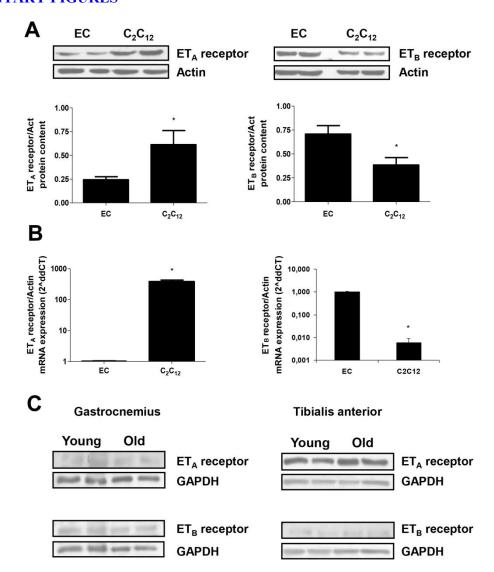
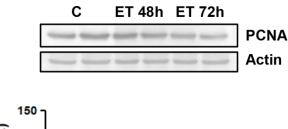
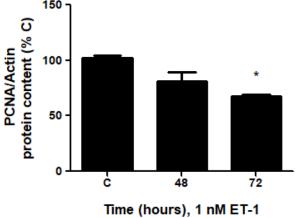
SUPPLEMENTARY FIGURES

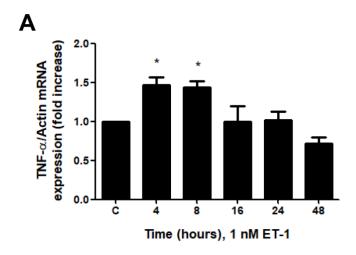


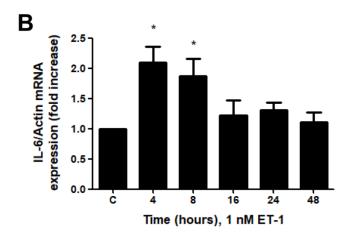
Supplementary Figure 1. Expression of ET receptors in mouse myoblasts (C_2C_{12}) and gastrocnemius and tibialis anterior muscles from mice. Expressions of ET_A and ET_B receptors were assessed in cultured cells, endothelial cells (EC) and mouse myoblasts (C_2C_{12}) (A, B), and mice muscles (C), by Western blot (A, C) and by RT-qPCR in cells (B). Representative Western blots are shown at the top of panel A with the densitometric analyses below. Values are the mean \pm SEM of 3 independent experiments, *p<0.05 vs. endothelial cells (EC).

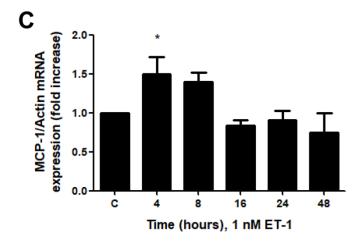




Supplementary Figure 2. Endothelin-1 reduces PCNA expression in mouse myoblasts (C_2C_{12}). Cells were incubated with 1 nM ET-1 at different times. Then, PCNA protein content was assessed by Western blot. A representative Western blot is shown at the top with the densitometric analysis below. Values are the mean \pm SEM of 3 independent experiments, *p<0.05 vs. control cells (C).







Supplementary Figure 3. Endothelin-1 increases pro-inflammatory cytokines expression in mouse myoblasts (C_2C_{12}). Cells were incubated with 1 nM ET-1 at different times. Then, TNF- α (A), IL-6 (B) and MCP-1 (C) mRNA expressions were assessed by real-time PCR. Values are the mean \pm SEM of 3 independent experiments, *p<0.05 vs. control cells (C).