M, Etzrodt M, Wildgruber M, Cortez-Retamozo V, Panizzi P, et al. Identification of splenic reservoir monocytes and their deployment to inflammatory sites. *Science* (80-). 2009;325(5940):612-6.
11. Butovsky O, Siddiqui S, Gabriely G, Lanser AJ, Dake B, Murugaiyan G, et al. Modulating inflammatory monocytes with a unique microRNA gene signature ameliorates murine ALS. *J Clin Invest*. 2012;122(9):3063-87.
12. Leuschner F, Rauch PJ, Ueno T, Gorbатов R, Marinelli B, Lee WW, et al. Rapid monocyte kinetics in acute myocardial infarction are sustained by extramedullary monocytopoiesis. *J Exp Med*. 2012;209(1):123-37.
13. Robbins CS, Chudnovskiy A, Rauch PJ, Figueiredo JL, Iwamoto Y, Gorbатов R,

- et al. Extramedullary hematopoiesis generates Ly-6C(high) monocytes that infiltrate atherosclerotic lesions. *Circulation*. 2012;125(2):364–74.
14. Tidball JG. Regulation of muscle growth and regeneration by the immune system. *Nat Rev Immunol*. 2017;17(3):165–78.
 15. Becher B, Tugues S, Greter M. GM-CSF: From Growth Factor to Central Mediator of Tissue Inflammation. *Immunity*. 2016;45(5):963–73.
 16. Buckley AF, Bossen EH. Skeletal muscle microvasculature in the diagnosis of neuromuscular disease. *J Neuropathol Exp Neurol*. 2013;72(10):906–18.
 17. Hulsmans M, Sager HB, Roh JD, Valero-Muñoz M, Houstis NE, Iwamoto Y, et al. Cardiac macrophages promote diastolic dysfunction. *J Exp Med*. 2018;215(2):423–40.
 18. Anzai A, Choi JL, He S, Fenn AM, Nairz M, Rattik S, et al. The infarcted myocardium solicits GM-CSF for the detrimental oversupply of inflammatory leukocytes. *J Exp Med*. 2017;214(11):3293–310.
 19. Mojumdar K, Liang F, Giordano C, Lemaire C, Danialou G, Okazaki T, et al. Inflammatory monocytes promote progression of Duchenne muscular dystrophy and can be therapeutically targeted via CCR2. *EMBO Mol Med*. 2014;6(11):1476–92.
 20. Zhao W, Wang X, Ransohoff RM, Zhou L. CCR2 deficiency does not provide sustained improvement of muscular dystrophy in mdx5cv mice. *FASEB J*. 2017;31(1):35–46.
 21. Wang H, Melton DW, Porter L, Sarwar ZU, McManus LM, Shireman PK. Altered macrophage phenotype transition impairs skeletal muscle regeneration. *Am J*

Pathol. 2014;184(4):1167–84.

22. Jin RM, Warunek J, Wohlfert EA. Chronic infection stunts macrophage heterogeneity and disrupts immune-mediated myogenesis. *JCI Insight*. 2018;3(18).
23. Marrocco V, Fiore P, Benedetti A, Pisu S, Rizzuto E, Musarò A, et al. Pharmacological Inhibition of PKC θ Counteracts Muscle Disease in a Mouse Model of Duchenne Muscular Dystrophy. *EBioMedicine*. 2017;

Figure legends

Figure 1

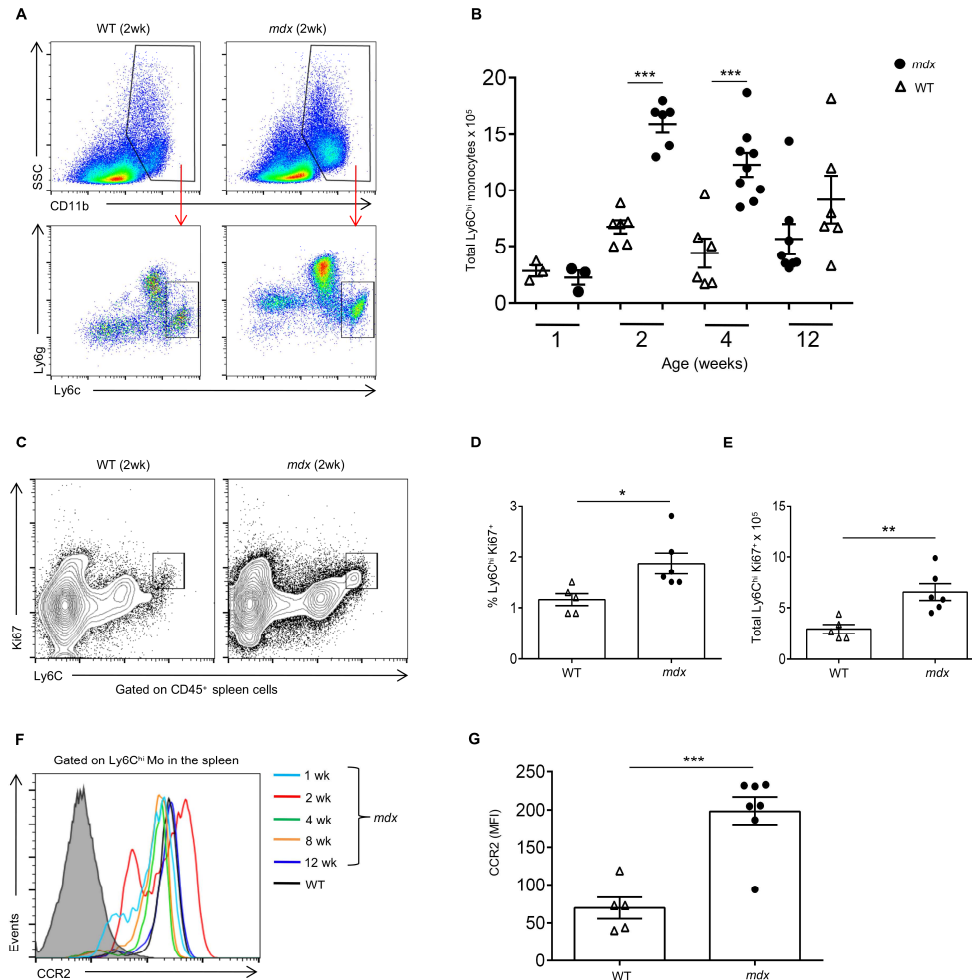


Figure 1. Expansion of the Ly6C^{hi} monocyte reservoir in *mdx* spleen. The number and kinetics of Ly6C^{hi} in *mdx* and WT spleen were analysed by flow cytometry at 1, 2, 4 and 12 weeks of age. (A) Representative flow cytometry plots showing the gating strategy for the identification of Ly6C^{hi} monocytes. Ly6C^{hi} monocytes in the spleen were identified as CD45⁺/CD11b⁺/Ly6g^{lo}/CCR2⁺/Ly6C^{hi} cells. (B) Quantification and kinetics of Ly6C^{hi} monocytes in *mdx* and WT spleen at 1 (n=3, *mdx* and n=3, WT), 2 (n=6, *mdx* and n=6, WT), 4 (n=9, *mdx* and n=6, WT) and 12 (n=8, *mdx* and n=6, WT) weeks of age. (C) Representative flow cytometry plots of proliferating (Ki67⁺) Ly6C^{hi} monocytes in *mdx* versus WT spleen at 2 weeks of age. (D) Frequency and (E) total number of proliferating Ki67⁺Ly6C^{hi} monocytes in *mdx* compared to WT spleen (n=6, *mdx* and n=5, WT) at 2 weeks of age. (F) Representative histogram overlays of flow cytometric analysis of CCR2 receptor expression by Ly6C^{hi} monocytes in *mdx* spleen at 1, 2, 4, 8 and 12 weeks and WT spleen at 2 weeks of age. (G) Mean fluorescence intensity (MFI) of CCR2 expression by Ly6C^{hi} monocytes in *mdx* (n=7) and WT (n=5) spleen as determined by flow cytometric analysis at 2 weeks of age. Data are presented as mean ± s.e.m. *P<0.05, **P<0.001, ***P<0.001 by Student's *t* test.

Figure 2

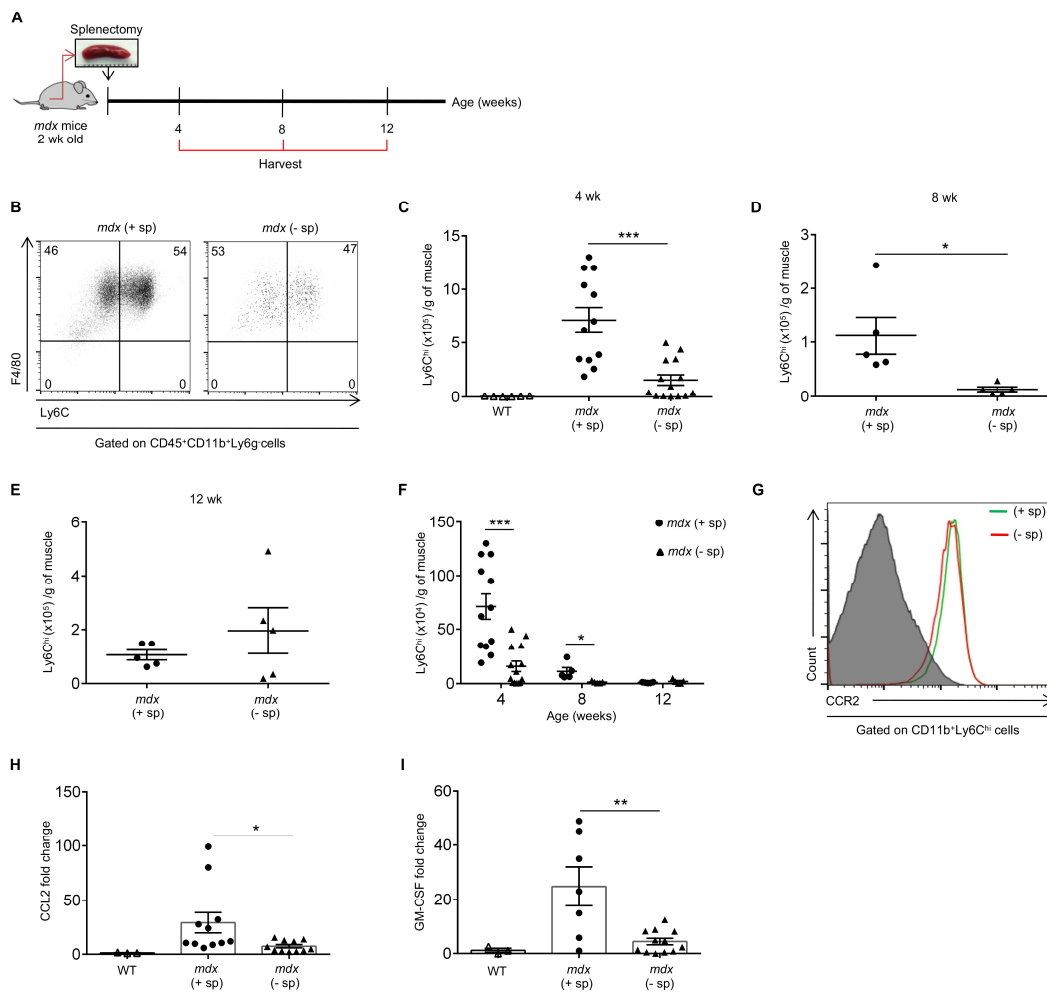


Figure 2. Splenectomy reduces the number of Ly6C^{hi} monocytes recruited to dystrophic muscle. (A) Schematic presentation of the experimental protocol. Mice were splenectomised at 2 weeks of age, and allowed to recover for at least 2 weeks before analysis. Control mice were treated the same but without removal of the spleen. Following splenectomy the Ly6C^{hi} monocyte infiltrate in hind limb muscle was analysed by flow cytometry at 4, 8 and 12 weeks of age. (B) Representative flow cytometry plots of Ly6C^{hi} monocytes in control (+ sp) and splenectomised (- sp) *mdx* muscle. Ly6C^{hi} monocytes in muscle were identified as CD45⁺/CD11b⁺/Ly6g⁻/F4/80⁺/CCR2⁺/Ly6C^{hi} cells. (C) Quantification of Ly6C^{hi} monocytes in dystrophic muscle of control and splenectomised *mdx* mice at 4 weeks of age normalised to muscle mass (n=12, control and n=14, splenectomised), (D) 8 weeks of age (n=5 for both groups), and (E) 12 weeks of age (n=5 for both groups). (F) Summary graph showing Ly6C^{hi} monocyte kinetics in muscle of control and splenectomised *mdx* mice. (G) Expression of the CCR2 chemokine receptor in muscle of control and splenectomised *mdx* mice at 4 weeks of age, analysed by flow cytometry and presented as histogram overlays. (H) Expression levels of CCL2 and (I) GM-CSF mRNA in muscle of control and splenectomised *mdx* mice at 4 weeks of age, assessed by quantitative real-time PCR (n=3, WT; n=11, *mdx* control; n=11, *mdx* splenectomised). Data are presented as mean

\pm s.e.m. * $P < 0.05$, ** $P < 0.01$, *** $P < 0.001$ by 1-way ANOVA (C, F, H and I) and Student's t test (D).

Figure 3

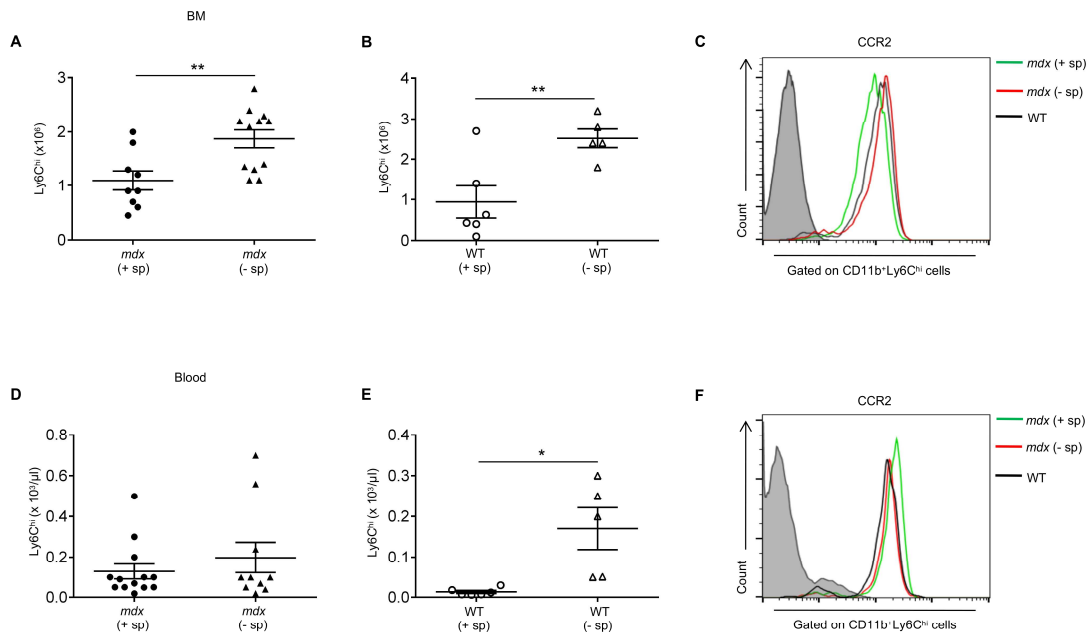


Figure 3. Effect of splenectomy on Ly6C^{hi} monocyte numbers in the bone marrow (BM) and blood. Following splenectomy at 2 weeks of age Ly6C^{hi} monocytes in the BM were analysed by flow cytometry at 4 weeks of age. (A) Quantification of Ly6C^{hi} monocytes in the BM (per leg) of control (+ sp) and splenectomised (- sp) *mdx* (n=9 *mdx* control; n=12, *mdx* splenectomised) and (B) WT mice (n=6, WT control; n=5 WT splenectomised) by flow cytometry. (C) Histogram overlay of CCR2 expression on Ly6C^{hi} monocytes in the BM of control and splenectomised *mdx* mice analysed by flow cytometry. (D) Quantification of Ly6C^{hi} monocytes in the blood of control and splenectomised *mdx* (n=13 *mdx* control; n=10, *mdx* splenectomised) and (E) WT mice (n=6, WT control; n=5 WT splenectomised) by flow cytometry. (F) Histogram overlay of CCR2 expression by circulating Ly6C^{hi} monocytes analysed by flow cytometry. Data are presented as mean \pm s.e.m. * $P < 0.05$, ** $P < 0.01$ by Student's t test.

Figure 4

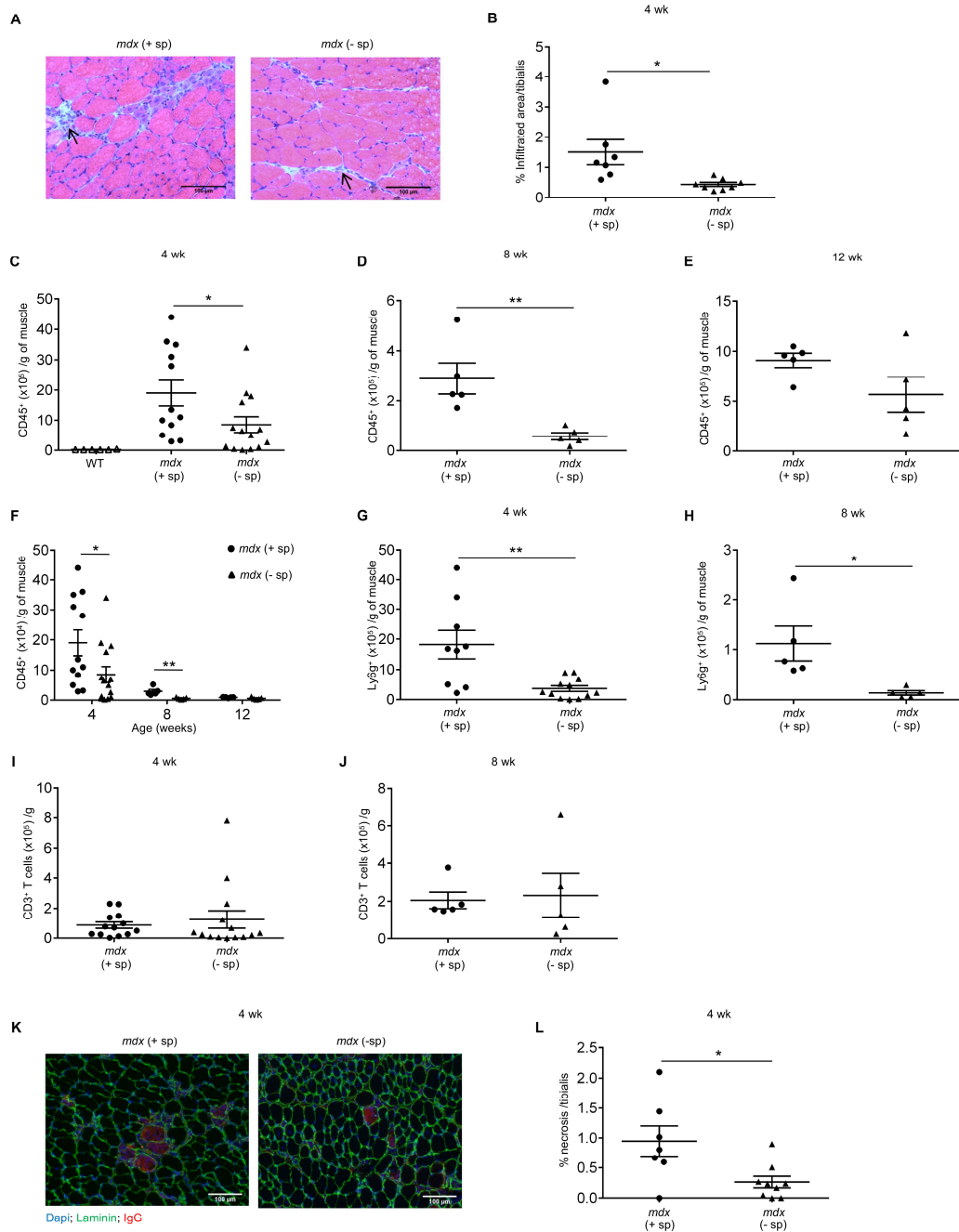


Figure 4. Splenectomy reduces dystrophic muscle inflammation and early necrosis. (A) Representative Haematoxylin and Eosin staining of tibialis anterior (TA) muscle sections showing reduced areas of infiltration (arrow) in splenectomised (- sp) *mdx* mice compared to control (+ sp) at 4 weeks of age. Scale bars: 100 μ m. (B) A summary graph of histological analysis showing percent infiltration area (n=7, *mdx* control; n=8, *mdx* splenectomised).

Following splenectomy at 2 weeks of age the inflammatory cell infiltrate in hind limb muscle was analysed by flow cytometry at 4, 8 and 12 weeks of age (C-J). (C) Total number of CD45⁺ cells normalised to muscle mass in control and splenectomised *mdx* mice at 4 weeks (n=6, WT; n=12, *mdx* control; n=14, *mdx* splenectomised), (D) 8 weeks (n=5 for all groups) and (E) 12 weeks of age (n=5 for all groups). (F) Summary graph showing the kinetics of CD45⁺ cells in muscle of control and splenectomised *mdx* mice at 4, 8 and 12 weeks of age. (G) Total number of Ly6g⁺ neutrophils in muscle of control and splenectomised *mdx* mice at 4 and (H) 8 weeks of age. (I) Total number of CD3⁺ T cells in muscle of control and splenectomised *mdx* mice at 4 and (J) 8 weeks of age. (K) Representative TA muscle sections stained with IgG (red) to reveal necrotic muscle fibers showing reduced necrosis in muscle of splenectomised *mdx* mice compared to control. Scale bars: 100 μ m. (L) Quantification of necrosis (n=7, control, n=9, splenectomised *mdx* mice) at 4 weeks. Data are presented as mean \pm s.e.m. *P<0.05, **P<0.01 by Student's *t* test (B, D, G, H and L) and 1-way ANOVA (C, F).

Figure 5

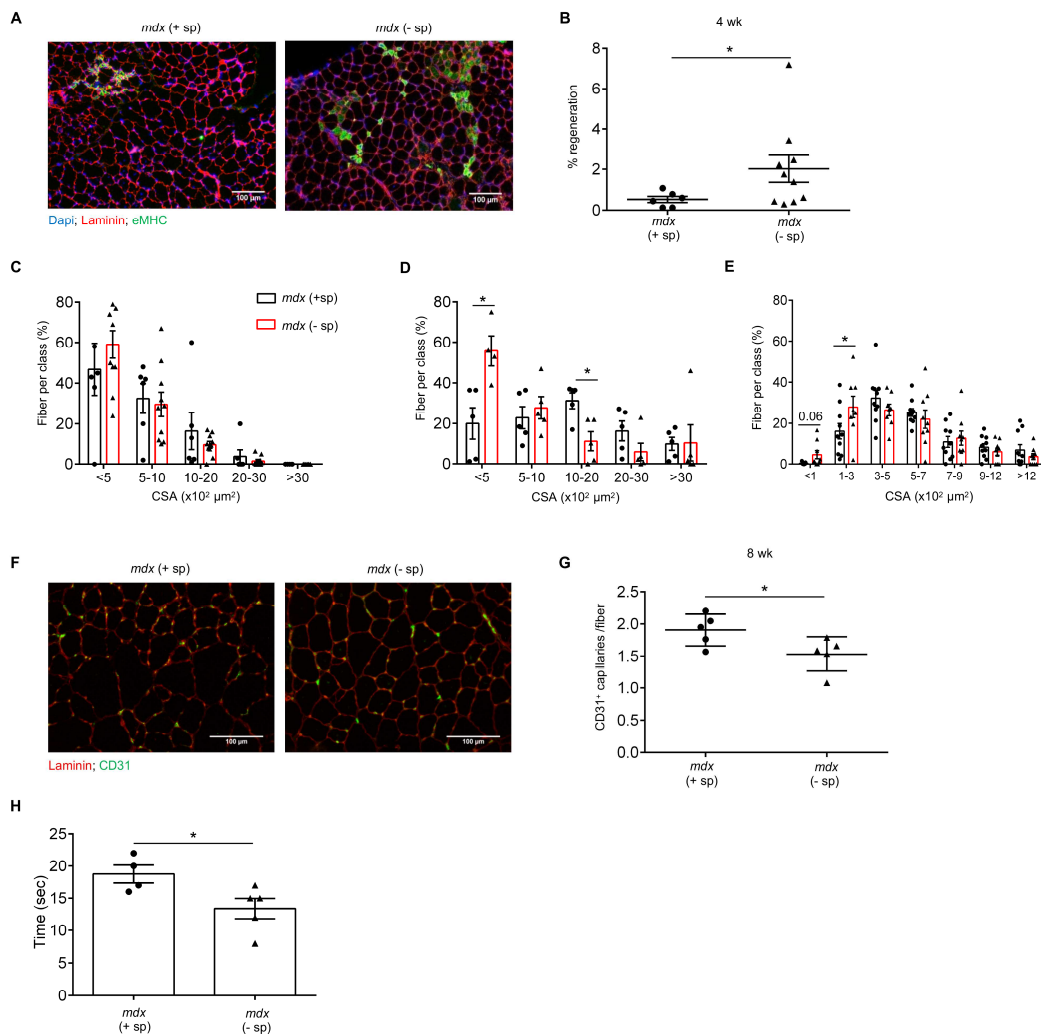


Figure 5. Splenic Ly6C^{hi} monocytes are necessary for efficient repair of dystrophic muscle fibers. (A) Representative TA muscle sections stained for embryonic myosin heavy chain (eMHC) to visualise regenerating muscle fibers (green) showing an increase in eMHC positive fibers in splenectomised (- sp) compared to control (+ sp) *mdx* mice at 4 weeks of age during the early phase of the pathology. Scale bars: 100 μ m. (B) Summary graph showing percent regenerating area (n=6, *mdx* control; n=10 *mdx* splenectomised). (C) Regenerating fiber size frequency distribution determined by measuring the cross sectional area (CSA) of regenerating fibers in TA muscle from control and splenectomised mice at 4 weeks of age (n=6, *mdx* control; n=10 *mdx* splenectomised), (D) 8 weeks of age (n=5 for both groups) and (E) 12 weeks of age (n=10, *mdx* control; n=9 *mdx* splenectomised). (F) Representative TA muscle sections stained for CD31 to reveal capillaries (green). Scale bars: 100 μ m. (G) Number of CD31+ capillaries per muscle fiber at 8 weeks of age in muscle of control and splenectomised *mdx* mice. (H) A graph showing a reduction in muscle strength in splenectomised compared to control *mdx* mice at 12 weeks of age, measured by a hanging wire test, which determines the time it takes for a mouse to fall off a metal wire following exhaustion. Data are presented as mean \pm s.e.m. *P<0.05 by Student's *t* test.

Figure 6

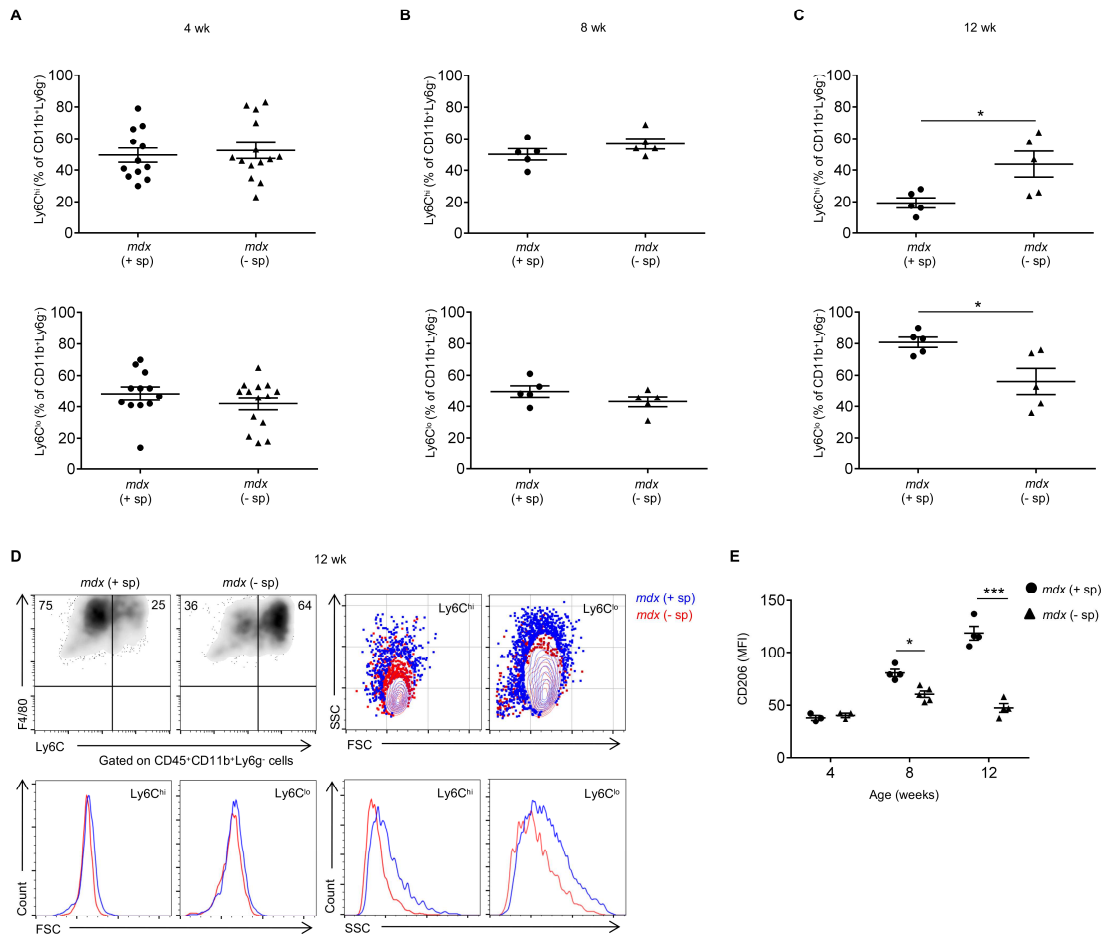


Figure 6. Splenectomy delays the phenotypic switch from pro-inflammatory Ly6C^{hi} to pro-regenerative Ly6C^{lo} macrophages in dystrophic limb muscle. (A) Frequency of Ly6C^{hi} and Ly6C^{lo} macrophages in *mdx* control and splenectomised muscle at 4, (n=12, control and n=14, splenectomised), (B) 8 (n=5 for both groups) and (C) 12 (n=5 for both groups) weeks of age analysed by flow cytometry. (D) Representative density and contour flow cytometry plots and histogram overlays of forward and side scatter parameters of Ly6C^{hi} and Ly6C^{lo} macrophages in control and splenectomised *mdx* muscle. (E) Kinetics of CD206 expression on Ly6C^{lo} macrophages assessed by flow cytometry. Shown is the mean fluorescence intensity (MFI) of CD206 expression, by Ly6C^{lo} macrophages in muscle from control and splenectomised *mdx* mice at 4 (n=3 for both groups), 8 (n=4, *mdx* control; n=5, *mdx* splenectomised) and 12 weeks of age (n=4 for both groups). Data are presented as mean \pm s.e.m. * $P < 0.05$, *** $P < 0.001$ by Student's *t* test.

