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Effect of 17 Days of Bed Rest on Peak Isometric Force and Unloaded Shortening Velocity of Human Soleus Fibers

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Abstract

The purpose of this study was to examine the effect of prolonged bed rest (BR) on the peak isometric force (P_o) and unloaded shortening velocity (V_o) of single Ca^{2+} -activated muscle fibers. Soleus muscle biopsies were obtained from eight adult males before and after 17 days of 6° head-down BR. Chemically permeabilized single fiber segments were mounted between a force transducer and position motor, activated with saturating levels of Ca^{2+} , and subjected to slack length steps. V_o was determined by plotting the time for force redevelopment vs. the slack step distance. Gel electrophoresis revealed that 96% of the pre- and 87% of the post-BR fibers studied expressed only the slow type I myosin heavy chain isoform. Fibers with diameter $>100\ \mu m$ made up only 14% of this post-BR type I population compared with 33% of the pre-BR type I population. Consequently, the post-BR type I fibers ($n = 147$) were, on average, 5% smaller in diameter than the pre-BR type I fibers ($n = 218$) and produced 13% less absolute P_o . BR had no overall effect on P_o per fiber cross-sectional area (P_o/CSA), even though half of the subjects displayed a decline of 9–12% in P_o/CSA after BR. Type I fiber V_o increased by an average of 34% with BR. Although the ratio of myosin light chain 3 to myosin light chain 2 also rose with BR, there was no correlation between this ratio and V_o for either the pre- or post-BR fibers. In separate fibers obtained from the original biopsies, quantitative electron microscopy revealed a 20–24% decrease in thin filament density, with no change in thick filament density. These results raise the possibility that alterations in the geometric relationships between thin and thick filaments may be at least partially responsible for the elevated V_o of the post-BR type I fibers.

in the rat, the absence of normal weight-bearing activity results in a rapid decline in the mass of the hindlimb extensor muscles. As little as 6 days of spaceflight or 7 days of ground-based hindlimb suspension is sufficient to reduce the soleus mass of this species by 25–35% (2, 8). These atrophied soleus muscles display altered contractile characteristics, including a reduction in peak force per unit muscle mass, an increase in maximal shortening velocity, and a reduction in peak power output (2, 8). At least a portion of this change in soleus function is the result of alterations in processes of muscle contraction that lie distal to sarcoplasmic reticulum Ca^{2+} release (9, 21, 22, 34).

Humans exposed to chronic non-weight bearing may also experience a reduction in lower limb muscular performance. Declines in voluntary peak isometric ankle extensor torque ranging from ~15–40% have been reported following long- and short-term spaceflight and prolonged bed rest (14, 19). However, due to the complexity of the intact neuromuscular system, it is often difficult to ascribe these changes in human muscle performance to specific physiological mechanisms. The development of effective interventions to combat reductions in muscle function will require a more complete understanding of how the physiological processes of muscle contraction are altered by chronic non-weight bearing.

In the present study, we used single chemically permeabilized muscle fiber segments, activated with saturating levels of Ca^{2+} , to investigate whether prolonged bed rest altered cross-bridge mechanisms of muscle contraction. We were specifically interested in whether the peak isometric force (P_o) and unloaded shortening velocity (V_o) of single soleus fibers were affected by bed rest, since P_o falls and V_o rises for rat soleus fibers

following non-weight bearing (9, 22, 34). The soleus was selected for study because, in the rat, single fibers from this slow-twitch muscle show the greatest functional responses to the absence of weight bearing activity (9). The duration of bed rest examined in this study, 17 days, corresponded with the duration of the National Aeronautics and Space Administration (NASA) Life and Microgravity Sciences (LMS) Space Shuttle mission (STS-78, June 20 to July 7, 1996). This was done to allow future evaluation of bed rest as a ground-based model of spaceflight, since similar single fiber experiments are being conducted on muscle fibers obtained before and after the LMS mission.

METHODS

Subjects

This study was approved by the Institutional Review Boards at Marquette University, the Medical College of Wisconsin, Ball State University, and NASA Ames Research Center. Eight adult males were selected from a pool of volunteers to serve as subjects. Each subject underwent a medical examination and provided written informed consent before participation. The mean ($\pm$ SE) age, height, and pre-bed rest body mass of these individuals were 43 ± 3 years, 182 ± 2 cm, and 82.2 ± 4.3 kg, respectively. For simplicity, we have identified these individuals as *subjects 1–8*. To facilitate comparison to other studies, the present identification numbers 1–8 correspond to NASA identification numbers 428, 439, 435, 249, 405, 457, 374, and 420, respectively.

Bed rest and tissue sampling procedures

Subjects were continuous residents at the Human Research Facility, NASA Ames Research Center (Moffett Field, CA) for 39 days. Their residence was divided into a 14-day ambulatory control period, a 17-day bed rest period, and a 8-day ambulatory recovery period. Subjects remained at a continuous 6° head-down tilt throughout the bed rest period. Compliance was monitored 24 h per day by the staff of the Human Research Facility. All activities, including eating, bathing, excretory functions, and physiological testing, were performed in this position using specially designed facilities and equipment.

One to three days before the start of the control period, a muscle sample was obtained from the left soleus of each subject using the needle biopsy procedure. During the control and bed rest periods, subjects underwent physiological testing using similar equipment and procedures that had been developed for the NASA LMS mission. This testing, which was performed by each subject on two separate days during the control period (*days 2 or 3 and days 7 or 8*) and on three separate occasions during bed rest (*days 2 or 3, days 8 or 9, and days 13 or 14*), included the determination of right ankle extensor isometric torque using a specially designed isokinetic dynamometer and measurement of maximal oxygen consumption during supine cycle ergometry. During each isokinetic testing session, subjects performed 175 voluntary contractions at angular velocities ranging from 0 to 300°/s, broken down as follows: 12 contractions at peak torque (isometric), ~26 contractions at >50% but ≤80% of peak torque, ~77 contractions at >25% but ≤50% of peak torque, and ~60 contractions at ≤25% of peak torque. A complete description of these testing procedures has been presented by Trappe et al. (32, 33). A post-bed rest biopsy sample was obtained from the right soleus on the final day of bed rest before reambulation.

Pre- and post-bed rest muscle biopsies were placed on saline-soaked gauze and divided into several portions. One portion, used for the single-fiber functional experiments described in this study, was immediately submerged in cold skinning solution [composition in mM: 125 potassium propionate, 20.0 imidazole (pH 7.0), 2.0 ethylene glycol-bis(β -aminoethyl ether)-*N,N,N',N'*-tetraacetic acid (EGTA), 4.0 ATP, 1.0 MgCl₂, and 50% glycerol (vol/vol)] and shipped overnight at 4°C to Marquette University, where on arrival it was stored at –20°C. A second portion was pinned at a mild stretch and immersion fixed in a 0.1 M cacodylate buffer (pH 7.2) consisting of 4% glutaraldehyde and 2% paraformaldehyde with 5 mM calcium chloride. This sample was shipped overnight

at 4°C to the Medical College of Wisconsin for osmium postfixation and embedding for electron microscopy as previously described (27).

Single fiber functional experiments

The compositions of the relaxing and activating solutions were determined with the computer program of Fabiato and Fabiato (4) and the stability constants (adjusted for temperature, pH, and ionic strength) compiled by Godt and Lindley (11). Each solution contained (in mM) 7 EGTA, 20 imidazole, 14.5 creatine phosphate, 1 free Mg^{2+} , 4 free MgATP, and sufficient KCl and KOH to produce a total ionic strength of 180 mM and a pH of 7.0. The free Ca^{2+} concentrations of the relaxing and activating solutions had pCa values (where $pCa = -\log$ free Ca^{2+} concentration) of 9 and 4.5, respectively.

All contractile measurements were conducted within the initial 28 days after the biopsy procedure. The methods used to mount the single fiber segments were the same as those used previously in this laboratory (9, 22, 34, 35). Briefly, a single fiber segment was isolated from a muscle bundle, transferred to an experimental chamber, and mounted between an isometric force transducer (Cambridge model 400; Cambridge Technology, Watertown, MA) and a direct current position motor (Cambridge model 300B; Cambridge Technology). The experimental apparatus was attached to the stage of an inverted microscope during the experiments. Sarcomere length was adjusted to 2.5 μm , and the fiber segment length (FL) was recorded. A Polaroid photograph was taken of the fiber while it was briefly suspended in air, and fiber width was measured at three points along the photograph. The mean of these measurements was defined as fiber diameter by assuming the fiber takes on a circular cross section when suspended in air (23). Prior to study, the fiber was briefly bathed in relaxing solution containing 0.5% Brij-58 (polyoxyethylene 20 cetyl ether; Sigma Chemical, St. Louis, MO).

The fiber was activated by transfer from its original chamber, containing relaxing solution, into an adjacent chamber containing activating solution. These solutions were maintained at 15°C throughout data collection. Output from the force transducer and position motor was directed to a digital storage oscilloscope before being amplified and interfaced to a personal computer. Custom software performed on-line analysis and stored data to disks.

Absolute peak force (in mN) was calculated as the difference between resting force, measured while the fiber was in relaxing solution, and the maximal force obtained during activation at pCa 4.5. Normalized peak force (in kN/m^2) was defined as the absolute peak force divided by the fiber cross-sectional area (CSA). Peak stiffness was determined by oscillating the fiber (1.5 kHz) at an amplitude that produced a 0.05% peak-to-peak change in FL (Δ length) as previously described (34). Peak elastic modulus (stiffness/fiber CSA) was calculated as $[(\Delta$ force in pCa 4.5 – Δ force in pCa 9.0)/(Δ length)] $\times$ (FL/fiber CSA).

V_o was determined by the slack test procedure as previously performed in this laboratory (9, 22, 34, 35). The times required for the redevelopment of force after five to six imposed slack steps (each $\leq 20\%$ of FL) were plotted against the corresponding slack length, and the points were fitted with a linear least squares regression line. The slope of this line was V_o that was normalized to the length of the fiber and expressed as FL per second.

Single fiber gel electrophoresis

After the contractile measurements were made, the fiber segment was solubilized in 10 μl of 1% sodium dodecyl sulfate sample buffer and stored at $-80^\circ C$. To determine fiber myosin heavy chain (MHC) composition, ~ 0.5 nl of fiber volume was run on a Hoefer SE 600 gel system consisting of a 3% (wt/vol) acrylamide stacking gel and a 5% (wt/vol) separating gel (21). To determine myosin light chain (MLC) expression, ~ 1 nl of fiber volume was loaded on a gel consisting of a 3.5% acrylamide stacking gel and a 12% acrylamide separating gel (21). All gels were silver stained as described by Giulian et al. (10). A flatbed scanner with a transparency adapter was used to store

an image of each gel on computer disk. Image analysis software (SigmaGel, Jandel Scientific Software) was used to quantify the relative levels of MLC₁, MLC₂, and MLC₃ in each fiber.

Electron microscopy

Longitudinal and thin cross sections (~60 nm) were cut from bundles of fibers and stained with uranyl acetate and lead citrate before examination and photographing in a JEOL 100 CXII electron microscope. Myofilament densities and sarcomere length were determined from cross and longitudinal sections, respectively. Final myofilament density values were adjusted for variations in sarcomere lengths on an individual subject basis (1). Interpretations were made on 40 pre- and 40 post-bed rest well-fixed fibers not damaged during dissection (5 pre- and 5 post-bed rest fibers per subject). Processing artifacts in pre-bed rest controls did not generate the ultrastructural changes observed for the post-bed rest tissues in this report.

Data analysis

Results are presented as mean $\pm$ SE. Pre- and post-bed rest means were compared with analysis of variance. Statistical significance was accepted at $P < 0.05$.

RESULTS

Single fiber MHC expression

Contractile properties and MHC composition were determined for a total of 395 single soleus fibers (227 pre-bed rest, 168 post-bed rest). As illustrated in *lane 1* of Fig.1, the 5% polyacrylamide gel system resolved the slow type I MHC isoform and the two fast MHC isoforms present in adult human skeletal muscle (29). Overall, 218 of the pre-bed rest fibers, or 96% of the group total, and 147 of the post-bed rest fibers, or 87% of the group total, expressed type I MHC exclusively. The remaining fibers expressed either a fast MHC isoform (7 pre- and 16 post-bed rest fibers with *subject 5* accounting for 11 of the fast post-bed rest fibers) or coexpressed slow and fast isoforms (2 pre- and 5 post-bed rest fibers). Typically, the fast isoform present in these fibers was type IIa, although a small number of post-bed rest fibers expressed the type IIx isoform. Because the vast majority of the pre- and post-bed rest fiber populations expressed the type I MHC exclusively, the present analysis focuses entirely on the contractile properties of these fibers.

Fig. 1. Representative 5% polyacrylamide gel illustrating myosin heavy chain (MHC) identification in human single muscle fibers. *Lane 1* was loaded with a human skeletal muscle myosin standard. Each of the remaining lanes was loaded with a single human post-bed rest soleus fiber. Values for unloaded shortening velocity (V_0) of fibers in *lanes 2–4* were 1.03, 0.72, and 0.65 fiber segment lengths (FL)/s, respectively.

Type I fiber diameter and absolute force

A positive correlation was observed between peak Ca^{2+} activated force and fiber diameter for both the pre-bed rest ($r = 0.80$, $P < 0.05$) and post-bed rest ($r = 0.61$, $P < 0.05$) populations (Fig.2). However, on average, the post-bed rest fibers were 5% smaller in diameter ($P < 0.05$) and produced 13% less peak absolute force ($P < 0.05$) than the pre-bed rest fibers (Table 1).

Fig. 2. Relationship between fiber diameter and peak Ca^{2+} -activated force. A: pre-bed rest type I fibers. B: post-bed rest type I fibers. Peak force (mN) was significantly correlated ($P < 0.05$) with fiber diameter for both groups (pre-bed rest, $r = 0.80$; post-bed rest, $r = 0.61$). Number of fibers per group is same as in Table 1.

Table 1. Diameter, peak absolute force, peak normalized force, and unloaded shortening velocity of pre- and post-bed rest type I fibers

Table 1. Diameter, peak absolute force, peak normalized force, and unloaded shortening velocity of pre- and post-bed rest type I fibers

	No. of Fibers	Diameter, μm	P_o , mN	P_o , kN/m^2	V_o , FL/s
Pre-bed rest	218	94 ± 1	0.99 ± 0.02	139 ± 2	0.86 ± 0.02
Post-bed rest	147	$89 \pm 1^*$	$0.86 \pm 0.02^*$	138 ± 3	$1.15 \pm 0.05^*$

Values are means $\pm$ SE. P_o , peak isometric force; V_o , unloaded shortening velocity; FL, fiber length.

Significant difference ($P < 0.05$) between pre-bed rest and post-bed rest means.

Figure 2 suggests that the reduction in the mean diameter of the post-bed rest population was due mainly to a decline in the number of large-diameter fibers making up this group. This is confirmed by the histograms in Fig. 3, in which fibers with diameters $>100\ \mu\text{m}$ made up only 14% of the post-bed rest population in comparison to 33% of the fiber population before bed rest. There was no indication that bed rest further reduced the diameter of the smallest fibers, as only three post-bed rest fibers, or 2% of the post-bed rest population, were observed with diameters that were less than the diameter of the smallest of the pre-bed rest fibers. Only five of the eight subjects showed fiber atrophy with bed rest (Fig.4A). However, the slope of the regression line in Fig.4A was considerably <1.00 , indicating that those subjects with the largest-diameter pre-bed rest fibers showed the greatest decline in fiber diameter with bed rest.

Fig. 3. Frequency distributions of fiber diameter. *A*: pre-bed rest type I fibers. *B*: post-bed rest type I fibers. Data were obtained by collapsing the individual data points in Fig. 2 into diameter bins of $5\ \mu\text{m}$. Note that fibers with diameters $>100\ \mu\text{m}$ made up 33 and 14% of the pre-bed rest and post-bed rest populations, respectively.

Fig. 4. Relationship between pre-bed rest (x-axes) and post-bed rest (y-axes) type I fiber diameter (A, in μm), absolute peak isometric force (P_0 , B, in mN), normalized P_0 (C, in kN/m^2), and V_0 (D, in FL/s) for the individual subjects. Symbols represent means $\pm$ SE values for a particular subject (with the no. adjacent to symbols indicating subject's identification number). Line of identity denoted by solid diagonal line. Significant correlations ($P < 0.05$) were observed between pre- and post-bed rest diameter ($r = 0.79$), absolute P_0 ($r = 0.78$), and normalized P_0 ($r = 0.82$). There was no significant relationship between pre- and post-bed rest V_0 . Slopes of the least squares regression lines (dotted lines) fit to the pre- and post-bed rest data were 0.59, 0.50, and 1.02 for A–C, respectively. Subject sample sizes were as follows (pre-bed rest, post-bed rest, respectively): *subject 1* = 33, 13; *subject 2* = 31, 19; *subject 3* = 27, 21; *subject 4* = 20, 22; *subject 5* = 27, 7; *subject 6* = 25, 19; *subject 7* = 32, 24; and *subject 8* = 23, 22.

Changes in peak absolute force followed a pattern similar to that observed for fiber diameter. Half of the fibers in the pre-bed rest population, but only 27% of the post-bed rest population, produced peak forces >1.0 mN (Fig. 5). Fibers from all but one subject produced less average force after bed rest vs. before (Fig. 4B).

Fig. 5. Frequency distributions of peak Ca^{2+} -activated force. *A*: pre-bed rest type I fibers. *B*: post-bed rest type I fibers. Data were obtained by collapsing the individual data points in Fig. 2 into force bins of 0.2 mN. Fibers with peak force >1 mN made up 51 and 27% of the pre-bed rest and post-bed rest populations, respectively.

Peak normalized fiber force

On average, peak absolute force declined in proportion to fiber atrophy and there was no change in peak normalized force following bed rest (Table 1). The linear regression line describing the relationship between pre- and post-bed rest mean values for individual subjects has a slope of 1.02 and almost overlaps the line of identity (Fig. 4C). However, four individuals displayed a 9–12% reduction in normalized force and one a drop of 4%.

Type I fiber peak stiffness

Stiffness was determined on 104 pre-bed rest and 89 post-bed rest fibers obtained from six of the eight subjects (*subjects 1–4, 7, and 8*). Peak elastic modulus decreased by 22% with bed rest (for pre-bed rest, $2.51 \pm 0.07 \text{ kN/m}^2 \times 10^4$; for post-bed rest, $1.95 \pm 0.06 \text{ kN/m}^2 \times 10^4$; $P < 0.05$). Because no difference in peak normalized force was observed between these 104 pre- and 89 post-bed rest fibers, the ratio of peak isometric tension to elastic modulus increased by 20% with bed rest (51 ± 1 for pre-bed rest and 61 ± 1 for post-bed rest; $P < 0.05$).

Type I fiber V_o

The average V_o of the post-bed rest type I soleus fibers was 34% greater than the pre-bed rest type I mean (Table 1). This overall increase in V_o was the result of a group of high-velocity type I fibers in the post-bed rest population that was absent in the pre-bed rest population (Fig. 6). A closer examination of Fig. 6 reveals that only one pre-bed rest type I fiber had a V_o that exceeded 1.80 FL/s. In contrast, 25 of the post-bed rest type I fibers, or 17%, had a $V_o > 1.80$ FL/s. Every subject showed an overall increase in V_o with bed rest (Fig. 4D). There was no significant relationship between pre-bed rest and post-bed rest V_o for the eight subjects.

Fig. 6. Frequency distributions of fiber V_o . A: pre-bed rest type I fibers. B: post-bed rest type I fibers. Number of fibers for each distribution is same as in Table 1. Only 1% of the pre-bed rest fibers had a $V_o > 1.8$ FL/s compared with 17% of the post-bed rest fibers.

Fiber MLC composition

Figure 7 is a representative 12% gel illustrating the MLC isoform expression of three soleus fibers. In this example, the two soleus fibers expressing type I MHC expressed only slow isoforms of MLC_1 and MLC_2 . Table 2 is a summary of the MLC composition of 182 pre- and 90 post-bed rest type I fibers, all of which expressed only the slow MLC_1 and MLC_2 isoforms. Note that the average V_o values of the pre- and post-bed rest fibers making up this subset were representative of the entire pre- and post-bed rest fiber population.

Fig. 7. Representative 12% gel illustrating myosin light chain (MLC) expression in human single muscle fibers. Single fiber *inright lane* expressed type II MHC and fast isoforms of MLC₁ and MLC₂. Two remaining fibers expressed type I MHC and slow isoforms of MLC₁ and MLC₂. Note lower level of MLC₃ expression in type I fibers.

Table 2. Myosin light chain isoform composition of pre-bed rest and post-bed rest type I fibers

Condition	No. of Fibers	MLC _{1s}	MLC _{2s}	MLC ₃	MLC ₃ /MLC _{2s}	V _o , FL/s
Pre-bed rest	182	0.525 ± 0.004	0.411 ± 0.004	0.064 ± 0.003	0.159 ± 0.009	0.87 ± 0.03
Post-bed rest	90	0.515 ± 0.004	0.407 ± 0.004	0.078 ± 0.004	0.198 ± 0.012*	1.22 ± 0.06*

Values are means ± SE. Myosin light chain (MLC) isoform content was expressed relative to the total MLC of the fiber.

*Significant difference ($P < 0.05$) between pre-bed rest and post-bed rest means.

There was no difference in the relative expression of MLC_{1s}, MLC_{2s}, or MLC₃ between pre- and post-bed rest fibers. Because each myosin S-1 expresses one regulatory MLC (MLC₂) and one essential MLC (either MLC₁ or MLC₃), we examined possible changes in the essential MLC composition of these single fibers by calculating the ratio of MLC₃ to MLC_{2s} (17). This ratio was elevated 25% in the post-bed rest population. However, there was no relationship between fiberV_o and the MLC₃-to-MLC_{2s} ratio either before or after bed rest (Fig.8).

Fig. 8. Relationship between V_o and the MLC_3 content of type I fibers. *A*: pre-bed rest fibers. *B*: post-bed rest fibers. MLC_3 expression is expressed relative to the regulatory light chain content (MLC_{2s}). No significant relationship ($P > 0.05$) was observed between V_o and MLC_3 expression for both the pre-bed rest and post-bed rest populations. Mean values for these data are compiled in Table 2.

Myofibrillar ultrastructure

Electron micrographs of longitudinally sectioned pre- and post-bed rest soleus fibers are presented in Fig. 9. After bed rest, the A bands of soleus myofibrils exhibited normal density, whereas the I bands appeared “moth-eaten.” This morphological alteration indicates that actin filaments were eliminated to a greater extent than myosin filaments with bed rest. An extensive ultrastructural quantitation of the thick and thin filament concentrations demonstrated no average change in thick filament density with bed rest but a 20% reduction in thin filament density in the I band and a 24% reduction in thin filament density in the A band (Table 3).

Fig. 9. Electron micrographs of longitudinal sections of pre-bed rest (A) and post-bed rest (B) soleus fibers. There is a decreased myofibril diameter in the post-bed rest fibers as indicated by the shorter Z lines. Sarcomeres in the post-bed rest fiber (B) are missing myofilaments in the I band regions (arrows), giving the fiber a moth-eaten appearance. Magnification: $\times 17,750$.

Table 3. Myofilament density of pre-bed rest and post-bed rest fibers

Condition	Filaments/ μm^2		
	Thin filaments, I band	Thin filaments, A band	Thick filaments
Pre	$3,027 \pm 130$	$2,559 \pm 60$	987 ± 24
Post	$2,426 \pm 118^{*150}$	$1,950 \pm 67^{*150}$	993 ± 39

Values are means $\pm$ SE of 40 pre- and 40 post-bed rest fibers (5 pre- and 5 post-bed rest fibers per subject).

Sarcomere length was 2.54 ± 0.15 and 2.63 ± 0.13 μm for pre- and post-bed rest fibers, respectively. Filament density values were normalized for variation in sarcomeric length as discussed in METHODS.

^{F3-150}Significant difference ($P < 0.05$) between pre-bed rest and post-bed rest means.

DISCUSSION

Effect of bed rest on fiber diameter and peak isometric force

After 17 days of bed rest, the average diameter of the type I fibers obtained from the soleus (measured at a consistent sarcomere length of 2.5 μm) was reduced by 5%. This change, which is equivalent to a 10% decline in fiber CSA, is almost identical to the 9% reduction in average fiber CSA obtained from the morphological analysis of sectioned muscle biopsy samples from these same eight subjects (32).

Interestingly, we observed a bed rest-induced reduction in the percentage of large-diameter soleus fibers, i.e., those with diameters >100 μm . Because membrane permeabilization is associated with a 20% increase in fiber diameter (12), these large fibers would have in vivo diameters of ~ 83 μm or greater. Related to this, we found that subjects with the largest-diameter pre-bed rest fibers displayed the greatest degree of post-bed rest fiber atrophy. Similarly, Edgerton et al. (3) noted that the greatest fiber atrophy after an 11-day spaceflight was experienced by those astronauts with the largest-diameter preflight fibers.

Our results suggest that the largest pre-bed rest type I fibers either were more susceptible to non-weight bearing and underwent the greatest atrophy or were more likely to express fast MHC isoforms after bed rest and, therefore, eliminated from our post-bed rest analysis. The fact that the percentage of type I fibers fell from

96 to 87% with bed rest would lend support to this second mechanism. However, 67% of the post-bed rest type II or I/II fibers were obtained from a single individual (*subject 5*). If these 12 post-bed rest fibers are eliminated from analysis, type I fibers then account for 95% of the post-bed rest population, a value virtually identical to the pre-bed rest result. The possibility that our post-bed rest type I fiber distributions were biased by this one individual is supported by a morphological analysis of >3,000 sectioned and histochemically stained fibers from these same subjects in which no pre- to post-bed rest change in fiber type distribution was observed (32). Furthermore, the three individuals who had the largest pre-bed rest fibers and displayed the most fiber atrophy (*subjects 2, 7, and 8*) accounted for a combined total of one pre-bed rest and two post-bed rest type II or type I/II fibers. These observations do not support the idea of a selective conversion of the largest type I pre-bed rest fibers to type II or I/II post-bed rest fibers but rather that the largest type I fibers were indeed more susceptible to atrophy.

Because peak Ca^{2+} -activated absolute force was directly correlated with fiber diameter before and after bed rest, the more pronounced atrophy of the largest fibers reduced the percentage of fibers producing relatively high force (>1.0 mN), with essentially no change in the proportion of fibers producing relatively low force. Overall, the average decline in peak force production was proportional to the loss of fiber CSA, although one-half of the subjects displayed reductions in normalized force of $\sim 10\%$.

Effect of bed rest on fiber V_o

On the basis of 5% polyacrylamide gel electrophoresis, all of the pre- and post-bed rest fibers making up this study expressed the adult type I MHC exclusively. The average V_o of these fibers rose by 34% with bed rest. This increase is quantitatively similar to the change in V_o noted for rat type I soleus fibers following 14–21 days of hindlimb suspension (9, 22, 34).

The average V_o of the pre-bed rest fibers, 0.86 FL/s, was greater than the value of 0.52 FL/s we previously reported for 61 human type I soleus fibers (35). However, an examination of individual mean values indicates that several subjects were in quite good agreement with our previous work (35) as well as our preliminary results from the astronauts of the LMS mission (Widrick and Fitts, unpublished observations). In contrast, some subjects (e.g., *subjects 2 and 4*) displayed values that were considerably greater than expected. Although we have no clear explanation for this wide range in pre-bed rest V_o , it should be noted that the eight subjects were extremely heterogeneous in terms of their occupational and recreational activity levels and their pre-bed rest fitness levels. These factors could have contributed to the wide variability in V_o observed in this study. Despite greater than expected variation in pre-bed rest type I fiber V_o , all subjects displayed an increase in this variable after bed rest. Furthermore, there was no significant relationship between a subject's pre-bed rest V_o and the extent to which this variable rose with bed rest (Fig. 4D).

V_o is thought to be limited by the rate of actomyosin cross-bridge detachment (16), a process that has been shown to vary with MHC (17, 26) and MLC (15, 30) isoform expression and the physical properties of the myofilament lattice (23). However, it is not clear which of these processes (or combination of processes) is responsible for the elevated post-bed rest V_o observed in this study.

In the laboratory rat, non-weight bearing is associated with an increase in the number of soleus fibers coexpressing type I and II MHCs (21). Therefore, one possibility is that our gel conditions failed to identify small amounts of fast MHC coexpressed by the post-bed rest fibers. However, Reiser et al. (25, 26) have concluded that V_o is a nonlinear function of the relative proportions of slow and fast MHC isoforms present in a fiber, since slower-cycling cross bridges act as an internal drag on the rate of detachment of faster cycling cross bridges and thereby negate some of the effects of faster-cycling cross bridges. Recent studies conducted on human type IIa, IIa/IIb, and IIb fibers (17, 18) and on rhesus monkey type I, I/IIa, and IIa fibers (R. H. Fitts, S. C. Bodine, J. G. Romatowski, and J. J. Widrick, unpublished observations) suggest that V_o remains relatively unchanged until the

expression of a faster MHC isoform reaches 20–30%, or more, of total MHC. Although it is possible that our electrophoretic conditions failed to detect $\leq 2.5\%$ of the total MHC present in a particular fiber (22), this appears to be an order of magnitude below that required to increase V_o by 34%.

A second possibility is that bed rest modified MLC composition, since alterations in regulatory and essential MLCs are known to have modulatory effects on fiber V_o (15, 30). This mechanism is not supported by the results of the present study. First, all of the type I pre- and post-bed rest fibers expressed similar levels of the slow MLC₂ isoform. Second, although MLC₃ content increased following bed rest, levels of this essential light chain were unrelated to V_o for both the pre- or the post-bed rest type I fibers. In this regard, our pre-bed rest results are in agreement with the work of Larsson and Moss (17), who also reported no relationship between MLC₃ levels and V_o in human slow-twitch muscle fibers.

Another possibility is that bed rest induced de novo or reexpression of an alternative slow MHC isoform. Such an alternative MHC isoform could be responsible for the elevated V_o of the post-bed rest fibers if this isoform had a greater intrinsic V_o than the adult β -MHC and comigrated with the adult β -MHC on our gels. However, even when we employed gel conditions previously shown to resolve two slow MHC isoforms in young adult rat whole soleus muscles (5), we observed only one slow MHC isoform. Thus it seems unlikely that the increased post-bed rest V_o was induced by the expression of a second slow MHC previously observed by Fauteck and Kandarian (5).

A final possibility pertains to the relationship between myofilament geometry and fiber shortening velocity. Riley et al. (27) have reported that, after hindlimb suspension, myofibrils of rat soleus fibers exhibit a moth-eaten appearance, suggesting a reduction in myofilament packing. Because it is known that fiber V_o is extremely sensitive to changes in myofilament lattice spacing (23), we quantified the density of thick and thin filaments in a separate group of histologically fixed pre- and post-bed rest fibers obtained from the original muscle biopsies. Whereas thick filament density remained constant, thin filament density declined by 20–24% in the post-bed rest fibers. In an idealized myofilament, this would be approximated by the loss of one thin filament for every six thin filaments, i.e., a single thick filament would be surrounded by five thin filaments instead of the normal six. Because the area occupied by these five filaments remains constant (no change in myosin density), plane geometry dictates that the average distance from thin to thick filaments increases.

One limitation of this approach is that we do not have myofilament density measurements and physiological data on the same fibers. However, the observed increase in V_o , the decline in elastic modulus, and the rise in the ratio of P_o to elastic modulus are all consistent with an increase in the distance between thin and thick filaments (13, 23). Furthermore, a relatively wide change in V_o coupled with a much smaller change in P_o is similar to results obtained from single fiber preparations in which filament lattice spacing is experimentally altered (23).

Comparison to previous human studies

To our knowledge, this is the first study to investigate the effects of bed rest on the functional properties of human soleus fibers. Previously, Larsson et al. (18) studied human vastus lateralis fibers obtained after 42 days of bed rest and found a reduction in peak normalized force with no change in type I fiber V_o . Comparisons between these two studies must take into account differences in experimental design, particularly the muscles studied and the duration of bed rest. For instance, the soleus and the vastus lateralis are under different anatomic constraints during the bed rest model of non-weight bearing. The subjects in the present study were observed to frequently extend and maintain the foot at a 40° plantar flexed position during bed rest (24). Consequently, the soleus was maintained at a shorter than normal length for much of the bed rest period. In contrast, the vastus lateralis is either at its normal weight-bearing length or stretched (if subjects flex the knee) during bed rest. It is well established that muscles maintained in neutral or lengthened positions do not display the same changes in protein metabolism (20), ultrastructure (27), or contractile function (28) observed for muscles maintained at short lengths. Single fibers obtained from muscles that are chronically shortened may

therefore display different contractile characteristics than fibers from muscles maintained at neutral or lengthened positions.

In the rat hindlimb suspension model, rates of protein degradation and synthesis follow distinct time courses (31). It seems likely that human skeletal muscle protein metabolism may also vary with the duration of bed rest. The fact that we observed no reduction in normalized peak force whereas Larsson et al. (18) observed a 40% reduction in this variable could be the result of an increase in net protein degradation as bed rest is extended beyond 17 days. Interestingly, one-half of our subjects showed a bed rest-induced reduction in normalized force of 9–12%, suggesting that, in many individuals, the loss in contractile protein had begun to exceed fiber atrophy by the 17th day of non-weight bearing.

Finally, to mimic the LMS space shuttle mission, the present subjects performed loaded contractile activity during the isokinetic and aerobic capacity testing sessions. On the basis of the observations of Edgerton et al. (3), it seems unlikely that the aerobic capacity test affected our results. The five sessions of isokinetic testing conducted between the pre- and post-bed rest biopsies averaged out to only 28 contractions per day. Furthermore, only 4 of these 28 contractions were performed above 50% of peak (voluntary) isometric force and of these 4 only 2 were performed as maximal isometric contractions. Clearly, this was insufficient activity to prevent the atrophy and functional changes noted in the single fibers of this study. However, we have no way of knowing whether the observed responses would have been exacerbated if this isokinetic testing activity had been absent.

Comparison to previous animal studies

The most widely studied model of non-weight bearing is ground-based rat hindlimb suspension, with the soleus being the most frequently studied muscle. The 34% increase in post-bed rest type I soleus fiber V_o in the present study is very similar to the 30–50% increase reported for rat type I soleus fiber after 14 days of hindlimb suspension (9, 22, 34). However, the decline in the absolute force of Ca^{2+} -activated rat type I fibers (22, 34) is four times greater than the absolute force deficits noted for the human fibers in the present study. In addition, rat type I soleus fibers show a significant reduction in peak normalized force after non-weight bearing (9, 22, 34). The greater force decline in rat fibers may be the result of interspecies differences in the net rates of protein degradation during non-weight bearing (6, 31).

Individual responses to bed rest

Subject 1 appeared to react to bed rest in a manner that was inconsistent with the responses of the other seven subjects. *Subject 1* displayed no atrophy and the largest increases in both normalized P_o and V_o with bed rest. This individual was also the only subject to show an increase in absolute P_o with non-weight bearing. It is unclear why this individual responded in this way. Regardless of the mechanism involved, elimination of *subject 1* had no effect on a reanalysis of the fiber diameter, P_o , and V_o data of the remaining seven individuals [for 185 pre- and 134 post-bed rest type I fibers, diameters were 96 ± 1 and $90 \pm 1 \mu\text{m}$ ($P < 0.05$), absolute P_o values were 1.02 ± 0.02 and $0.85 \pm 0.02 \text{ mN}$ ($P < 0.05$), normalized P_o values were 140 ± 2 and $137 \pm 3 \text{ kN/m}^2$ ($P > 0.05$), and V_o values were 0.85 ± 0.03 and 1.10 ± 0.05 ($P < 0.05$), respectively].

Summary and conclusions

Seventeen days of bed rest produced an average decline in type I fiber diameter of 5% and a reduction in peak Ca^{2+} -activated force of 13%, with relatively large-diameter, and therefore high-force-producing, fibers being most susceptible to atrophy. On average, fiber atrophy was proportional to reductions in Ca^{2+} -activated force so that peak force per fiber CSA was unchanged. After bed rest, V_o and MLC_3 content were elevated 34 and 25%, respectively, in single fibers expressing type I MHC. However, MLC_3 content was unrelated to V_o in these slow-twitch fibers. The mechanism underlying the elevated post-bed rest V_o is unclear. Possibilities include the

expression of an alternative type I MHC or changes in the geometric properties of the myofilament lattice. Support for the second of these mechanisms comes from our finding of a 20–24% reduction in thin filament density following bed rest.

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AUTHOR NOTES

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