

Original

ARTÍCULO EN INGLÉS

## Effect of whole body vibration (WBV) on PTH in elderly subjects

G. Martín<sup>a</sup>, Y. de Saa<sup>a</sup>, M. E. Da Silva-Grigoletto<sup>b</sup>, D. Vaamonde<sup>c</sup>, S. Sarmiento<sup>a</sup> and J. M. García-Manso<sup>a</sup>

<sup>a</sup>Department of Physical Education. University of Las Palmas de Gran Canaria. Las Palmas de Gran Canaria. Spain.

<sup>b</sup>Andalusian Center of Sports Medicine. Córdoba. Spain.

<sup>c</sup>Morphological Sciences Department. School of Medicine. Universidad de Córdoba. Spain.

### History of the article:

Received June 22, 2008.

Accepted November 13, 2008.

### Key words:

Vibration training

Hormones.

Elderly.

### Palabras clave:

Entrenamiento con vibraciones mecánicas.

Hormonas.

Ancianos.

### Correspondence:

J. M. García-Manso.

Departamento de Educación Física.

Facultad de Ciencias de la Actividad Física

y el Deporte.

Universidad de Las Palmas de Gran Canaria.

35017 Islas Canarias. España.

E-mail: jgarcia@def.ulpgc.es

### ABSTRACT

**Purpose.** The present study aims to analyze, in elderly women, the response of parathormone (PTH) to mechanical vibration stimuli and to assess its effect on basal calcium and phosphate concentrations after a low- and very low- intensity short-term training.

**Methods.** Sixteen elderly females (mean age  $69.64 \pm 4$  years) participated from the study. Both experimental (EG;  $n = 8$ ) and control group (CG;  $n = 8$ ) were exposed to whole body vibration on a vibration platform for 24 sessions (three times per week). On the platform, both groups (EG, CG) performed three exercises (two of them were dynamic and one was static). The vibration frequency was progressively increased in GE (20-32 Hz) while CG always received the same frequency (10 Hz). Outcome measures were blood concentrations of PTH, calcium, phosphate and  $\beta$ -cross lap. Likewise, other complementary tests used were anthropometry, and the 30-second chair stand test.

**Results.** PTH blood concentration increased significantly ( $p < 0.05$ ) in EG by 44.3%; whereas the responses of blood calcium, phosphate, and  $\beta$ -cross lap showed no significant increase ( $p > 0.05$ ). The 30-second chair stand test showed significant changes ( $p < 0.05$ ) in strength levels for both groups, with changes being more marked for EG. Anthropometric tests also showed training was more useful for EG.

**Conclusions.** The results from the present study suggest that low-intensity whole body vibration training leads to positive hormonal profile on PTH, which can benefit the osseous construction processes on elderly people.

© 2009 Revista Andaluza de Medicina del Deporte.

### RESUMEN

#### Efecto del entrenamiento por estimulación neuromuscular mecánica (WBV) sobre la PTH en personas mayores

**Objetivo.** El presente estudio pretende analizar, en mujeres de edad avanzada, la respuesta de la parathormona (PTH) al estímulo de vibración mecánica y evaluar su efecto sobre las concentraciones basales de calcio y fosfato tras un corto periodo de entrenamiento de baja y muy baja intensidad.

**Método.** Dieciséis mujeres de edad avanzada ( $69,64 \pm 4$  años) participaron en este estudio. El grupo experimental (GE;  $n = 8$ ) y el grupo control (GC;  $n = 8$ ) realizaron tres ejercicios, dos estáticos y uno dinámico sobre una plataforma de vibración durante 24 sesiones (3 días/semana). La frecuencia de vibración se incrementó progresivamente en GE (20 Hz-32 Hz) mientras que en GC se mantuvo constante a 10 Hz. Los parámetros bioquímicos determinados pre- y post- entrenamiento fueron la PTH, calcio, fosfato y  $\beta$ -cross lap. Otras pruebas complementarias usadas fueron la antropometría y el test de sentarse y levantarse.

**Resultados.** La concentración de PTH tras el periodo de entrenamiento aumentó significativamente ( $p < 0,05$ ) en GE en un 44,3%. Tras el entrenamiento no se observaron cambios significativos ( $p > 0,05$ ) en las concentraciones de calcio, fosfato y  $\beta$ -cross lap en ninguno de los grupos. La prueba de sentarse y levantarse mostró cambios significativos ( $p < 0,05$ ) en los niveles de fuerza para ambos grupos, siendo más notables para el GE. Las pruebas antropométricas también demostraron que el entrenamiento fue más eficaz en el GE.

**Conclusión.** Los resultados de este estudio sugieren que el entrenamiento con vibraciones mecánicas (WBV) de baja intensidad promueve un perfil hormonal positivo sobre la PTH, el cual puede beneficiar los procesos de construcción ósea en personas mayores.

© 2009 Revista Andaluza de Medicina del Deporte.

## Introduction

It is estimated that, by mid-21st century, one in five people will be above 65 and, thus, it will be the state's responsibility to find formulas to improve its citizens' quality of life. Physical exercise has frequently been presented as a useful means to induce bone mass gaining<sup>1-3</sup> and, therefore, to prevent osteoporosis<sup>4-6</sup>. Osteoporosis and osteopenia are two of the greatest health problems that modern society faces, especially in a moment in which population is significantly increasing its life expectancy.

The continuous remodeling processes (formation and resorption) of bone structures are affected by many factors, among which, age, diet, metabolic disorders, hormonal imbalances, lactation, menopause or exercise are included as the most relevant<sup>7-11</sup>.

Tensions resulting from mechanical loads, as generated by physical exercise, are an important regulator of remodeling in some parts of the skeleton. The introduction of new technologies, applied to physical condition improvement in sportsmen, has led to the development of new methodologies that, years later, have been transferred to other population sectors (sedentary, sick people or special populations). One of the most recent and popular innovations is the use of neuromuscular activation by means of vibrating mechanical stimuli<sup>12,13</sup>.

Whole body stimulation (Whole Body Vibration [WBV]) is, from all the different applications this technology (vibration systems applied to traditional weights machines, upper limb vibration, lower limb vibration or whole body vibration) allows, the one that has reached greatest popularity in recent years and the one generating the greatest body of knowledge. This technology generates controlled vibrations that, originating on the person's foot sole, are transmitted to the caudal body area activating, in the neuromuscular system, what is known as Tonic Vibration Reflex (TVR)<sup>14-19</sup>. TVR's response intensity depends on four factors: localization of the vibration application (on muscle or tendon), muscle's initial length (the more stretched, the greater the response will be), CNS's excitability state and/or vibrating stimulus parameters<sup>20</sup>.

The impact produced by the stimulus on the musculature depends on factors such as vibration frequency (Hz), displacement amplitude (mm), movement magnitude ( $\text{m}\cdot\text{s}^{-1}$ ), direction (linear and/or rotational), exposure duration (min or sec), number of exposures, recovery time between each exposure (sec or min) and number of sessions<sup>21</sup>.

Recently, it has been proposed that this type of activity may result in a convenient and simple alternative to be used by a population suffering from problems in bone structure mineralization or that present with risk of fracture; in fact, WBV use is especially interesting when subjects have part of their motor autonomy compromised<sup>22-27</sup>. Some of the functional responses that would be affected by vibration would be related to calcium metabolism and to the hormonal response that regulates it. Parathormone (PTH), 1,25 dihydroxy-vitamin D (1,25(OH)<sub>2</sub>D<sub>3</sub>) and calcitonin are some of the compounds related to the osseous formation and resorption processes, offering us an idea on how bone construction mechanisms behave during the application of these exercises<sup>25</sup>.

PTH contributes to several physiological processes such as: maintaining calcium concentration in extracellular fluids within physiological limits<sup>28</sup>, controlling intracellular  $\text{Ca}^{2+}$  homeostasis<sup>29</sup>, regulating 1,25 (OH)<sub>2</sub>D<sub>3</sub> formation by the liver<sup>30</sup> or, by means of this other hormone, incrementing  $\text{Ca}^{2+}$  resorption and decrease phosphate excretion. Such facts make this hormone a key factor in the control of osseous construction. The process entails that when extracellular  $\text{Ca}^{2+}$  levels decrease (< 8 mg/dl) PTH release by the parathyroid cells is increased<sup>31</sup>.

While in the long run the hormone's function is to favor osseous re-sorption mobilizing both calcium and phosphate, when there are high levels of calcemia (> 9.5 to 10 mg/dl), PTH's function might be the opposite favoring bone construction<sup>32-35</sup>. The anabolic mechanism is linked to osteoblast-produced plasminogen<sup>36</sup>, with PTH reducing the activity of this protein's inhibitor and favoring plasmin production, which, in turn, favors the synthesis of local growth factors (*Insulin like Growth Factor* [IGF-II] and possibly *Transforming Growth Factor- $\beta$*  [TGF $\beta$ ]). Such growth factors favor the release and proliferation of osteoblasts, responsible of initiating osseous production.

There is a relatively abundant literature on the acute response of PTH to physical exercise<sup>37-44</sup>, though none of the studies was performed using elderly subjects. From the studies, it is derived that both intensity and duration of exercise are determining factors to achieve an intense response of PTH to exercise, postulating the existence of a load threshold from which the aforementioned response is provoked<sup>43,45</sup>. On the contrary, the studies analyzing the short- and long-term responses of PTH to training loads are less numerous; however, in such case, the population was composed of elderly people. Among these, Zerath et al<sup>46</sup> find a statistically significant increase in PTH levels after training without observing any changes in osteocalcin.

We hypothesize that a short term WBV training provokes an endocrine response in elderly subjects that results favorable to the osseous construction processes. Thus, the objective of the present study was to analyze, in elderly women, the response of PTH to mechanical vibration stimuli and to assess its effect on basal calcium and phosphate concentrations after a low- and very low- intensity short-term training.

## Methods

### Subjects

Sixteen elderly females (age:  $69.64 \pm 4$  years; height:  $156.24 \pm 4.42$  cm; body mass index:  $28.29 \pm 1.97$   $\text{kg}\cdot\text{m}^{-2}$ ) participated in the study. The inclusion criteria were related to: age, health condition, and physical condition; it was decided to choose people with a medium level of physical activity, which was determined by means of a typical simplified questionnaire. In order to be included, subjects could not be taking any medication that could affect calcium metabolism, or hormonal reposition. Therefore, exclusion criteria were related to not meeting any of the above-mentioned inclusion criteria and/or having missed more than one training sessions. None of the subjects had any previous experience in vibration platforms training. Subjects were asked to maintain their regular diet during the whole study period; this feature was controlled by means of alimentary records. All the participants gave written consent according to the Helsinki Declaration guidelines for research on humans. The sample was randomly divided into two groups: Experimental Group (EG),  $n = 8$ ; and Control Group (CG),  $n = 8$  (fig. 1). Sample size was estimated as described by Cohen<sup>47</sup> for ANOVA by using the PTH variable and data from pilot studies.

### Materials

A vibration platform (NEMES, Ergotest, Rome, Italy) allowing work frequencies of 10-50 Hz, and vertical displacements of 4 mm was used. For the anthropometric assessment and for height, a skinfold calliper, and a stadiometer were used, respectively (Holtain Ltd., Dyfed, UK); for peri-

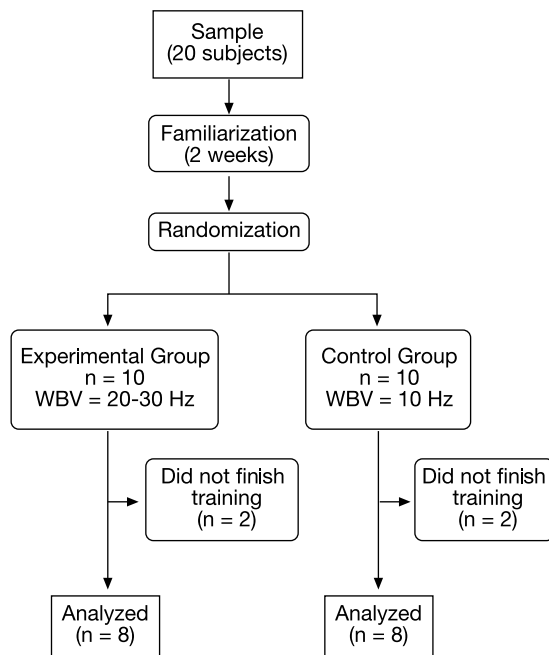


Fig. 1. Flow diagram of participants. WBV: whole body vibration.

meter measurements and weight measurements, a metallic metric tape (Holtain Ltd., Dyfed, UK), and a commercial digital scale were employed, respectively.

#### Anthropometric measurements and 30-second chair stand test

Anthropometric measurements were performed according to ISAAC's instructions<sup>48</sup>, assessing weight, height, body mass index, skinfolds (triceps, subscapular, suprailliac, abdominal, upper-thigh, and mid-calf), and thigh and calf perimeters (to estimate muscle area).

In order to estimate lower limb muscular strength, the 30-sec chair stand test was performed as described by Rikli and Jones<sup>49</sup>; briefly, the test started with the subject sitting down with the back resting on the chair and feet on the floor. When the researcher indicated, the subject would stand up and go back to the initial position. The subject was encouraged to complete the highest possible number of repetitions in a 30 second-period. Before commencing the test, the researcher showed how to perform the exercise correctly, with the subjects practicing it one to three times in order to become familiar with the task, and starting the test immediately after.

#### Procedure

All subjects participated from the familiarization period during which they learnt to correctly perform the exercises without the use of the vibration platform. In a second phase, the exercises were repeated on a vibration platform using a 10 Hz vibration.

For the experimental phase, subjects underwent 24 training sessions (3 days per week<sup>-1</sup>). Each session consisted of performing three exercises according to the protocol described in table 1:

- 1) Static standing position.
- 2) Dynamic squatting exercises.
- 3) Static squatting position.

Briefly, the technical characteristics of the exercises were the following: a) static standing position: the hands should grip the platform handle in a relaxed way. The head had to be kept in the anatomical position, the legs in a 180° knee extension, and feet had to be resting on the metatarsus with a slight heel elevation; b) dynamic squatting exercise: with the hands on the handle, a flexion of the knee joint was performed, with a range of motion from 120 to 180°. The execution speed, four seconds for each repetition, was marked by means of a digital metronome; c) static squatting position: with the hands on the handle, a sustained flexion of the knee joint is performed at approximately 160-170°. For all exercises and sessions, the used work/recovery ratio was 1/1.

Previous to each session, a warm-up phase, consisting of articular mobility exercises and muscular groups' activation exercises, was performed. At the end of the session, a series of static stretch exercises for plantar flexor, knee extensor and hip flexor muscles was performed.

#### Clinical assessment

Before and after the training protocol, all subjects underwent a clinical analysis test (blood and urine) to determine the following biochemical parameters: PTH (Immulite 2000 intact PTH) by solid-phase; enzyme-labelled, chemiluminescent sequential immunometric assay; calcium (Roche/Hitachi Modular) by color test with final point and white sample determination; inorganic phosphate (Roche/Hitachi Phosphorus) and  $\beta$ -crosslap ( $\beta$ -CTX) by electrochemiluminescence immunoassay (ECLIA, Roche Eclesys automated analyzers/E170 module). The analysis were performed at Teror Health Center after overnight fasting, both one day before familiarization and two days after finishing the training program sessions.

#### Statistical analysis

A basic descriptive statistics (mean and standard deviation) and a comparative statistics ANOVA (2 x 2) with a 95% confidence interval were

Table 1

Whole body vibration (WBV) training sessions used for the experimental group (EG; n = 8) and the control group (CG; n = 8)

Session (number)	EG frequency (Hz)	CG frequency (Hz)	Exposure time (sec)	Recovery time* (sec)
1	20	10	30	75
2	20	10	30	75
3	20	10	30	75
4	22	10	30	75
5	22	10	30	75
6	22	10	30	75
7	22	10	35	75
8	22	10	35	75
9	22	10	35	75
10	22	10	40	90
11	22	10	40	90
12	22	10	40	90
13	24	10	45	90
14	24	10	45	90
15	24	10	45	90
16	27	10	50	90
17	27	10	50	90
18	27	10	50	90
19	30	10	55	90
20	30	10	55	90
21	30	10	55	90
22	32	10	60	90
23	32	10	60	90
24	32	10	60	90

\*: recovery between exercises.

performed. In addition, the effect size has been calculated as described by Cohen<sup>47</sup>. The statistical package SPSS 12.0 was used for all statistical tests.

## Results

The results of blood PTH, calcium, phosphate, and  $\beta$ -crosslap concentrations are shown in table 2. There were no changes observed for the values of calcium, phosphate, and  $\beta$ -crosslap at the end of the training program for any of the groups. We should highlight that the measured calcium corresponds to total blood calcium, not ionized calcium. On the contrary, PTH shows a significant improve for EG ( $p < 0.05$ ). None of the subjects commented on having experienced any negative side effects.

By means of the complementary pre-post tests applied for the assessment of physical condition and anthropometry (table 3), it is confirmed that, after training, there were significant changes ( $p < 0.05$ ) in knee extensor musculature strength levels for both groups, with changes be-

**Table 2**

Values of parathormone (PTH), calcium, phosphate, and  $\beta$ -crosslap for experimental group (EG;  $n = 8$ ) and control group (CG;  $n = 8$ ) before and after the 8-week WBV training are given as mean  $\pm$  standard deviation

Group	Mean ( $\pm$ SD)		Effect Size	Reference
	Pre	Post		
PTH				
EG	36.51 ( $\pm$ 10.25)	52.68 ( $\pm$ 15.54)*.#	1.58	0-68.2 pg/ml
CG	39.02 ( $\pm$ 11.13)	45.79 ( $\pm$ 12.11)	0.61	
Calcium				
EG	9.77 ( $\pm$ 0.39)	9.72 ( $\pm$ 0.38)	0.13	8.2-10.5 mg/dl
CG	9.66 ( $\pm$ 0.30)	9.61 ( $\pm$ 0.33)	0.17	
Phosphate				
EG	4.04 ( $\pm$ 0.38)	3.87 ( $\pm$ 0.31)	0.45	2.7-4.5 mg/dl
CG	3.72 ( $\pm$ 0.34)	3.89 ( $\pm$ 0.26)	0.50	
$\beta$ -crosslap				
EG	0.31 ( $\pm$ 0.15)	0.39 ( $\pm$ 0.14)	0.53	0.104-1.008 ng/dl
CG	0.36 ( $\pm$ 0.14)	0.38 ( $\pm$ 0.17)	0.14	

\*: Significant difference ( $p < 0.05$ ) between pre and post values; #: significant difference ( $p < 0.05$ ) between EG and CG.

**Table 3**

Morphofunctional parameters for experimental group (EG;  $n = 8$ ) and control group (CG;  $n = 8$ ) before and after the 8-week whole body vibration training

Group	Mean ( $\pm$ SD)		Effect size
	Pre	Post	
30-sec chair stand test			
EG	9.89 ( $\pm$ 2.17)	13.13 ( $\pm$ 1.96)**	1.49
CG	10.86 ( $\pm$ 1.81)	12.57 ( $\pm$ 1.50)*	0.94
Calf muscle area			
EG	72.43 ( $\pm$ 13.61)	84.68 ( $\pm$ 12.50)*.#	0.90
CG	69.73 ( $\pm$ 11.81)	71.97 ( $\pm$ 8.20)	0.19
Thigh muscle area			
EG	174.48 ( $\pm$ 15.44)	189.01 ( $\pm$ 14.23)*	0.94
CG	172.54 ( $\pm$ 17.21)	181.07 ( $\pm$ 15.82)	0.27
$\Sigma$ skinfolds			
EG	127.89 ( $\pm$ 19.78)	100.87 ( $\pm$ 18.44)*.#	1.37
CG	151.47 ( $\pm$ 17.77)	132.27 ( $\pm$ 14.60)	1.08

Values given as mean  $\pm$  standard deviation (SD).

\*: significant difference ( $p < 0.05$ ) between pre and post values; \*\*: significant difference ( $p < 0.001$ ) between pre and post values; #: significant difference ( $p < 0.05$ ) between EG and CG.

$\Sigma$  skinfolds: triceps, subscapular, suprailiac, abdominal, upper-thigh, and mid-calf. For this variable, only 11 subjects were analyzed (EG;  $n = 6$  and CG;  $n = 5$ ) since, in five subjects, the abdominal skinfold measurements did not pass the reproducibility testing.

ing more marked for EG in comparison to CG (see effect size). In addition, EG also showed a significant increase in calf and thigh muscle area (16.9%,  $p < 0.05$ , and 8.3%,  $p < 0.05$ ; respectively); conversely, CG showed only small increments that did not reach statistical significance. The mean skinfold sum showed a statistical significant decrease in EG albeit no statistically significant changes were observed in body weight in any of the analyzed groups.

## Discussion

The obtained results indicate that, after finishing the WBV training program, PTH concentration significantly increased in EG. Thus, it seems there is a relationship between the PTH production increase and the vibrating stimuli from the WBV platforms.

The lesser-intensity (very low) vibration stimulus (frequency 10 Hz) used in CG also provoked an increase in PTH although it was not statistically significant. The increases observed were not accompanied by any changes in calcium or phosphate values in neither one of the groups (EG and CG). Normally, calcium concentration in extracellular fluids is regulated in a very precise manner, and very rarely varies in a small percentage from its reference values (8.2-10.5 mg  $\cdot$  dl<sup>-1</sup>), with daily variations that are not above 5%. Minimal changes in this parameter are rapidly detected and influence PTH. In our case, calcium levels of the analyzed subjects were high, as expected in subjects of such age range, staying elevated until the end of the study (table 2). Phosphate, which normal blood concentration oscillates between 3 and 4.5 mg  $\cdot$  dl<sup>-1</sup> in adults, showed a tendency towards a slight decrease in EG (4.2%; effect size: 0.45) although values were always within normal range. Such tendency was not observed in CG, which even showed increasing values.

The regulation of PTH secretion is basically related to blood calcium concentration. However, during physical exercise, other factors, in addition to calcium, may modify PTH secretion; such factors are catecholamines, and acidosis, which are influenced by the training load. While PTH secretion is related to blood calcium concentration, several studies show that the variations in PTH response during exercise are independent from calcium concentration<sup>39,50,51</sup>. The adrenergic system is activated during exercise<sup>52</sup>, and it has been proved that this system influences the regulation of PTH secretion<sup>53</sup>. Elevated values of blood PTH concentration upon finishing training seem to be more related to this hormone's anabolic function<sup>39</sup>. Although the complex mechanisms by which PTH may promote osseous mass gain are not clearly understood<sup>34</sup>, there are several studies showing that an intermittent infusion of PTH increases bone formation more than it increases bone resorption<sup>54,55</sup>, leading to an increased bone mass<sup>54,56-58</sup>.

In the present study, serum PTH values increased in both groups (EG: 44.3%; CG: 17.3%) although the increase only reached statistical relevance in EG. However, it should be noted that the increase in PTH levels does not surpass the normality range estimated for this population (0-68.2 mg  $\cdot$  dl<sup>-1</sup>). In our study, we cannot accurately conclude that the increases in PTH levels might result in an osseous remodeling. However, the described behaviour and the moderate decrease in phosphate levels could make us think it might be happening indeed. We could infer that there are no significant changes in pro-collagen I levels since  $\beta$ -crosslap, a resorption marker, which indicates that the degraded type I collagen fragments appear as a consequence of the osteoclasts' action during the degradation of collagen in bones<sup>59</sup> did not experience any significant modification in its values (table 2). Since no changes were observed in

this marker, this could indicate that, while osseous degradation has not been increased, other construction processes have done it indeed.

It is worth mentioning, due to the possible important repercussion of the finding, that most of the participants subjectively referred to having experienced an increase in strength in lower limbs. Complementary pre-post tests applied for assessing physical condition and anthropometry, showed that: a) in spite of both CG and EG significantly improving lower limb muscle strength after the training period, such increase was greater in EG than in CG; by the effect size calculation (1.49 vs 0.94, respectively), we could postulate that, when increasing sample size, the difference among groups could reach statistical significance; b) thigh and calf muscle area were significantly higher in EG as compared to CG after training; c) the skinfold sum showed that EG experienced a greater decrease in fat mass than CG after training. Both the increase in muscle strength and in muscle area are substantially important for improving the quality of life in the elderly, in aspects such as a greater autonomy, and, possibly, less sarcopenia<sup>60</sup>.

In conclusion, it seems that low-intensity training programs using the vibration platform, in elderly subjects, generate an endocrine response on PTH which results favorable to osseous construction processes. However, the development of more exhaustive studies, assessing a bigger sample, are needed in order to advance the knowledge on the effect the vibration stimulus produces on osseous remodeling markers.

## Acknowledgements

We would like to thank all the participants in the present study for having volunteered and adhered to the whole study protocol. Likewise, we would like to thank Dr. Prof. Clodoaldo de Sá for his numerous and valuable suggestions and contributions during the correction and editing process of the manuscript.

## References

- Maïmoun L, Lumbroso S, Manetta J, Paris F, Leroux JL, Sultan C. Testosterone is significantly reduced in endurance athletes without impact on bone mineral density. *Horm Res*. 2003;59:285-92.
- Eliakim A, Beyth Y. Exercise training, menstrual irregularities and bone development in children and adolescents. *J Pediatr Adolesc Gynecol*. 2003;16:201-6.
- Vainionpää A, Korpelainen R, Vihriala E, Rinta-Paavola A, Leppaluoto J, Jamsa T. Intensity of exercise is associated with bone density change in premenopausal women. *Osteoporos Int*. 2006;17:455-63.
- Dalsky GP. Exercise: its effect on bone mineral content. *Clin Obstet Gynecol*. 1987;30:820-32.
- Frost HM. Vital biomechanics: proposed general concepts for skeletal adaptations to mechanical usage. *Calcif Tissue Int*. 1988;42:145-56.
- Snow-Harter C, Marcus R. Exercise, bone mineral density, and osteoporosis. *Exerc Sport Sci Rev*. 1991;10:351-88.
- Robinson CJ, Hall J, Beshir SO. Hormonal modulation of mineral metabolism in reproduction. *Proc Nutr Soc*. 1983;42:169-80.
- Garel JM. Hormonal control of calcium metabolism during the reproductive cycle in mammals. *Physiol Rev*. 1987;67:1-66.
- O'Flaherty EJ. Physiologically based models for bone-seeking elements. I. Rat skeletal and bone growth. *Toxicol Appl Pharmacol*. 1991;111:299-312.
- O'Flaherty EJ. Physiologically based models for bone-seeking elements. II. Kinetics of lead disposition in rats. *Toxicol Appl Pharmacol*. 1991;111:313-31.
- O'Flaherty EJ. Physiologically based models for bone-seeking elements. III. Human skeletal and bone growth. *Toxicol Appl Pharmacol*. 1991;111(2):332-41.
- Da Silva-Grigoletto ME, Vaamonde D, Padullés JM. Efectos del entrenamiento con vibraciones mecánicas sobre la "performance" neuromuscular. *Apunts Educación Física y Deportes*. 2006;84:39-46.
- Da Silva-Grigoletto ME, Vaamonde D, Padullés JM. Entrenamiento con vibraciones mecánicas y salud: efectos sobre los sistemas óseo, endocrino y cardiovascular. *Apunts Educación Física y Deportes*. 2006;84:47-57.
- Eklund G, Hagbarth KE. Motor effects of vibratory muscle stimuli in man. *Electroencephalogr Clin Neurophysiol*. 1965;19:619.
- Eklund G, Hagbarth KE. Normal variability of tonic vibration reflexes in man. *Exp Neurol*. 1966;16:80-92.
- Johnston RM, Bishop B, Coffey GH. Mechanical vibration of skeletal muscles. *Phys Ther*. 1970;50:499-505.
- De Gail P, Lance WP, Neilson PD. Differential effects on tonic and phasic reflex mechanics produced by vibration of muscles in man. *J Neurol Neurosurg Psychiatry*. 1966;29:1-11.
- Hagbarth KE. EMG studies of stretch reflex in man. *Electroencephalogr Clin Neurophysiol Suppl*. 1967;25:74-9.
- Marsden CE, Meadows JC, Hodgson HJF. Observations of the reflex response to muscle vibration in man and its voluntary control. *Brain*. 1969;92:829-46.
- Tous J, Moras G. Entrenamiento por medio de vibraciones mecánicas: revisión de la literatura. *Lecturas de Educación Física y Deporte. Revista Digital EF Deportes*. 2004;79 [disponible en: <http://www.efdeportes.com/efd79/vibrac.htm>].
- Luo J, McNamara B, Moran K. The use of vibration training to enhance muscle strength and power. *Sports Med*. 2005;35(1):23-41.
- Flieger J, Karachalios T, Khaldi L, Raptou P, Lyritis G. Mechanical stimulation in the form of vibration prevents postmenopausal bone loss in ovariectomized rats. *Calcif Tissue Int*. 1998;63(6):510-4.
- Russo CR, Lauretani F, Bandinelli S, Bartali B, Cavazzini C, Guralnik JM, et al. High-frequency vibration training increases muscle power in postmenopausal women. *Arch Phys Med Rehabil*. 2003;84:1854-7.
- Verschueren SM, Roelants M, Delecluse C, Swinnen S, Vanderschueren D, Boonen S. Effect of 6-month whole body vibration training on hip density, muscle strength, and postural control in postmenopausal women: a randomized controlled pilot study. *J Bone Miner Res*. 2004;19(3):352-9.
- Rubin C, Recker R, Cullen D, Ryaby J, McCabe J, McLeod K. Prevention of postmenopausal bone loss by a low-magnitude, high-frequency mechanical stimuli: a clinical trial assessing compliance, efficacy, and safety. *J Bone Miner Res*. 2004;19(3):343-51.
- Kawanabe K, Kawashima A, Sashimoto I, Takeda T, Sato Y, Iwamoto J. Effect of whole-body vibration exercise and muscle strengthening, balance, and walking exercises on walking ability in the elderly. *Keio J Med*. 2007;56(1):28-33.
- Gusí N, Raimundo A, Leal A. Low-frequency vibratory exercise reduces the risk of bone fracture more than walking: a randomized controlled trial. *BMC Musculoskelet Disord*. 2006;7:92.
- Lane NE, Sánchez S, Modin GW, Genant HK, Pierini E, Arnaud CD. Parathyroid hormone treatment can reverse corticosteroid-induced osteoporosis. Results of a randomized controlled clinical trial. *J Clin Invest*. 1998;102:1627-33.
- Rasmussen H, Pechet M, Fast D. Effect of dibutyl cyclic adenosine 3', 5'-monophosphate, theophylline, and other nucleotides upon calcium and phosphate metabolism. *J Clin Invest*. 1968;47(8):1843-50.
- Poole KE, Reeve J. Parathyroid hormone - a bone anabolic and catabolic agent. *Curr Opin Pharmacol*. 2005;5(6):612-27.
- Prieto S. Control del metabolismo del calcio, fósforo y magnesio. En: Tresgüeres JAF, editor. *Fisiología Humana*. 2ª ed. Madrid: McGraw-Hill-Interamericana; 1999. p. 979-93.
- Finkelstein JS, Klubanski A, Schaefer EH, Hornstein MD, Schiff I, Neer RM. Parathyroid hormone for the prevention of bone loss induced by estrogen deficiency. *N Engl J Med*. 1994;331(24):1618-24.
- Oxlund H, Ejersted C, Andreassen TT, Tørring O, Nilsson MH. Parathyroid hormone (1-34) and (1-84) stimulate cortical bone formation both from periosteum and endosteum. *Calcif Tissue Int*. 1993;53(6):394-9.
- Hock JM. Anabolic actions of PTH in the skeletons of animals. *J Musculoskelet Neuronal Interact*. 2001;2:1-15.
- Rubin MR, Cosman F, Lindsay R, Bilezikian JP. The anabolic effects of parathyroid hormone. *Osteoporos Int*. 2002;13:267-77.
- Li JY, Specker BL, Ho ML, Tsang RC. Bone mineral content in black and white children 1 to 6 years of age: early appearance of race and sex differences. *Am J Dis Child*. 1989;143:1346-49.
- Ljunghall S, Joborn H, Roxin LE, Rastad J, Wide L, Akerstrom G. Prolonged low-intensity exercise raises the serum parathyroid hormone levels. *Clin Endocrinol*. 1986;25:535-42.
- Grimston S, Willows N, Hanley D. Mechanical loading regime and its relationship to bone mineral density in children. *Med Sci Sport Exerc*. 1993;25(11):1203-10.
- Salvesen H, Johansson AG, Foxdal P, Wide L, Pihl-Aulin K, Ljunghall S. Intact serum parathyroid hormone levels increase during running exercise in well-trained men. *Calcif Tissue Int*. 1994;54:256-61.
- Lu KC, Shieh SD, Li BL, Chu P, Jan SY, Lin YF. Rapid correction of metabolic acidosis in chronic renal failure: effect on parathyroid hormone activity. *Nephron*. 1994;67:419-24.
- Brahm H, Pihl-Aulin K, Ljunghall S. Bone metabolism during exercise and recovery: the influence of plasma volume and physical fitness. *Calcif Tissue Int*. 1997;61:192-8.

42. Movilli E, Zani R, Carli O, Sangalli L, Pola A, Camerini C, et al. Direct effect of the correction of acidosis on plasma parathyroid hormone concentrations, calcium and phosphate in hemodialysis patients: a prospective study. *Nephron*. 2001;87:257-62.
43. Maimoun L, Manetta J, Couret I, Dupuy AM, Mariano-Goulart D, Micallef JP, et al. The intensity level of physical exercise and the bone metabolism response. *Int J Sports Med*. 2006;27:105-11.
44. Thorsen K, Kristoffersson A, Hultdin J, Lorentzon R. Effects of moderate endurance exercise on calcium, parathyroid hormone, and markers of bone metabolism in young women. *Calcif Tissue Int*. 1997;60:16-20.
45. Bouassida A, Zalleg D, Zaouali-Ajina M, Gharbi N, Duclos M, Richalet JP, et al. Parathyroid hormone concentrations during and after two periods of high intensity exercise with and without an intervening recovery period. *Eur J Appl Physiol*. 2003;88:339-44.
46. Zerath E, Holy X, Douce P, Guezennec CY, Chatard JC. Effect of endurance training on postexercise parathyroid hormone levels in elderly men. *Med Sci Sports Exerc*. 1997;29(9):1139-45.
47. Cohen J. *Statistical power analysis for the behavioral sciences*. 2nd ed. Hillsdale, NJ: L. Erlbaum Associates; 1998.
48. Norton K, Olds T. *Anthropometricia*. Sydney: University of New South Wales Press; 1996.
49. Rikli R, Jones J. Development and validation of a functional fitness test for community-residing older adults. *J Aging Phys Act*. 1999;7:129-61.
50. Henderson SA, Graham HK, Mollan RAB, Riddoch C, Scheridan B, Johnston H. Calcium homeostasis and exercise. *Int Orthop*. 1989;13:69-73.
51. Rong H, Berg U, Torring O, Sundberg CJ, Granberg B, Bucht E. Effect of acute endurance and strength exercise on circulating calcium-regulating hormones and bone markers in young healthy males. *Scand J Med Sci Sports*. 1997;7(3):152-9.
52. Sagnol M, Claustre J, Cottet-Emard JM, Pequignot JM, Fellman N, Coudert J, et al. Plasma free and sulphated catecholamines after ultra-long exercise and recovery. *Eur J Appl Physiol Occup Physiol*. 1990;60:91-7.
53. Joborn H, Hjemdahl P, Larsson PT, Lithell H, Olsson G, Wide L, et al. Effects of prolonged adrenaline infusion and of mental stress on plasma minerals and parathyroid hormone. *Clin Physiol*. 1990;10:37-53.
54. Hock JM, Gera I. Effects of continuous and intermittent administration and inhibition of resorption on the anabolic response of bone to parathyroid hormone. *J Bone Miner Res*. 1992;7:65-72.
55. Dempster DW, Cosman F, Parisien M, Shen V, Lindsay R. Anabolic actions of parathyroid hormone on bone. *Endocr Rev*. 1993;14:690-709.
56. Reeve J, Bradbeer JN, Arlot M, Davies UM, Green JR, Hampton L, et al. hPTH 1-34 treatment of osteoporosis with added hormone replacement therapy: biochemical, kinetic and histological responses. *Osteoporos Int*. 1991;1:162-70.
57. Slovik DM, Rosenthal DI, Doppelt SH, Potts JT Jr, Daly MA, Campbell JA, et al. Restoration of spinal bone in osteoporotic men by treatment with human parathyroid hormone (1-34) and 1,25-dihydroxyvitamin D. *J Bone Miner Res*. 1986;1:377-381.
58. Neer R, Slovik D, Daly M, Lo C, Potts J, Nussbaum S. Treatment of postmenopausal osteoporosis with daily parathyroid hormone plus calcitriol. In: Christiansen C, Overgaard K, editors. *Osteoporosis 1990: proceedings of the Third International Symposium on Osteoporosis*. Vol. 3. Copenhagen, Denmark: Osteopress ApS; 1990. p. 1314-7.
59. Woitge HW, Pecherstorfer M, Li Y, Keck AV, Horn E, Ziegler R, et al. Novel serum markers of bone resorption: clinical assessment and comparison with established urinary indices. *J Bone Miner Res*. 1999;14(5):792-801.
60. Asikainen TM, Kukkonen-Harjula K, Miilunpalo S. Exercise for health for early postmenopausal women: a systematic review of randomised controlled trials. *Sports Med*. 2004;34(11):753-78.