The determining role of silicon in bone formation - review and clinical results

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SHORT COMMUNICATION

Bone tissue consists of cells that reabsorb bone (osteoclasts) and cells that form and mineralize bone (osteoblasts). Osteoblasts produce collagen and hydroxyapatite, which together are the constituents of the mineral part of the bone.

The bone tissue is subjected to a continuous remodeling process, summarized below:

Silicon activates the prolyl-hydroxylase enzyme, present in the granular endoplasmic reticulum of the osteoblast cytoplasm. The enzyme governs the formation and bone mineralization.

This process is always present, for a lifetime

- with a positive balance up to around 30 years
- with negative balance subsequently (with a 2% drop every year)

This physiological decline mainly affects calcium and can be increased by other favourable factors:

- poor calcium and silicon diet or excessive sodium intake
- sedentary life
- high and prolonged intake of alcohol
- high and protracted coffee intake
- cigarette smoke
- the reduction of gonadal function that occurs in menopause involves a further increase in calcium loss (osteoporosis)
- even bad oral hygiene can adversely affect bone remodeling at the oral level
Silicon activates and ensures the normal and physiological synthesis of bone tissue

Common experience indicated that the diet enriched with *Equisetum arvense* had positive effects on the growth of farmed animals. So starting from the 60s it began to study which was the reason and it was highlighted that silicon was responsible for this result. In the cartilaginous and bone tissues there was a greater amount of silicon in the phase of growth compared to that present in the phases of quiescence and stability (1-5).

In all the cells involved in the development of the structural tissues (dermal matrix, cartilage and bone) is present the prolyl-hydroxylase enzyme which is determinant for their specific function. Through the activity of prolyl-hydroxylase the fibroblasts produce collagen for the intracellular matrix, the chondroblasts form the cartilage and the osteoblasts are able to produce collagen and hydroxyapatite, which are at the base of the formation of the mineral part of the bone tissue.(6).

Silicon, in fact, is indispensable in the early stages of collagen synthesis, conditioned by the activity of the enzyme prolyl-hydroxylase, silicon-dependent enzyme, present in the endoplasmic reticulum of cells that synthesize collagen (fibroblasts, chondroblasts and osteoblasts) (7-8).

Other studies have also documented that the addition of silicon from *Equisetum arvense* in the diet is able to restore and enhance the activity and functionality of these cells, improving any pathological conditions (9-11).

Consequently, the use of silicon supplementation can constitute a preventive and curative treatment, in all quantitative alterations of bone mass (osteoporosis, osteoarthritic or post-traumatic bone degeneration, bone fractures, orthopaedic and dental surgery, and guided bone regeneration): in fact, silicon promotes and activates, qualitatively and quantitatively, the bone remodelling processes (12-17).

In particular, the study by Mattioli-Belmonte et al. (18) has documented *in vitro* the efficacy of OSTEOSIL Calcium in improving the viability and metabolic function of osteoblasts. OSTEOSIL Calcium (Ghimas SpA, Casalecchio di Reno – Bologna, Italy) is a dietary supplement based on calcium and silicon present in *Equisetum arvense*.

Osteoblasts were cultured with OSTEOSIL Calcium in an increasing concentration of 2.5 μg/ml to 10 μg/ml and it was observed at SEM that the cell morphology was directly proportional to the concentration of OSTEOSIL Calcium. In cell cultures treated with the highest concentration, cell morphology was the best (good adhesion, cell viability and absence of degenerative aspects).

The concentrations between 2.5 and 10 μg/ml, achieved by the recommended dosage of OSTEOSIL Calcium, thus represent the optimal treatment for a balanced supply of silicon and calcium, even for prolonged periods of administration.

The EDAX micro analytic investigation also confirms that silicon and calcium administered with OSTEOSIL Calcium are usefully used by osteoblasts in their metabolism aimed at bone production.

Silicon activates prolyl-hydroxylase, which synthesizes collagen, a precursor of bone

Silicon, in an orally absorbable form, optimizes the activity of the silicon-dependent prolyl-hydroxylase enzyme, enzyme present in the endoplasmic reticulum of the osteoblast, which synthesizes the collagen, precursor of the bone, more rapidly and copiously.

The osteoblasts activated by the silicon are able to make better use of the blood calcium, which would otherwise be clinically very useful.

It is known that calcium, as it ages, undergoes its physiological decline, which often increases due to the concomitant presence of unfavourable conditions.
The available calcium or that taken with the diet, however, in the presence of silicon instead becomes more bioavailable and used for the phases of bone remodelling.

The functional synergism between osteoblasts and osteocytes allows to strengthen the formation of new bone and to improve its mineralization, obtaining a bone, which in implantology makes the implant stable.

Greater quantity and better bone quality in implantology with OSTEOSIL Calcium

This is the rationale underlying the clinical outcome observed after supplementation with OSTEOSIL Calcium in 124 subjects (45 females and 79 males, age 61.4 ± 14.6 years, range 41-79 years) who needed an implant surgery. The basal CT scan showed bone D1 in 15 cases, bone D2 in 37 cases, bone D3 in 54 cases and bone D4 in 18 cases.

After supplementation of OSTEOSIL Calcium to the daily dosage of two tablets for three months, the implant site showed an optimal bone for both quality and quantity during implant insertion, ensuring better implant stability in all cases.

The CT scan performed before implant insertion indicated a sharp increase in bone density (about 1.5% improvement in three months of OSTEOSIL Calcium supplementation): bone D1 in 17 cases, bone D2 in 67 cases and bone D3 in 40 cases, with disappearance of 18 cases with bone D4 (P <0.001 at the Mann Whitney U test).

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<th>CT (HU)</th>
<th>Micro CT (BV%)</th>
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<tbody>
<tr>
<td></td>
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<td>after</td>
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<tr>
<td>Num.</td>
<td>124</td>
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<tr>
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<td>741</td>
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<tr>
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<td>204</td>
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<td>1123</td>
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<tr>
<td>var. %</td>
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<td>1.32%</td>
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Variation of CT scan values before and after supplementation with OSTEOSIL Calcium. Descriptive statistical analysis (mean, SD, es, minimum and maximum value). The density of bone was first measured in Hounsfield units (HU) using computed tomography (CT) and also in bone volume % (BV%) using micro-CT scan. The percentage change between before and after is also indicated.

In implant surgery, it will thus be easy to find during supplementation with OSTEOSIL Calcium that the bone of the implant site is more hard and compact.

Silicon and calcium thus favour better calcification and a quicker mineralization of bone tissue, which in implantology results in faster bone formation.

It follows the logical possibility of being able to "load" the plants in shorter times with great consequent advantages both with the technique of immediate loading and in that of the early load, obtaining, in addition, an increase in the density of the newly formed bone.
Increase in the amount of bone and consequently in bone quality after supplementation with OSTEOSIL Calcium. Only calcium intake is not sufficient to induce a significant increase in the quantity of bone. The joint assumption of silicon, which activates the processes of the new bone formation, is decisive for the optimal reconstitution of the mineral share.

REFERENCES