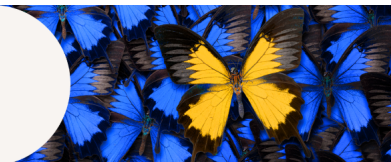




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On the cover: Mutation of IFN- γ in the *mdx* mouse model of Duchenne muscular dystrophy shifts macrophages toward an M2 phenotype and reduces damage, increases regeneration, and improves function of muscle. The image shows a section of IFN- γ null *mdx* muscle where M2 macrophages (red) are enriched in endomysium between regenerative muscle fibers. Villalta, S. A., B. Deng, C. Rinaldi, M. Wehling-Henricks, and J. G. Tidball. 2011. IFN- γ promotes muscle damage in the *mdx* mouse model of Duchenne muscular dystrophy by suppressing M2 macrophage activation and inhibiting muscle cell proliferation. *J. Immunol.* 187: 5419–5428.

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