# Cycle training induces muscle hypertrophy and strength gain: strategies and mechanisms (Review)

# H Ozaki<sup>1, 2, 3</sup>, JP Loenneke<sup>4</sup>, RS Thiebaud<sup>5</sup>, T Abe<sup>4</sup>

 <sup>1</sup>Graduate School of Medicine, Juntendo University, Tokyo, Japan
 <sup>2</sup>School of Sports and Health Science, Juntendo University, Inzai, Japan
 <sup>3</sup>Research fellow of the Japan Society for the Promotion of Science, Japan
 <sup>4</sup>Department of Health, Exercise Science, and Recreation Management, School of Applied Science, The University of Mississippi, University, MS, USA
 <sup>5</sup>Department of Kinesiology, School of Education, Texas Wesleyan University, Fort Worth, TX, USA

> Received: February 5, 2014 Accepted after revision: June 26, 2014

Cycle training is widely performed as a major part of any exercise program seeking to improve aerobic capacity and cardiovascular health. However, the effect of cycle training on muscle size and strength gain still requires further insight, even though it is known that professional cyclists display larger muscle size compared to controls. Therefore, the purpose of this review is to discuss the effects of cycle training on muscle size and strength of the lower extremity and the possible mechanisms for increasing muscle size with cycle training. It is plausible that cycle training requires a longer period to significantly increase muscle size compared to typical resistance training due to a much slower hypertrophy rate. Cycle training induces muscle hypertrophy similarly between young and older age groups, while strength gain seems to favor older adults, which suggests that the probability for improving in muscle quality appears to be higher in older adults compared to young adults. For young adults, higher-intensity intermittent cycling may be required to achieve strength gains. It also appears that muscle hypertrophy induced by cycle training results from the positive changes in muscle protein net balance.

Keywords: aerobic exercise, muscular adaptation, lower body, cycling, ergometer

Endurance training is a major part of any exercise program seeking to improve aerobic capacity and cardiovascular health. Another major part of exercise programming is strength/ resistance training, which improves muscle morphology. Thus to improve muscular strength and cardiovascular fitness in young, middle-aged and older populations, the American College of Sports Medicine recommends combining training intensity, volume, and frequency to optimize muscle hypertrophy and strength gain as well as aerobic capacity ( $\dot{VO}_2max$ ) (23). However, the vigorous training intensity and/or high training frequency might hinder some older adults from participating in this type of training program. Interestingly, recent studies have reported concurrent improvements in  $\dot{VO}_2max$  and muscle hypertrophy in young and older populations after single exercise training (27, 62, 64, 65). These single exercise modes include ambulatory exercise (walking, jogging, and running), cycling, and swimming.

Corresponding author: Hayao Ozaki

School of Sports and Health Science, Juntendo University

<sup>1-1</sup> Hiragagakuendai, Inzai, Chiba, Japan

Phone: (+81) 47698-1001; Fax: (+81) 47698-1030; E-mail: ozaki.hayao@gmail.com

Recently, we have summarized whether or not ambulatory exercise produces muscle hypertrophy and strength gain in the lower extremities (65). According to the literature, it seems that relatively long periods, over half a year, of walking and jogging can increase leg muscle size among older adults. However, competitive marathon running and regular high-intensity distance running may not produce leg muscle hypertrophy in young and middle-aged adults, which might be related to insufficient recovery from the muscular damage caused by repeated eccentric contractions during running. Meanwhile, cycling exercise involves mainly concentric contractions and therefore muscular damage is lower in cycling compared with running (59). With respect to muscle damage, cycle training may therefore be better suited for improving muscle size and function compared to running.

To discuss the effect of cycle training on muscle size and strength, cross-sectional and longitudinal studies have been used. In cross-sectional studies, it is known that professional cyclists have larger thigh muscle size compared to controls (36, 52). Hug et al. (36) have shown that total thigh muscle cross-sectional area (CSA), especially vastus lateralis (VL) and biceps femoris (BF) muscle CSA, is larger for professional cyclists than for recreationally-active sport science students. Maximum isometric strength of knee extension is greater in track sprint cyclists than in untrained subjects (52). In addition, both fast- (FT) and slow-twitch (ST) muscle fiber areas in VL are larger in cyclists than that of untrained subjects (25, 52). It is unclear, however, whether the greater muscle size and strength were induced exclusively by cycle training because elite cyclists perform other types of exercise training such as resistance training. Moreover, the influence of genetic factors may have also confounded the observed differences. Unfortunately, it is difficult to differentiate the effect of cycle training on muscle size and strength from these confounding factors with a cross-sectional study design. Therefore, the results of training studies employing untrained subjects needs to be reviewed.

A training intensity of more than 60% of one's concentric repetition maximum (1RM) is commonly considered as the minimum intensity required to achieve muscle hypertrophy under work matched conditions (23). However, in recent years, it has been established that, when performed repetitively or until volitional failure, a low exercise intensity such as 30% 1RM can lead to an increase in myofibrillar protein synthesis (9). These results suggest that high external loads are not a prerequisite for increasing muscle protein synthesis or muscle size (60). Peak muscular activation in VL and vastus medialis (VM) during cycling corresponded to approximately 50% of maximum voluntary contraction (MVC) (19). Therefore, cycle training that consists of repetitive movements may suffice as a minimum stimulus required to increase muscle protein synthesis. In fact, previous studies have shown that protein synthesis acutely (28) and chronically (77) can be stimulated by cycling in untrained subjects. Furthermore, muscle hypertrophy by cycle training is frequently observed when cycle training has been performed for relatively long periods (24, 58, 62). Thus, it is plausible that cycle training does not increase muscle size during short periods (34, 35) but that cycle training requires relatively long periods to induce significant muscle hypertrophy (5, 58).

The primary purpose of this review is to discuss the effect of cycle training on muscle size and strength of the lower extremity, especially thigh muscle mainly activated during pedaling, with three groups of subjects: untrained and healthy young adults, older adults and patients. Furthermore, we also discuss the possible mechanism of muscle hypertrophy induced by cycle training.

#### Methods

#### Literature search

Typical online search using MEDLINE, Web of Science and SPORTDiscuss was performed with the following keywords to obtain relevant articles: 'endurance training', 'cycling', 'cycle', 'ergometer', 'training', 'muscle', 'muscle strength', 'muscle size', 'muscle cross-sectional area', 'protein synthesis', 'concurrent resistance and endurance training', 'concurrent strength and aerobic training', 'combined resistance and endurance training' and 'combined strength and aerobic training'. References from pertinent articles and names of the authors cited were cross-referenced to locate any further relevant articles not found with the initial search.

#### Inclusion criteria

To be included, a study needed to meet the following criteria: (a) Study population: Subjects were untrained healthy young (20–40 years) and older (more than 60 years) adults and untrained patients (more than 20 years) defined as individuals with a cardiovascular and/or muscular disease. Young and older adults could be physically active but could not be participating in regular strength and endurance training. (b) Outcome measures: The study needed to investigate whole muscle size, muscle fiber size, fat-free mass (FFM) and/or muscle strength (1RM, isokinetic and/or isometric strength). FFM and muscle volume estimated by skinfold measurements were excluded. (c) Language: The search was limited to original research that was written in English. Furthermore, to investigate the effect of typical cycle training on muscle size and strength, studies were excluded if cycling was performed with one-leg. Studies were also excluded if cycle training was combined with other interventions such as nutritional and/or blood flow restriction to an exercised muscle. We also discuss the possible mechanism of muscle hypertrophy induced by cycle training, including the articles which were not collected by means of the aforesaid online search procedures.

## Analysis of effect size

Analysis of effect size was performed by reference to previous studies (73, 89) to investigate the magnitude of muscle hypertrophy and strength gain with cycle training. Effect size (ES) was calculated with the following formula: [(posttest mean – pretest mean) / pretest standard deviation], using the data of the searched articles which clearly demonstrated pretest and posttest mean and standard deviation (SD) or standard error (SE) in terms of muscle size and strength. When only SE was reported, SD was calculated from the SE. Differences of ES among the three subject groups and within training design variables (less than vs. more than 40 training sessions, continuous vs. interval training) were evaluated with one-way ANOVA and unpaired *t*-test, respectively. Statistical significance was set at  $p \leq 0.05$ .

## Changes in muscle size and strength induced by cycle training Overall effect size for muscle hypertrophy and strength gain

The ES is presented in Table I and Fig. 1. The 31 ESs for lower limb muscle hypertrophy and 22 ESs for lower body strength development were obtained from 39 studies. The mean ES for muscle hypertrophy was 0.40 (95% confidence interval [CI]: 0.10, 0.71; the number of ESs [*n*]: 18) for young adults, 0.28 (95% CI: -0.31, 0.87; *n*: 6) for older adults and 0.69 (95% CI: -0.07, 1.44; *n*: 7) for patients. A significant difference was not found among the three groups. Meanwhile, the mean ES for strength gain was 0.16 (95% CI: -0.06, 0.39; *n*: 12) for young

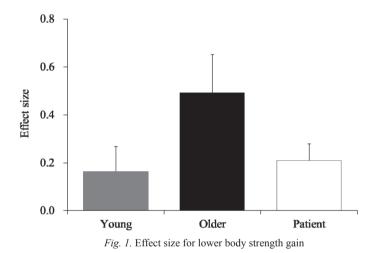
#### Ozaki H et al.

adults, 0.49 (95% CI: -0.01, 1.00; *n*: 4) for older adults and 0.21 (95% CI: 0.03, 0.39; *n*: 6) for patients. Although the value of the older adults tended to be higher compared to the other two groups, a significant difference was not found among the three groups.

	Overall		Num	ber of t	training sessions		
			< 40		40 ≦		
	Mean (95% CI)	N	Mean (95% CI)	N	Mean (95% CI)		N
Young (Y)	0.40 (0.10, 071)	18	0.21 (-0.13, 0.54)	13	0.91 (0.34, 1.48)	*	5
Older (O)	0.28 (-0.31, 0.87)	6	ID		0.41 (-0.63, 1.45)		4
Y+O	0.37 (0.12, 0.62)	24	0.18 (-0.11, 0.47)	15	0.69 (0.24, 1.13)	*	9
Patient (P)	0.69 (-0.07, 1.44)	7	0.63 (-0.32, 1.59)	5	ID		
Y+O+P	0.44 (0.20, 0.68)	31	0.29 (0.01, 0.58)	20	0.71 (0.27, 1.15)		11

Table I.	Effect	size	for	muscle	hy	pertrop	ohy

CI: confidence interval; N: number of effect sizes; ID: insufficient data (< 4 Effect sizes); \* p < 0.05, vs < 40



## Untrained young adults

With respect to the probability of muscle hypertrophy in young adults, 8 out of 22 studies evaluating muscle size have reported that cycle training induced thigh muscle hypertrophy at the whole muscle level and/or muscle fiber level for young adults. It is suspected that training design variables could be the key determinants for these hypertrophic effects. One study has shown that muscle fiber hypertrophy was observed in cycle training at 75–85% HRmax for 30–60 min, 4 days per week, (62) whereas another study demonstrated no muscle hypertrophy with cycle training at an exercise intensity varying from the ventilatory threshold to 90%  $VO_2max$  for 21–42 min, 3 days per week (5). There were not large differences in exercise intensity, duration and frequency between the previous two studies but the program period of the former was approximately two times longer than that of the latter. As summarized in

Table II, muscle hypertrophy is more likely to take place when the training period or the total number of training sessions is greater. Moreover, this trend is kept consistent regardless of the exercise protocol used, although the previous studies investigating the effect of cycle training on muscle size are broadly divided into two types; continuous or interval training. For continuous cycling, 2 of 9 studies with less than 40 training sessions have shown muscle hypertrophy at the whole muscle or muscle fiber level, while 3 of 4 studies with equal to or more than 40 training sessions have observed it. Muscle fiber area tended to increase in both ST (17%, ES: 2.09) and FTa (11%, ES: 0.77) fibers in the only study without muscle hypertrophy (15). Meanwhile, for interval cycling, both studies with equal to or more than 40 training sessions have shown muscle hypertrophy but no studies with less than 40 training sessions have observed significant changes. Therefore, program period would appear to be the key determinant for the probability of muscle hypertrophy with cycle training in young adults. Meanwhile, similar to the probability of muscle hypertrophy, program period appears to be the key determinant for the magnitude of change. The mean ES for muscle hypertrophy in previous studies with less than 40 training sessions was 0.21 (95% CI: -0.13, 0.54; n: 13), whereas that with equal to or more than 40 training sessions was 0.91 (95% CI: 0.34, 1.48; *n*: 5), and the value of the latter was significantly higher than that of the former.

To better determine the reason why cycle training requires a relatively longer period of training until increases in muscle size are observed, we compared the magnitude of muscle hypertrophy between cycle and resistance training. Mikkola et al. (58) compared the percentage of muscle hypertrophy between resistance training, continuous cycle training and the combination of both. As a result, muscle CSA of the quadriceps femoris significantly increased by 6% for the resistance training group and only by 2% for the cycle training group after the same training period (42 training sessions). In other words, the magnitude of muscle hypertrophy with cycle training appears to be one third of that of resistance training. However, the calculation of percentage increases cannot be accurately compared either within or across research studies because percent change does not take into consideration the variance of muscle hypertrophy among subjects (73). Therefore, we calculated the ES for muscle hypertrophy with the previous data of McCarthy et al. (57) and Bell et al. (5). The ES was 0.56–1.17 for lower body resistance training whereas only 0.16–0.39 for continuous cycle training despite a similar training period and frequency. Therefore, it is plausible that cycle training requires a longer training period than typical resistance training until significant increases in muscle size can be observed because of a much slower hypertrophy rate. It is also possible and likely that there are differences in the intrinsic muscular environment between resistance and cycle training such that although increases in muscle size occur with cycling, those changes may never reach the magnitude observed with resistance training.

In addition to the probability of muscle gain for young adults, significant increases were observed in 1RM and/or isokinetic and/or isometric strength in 4 out of 11 studies. This is similar to that found for changes in muscle size. However, the key determinant for the strength gain is unlikely to be identical to that of muscle hypertrophy. While the occurrence of muscle hypertrophy depends on the program period or the total number of training sessions, strength gain following cycle training is more likely to be influenced by the exercise type and intensity. In previous studies evaluating muscle strength, 3 out of 4 studies using maximal or submaximal interval cycling resulted in strength gain in the lower limb muscle. However, with continuous cycling, this adaptation was observed only for 1 out of 7 studies using continuous cycling. Therefore, it appears that exercise intensity or effort, rather than the training period, is the key factor for strength gain after cycle training in untrained young

ing adults
ned you
ı untrain
<sub>2</sub> max in u
0
20
as
ell
ž
~
as
ų
ad
en
tt.
s
and
size
e
5
n muscl
Ē
l nc
ã
Ē
E
a
Ħ
en
:=
S.
<u>ي</u> :
of
ts e
ct
ffe
ef
le
Th
Ш.
le
ab
$T_{\ell}$

mass         II           aak         -           ined         ns           x         -           x         -           x         -           x         -           ns (-2)         SQ           ns (1)         -           x         ns (0)           x         ns (0)           ns (-1)         ns (-1)           ns (-1)         -	Training	Body	M	Muscle strength (%)	(%)			Mu	Muscle size (%)	(%)		<b>∀O₂max</b>
is cycling         (sessions)         (sessions) $vic cycling$ $vic cycling$ $vic cycling$ $vic call$ $M$ $8 wk$ $54-65\%$ HRpeak $$ $vic call$ $M$ $8 wk$ $54-65\%$ HRpeak $$ $vic call$ $M$ $8 wk$ $54-65\%$ HRpeak $$ $vic call$ $M$ $6 wk$ $15-30 \min$ $$ $vic call$ $M$ $6 wk$ $72\%$ $VO_2 max$ $$ $vic call$ $M$ $6 wk$ $72\%$ $VO_2 max$ $$ $vic call$ $M$ $6 wk$ $70\%$ HRR $ns (-2)$ $SQ$ $vic call$ $M$ $8 wk$ $30 \min$ $$ $$ $vic call$ $M$ $8 wk$ $30 \min$ $$ $$ $vic all$ $M$ $8 wk$ $30 \min$ $$ $$ $vic all$ $M$ $8 wk$ $30 \min$ $$ $$ $vic all$ $M$ $8 wk$ $80\%$ $00\%$ $$ <		mass	1RM	Isokinetic	Isoi	Isometric	mCSA		mfCSA (VL)	/L)	FFM	(%)
uns cycling         i       (24) $2-5 d/wk$ $54-65\%$ HRpeak          i       (24) $2-5 d/wk$ $15-30 min$ i       (24) $2-5 d/wk$ $15-30 min$ i       (24) $2-5 d/wk$ $15-30 min$ i       (28)       (30) $30 min$ i       (28)       (30) $30 min$ i $6 wk$ $72\%$ $VO_2 max           i       6 wk 72\% VO_2 max           i       6 wk 72\% VO_2 max           i       6 wk 72\% VO_2 max       ns (-2)       SQ         i       30 min            i       (29) (30) 3 d/wk 40 min          i       (21) 4 d/wk 40 min ns (-1)          i       (21) (22) 5 d/wk 60\% VO_2 max ns (0)          i       (22) 5 d/wk 60\% VO_2 max ns (0)       <$											(DEXA)	
matrix         M         8 wk         54-65% HRpeak            (24)         2-5 d/wk         15-30 min         -           (28)         (28)         (28)         maximal maintained         ns           (1)         (24)         2-5 d/wk         maximal maintained         ns           (28)         (30)         30 min         -         -           (29)         (30)         30 min         -         -           (20)         (30)         30 min         -         -           (21)         4 d/wk         45 min         -         -           (21)         3 d/wk         40 min         -         -           (21)         4 d/wk         60 min         -         -           (21)         (32)         60% VO_2max         ns (1)         -           (22)         5 d/wk         60 min         -         -												
) (24) $2-5 d/wk$ I5-30 min ret al. M 6 wk maximal maintained ns F 5 $d/wk$ power (28) (30) 30 min et al. M 6 wk 72% $VO_2max$ (29) (30) 30 min (29) (30) 30 min y et M 10 wk 72% $VO_2max$ (30) 30 min (29) (30) (30) 10 wk 70% HRR ns (-2) SQ (30) 30 min (32) $d/wk$ 45 min (32) $d/wk$ 45 min (32) $d/wk$ 45 min (32) $d/wk$ 60% $VO_2max$ ns (1) (32) $d/wk$ 60 min (33) $d/wk$ 60 min (35) F (22) 5 $d/wk$ 60 min (84) (24) 5 $d/wk$ 1 hour (84) (24) 5 $d/wk$ 1 hour (35) F (22) M 12 wk $VT-90\% VO_2max$ ns (0) (35) F (35) $f$ (35) (35) (35) (35) (35) (35) (35) (35)					KE	su				ns		ns (–10)
Tet al.         M $6 \text{ wk}$ maximal maintained         ns           (28)         (30)         30 min         90wer          -           (28)         (30)         30 min          -         -           (29)         (30)         3 d/wk         45 min         ns (-2)         SQ           2002         (27)         3 d/wk         40 min         ns (-2)         SQ           1 et al.         M         8 wk         80% VO2max         ns (1)         -           al.         M         7 wk         60 min         ns (1)         -           (21)         4 d/wk         40 min         ns (1)         -         -           (32)         5 d/wk         60 min         ns (1)         -         -           (33)         F         -         -         -         -         -           sky et         M         7 wk         60 win<												
) F 5 $d/wk$ power et al. M 6 $wk$ 72% $VO_2max$ - (28) (30) 30 min F 5 $d/wk$ 30 min (29) (30) (30) 30 min y et M 10 wk 72% $VO_2max$ - (30) 3 $d/wk$ 45 min (31) 4 $d/wk$ 45 min (32) 4 $d/wk$ 40 min (32) 5 $d/wk$ 60% $VO_2max$ ns (1) (32) 5 $d/wk$ 60 min (33) 7 $wk$ 60% $VO_2max$ ns (1) ns (-1) (33) 7 $wk$ 60% $VO_2max$ ns (0) ns (-1) (33) 7 $wk$ 60% $VO_2max$ ns (0) ns (-1) (32) 5 $d/wk$ 10 min (84) (24) 5 $d/wk$ 10 min F $M$ 12 $wk$ $VT-90\% VO_2max$ ns (0) ns (-1) (35) 7 $d/wk$ 10 min ns (-1) $ws$ (-1)	-	ns							ns	ns (–3)		14
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$												
mtet al.         M         6 wk         72% VO <sub>2</sub> max          F         5 d/wk         30 min          F         5 d/wk         5 d/wk         45 min         F         SQ												
) F 5 d/wk 30 min yet M 10 wk 70% HRR $ns(-2)$ SQ 2002 (27) 3 d/wk 45 min 2002 (27) 3 d/wk 40 min (21) 4 d/wk 40 min (21) 4 d/wk 60 min (22) 5 d/wk 60 min (33) $ns(-1)$ (35) $ms(-1)$ (35) $ms(-1)$ (36) $ms(-1)$ (37) $ms(-1)$ (37) $ms(-1)$ (38) $ms(-1)$ (38) $ms(-1)$ (38) $ms(-1)$ (38) $ms(-1)$ (38) $ms(-1)$ (38) $ms(-1)$ (38) $ms(-1)$ (37) $ms(-1)$ (38) $ms(-1)$									I	ns		14
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$								Π		ns		
y et         M         10 wk $70\%$ HR         ns (-2)         SQ           2002         (27)         3 d/wk         45 min         20         2           1         (30)         3 d/wk         45 min         6         10         10           1 et al.         M         8 wk         80% $VO_2max$ ns (1)         10           1 et al.         M         8 wk         80% $VO_2max$ ns (1)         10           1 d/wk         60% $VO_2max$ ns (1)         10         10         10         10           1 d/wk         60 min         60 min         10 <td< td=""><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td>II B</td><td>ns</td><td></td><td></td></td<>									II B	ns		
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	70% HRR			KE ns (-5)	) KE	ns (–2)	QF	Э	I ns	ns (4)		18
1 et al.     M     8 wk     80% $VO_2 max$ ns (1)       1 et al.     M     8 wk     80% $VO_2 max$ ns (1)       (21)     4 d/wk     40 min     (1)       (32)     5 d/wk     60% $VO_2 max$ ns (0)       (32)     5 d/wk     60 min     (1)       (35)     7 wk     60 min     (1)       (22)     5 d/wk     1 hour     (1)       (84)     (24)     5 d/wk     1 hour       (82)     (23)     1 hour     (1)       F     3 d/wk     7.4.2 min     (1)									II ns	ns (5)		
et al. M 8 wk 80% $\dot{V}O_2$ max ns (1) (21) 4 $d/wk$ 40 min (32) $(32)$ $\delta d/wk$ 60% $\dot{V}O_2$ max ns (0) F (22) $5 d/wk$ 60% $\dot{V}O_2$ max ns (0) F (35) $60 min$ ns (-1) (35) $7 wk$ 60% $\dot{V}O_2$ max ns (0) ky et M 7 wk 60% $\dot{V}O_2$ max ns (0) F (22) (35) $F$ (35) $r (-1)$ ns (-1) F (22) $3 d/wk$ 1 hour ns (0) F (22) $3 d/wk$ $VT-90\% \dot{V}O_2$ max - KE												
$ \begin{array}{c ccccccccccccccccccccccccccccccccccc$	80% VO,max	1s (1)							I ns	(20)		16
al. M 7 wk $60\% VO_2 max$ ns (0) F (22) 5 d/wk $60\% VO_2 max$ ns (0) F (35) 7 d/wk $60 \min$ ns (-1) (35) 84) 7 wk $60\% VO_2 max$ ns (0) ky et M 7 wk $60\% VO_2 max$ ns (0) F (35) 1 hour ns (0) F (35) 7 mour ns (0) F								П		17		
al. M 7 wk 60% VO <sub>2</sub> max ns (0) F (35) 5 d/wk 60 min (35) (35) (35) (35) (35) (35) (35) (35) (35) (35) ky et M 7 wk 60% VO <sub>2</sub> max ns (0) ky (24) 5 d/wk 1 hour (35) (35) (35) (35) F (35) (35) (35) (35) F (35) (35) (35) (35) (35) (35) F (35) (35) (35) (35) (35) (35) (35) (35)								Π	II B	28		
(22)         5 d/wk         60 min         ns (-1)           F         (35)         (35)         ns (-1)           ky et         M         7 wk         60% VO <sub>2</sub> max         ns (0)           ky et         M         7 wk         60% VO <sub>2</sub> max         ns (0)           84)         (24)         5 d/wk         1 hour         ns (0)           F         (35)         1 hour         ns (-1)           F         3 d/wk         71-40% VO <sub>2</sub> max         ns (-1)	60% VO <sub>2</sub> max	1s (0)							ST ns	ns (31)	ns (0)	17
$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$								_		ns (19)		
$ \begin{array}{ c c c c c c c c c c c c c c c c c c c$		:										
et M 7 wk 60% VO <sub>2</sub> max ns (0) (24) 5 d/wk 1 hour F (35) R (22) M 12 wk VT-90% VO <sub>2</sub> max - KE	n	s (-1)							ns	ns (16)	ns (0)	30
$ \begin{array}{cccccccccccccccccccccccccccccccccccc$		(0)									(0) 54	c
(27)         5 www.         1 nour           F         (35)         1 nour           (35)         (35)         ns (-1)           M         12 wk         VT-90% VO_2max         -           F         3 d/wk         21-43 min         -	$60\% VO_2 max$	(U) St									(0) su	у
F (22) (22) M 12 wk VT-90% VO <sub>2</sub> max - KE 7.47 min												
(22) (22) (22) (22) (22) (22) (22) (22)	1	s (-1)									ns (0)	14
L. M 12 wk VT–90% VO <sub>2</sub> max – KE F 3 d/wk 21–42 min		~									~	
F 3 d/wk 21_42 min										ns (11)		5
			ns (14)						II ns	ns (11)		13
(22) (36)												

<b>ỳO₂max</b>	(%)			12	6	5	15	20	13		13	
	FFM	(DEXA)										
ize (%)	mfCSA (VL)		su su				ns (17) ns (11)	20 8 8	23 ns (–8)		ns (6) ns (12) ns (12)	ns (10) ns (6)
Muscle size (%)	mfCS		I II A				ST FTa	I II A II B	ST FT		I II A II B	ST FT
	SA					7						
	mCSA					QF						
	Isometric				SD	ns(4)						
(0)	Iso				KE	KE						
Muscle strength (%)	Isokinetic							14				
scle st	Isok							KE				
Mu	1RM			ns ns		ns(1)						
	-			LP KE		KE						
Body	mass		I	I	I	ns (1)		ns (–3)	I		ns (0)	I
Training			VT–90% VO <sub>2</sub> max 21–42 min	60–80% VO <sub>2</sub> max 30–90 min	35-70% maximal load 30 min	A Aer–AnT 30–90 min	70–80% VO <sub>2</sub> max 60 min	75–85% HRmax 30–60 min	75–90% VO <sub>2</sub> max 1 hour		75 g/kg TBM 3×30 s supramaxi- mal sprint	75 g/kg body weight 2-6×15 s + 2-6×30 s maximal sprint
Period	frequency	(sessions)	12 wk 3 d/wk (36)	12 wk 3 d/wk (36)	10 wk 4 d/wk (40)	21 wk 2 d/wk (42)	20 wk 4 d/wk (80)	20 wk 4 d/wk (80)	20 wk 4 d/wk (80)		6 wk 2–3 d/wk (15)	6 wk A 2.5/wk (A 15)
Sex	(Age)		M (23) F (22)	M (23)	M (22)	M (37)	M (22)	M (30)	M (28– 40)		M (23)	M F (22)
Author			Putman et al. 2004 (72)	Parcell et al. 2009 (66)	Kaljumae et al. 1994 (41)	Mikkola et al. 2012 (58)	Denis et al. 1986 (15)	Nelson et al. 1990 (62)	Gollnick et al. 1973 (24)	Interval cycling	Allemeier et al. 1994 (1)	Jacobs et al. 1987 (39)

Table II. (cont.)

ХВ											
<b>ỳO₂max</b>	(%)				I	Ι	1	ns (-1)	10	1	Ι
	FFM	(DEXA)									
ize (%)	mfCSA (VL)						ns (7)	-11 ns (-6)		ns (9) ns (16) 45	SI
Muscle size (%)	mfCS							ST FT		ST FTa FTb	П
	mCSA							ns (3) ns (-1)	(9–)	6 5	
	mC							TH R L	QF	Ex VL	
	Isometric		7 ns (2)	~ ~	ns (3) ns (-5)	ns (5) ns (5)	٢				
(%	Iso		HE KE				KE				
Muscle strength (%)	Isokinetic								18		ns ns
iscle st	Isok								KE /CSA		KE LP
Mı	1RM										
	1R										
Body	mass					I	1	1	1	ns (2)	ns (1)
Training			5 maximal efforts loads equal 10% of body weight	5 maximal efforts loads equal 5% of body weight	5×3 min 250 W, 80 rpm	5×3 min 250 W, 45 rpm	6% of body mass 3×3-s maximal sprints 8-16 sets	2×8–13×5 s maxi- mal sprint	90% ÝO <sub>2</sub> max 500 kcal/d	8% of body mass 2×15×5 s maximal sprint	65 g/kg of body mass 2-4×4-5×10 s maximal sprint
Period	frequency	(sessions)	4 wk 4 d/wk (16)				6 wk 4 d/wk (24)	7 wk 4 d/wk (28)	7 wk 5 d/wk (35)	10 wk 4 d/wk (40)	14 wk 3 d/wk (42)
Sex	(Age)		22			- 23	M (22)	M F (22)	M (23)	M (20)	M (20- 28)
Author			Busko et al. 2008 (10)				Harridge et al. 1998 (29)	Linossier et al. 1993 (47)	Tabata et al. 1990 (83)	Linossier et al. 1997 (48)	Sleivert et al. 1995 (79)

Table II. (cont.)

Author	Sex	Period	Training	Body		Mu	Muscle strength (%)	(%)			M	Muscle size (%)	ze (%)		<b>ỳO₂max</b>
	(Age)	(Age) frequency		mass	11	1RM	Isokinetic	Iso	Isometric	mCSA	V	mfCSA (VL)	V(VL)	FFM	(%)
		(sessions)							_					(DEXA)	
Others (Continuous + interval training)	ous + int	erval training	(												
Farup et al. 2012 (20)	M (23)	10 wk 3 d/wk (30)	<ol> <li>(1) 60-75% of WM 30-45 min</li> <li>(2) 70-80% of WM 2×20 min</li> <li>(3) 80-90% of WM 8×4 min</li> </ol>	su	LP	SI		KF	su su	HT	su		Su		SI
Simoneau et al. 1985 (78)	M F (19– 30)	15 wk 4–5 d/wk (60)	(1) continuous (2) 10–15×15–30 s (3) 4–5×60–90 s	1								I II A II B	21 ns (12) 24		1
M: male, F: female, m: muscle, mf: 1	le, m: mu	scle, mf: mus	muscle fiber, TBM: total body mass, VT: ventilatory threshold, VO <sub>2</sub> max: maximal oxygen uptake, HRmax: maximal heart rate, HRR:	dy mass,	VT: ve	ntilatory t	threshold, VO <sub>2</sub>	max: m	aximal ox	ygen up	take, H	Rmax: r	naximal ł	neart rate, I	HRR:

Table II. (cont.)

vastus lateralis, Ex: extensor, TH: thigh, R: right, L: left, I: type II, AerT: aerobic threshold, AnT: anaerobic threshold, FFM: fat-free mass, DEXA: dual energy X-ray heart rate reserve, WM: watt-max, KE: knee extension, KF: knee flexion, LP: leg press, SQ: squat, HE: hip extension, QF: quadriceps femoris, CSA: cross-sectional area, VL: absorptiometry, IRM: one repetition maximum, ST: slow-twitch fibers, FT: fast-twitch fibers, ns: not significant, A: about, SI: significant increase, SD: significant decrease subjects. Furthermore, we compared the mean ES for strength gain between both types of training to better determine the differences in the magnitude of strength development. As a result, the ES was 0.19 (95% CI: -0.17, 0.56; *n*: 7) for continuous training, whereas 0.13 (95% CI: -0.30, 0.56; *n*: 5) for interval training. Significant differences were not found between both types. Therefore, in young adults, the magnitude of strength gain with cycle training is low and might not necessarily differ between two types though further studies are needed to verify this because of the small number of ESs.

#### Untrained older adults

With respect to the probability of muscle hypertrophy in older adults, 4 out of 8 studies evaluating muscle size have shown thigh muscle hypertrophy at the whole muscle and/or muscle fiber level in untrained older adults. In agreement with young adults, the increase in muscle size in older subjects consistently occurred more when the total number of training sessions was high. As summarized in Table III, for continuous cycle training, all 4 studies with less than 40 training sessions showed no muscle hypertrophy, but 2 out of 3 studies with more than 40 training sessions observed muscle hypertrophy. For interval cycle training, both studies with over 40 training sessions have shown a significant increase in muscle fiber area. No studies were found that used less than 40 training sessions. The probability of muscle hypertrophy after cycle training tended to be higher for older adults compared to young adults. However, it is unlikely that this difference resulted from physiological variance. Simply, there are so many studies that used less than 40 training sessions for young subjects that the probability of muscle hypertrophy is reduced. Indeed, of the searched articles, 5 out of 8 older adult studies used the total number of training sessions equal to or more than 40. while only 7 out of 22 young subject studies satisfied this number. Meanwhile, similar to the probability of muscle hypertrophy, program period might also be the key determinant for the magnitude of change. When the ESs of both young and older adults were pooled together because of insufficient data within older adults to draw conclusions based on age, the mean ES for muscle hypertrophy in studies with equal to or more than 40 training sessions (ES: 0.69; 95% CI: 0.24, 1.13; n: 9) was significantly higher than that with less than 40 training sessions (ES: 0.18; 95% CI: -0.11, 0.47; n: 15). Also, the ES was 0.91 (95% CI: 0.34, 1.48; n: 5) in young adult studies with equal to or more than 40 training sessions, whereas 0.41 (95% CI: -0.63, 1.45; n: 4) in older adult studies with it, and a significant difference was not found between both age groups. Therefore, as long as the total number of training sessions exceeded 40, the magnitude of hypertrophy with cycle training between young and older age groups appears similar, although this needs to be verified with more studies.

Regarding the probability of strength gain in older adults, a significant increase was observed in 1RM and/or isokinetic and/or isometric strength in 7 out of 8 studies evaluating muscle strength. The probability of strength gain after cycle training is higher in older adults than in young adults. Unlike young adults, most of the studies that used continuous cycle training showed significant strength gain in older adults as summarized in Table III. This suggests that significant muscle strength gain can be achieved regardless of exercise type or intensity for untrained older adults. Furthermore, the mean ES for strength gain in older subjects (ES: 0.49; 95% CI: -0.01, 1.00; n: 4) was approximately three times higher than that in young subjects (ES: 0.16; 95% CI: -0.06, 0.39; n: 12). Therefore, it appears that strength gain favors older adults. It is unclear whether these findings suggest a physiologic difference with age or if the difference in strength may partially be explained by an attenuated baseline strength test due to an inability to maximally produce force under novel conditions such as

<b>∀0₂max</b>	(%)			9					20		24			00	00		15		23			20	
¢		(V)															7						
	FFM	(DEXA)															nr						
Muscle size (%)	mfCSA (VL)				ns (1)	ns (-3) ns (-8)						ns (–1)	ns (3) ns (17)	(/1) (11	16	su			ns (–5)				
<b>Auscle</b>	mfC			I	II a	Пb					I	ΠA	ΠB	-	I П а	3							
	mCSA						ns (4)	,						5	71								
	)m						QF							E C	Ξγ A								
	Isometric								su					30	cc							13	
(%)									KE					12	PE N							KE	
Muscle strength (%)	Isokinetic						-																
iscle st	Isol																						
Mu	1RM						11		25								18						
	Ē						SQ		KE								SQ						
Body	mass			ns (0)	~		ns (0)		ns (–1)		ns (1)			(1)	IIS (-1 )		- 1		Ś			ns (–1)	
Training				70-80% HRmax	60 min		70–90% HRmax	30-40 min	80–100% VT,	A 20–30 min <sup>2</sup>	70–80% VO,peak	30 min		000 07 AU	00-00% HKK 20-45 min		50–70% VO <sub>2</sub> max	30-45 min	70-80% HRR	30 min		50–80% VO <sub>2</sub> max	60 min
Period	frequency	(sessions)		6 wk	4 d/wk	(24)	16 wk	2 d/wk (32)	12 wk	3 d/wk (36)	12 wk	3 d/wk	(36)	-101	12 WK 3 5 d/w/b	(42)	16 wk	3 d/wk (48)	18 wk	3 d/wk	(54)	18 wk	3 d/wk (54)
Sex	(Age)		Bı	Μ	(63)		Σ	(68)	Μ	(64)	ц	(69)		F	F (71)		Μ	(75)	Μ	(68)		Μ	(64)
Author			Continuous cycling	Freyssenet et al.	1996 (22)		Izquierdo et al.	2004 (37)	Cadore et al.	2010 (11)	Ferketich et al.	1998 (21)		I Tank an at al	7009 (27)		Lovell et al.	2010 (50)	Hepple et al.	1997 (33)		Okazaki et al.	2002 (63)

Table III. The effects of cycling training on muscle size and strength as well as VO<sub>2</sub>max in untrained older adults

		NOC N	reriod	Training	Body		MUSC	le streng	Muscle strength (%)			I	Muscle s	Muscle size (%)		VO <sub>2</sub> max
M         24 wk         60% VO_max         ns (1)         LP         ns         13           F         3 dwk         15-40 min         15-40 min         15         10         11         11           (75)         (72)         14 wk         1 min (75-85% ns (-2))         KE         12         KE         13         QF         ns(2)         1           M         14 wk         1 min (75-85% ns (-2))         KE         12         KE         13         QF         ns(2)         1           M         14 wk         1 min (75-85% ns (-2))         KE         12         KE         13         QF         ns(2)         1           M         14 wk         1 min (85-95%          KE         12         KE         13         QF         ns(2)         1           M         14 wk         1 min (85-75%          N </th <th></th> <th>(Age)</th> <th>frequency (sessions)</th> <th></th> <th>mass</th> <th>IR</th> <th>7</th> <th>Isokine</th> <th></th> <th>ometric</th> <th></th> <th>CSA</th> <th>mfCS</th> <th>(JV) A(</th> <th>FFM (DEXA)</th> <th>(%)</th>		(Age)	frequency (sessions)		mass	IR	7	Isokine		ometric		CSA	mfCS	(JV) A(	FFM (DEXA)	(%)
M         14 wk         1 min (73–85%         ns (-2)         KE         12         KE         13         QF         ns (2)         1           (73)         3 dwk         H max)         (42)         4 min (80–95%         ns (-2)         H         ns (0)         11a           (73)         3 dwk         H max)         (42)         4 min (80–95%         -         -         1         -         1         -         1         -         1         -         1         -         1         -         -         -         -         -         -         1         -         1         -         1         -         1         -         1         -         1         -         1         -         -         -         -         1         1         -         1         1         -         1         -         -         -         1         1         1         -         -         -         1         1         1         1         1         -         -         1         1         1         1         1         1         1         1         1         1         1         1         1         1         1         1 <th>Strasser et al. 2009 (81)</th> <th>M F (76)</th> <th>24 wk 3 d/wk (72)</th> <th>60%</th> <th>ns (1)</th> <th>LP</th> <th>ns (11)</th> <th></th> <th></th> <th></th> <th></th> <th></th> <th></th> <th></th> <th></th> <th>ns(6)</th>	Strasser et al. 2009 (81)	M F (76)	24 wk 3 d/wk (72)	60%	ns (1)	LP	ns (11)									ns(6)
M         14 wk         1 min (75–85%         ns (-2)         KE         12         KE         13         QF         ns (2)         1           (73)         3 d/wk         HRmax)         4 min (80–95%         ns (-2)         KE         12         KE         13         QF         ns (2)         1           (73)         3 d/wk         HRmax)         10–12 min×3 sets           1          1         n         10         1         ns (0)         11 a          1         ns (0)         11 a         1         n         1         ns (0)         11 a         1         n         1         n         1         n< (0)	Interval cycling								-	-	_				-	_
(73)       3 dwk       Hkmax)         (42)       4 min (80–95%       H       ms (0)         1       Hkmax)       10–12 min × 3 sets       H       ms (0)         M       14 wk       1 min (85–95%       -       H       ms (0)         M       14 wk       1 min (85–95%       -       H       ms (0)       H         (73)       4 dwk       Hkmax)       -       -       I       I         (73)       4 dwk       Hkmax)       -       -       I       I         (73)       4 dwk       1 min (55–75%       -       -       -       I       I         (73)       4 dwk       4 ms (0)       -       -       -       -       -       -       I       I         (73)       4 dwk       4 ms (0)       -       -       -       -       -       -       -       -       -       -       -       -       I       I       I       I       -	Verney et al.	Σ	14 wk	1 min (75–85%	ns (–2)				<u> </u>		QF	ns (2)	- ;	3		10
al. M I4 wk Imix 3 sets al. M I4 wk Imix (85-95% 10-12 mix 3 sets) (73) 4 d/wk Imix (85-95% 10-12 mix 3 sets) (56) 4 mix (65-75% 10 10 10 10 10 10 10 10 10 10 10 10 10	2006, 2008 (85,   86)	(73)	3 d/wk (42)	HRmax) 4 min (80–95%							H	ns (0)	Пa	ns (8) 13		
al. M 14 wk 1 min (85–95% – 1 (73) 4 d/wk HRmax) (73) 4 d/wk HRmax) (56) 4 min (65–75% – 1 [1 a] (56) 4 min (65–75% HRmax) (56) 3 d/wk 8 sets 5 min×9 sets F 16 wk 40% of 2 RM ns F 16 wk 8 sets×16 PR ns F 16 wk 8 sets×16 PR ns F 1 4 sets×16 PR ns F 2) 80% of 2 RM ns F 3 R	(22		Ì	HRmax)										1		
al. M 14 wk 1 min (85–95% – 1 (73) 4 d/wk HRmax) (56) 4 min (65–75% HRmax) (56) 4 min (65–75% HRmax) (56) 4 min (65–75% HRmax) 5 min×9 sets (69) 3 d/wk 8 sets×16 PR ns (69) 3 d/wk 8 sets×16 PR ns (69) 3 d/wk 8 sets×16 PR ns (1) 40% of 2 RM ns (1) 40% of 2 RM ns (2) 80% of 2 RM ns (2) 80% of 2 RM ns (2) 80% of 2 RM ns				10–12 min×3 sets												
(73)       4 d/wk       HRmax)       II a         (56)       4 min (65–75%       1 min (65–75%       1 min (65–75%         (56)       4 min (65–75%       1 min (65–75%       1 min (65–75%         (50)       3 min×9 sets       5 min×9 sets       1 min (65–75%         etal.       F       16 wk       40% of 2 RM       ns         etal.       F       16 wk       8 sets×16 PR       1 ms         etal.       F       16 wk       8 sets×16 PR       1 ms         etal.       F       16 wk       8 sets×16 PR       1 ms         etal.       F       10 40% of 2 RM       ns       1 ms         etal.       F       10 40% of 2 RM       ns       1 ms         etal.       F       10 40% of 2 RM       ns       1 ms         (1)       4 sets×16 PR +       1 ms       1 ms       1 ms         (2)       80% of 2 RM       ns       1 ms       1 ms       1 ms	Charifi et al.	Μ	14 wk	1 min (85–95%									Ι			14
(56)     4 min (65–75%     1 min (65–75%       HRmax)     5 min×9 sets     1 max)       5 min×9 sets     1 max)       et al.     F     16 wk       (69)     3 d/wk     8 sets×16 PR       (69)     3 d/wk     8 sets×16 PR       8 sets×8 PR     ns       8 sets×8 PR     Si       4 sets×16 PR +     Si       (2) 80% of 2 RM     ns	2003 (13)	(73)	4 d/wk	HRmax)									Па	ns (6)		
HRmax)     HRmax)       5 min×9 sets     5 min×9 sets       et al.     F       16 wk     40% of 2 RM       ad/wk     8 sets×16 PR       (69)     3 d/wk       8 sets×16 PR       8 sets×8 PR       8 sets×6 PR       11 40% of 2 RM       ns       4 sets×16 PR +       (2) 80% of 2 RM			(56)	4 min (65–75%										23		
et al.     F     16 wk     40% of 2 RM     ns       et al.     F     16 wk     40% of 2 RM     ns       (69)     3 d/wk     8 sets×16 PR     KE       (48)     80% of 2 RM     ns     LP       8 sets×8 PR     ns     4 sets×16 PR +     13       (1)     4 sets×16 PR +     12     8				HRmax)												
et al. F 16 wk 40% of 2 RM ns KE (69) 3 d/wk 8 sets×16 PR 12 M ns 12 LP (48) 80% of 2 RM ns 8 sets×16 PR 12 LP 8 sets×8 PR 13 8 sets×8 PR 13 13 13 13 13 13 13 13 13 13 13 13 13				5 min×9 sets	_											
et al. F 16 wk 40% of 2 RM ns KE (69) 3 d/wk 8 sets×16 PR (48) 80% of 2 RM ns LP (48) 80% of 2 RM ns 8 sets×8 PR 8 sets×8 PR (1) 40% of 2 RM ns 4 sets×16 PR + (2) 80% of 2 RM ns (2) 80% of 2 RM	Other															
(69)     3 d/wk     8 sets×16 PR     LP       (48)     80% of 2 RM     ns     LP       (10)     8 sets×8 PR     ns     100% of 2 RM       (10)     40% of 2 RM     ns     100% of 2 RM       (10)     4 sets×16 PR +     100% of 2 RM	Macaluso et al.	ы	16 wk	40% of 2 RM	ns				KI							
IIS	2003 (51)	(69)	3 d/wk (48)	8 sets×16 PR												
su				80% of 2 RM	SU					SI						
IIS				8 sets×8 PR						SI						
				(1) 40% of 2 RM	SU					SI						
(2) 80% of 2 RM				4 sets×16 PR +						SI						
				(2) 80% of 2 RM												
4 sets×8 PK				4 sets×8 PK												

sectional area, VL: vastus lateralis, I: type II, FFM: fat-free mass, DEXA: dual energy X-ray absorptiometry, UL: upper leg, 1RM: one repetition maximum, ns: not

significant, SI: significant increase

Table III. (cont.)

Acta Physiologica Hungarica 102, 2015

12

#### Ozaki H et al.

Author	Sex	Subject	Period	Training	Body		Muscl	e stren	Muscle strength (%)			Mu	Muscle size (%)	(0)	<b>ỳO</b> 2peak
	(Age)		irequency (sessions)		mass	11	1RM	Isokinetic		Isometric	-	mfCSA (VL)		FFM	(%)
Continuous cycling	00						1				-				
Petersen et al.	М	hemodialysis	6 wk	50-80% VO, peak ns (0)	ns (0)			KE	ns K	KE	ns				ns (2)
2009 (67)	Н		3 d/wk	20  min											r.
	(42)		(18)												
Steiner et al.	Σ	coronary	8 wk	~60% VO,peak	1			KE	ns K	KE	ns	19	DEXA	3	1
2004 (80)	(56)	artery disease	3d/wk (24)	$30 \min_{z}$											
Pitta et al.	Μ	chronic	8 wk	80% HRpeak									imped-	- ns (0)	
2004 (69)	Ц	obstructive	3d/wk	30 min									ance		
	(64)	pulmonary	(24)												
		disease													
Belardinelli et al.	Μ	chronic heart	8 wk	40% VO,peak	Ι						I	24			17
1995 (4)	Ч	failure	3 d/wk	$30 \min$							Π				
	(56)		(24)												
Haykowsky et al.	ц	chronic heart	12 wk	60-70% HRR	1	Γb	13								12
2005 (32)	(72)	failure	2 d/wk	15-45 min											
			(24)												
Lee et al.	Μ	stroke	10–12 wk	50-70% VO,peak	Ι	LB	ns (3)								12
2008 (46)	ц		3 d/wk	$30 \min$											
	(67)		(30)												
Kiilavuori et al.	М	chronic heart	12 wk	50-60% VO.peak	1				×	KE	ns ST	F ns (10)	(0		I
2000 (42)	(52)	failure	3 d/wk	$30 \min$							FT				
			(36)												
Preisler et al.	Ι	Kennedy	12 wk	65–70% VO <sub>2</sub> max					k	KE ns	ns (1)		DEXA	ns (–1)	su
2009 (71)	(56)	disease	2-5 d/wk	$30 \min$					Ŧ	HE	ns				
			(42)							<u>.</u>	-5)				

Table IV. The effects of cycling training on muscle size and strength as well as VO2max in patients

Г

T

Author	Sex (A GP)	Subject	Period	Training	Body		Musc	Muscle strength (%)	h (%)			Muscle	Muscle size (%)		ÝO₂peak
	(250)		(sessions)			1R	1RM	Isokinetic Isometric	c Is	ometric	mfCS	mfCSA (VL)	H	FFM	(0/)
Sveen et al. 2008 (82)	M (32)	Becker muscular dystrophy	12 wk ~5 d/wk (50)	65% VO <sub>2</sub> max 30 min	ns (-1)				KE HE	KE ns (5) HE ns (2)	I	ns (39) ns (42)	DEXA	DEXA ns (-1)	SI
Interval cycling															
Bouchla et al. 2011 (7)	Ъч	chronic heart failure	12 wk 3 d/wk	~40 min (1) 30 sec: 50%		KE 2 RM	10						DEXA	ns (1)	8
~	(51)		(36)	Wpeak (2) 60 sec: rest											
Delagardelle et	M	congestive	12 wk	40 min	su			KE 3							ns (0)
al. 2002 (14)	(09)	heart failure	3-4 d/wk (40)	<ul> <li>(1) 2 min</li> <li>(50% VO<sub>2</sub>peak)</li> <li>(2) 2 min</li> <li>(75% VO<sub>2</sub>peak)</li> </ul>	(-1)			KF 11							
El Mhandi et al. 2008 (18)	M (20- 44)	Charcot-Ma- rie-Tooth	24 wk 3 d/wk (72)	6×5 min (1) 4 min (40% Pmax)	1			KE 10 KF 13	KF KF	ns (2) ns (4)					10
				(vnii i) 11111 i (7)											

VL: vastus lateralis, I: type I, II: type II, FFM: fat-free mass, DEXA: dual energy X-ray absorptiometry, 1RM: one repetition maximum, ST: slow-twitch fibers, FT: fast-twitch Pmax: maximal aerobic power, Wpeak: peak workload, KE: knee extension, KF: knee flexion, HE: hip extension, LP: leg press, LB: lower body, CSA: cross-sectional area, M: male, F: female, m: muscle, mf: muscle fiber, VO<sub>2</sub>max: maximal oxygen uptake, VO<sub>2</sub>peak: peak oxygen uptake, HRpeak: peak heart rate, HRR: heart rate reserve, fibers, ns: not significant, SI: significant increase

Table IV. (cont.)

Ozaki H et al.

exercise (e.g. fear of injury). Meanwhile, in the only study without strength gain after continuous cycle training, training intensity was constant throughout the training period, while in the other studies observing significant strength gain gradually increased exercise training intensity over the training period. This suggests that it is important that workload is adjusted to maintain sufficient mechanical stress to skeletal muscle.

#### Patients

Generally, fitness levels are lower in patients compared to healthy adults because of lower levels of daily activity, which may influence muscle adaptations to cycle training. Therefore, we will discuss the effect of cycle training on muscle size and strength for patients separately from healthy adults (Table IV). With respect to the probability of muscle hypertrophy, 2 out of 4 studies evaluating muscle size have shown muscle fiber hypertrophy of the VL after continuous cycle training in patients. In patients, muscle hypertrophy following cycle training appears to be influenced more by the initial value rather than training design variables. Both type I and type II fiber size significantly increased after 8 weeks of continuous cycle training even at an exercise intensity of 40% VO<sub>2</sub>peak (4) but did not change after 12 weeks of cycle training at 50-60% VO<sub>2</sub>peak (42) in patients with chronic heart failure. Thus, a longer training period and a higher exercise intensity do not necessarily contribute to muscle fiber hypertrophy after continuous cycle training in patients. Meanwhile, with respect to initial level of muscle size, muscle fiber area was much lower in both studies observing an increase in muscle size (approximately  $2500-4000 \ \mu m^2$ ) (4, 80) compared to the other 2 studies where no change in muscle size occurred (approximately 4000–6000  $\mu$ m<sup>2</sup>) (42, 82). Therefore, it appears that muscle fiber hypertrophy is observed especially in patients with lower muscle fiber area. Furthermore, the mean ES tended to be higher in patients (ES: 0.69: 95% CI: -0.07, 1.44; n: 7) compared to untrained healthy young (ES: 0.40; 95% CI: 0.10, 0.71; n: 18) and older (ES: 0.28; 95% CI: -0.31, 0.87; n: 4) adults, which may be also related to lower muscle size for patients compared to healthy adults. Meanwhile, to the best of our knowledge, there is no study evaluating whole muscle size measured by MRI and CT in patients before and after cycle training. Future research needs to determine whether cycle training elicits muscle hypertrophy at the whole muscle level in patients.

Regarding the probability of strength gain, only 1 out of 7 previous studies using continuous cycle training have shown increased strength in patients, whereas strength gain was observed in all 3 studies using interval training. Interval cycle training could result in a greater training stimulus to the exercising muscle because interval training allows patients to exercise for longer total periods of time at a higher exercise intensity (74). Therefore, it appears that interval cycle training may be more suitable to improve muscle strength in patients than continuous cycle training. One of the reasons that significant strength gain is frequently observed in continuous training for older adults may be the lower initial value of strength compared to young adults. Thus, even continuous cycle training is likely to induce significant strength gain for patients because they generally have lower values of strength compared to untrained healthy adults. However, continuous cycle training does not appear to be an effective method for strength gain in patients. For example, the mean ES for strength gain tended to be lower in patients (ES: 0.21; 95% CI: 0.03, 0.39; n: 6) compared to older adults (ES: 0.49; 95% CI: -0.01, 1.00; n: 4). In contrast to older adults, patients may not increase strength after continuous training for several reasons. To illustrate, in previous studies employing patients, the training period, exercise time and intensity were 6–12 weeks, 20–30 min and 50–80%  $\dot{VO}_2$ peak, respectively, which appears to be shorter or lower compared to untrained healthy subjects. Furthermore, a relative mechanical stimulus to skeletal muscle might be lower in patients than that in untrained healthy adults even if the relative exercise intensity (% $\dot{VO}_2$ peak) was the same between both adult groups because exercise intensity was frequently set using  $\dot{VO}_2$ peak measured by an incremental symptom-limited exercise test. Therefore, a lack of strength gain found after continuous cycle training for the patients may be related to a lower exercise intensity, shorter training period and shorter exercise time compared to the studies employing untrained healthy adults.

## Possible mechanisms for cycling-induced muscle hypertrophy

In this section, we discuss the possible mechanisms for continuous cycling-induced muscle hypertrophy due to the insufficient data regarding interval cycling. The following mechanisms may be common to interval cycling because cycling training can induce muscle hypertrophy regardless of exercise type as mentioned previously. However, it should be acknowledged that there may be distinct or subtle differences between continuous and interval cycle exercise.

#### Muscle activity during pedaling

Numerous studies have reported muscular activation during pedaling using EMG analysis (19, 54, 76). For example, Ericson et al. (19) have quantified the activation of thigh muscles during ergometer cycling as recorded by EMG in recreationally-active students. Peak muscular activation (normalized by EMG recorded during MVC (%MVC)), when performing cycle exercise at 120 W and 60 rpm, was the following values for each muscle: 12% in RF, 54% in VM, 50% in VL, 12% in BF and 10% in medial hamstring (SM and ST). VM and VL muscles were especially activated by cycling exercise and corresponded to ~50% MVC. Meanwhile, Marsh and Martin (54) found that the average %MVC during cycling was approximately 30% in VL at 200 W and 110 rpm in young non-cyclists. In recent years, it has been established that a very low exercise intensity such as 30% 1RM can lead to an increase in myofibrillar protein synthesis when performed repetitively or until volitional failure (9). Therefore, cycle training that consists of repetitive movements may suffice as a minimum stimulus required to increase muscle fiber activation high enough to result in favorable physiologic increases in muscle protein synthesis.

#### Muscle cell swelling

It is empirically known that a bout of high-intensity cycle exercise induces a temporary increase in thigh size similar to resistance training, which is likely due to a fluid shift from the plasma into the muscle cell. According to hypothetical model for cell swelling introduced by Haussinger (30), cell swelling may affect protein metabolism, gene expression and proteolysis through the activation of MAPK (49). These acute changes in muscle size are considered an indirect measure of muscle cell swelling although the possibility of the increase in muscle size induced by just an increase in interstitial fluid cannot be completely ruled out (49). Ploutz-Snyder et al. (70) have shown that the CSA of vasti and adductor muscle groups increased 10% and 5%, respectively, whereas plasma volume decreased immediately after squat exercise and the reduction of plasma volume was correlated with the increase in muscle CSA. We have confirmed that muscle thickness significantly increased 8% in RF and VL after a bout of 5 min pedaling exercise at 90% VO<sub>2</sub>max (unpublished data), the value of which is not largely different from that after a bout of resistance training. Therefore, muscle cell swelling may be one of factors affecting muscle protein metabolism after a bout of cycle exercise.

## Muscle protein synthesis (mTOR and MAPK signaling pathway)

Skeletal muscle hypertrophy results from a prolonged shift of muscle protein turn-over towards synthesis rather than breakdown (75). The translation of messenger ribonucleic acid (mRNA) plays a prominent role in protein synthesis following an exposure to exercise stimuli (43, 61). In translation, the mechanistic target of rapamycin (mTOR) enhances mRNA translation through the phosphorylation of eukaryotic translation initiation factor 4E binding protein 1 (4E-BP1) and ribosomal protein S6 kinase 1(S6K1) (8, 9, 16, 17), which in turn results in an increase in muscle protein synthesis. Several mitogen-activated protein kinase (MAPK) signaling pathways such as extracellular signal-regulated kinase (ERK) and p38-MAPK signaling pathways also play an important role in muscle protein synthesis (16, 53). Some studies have shown that a bout of cycle exercise activated both mTOR (55) and MAPK (87) signaling pathway as well as muscle protein synthesis (28). However, the magnitude of increase appears to be smaller for cycle exercise compared to resistance exercise. For example, S6K1 phosphorylation increased above basal values immediately after both cycle and resistance exercise (88). However, after 4 hours, S6K1 phosphorylation remained above basal values only for the resistance exercise (88). Furthermore, one study has shown that myofibrillar protein synthesis was stimulated over the 4-hour period following resistance exercise but not after cycle exercise (88). These results suggest that increased myofibrillar protein synthesis rate after cycle exercise is smaller or slower compared to resistance exercise. or that cycle exercise may not necessarily stimulate myofibrillar protein synthesis. A smaller or slower increase in myofibrillar protein synthesis may provide some explanation for why many training sessions are required for cycle training to induce muscle hypertrophy as mentioned earlier.

#### Proteolytic gene expression

In addition to protein synthesis, it is known that three proteolytic systems are involved in muscle protein degradation: ubiquitin-proteasome system (UPS), cytosolic calciumdependent calpain system and the lysosomal system. Among these systems, UPS plays a prominent role in muscle protein breakdown (3, 38) through the elevated expressions of muscle-specific ubiquitin ligases: Atrogin-1 and muscle ring finger 1 (MuRF1) (6, 26). Therefore, it appears that a reduced expression following exercise results in muscle hypertrophy through a positive change in muscle net protein balance (90). Konopka et al. (27, 44) have shown that 12-week cycle training induced muscle hypertrophy and significantly reduced mRNA expression of Forkhead transcription factor 3A (FOXO3A) at rest with a trend for its downstream targets, Atrogin-1 and MuRF1, to also be reduced. Furthermore, FOXO3A significantly decreased 6 hours after an acute bout of 60-minute cycle exercise compared to rest (28). Therefore, following a bout of cycle exercise, the reduction of proteolytic mRNA expression may be another factor contributing to muscle hypertrophy. However, to better determine the overall contribution of reduced proteolytic systems to muscle hypertrophy, more research investigating the change in muscle protein degradation rate following cycle exercise is needed.

#### Satellite cell

The proliferation of satellite cells may also contribute substantially to muscle hypertrophy (40, 68). Although muscle fibers have multiple myonuclei, the addition of new myonuclei are thought to be important for substantial long term increases in human skeletal muscle mass (31). Satellite cells are ordinarily in a quiescent state but they can increase muscle fiber area

through differentiation and ultimate fusion with a muscle fiber if they are activated to enter the cell proliferation cycle when a muscle is injured or subjected to mechanical stress (31). Charifi et al. (13) and Verney et al. (85) have investigated the effect of cycle training on muscle size and satellite cells in older men. They demonstrated that cycle training induced a 12-23% increase in type II a fiber area of VL muscle and increased the number of satellite cells per fiber. However, the number of myonuclei per fiber did not change in either study (13, 85). This is likely because new myonuclei are not required when the magnitude of muscle fiber hypertrophy does not exceed 26% (40) or a myonuclear domain of ~2000  $\mu$ m<sup>2</sup> (68). Therefore, it is likely that levels of muscle hypertrophy induced by cycle training can be observed independent of changes in myonuclei.

## Conclusion

Cycle training appears to be capable of inducing muscle hypertrophy as well as increased aerobic capacity. However, it is plausible that cycle training requires a longer training period than typical resistance training until an increase in muscle size can be observed due to a much slower hypertrophy rate.

#### Practical applications

Our research suggests that cycle training elicits muscle hypertrophy of the thigh similarly between healthy untrained young and older adults, while strength gain seems to favor older adults. Thus, this suggests that improving muscle quality may be higher in older adults than in young adults. Cycle training is a training mode to produce muscle hypertrophy and strength gain but may not be the most effective way. Cycle training appears to require a relatively longer period of time to promote significant increases in muscle size compared to traditional resistance training. Therefore, trainers and therapists need to select the most suitable training method based on the preference and purpose of exercise training of their clients. Furthermore, future research is needed to determine an optimal training design to maximize the hypertrophic effect and/or strength gain.

## Acknowledgement

There was no funding received. No potential conflicts of interest were disclosed.

#### REFERENCES

- 1. Allemeier CA, Fry AC, Johnson P, Hikida RS, Hagerman FC, Staron RS: Effects of sprint cycle training on human skeletal muscle. J. Appl. Physiol. 77, 2385–2390 (1994)
- 2. Andersen P, Henriksson J: Capillary supply of the quadriceps femoris muscle of man: adaptive response to exercise. J. Physiol. 270, 677–690 (1977)
- Baar K, Nader G, Bodine S: Resistance exercise, muscle loading/unloading and the control of muscle mass. Essays Biochem. 42, 61–74 (2006)
- Belardinelli R, Georgiou D, Scocco V, Barstow TJ, Purcaro A: Low intensity exercise training in patients with chronic heart failure. J. Am. Coll. Cardiol. 26, 975–982 (1995)
- Bell GJ, Syrotuik D, Martin TP, Burnham R, Quinney HA: Effect of concurrent strength and endurance training on skeletal muscle properties and hormone concentrations in humans. Eur. J. Appl. Physiol. 81, 418–427 (2000)

- Bodine SC, Latres E, Baumhueter S, Lai VK, Nunez L, Clarke BA, Poueymirou WT, Panaro FJ, Na E, Dharmarajan K, Pan ZQ, Valenzuela DM, DeChiara TM, Stitt TN, Yancopoulos GD, Glass DJ: Identification of ubiquitin ligases required for skeletal muscle atrophy. Science 294, 1704–1708 (2001)
- Bouchla A, Karatzanos E, Dimopoulos S, Tasoulis A, Agapitou V, Diakos N, Tseliou E, Terrovitis J, Nanas S: The addition of strength training to aerobic interval training effects on muscle strength and body composition in CHF patients. J. Cardiopulm. Rehabil. Prev. 31, 47–51 (2011)
- Burd NA, Holwerda AM, Selby KC, West DWD, Staples AW, Cain NE, Cashaback JGA, Potvin JR, Baker SK, Phillips SM: Resistance exercise volume affects myofibrillar protein synthesis and anabolic signalling molecule phosphorylation in young men. J. Physiol. 588, 3119–3130 (2010)
- Burd NA, West DWD, Staples AW, Atherton PJ, Baker JM, Moore DR, Holwerda AM, Parise G, Rennie MJ, Baker SK, Phillips SM: Low-load high volume resistance exercise stimulates muscle protein synthesis more than high-load low volume resistance exercise in young men. Plos One 5, e12033 (2010)
- Busko K, Madej A, Mastalerz A: Changes of muscle torque after sprint and endurance training performed on the cycle ergometer. Biol. Sport 25, 275–294 (2008)
- Cadore EL, Pinto RS, Lhullier FL, Correa CS, Alberton CL, Pinto SS, Almeida AP, Tartaruga MP, Silva EM, Kruel LF: Physiological effects of concurrent training in elderly men. Int. J. Sports Med. 31, 689–697 (2010)
- Carter SL, Rennie CD, Hamilton SJ, Tarnopolsky MA: Changes in skeletal muscle in males and females following endurance training. Can. J. Physiol. Pharmacol. 79, 386–392 (2001)
- Charifi N, Kadi F, Feasson L, Denis C: Effects of endurance training on satellite cell frequency in skeletal muscle of old men. Muscle Nerve 28, 87–92 (2003)
- 14. Delagardelle C, Feiereisen P, Autier P, Shita R, Krecke R, Beissel J: Strength/endurance training versus endurance training in congestive heart failure. Med. Sci. Sports Exerc. 34, 1868–1872 (2002)
- Denis C, Chatard JC, Dormois D, Linossier MT, Geyssant A, Lacour JR: Effects of endurance training on capillary supply of human skeletal muscle on two age groups (20 and 60 years). J. Physiol. (Paris) 81, 379–383 (1986)
- Drummond MJ, Dreyer HC, Fry CS, Glynn EL, Rasmussen BB: Nutritional and contractile regulation of human skeletal muscle protein synthesis and mTORC1 signaling. J. Appl. Physiol. 106, 1374–1384 (2009)
- Drummond MJ, Fry CS, Glynn EL, Dreyer HC, Dhanani S, Timmerman KL, Volpi E, Rasmussen BB: Rapamycin administration in humans blocks the contraction-induced increase in skeletal muscle protein synthesis. J. Physiol. 587, 1535–1546 (2009)
- El Mhandi L, Millet GY, Calmels P, Richard A, Oullion R, Gautheron V, Feasson L: Benefits of interval-training on fatigue and functional capacities in Charcot-Marie-Tooth disease. Muscle Nerve 37, 601–610 (2008)
- Ericson MO, Nisell R, Arborelius UP, Ekholm J: Muscular activity during ergometer cycling. Scand. J. Rehabil. Med. 17, 53–61 (1985)
- Farup J, Kjolhede T, Sorensen H, Dalgas U, Moller AB, Vestergaard PF, Ringgaard S, Bojsen-Moller J, Vissing K: Muscle morphological and strength adaptations to endurance vs. resistance training. J. Strength Cond. Res. 26, 398–407 (2012)
- Ferketich AK, Kirby TE, Alway SE: Cardiovascular and muscular adaptations to combined endurance and strength training in elderly women. Acta Physiol. Scand. 164, 259–267 (1998)
- Freyssenet D, Berthon P, Denis C, Barthelemy JC, Guezennec CY, Chatard JC: Effect of a 6-week endurance training programme and branched-chain amino acid supplementation on histomorphometric characteristics of aged human muscle. Arch. Physiol. Biochem. 104, 157–162 (1996)
- 23. Garber CE, Blissmer B, Deschenes MR, Franklin BA, Lamonte MJ, Lee IM, Nieman DC, Swain DP: American College of Sports Medicine position stand. Quantity and quality of exercise for developing and maintaining cardiorespiratory, musculoskeletal, and neuromotor fitness in apparently healthy adults: guidance for prescribing exercise. Med. Sci. Sports Exerc. 43, 1334–1359 (2011)
- Gollnick PD, Armstrong RB, Saltin B, Saubert CW 4th, Sembrowich WL, Shepherd RE: Effect of training on enzyme activity and fiber composition of human skeletal muscle. J. Appl. Physiol. 34, 107–111 (1973)
- Gollnick PD, Armstrong RB, Saubert CW 4th, Piehl K, Saltin B: Enzyme activity and fiber composition in skeletal muscle of untrained and trained men. J. Appl. Physiol. 33, 312–319 (1972)
- Gomes MD, Lecker SH, Jagoe RT, Navon A, Goldberg AL: Atrogin-1, a muscle-specific F-box protein highly expressed during muscle atrophy. Proc. Natl. Acad. Sci. U. S. A. 98, 14440–14445 (2001)
- Harber MP, Konopka AR, Douglass MD, Minchev K, Kaminsky LA, Trappe TA, Trappe S: Aerobic exercise training improves whole muscle and single myofiber size and function in older women. Am. J. Physiol. Regul. Integr. Comp. Physiol. 297, R1452–R1459 (2009)

- Harber MP, Konopka AR, Jemiolo B, Trappe SW, Trappe TA, Reidy PT: Muscle protein synthesis and gene expression during recovery from aerobic exercise in the fasted and fed states. Am. J. Physiol. Regul. Integr. Comp. Physiol. 299, R1254–R1262 (2010)
- Harridge SDR, Bottinelli R, Canepari M, Pellegrino M, Reggiani C, Esbjornsson M, Balsom PD, Saltin B: Sprint training, in vitro and in vivo muscle function, and myosin heavy chain expression. J. Appl. Physiol. 84, 442–449 (1998)
- 30. Haussinger D: The role of cellular hydration in the regulation of cell function. Biochem. J. 313, 697-710 (1996)
- Hawke TJ, Garry DJ: Myogenic satellite cells: physiology to molecular biology. J. Appl. Physiol. 91, 534–551 (2001)
- Haykowsky M, Vonder Muhll I, Ezekowitz J, Armstrong P: Supervised exercise training improves aerobic capacity and muscle strength in older women with heart failure. Can. J. Cardiol. 21, 1277–1280 (2005)
- 33. Hepple RT, Mackinnon SL, Goodman JM, Thomas SG, Plyley MJ: Resistance and aerobic training in older men: effects on VO<sub>2</sub>peak and the capillary supply to skeletal muscle. J. Appl. Physiol. 82, 1305–1310 (1997)
- 34. Hoppeler H, Howald H, Conley K, Lindstedt SL, Claassen H, Vock P, Weibel ER: Endurance training in humans: aerobic capacity and structure of skeletal muscle. J. Appl. Physiol. 59, 320–327 (1985)
- 35. Howald H, Hoppeler H, Claassen H, Mathieu O, Straub R: Influences of endurance training on the ultrastructural composition of the different muscle fiber types in humans. Pflügers Arch. 403, 369–376 (1985)
- Hug F, Marqueste T, Le Fur Y, Cozzone PJ, Grelot L, Bendahan D: Selective training-induced thigh muscles hypertrophy in professional road cyclists. Eur. J. Appl. Physiol. 97, 591–597 (2006)
- Izquierdo M, Ibanez J, Hakkinen K, Kraemer WJ, Larrion JL, Gorostiaga EM: Once weekly combined resistance and cardiovascular training in healthy older men. Med. Sci. Sports Exerc. 36, 435–443 (2004)
- Jackman RW, Kandarian SC: The molecular basis of skeletal muscle atrophy. Am. J. Physiol. Cell Physiol. 287, C834–C843 (2004)
- Jacobs I, Esbjornsson M, Sylven C, Holm I, Jansson E: Sprint training effects on muscle myoglobin, enzymes, fiber types, and blood lactate. Med. Sci. Sports Exerc. 19, 368–374 (1987)
- Kadi F, Schjerling P, Andersen LL, Charifi N, Madsen JL, Christensen LR, Andersen JL: The effects of heavy resistance training and detraining on satellite cells in human skeletal muscles. J. Physiol. 558, 1005–1012 (2004)
- Kaljumae U, Hanninen O, Airaksinen O: Knee extensor fatigability and strength after bicycle ergometer training. Arch. Phys. Med. Rehabil. 75, 564–567 (1994)
- 42. Kiilavuori K, Naveri H, Salmi T, Harkonen M: The effect of physical training on skeletal muscle in patients with chronic heart failure. Eur. J. Heart Fail. 2, 53–63 (2000)
- Kimball SR, Jefferson LS: Control of translation initiation through integration of signals generated by hormones, nutrients, and exercise. J. Biol. Chem. 285, 29027–29032 (2010)
- 44. Konopka AR, Douglass MD, Kaminsky LA, Jemiolo B, Trappe TA, Trappe S, Harber MP: Molecular adaptations to aerobic exercise training in skeletal muscle of older women. J. Gerontol. A Biol. Sci. Med. Sci. 65, 1201–1207 (2010)
- LaStayo PC, Pierotti DJ, Pifer J, Hoppeler H, Lindstedt SL: Eccentric ergometry: increases in locomotor muscle size and strength at low training intensities. Am. J. Physiol. Regul. Integr. Comp. Physiol. 278, R1282–R1288 (2000)
- 46. Lee MJ, Kilbreath SL, Singh MF, Zeman B, Lord SR, Raymond J, Davis GM: Comparison of effect of aerobic cycle training and progressive resistance training on walking ability after stroke: A randomized sham exercise-controlled study. J. Am. Geriatr. Soc. 56, 976–985 (2008)
- 47. Linossier MT, Denis C, Dormois D, Geyssant A, Lacour JR: Ergometric and metabolic adaptation to a 5-s sprint training program. Eur. J. Appl. Physiol. Occup. Physiol. 67, 408–414 (1993)
- Linossier MT, Dormois D, Geyssant A, Denis C: Performance and fibre characteristics of human skeletal muscle during short sprint training and detraining on a cycle ergometer. Eur. J. Appl. Physiol. Occup. Physiol. 75, 491–498 (1997)
- Loenneke JP, Fahs CA, Thiebaud RS, Rossow LM, Abe T, Ye X, Kim D, Bemben MG: The acute muscle swelling effects of blood flow restriction. Acta Physiol. Hung. 99, 400–410 (2012)
- 50. Lovell DI, Cuneo R, Gass GC: Can aerobic training improve muscle strength and power in older men? J. Aging Phys. Act. 18, 14–26 (2010)
- Macaluso A, Young A, Gibb KS, Rowe DA, De Vito G: Cycling as a novel approach to resistance training increases muscle strength, power, and selected functional abilities in healthy older women. J. Appl. Physiol. 95, 2544–2553 (2003)

- Mackova E, Melichna J, Havlickova L, Placheta Z, Blahova D, Semiginovsky B: Skeletal muscle characteristics of sprint cyclists and nonathletes. Int. J. Sports Med. 7, 295–297 (1986)
- Mahoney SJ, Dempsey JM, Blenis J: Cell signaling in protein synthesis ribosome biogenesis and translation initiation and elongation. Prog. Mol. Biol. Transl. Sci. 90, 53–107 (2009)
- Marsh AP, Martin PE: The relationship between cadence and lower extremity EMG in cyclists and noncyclists. Med. Sci. Sports Exerc. 27, 217–225 (1995)
- Mascher H, Andersson H, Nilsson PA, Ekblom B, Blomstrand E: Changes in signalling pathways regulating protein synthesis in human muscle in the recovery period after endurance exercise. Acta Physiol. (Oxf) 191, 67–75 (2007)
- McCarthy JP, Agre JC, Graf BK, Pozniak MA, Vailas AC: Compatibility of adaptive responses with combining strength and endurance training. Med. Sci. Sports Exerc. 27, 429–436 (1995)
- McCarthy JP, Pozniak MA, Agre JC: Neuromuscular adaptations to concurrent strength and endurance training. Med. Sci. Sports Exerc. 34, 511–519 (2002)
- Mikkola J, Rusko H, Izquierdo M, Gorostiaga EM, Hakkinen K: Neuromuscular and cardiovascular adaptations during concurrent strength and endurance training in untrained Men. Int. J. Sports Med. 33, 702–710 (2012)
- 59. Millet GY, Lepers R: Alterations of neuromuscular function after prolonged running, cycling and skiing exercises. Sports Med. 34, 105–116 (2004)
- Mitchell CJ, Churchward-Venne TA, West DWD, Burd NA, Breen L, Baker SK, Phillips SM: Resistance exercise load does not determine training-mediated hypertrophic gains in young men. J. Appl. Physiol. 113, 71–77 (2012)
- Ishii N, Ogasawara R, Kobayashi K, Nakazato K: Roles played by protein metabolism and myogenic progenitor cells in exercise-induced muscle hypertrophy and their relation to resistance training regimens. Jpn. J. Phys. Fitness Sports Med. 1, 83–94 (2012)
- Nelson AG, Arnall DA, Loy SF, Silvester LJ, Conlee RK: Consequences of combining strength and endurance training regimens. Phys. Ther. 70, 287–294 (1990)
- Okazaki K, Kamijo YI, Takeno Y, Okumoto T, Masuki S, Nose H: Effects of exercise training on thermoregulatory responses and blood volume in older men. J. Appl. Physiol. 93, 1630–1637 (2002)
- Ozaki H, Loenneke JP, Thiebaud R, Abe T: Resistance training induced increase in VO<sub>2</sub>max in young and older subjects. Eur. Rev. Aging Phys. Act. 10, 107–116 (2013)
- Ozaki H, Loenneke JP, Thiebaud RS, Stager JM, Abe T: Possibility of leg muscle hypertrophy by ambulation in older adults: a brief review. Clin. Interv. Aging 8, 369–375 (2013)
- Parcell AC, Woolstenhulme MT, Sawyer RD: Structural protein alterations to resistance and endurance cycling exercise training. J. Strength Cond. Res. 23, 359–365 (2009)
- Petersen AC, Leikis MJ, McMahon LP, Kent AB, McKenna MJ: Effects of endurance training on extrarenal potassium regulation and exercise performance in patients on haemodialysis. Nephrol. Dial. Transplant. 24, 2882–2888 (2009)
- Petrella JK, Kim JS, Cross JM, Kosek DJ, Bamman MM: Efficacy of myonuclear addition may explain differential myofiber growth among resistance-trained young and older men and women. Am. J. Physiol. Endocrinol. Metab. 291, E937–E946 (2006)
- 69. Pitta F, Brunetto AF, Padovani CR, Godoy I: Effects of isolated cycle ergometer training on patients with moderate-to-severe chronic obstructive pulmonary disease. Respiration 71, 477–483 (2004)
- Ploutz-Snyder LL, Convertino VA, Dudley GA: Resistance exercise-induced fluid shifts: change in active muscle size and plasma volume. Am. J. Physiol. 269, R536–R543 (1995)
- Preisler N, Andersen G, Thogersen F, Crone C, Jeppesen TD, Wibrand F, Vissing J: Effect of aerobic training in patients with spinal and bulbar muscular atrophy (Kennedy disease). Neurology 72, 317–323 (2009)
- Putman CT, Xu X, Gillies E, MacLean IM, Bell GJ: Effects of strength, endurance and combined training on myosin heavy chain content and fibre-type distribution in humans. Eur. J. Appl. Physiol. 92, 376–384 (2004)
- Rhea MR: Determining the magnitude of treatment effects in strength training research through the use of the effect size. J. Strength Cond. Res. 18, 918–920 (2004)
- 74. Sabapathy S, Kingsley RA, Schneider DA, Adams L, Morris NR: Continuous and intermittent exercise responses in individuals with chronic obstructive pulmonary disease. Thorax 59, 1026–1031 (2004)
- 75. Sandri M: Signaling in muscle atrophy and hypertrophy. Physiology (Bethesda) 23, 160-170 (2008)
- Sarre G, Lepers R, Maffiuletti N, Millet G, Martin A: Influence of cycling cadence on neuromuscular activity of the knee extensors in humans. Eur. J. Appl. Physiol. 88, 476–479 (2003)
- Short KR, Vittone JL, Bigelow ML, Proctor DN, Nair KS: Age and aerobic exercise training effects on whole body and muscle protein metabolism. Am. J. Physiol. Endocrinol. Metab. 286, E92–E101 (2004)

#### Ozaki H et al.

- Simoneau JA, Lortie G, Boulay MR, Marcotte M, Thibault MC, Bouchard C: Human skeletal muscle fiber type alteration with high-intensity intermittent training. Eur. J. Appl. Physiol. Occup. Physiol. 54, 250–253 (1985)
- Sleivert GG, Backus RD, Wenger HA: The influence of a strength sprint training sequence on multijoint power output. Med. Sci. Sports Exerc. 27, 1655–1665 (1995)
- Steiner R, Meyer K, Lippuner K, Schmid JP, Saner H, Hoppeler H: Eccentric endurance training in subjects with coronary artery disease: a novel exercise paradigm in cardiac rehabilitation? Eur. J. Appl. Physiol. 91, 572–578 (2004)
- Strasser B, Keinrad M, Haber P, Schobersberger W: Efficacy of systematic endurance and resistance training on muscle strength and endurance performance in elderly adults – a randomized controlled trial. Wien. Klin. Wochenschr. 121, 757–764 (2009)
- Sveen ML, Jeppesen TD, Hauerslev S, Kober L, Krag TO, Vissing J: Endurance training improves fitness and strength in patients with Becker muscular dystrophy. Brain 131, 2824–2831 (2008)
- Tabata I, Atomi Y, Kanehisa H, Miyashita M: Effect of high-intensity endurance training on isokinetic muscle power. Eur. J. Appl. Physiol. Occup. Physiol. 60, 254–258 (1990)
- Tarnopolsky MA, Rennie CD, Robertshaw HA, Fedak-Tarnopolsky SN, Devries MC, Hamadeh MJ: Influence of endurance exercise training and sex on intramyocellular lipid and mitochondrial ultrastructure, substrate use, and mitochondrial enzyme activity. Am. J. Physiol. Regul. Integr. Comp. Physiol. 292, R1271–R1278 (2007)
- Verney J, Kadi F, Charifi N, Feasson L, Saafi MA, Castells J, Piehl-Aulin K, Denis C: Effects of combined lower body endurance and upper body resistance training on the satellite cell pool in elderly subjects. Muscle Nerve 38, 1147–1154 (2008)
- Verney J, Kadi F, Saafi MA, Piehl-Aulin K, Denis C: Combined lower body endurance and upper body resistance training improves performance and health parameters in healthy active elderly. Eur. J. Appl. Physiol. 97, 288– 297 (2006)
- Widegren U, Wretman C, Lionikas A, Hedin G, Henriksson J: Influence of exercise intensity on ERK/MAP kinase signalling in human skeletal muscle. Pflügers Arch. 441, 317–322 (2000)
- Wilkinson SB, Phillips SM, Atherton PJ, Patel R, Yarasheski KE, Tarnopolsky MA, Rennie MJ: Differential effects of resistance and endurance exercise in the fed state on signalling molecule phosphorylation and protein synthesis in human muscle. J. Physiol. 586, 3701–3717 (2008)
- Wilson JM, Marin PJ, Rhea MR, Wilson SM, Loenneke JP, Anderson JC: Concurrent training: a meta-analysis examining interference of aerobic and resistance exercises. J. Strength Cond. Res. 26, 2293–2307 (2012)
- 90. Zanchi NE, de Siqueira MV, Lira FS, Rosa JC, Yamashita AS, Carvalho CRD, Seelaender M, Lancha AH: Chronic resistance training decreases MuRF-1 and Atrogin-1 gene expression but does not modify Akt, GSK-3 beta and p70S6K levels in rats. Eur. J. Appl. Physiol. 106, 415–423 (2009)