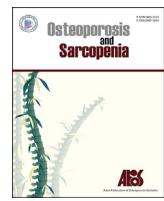




Osteoporosis and Sarcopenia

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Response to the editor

Reply on "Coffee consumption and bone health: A risk assessment"



To the editor

As pointed by Kawada [1] the mechanisms of caffeine consumption involved in bone health affect bone remodeling and falls risk.

Evidence of observational studies has shown an association between caffeinated beverages consumption and fragility fracture in women [2]. The effect of different drinks containing caffeine has been implicated as a cause of osteoporosis and fragility fracture [3]. Due to the information of observational studies in favor and against this association, there is controversy [4].

Some studies have suggested that the effect of coffee on bone health may be mediated by different mechanisms:

- Deleterious effect on osteoblasts [5,6].
- Increased urinary calcium excretion [7–9].
- Decrease in the efficiency of intestinal absorption [10].
- Low calcium intake in coffee drinkers [11,12].

The cytotoxicity of caffeine has been linked to apoptosis [5,13]. Caspases [14] and Bcl-2 family [15] play an important role in apoptosis, they regulate mitochondrial membrane potential changes and the release of cytochrome C by modulating the permeability of the outer mitochondrial membrane.

Heavy caffeine intake increases the urinary excretion of calcium, whereas moderate coffee consumption (1–2 cups per day) does not have a significantly impact on calcium imbalance in postmenopausal women [8]. It has been associated to a bone mineral density (BMD) loss between 2% and 4% depending on site, but with low risk for osteoporosis (odds ratio, 1.28; 95% confidence interval, 0.88–1.87). Thus, heavy ≥ 4 cups per day versus low (< 1 cup per day) coffee intake might be associated to slight decrease on BMD, without increasing the rate of fractures in women [4]. In men, according to a large cohort assessed by a self-administered food frequency questionnaire, a high coffee consumption was not associated with an increased rate of fractures [16].

At the same time, recent evidence suggests that coffee consumption can help reduce the risk of several diseases: type 2 diabetes, Parkinson disease, Alzheimer disease, cardiovascular disease, and cancer [9]. In the same way, it has been suggested that it might have a protective effect for disability in women with hypertension, obesity or diabetes, and on the risk of falls in elderly [17,18].

Recent information from a study that managed memory bias observed in self-reported questionnaire for coffee consumption and individual metabolism assessing coffee consumption through

metabolomics.

Conversely, a recent study that assessed coffee consumption through an individual assessment of metabolomics, reported a positive association of coffee metabolites, AFMU (5-acetylaminoo-6-formylamino-3-methyluracil), trigonelline, and 3-hydroxyhippurate, with BMD. Likewise, they reported that coffee consumption was directly associated with low rate of hip fractures [19].

Coffee may exert beneficial effects on bone health due to its high polyphenols composition; this impact may be especially prominent in men, who are resistant to caffeine-induced bone loss [20,21]. Recent evidence supports this assumption on younger [22] and older [16] males.

Most information in this matter derives from large epidemiological studies, which assessed coffee intake through self-applied questionnaires, introduction of metabolomics to this problem may bring some light to this matter. But still remains the need to establish the mechanisms of these effects. In the meantime, information about the effect of coffee intake on bone health should be taken with precautions, metabolites involved in this practice may have different effects on osteoporosis risk, bone resistance and risk of fracture.

Declaration of competing interest

No potential conflict of interest relevant to this article was reported.

References

- [1] Kawada T. Coffee consumption and bone health: a risk assessment. *Osteoporos Sarcopenia* 2020;6:33.
- [2] Hallström H, Wolk A, Glynn A, Michaélsson K. Coffee, tea and caffeine consumption in relation to osteoporotic fracture risk in a cohort of Swedish women. *Osteoporos Int* 2006;17:1055–64.
- [3] Samelson EJ, Hannan MT. Epidemiology of osteoporosis. *Curr Rheumatol Rep* 2006;8:76–83.
- [4] Hallström H, Byberg L, Glynn A, Lemming EW, Wolk A, Michaélsson K. Long-term coffee consumption in relation to fracture risk and bone mineral density in women. *Am J Epidemiol* 2013;178:898–909.
- [5] Lu PZ, Lai CY, Chan WH. Caffeine induces cell death via activation of apoptotic signal and inactivation of survival signal in human osteoblasts. *Int J Mol Sci* 2008;9:698–718.
- [6] Tsuang YH, Sun JS, Chen LT, Sun SC, Chen SC. Direct effects of caffeine on osteoblastic cells metabolism: the possible causal effect of caffeine on the formation of osteoporosis. *J Orthop Surg Res* 2006;1:7.
- [7] Massey LK, Opryszek AA. No effects of adaptation to dietary caffeine on calcium excretion in young women. *Nutr Res* 1990;10:741–7.
- [8] Hasling C, Søndergaard K, Charles P, Mosekilde L. Calcium metabolism in postmenopausal osteoporotic women is determined by dietary calcium and coffee intake. *J Nutr* 1992;122:1119–26.
- [9] Yang P, Zhang XZ, Zhang K, Tang Z. Associations between frequency of coffee consumption and osteoporosis in Chinese postmenopausal women. *Int J Clin Exp Med* 2015;8:15958–66.

- [10] Barger-Lux MJ, Heaney RP. Caffeine and the calcium economy revisited. *Osteoporos Int* 1995;5:97–102.
- [11] Barrett-Connor E, Chang JC, Edelstein SL. Coffee-associated osteoporosis offset by daily milk consumption. The Rancho Bernardo Study. *J Am Med Assoc* 1994;271:280–3.
- [12] Harris SS, Dawson-Hughes B. Caffeine and bone loss in healthy postmenopausal women. *Am J Clin Nutr* 1994;60:573–8.
- [13] Fernández MJ, López A, Santa-Maria A. Apoptosis induced by different doses of caffeine on Chinese hamster ovary cells. *J Appl Toxicol* 2003;23:221–4.
- [14] Martins LM, Kottke T, Mesner PW, Basi GS, Sinha S, Frigon Jr N, et al. Activation of multiple interleukin-1beta converting enzyme homologues in cytosol and nuclei of HL-60 cells during etoposide-induced apoptosis. *J Biol Chem* 1997;272:7421–30.
- [15] Thornberry NA, Rano TA, Peterson EP, Rasper DM, Timkey T, Garcia-Calvo M, et al. A combinatorial approach defines specificities of members of the caspase family and granzyme B. Functional relationships established for key mediators of apoptosis. *J Biol Chem* 1997;272:17907–11.
- [16] Hallström H, Wolk A, Glynn A, Michaëlsson K, Byberg L. Coffee consumption and risk of fracture in the Cohort of Swedish Men (COSM). *PLoS One* 2014;9:e97770.
- [17] Machado-Fragua MD, Struijk EA, Ballesteros JM, Ortolá R, Rodríguez-Artalejo F, Lopez-García E. Habitual coffee consumption and risk of falls in 2 European cohorts of older adults. *Am J Clin Nutr* 2019;109:1431–8.
- [18] Machado-Fragua MD, Struijk EA, Graciani A, Gualar-Castillón P, Rodríguez-Artalejo F, Lopez-García E. Coffee consumption and risk of physical function impairment, frailty and disability in older adults. *Eur J Nutr* 2019;58:1415–27.
- [19] Chau YP, Au PCM, Li GHY, Sing CW, Cheng VKF, Tan KCB, et al. Serum metabolome of coffee consumption and its association with bone mineral density: the Hong Kong Osteoporosis Study. *J Clin Endocrinol Metab* 2019 Nov 21. <https://doi.org/10.1210/clinem/dgz210> [Epub]. pii: dgz210.
- [20] Yano K, Heilbrun LK, Wasnich RD, Hankin JH, Vogel JM. The relationship between diet and bone mineral content of multiple skeletal sites in elderly Japanese-American men and women living in Hawaii. *Am J Clin Nutr* 1985;42:877–88.
- [21] Holbrook TL, Barrett-Connor E, Wingard DL. Dietary calcium and risk of hip fracture: 14-year prospective population study. *Lancet* 1988;2:1046–9.
- [22] Choi MK, Kim MH. The association between coffee consumption and bone status in young adult males according to calcium intake level. *Clin Nutr Res* 2016;5:180–9.

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