Letter to the Editor



Influence of Osteoporosis on Blood Lead Levels in Postmenopausal Korean Women

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Dear Editor-in-Chief

Osteoporosis is a significant global public health issue affecting over 200 million people worldwide and the prevalence of osteoporosis among women aged 50 and older ranges from 15.4% to 16.5% in the United States (1). Potential risk factors for osteoporosis encompass older age, low body mass index (BMI), cigarette smoking, and excessive alcohol consumption (2). Notably, estrogen deficiency is a substantial risk factor for osteoporosis in postmenopausal women (3). Among environmental pollutants, lead (Pb) is emerging as a significant risk factor for osteoporosis (4). Among adults, 80-95 % of retained Pb is stored in bone, with a half-life of approximately 20-30 years (5). These bone Pb deposits are released into the bloodstream during periods of heightened bone resorption, such as menopause, potentially serving as an endogenous source of Pb supply (6). Postmenopausal women face an elevated risk of bone Pb release due to hormonal and age-related alterations in bone metabolism. Hence, to evaluate the influence of osteoporosis on blood Pb levels in postmenopausal women, data including sociodemographic factors and blood Pb levels were extracted from the KNHANES 2016–2017 databases.

Of the 1,301 participants, the geometric mean blood Pb level was 1.76 μ g/dL; the Pb concentration among postmenopausal women with osteoporosis was significantly lower than that among those without osteoporosis (P = 0.016) (Table 1).

Characteristics	N	Osteoporosis group (N=281)	None osteoporosis group (N=1,020)	P-value*
Total Age (yr)	1301	1.68 (1.61–1.76)	1.79 (1.75–1.83)	0.016
50-59	528	1.56 (1.39-1.73)	1.77 (1.71–1.83)	0.024
60–69	409	1.69 (1.59–1.80)	1.80 (1.73–1.88)	0.112
≥ 70	364	1.72 (1.60–1.86)	1.81 (1.73–1.89)	0.294

Table 1: Geometric mean Pb [μ g/dL (95% CI)] by the prevalence of osteoporosis in Korean women aged \geq 50

*P-Values from t-test for difference in geometric mean lead concentrations between groups



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Participants with osteoporosis consistently had lower blood lead concentrations than those without osteoporosis across all age groups, with the most notable difference being the 50–59-year-old group which displayed a statistically significant difference (P = 0.024). The prevalence and ORs of osteoporosis demonstrated clear association with blood Pb levels. The OR for osteoporosis exhibited a significant decrease with increasing blood Pb concentration in both ageadjusted (Model 1) and fully-adjusted (Model 2) models (Table 2).

Table 2: Prevalences and adjusted odds ratios (95% CI) of osteoporosis by tertile of blood Pb in Korean wom-
en aged ≥ 50 years

Osteoporosis —	Tertile blood Pb level (µg/dL)			P for trend
	Tertile 1 (< 1.50)	Tertile 2 (1.50–2.05)	Tertile 3 (> 2.05)	
OR (95%				
CI)*				
Model 1	1.00 (reference)	0.88 (0.58-1.33)	0.59 (0.38-0.91)	< 0.001
Model 2	1.00 (reference)	0.87 (0.56–1.34)	0.55 (0.35-0.86)	< 0.001

*Model 1: adjusted for age. Model 2: adjusted for age, BMI, education, income, cigarette smoking status, and alcohol drinking status

The relationship between osteoporosis and blood metal levels is complex and depends on various factors including age, sex, lifestyle habits, and race/ethnicity. Previous studies have reported that the association between blood Pb levels and bone mineral density is influenced by factors such as race and menopausal status (7). In osteoporosis, accelerated bone resorption releases stored metals like zinc, copper, and iron back into the bloodstream. This increase in circulating metals can ultimately lead to their enhanced excretion through urine, feces, and sweat, potentially lowering their blood concentrations (8). However, it is important to note that this effect may not be uniform for all metals, and further research is needed to fully understand the specific impacts of osteoporosis on individual metal levels.

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Conflict of Interest

The authors declare that they have no conflict of interest.

References

- Shen Y, Huang X, Wu J, et al (2022). The global burden of osteoporosis, low bone mass, and its related fracture in 204 countries and territories, 1990-2019. *Front Endocrinol (Lausanne)*, 13:882241.
- 2. Kanis JA, Cooper C, Rizzoli R, et al (2019). European guidance for the diagnosis and management of osteoporosis in postmenopausal women. Osteoporos Int, 30(1):3-44.
- North American Menopause Society (2021). Management of osteoporosis in postmenopausal women: the 2021 position statement of The North American Menopause Society. *Menopause*, 28(9):973-97.

- Shahida S, Rehman S, Ilyas N, et al (2021). Determination of blood calcium and lead concentrations in osteoporotic and osteopenic patients in Pakistan. ACS Omega, 6(42):28373-8.
- Patrick L (2006). Lead toxicity, a review of the literature. Part 1: Exposure, evaluation, and treatment. *Altern Med Rev*, 11(1):2-22.
- Machida M, Sun SJ, Oguma E, Kayama F (2009). High bone matrix turnover predicts blood levels of lead among perimenopausal women. *Environ Res*, 109(7):880-6.
- Campbell JR, Auinger P (2007). The association between blood lead levels and osteoporosis among adults--results from the third national health and nutrition examination survey (NHANES III). *Environ Health Perspect*, 115(7):1018-22.
- 8. Zheng J, Mao X, Ling J, He Q, Quan J (2014). Low serum levels of zinc, copper, and iron as risk factors for osteoporosis: a meta-analysis. *Biol Trace Elem Res*, 160(1):15-23.