An overview on the correlation between blood zinc, zinc intake, zinc supplementation and bone mineral density in humans

Una visión general sobre la correlación entre el zinc en la sangre, la ingesta de zinc, la suplementación de zinc y la densidad mineral ósea en los seres humanos

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ABSTRACT. Introduction: In case of zinc (Zn) deficiency, this mineral becomes a nutrient limiting muscle and bone synthesis. The study in humans on zinc and bone health are few and no reviews have been published on this topic. So, the aim of this narrative review was to consider the state of the art on the correlation between blood zinc, daily zinc intake, zinc supplementation and bone mineral density.

Material and methods: A narrative review was performed.

Results: This review included 16 eligible studies: eight studies concern Zn blood; three studies concern Zn intake and five studies concern Zn supplementation.

Conclusion: Blood zinc levels seem to be lower in subjects with pathology related to bone metabolism. Regarding daily zinc intake, a high proportion of the population, more than 20%, seems to be at risk of having inadequate zinc intake. The literature suggests that an insufficient zinc intake (less than 3 mg/day) could be a risk factor for fractures and for development of osteopenia and osteoporosis. Zinc supplementation (40-50 g/day) could have beneficial effects on bone health in terms of maintaining bone mineral density and faster healing in the event of fractures, with even better results in situations of reduced intake zinc through food.

RESUMEN. Introducción: En caso de deficiencia de zinc, este mineral se convierte en un nutriente limitante de la síntesis muscular y ósea. Los estudios en humanos sobre zinc y salud ósea son pocos y no se han publicado comentarios sobre este tema. Por lo tanto, el objetivo de esta revisión narrativa es considerar el estado de la técnica sobre la correlación entre el zinc en la sangre, la ingesta diaria de zinc, la suplementación de zinc y la densidad mineral ósea. Material y métodos: Se realizó una revisión narrativa.

Resultados: Esta revisión incluyó 16 estudios elegibles: ocho se refieren al zinc en sangre; tres estudios se refieren a la ingesta diaria de zinc; y cinco estudios se refieren a la suplementación de zinc.

Conclusión: Los niveles de zinc en sangre parecen ser más bajos en sujetos con patología relacionada con el metabolismo óseo. En cuanto a la ingesta diaria de zinc, una alta proporción de la población, más del 20%, parece estar en riesgo de tener una ingesta insuficiente. La literatura sugiere que una ingesta insuficiente de zinc (menos de 3 mg/día) podría ser un factor de riesgo para fracturas y para el desarrollo de osteopenia y osteoporosis. La suplementación con zinc (40-50 g/día) podría tener efectos beneficiosos sobre la salud ósea, mantenimiento de la densidad mineral ósea y curación más rápida en caso de fracturas.
Introduction

Zinc is an essential component for our body. Over 85 percent of body zinc total is found in skeletal muscles and bones, while zinc contained in plasma represents only 0.1 percent of the total and its concentration, strictly regulated, varies from about 10 to 15 μmol/l. Zinc plasma concentrations are maintained without significant changes even when zinc intake has decreased or increased, unless these changes in intake are severe and prolonged.2

It is widely distributed in food, but the best food sources are meat, eggs, fish, cheeses and cereals.3

The RDA for adult men is 8 mg per day while for adult women it is 11 mg per day.4 In vitro it has been shown that the proliferation of osteoblastic cells has been stimulated after zinc culture with an inhibitory effect on the formation of osteoclastic cells.11,12,13,14,15,16 The same anabolic effect is confirmed in animal studies.18,19 as well as the role of osteoblastic stimulation and osteoclastic inhibition is confirmed.20,21,22 Also in animals it has been shown that zinc deficiency seems to interfere with bone metabolism with consequent reduction of bone formation23 and causes criticalities in bone consolidation in the spine;24 in another study it is highlighted how zinc deficiency can lead to a reduction in serum calcium concentration and to an increase in parathyroid hormone with subsequent bone fragility.25

Despite this background, the study in humans on zinc and bone health are few and no reviews have been published on this topic. So, the aim of this narrative review was to consider the state of the art on the correlation between blood zinc, daily zinc intake, zinc supplementation and bone mineral density.

Material and methods

The present narrative review was performed following the steps by Egger et al.26 as follows:

1. Configuration of a working group: three operators skilled in clinical nutrition (one acting as a methodological operator and two participating as clinical operators).
2. Formulation of the revision question on the basis of considerations made in the abstract: «the state of the art on the correlation between human blood zinc concentrations, daily zinc intake with food, zinc supplementation and bone mineral density».
3. Identification of relevant studies: a research strategy was planned on PubMed (Public Medline run by the National Center of Biotechnology Information [NCBI] of the National Library of Medicine of Bethesda [USA]) as follows: (a) Definition of the keywords (zinc, bone, dietary supplementation, bone mineral density), allowing the definition of the interest field of the documents to be searched, grouped in quotation marks.
and used separately or in combination; (b) use of: the Boolean (a data type with only two possible values: true or false) AND operator, that allows the establishments of logical relations among concepts; (c) Research modalities: advanced search; (d) Limits: time limits: papers published in the last 30 years; humans; adults; languages: English; (e) Manual search performed by the senior researchers experienced in clinical nutrition through the revision of articles on the state of the art on the correlation between human blood zinc concentrations, daily zinc intake with food, zinc supplementation and bone mineral density.

4. Published in journals qualified in the Index Medicus.

5. Analysis and presentation of the outcomes: we create paragraphs about the state of the art on the correlation between human blood zinc concentrations, daily zinc intake with food, zinc supplementation and bone mineral density, and the data extrapolated from the «revised studies» were collocated in tables; in particular, for each study we specified the author and year of publication and study characteristics.

6. The analysis was carried out in the form of a narrative review of the reports. At the beginning of each section, the keywords considered and the type of studies chosen are reported. We evaluated, as is suitable for the narrative review, studies of any design which considered the the state of the art on the correlation between human blood zinc concentrations, daily zinc intake with food, zinc supplementation and bone mineral density. Figure 1 shows the flow chart of literature research.

Results

Blood zinc concentrations in relation to bone metabolism

This research was conducted based on the keywords: «zinc» AND «zinc blood concentrations» and «bone» and «humans». For the present review we have analyzed a total of eight studies: five cross-sectional studies, two case-control studies and one clinical trial.

The results of these eight studies have been shown in Table 1.

Data from three studies, from 1983 to 2007, agree on a reduction in zinc values in osteoporotic women compared to healthy controls. In 1983 Atik took into consideration 22 women aged between 48 and 86 and found significantly lower blood zinc values in women with osteoporosis compared to healthy controls.27

In a study from the early 2000s, the same two groups of postmenopausal women (70 with osteoporosis and 30 healthy) were compared to the same conclusion as the previous study.28

In 2007 Mutlu et al. added osteopenic subjects to the comparison, for a total of 120 menopausal women divided into three groups (osteoporotic, osteopenic, healthy), concluding that the blood zinc levels in women with osteoporosis were significantly lower than osteopenic and healthy women, and which in turn women with osteopenia had significantly decreased blood zinc levels compared to healthy controls.29 In the 2013 study by Okyay, zinc

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**Figure 1:** Flow chart of literature research.

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Table 1: Studies cover blood zinc levels.

<table>
<thead>
<tr>
<th>First author, year</th>
<th>Study design</th>
<th>Setting</th>
<th>Inclusion criteria</th>
<th>Exclusion criteria</th>
<th>Number of subjects (M-F)</th>
<th>Micronutrient serum concentration osteoporosis</th>
<th>Micronutrient serum concentration osteopenia</th>
<th>Micronutrient serum concentration normal</th>
<th>Micronutrient serum reference value</th>
<th>Primary outcomes</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gur A, 2002</td>
<td>Clinical trial</td>
<td>Department of Physical Therapy and Rehabilitation of Dicle University Hospital</td>
<td>Women were eligible for our study if they were 50 years of age or older and in good general health as determined by medical history and routine clinical blood analysis (complete blood counts and differential count)</td>
<td>Women were excluded if they (1) had used any drug or had any disease or condition known to affect bone or calcium metabolism; (2) had taken corticosteroid medications during the previous 6 months; (3) had a history of chronic renal, hepatic, or gastrointestinal disease or lumbar compression fracture; or (4) had evidence of collapsed or focal vertebral sclerosis</td>
<td>100 postmenopausal women: 70 osteoporotic and 30 non-osteoporotic</td>
<td>Zinc serum level : 0.61 ± 0.425 (test pre supplementation of calcitonin)</td>
<td>Zinc serum level : 1.22 ± 0.31 (test pre supplementation of calcitonin)</td>
<td></td>
<td></td>
<td>To determine whether the mineral profile was different between 70 osteoporotic and 30 non-osteoporotic postmenopausal women and to evaluate the efficacy of calcitonin therapy for 6 months on these trace minerals in postmenopausal osteoporotic women</td>
<td>Zn levels in the serum of patients with postmenopausal osteoporosis were lower than those in the control group</td>
</tr>
<tr>
<td>Okyay E, 2013</td>
<td>Cross-sectional study</td>
<td>Division of Reproductive Endocrinology and Infertility, Department of Obstetrics and Gynecology at Dokuz Eylul University School of Medicine, Izmir, Turkey</td>
<td>Postmenopausal women between age 45 and 80 were included in the study</td>
<td>Women with a history of drug abuse or alcohol consumption (to drink at least ≥ 2 days per week z), and highly intake of caffeine and coffee (≥ 2 cups per day), laboratory tests or radiography of any bone metabolism disorder were excluded. Any other disease or drugs that effect bone metabolism were excluded</td>
<td>728 postmenopausal women</td>
<td>Women at 45-59 years: (p value &lt; 0.05) - L1-L4 OP</td>
<td>Zinc (μg/ml): 82.6 ± 21.7</td>
<td></td>
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<td></td>
<td>To determine the relationship between serum main minerals and postmenopausal osteoporosis.</td>
</tr>
<tr>
<td>Mutlu M, 2007</td>
<td>Cross-sectional study</td>
<td>Orthopaedics Department of the Erciyes University Medical Faculty</td>
<td>Women postmenopausal if they were &gt; 55 years of age and there had been no menstruation for ≥ 6 months prior to entry into the study.</td>
<td></td>
<td>120 postmenopausal women</td>
<td>Zinc (mg/l) 0.47 ± 0.1</td>
<td></td>
<td>Zinc (mg/l) 0.63 ± 0.09</td>
<td></td>
<td></td>
<td>To investigate the changes of Zinc in osteoporotic, osteopenic and normal postmenopausal women</td>
</tr>
</tbody>
</table>
### Continue Table 1: Studies cover blood zinc levels.

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<tr>
<td>Arikan DC, 2011</td>
<td>Case-control study</td>
<td>Department of Gynecology and Obstetrics of the Medical Faculty at Kahramanmaras Sutcu Imam (Kahramanmaras, Turkey)</td>
<td>Women 50-55 years of age were classified as postmenopausal if their plasma follicle stimulating hormone (FSH) level was &gt; 50 IU/l and their plasma estradiol concentration was &lt; 100 pmol/l</td>
<td>Surgical menopause and secondary osteoporosis or other medical conditions that might affect bone metabolism or trace element status such as kidney disease, diabetes mellitus or drug use (e.g. diuretics). Patients who were treated with bisphosphonates, calcitriol, anabolic steroids, hormone replacement therapy, calcium or vitamin D up to six months before the investigation were also excluded</td>
<td>107</td>
<td>Zn (μg/dl) 106.25 ± 36.45</td>
<td>Zn (μg/dl) 116.48 ± 35.46</td>
<td>Zn (μg/dl) 127.53 ± 45.04</td>
<td>Zn (μg/dl) 106.25 ± 36.45</td>
<td>To investigate serum zinc (Zn) levels in postmenopausal women with osteoporosis, osteopenia and in healthy controls, and to determine the relationship between Zn and BMD</td>
<td>Plasma Zn levels were higher in the healthy group when compared to the osteopenic and osteoporotic groups but the difference was not statistically significant (p &gt; 0.05)</td>
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<tr>
<td>Mahdavi-Roshan, 2015</td>
<td>Cross-sectional study</td>
<td>Rheumatology clinic in Tabriz, Islamic Republic of Iran</td>
<td>women &gt; 50 years old who had been no menstruation for ≥ 6 months prior to entry into the study, having no history of hormone replacement therapy, other bone disease, kidney stones, endocrine disorders or any medical conditions that could influence on the mineral status</td>
<td>use of mineral supplements, having history of hormone replacement therapy, bone disease, kidney stones, endocrine disorders or any medical conditions that could influence on the mineral status</td>
<td>A total of 51 post-menopausal women</td>
<td>Serum Zinc (μg/dl) 70.44 ± 4.5</td>
<td>Serum Zinc (μg/dl) 63.3 ± 4.8</td>
<td>Serum Zinc (μg/dl) 70.44 ± 4.5</td>
<td>Serum Zinc (μg/dl) 63.3 ± 4.8</td>
<td>Serum Zinc (μg/dl) 70.44 ± 4.5</td>
<td>Investigate and compare the mineral status between osteopenic and osteoporotic postmenopausal women in Tabriz, Islamic Republic of Iran.</td>
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<th>Micronutrient serum reference value</th>
<th>Primary outcomes</th>
<th>Results</th>
</tr>
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<tbody>
<tr>
<td>Atik OS, 1983</td>
<td>Case-control study</td>
<td>Department of Orthopedic Surgery of Hacettepe University Hospital.</td>
<td>Osteoporosis for cases and non osteoporosis for controls</td>
<td>22 patients (with senile Osteoporosis and controls)</td>
<td>Zinc in Serum (μg/dl) 53.5 ± 2.8</td>
<td>Zinc in Serum (μg/dl) 75.9 ± 4.1</td>
<td>to determine the zinc ion levels in serum and bone tissue of patients with senile osteoporosis.</td>
<td>zinc levels in serum of the patients with senile osteoporosis were lower than those of the control group</td>
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<tr>
<td>Relea P, 1995</td>
<td>Cross-sectional study</td>
<td>Clinic of the Rheumatology Unit of the University Hospital “Príncipe de Asturias”, Madrid</td>
<td>No pharmacological treatment, haven’t any condition that might affect calcium metabolism, such as liver disease, diabetes or renal dysfunction</td>
<td>60 postmenopausal women (30 controls and 30 with osteoporosis)</td>
<td>Zinc serum: (mg/dl) 72.7 ± 9.9 [Urinary zinc (μg/g Cr): 5.5 ± 1.9]</td>
<td>Zinc serum: (mg/dl) 74.9 ± 18.4 [Urinary zinc (μg/g Cr): 4.0 ± 2.0]</td>
<td>To evaluate the correlation between the concentrations of plasmatic zinc and urinary zinc with bone mass</td>
<td>Plasma zinc levels did not differ between the women with postmenopausal osteoporosis and the healthy postmenopausal controls, but urinary zinc excretion was higher (p=0.002) in the woman with postmenopausal osteoporosis</td>
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<tr>
<td>LIU SZ, 2009</td>
<td>Cross-sectional study</td>
<td>Xi’an urban area</td>
<td>45 to 65-year-old females of Chinese Han Nationality who had lived in the Xi’an urban area more than 30 years, and had been in natural menopause for more than half a year, with no diseases which might influence bone metabolism, no other severe chronic diseases which needed long-term therapy, no gynaecological diseases which could influence the secretion of female sex hormones, and no hormone drugs intake and osteoporosis treatment six months before investigation</td>
<td>290 women</td>
<td>Zn serum (mg/l): 0.9168 ± 0.2557</td>
<td>Zn serum (mg/l): 0.9181 ± 0.3177</td>
<td>Zn serum (mg/l): 0.9345 ± 0.2726</td>
<td>to investigate the correlation between serum macroelement and trace element contents and bone mineral density (BMD) as well as the occurrence of osteoporosis</td>
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<td>There exist significant correlations between the contents of serum elements such as calcium, phosphonium, sodium, potassium, magnesium, zinc, iron, copper, and selenium, but no significant differences in these elements contents between the osteoporosis group, osteopenia group, and healthy group</td>
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</table>
deficiency is directly defined as an independent risk factor for developing osteoporosis and in the research it is specified how this deficiency is related to lumbar osteoporosis both in the 45-59 years and 60-80 age groups years.

At an intermediate level there are two other studies, from 2009 and 2011. Liu et al in 2009 define the existence of a difference in blood zinc levels between osteoporotic, osteopenic and healthy women, but that however this difference is not statistically significant. Two years later, in 2011 Arikan reaches the same conclusion by comparing the same three groups.

Unlike these studies, in 1995 Relea et al, found statistically significant differences between osteoporotic and healthy women, not in the blood zinc levels (which are not different in this study), but in the urinary zinc excretion which is increased in a way significant in women with osteoporosis compared to healthy controls.

The study by Mahdavi-Roshan published in 2015 is totally opposite, where there is no difference in plasma zinc levels in the comparison between women with osteopenia and healthy women.

Zinc intake in relation to bone metabolism

This research was conducted based on the keywords: «zinc» and «zinc intake» and «bone» and «humans».

For the present review we have analyzed a total of three studies: two cohort studies and one cross-sectional study.

The results of these three studies have been shown in Table 2.

The daily zinc intake with food in relation to bone metabolism is analyzed in three studies, two of which substantially agree in the results.

In 1998, Elmstahl and colleagues study a large male population, consisting of 6,576 Swedish men aged between 46 and 68 years, concluding that low zinc intakes are a risk factor for fractures and that about 20% of the population studied took on inadequate quantities. The same results emerged from study by Hyun’s 2004 conducted on 396 men, where zinc intake levels are significantly reduced in subjects with osteoporosis.

Different results emerge from the 2015 Mahdavi-Roshan study on 51 menopausal women, where no difference in zinc intake was observed between women with osteoporosis, osteopenia and healthy; however, the study shows a general zinc intake lower than the values recommended by the RDA.

Zinc supplementation in relation to bone metabolism

This research was conducted based on the keywords: «zinc» and «zinc supplementation» and «bone» and «humans».

For the present review we have analyzed a total of five studies: three double-blind placebo controlled trials, one clinical trial and one randomized controlled trial.

The results of these five studies have been shown in Table 3.

As regards zinc supplementation, both alone and in associations with other nutrients, all the studies identified are substantially in agreement in defining beneficial effects for the bone (Table 3).

Already in 1974, in a study carried out on adolescents for 18 months, comparing an integration with 40 mg of zinc against placebo, there was an increase in bone age and bone development in the integrated group, with better results especially after 12 months of integration.

The effect of a zinc-only supplement on 60 men and women with fractures is also evaluated in the study by Sadighi in 2008; the subjects were divided into two groups destined to receive 220 g of zinc sulphate (corresponding to 50 g of zinc) or placebo and the final results show that in the group of subjects treated with zinc there was a faster healing of the fracture and significant change in bone callus formation 60 days after fracture.

In other three papers the effects of zinc in association with other nutrients are studied.

The study by Strause in 1994 involved healthy postmenopausal women divided into four groups (calcium supplement + micronutrient supplement, calcium supplement + micronutrient placebo, calcium placebo + micronutrient supplement, calcium placebo + micronutrient placebo), where the group that received both calcium and micronutrient supplementation, including zinc, maintained lumbar bone mineral density with a significant difference compared to the group that received only placebo; the remaining two groups positioned themselves at an intermediate level, without showing significant differences with the treated group or with the placebo group.

Subsequently, in 2011, the study by Nielsen and colleagues compares calcium supplementation versus calcium supplementation associated with zinc and copper in a group of menopausal women; the results confirm that zinc could bring beneficial effects on bone health only if the intake of zinc with diet is reduced (< 8 mg per day), while there were no significant beneficial effects with adequate zinc intake.

Always the same author, a few years earlier, wanted to check whether zinc supplementation could lead to changes in copper metabolism such as to lead to changes in bone turnover. The results did not lead to defining significant changes in copper metabolism even with high zinc supplements (53 mg per day), while this supplement led to an excessive excretion of magnesium. Low doses of zinc (3 mg per day) have instead caused unwanted changes in circulating osteocalcin and calcitonin.

Conclusion

We can define that blood zinc levels seem to be lower in subjects with pathology related to bone metabolism. The literature suggests that an insufficient daily intake of
### Table 2: Studies involving daily zinc intake in humans.

<table>
<thead>
<tr>
<th>First author, year</th>
<th>Study design</th>
<th>Setting</th>
<th>Inclusion criteria</th>
<th>Exclusion criteria</th>
<th>Number of subjects (M-F)</th>
<th>Lowest quintile intake/RDA or EAR</th>
<th>% subject in lowest quintile intake/% subject &lt; RDA or EAR</th>
<th>Highest Quintile intake</th>
<th>% subject in highest quintile intake</th>
<th>Primary outcomes</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hyun, 2004</td>
<td>Cohort study</td>
<td>All surviving from the original Rancho Bernardo cohort who still resided in southern California</td>
<td>≥ 45 years</td>
<td>396 men</td>
<td>The intakes in the lowest quintiles were 10 mg for zinc</td>
<td>A high proportion of the population, more than 20%, seems to be at risk of having inadequate dietary habits with respect to zinc</td>
<td>A lower fracture risk was noted in men with zinc intake in the second quintile [RR = 0.58 (0.34-0.99)] and fifth quintile [RR = 0.47 (0.25-0.89)] compared with the lowest quintile intake.</td>
<td>To examine the independent association between dietary zinc and plasma zinc and the association of each with bone mineral density (BMD)</td>
<td>Age- and BMI-adjusted dietary and total zinc intakes were significantly lower in the men with osteoporosis at the spine than in men without osteoporosis at that location</td>
<td>To determine dietary risk factors for fracture in men aged 46-68 years</td>
<td>Inadequate intakes of zinc are important risk factors for fracture</td>
</tr>
<tr>
<td>Elmstahl, 1998</td>
<td>Population-based prospective cohort study</td>
<td>City of Malmo, in the southern part of Sweden</td>
<td>Aged 46-68 years</td>
<td>6576 men</td>
<td>The intakes in the lowest quintiles were 10 mg for zinc</td>
<td>The intakes in the lowest quintiles were 10 mg for zinc</td>
<td>The intakes in the lowest quintiles were 10 mg for zinc</td>
<td>A lower fracture risk was noted in men with zinc intake in the second quintile [RR = 0.58 (0.34-0.99)] and fifth quintile [RR = 0.47 (0.25-0.89)] compared with the lowest quintile intake</td>
<td>To determine dietary risk factors for fracture in men aged 46-68 years</td>
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</tr>
<tr>
<td>Mahdavi-Roshan, 2015</td>
<td>Cross-sectional study</td>
<td>Rheumatology Clinic in Tabriz, Islamic Republic of Iran</td>
<td>Women &gt; 50 years old who had been no menstruation for ≥ 6 months prior to entry into the study, having no history of hormone replacement therapy, other bone disease, kidney stones, endocrine disorders or any medical conditions that could influence the mineral status</td>
<td>A total of 51 post-menopausal women</td>
<td>The mean dietary intake (and percent from RDA) of zinc in post-menopausal women with low bone density was 3.82 ± 0.19 mg/day (48 ± 2.41% RDA)</td>
<td>To investigate and compare the mineral status between osteopenic and osteoporotic postmenopausal women in Tabriz, Islamic Republic of Iran</td>
<td>The mean dietary intake of zinc was significantly lower than recommended dietary allowance (RDA). No statistically significant differences were observed between the osteopenic and osteoporotic groups with respect to dietary intake of zinc</td>
<td>To examine the independent association between dietary zinc and plasma zinc and the association of each with bone mineral density (BMD)</td>
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<tr>
<td>First author, year</td>
<td>Study design</td>
<td>Setting</td>
<td>Inclusion criteria</td>
<td>Exclusion criteria</td>
<td>Intervention</td>
<td>Parallel treatments</td>
<td>Number of subjects (M-F)</td>
<td>Duration of the intervention</td>
<td>Primary outcomes</td>
<td>Secondary outcomes</td>
<td>Results</td>
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<tr>
<td>Sadighi A, 2008</td>
<td>Randomized, double blind, placebo controlled clinical trial</td>
<td>Shohada Hospital of Tabriz, Iran</td>
<td>Men and women, aged 20-50 years old with traumatic long bone fracture</td>
<td>No history of osteoporosis, osteoarthritis, kidney stones, diabetes, and other endocrine disorders. Taking any medication or supplementation known to influence bone metabolism or zinc status</td>
<td>One capsule of 220 mg zinc sulfate contain 50 mg zinc</td>
<td>Control group receiving placebo contain starch</td>
<td>60 (39 M, 21 F)</td>
<td>60 days</td>
<td>Determine the effect of zinc supplementation on fracture healing</td>
<td>Determine the relation between callus formation with zinc and alkaline phosphatase activity in serum</td>
<td>Significant change in callus formation if the group will compare to the control group after 60 days (Figure 3), and fracture healing was faster in the supplement group than control group</td>
</tr>
<tr>
<td>Nielsen FH, 2011</td>
<td>Double-blind, placebo-controlled design</td>
<td>Postmenopausal women aged 51-80 years, BMI ≤ 32 kg/m², bone mineral density not more than 2.5 standard deviations below that for young adults; no collapsed/compressed vertebrae determined by using dual-energy X-ray absorptiometry (DXA); history of no menses for at least five years; and a circulating follicle-stimulating hormone concentration, 40 IU/l Eligible applicants were invited to an information meeting</td>
<td>A positive Pap smear or mammogram during the previous year, any disease or condition known to affect bone or calcium metabolism, a history of chronic renal, hepatic or gastrointestinal disease, evidence of collapsed or focal vertebral sclerosis</td>
<td>Groups 2) placebo calcium, active trace minerals; groups 3) active calcium, placebo trace minerals; and groups 4) active calcium, active trace minerals Subjects received placebo or 1,000 mg elemental calcium/d in the form of calcium citrate malate</td>
<td>Supplement containing 600 mg Ca plus a maize starch placebo</td>
<td>649 M</td>
<td>2 years</td>
<td>Determine whether increased Zn intakes would reduce the risk for bone loss</td>
<td>The findings indicate that Zn supplementation may be beneficial to bone health in postmenopausal women with usual Zn intakes &lt; 8.0 mg/d but not in women consuming adequate amounts of Zn</td>
<td></td>
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</tr>
<tr>
<td>Strause L, 1994</td>
<td>Double-blind, placebo-controlled trial</td>
<td>San Diego greater Metropolitan area</td>
<td>&gt; 50 y old and in good general health</td>
<td>Evaluate the impact of supplementary calcium with and without the addition of a combination of copper, manganese and zinc on spinal bone loss in healthy older post</td>
<td>Groups: 1) placebo calcium, placebo trace minerals</td>
<td>59 F</td>
<td>2 years</td>
<td>The findings indicate that Zn supplementation may be beneficial to bone health in postmenopausal women with usual Zn intakes &lt; 8.0 mg/d but not in women consuming adequate amounts of Zn</td>
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</tbody>
</table>

Table 3: Studies regarding zinc supplementation and bone metabolism.
Continue Table 3: Studies regarding zinc supplementation and bone metabolism.

<table>
<thead>
<tr>
<th>First author, year</th>
<th>Study design</th>
<th>Setting</th>
<th>Inclusion criteria</th>
<th>Exclusion criteria</th>
<th>Intervention</th>
<th>Parallel treatments</th>
<th>Number of subjects (M-F)</th>
<th>Duration of the intervention</th>
<th>Primary outcomes</th>
<th>Secondary outcomes</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nielsen FH, 2004</td>
<td>Randomized controlled trial</td>
<td>The metabolic unit of the Grand Forks Human Nutrition Research Center, Grand Forks, ND, USA</td>
<td>No underlying disease</td>
<td></td>
<td>Each active supplement contained 15.0 mg of zinc as sulfate salt, 2.5 mg of copper, and 5.0 mg of manganese as gluconate salts</td>
<td>25 postmenopausal women</td>
<td>200 days</td>
<td></td>
<td>menopausal women</td>
<td>Bone density Bone losses in the groups supplemented with trace mineral alone and with calcium alone were intermediate, but not significantly different from loss for either the placebo group or the group receiving calcium plus trace minerals</td>
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<tr>
<td>Ronaghy 1974</td>
<td>Clinical trial</td>
<td>Southern Iran</td>
<td>13-year-old prepubertal schoolboys</td>
<td></td>
<td>Grupo C) Zinc carbonate 40 mg supplement of egg-white protein (10 g daily), corn oil, minerals, and vitamins</td>
<td>Grupo A) placebo</td>
<td>Group B) supplement of egg-white protein (10 g daily), corn oil, minerals, and vitamins</td>
<td>18 months</td>
<td>To determine whether moderately high or low intakes of zinc adversely affect the copper status of postmenopausal women to result in unfavorable changes in calcium and magnesium metabolism and other indicators of bone turnover</td>
<td>Low dietary zinc (45.9 mmol/day; 3 mg/day) apparently resulted in undesirable changes in circulating calcitonin and osteocalcin. The moderately high intake compared to the low intake of zinc increased the excretion of magnesium in the feces and urine, which resulted in a decreased magnesium balance.</td>
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<td></td>
<td>Grupo C) Zinc carbonate 40 mg supplement of egg-white protein (10 g daily), corn oil, minerals, and vitamins</td>
<td>Grupo A) placebo</td>
<td>Group B) supplement of egg-white protein (10 g daily), corn oil, minerals, and vitamins</td>
<td>18 months</td>
<td>To learn whether these failures could have been in part the result of administration of insufficient quantities of zinc as a dietary supplement</td>
<td>Significantly increased heights, weights, and bone ages occurred in those receiving the supplementary zinc. During the 2nd year, bone development of the zinc-supplemented group surpassed that of the other groups by a substantial and statistically significant margin.</td>
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</tr>
</tbody>
</table>
zinc through nutrition (less than 3 mg/day) could be a risk factor for fractures and for the development of osteopenia and osteoporosis. A high proportion of the population, more than 20%, seems to be at risk of having inadequate dietary habits with respect to zinc. The supplementation of zinc in an amount equal to 40-50 g, on the other hand, could have beneficial effects on bone health in terms of maintaining bone mineral density and faster healing in the event of fractures, with even better results in the situation of reduced zinc intake through food.

References


Conflict of interest: None.