Electrical stimulation induces IL-6 in skeletal muscle through extracellular ATP by activating Ca\(^{2+}\) signals and an IL-6 autocrine loop

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Electrical stimulation induces IL-6 in skeletal muscle through extracellular ATP by activating Ca\(^{2+}\) signals and an IL-6 autocrine loop. Am J Physiol Endocrinol Metab 306: E869–E882, 2014. First published February 11, 2014; doi:10.1152/ajpendo.00450.2013.—Interleukin-6 (IL-6) is an important myokine that is highly expressed in skeletal muscle cells upon exercise. We assessed IL-6 expression in response to electrical stimulation (ES) or extracellular ATP as a known mediator of the excitation-transcription mechanism in skeletal muscle. We examined whether the canonical signaling cascade downstream of IL-6 (IL-6/JAK2/STAT3) also responds to muscle cell excitation, concluding that IL-6 influences its own expression through a positive loop. Either ES or exogenous ATP (100 \(\mu\)M) increased both IL-6 expression and p-STAT3 levels in rat myotubes, a process inhibited by 100 \(\mu\)M suramin and 2 U/ml apyrase. ATP also evoked IL-6 expression in both isolated skeletal fibers and extracts derived from whole FDB muscles. ATP increased IL-6 release up to 10-fold. STAT3 activation evoked by ATP was abolished by the JAK2 inhibitor HBC. Blockade of secreted IL-6 with a neutralizing antibody or preincubation with the STAT3 inhibitor VIII reduced STAT3 activation evoked by extracellular ATP by 70%. Inhibitor VIII also reduced by 70% IL-6 expression evoked by ATP, suggesting a positive IL-6 loop. In addition, ATP increased up to 60% the protein levels of SOCS3, a negative regulator of the IL-6 signaling pathway.

On the other hand, intracellular calcium chelation or blockade of IP\(_3\)-dependent calcium signals abolished STAT3 phosphorylation evoked by either extracellular ATP or ES. These results suggest that expression of IL-6 in stimulated skeletal muscle cells is mediated by extracellular ATP and nucleotide receptors, involving IP\(_3\)-dependent calcium signals as an early step that triggers a positive IL-6 autocrine loop.

myokines; muscle plasticity; exercise; signal transducer and activator of transcription 3; purinergic signaling

Interleukin-6 (IL-6) is a proinflammatory cytokine that has been related to several processes in skeletal muscle cells, including in vitro proliferation and differentiation, regeneration of damaged adult fibers, atrophy, and hypertrophy, among others (7, 35, 71, 86). Variations of IL-6 expression levels have been related to metabolic changes in skeletal muscle undergoing different exercise protocols (17, 66, 77). It is known that IL-6 plasma concentration in volunteers performing exercise can reach increases of up to 100-fold, depending on the exercise characteristics (endurance/strength) and duration (reviewed in Ref. 61). The main source of plasma IL-6 during exercise is the skeletal muscle fibers themselves, although other cell types such as cells from the immune system and subcutaneous adipose tissue cannot be ruled out as IL-6 producers (2, 8, 41, 58).

Several intracellular signaling pathways have been related to IL-6 expression in skeletal muscle. It has been suggested that intracellular free Ca\(^{2+}\), acting as a second messenger, regulates the expression of IL-6 in skeletal muscle, possibly through the Ca\(^{2+}\)-dependent phosphatase calcineurin (5, 8, 87). Also, another Ca\(^{2+}\)-dependent process, possibly through the action of p38 MAPK and Ca\(^{2+}/\)calmodulin-dependent kinase, has been reported to play a role on IL-6 expression related to intracellular glycogen content (17, 87). The role for NF-\(\kappa\)B, a classic activator of proinflammatory interleukin expression, in IL-6 expression mediated by muscle activity is less clear, as we and others have obtained disparate results (4, 45, 89). Whereas we reported a NF-\(\kappa\)B activation in C2C12 cells after a 45-Hz electrical stimulation (ES) (4, 45), Whitham et al. (89) did not find any differences in IL-6 expression between control and cells pharmacologically treated for IKK inhibition after 1-Hz ES. This discrepancy might represent a fine-tuned regulation of IL-6 expression that is strongly dependent on a stimulation pattern that might have a role in the muscle plasticity process. On the other hand, AP-1 appears to have a major role in IL-6 expression. Recently, it has been shown that the JNK/AP-1 pathway plays an important role in IL-6 expression in muscle cells (89). Whitham et al. (89) demonstrated that contraction of skeletal muscle cells induced by electric pulse stimulation increases the JNK phosphorylation, the activity of an AP-1 luciferase reporter, and the expression of IL-6. All of these effects were abolished upon pharmacological inhibition of JNK. The same results were obtained when they analyzed p-JNK levels and IL-6 expression in control and knockout mice for JNK1 at rest and after 30 min of exercise (89). In agreement with the role of AP-1 on IL-6 expression, we have demonstrated previously that depolarization of primary skeletal muscle cells from rats, as well as C2C12 cells, increases both IL-6 mRNA levels and activity of an IL-6 reporter. The mutation of the AP-1 response element in IL-6 reporter fully abolished reporter activity after membrane depolarization, reinforcing the major role of AP-1 in IL-6 expression. (45). Furthermore, the increase in IL-6 expression as a consequence of membrane depolarization was abolished using inositol 1,4,5-trisphosphate (IP\(_3\)) pathway inhibitors, indicating a role for Ca\(^{2+}\) from intracellular deposits (45).

Nerve activity over skeletal muscle, in addition to promoting contraction of muscle fibers, drives a process known as excitation-transcription coupling (ETC) that induces changes in transcription of genes of several pathways (metabolic, structural, endocrine, etc.) (6, 59). We have demonstrated that the
ETC process involves the participation of dihydropyridine receptor (Cav1.1) as the membrane voltage sensor coupled to ATP release from skeletal muscle cells through pannexin-1 hemichannels (6, 11, 24). We have established that extracellular ATP is a relevant mediator between membrane depolarization and signaling pathways leading to gene expression both in rat newborn-derived myotubes and in mouse adult skeletal fibers (11, 16, 42, 43). Extracellular ATP activates metabotropic P2Y receptors; β/γ-subunits of the attached heterotrimeric G protein subsequently activate phosphatidylinositol 3-kinase (PI3K) and phospholipase C (PLC), increasing the intracellular IP3 levels and cytosolic Ca2+ concentration via IP3 receptor (IP3R) activation in the sarcoplasmic reticulum membrane (11, 23, 24). This calcium signal is mainly nuclear in distribution (12) and has been related to expression of early genes (Fos/Jun, Egr1), late genes as IL-6, and structural genes as Tropomin I via activation of several signal transduction cascades (ERK1/2, CREB, NF-κB, AP-1) in skeletal muscle cells (11–13, 16, 45, 62, 85). We have also reported that direct stimulation of rat myotubes with exogenous ATP induces an increment in IL-6 mRNA levels (11). Beyond the role of the depicted transcription factors, we cannot rule out the participation of other players in IL-6 expression in muscle cells. A number of studies in different tissues have demonstrated that expression of IL-6 can be the result of the action of extracellular IL-6 itself through an autoregulatory mechanism (30, 48, 73, 87). After receptor binding, IL-6 induces the activation of Janus-activated kinase 2 (JAK2) tyrosine kinase and the signal transducer and activator of transcription 3 (STAT3) (74). This signaling cascade has been related directly to the expression of IL-6, since pharmacological inhibition of the JAK2/STAT3 pathway in the malignant fibrous histiocytoma cell line blocks both the expression and secretion of IL-6 (73). The same result was obtained when the suppressor of cytokine signaling 3 (SOCS3), a natural inhibitor of the pathway, was overexpressed (73). The expression of IL-6 was also promoted in rat osteoblasts by 100 ng/ml of exogenous IL-6; however, this induction is dependent on the presence of a soluble IL-6 receptor. Moreover, intact response elements on IL-6 promoter for NF-kB, NF-IL-6, and CREB transcription factors are necessary for IL-6 expression, indicating that these proteins play a major role on the expression of the cytokine (30). In muscle tissue, positive feedback for IL-6 has been demonstrated in vivo as well as in vitro (48, 87). In healthy people, an IL-6 infusion right into the femoral artery was demonstrated in vivo as well as in vitro (48, 87). In healthy people, an IL-6 infusion right into the femoral artery was demonstrated in vivo as well as in vitro (48, 87). In healthy people, an IL-6 infusion right into the femoral artery was demonstrated in vivo as well as in vitro (48, 87). In healthy people, an IL-6 infusion right into the femoral artery was demonstrated in vivo as well as in vitro (48, 87). In healthy people, an IL-6 infusion right into the femoral artery was demonstrated in vivo as well as in vitro (48, 87). In healthy people, an IL-6 infusion right into the femoral artery was demonstrated in vivo as well as in vitro (48, 87).

The aim of this work was to confirm that IL-6 expression responds to the events described for the mechanism of ETC in skeletal muscle. Additionally, we examined whether the canonical signaling cascade downstream of IL-6 (IL-6/JAK2/STAT3) also responds to ETC and whether it participates in the autocrine regulation of the cytokine expression. We demonstrated that depolarization of skeletal muscle cells induces not only expression of IL-6 mediated by ATP signaling but also secretion of the cytokine to extracellular medium. Furthermore, expression of IL-6 depends on an IP3-derived Ca2+ signal, acting as an early step to promote a positive IL-6 loop via the JAK2/STAT3 pathway. These results expand the understanding of the ETC mechanisms in skeletal muscle and the regulation of IL-6 expression due to muscle activity.

**EXPERIMENTAL PROCEDURES**

**Reagents**

ATP, ADP, UTP, UDP, apyrase grade VII from potato, suramin, cytosine arabinoside, penicillin, streptomycin, amphotericin B. LY-290042, U-73122, cycloheximide, actinomycin D, and mouse anti-β-actin antibody were obtained from Sigma-Aldrich (St. Louis, MO). Dulbecco’s modified Eagle’s medium–F-12, bovine serum, and fetal calf serum were from Invitrogen (Carlsbad, CA). Collagenase type II was from Worthington Biochemical (Lakewood, NJ). Recombinant rat IL-6 was from PeproTech (Rocky Hill, NJ). Complete Mini Protease Inhibitors were from Roche Applied Science (Indianapolis, IN). Antibodies against p-Tyr705/STAT3 and SOCS3 were from Cell Signaling Technology (Beverly, MA). Anti-rat IL-6-neutralizing antibody was from R & D Systems (Minneapolis, MN). Secondary horseradish peroxidase-conjugated anti-rabbit and anti-mouse antibodies were from Pierce Biotechnology (Rockford, IL). Enhanced chemiluminescence (ECL) reagents were from Amersham Biosciences (Piscataway, NJ). Pharmacological inhibitors 2-aminooxyethyldiphenyl borate (2-APB) and 1,2,3,4,5,6-hexabromocyclohexane (HBC) were from Tocris Bioscience (Bristol, UK). STAT3 inhibitor VIII (5,15-diohenylporphirin) was from Santa Cruz Biotechnology (Dallas, TX). Xestospongin B was kindly donated by Dr. Jordi Molgò (Laboratoire de Neurobiologie Cellulaire et Moléculaire, Institut Fédératif de Neurobiologie Alfred Fessard, CNRS, France). The cell permeant chelator BAFTA-AM was from Molecular Probes (Eugene, OR). Plasmid coding for parvalbumin protein with cytosolic localization (PV-NES-DsRed) was kindly provided by Dr. Manuel Estrada (25). p-DsRed-Monomer control plasmid was purchased from Clontech Laboratories, (Mountain View, CA).

**Newborn-Derived Rat Myotube Culture**

Animal care, manipulation, and procedures were in agreement with protocols approved by the Bioethical Committee of the Facultad de Medicina, Universidad de Chile. Neonatal derived rat myotubes were cultured as described previously (40). Briefly, muscle tissue from the hindlimbs of 12- to 24-h postnatal Sprague-Dawley rat pups was dispersed mechanically and then treated with 0.2% (wt/vol) collage-nase for 15 min with mild agitation. The suspension was filtered through a Nytex membrane or lens tissue paper and spun down at low speed. Ten to fifteen minutes of predesting was performed for the enrichment of myoblasts; cells were plated at densities of 3.5 × 10^3 cells/dish (35 mm). The plating medium was Dulbecco’s modified Eagle’s medium–Ham’s F-12, 10% bovine serum, 25% fetal calf serum, 100 mg/l penicillin, and 50-mg/l streptomycin. After 36 h in culture, fetal calf serum concentration was reduced to 1.8% to induce differentiation. Myotubes in the dish, some spontaneously contracting, with an estimated purity of 90% were visible after the 5th day of culture; these were used for experiments after 5–7 days in culture. When required, transfections were performed on myoblasts on the 4th day. Two micrograms of plasmids was used to transfect cells with Lipofectamine 2000 (Invitrogen) according to the supplier’s instructions.
EXTRACELLULAR ATP EVOKES MUSCLE IL-6 EXPRESSION

Whole Muscle Dissection and Skeletal Fiber Isolation

Flexor digitorum brevis (FDB) muscles were dissected from 5- to 7-wk-old BalbC mice. Either whole muscle treatment with exogenous ATP or skeletal fiber isolation was carried out. Isolated muscle fibers were obtained by enzymatic digestion with collagenase type II, as described by Casas et al. (16). Isolated fibers were seeded in matrigel-coated dishes and used 20 h after seeding.

Cell Treatments

Depolarization assays were performed as reported previously (24). In brief, cells were washed and incubated in Krebs buffer (145 mM NaCl, 5 mM KCl, 1 mM CaCl₂, 1 mM MgCl₂, 5.6 mM glucose, and 10 mM HEPES, pH 7.4) for 30 min. Then, electrical field stimulation (ES) of the whole dish was carried out with a handmaided stimulation device connected to a GRASS S48 stimulator. The tetanus protocol used was 45 Hz, 400 pulses, 1 ms each (24). Alternatively, 0.1–500 µM of exogenous nucleotides (ATP, ADP, UTP, or UDP) was added to culture media by the indicated times. The blockers and inhibitors were preincubated by different time periods and maintained during the stimulus (ATP, ES, or recombinant IL6), as follows: apyrase, 5 min; suramin, 30 min; cycloheximide or actinomycin D, 2 h; HBC or inhibitor VIII, overnight.

Western Blot Analysis

Stimulated cells were lysed in 60 µl of ice-cold lysis buffer (20 mM Tris-HCl, pH 7.4, 1% Triton X-100, 2 mM EDTA, 10 mM Na3VO4, 20 mM NaF, 10 mM sodium pyrophosphate, 150 mM NaCl, 1 mM PMSF, and a protease inhibitor mixture). Cell lysates were separated in 10% SDS-polyacrylamide gels and transferred to polyvinylidene fluoride membranes (Millipore). Membranes were blocked at room temperature for 1 h in Tris-buffered saline containing 3% fat-free milk, with or without 0.5% Tween-20, and then incubated overnight with the appropriate primary antibody. Membranes were incubated with the secondary antibody at room temperature for 1.5 h. The immunoreactive proteins were detected using ECL reagents according to the manufacturer’s instructions. For loading control, membranes were stripped in buffer containing 0.2 M glycine (pH 2) and 0.05% Tween-20 at room temperature for 30 min, blocked as described above, and assessed with the corresponding control antibody.

mRNA Determinations

Total RNA from either skeletal myotubes, isolated fibers, or whole muscles was extracted with TRIzol reagent (19). The reverse transcription (RT) reaction was performed with 1 µg of total RNA using an oligo(dT) primer. Conventional PCR (semi-quantitative) was carried out using forward and reverse primers specific for IL-6, GAPDH, and β-actin. Conventional PCR (semiquantitative) described above, and assessed with the corresponding control anti-GAPDH or β-actin reverse primer was 5'-CCAATTTCCAAATGCTTCT-3'; IL-6 reverse primer, 5'-ACCAAGTGGAGGATTGCAA-3'. GAPDH mRNA amplification was used as the internal control: GAPDH forward primer, 5'-CAACTTTGCGACCTGTTGGAAG-3'; GAPDH reverse primer, 5'-TGGGACACACCCACATCCTT-3'. After an initial 10-min denaturing at 94°C, amplifications were carried out for 25–30 cycles as follows: denaturing at 94°C for 30 s, annealing at 56°C for 30 s, and extension at 72°C for 30 s. After completion of the cycles, a final 10-min extension at 72°C was carried out. PCR products were analyzed by electrophoresis in 1.5% agarose gels. Amplifications without the RT step were made to exclude possible contamination with genomic DNA. Quantitative PCR (qPCR) was performed in Mx3000P Thermocycler (Stratagene, La Jolla, CA) using the Brilliant SYBR Green QPCR Core Reagent Kit, also from Stratagene. The primers used to amplify IL-6 or GAPDH mRNA were the same used in conventional RT-PCR experiments. PCR amplification of the housekeeping gene GAPDH or β-actin was performed as a control. The β-actin forward primer was 5'-CTACAAATGACGTGGTGTTG-3', and the β-actin reverse primer was 5'-TACATGGCGGGGTGTTGA-3'. Expression values were calculated using the 2-ΔΔCt method (55).

ELISA

The concentration of IL-6 released to the culture media at different times was assessed by rat IL-6 Enzyme-Linked Immuno Sorbent Assay Quantikine Rat IL-6 (R & D Systems) according to the manufacturer’s instructions. The absorbance was read at 450 nm (corrected at 540 nm) in a Synergy 2 Multi-Mode Microplate Reader (Biotek). Results were expressed as total picograms of IL-6 at the supernatant per total milligrams of protein in the cell extract (pg IL-6/mg protein).

Statistical Analysis

Data of n experiments were expressed as means ± SE. The significance of difference among treatments was evaluated using a t-test for unpaired data or analysis of variance followed by Dunnett’s posttest for multiple comparisons or by one-way ANOVA test followed by Bonferroni’s posttest. A P value of <0.05 was considered statistically significant.

RESULTS

IL-6 Expression Evoked By ES in Rat Myotubes: Dependence on Extracellular ATP and Nucleotide Receptor Activation

ES of skeletal myotubes evoked a significant and transient increase in IL-6 mRNA levels detected by conventional RT-PCR. Two hours after the stimulus, IL-6 mRNA doubled its basal level (Fig. 1A). Previously, we demonstrated that ATP is endogenously released during myotube or adult fiber ES, acting as a relevant mediator between membrane depolarization and cell signaling leading to gene expression (11, 42). In light of those results, we looked for a possible role of extracellular ATP-mediating IL-6 mRNA expression evoked by ES. Both extracellular ATP/ADP metabolism using apyrase (2 U/ml) and P2Y/P2X nucleotide receptor blockade using the general blocker suramin (100 µM) strongly reduced the rise in IL-6 mRNA expression evoked by ES (Fig. 1, B and C). These results suggest that extracellular ATP, activating P2Y/P2X receptors, is a mediator between membrane depolarization and IL-6 expression changes in skeletal cells. Intereperiment variability in ES-evoked IL-6 mRNA increase was observed, as can be seen comparing Fig. 1, A and B, with Fig. 1C. For that reason, we performed proper controls for each experimental set and compared treatments with their own controls.

Subsequently, we studied the effect of exogenous ATP addition on IL-6 expression detected by conventional RT-PCR. ATP rose IL-6 mRNA significantly in a concentration-dependent manner (Fig. 2, A–C). One-hundred micromolars of ATP increased IL-6 mRNA in just 15 min, reaching a rise of up to sixfold after 45 min of exposure (Fig. 2A). The rise was maintained >90-min incubation with ATP. In the concentration-response curve, 10–500 µM ATP incubated for 30 min significantly increased IL-6 mRNA levels (Fig. 2, B and C). Considering that ATP has the ability to activate all the P2Y/P2X receptor subtypes, we used a pharmacological strategy, tending to discriminate between receptor subtypes involved in IL-6 expression in rat myotubes. ADP, an agonist that activates only some P2Y receptor subtypes (P2Y1, P2Y12, P2Y13), evoked a concentration-dependent rise in IL-6 mRNA, reach-
ES or Exogenous ATP Activates the Canonical IL-6 Signaling Pathway Through IP$_7$-Dependent Calcium Transients and an Autocrine IL-6-Positive Loop

ATP evokes IL-6 secretion that can activate a positive IL-6 loop through the canonical IL-6 receptor pathway. The canonical IL-6 signaling pathway considers sequential activation of plasma membrane IL-6 receptors, JAK2 kinase, and the transcription factor STAT3 (69, 74). An autocrine-positive loop of IL-6 has been widely described in several cell types, where released IL-6 controls its own expression by activation of its canonical pathway (30, 48, 73, 87). In our model, we demonstrated that 100 μM ATP evoked a strong increase in extracellular IL-6 in 1 h, reaching a 10-fold peak after 3 h of incubation (Fig. 3A). To assess whether released IL-6 involves de novo synthesis or a preformed pool, we measured ATP-evoked IL-6 release after 2 h of preincubation with 30 μM cycloheximide (translation inhibitor) or 0.5 μM actinomycin D (transcription inhibitor). Actinomycin D abolished ATP-evoked IL-6 release (from 1 to 4 h); cycloheximide abolished ATP-induced IL-6 at 1 and 2 h, maintaining just a 15–25% release at longer times (Fig. 3A). It can be seen that neither cycloheximide nor actinomycin D altered the extracellular IL-6 content at rest (Fig. 3B) at the maximal incubation time (6 h, corresponding to 2-h preincubation and 4-h treatment). Although 6-h actinomycin D did not modify total protein content of skeletal myotubes, 6-h incubation with cycloheximide reduced it by 15% (Fig. 3C). Four-hour ATP evoked a fivefold increase in IL-6 mRNA, as measured by qPCR, that was totally abolished by actinomycin D (Fig. 4D). Actinomycin D also reduced expression levels at rest (Fig. 4D). Surprisingly, 6-h cycloheximide increased ξ40-fold IL-6 mRNA levels at rest and rose 120-fold ATP-evoked IL-6 mRNA expression (Fig. 4D), suggesting that translation blockade triggers a compensatory response by increasing IL-6 expression levels. Total RNA content of rat myotubes was unaffected by 6-h cycloheximide but reduced by 30% after 6 h with actinomycin D (Fig. 4E).

Considering the IL-6 release, we explored whether a positive IL-6-loop could be controlling IL-6 expression in skeletal muscle cells. In adult mouse isolated skeletal fibers, 0.2–2 ng/ml recombinant IL-6 (rIL-6) increased IL-6 mRNA levels significantly ξ4.5-fold, as measured by qPCR assays (Fig. 4A). To study the canonical pathway involved in IL-6-evoked IL-6 expression in our model, we first assessed the activation of the transcription factor STAT3 by detecting its phosphorylated form (p-Tyr). Addition of rIL-6 to rat myotubes evoked a transient and concentration-dependent STAT3 phosphorylation, reaching a 70% increase after 30 min with 20 ng/ml rIL-6 (Fig. 4, B and C). STAT3 activation evoked by rIL-6 was abolished when myotubes were preincubated with the JAK2 inhibitor HBC (50 μM, overnight), demonstrating a role of this kinase in the signaling pathway activated by this stimulus (Fig.
Control of HBC alone did not change STAT3 phosphorylation levels (not shown).

These results suggest that ATP stimulation evokes de novo synthesis and release of IL6, which could control its own expression, by activating the canonical IL-6 receptor signaling pathway.

ES evokes canonical IL-6 pathway activation mediated by extracellular ATP. ES doubled the STAT3 phosphorylation level after 60 min (Fig. 5A). One-hundred micromolars of suramin abolished STAT3 activation evoked by ES (Fig. 5B), reinforcing the role of extracellular ATP as a mediator between ES and cell signaling related to IL-6 expression. Exogenous ATP (100 μM, 60 min) also doubled the STAT3 phosphorylation level after 4 h evoked a 2-fold increase in IL-6 expression. Incubation of isolated mice flexor digitorum brevis (FDB) fibers with 100 μM ATP for 4 h evoked a 10-fold increase in IL-6 expression. Incubation of mouse whole FDB muscles with 500 μM ATP for 4 h evoked a 2-fold increase in IL-6 expression. In D and E, data were normalized to GAPDH mRNA levels and expressed as fold increase related to nonstimulated conditions (control) (n = 6). Values are expressed as means ± SE. *P < 0.05, **P < 0.01, and ***P < 0.001, analysis of variance followed by Dunnett’s multiple comparison test.

It has been described that the final step of the IL-6 signaling pathway is the STAT3-dependent increase in SOCS3 expression, which acts as a negative regulator of the IL-6 pathway and stops the signaling, avoiding further JAK2 activation (38, 74). We assessed the protein levels of SOCS3 in skeletal muscle cells after stimulation with exogenous ATP at different times. Rat myotubes incubated with 100 μM ATP increased SOCS3 levels in a time-dependent manner, with a peak of 60% increase at 2 h and a complete return to basal levels after 6 h (Fig. 6E).
Fig. 3. Extracellular ATP evoked IL-6 de novo synthesis and secretion. A: ATP increased IL-6 extracellular levels ≤10-fold, depending on transcription and translation processes. Rat myotubes were stimulated with 100 μM ATP after 2-h preincubation, with 30 μM cycloheximide (translation blocker), with 0.5 μM actinomycin D (transcription blocker), or without (w/o) treatment. At different times, 50 μl of supernatant was removed, and IL-6 was quantified by ELISA. B: resting levels of extracellular IL-6 were not affected by 6-h treatment with either cycloheximide or actinomycin D. C: total protein levels from myotube lysates were significantly reduced by 6-h treatment with cycloheximide but not actinomycin D. D: IL-6 mRNA increase evoked by 100 μM ATP (4 h) was abolished by actinomycin D treatment but largely increased after cycloheximide. Cycloheximide treatment increased IL-6 mRNA levels at rest by 40-fold, suggesting that translation blockade activates a positive loop of IL-6 transcription. IL-6 mRNA was measured by quantitative PCR, normalized to GAPDH, and expressed as fold increase related to nonstimulated condition. E: total mRNA levels form myotube extracts were significantly reduced by 6-h actinomycin D treatment but unaffected by 6-h cycloheximide. Values (n = 3) are expressed as means ± SE. **P < 0.01, ***P < 0.001, analysis of variance followed by Dunnett’s multiple comparison test against each “without treatment” value (A–C and E) or Bonferroni’s test (D). NS, not significant.

All of these results suggest that ES, through extracellular ATP, increases IL-6 expression at least in part by activating a positive IL-6 loop.

**IP3-dependent calcium transients are relevant for STAT3 activation promoted by ATP or ES.** Previously, we have demonstrated the relevance of the slow IP3-dependent calcium signal in gene expression evoked by ES in rat myotubes and adult skeletal fibers (16, 43, 45). In the current work, we studied the dependence of STAT3 activation on these calcium signals using molecular and pharmacological approaches. The rise in free intracellular Ca2+ evoked by 40 mM caffeine activated STAT3 at levels similar to those observed previously with tetanic stimulation or ATP (Fig. 7A). To assess whether free intracellular Ca2+ signals were required for STAT3 phosphorylation, myotubes were transfected with a plasmid coding for parvalbumin protein with cytosolic localization (PV-NES-DsRed; Fig. 7B). This tool, used as Ca2+ chelator, strongly reduced basal STAT3 phosphorylation levels as well as ES-evoked STAT3 activation (Fig. 7B). We also tested the blockade of other critical mediators of the slow calcium signal such as PLC, PI3K, and IP3R (11, 16, 23, 24, 39). Ten micromolars of U-73122, a PLC inhibitor, abolished STAT3 activation evoked by ES (Fig. 7C). On the other hand, STAT3 activation evoked directly by exogenous ATP was abolished when free intracellular Ca2+ was chelated with 50 μM BAPTA-AM (Fig. 7D). Either 10 μM U-73122 or 40 μM LY-290042 (PI3K inhibitor) totally blocked ATP-evoked STAT3 phosphorylation (Fig. 7E). STAT3 activation induced by extracellular ATP was also abolished when myotubes were preincubated with 50 μM 2-APB (Fig. 7F), which has been used in previous studies as a blocker of the IP3-dependent slow calcium signal and gene expression in skeletal myotubes (15, 43, 45, 62). The specific blocker of IP3R xestospongion B (5 μM) reduced by 40% the STAT3 phosphorylation evoked by ATP (Fig. 7F). All of these data suggest that the IP3-dependent calcium signal is a critical step in the pathway for STAT3 activation promoted by either ATP or ES.

**DISCUSSION**

In this work, as schematically depicted in Fig. 8, we have demonstrated that electrical stimulation of cultured myotubes induces both expression and secretion of IL-6 mediated by extracellular ATP and the consequent IP3-dependent intracellular Ca2+ signal. IL-6 released into the culture medium would induce the activation of the IL-6 receptor-α (IL-6Rα) through an autocrine mechanism, leading to the activation of transcription factor STAT3. In turn, STAT3 would modulate IL-6 signaling at the transcriptional level either by increasing the synthesis of the cytokine or by promoting the expression of the negative regulator of IL-6 SOCS3.
Muscle contraction depends on the depolarization of the plasma membrane (T-tubules), where Cav1.1 acts as a voltage sensor (63). Membrane depolarization is the primary event for excitation-transcription coupling (9, 22) that triggers Cav1.1 activation and the IP3-dependent Ca2+ pathway (6, 24). Previous results from our laboratory have placed the activation of gene expression, particularly of IL-6, as an event downstream of the myotube membrane depolarization (15, 45, 46). Depolarization of cultured myotubes using a high K+ solution induces the expression of IL-6 (45). By using a more physiological stimulus, we now show expression of IL-6 induced by tetanic electrical stimulation. The kinetics of activation of IL-6 mRNA expression after this stimulus is similar to that obtained with K+ (45). The expression of IL-6 in response to membrane electrical activity has also been studied in other cell types (47, 67, 79). Depolarization of neurons in culture using an extracellular solution of 45 mM K+ evokes an increment in IL-6 expression. The same effect was observed in neuronal-type PC12 cells (67). Furthermore, it has been observed that ischemia-reperfusion injury of rat brain induces expression of IL-6 via activation of voltage-dependent Ca2+ channels (79). The expression of IL-6 in neurons and glia would have a neuroprotective function against damage caused by hypoxia or inflammation (31, 79). All these data suggest that the expression of IL-6 appears to be a general physiological phenomenon in excitable cells, responding to specific demands of each tissue such as the energy stress in skeletal muscle (17, 77) or the need for neuronal protection against ischemia-reperfusion injury, epilepsy, or other brain diseases (47).

Our laboratory has described that electrical stimulation of skeletal muscle cells evokes the release of ATP, subsequent activation of metabotropic purinergic receptors, and the generation of IP3-dependent Ca2+ transients (6, 11, 23, 40). In this work, we have determined that this pathway is required for IL-6 expression. Blockade of the purinergic receptors with the general antagonist suramin, as well as extracellular nucleotide metabolization using apyrase, abolished the increase in IL-6 mRNA induced by depolarization. These results convincingly suggest that the cellular signaling that leads to the expression of IL-6 in electrically stimulated myotubes is mediated by extracellular ATP, indicating that membrane depolarization, ATP signaling, and the expression of IL-6 are part of the same signaling pathway. Numerous reports have related the expression and secretion of IL-6 with extracellular nucleotides (10, 31, 60). Previously, we have reported the expression of IL-6 in skeletal myotubes after 30-min incubation with 500 µM ATP (11). Our results indicate that ATP-induced IL-6 expression is concentration dependent in rat myotubes. In addition, we know that the release of ATP depends on the activity of pannexin-1 channels, suggesting a regulated process of ATP release (11). The results obtained in this study reinforce the idea of a specific and physiological response to ATP in myotubes to promote the expression of IL-6. In this work we did not address the question of which purinergic receptor subtype commands IL-6 expression; however, pharmacological analysis and the fact that the expression of IL-6 and STAT3 phosphorylation depends on IP3/Ca2+ signaling strongly suggest that a P2Y receptor subtype is involved. Four P2Y receptors, P2Y1, P2Y2, P2Y4, and P2Y11, are expressed mainly in myotubes, with P2Y1 and P2Y2 being expressed in higher amounts (11). In addition, recent findings in our laboratory showed that P2Y1 and P2Y4 are also the predominant P2Y receptor subtypes in skeletal fibers isolated from adult mice FDB; both of them are putatively activated by ATP and ADP (28).

We demonstrated an increase in IL-6 mRNA after 100–500 µM ATP in either adult FDB-isolated fibers or adult whole FDB muscle. ATP concentrations at the micromolar range (1) and the measured interstitial ATP concentration of contracting muscles (2 µM) (36). However, it is relevant to note that released ATP plays a role in autocrine...
signaling, acting very locally inside the T-tubule space and at the pericellular space, being rapidly metabolized by ectonucleotidases, which precludes a significant convection into the bulk milieu (44, 90). So the ATP locally released at the T-tubes is probably much higher than what we could measure. For the same reason, to allow ATP diffusion to the T-tubes and surpass ectonucleotidases metabolism, we need to use higher ATP concentrations and time exposures for whole muscle experiments (500 μM, 4 h) than for newborn-derived myotubes (10–100 μM for 15–60 min). All the studies relating P2Y receptor activity with skeletal muscle physiology, using varying experimental models, have used high μM ATP concentrations (50–180 μM) (18, 20, 21, 56, 57, 80). It is relevant to note that, considering that ATP release is a local event, only ≤1% intracellular ATP pool needs to be released to maximally activate all receptors. Thus, extracellular ATP signaling can occur without compromising cell metabolism or essential enzymatic reactions (reviewed in Ref. 70).

It has been described that IL-6 plasma levels range between 1 pg/ml at rest and ~120 pg/ml after strenuous physical performance such as a marathon (27, 29, 49, 76). However, interstitial IL-6 concentration in skeletal muscle is 100-fold higher than plasma levels, ranging between 300 pg/ml at rest and 2,000 pg/ml during and after exercise (49, 65). Very interestingly, concentrations of recombinant IL-6 widely described for in vitro assays are even higher, between 20 ng/ml in C2C12 and L6 myotubes and 120 ng/ml in skeletal muscle strips from patients (14, 32, 88). Considering that in vivo IL-6 is probably secreted from skeletal fibers to the highly packed T-tubule, it is possible that the local IL-6 concentration for autocrine signaling is higher than that measured by microdialysis in muscle interstitium. We assessed from 0.2 to 20 ng/ml rIL6 in our systems and obtained STAT3 phosphorylation starting from 2 ng/ml and IL-6 mRNA expression from 0.2 ng/ml.

From the results obtained in this work, it appears that STAT3 activation evoked by myotube depolarization is totally secondary to IP3-dependent Ca2+ transients. Previous studies have indirectly shown cytosolic Ca2+–induced phosphorylation of STAT3. Gong et al. (34) suggest a link between impaired levels of intracellular Ca2+, caused by a protein from hepatitis C virus, and the activity of STAT3. In another report, the activation of L-type Ca2+ channels by ischemia in rat hippocampus results in the activation of STAT3 (52). Finally, Shi and Kehrl (72) stated the involvement of Ca2+-dependent tyrosine kinase Pyk2 in potentiating the activity of STAT3 induced by epidermal growth factor. However, in none of these works did the authors identify the Ca2+ pool that would participate in the activation of STAT3. Here we show that intracellular free Ca2+ is required for STAT3 activation by either ATP or tetanic stimulation. Considering recent reports suggesting that IL-6 release evoked by skeletal muscle contraction is independent of calcium transients (33, 50), in our system Ca2+ would probably be required for an initial step in IL-6 expression but not for the IL-6 release mechanism. Our results suggest that STAT3 activation mediated by Ca2+ transients is an indirect effect: 1) both electrical stimulation and stimulation with ATP evoke transient IP3-dependent Ca2+ signals (11, 24, 40), 2) both depolarization and ATP induce the expression of IL-6 in skeletal muscle (this work and Refs. 11 and 45), 3) the increased expression of IL-6 by depolarization is dependent on IP3-dependent Ca2+ transients (45), and 4) inhibition of IL-6 signaling cascade blocks STAT3 phosphorylation. Together, these results suggest that intracellular free Ca2+ would be necessary for the expression of IL-6 and that
the cytokine could act as a mediator between cell stimulation and STAT3 phosphorylation.

The fact that STAT3 activation is a late event (1 h post-stimulation) also suggests that there should be a secondary mediator between stimulation of the myotubes and activation of the transcription factor, which is probably IL-6. In other cell types, a late activation of STAT3 as a result of stimuli as varied as the epidermal growth factor (EGF)-like growth factor binding to heparin, isoproterenol, or angiotensin II (51, 68, 91) has been observed. In each of these cases, STAT3 activation was due to prior expression and release of IL-6 to the culture medium and the establishment of an autocrine signaling pathway for the cytokine. It is interesting to note that, in our experiments, when the autocrine IL-6 loop or the canonical IL-6 pathway was blocked using neutralizing IL-6 antibodies or a STAT3 dimerization inhibitor, a 30% residual-activated STAT3 or IL-6 expression evoked by ATP was maintained. These results reinforce the idea of a parallel signaling pathway, most probably Ca2+ dependent, promoting IL-6-expression as a first step to evoke the autocrine IL-6 loop. Indeed, we have

Fig. 6. Extracellular ATP activated STAT3 and increased IL-6 expression partially through a positive IL-6 loop. A: blockade of released IL-6 abolishes STAT3 activation evoked by ATP. Myotubes were incubated with 100 μM ATP for 1 h in the absence or presence of an anti-rat IL-6-neutralizing antibody (aIL6, 1 μg/ml, 30 min before and during the ATP stimuli; n = 4). B: ATP induced STAT3 phosphorylation via JAK2. Rat myotubes were incubated overnight with 50 μM HBC or vehicle. Stimulation with 100 μM ATP for 1 h was performed in the presence of HBC. After that, total protein extracts were obtained (n = 3). C: STAT3 inhibitor VIII reduced STAT3 phosphorylation evoked by ES or ATP. Rat myotubes were incubated overnight with 50 μM inhibitor VIII or vehicle; 1 h after ES or 100 μM ATP addition, total protein extracts were obtained, and phosphorylated STAT3 was detected by WB (n = 4). D: STAT3 inhibitor VIII reduced IL-6 expression evoked by ATP. Myotubes were stimulated as in C but processed for total RNA extraction. IL-6 mRNA expression was assessed by conventional semiquantitative RT-PCR, normalized to GAPDH expression, and presented as fold increase of untreated control cells (n = 3). E: ATP evoked expression of suppressor of cytokine signaling 3 (SOCS3), a negative regulator of the JAK/STAT3 pathway. Myotubes were incubated with 100 μM ATP for different times; total protein extracts were obtained, and SOCS3 expression was assessed by WB (n = 4). Values are expressed as means ± SE. *P < 0.05 and **P < 0.01 (as indicated); #P < 0.05 and ##P < 0.01 (compared against condition with only inhibitor VIII), analysis of variance followed by Dunnett’s multiple comparison test.
demonstrated that IL-6 release evoked by ATP involves de novo synthesis. In our case, the time course of events ending in STAT3 phosphorylation would be as follows:

1) the depolarization of cultured myotubes induces the release of ATP into the culture medium reaching a maximum extracellular ATP 15 s after depolarization (11);
2) the ATP released by depolarization activates P2Y receptors and induces a transient IP3-dependent Ca2+/H11011 signal, which remains high for seconds to minutes in the cytoplasm (11);
3) these Ca2+/H11011 signals induce the activation of transcription factors like AP-1 and NF-kB (15–90 min), which participate in the expression of IL-6 (10 min to 2 h) (15, 85);
4) IL-6 is released into the extracellular medium (hours), reaching an increase of 4.5-fold over baseline 1 h after stimulation with ATP; and
5) finally, IL-6 activates IL-6R, triggering JAK2 phosphorylation, which induces STAT3 phosphorylation and IL-6 expression (hours post-stimulus).

Very interestingly, we noted that cycloheximide treatment intended for protein translation blockade increased resting STAT3 phosphorylation similar to ATP (P < 0.05 vs. control). The participation of IP3-related calcium signaling on phosphorylation of STAT3 was confirmed by experiments using pharmacological inhibitors of the IP3-dependent Ca2+/H11011 pathway (Figure 7).

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Fig. 7. Inositol trisphosphate (IP3)-related calcium signaling mediates STAT3 phosphorylation by ES and ATP. To determine the participation of intracellular Ca2+ signaling on phosphorylation of STAT3, myotubes were incubated with 40 mM caffeine (A) and 100 μM ATP (D–F), or electrically stimulated, as described in EXPERIMENTAL PROCEDURES (B and C) and different inhibitors or blockers of the IP3-dependent Ca2+ pathway were used. A–F, top: representative Western blot of p-STAT3 and β-actin used as loading control. A–F, bottom: bar graph showing normalized levels of phosphorylated STAT3 compared with basal. A: myotubes were preincubated in Krebs buffer for 30 min. The cells were stimulated with 40 mM caffeine by 9 s in the same buffer. After the times indicated, total protein extracts were obtained and Western blot performed (n = 3–4). B: myotubes expressing PV-NES-DsRed plasmid or a control DsRed plasmid were electrically stimulated, and total protein extracts were obtained after 60 min to perform Western blot (n = 3). Cell micrograph shows staining of PV-NES-DsRed (left) or LAP2 immunofluorescence (right) in transfected myotubes. C: STAT3 activation by electrical stimulation is dependent on IP3-signaling pathway. Myotubes were preincubated with 10 μM U-73122 for 20 min, and ES was performed. After 60 min, total protein extracts were obtained to perform Western blot (n = 3). D: myotubes were preincubated in Krebs buffer containing 50 μM BAPTA-AM. The cells were washed twice in the same buffer, and 100 μM ATP was added. After 1 h, the proteins were obtained and Western blot performed. The values are the mean of 2 different experiments. E: to further investigate the participation of IP3 pathway on STAT3 activation by ATP, phosphatidylinositol 3-kinase and phospholipase C inhibitors were used. Cells were preincubated with 40 μM LY-290042 for 30 min or with 10 μM U-73122 for 20 min. Then, the cells were stimulated with 100 μM ATP. The analysis of STAT3 phosphorylation was carried out by Western blot (n = 3–4). F: to see whether STAT3 activation by ATP was also dependent on IP3-Ca2+ pool, cultured myotubes were preincubated or not with the IP3 receptor inhibitor xestospongin B (5 μM, 30 min) or with the IP3 pathway inhibitor 2-aminoethoxydiphenyl borate (2-APB; 50 μM, 30 min). After that, 100 μM ATP was added, and total protein extracts were obtained after 60 min (n = 4). *P < 0.05, **P < 0.01, and ***P < 0.001 vs. basal nonstimulated cells; #P < 0.05 and ##P < 0.01 vs. ATP stimulated without pharmacological inhibitors. Analysis of variance followed by Dunnett’s multiple comparison test (A, E, and F) or Bonferroni’s test (B and C).
levels of IL-6 mRNA ≤40-fold, and ATP-evoked IL-6 expression ≤120-fold. Previously, some authors have reported IL-6 superinduction evoked by blockade of protein synthesis in epithelial cells through increases in both IL-6 mRNA synthesis and stability (26, 37, 64). In the light of our results, considering that actinomycin D totally blocked ATP-evoked IL-6 expression and secretion, the effect over mRNA synthesis is probably more important than that over mRNA stability. This mechanism has never been described in skeletal muscle cells, and it could be a very important control point for IL-6 expression regulation in physiopathological conditions.

We also demonstrated that stimulation with ATP induced a 65% increase in expression of SOCS3 protein. The expression of SOCS3 in response to exercise has been shown previously (82, 83). Analysis performed on young people and older adults shows that the expression of SOCS3 is also much higher in young people; this correlates well with the fact that phosphorylation of STAT3 is higher in older adults (83). Those authors suggest that these alterations in cytokine signaling are explained by reduced expression of SOCS3 in older people. Furthermore, those authors speculate that this signaling potentiation by IL-6 would be involved in impaired muscle regenerative capacity observed in older adults through the establishment of a proinflammatory state. In our case, the expression of SOCS3 reaches the peak after 3 h in stimulated cells, which temporally coincides with the decrease in IL-6 present in the environment, suggesting that SOCS3 is negatively regulating the expression of IL-6. Thus, the negative regulator could have a function of terminating signal to avoid a chronic IL-6 response to exercise.

It is worth noting the fact that myotube depolarization evokes a late (2 h) expression of IL-6 compared with that obtained with ATP (30 min), whereas STAT3 phosphorylation by electrical stimulation occurs at the same time with both stimuli. This point that appears as a discrepancy could be explained by the fact that although we know that electrical stimulation of myotubes, as well as exercise or the activity of motor neurons, causes ATP release from the muscle cells (11, 53, 54, 75), we do not know the effective amount of ATP reaching purinergic receptors on muscle. This is very important considering the different affinity (EC50) that the purinergic receptor subtypes have for ATP and ADP (78, 81). One possibility then is that the purinergic receptors that are activated by exogenous ATP (100 μM) are different (in expression levels or receptor subtype) from the ones activated by ATP released from the depolarized myotubes, thus generating different responses. Another possibility is that exogenous ATP activates purinergic receptors located both at the sarcolemma and the T-tubules, triggering signaling pathways parallel to those evoked by endogenous ATP released at the T-tubules during depolarization. STAT3 phosphorylation prior to increased expression of IL-6 by depolarization could correspond to preformed IL-6, which is stored in vesicles within the cytoplasm of muscle cells, since these vesicles merge with plasma membrane and release their contents upon muscular contraction.

The results presented here contribute to a wider understanding of the ways by which electrical activity of skeletal muscle cells could drive gene expression. IL-6 is a good model to study ETC not only because it responds to membrane depolarization but also because it does in response to IP3-dependent calcium signals that we have related previously to gene expression. IL-6 has a major role in exercise physiology, acting as an energy sensor for skeletal muscle and also acting sys-
terically, helping to avoid metabolic syndrome (66). Future research will shed light on the role of IL-6 in energetic and metabolic balance of skeletal muscle and further unravel details of its regulation.

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AUTHOR CONTRIBUTIONS


REFERENCES


