The good, the bad, and the ugly of calcium supplementation: a review of calcium intake on human health

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Abstract: Calcium is an important integrative component of the human body and critical for human health. It has been well established that calcium intake is helpful in the prevention and treatment of osteoporosis, which has become one of the most serious public health problems across the world. However, community-dwelling adults with and without osteoporosis are rarely concerned or even not aware of the potential side effects of high or inappropriate doses of calcium intake. Some recent studies have revealed that excessive calcium intake might increase the risks of cardiovascular diseases. The purpose of this article was to review the health benefits, costs, and consequences of calcium supplementation on osteoporosis/osteoporotic fractures, cardiovascular events, kidney stones, gastrointestinal diseases, and other important diseases. In the end, we suggest that calcium supplementation should be prescribed and taken cautiously, accounting for individual patients’ risks and benefits. Clearly, further studies are needed to examine the health effects of calcium supplementation to make any solid recommendations for people of different genders, ages, and ethnicities.

Keywords: calcium, osteoporosis, osteoporotic fractures, cardiovascular diseases, kidney stones, gastrointestinal diseases

Introduction
Calcium supplementation is a widespread practice in different age-groups and has been promoted widely to improve bone density.1 The public generally believes that calcium is always good for health, owing largely to successful marketing and various clinical practice guidelines (Table 1). Based on the published data, 61% of women aged >60 years received calcium supplements in 2003–2006 in the United States.2 As a result, calcium supplementation has become a billion dollar market in recent years and has been taken by millions of both men and women, children, adults, and the elderly wishing to improve their skeletal health.

People may get more calcium from dietary intake or calcium supplementation. Due to the limited availability of calcium-rich foods and dietary habits, it is not always possible for people to take more and sufficient dietary calcium. Thus, there is a role for calcium supplementation for patients with inadequate dietary calcium intake. Initial studies suggested that dietary calcium had a greater effect on bone building due to its better absorption than supplemental calcium, but further studies are needed to fully elucidate whether one is better than the other to improve bone density.3

Recently, increasing concerns have been raised regarding whether indiscriminate calcium supplementation is safe. A meta-analysis by Bischoff-Ferrari et al demonstrated...
an increased hip fracture risk with calcium supplementation.\textsuperscript{4} A recent meta-analysis of randomized clinical trials showed that calcium supplementation was not associated with a low risk of fractures among community-dwelling older adults.\textsuperscript{5} There has been a concern for a significant increase in the risk of cardiovascular diseases with high supplemental calcium intake in men.\textsuperscript{6–8}

Given the emerging dilemma regarding calcium supplementation, the present article reviews discusses the potential benefits and side effects of calcium supplementation on health and diseases such as osteoporosis/osteoporotic fractures, cardiovascular events, kidney stones, gastrointestinal diseases, and others.

The basics: the role of calcium

It is well established that calcium is an extremely important and integrative component of human body, with 99\% of the body’s calcium being contained within the skeleton.\textsuperscript{9} Plasma calcium homeostasis plays a vital role in maintaining human life activities, such as maintenance of the skeleton, regulation of hormonal secretion, transmission of nerve impulses, and vascular activities.\textsuperscript{10–12} As shown in Figure 1, dietary calcium is absorbed mainly in the small intestine. Through circulation, it is mostly deposited into bones. Excessive or unabsorbed calcium is excreted in urine and feces.\textsuperscript{10} The homeostasis of calcium is mainly maintained by both parathyroid hormone (PTH) and calcitonin.\textsuperscript{13} When serum calcium level drops, PTH promotes calcium releases from the bones and stimulates reabsorption of calcium by the kidney tubules. In addition, PTH indirectly increases calcium absorption in the intestine via calcitriol, which is produced by the kidney and is the hormonal form of vitamin D. On the other hand, when serum calcium level rises, calcitonin would suppress calcium release from the bones\textsuperscript{10} and reduce calcium reabsorption by the kidney tubules.\textsuperscript{14}

### Potential beneficial effects of calcium supplementation

#### Osteoporosis

As calcium is one of the constituents of bone, the majority of researches considered that high calcium intake may result in higher bone density.\textsuperscript{1} According to the current status of our knowledge and understanding, improving accrual of peak bone mass and slowing the loss of bone after reaching the peak are two key factors in the prevention of osteoporosis. Moreover, bone mass grows most quickly during adolescence;\textsuperscript{15} thus, the prevention of osteoporosis should ideally start from this time.\textsuperscript{16} Importantly, calcium supplementation, together with vitamin D supplementation, is supported by the accumulating evidence for patients at high risk of calcium and vitamin D insufficiency and for those who receive treatment for osteoporosis.\textsuperscript{17}

Interestingly, some studies illustrated that calcium supplementation played a protective role for bone health, improving bone mass density (BMD) and decreasing morbidity of osteoporosis and osteoporotic fractures in different genders and age-groups.\textsuperscript{18} For children or adults, best evidence (grade A) is available for positive effects of calcium intake on the prevention of osteoporosis according to the National Osteoporosis Foundation (NOF).\textsuperscript{16} In addition, the administration of calcium was also shown to stave off fracture.\textsuperscript{19} Radford et al investigated the effects of calcium on fracture incidence and BMD after calcium supplement discontinuation; they concluded that the fractures in forearm and vertebrae were significantly decreased, but the positive benefits on BMD observed in the original

### Table 1 Calcium intake clinical practice guidelines for osteoporosis in different populations

<table>
<thead>
<tr>
<th>Source</th>
<th>Year published</th>
<th>Populations</th>
<th>Recommended intake (mg/day)</th>
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<td>Adults Postmenopausal women Old people</td>
<td>800 1,000 1,000</td>
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<td>2010</td>
<td>People aged &gt;50 years</td>
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Figure 1 Schema of calcium metabolism and homeostasis in human body.

Notes: Dietary calcium is absorbed mainly in the small intestine and mostly deposited into bones. Excessive or unabsorbed calcium is excreted in urine and feces. Plasma calcium level is mainly maintained by both PTH and calcitonin. When serum calcium level drops, PTH promotes calcium releases from the bones and stimulates reabsorption of calcium by the kidney tubules. In addition, PTH indirectly increases calcium absorption in the intestine via calcitriol. On the other hand, when serum calcium level rises, calcitonin would suppress calcium release from the bones and reduce calcium reabsorption by the kidney tubules.

Abbreviation: PTH, parathyroid hormone.

trial did not persist when the calcium supplementation was discontinued. Taking all the evidence into account, these studies all suggested that calcium supplementation, either in the form of calcium supplements or dietary calcium, played a positive role in people of different ages and genders, even though those results were obtained from studies of different sample sizes.

Postmenopausal women

A randomized, double-blind, placebo-controlled study performed on 1,471 postmenopausal women with 5-year follow-up, receiving either 1,000 mg/d calcium supplements or identical placebo, showed that calcium supplements could decrease bone loss. One meta-analysis, which included 29 randomized trials with men and women aged >50 years (including menopausal women), indicated that adequate calcium intake with/without vitamin D is associated with a reduced risk of fractures of any type (the RR =0.88, 95% CI =0.83–0.95) as well as provided a modest benefit on reducing the rate of bone loss, with 0.54% reduction of bone loss in the hip and 1.19% reduction of bone loss in the spine. In addition, there was a 24% risk reduction in osteoporotic fractures in people with high compliance rate for calcium supplementation. Furthermore, this study also suggested that a minimum dose of 1,200 mg/d calcium with 800 IU/d vitamin D was the best treatment for osteoporosis. Most studies involving postmenopausal women did not conduct data stratification to address whether calcium supplementation had similar or different effects in those women with or without osteoporosis. A Spanish study demonstrated a high prevalence of low calcium intake and vitamin D deficiency in postmenopausal women with osteoporosis. Further studies are needed to determine whether higher dose of calcium supplementation would be beneficial in this patient population. In one study performed exclusively in the postmenopausal women with no history of osteoporosis, 187 women aged >65 years were enrolled in a descriptive cross-sectional trial, revealing that adequate calcium intake along with proper exercise can effectively prevent osteoporosis. The recommended dosage of calcium supplementation for postmenopausal women with or without osteoporosis has been supported by the results of some large studies; 36,282 postmenopausal women were randomized into various groups with 1,000 mg/d calcium and vitamin D or placebo and followed for 7 years in the largest study, suggesting that hip bone density was improved with calcium supplements.
**Premenopausal women**
The current evidence is not sufficient to recommend calcium supplementation for the primary prevention of fractures in community-dwelling, asymptomatic premenopausal women. However, for those with a history of osteoporotic fractures, a diagnosis of osteoporosis, vitamin D deficiency, or a high risk for osteoporosis (eg, primary ovarian insufficiency), calcium supplementation may be considered.

**Pregnant and lactating women**
There is still a lack of clinical data to recommend calcium supplementation in pregnant or lactating women for the skeletal health of fetus and mother. A Cochrane database systemic review found that calcium supplementation during pregnancy did not have any clear additional benefits in the prevention of preterm birth or low infant birthweight. However, in pregnant and lactating women with low calcium intakes, calcium supplementation may be considered. Interestingly, in a study in pregnancy women in Gambia, West Africa, with low calcium intakes, calcium supplement resulted in significantly lower bone mineral content, bone area, and BMD at the hip throughout 12-month lactation. The women also had greater decreases in bone mineral during lactation at the lumbar spine and distal radius and had biochemical changes consistent with greater bone mineral mobilization.

**Men**
Although the evidence for preventing osteoporosis and improving BMD has been mostly generated in studies of postmenopausal women, a study back in 2008 compared the effects of two dosages of calcium citrate (600 mg/d, 1,200 mg/d, or placebo) on BMD in healthy men. The double-blind, randomized controlled trial performed on 323 healthy men aged >40 years for a 2-year period revealed that the 1,200 mg/d group showed increased BMD; however, it needs to be noted that urine calcium tended to increase by 57% correspondingly. In one study recently performed in Korean subjects, including 5,955 men, 1,256 premenopausal women and 4,494 postmenopausal women, it was revealed that increasing dietary calcium intake significantly improved BMD in both men and women.

**Children**
In a study conducted in rural Gambian children accustomed to a low-calcium diet, calcium supplementation resulted in higher bone mineral content and BMD. For healthy children, there has been no recommendation for routine calcium supplementation. In clinical practice, children with a high risk of osteoporosis (eg, celiac disease, inflammatory bowel disease, or congenital bone disorder) or low calcium intake may benefit from calcium supplementation.

**Cardiovascular disease**
There is evidence suggesting that cardiovascular disease might be protected by calcium supplementation. A prospective cohort study conducted among an elderly Chinese population indicated that dietary calcium intake (mean =600 mg/d) could reduce the risk of death from all causes and cardiovascular diseases. Despite the limitations (a self-administered questionnaire was used), its results added to the body of evidence that calcium supplementation decreases the mortality risk of cardiovascular diseases. Another prospective cohort study in Europe indicated that a daily administration of a mean of 820 mg dietary calcium has an effect of up to a 30% decrease in the risk of myocardial infarction. Similarly, in a prospective cohort study performed exclusively in >34,000 postmenopausal women, both the high intake of dietary and supplemental calcium were associated with a decrease in the mortality of ischemic heart disease. It was even observed that adverse cardiovascular effects were stopped after calcium supplements discontinued. Findings of these studies are likely to suggest that positive effects between calcium and cardiovascular diseases exist and that there is generally favorable evidence to imply that calcium influences cardiovascular disease occurrence. Calcium may slightly adjust lipid profiles and reduce blood pressure and – more importantly – the risk of developing hypertension. In prospective studies, the risks for developing hypertension were reduced with calcium supplements in men and women. In addition, preeclampsia is characterized by hypertension and would lead to perinatal morbidity and mortality of both mothers and infants. In 2013, the World Health Organization published a strong recommendation that pregnant women should take calcium supplements to prevent preeclampsia. A randomized controlled trial in women suggested that calcium supplementation >1,000 mg/d is associated with a significant reduction in the risk of preeclampsia.

**Gastrointestinal disease**
The effect of dietary calcium intake on the colorectal cancer and a comparison with the effect of higher and lower level of dietary calcium have been investigated by Galas et al. They concluded that an increase of 100 mg dietary calcium intake per day lowered by 5% the risk of colorectal cancer. Another observational study found that calcium consumption was inversely associated with the risk of developing colorectal cancer in a large population-based cohort follow-up. However, Galas et al. have observed that adverse cardiovascular effects were stopped after calcium supplements discontinued.
cancer (OR = 0.95, 95% CI = 0.92–0.99), and an increase of 1,000 mg/d dietary calcium consumption decreased incidence by 37% of colorectal cancer and the risk of colon cancer by 50%. Simultaneously, dietary calcium >1,000 mg/d decreased 46% incidence of colon cancer by 46% and the risk of colorectal cancer by 30%. A prospective cohort study revealed that the greatest benefits were observed with subjects who had highest dietary calcium intake (median = 914 mg/d) compared with subjects who had lowest calcium intakes at a median level of 486 mg/day.

A similar effect was observed with calcium supplement. A randomized, double-blind trial performed on people with a history of colorectal adenomas, receiving either 1,200 mg/d elemental calcium or placebo, revealed a significant reduction in the risk of recurrent colorectal adenomas with calcium supplementation. The explanation behind those findings is the ability of calcium to combine with bile acids in the intestines, reducing the rectal epithelial proliferation rate.

Most of the risk reduction benefit was achieved by calcium intake of at least 900 mg/day. Lower calcium intake was reported to increase the risk of colorectal cancer in Korean individuals with the CC genotype of the CASR gene in the single nuclear polymorphism of rs2270916, compared with those in higher calcium intake with the TT genotype (OR = 2.11, 95% CI = 1.27–3.51).

Kidney stone

Kidney stones are a common disease of the urinary system and are partially formed by calcium phosphate and calcium oxalate. Taylor and Curhan reported a relationship between calcium intake and urinary oxalate levels in a cross-sectional study. They concluded that high intake of dietary calcium or supplemental calcium (>500 mg/d) could lower urinary oxalate levels, subsequently decreasing the risk of kidney stones formation. Interestingly, a 12-year prospective study of 91,731 women without a history of kidney stones with 12-year follow-up suggested that high dietary calcium could reduce the risk of kidney stones, whereas the risk would be increased in women with high supplemental calcium. The mechanism of the different results caused by different types of calcium on kidney stones has not been definitively determined, but it is likely that dietary calcium decreases the absorption of oxalate and is probably associated with the timing of calcium ingestion relative to the amount of oxalate consumed. Consequently, it is reasonable to speculate that the effect of supplemental calcium on kidney stone risk depends on whether supplements are received with or between meals.

Aside from its beneficial roles on people, calcium is also used in the management of hypoparathyroidism and osteomalacia. Calcium supplements are considered a standard treatment for hypocalcemia associated with postsurgical hypoparathyroidism and achieves better clinical results.

Potential adverse effects of calcium supplementation

Despite the abovementioned studies providing powerful evidence for beneficial health-related effects of taking calcium, some recent evidence has implied that calcium supplementation might increase the risk for cardiovascular diseases and malignancy. What is more notable about these adverse effects is that, under the recommended dosage, calcium would produce some unwelcome reactions; therefore, a more detailed consideration of the safety and health benefits of calcium supplementation would be warranted. A number of adverse events are possibly influenced by calcium supplementation; these include myocardial infarction, constipation, colorectal neoplasms, and kidney stone. For example, in the Kuopio Osteoporosis Study, where 10,555 women were followed for 7 years, the HR for coronary heart disease in those who were taking calcium was 1.24 compared with nonusers. Thus, a doubt was for the first time raised in the beneficial effects of calcium supplementation on bone due to its adverse cardiovascular effects by this randomized control study. Moreover, a total of 1,460 postmenopausal women were randomized to taking calcium or a placebo and followed over 5 years, and the HR was found to be 1.12 for incident ischemic heart disease (95% CI 0.77–1.64). These surprising findings from the large related studies provide a stimulus for evaluating the effects of calcium on some disease events in other trials, which we describe below.

Osteoporosis

Although most studies indicate that calcium supplementation has a beneficial effect on osteoporosis and fractures, this statement is not confirmed by some other researches. Warenşjö et al found that dietary calcium intake <751 mg/d raised the risk of osteoporosis-related fractures and osteoporosis; meanwhile, intake >1,137 mg/d would increase the risk of hip fractures in women. A meta-analysis with seven cohorts totaling 170,991 women reported that calcium supplementation does not reduce the risk of hip fractures, and on the contrary, an increase in risk is possible. From these studies, the benefits of calcium supplementation become unclear, and there may be a therapeutic window where the greatest benefits on skeletal health are gained. Further research is needed to
clarify this, with respect to dietary intake, ethnicity, race, age, and gender.

**Cardiovascular diseases**

The side effects of calcium have long been focused on cardiovascular diseases as the principal concern. Cardiovascular diseases, especially myocardial infarction and myocardial ischemia, are among the leading causes of death around the world. It is estimated by a meta-analysis (including eleven randomized controlled trials) that calcium supplements have up to 30% increase risk for myocardial infarction.\(^5\) It was noteworthy that men or women aged >50 years were recommended to take ≥1,000 mg/d calcium according to the guidelines;\(^56\) however, the risk of cardiovascular diseases appears to be particularly increased when those doses are actually used. A prospective study found that supplemental calcium intake of >1,000 mg/d may raise the risk of death from cardiovascular diseases among men, but not among women.\(^6\) A randomized, placebo controlled trial performed on a population of 1,471 postmenopausal healthy women, with a follow-up that lasting 5 years, revealed that 1,000 mg/d elemental calcium supplements increased the incidence of cardiovascular events.\(^57\) In addition, a prospective observational study suggested that the use of calcium supplements in a long term would raise the risk of hospitalization in patients with chronic heart failure; for patients with diabetes, that risk was doubled.\(^58\) It is a remarkable fact that the effects of cardiovascular diseases between dietary and supplements of calcium lead to disparate results. A study indicated that the risk of myocardial infarction remarkably increased by taking calcium supplements compared with nonusers of any supplements (HR =1.86, 95% CI =1.17–2.96).\(^59\) The finding of an adverse trend in cardiovascular diseases with calcium supplementation is not necessarily surprising, as calcium supplements potentially contribute to elevated serum calcium levels and possibly accelerated cardiovascular calcification in the long run,\(^59\) which is predictive of cardiovascular event rates.\(^60\) Because of the high incidence of cardiovascular disease, any effects of calcium supplements on vascular health could be as important in terms of their effects on morbidity and mortality as their effects on osteoporosis or bone health.\(^61\) As these adverse effects are confirmed, a comprehensive reappraisal of taking calcium supplements may be required.

**Gastrointestinal diseases**

The second major safety concern with respect to calcium has been gastrointestinal diseases. The conclusion regarding the effects of calcium supplements on the gastrointestinal diseases is not totally consistent. Calcium supplements may increase the incidence of constipation, severe diarrhea, and abdominal pain.\(^1\)\(^2\) It highlights that calcium carbonate is more often associated with gastrointestinal side effects, including constipation, flatulence, and bloating.\(^1\)\(^8\) Of the 92,000 adverse events recorded, the occurrence of constipation was increased with calcium carbonate treatment (1,200 mg/d) in a 5-year, double-blind, placebo-controlled study.\(^53\) Moreover, some studies demonstrated a negative correlation between the intake of calcium and colorectal adenoma. A meta-analysis of 24 studies exploring the relationship between calcium supplementation and colorectal adenoma implied that the pooled RR was 1.13 for the occurrence of colorectal adenoma (95% CI =0.91–1.39).\(^63\) As stated previously, these findings are inconsistent as other research has shown reduction in the risk of colorectal cancer with calcium supplements.

**Kidney stone**

The third major concern regarding the safety of calcium has been the occurrence of kidney stones. Despite that calcium has positive effects on kidney stones by some researches, the literature is not consistent regarding the relationship between calcium and kidney stone formation. It was found that the abnormality of urinary supersaturation is one of the main factors for the formation of kidney stones.\(^64\) Specifically, levels of supersaturation above 1 can easily result in crystal deposits and growth and are considered a high-risk factor in kidney stone formation. The formation of urinary calcium oxalate stones is probably related to abnormal oxalate metabolism.\(^32\) Of note, people lacking adequate calcium intake should carefully consider the intake of oxalate-rich foods, such as spinach, chocolates, and teas.\(^33\) While studies indicate the potential beneficial effects of dietary calcium intake on the reduction of the risk of kidney stones, individuals who consumed any amount of supplemental calcium had an increased risk for kidney stones compared with individuals who did not consume supplemental calcium.\(^50\) A study performed in 53 postmenopausal women with 1,000 mg/d calcium (with 400 IU/d vitamin D) for 1 year showed a weak relationship with kidney stones formation; only one patient with urinary calculi was proved by ultrasonography. However, the small sample size and short clinical trial duration may contribute to lack of significant findings in this study.\(^65\) Nonetheless, a study conducted in 2006, including 36,282 postmenopausal women aged 50–70 years followed up for an average of 7 years, receiving either 1,000 mg/d elemental calcium or placebo, found a 17% increased risk of developing kidney
stones, with 449 women reporting kidney stones in the calcium with vitamin D group vs 381 women in the placebo group.\textsuperscript{24} Normally, stone formation in a normal kidney is relatively rare and is probably due to a combination of factors, such as health condition, urinary acidity, dietary habit, and variation in intake of calcium supplements (calcium oxalate, calcium citrate, and calcium carbonates). Dietary calcium is more beneficial for health in the aspect of kidney stone development.\textsuperscript{66}

Other diseases

In addition, adverse effects of calcium on other diseases such as age-related macular degeneration (AMD) and metabolic syndrome were reported. A recent cross-sectional study found that >800 mg/d of calcium consumption in people aged >67 years may increase the risk of AMD compared with those who do not take calcium (OR=2.63, 95% CI =1.52–4.54).\textsuperscript{67} A study in Korea found that men taking dietary calcium >1,200 mg/d had increased the prevalence of metabolic syndrome compared with those taking <1,200 mg/d, but no similar correlation was found in women.\textsuperscript{32}

For ease of reference, we summarized the above most salient points for the benefits and adverse effects in Table 2.

Conclusion

The issue of safety in people receiving calcium has long been a matter of debate. There is an abundance of data for beneficial effects and side effects in the literature on the role of calcium. All taken together, it is important to balance the advantages and disadvantages of calcium supplementation on human health. Based the literature, calcium supplementation is a double-edged sword in promoting bone formation or preventing osteoporosis, as it also may have a potential negative impact.

Table 2 The effects of calcium supplementation in diseases

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Questions remain as to whether taking extra calcium as calcium supplements is helpful to the general population. What is the real value of calcium intake by supplements per day for different individuals? Could it do more harm rather than good, particularly taking into consideration the increased risk of other diseases? More studies are needed to examine these effects for making better recommendations in different genders, ages, and ethnicities.

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Author contributions
HWD contributed to study conception, initiation, general development, and design. KL, XFW, YCC, YFG, and RZ collected data. XFW, DYL, YCC, XGL, LJZ, and YFG drafted the manuscript. KL, XFW, DYL, YCC, XGL, LJZ, JS, and HWD revised the final manuscript. All authors contributed to data analysis, drafting and revising the article, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Disclosure
The authors report no conflicts of interest in this work.

References


