Resistance exercise load does not determine training-mediated hypertrophic gains in young men

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Running title: Resistance exercise load and hypertrophy

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Abstract

We have reported that the acute post-exercise increases in muscle protein synthesis rates, with differing nutritional support, are predictive of longer-term training-induced muscle hypertrophy. Here, we aimed to test whether the same was true with acute exercise-mediated changes in muscle protein synthesis.

Eighteen men (21±1 yr, 22.6±2.1 kg•m⁻² means±SE) had their legs randomly assigned to two of three training conditions that differed in contraction intensity (% of maximal strength [1RM]) or contraction volume (1 or 3 sets of repetitions): 30%-3, 80%-1 and, 80%-3. Subjects trained each leg with their assigned regime for a period of 10wk, 3 times/wk. We made pre- and post-training measures of strength, muscle volume by magnetic resonance (MR) scans, as well as pre- and post-training biopsies of the vastus lateralis, and a single post-exercise (1h) biopsy following the first bout of exercise, to measure signalling proteins. Training-induced increases in MR-measured muscle volume were significant (P<0.01), with no difference between groups: 30%-3 = 6.8±1.8%, 80%-1 = 3.2±0.8%, and 80%-3= 7.2±1.9%, P=0.18. Isotonic maximal strength gains were not different between 80%-1 and 80%-3, but were greater than 30% -3 (P=0.04), whereas training-induced isometric strength gains were significant but not different between conditions (P =0.92). Biopsies taken 1h following the initial resistance exercise bout showed increased phosphorylation (P<0.05) of p70S6K only in the 80%-1 and 80%-3 conditions. There was no correlation between phosphorylation of any signalling protein and hypertrophy. In accordance with our previous acute measurements of muscle protein synthetic rates a lower load lifted to failure resulted in similar hypertrophy as a heavy load lifted to failure.

Keywords: Skeletal muscle, protein synthesis, motor unit, loading
Introduction

Heavier loading (usually expressed as percentage of a person’s maximal strength or single repetition maximum – 1RM) is often recommended as the optimal way to maximize muscle hypertrophy with resistance training (1). However, there is very little empirical evidence to support this supposition and it is unclear as to the physiological mechanisms by which heavier training loads would provide a signal for greater muscle hypertrophy as compared to, for example, a lighter load lifted to the point of fatigue; both conditions would result in a large amount of muscle fibers being recruited. As proof-of-principle, we recently tested this idea and demonstrated (9) that a single bout of resistance exercise performed at 30% of 1RM to the point of momentary muscle fatigue (failure), was equally as effective in stimulating myofibrillar protein synthesis rates (MPS) as loads lifted at 90% of 1RM (also lifted to fatigue). In addition, the 30%-1RM condition resulted in a more prolonged muscle protein synthetic response with a greater elevation of myofibrillar protein synthesis rates than the 90% of 1RM condition at 24h after exercise (9). We have proposed that the acute exercise-induced increases in MPS that are further augmented with protein ingestion summate and lead to muscular hypertrophy (27). If such a thesis is valid then acute measures of protein synthesis would be, at least qualitatively if not quantitatively, predictive of long-term gains in muscle protein. There is support for this concept as we have shown that measures of acute post-exercise MPS, with differing nutritional support (44), are qualitatively predictive of the training-induced phenotypic outcome (17) between young men consuming milk or soy. Therefore our previous results (9) suggest that resistance training performed at 30% of 1RM to fatigue should result in muscle hypertrophy after chronic resistance training that, based on its prolonged stimulation of myofibrillar protein synthesis, is at least equivalent or greater than the degree of hypertrophy resulting from training with heavy loads. Support for this thesis exists from previous training studies and from a number of studies using low loads with vascular occlusion showing equivalent hypertrophy as that with high loads (36-38).
Other than relative training load, another resistance training variable that is often considered important is the volume of work performed [1, 7, 25]. Since fatigue limits the number of repetitions that can be completed in a set at a given load, varying the number of sets is a common way to adjust training volume. There is currently disagreement concerning the benefit of additional sets for increases in muscle hypertrophy [11, 29, 34]. We have generated acute protein synthesis data from young men showing that 3 sets performed at 70% of 1RM to failure resulted in a greater and more prolonged myofibrillar protein synthetic rate as compared to a single set condition [7]. We speculate that if the model of summative changes leading to hypertrophy [27] is correct, then our data [7] would mean a greater hypertrophic response with 3 sets versus 1 set of resistance exercise.

We also wished to test the thesis that early post-exercise signalling responses, in particular that of p70S6-kinase (p70S6K), would be related to hypertrophy as has been shown in humans [39] and rodents [4]. This is of interest since if the phosphorylation of one single protein is truly predictive of hypertrophy and strength gains then it is certainly worthy of great attention as to its exact mechanistic role in muscle hypertrophy. The overall purpose of the present study was to test whether the acute results we had observed previously [7, 9] would be reflected with longer-term resistance training adaptations. We employed acute measurements of protein signalling to evaluate the relevance of these variables in predicting phenotype.

**Methods**

*Subjects.* Eighteen healthy young men (21 ± 0.8 years, 1.76 ± 0.04 m, 73.3 ± 1.4 kg; means ± SE) volunteered to participate in the study after being informed of the procedures and potential risks involved in the investigation. Subjects were recreationally active with no formal weight-lifting experience or regular weight-lifting activity over the last year. The protocol was approved by the...
Research Ethics Board of Hamilton Health Sciences and McMaster University and was written in accordance with standards set by the *Declaration of Helsinki*.

**Experimental design.** Participants completed 10 weeks of unilateral knee extension resistance training. Each leg was randomly assigned in counter-balanced fashion to one of three possible unilateral training conditions: one set of knee extension performed to voluntary failure at 80% of 1RM (80%-1); three sets of knee extension performed to the point of fatigue at 80% of 1RM (80%-3); or three sets performed to the point of fatigue with 30% of 1RM (30%-3). Each participant trained both legs and was therefore assigned to two of the three possible training conditions. Immediately after each training session subjects consumed a source of high quality protein (PowerBar Protein Plus, 360 kcal, 3.5g leucine 30g protein, 33g carbohydrate, 11g fat; Nestle Nutrition, Florham Park, NJ) in conjunction with ~300ml of water to standardize the postexercise meal and maximize training adaptations.

Before and after the training program, whole muscle volume was measured using magnetic resonance imaging (described below) and changes in muscle fiber area were determined by fiber planimetry with myosin ATPase histochemistry (described below). Knee extension performance was measured by 1RM, maximal voluntary isometric contraction (MVC), rate of isometric force development (RFD) and peak power (described below).

Prior to the first training session, participants fasted overnight and consumed a standard liquid meal (480 kcal, 20g protein, 82g carbohydrate, 8g fat) two hours prior to having resting bilateral muscle biopsies taken from the vastus lateralis. Biopsies were performed with a Bergström needle that was custom-modified for manual suction under local anaesthesia (2% xylocaine). Tissue from this biopsy (~80mg) was used for myosin ATPase histochemistry and Western blotting analysis. Following these initial biopsies, subjects completed their prescribed training session for each leg and immediately consumed a PowerBar Protein Plus bar (Nestle Nutrition, Florham Park, NJ) and 300ml of water. One hour following the completion of the exercise, bilateral biopsies were taken from the vastus lateralis,
for the measurement of acute changes in anabolic signalling molecule phosphorylation status via
Western blot. Following 10 weeks of training a third set of muscle biopsies was taken from each
vastus lateralis muscle and used for fiber area quantification.

*Western blots.* A piece of wet muscle (≈20 mg) was homogenized by hand on ice using a Teflon pestle
in a standard Western blotting homogenization buffer (10 μl mg⁻¹): 25 mM Tris (pH 7.2) buffer
containing 1 mM Na₃VO₄, 50 mM NaF, 40 mM β-glycerolphosphate, 20 mM sodium pyrophosphate,
0.5% v/v Triton X-100, and Complete Protease Inhibitor Mini-Tabs (Roche, Indianapolis, IN, USA)
was used. The samples were centrifuged at 1500 g at 4°C for 10 min. The resulting supernatant was
removed and protein content was determined by the Bradford assay. Equal aliquots of protein were
boiled in Laemmli sample buffer (250mM Tris-HCl, pH 6.8; 2% SDS; 10% glycerol; 0.01%
bromophenol blue; 5%β-mercaptoethanol) for 5 min. Samples (20 μg of protein) were loaded onto 7.5
– 10% SDS-polyacrylamide gels and run for 1.5 h at 150 volts. Gels were then transferred to a PVDF
membrane at 100 volts for 1 h. Membranes were blocked with 5% skim milk powder (w/v) in Tris-
buffered saline with 0.1% Tween (v/v) (TBST). Membranes were then incubated overnight in primary
antibody at 4°C. The following phosphorylation sites were determined: p70S6K-1Thr³⁸⁹ (Santa Cruz
Biotechnology, Inc., Santa Cruz, CA, USA; no. 11759, 1:500 dilution in TBST) mTORSer²⁴⁴⁸ (Cell
Signaling Technology, no. 2971; 1:1000 dilution in TBST), AktSer⁴⁷³ (Cell Signaling Technology, no.
4056; 1:1000 dilution in TBST). After washing in TBST, membranes were incubated in horseradish
peroxidase-linked anti-rabbit IgG secondary antibody (GE Healthcare, Amersham Biosciences,
Piscataway, NJ, USA; no. NA934VS, 1:15,000 dilution in TBST), washed with TBST, and detected
by chemiluminescence (SuperSignalWest Dura Extended Duration Substrate, Thermo Scientific, no.
34075). Images were developed using FluorChem SP Imaging system and quantified by spot
densitometry using ImageJ software. All signalling protein phosphorylation responses were
normalized to their respective total protein.
**Magnetic resonance imaging.** Subjects rested in the supine position for 1 h prior to scanning to prevent the influence of fluid shifts on muscle volume. Also, no strenuous activity was allowed within 24 h of the scanning. MR imaging was performed in a 3T HD scanner (Signa MRI System; GE Medical, Milwaukee, WI) at the Brain–Body Institute, Imaging Research Centre, St. Joseph’s Healthcare (Hamilton, Ontario). Image acquisition was carried out using T1 fluid attenuation inversion recovery (FLAIR) in the axial plane with the following parameters: repetition time/echo time = 2574 ms/6.7 ms; field of view = 25–30 cm; matrix size = range from 320/320 to 512/512 phase/frequency; inversion time = 958 ms; slice thickness = 5 mm. Thigh image acquisition utilized an eight-channel torso coil with two excitations. There was a 10 mm gap between slices. Quadriceps volume was calculated by multiplying the slice area by the distance between slices. Volume was measured from the first slice where the rectus femoris was visible to the first slice where the gluteus maximus was visible. Image J software (U. S. National Institutes of Health, Bethesda, Maryland, USA) was used to determine the area of each slice. Pre- and post scans were performed at the same time of day and joint angle and leg compression was controlled.

**Muscle function.** Subjects completed two testing sessions to assess knee extensor function before and after the 10 week training period. The two sessions occurred in randomized order and were separated from each other and the first training session by three days. Each session started with two sets of submaximal dynamic knee extensions with a load designed not to be fatiguing. Session one consisted of three 5 s unilateral knee extension MVCs conducted on a Biodex dynamometer (Shirley, NY) with one minute of rest between contractions. A knee angle of 90° was used. The highest recorded torque for each leg was taken as the MVC. The analog torque signal for the Biodex was sampled at 2000 Hz with PowerLab 3 data acquisition system (ADInstruments, Bella Vista, Australia) and the RFD was calculated off-line by taking the first derivative with respect to time. The dynamometer was then set in isotonic mode and subjects were instructed to move the load as quickly as possible. Three trials at 20,
30, 40 and 50% of MVC were completed in a random order and the highest instantaneous power achieved in any of the trials was recorded as peak power. Subjects then completed a single set of knee extension to muscle fatigue for each leg. The load used for this test was 30% of 1RM for legs assigned to train with 80% of 1RM and 80% of 1RM for legs assigned to train with 30% of 1RM (to assess the endurance of each subject at their non-trained load). The second visit required subjects to complete a single set to fatigue using the percentage of 1RM which they would train with for the 10 week study. Both the total number of repetitions completed and the total work (the product of repetitions and load) completed were recorded for each test.

Muscle fiber cross-sectional area (CSA). Muscle fibers were oriented vertically by visual inspection and embedded in optimal cutting temperature (OCT) medium. The mounted muscle was frozen in isopentane cooled by liquid nitrogen and stored at −80°C until processing for cross-sectional area analysis. Cross sections (10 μm thick) were cut, mounted on glass slides, and stained using a myofibrillar ATPase histochemistry procedure that uses an acid preincubation pH of 4.6 to distinguish type I and type II fibers as described previously (24, 33). Pictures of the stained slides were taken using a light microscope and NIS Elements 3.0 Imaging Software (Nikon, NY). An average of 55 fibers of each type (type I and II) were outlined using Image J software (U. S. National Institutes of Health, Bethesda, Maryland, USA) for each subject.

Statistics. Our mixed design did not permit us to make within subject comparisons; therefore, between-condition differences (muscle hypertrophy, anabolic signaling, and performance) were tested with a blocked two-factor (condition × time) analysis of variance with repeated measures on time, where applicable. A Tukey’s post hoc test was used to test for significant interactions. For all analyses differences were considered significant at P<0.05. All results are presented as means ± SEM.

Results
Muscle Hypertrophy. Prior to training, quadriceps muscle volume was 39,775 ± 1,836, 40,328 ± 1,640 and 38,189 ± 1,483 cm$^3$ in the 30%-3, 80%-1 and 80%-3 groups, respectively (no differences between conditions at baseline). After 10 weeks of training, the quadriceps muscle volume increased significantly in all groups (P<0.001) to 42,198 ± 1,836, 41,567 ± 1,669 and 40,808 ± 1,410 cm$^3$ in the 30%-3, 80%-1 and 80%-3 groups, respectively. Figure 1 depicts these data expressed as percentage change from baseline. Average type I and type II muscle fiber area increased with training (both P<0.05), irrespective of training condition with no significant between group differences (Table 1).

Western blots. Phosphorylation of Akt at Ser473 was not elevated in any of the conditions. Phosphorylation at mTOR at Ser2448 was elevated above rest at 1 hour post exercise in all conditions (P = 0.0002). p70S6K phosphorylation at Thr389 was elevated 1 hour post exercise in the 80%-1 and 80%-3 groups, but not 30%-3. There was no correlation between the degree of p70S6K phosphorylation at Thr389 changes in muscle volume within any training condition (all P>0.3). Overall, there was no correlation between the degree of p70S6K phosphorylation at Thr389 and the magnitude of quadriceps hypertrophy (r = -0.03, P=0.88; Figure 2, panel D). There was also no correlation between the degree of p70S6K phosphorylation at Thr389 and the magnitude of fiber hypertrophy for either fiber (data not shown).

Muscle Function. Initial unilateral knee extension 1RM was 71 ± 2 kg in the 30%-3 condition, 76 ± 2 kg in the 80%-1 condition, and 73 ± 2 kg in the 80%-1 condition. After the training period all conditions significantly increased in 1RM strength. The increase in 1RM strength was greater in the 80%-1 and 80%-3 conditions compared to the 30%-3 condition (P=0.04; Figure 3). MVC force, knee extension maximal power output and RFD increased in all conditions with no between condition differences (Table 1). The total work that could be completed with 30% of the subject’s 1RM increased with no between condition differences (Figure 4). The total work that could be completed with 80% of the subject’s 1RM increased in all groups. The magnitude of the increase was...
significantly less in the 30%-3 condition compared to the other conditions. The number of repetitions that could be performed with 80% of their current 1RM increased in all groups from 30%-3: 10 ± 1, 80%-1: 10 ± 1, 80%-3: 11 ± 1 pre training to 30%-3: 12 ± 1, 80%-1: 13 ± 1, 80%-3: 12 ± 1 with no between condition differences in the magnitude of the increase. The number of repetitions that could be performed with 30% of 1RM increased in the 30%-3 condition the other conditions did not increase.

Discussion

We have advocated a model of resistance exercise-induced human skeletal muscle hypertrophy that, when supported by adequate nutrition, arises over time due to summed incremental acute increases in muscle protein synthesis that occur after each training session (27, 28). Thus, as a proof-of-principle of this model we tested here whether the acute changes we observed in two previous studies comparing different volumes of work (1 set versus 3 sets of exercise) (7) and divergent intensities of work (high-load versus low-load lifting) (9) would be borne out in a long-term study. We discovered that there was no difference in the magnitude of quadriceps muscle hypertrophy, as determined by both MRI and muscle fiber area, between legs that trained at 30% or 80% of 1RM after ten weeks of knee-extensor exercise. Interestingly, there was no statistical difference in the degree of quadriceps hypertrophy between the 80%-1 and 80%-3 conditions, despite a mean gain in quadriceps volume of ~7% in the 80%-3 condition and only ~3% in the 80%-1 condition (p = 0.18). However, the 80%-3 and 30%-3 showed more than double the average hypertrophy of the 80%-1 condition. These results, while not quantitatively congruent with our acute data (7, 9), are, we propose, broadly supportive of the framework we have proposed of how muscle hypertrophy arises (27, 28). Moreover, our results are actually congruent with a number of other lines of evidence showing that lifting lighter loads, so long as fatigue is induced, lead to roughly equivalent hypertrophy and strength gains (23, 36-38). There are
of course a number of factors, beyond acute changes in muscle protein synthesis, which contribute to hypertrophy. In fact, when subjects are stratified as high and low responders 20-25% subjects exhibit a very limited hypertrophic response whereas the top 20-25% show robust muscle hypertrophy that is four to five times greater than that seen in low responders (17, 26). To date, factors such as changes in microRNA expression, (12) satellite cell number, (26) and intramuscular anabolic signalling protein activation (39) have been shown to be related to the variability in training response, but systemic hormonal factors do not play a role (41-43).

Perhaps the most interesting finding from our work is that hypertrophy in the 80%-3 and 30%-3 conditions was equivalent, which is in contrast to the range of lifting intensities usually prescribed to promote muscle hypertrophy (1, 10). However, the current recommendations (1, 10) ignore a large body of evidence showing that lower loads, when combined, with vascular occlusion, promote equivalent hypertrophy and strength gains as that observed with conventional heavy training (36-38). These results indicate that at least in principle lower load are effective at inducing muscle hypertrophy. Moreover, most work which has investigated different relative resistance training loads has focused on muscle function measurements such as 1RM and relative endurance (5, 6, 13, 35). In one study by Campos and colleagues, (10) eight weeks of training in a 20-28 repetition range did not elicit hypertrophy despite increases in 1RM strength and the number of repetitions that could be completed with 60% of 1RM (10). However, in a subsequent study in which the training methods employed previously by Campos et al. (10) were replicated, equivalent hypertrophy was found in high and low load training groups (23). It is often claimed (1, 10) that high relative training loads are necessary to induce hypertrophy as they are associated with full muscle fiber recruitment and activation of the type II fibers which are known to be more ‘responsive’ to hypertrophy (40). This statement is, however, only accurate during a single effort (repetition) since Henneman’s size principle of motor unit activation states that motor units are recruited in an orderly fashion from smallest to largest with
increasing requirement for force generation (18, 19). Thus, it is true that an isometric contraction performed at 30% 1RM will recruit less muscle than a contraction performed at 80% of 1RM (2). In agreement, it has also been shown that non-fatiguing acute resistance exercise or chronic resistance training results in lower rates of MPS (21) and hypertrophy (20) when compared to heavier resistance exercise loads, respectively. However, when a sub-maximal contraction is sustained, motor units that were initially recruited will fatigue (produce less force) or cease firing completely necessitating the recruitment of additional motor units (15) to sustain force generation. In this way, as the repetitions at lighter loads are repeated, the point of failure/fatigue ultimately necessitates near maximal motor unit recruitment to sustain muscle tension (16). Thus, relatively lighter loads lifted to the point of failure would result in a similar amount of muscle fiber activation as compared to heavier loads lifted to failure (18, 31); however, we acknowledge it is difficult to experimentally verify this motor unit recruitment strategy during voluntary dynamic contractions in humans. In the present study, the average area of both type I and II fibers increased equally with heavy and light relative loads, which is suggestive that both fiber types were recruited during training and to a roughly equivalent extent, at least insofar as the phenotypic hypertrophy response is an indication of this.

The regulation of muscle mass is the result of small changes in net muscle protein balance over time. MPS can be increased by resistance exercise alone however, adequate and properly timed protein consumption is required for a positive net muscle protein balance and thus muscle hypertrophy (8). It is not completely clear how resistance exercise results in increases in MPS as this process is multifaceted. One potentially important step is the activation of p70S6K, which is a downstream protein target of mTORC1 that when activated up regulates initiation of mRNA translation and subsequently increases muscle protein synthesis rates (14). It has been reported that heavy resistance exercise results in an increase in p70S6K phosphorylation (7, 9, 42); our data, at least in the 80%-1 and 80%-3 conditions, broadly agree with these findings. We did not observe an increase in p70S6K
phosphorylation in the 30%-3 condition at 1h post-exercise, but we have observed elevated phosphorylation of p70S6K at 4h after exercise when subjects have completed the same protocol (9). Taken together these results suggest that heavy and light relative loads lifted until the point of failure may result in a different time course of anabolic signalling, with p70S6K phosphorylation occurring later after exercise with light compared to heavy relative loads. Previously, a significant correlation between early (1 h post-exercise) phosphorylation of p70S6K and muscle hypertrophy was observed (39); our results, which include a larger sample size than the previous investigation, failed to support a similar relationship (Figure 2D).

Although relative training load did not impact the magnitude of the hypertrophic response, it did have a clear impact on voluntary isotonic strength gains. Both the 80%-1 and 80%-3 conditions demonstrated a larger increase in 1RM strength compared to the 30%-3 group. These results suggest that practice with a heavy relative load is necessary to maximize gains in 1RM strength of the trained movement. These observations are in line with previous work which has shown that strength gains are specific to the movement that is trained (30) and strength gains are due to a combination of muscle hypertrophy and neural adaptations (32). It should be noted, however, that similar gains in MVC strength, maximal instantaneous power output, and rate of force development were seen across all three training conditions. These data show that hypertrophy is generally beneficial to all strength and power tests that engage the larger muscle. However, it appears that neural adaptations are largely specific to the movement and load used in training (32).

One potential limitation to the unilateral training model employed in this study is the cross education effect where resistance training of one muscle can lead to neurally mediated strength gains in the untrained contralateral muscle (22). The magnitude of the effect varies widely but averages ~7% (22). However, to the author’s knowledge there has been no attempt to quantify this effect when both limbs are training will different protocols. We found no correlation between isotonic strength gains in
the left and right legs ($r=.33$, N.S), suggesting that the cross education effect is minimal or non-existent when both limbs are training with different protocols.

In summary, we report that similar resistance training-induced muscle hypertrophy can result from lifting loads to failure with higher (80% of 1RM) and lower (30% of 1RM) loads than are currently recommended for novice lifters (1). The results from our study also suggest that additional training volume in the form of more sets may result in greater muscle hypertrophy; however, due to the inherent variability in the individual response to resistance training, it appears that longer-term training studies may be required to manifest these differences more clearly. Importantly, these data support the concept that acute increases in rates of MPS are reasonable qualitative indices of the amount of muscle protein gain with similar training as they appear to be with nutrition. However theses gains in muscle mass may be dependent in the adequate provision of amino acids (3). Finally, despite that lack of support for the idea of a hypertrophy-specific load and repetition range, these data confirm the specificity principal of training in regards to muscle strength and endurance.

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443 strength and hypertrophy in a large cohort after weight training. Eur J Appl Physiol. (November 22,

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446 Phillips SM. Consumption of fluid skim milk promotes greater muscle protein accretion after
447 resistance exercise than does consumption of an isonitrogenous and isoenergetic soy-protein beverage.
Table 1. Changes in Knee Extension Performance and Vastus Lateralis Fiber Area Following 10 Weeks of Resistance Training. *Significant main effect pre vs post (p<0.05). RFD= rate of force development, MVC= maximal voluntary contraction. 12 legs in each conditions for all comparisons. Values are presented as means ± SEM.

<table>
<thead>
<tr>
<th>Group</th>
<th>MVC (Nm)</th>
<th>RFD (Nm/s)</th>
<th>Average Type I Fiber area (µm²)</th>
<th>Average Type II Fiber area (µm²)</th>
<th>Percentage Change in Fiber size</th>
<th>Knee Extension Peak Power (W)</th>
</tr>
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<tbody>
<tr>
<td></td>
<td>Pre</td>
<td>Post*</td>
<td>Pre</td>
<td>Post*</td>
<td>Pre</td>
<td>Post*</td>
</tr>
<tr>
<td>80% 1 set</td>
<td>192 ±37</td>
<td>248 ±52</td>
<td>768 ±150</td>
<td>1259 ±256</td>
<td>4532 ±739</td>
<td>16 ± 7%</td>
</tr>
<tr>
<td>80% 3 sets</td>
<td>188 ±42</td>
<td>255 ±47</td>
<td>817 ±101</td>
<td>1110 ±234</td>
<td>4268 ±701</td>
<td>17 ±4%</td>
</tr>
<tr>
<td>30% 3 sets</td>
<td>219 ±37</td>
<td>278 ±52</td>
<td>747 ±186</td>
<td>1248 ±335</td>
<td>4315 ±685</td>
<td>30 ± 12%</td>
</tr>
</tbody>
</table>
Figure 1. Percentage Change in Quadriceps Muscle Volume Following 10 Weeks of Resistance Training. There was a significant main effect for time (increase in quadriceps volume pre to post, P<0.0001). N=12 legs in each condition.

Figure 2. Phosphorylated Akt, mTOR and P70S6K at Rest and 1 Hour Following Resistance Exercise. (A) Akt phosphorylated at Ser 473 expressed relative to total Akt. There are no significant differences between conditions. (B) mTOR phosphorylated at Ser 2448 relative to total mTOR. *Main effect for greater phosphorylated mTOR 1 hour following resistance exercise (P<0.05). (C) p70S6K phosphorylated at Thr 389 relative to total p70S6K. *Significant increases in phosphorylated p70S6K 1 hour after 1 or 3 sets at 80% 1 RM but not 3 sets at 30% of 1RM (p<0.05) (D) Correlation between p70S6K phosphorylation and muscle hypertrophy, r = -0.03 (P=0.88). N=12 legs in each condition.

Figure 3. Knee extension Strength. (A) The maximal load which could be lifted prior to training and after 10 weeks of training. *Significantly greater than prior to training (P<0.05). †Significantly greater than the 3-set at 30% of 1RM condition. (B) Single leg isometric knee extension torque before and after 10 weeks of training. *Significantly greater then prior to training (P<0.05). N=12 legs in each condition.

Figure 4. Maximal Work and Repetitions with 30 and 80% of 1 RM before and After 10 Weeks of Resistance Training. Load product (in arbitrary units) is the product of load in kilograms and repetitions completed. (A) Work completed at 80% of the subjects 1RM (determined within a week of the test). *Significant main effect for an increase in work performed with 80% of 1RM pre to post training (P<0.05). †Significantly less work at 80% 1RM after training in the 3-set at 30% of 1RM condition than the other two groups (P<0.05). (B) Work completed with 30% of the subjects 1RM.
*Significant main effect for an increase in work performed with 30% of 1RM pre to post training (p<0.05). (C) Maximal number of repetitions completed with 80% of 1RM (determined within a week of the test). *Significant main effect for an increase in the number of repetitions performed with 80% of 1RM pre to post training (P<0.05). (D) Maximal number of repetitions completed with 30% of 1RM. *3-set at 30% of 1RM condition completed more repetitions after training than before (P<0.05). †3-set at 30% of 1RM condition completed more repetitions after training then either of the other groups. N=12 legs in each condition.
Figure 2

A. Akt Ser473 (a.u.)

B. mTOR Ser2448 (a.u.)

C. p70S6K Thr389 (a.u.)

D. Quadriceps volume (% change)

Rest 1 Hour

30%-3 80%-1 80%-3

r = -0.03
Figure 3

A

[Graph showing 1RM (kg) with Pre and Post conditions for 30%-3, 80%-1, and 80%-3]

B

[Graph showing Torque (Nm) with Pre and Post conditions for 30%-3, 80%-1, and 80%-3]
Figure 4

- **A**
  - Load Product 80% 1RM (kg * reps)
  - Pre vs. Post
  - 30%-3, 80%-1, 80%-3

- **B**
  - Load Product 30% 1RM (kg * reps)
  - Pre vs. Post
  - 30%-3, 80%-1, 80%-3

- **C**
  - Repetitions at 80% 1RM
  - Pre vs. Post
  - 30%-3, 80%-1, 80%-3

- **D**
  - Repetitions at 30% 1RM
  - Pre vs. Post
  - 30%-3, 80%-1, 80%-3