

OSTEOPOROSIS TREATMENT GAP – A SYSTEMATIC LITERATURE REVIEW

VRZEL V ZDRAVLJENJU OSTEOPOROZE - SISTEMATIČNI PREGLED LITERATURE

Kaja GRIL ROGINA ^{1,2*} , Luka PETRAVIČ ^{2,3} , Christos OIKONOMIDIS ¹ 

¹ National Institute of Public Health - Regional Unit Maribor, Ljubljanska ulica 4, 2000 Maribor, Slovenia

² University of Ljubljana, Faculty of Medicine, Vrazov trg 2, 1000 Ljubljana, Slovenia

³ University Medical Center Maribor, Ljubljanska ulica 5, 2000 Maribor, Slovenia

Received: Oct 30, 2025

Accepted: Jun 03, 2026

Review article

ABSTRACT

Keywords

Osteoporosis

Fragility fractures

Healthcare disparities

Treatment gap

Care gap

Introduction

Osteoporosis is a common metabolic bone disease that increases fracture risk, particularly among older adults, and imposes a substantial global burden. Despite advances in risk assessment and treatment, a large treatment gap remains, with many high-risk individuals not receiving care.

Methods

We conducted a systematic review of English-language studies published up to April 2025, identified through searches in multiple databases (PubMed, Scopus, Web of Science, ScienceDirect, Europe PMC). 43 studies were included. Due to the heterogeneity of study designs and treatment gap definitions, a narrative thematic synthesis was performed.

Results

Reported treatment gap estimates varied widely (38.8-94.6%) across countries and study populations. Lower treatment rates were associated with patient factors (male sex, lower socio-economic status, comorbidities) and healthcare system factors (limited physician awareness, institutional barriers, poor interprofessional communication). Higher treatment rates were associated with older age, family history of osteoporosis, prior diagnosis or bone mineral density testing, and participation in structured care programmes.

Conclusions

Substantial global treatment gaps in osteoporosis care persist, particularly following fragility fractures. These gaps arise from multiple patient, physician, and system-level factors, with persistent disparities by sex, ethnicity, comorbidities, and socio-economic status. Findings highlight the need for standardised definitions of the treatment gap and suggest areas for future research and intervention development.

IZVLEČEK

Ključne besede

osteoporoza

osteoporotični zlomi

vrzeli v zdravljenju

neenakosti v obravnavi

Uvod

Osteoporoza je pogosta presnovna bolezen kosti, ki povečuje tveganje za zlome, zlasti pri starejših odraslih, in predstavlja pomembno globalno zdravstveno breme. Kljub napredku pri oceni tveganja za zlome in razpoložljivosti učinkovitih možnosti zdravljenja, vrzel v zdravljenju osteoporoze vztraja, saj številni posamezniki z visokim tveganjem, predvsem starostniki, ne prejemajo ustrezne oskrbe. Namen tega sistematičnega pregleda je kategorizirati vzroke za vrzel v zdravljenju osteoporoze.

Metode

Opravili smo sistematični pregled študij v angleškem jeziku, objavljenih do aprila 2025 v podatkovnih bazah PubMed, Scopus, Web of Science, ScienceDirect, Europe PMC. V analizo je bilo vključenih 43 študij. Zaradi heterogenosti zasnov študij in definicij vrzeli je bila izvedena narativna tematska sinteza.

Rezultati

Ocene vrzeli v zdravljenju se med državami in okolji gibljejo med 38,8 % in 94,6 %. Dejavniki, povezani z nižjimi stopnjami zdravljenja, vključujejo moški spol, nižji socialno-ekonomski status, določene pridružene bolezni, pa tudi dejavnike, povezane z zdravstvenim sistemom, kot so pomanjkanje ozaveščenosti pri zdravnikih, institucionalne ovire ter nezadostno medpoklicno sodelovanje. Višje stopnje zdravljenja so povezane s starostjo, družinsko anamnezo osteoporoze, predhodno diagnozo ali meritvijo kostne gostote ter učinkovito usklajeno oskrbo v okviru specializiranih programov.

Zaključki

Kljub učinkovitim diagnostičnim in terapevtskim možnostim po vsem svetu, vrzel v zdravljenju osteoporoze vztraja, zlasti pri ljudeh po zlomu. Ta vrzel izvira iz dejavnikov na ravni bolnika, zdravstvenih delavcev in zdravstvenega sistema ter se kaže v doslednih razlikah glede spola, etnične pripadnosti, pridruženih bolezni in socialno-ekonomskega položaja. Reševanje teh izzivov bo zahtevalo integrirane poti oskrbe, izboljšano klinično izobraževanje in razvoj modelov za usklajeno obravnavo po zlomu.

*Correspondence: grilroginakaja@gmail.com

Published under Creative Commons Attribution 4.0 International (CC BY 4.0)

© 2026 The Authors

1 INTRODUCTION

Osteoporosis is the most prevalent metabolic bone disease, characterised by decreased bone mass and microarchitectural deterioration of bone tissue, which increases fracture risk (1). Approximately one in three women and one in six men will experience an osteoporotic fracture during their lifetime. In the European Union, more than 23 million individuals are at high risk of such fractures (1, 2). Globally, 178 million new fractures occurred in 2019, representing a 33.4% increase since 1990 and accounting for 25.8 million years lived with disability (YLDs), a 65.3% increase over the same period (2).

Osteoporotic fractures predominantly affect older adults, particularly women, and typically result from low-energy trauma (2, 3). The most common sites include the hip, vertebrae, distal radius, and proximal humerus. These fractures are associated with chronic pain, functional decline, loss of independence, and reduced quality of life (3). Hip fractures in particular frequently require hospitalisation and are associated with complications such as thrombosis (27%), urinary tract infections (12-61%), and pneumonia (7%) (4). Osteoporosis is diagnosed based on low bone mineral density measured by DXA or the presence of a fragility fracture, particularly at the hip or vertebra. Treatment decisions are additionally informed by fracture risk assessment tools such as FRAX (1, 4).

Despite advances in fracture risk assessment, preventive strategies, and clinical guidelines, a large proportion of individuals at high risk remain untreated (1, 5). The SCOPE 21 report estimates an average treatment gap of 71% across Europe among women at high fracture risk (predominantly postmenopausal), ranging from 87% in Bulgaria to 32% in Ireland; in Slovenia, it is approximately 57% (1). The treatment gap is widest among older adults, particularly those in long-term care (5). This gap contrasts with other areas of secondary prevention, such as post-myocardial infarction care, where treatment rates have improved substantially over time (6). Reducing the osteoporosis treatment gap is essential, given its substantial contribution to fracture burden and associated healthcare costs (1).

This systematic review aims to identify and categorise the reported reasons for the osteoporosis treatment gap, intending to inform future interventions to address this issue.

2 METHODOLOGY

This review was conducted as a systematic review of observational studies addressing determinants of the osteoporosis treatment gap and is reported according to PRISMA guidelines (7). The study protocol was registered in PROSPERO (8) on the 3rd of April 2025.

2.1 Search strategy and information sources

The electronic search strategy was developed using combinations of MeSH terms and free-text keywords related to osteoporosis and treatment gaps. A representative PubMed (9) search strategy was as follows: (“Osteoporosis”[Mesh] OR osteoporosis[tiab]) AND (“treatment gap”[tiab] OR “care gap”[tiab] OR “treatment disparity”[tiab] OR “under-treatment”[tiab] OR “under-diagnosis”[tiab]). Equivalent search strategies were adapted for Scopus (10), Web of Science (11), ScienceDirect (12), and Europe PMC (13). Searches included studies published up to the 7th of April 2025. Full search strategies for all databases are available from the authors upon reasonable request. Grey literature was not systematically searched; however, reference lists of included studies were screened for additional eligible publications. A total of 1457 references were imported to the Systematic Review Accelerator (SRA) platform (14), where duplicates were removed using the Deduplicator Tool (15). Non-English studies were excluded during initial screening due to feasibility constraints.

2.2 Eligibility criteria

Studies were eligible if they met the following criteria:

1. Addressed the osteoporosis treatment gap;
2. Investigated reasons for the treatment gap;
3. Possibly provided an estimate of the treatment gap size.

Adult populations at risk of osteoporosis, with or without prior fragility fractures, were included. “Treatment” was defined as pharmacological therapy for osteoporosis as reported in individual studies. There is no universally accepted definition of the “treatment gap” in osteoporosis. Across studies, it variably refers either to non-treatment following a fragility fracture or to non-treatment among individuals at high fracture risk (based on FRAX, with or without DXA). We therefore did not impose a single definition, but extracted and reported study-specific definitions and interpreted findings within their respective contexts.

2.3 Screening and selection process

Two authors (K.G.R. and L.P.) independently screened titles and abstracts. Discrepancies were resolved through discussion. Full texts of potentially eligible studies were then retrieved and assessed independently for inclusion by the same reviewers. Non-original research articles and non-English publications were excluded. The selection process was recorded in sufficient detail to complete a PRISMA (7) flow diagram (Figure 1).

2.4 Data extraction

A standardised data extraction form was developed and piloted on a subset of included studies. Data were independently

extracted by two reviewers (K.G.R. and L.P.) using Microsoft Office 365 Excel (16). Missing data were not sought from the study authors. Given the heterogeneity of included studies, a narrative thematic synthesis was conducted.

2.5 Risk of Bias Assessment

Risk of bias (RoB) was assessed using the Newcastle-Ottawa scale for cohort (17) and cross-sectional studies (18) and the Critical Appraisal Skills Programme (CASP) checklist for qualitative studies (19).

3 RESULTS

A total of 1,457 records were identified across five databases. After screening, 43 original research studies were included (20-62). The PRISMA flow diagram is presented in Figure 1 (7).

3.1 Study analysis

Of the included studies, 5 were prospective cohort studies (20, 29, 37, 41, 43), 22 were retrospective cohort studies (21-28, 30-36, 38-40, 42, 58, 61-62), 15 were cross-sectional studies (44-57, 60), and 1 was a qualitative study (59). 11 studies were conducted in European countries (27-31, 45-48, 50, 57), 17 in the United States and Canada (24, 33-34, 36-43, 51-53, 58, 60-61), 9 in Asian countries (21-23, 25-26, 32, 35, 49, 62), 3 in Australia (20, 56, 59), 2 in northern Africa (44, 54) and 1 study combined data from the US, Canada and India (55) (Figure 2). Most studies were conducted in secondary or tertiary care settings (21-24, 31-32, 37, 41-42, 47, 53) or in mixed healthcare settings (20, 25-27, 29-30, 33-36, 38-40, 43-44, 46, 49, 51-52, 54-59, 61-62), while five studies were performed exclusively in primary care settings (28, 45, 48, 50, 60). The majority of studies included both male and female participants (20-28,

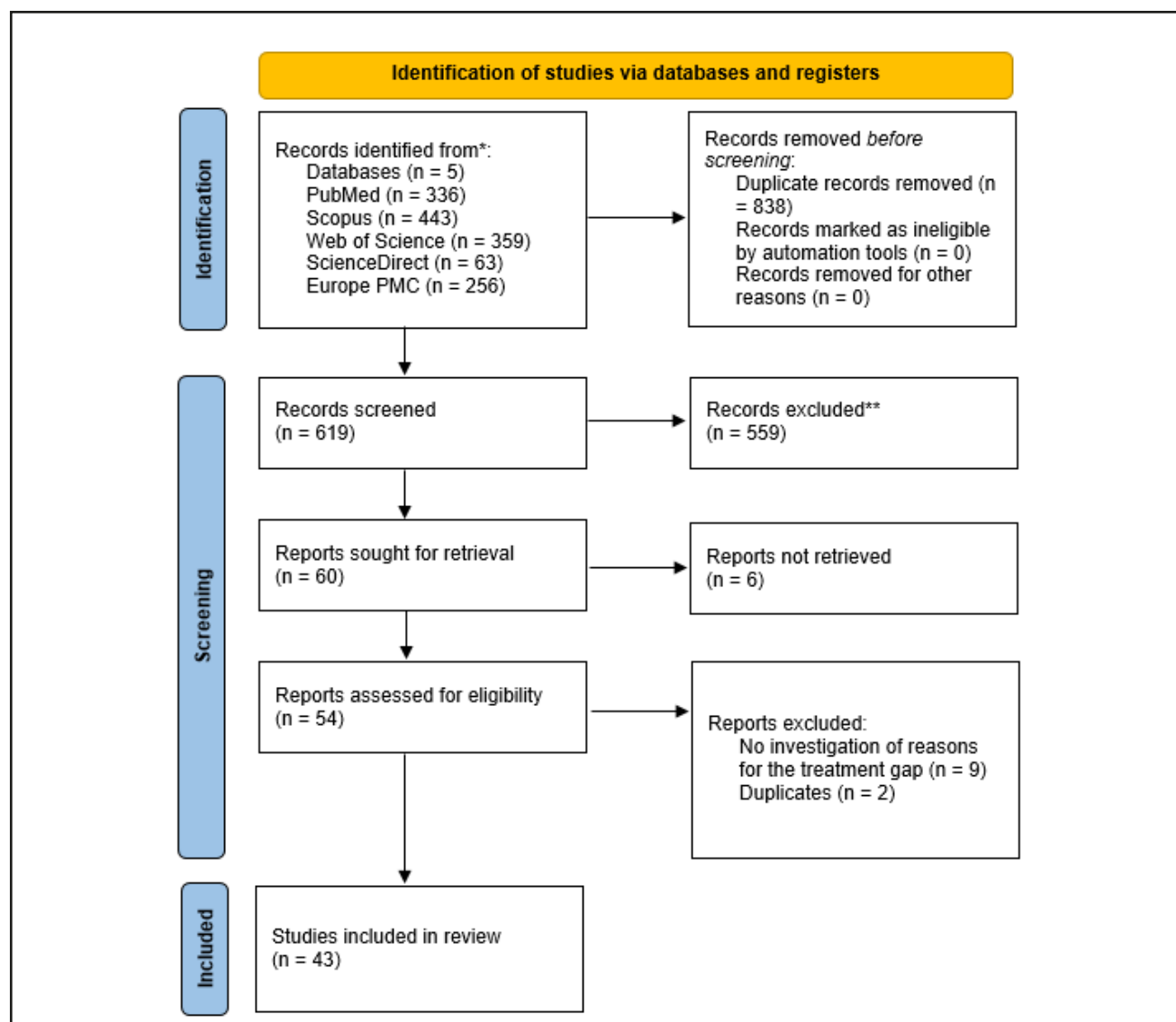


Figure 1: PRISMA 2020 flow diagram (7) representing the selection of 43 (20-62) original research studies included in the systematic review.

30-32, 35-37, 39-40, 42, 44-47, 50-52, 58-60, 62). Seven studies focused specifically on postmenopausal women (29, 33-34, 38, 41, 48-49), one included only male patients (43), and one examined patients with cancer of both sexes (61). Several studies were survey-based and targeted healthcare professionals (53-57). The summary data from the 43 included studies were organised by outcomes.

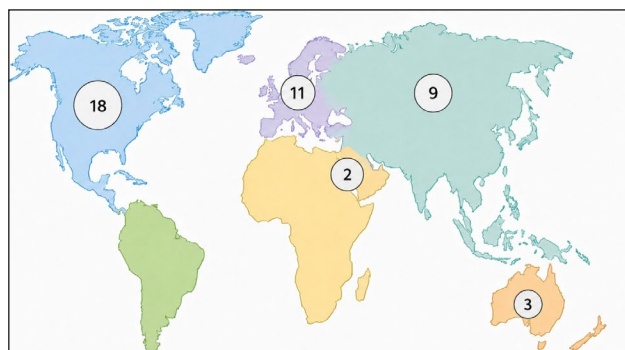


Figure 2: Geographical distribution of included studies (20-62).

3.2 Primary outcome

The factors contributing to the treatment gap were categorised (Table 1). Patient-level factors were defined broadly to include both demographic characteristics and clinical or disease-related factors, as these were not consistently distinguished across studies.

3.3 Secondary outcome

The treatment gap estimates are presented in Table 2. Because treatment gap estimates were derived from clinically distinct populations, including post-fracture secondary prevention cohorts and broader osteoporosis/high-risk populations, direct comparison of estimates across these categories should be interpreted with caution.

3.4 Risk of bias assessment

For cohort studies, NOS scores (17) of 7-9 were considered “good”, 5-6 “fair”, and ≤ 4 “poor”. For cross-sectional studies, adapted NOS scores (18) of 8-10 were considered “good”, 6-7 “fair”, and < 6 “poor”. CASP checklist (19) gives a descriptive rather than a numerical grade. Overall, 32 of 43 studies were rated as “good” quality (21-30, 33, 35-42, 44-52, 56-59, 61), 8 as “fair” (20, 32, 34, 43, 53, 55, 60), and 4 as “poor” (31, 54, 57, 62). The use of non-validated measurement tools, the lack of sample size justification, and inadequate reporting of response rates limited the two cross-sectional studies, which were rated as poor quality (54, 57). The two cohort studies rated as poor quality (31, 62) were affected by issues including non-representative sampling, failure to ensure absence of the outcome at baseline, reliance on self-reported data, absence of a non-exposed cohort, and insufficient adjustment for confounding factors. Risk of bias assessments informed interpretation, with greater emphasis placed on higher-quality studies. Determinants from studies with a higher risk of bias were generally consistent with those from higher-quality studies; however, all findings were interpreted with caution.

Table 1: Overview of patient-related (demographic and clinical), physician-related, system-level, and organisational factors associated with variations in osteoporosis treatment rate (20–62).

Dimensions		Factors contributing to lower treatment rates	Factors contributing to higher treatment rates
Patient-level factors (demographic and clinical)	Patient Demographics	- Male sex (22, 24–26, 30–32, 34–37, 40, 45–46, 50–52, 61) - Age: in most studies (21, 28–29, 35, 37, 40–41, 46, 49, 50–51, 61), older people are more likely to be treated; some studies show the opposite (26, 31, 38) - Body mass index of 30 and over (34, 51) - Minority racial and ethnic groups (marginalised groups) (21, 33–34, 39, 50–52)	
	Social History	- Lower socio-economic status (40, 61) - Lower education levels (23) - Lower annual income (49) - Socioeconomically disadvantaged residential areas (40) - Current tobacco use (34)	Married status (49)
	Family History		Family history of osteoporosis or fragility fractures (31, 51)
	Fracture Type	- No documented previous fragility fractures (31) - Non-vertebral fractures (26–27, 33, 58, 61) - Non-major osteoporotic fracture (21, 37)	
	Past Medical History and Investigations	- Dementia (22, 33) - Diabetes (28, 31, 38) - Chronic liver disease (28) - Chronic kidney disease (28, 33, 62) - Cardiovascular disease and hypertension (28, 38) - Arthritis (34) - Gastrointestinal troubles (38) - Comorbidity (20, 45, 60), high Charlson comorbidity index score (23, 33, 58)	- Rheumatoid arthritis and systemic lupus (28, 51) - Diagnosis of osteoporosis or bone mineral density testing before a fragility fracture (21, 29, 36, 43–44, 48, 58) - Low bone mineral density score (25, 41)
	Pharmacotherapy	- Polypharmacy (26, 45, 57) - Oestrogen-only hormone replacement therapy (28)	- Corticosteroids (28, 51, 58) - Prior vitamin D and calcium prescriptions (41)
	Economic Factors	Cost of treatment (49, 54, 60)	
	Patient Belief & Behaviour	- Low adherence (47) - Refusal of treatment (60) - Belief that osteoporosis is not serious (49) - More frequent doctor visits (51)	

Dimensions		Factors contributing to lower treatment rates	Factors contributing to higher treatment rates
Healthcare professionals and healthcare system-level factors	Physician-related factors	• Lack of osteoporosis awareness (54) • Limited familiarity with treatment (47, 53) • Fear of anti-osteoporotic medications side effects (as perceived barrier by physicians) (54, 60) • Workload concerns (53) • Differences in prescription practices (46, 61-62) • Differences in referral practices (62) • Risk assessment not routine practice (44, 47, 55) • Failure to investigate (47, 56) • Knowledge gaps and lack of confidence (53) • Lack of comfort with making treatment changes (60) • Intentions to modify the treatment plan (60) • Disagreement with recommendations (60), reservations about protocol-based approaches (53) • Preference for Fracture Liaison Service (FLS) programmes over standard care (53) • Perception by general practitioners that preventive therapy is a specialist's responsibility (57) • Interprofessional communication issues and role ambiguity (44, 59) • Belief that the current care plan was appropriate (60) • Specialist-prescribed treatment plans limiting other providers' interventions (60)	Both physician and system-related factors: - Transfer to a rehabilitation (42) or a geriatric unit (42) - Preference for FLS (53)
	System/ Institutional factors	• Lack of patient education and awareness (44, 49, 55, 59) • Long intervals between fracture and outpatient visit (25) • Length of hospital stay (42) • Bone-strengthening or community-based physical activity programmes rare (55)	

Table 2: Country-wise summary of treatment gap estimates in included studies (20-41, 43-48, 51-52, 56, 58, 60-62).

Author & Year	Country	Treatment Gap Estimate	Treatment Gap Type*	Author & Year	Country	Treatment Gap Estimate	Treatment Gap Type*
Feldstein 2005 (58)	USA	84% (post-hip or vertebral fracture)	Post-fracture secondary prevention	Leslie 2020 (61)	Canada	77.1% of cancer patients and 78.2% of non-cancer patients	Post-fracture secondary prevention
Papaioannou 2008 (43)	Canada	90%	Post-fracture secondary prevention	McCloskey 2020 (48)	Belgium, France, Germany, Ireland, Poland, Slovakia, UK	74.6% (varying from 53% in Ireland to 91% in Germany)	Osteoporosis/ high-risk population
Bessette 2009 (41)	Canada	84%	Post-fracture secondary prevention	Skjødt 2020 (30)	Denmark, Spain (Catalonia), UK	Denmark: 88-90% (2005-2014)	Post-fracture secondary prevention
Curtis 2009 (52)	USA	74% even among the highest risk group (Caucasian men with prior fracture)	Osteoporosis/ high-risk population	Malle 2021 (47)	Austria	82% in women, 92% in men	Post-fracture secondary prevention
Leslie 2011 (40)	Canada	93.9% in 1996/1997, 87.7% in 2001/2002, 94.1% in 2007/2008	Post-fracture secondary prevention	Nakatoh 2021 (26)	Japan	68.4% (hip fracture), 38.3% (vertebral fracture)	Post-fracture secondary prevention
Leslie 2012 (39)	Canada	94.6% in First Nations peoples vs 87.5% in non-Indigenous	Post-fracture secondary prevention	Papaioannou 2021 (60)	Canada	63%	Osteoporosis/ high-risk population
Roux 2013 (37)	Canada	81.2% in the standard care group	Post-fracture secondary prevention	Skjødt 2021 (27)	Denmark	85% (2005), 79% (2014)	Post-fracture secondary prevention
Siris 2013 (38)	USA	64.3%	Osteoporosis/ high-risk population	Walsh 2021 (28)	Ireland	79%	Osteoporosis/ high-risk population
Cunningham 2014 (51)	USA	38.8%	Osteoporosis/ high-risk population	Hoit 2022 (24)	USA	74% (30 days after operation)	Post-fracture secondary prevention
Yusuf 2016 (36)	USA	85.1%	Post-fracture secondary prevention	Martinez-Laguna 2022 (45)	Spain	56.8%	Post-fracture secondary prevention
Gani 2017 (32)	Singapore	68.5%	Post-fracture secondary prevention	Wu 2022 (25)	Taiwan	64.4%	Post-fracture secondary prevention

Author & Year	Country	Treatment Gap Estimate	Treatment Gap Type*	Author & Year	Country	Treatment Gap Estimate	Treatment Gap Type*
Keshishian 2017 (33)	USA	72.2%	Post-fracture secondary prevention	Yavropoulou 2022 (46)	Greece	58%	Osteoporosis/high-risk population
Mendis 2017 (56)	Australia	87%	Post-fracture secondary prevention	Bliuc 2023 (20)	Australia	78% (women), 86% (men)	Post-fracture secondary prevention
Sattari 2017 (34)	USA	78.4%	Osteoporosis/high-risk population	Cheah 2023 (21)	Malaysia	82.9%	Post-fracture secondary prevention
Wang 2017 (35)	Taiwan	> 60% for women, 90% for men	Osteoporosis/high-risk population	El Miedany 2023 (44)	Egypt	82.1%	Post-fracture secondary prevention
Cheung 2018 (62)	Hong Kong	From 91% to 85% between 2009 and 2012	Post-fracture secondary prevention	Fujii 2023 (22)	Japan	60%	Post-fracture secondary prevention
Bougioukli 2019 (31)	Greece	78.6%	Post-fracture secondary prevention	Mahaisavariya 2023 (23)	Thailand	68.4%	Post-fracture secondary prevention
Iconaru 2020 (29)	Belgium	74.6%-76.5%	Post-fracture secondary prevention				

*Post-fracture secondary prevention refers to treatment eligibility assessed following a fragility fracture. Osteoporosis/high-risk population refers to individuals identified through osteoporosis diagnosis, low bone mineral density, fracture-risk assessment tools (FRAX or CAROC), or population-based estimates of osteoporosis prevalence.

4 DISCUSSION

Despite growing awareness of the importance of fracture prevention (63), substantial treatment gaps in osteoporosis care persist globally. These gaps are consistently driven by patient-, physician-, and system-level factors, in line with previous reviews (64-65).

4.1 Treatment gap estimates

This systematic review found reported treatment rates ranging from 38.8% to 94.6% across studies. However, comparisons between studies are limited by substantial methodological heterogeneity. To facilitate interpretation, treatment gap estimates were classified according to whether they were derived from post-fracture secondary prevention populations or broader osteoporosis/high-risk populations. Nevertheless, direct comparisons between these categories should be interpreted with caution due to differences in study populations, eligibility criteria, and treatment gap definitions.

There is no uniform definition or approach to calculating the osteoporosis treatment gap. Some studies assessed treatment following fragility fractures as a form of secondary

prevention (20-32, 36-37, 39-41, 43, 47, 56, 58, 61-62), while others included broader high-risk populations, including individuals defined by DXA criteria or FRAX-based risk (29). In several studies, the timing of fracture or treatment initiation was unclear (34-35, 38, 44-46, 48, 50-52).

Despite these differences, similar treatment gaps were observed across clinically distinct groups. For example, one study reported comparable gaps in postmenopausal women defined by T-score ≤ -2.5 and those defined by prior major osteoporotic fracture (29), suggesting that undertreatment is systemic across both diagnosed osteoporosis and high-risk populations.

Overall, variability in definitions and methodologies limits direct comparison and highlights the need for standardised approaches to defining and reporting the treatment gap.

4.2 Factors contributing to the treatment gap

The identified determinants were grouped into patient-, physician-, and system-level factors to aid interpretation; this categorisation is descriptive and does not imply causal relationships.

4.2.1 Patient-level factors (demographic and clinical)

Male sex, comorbidity burden, socio-economic disadvantage, and reduced health literacy were consistently associated with lower treatment rates. Male sex was repeatedly linked to lower treatment likelihood despite substantial fracture burden in men (22, 24-26, 30-32, 34-37, 40, 45-46, 50-52, 61), consistent with evidence that up to 40% of osteoporotic fractures occur in men (66). Age showed a non-linear association with treatment. Most studies reported higher treatment rates with increasing age (21, 28-29, 35, 37, 40-41, 46, 49, 50-51, 61), likely reflecting greater recognition of increased fracture risk. However, very advanced age and multimorbidity were sometimes associated with lower treatment rates (23, 26, 31, 38), suggesting competing clinical priorities and concerns regarding frailty, polypharmacy, and life expectancy. Comorbid conditions, especially dementia, diabetes, cardiovascular and chronic illnesses, were consistently associated with lower treatment rates (20, 22-23, 28, 31, 33-34, 38, 45, 60, 62). The same can be said for polypharmacy (26, 45, 57), with physicians often withholding treatment due to concerns over side effects, drug interactions, and life expectancy considerations.

In multiple studies (21, 33-34, 39, 50-52), minority and Indigenous populations were consistently less likely to receive treatment compared to white/Caucasian peers, even after a fracture. This disparity highlights structural inequities and cultural or healthcare system barriers in osteoporosis care. Racial and ethnic characteristics are inextricably linked to socio-economic status (SES). Lower SES (40, 61), lower education (23) and lower income (49) were strongly associated with osteoporosis treatment gaps. Cost concerns (49, 54, 60) were explicitly cited by patients as reasons for treatment refusal, especially among lower-income groups. A lower SES is indicative of less frequent and lower-quality care and reduce access to care (67). Additionally, health professionals' implicit bias against patients with low SES has a negative effect on clinical decision-making (68).

Additionally, limited health literacy and awareness of osteoporosis and fracture risk were pervasive across both high- and low-income countries (31, 40, 47, 49, 51). Patient perceptions, attitudes towards osteoporosis, and health behaviours were frequently cited as barriers to assessment and treatment initiation. Malle et al. (47) found that patients lacked an understanding of the indications for treatment or its potential to prevent future fractures. Patients often discontinued medications early due to side effects or a misunderstanding of benefits. Commonly reported issues included a belief that osteoporosis is not a serious condition, particularly in the absence of symptoms or fractures, leading many to decline treatment offers. Roh et al. (49) highlighted that women who believed osteoporosis was a minor health concern or inconvenient to assess were significantly less

likely to undergo evaluation or treatment, even when at high risk. A study of postmenopausal women treated with monthly bisphosphonates (69) found that patients' belief in the importance of osteoporosis treatment was a key predictor of adherence.

Clinical severity influenced treatment likelihood. Vertebral and major osteoporotic fractures were more likely to trigger treatment than non-major fractures (21, 26-27, 33, 37, 58, 61). Prior diagnosis of osteoporosis, lower BMD, or prior fractures also increased the likelihood of treatment (21, 25, 29, 31, 36, 41, 43-44, 48, 58).

4.2.2 Physician and system-level factors

Healthcare professionals and system factors emerged as major contributors to the treatment gap. Physician-related barriers were frequently identified, ranging from limited familiarity with osteoporosis management guidelines to concerns about medication side effects, particularly in older or comorbid patients. Beshyah et al. (54) reported that fear of medication adverse effects, workload constraints, and discomfort with protocol-based management approaches were commonly cited reasons. Papaioannou et al. (60) found that many primary physicians intentionally deferred treatment decisions due to perceived appropriateness of current care plans, concerns about intolerance, or uncertainty about modifying specialist-initiated regimens. Interprofessional communication challenges also negatively impacted care continuity. One study (59) highlighted role ambiguity and poor information transfer between system levels, contributing to delayed or missed treatment opportunities. A lack of integrated care pathways and post-fracture assessment services further exacerbated care gaps.

System-level barriers included logistical obstacles such as long waiting times for specialist consultations, limited access to bone density testing, and regional disparities in service provision. Leslie et al. (40) demonstrated marked geographic differences in treatment rates across urban, rural, and remote populations. Furthermore, logistical difficulties, including transportation challenges in older and disabled patients (23) and the absence of community-based fracture prevention programmes (55), compound inequities in access to osteoporosis care.

The observational nature of the included studies limits the interpretation of these determinants. Reported associations do not allow clear separation of confounding from causation, particularly for variables such as age, comorbidity burden, and polypharmacy, which may reflect underlying clinical complexity or competing care priorities rather than independent effects. Furthermore, the distinction between modifiable and non-modifiable factors is not always clear-cut, as observed differences (by sex, age, socio-economic status, or other factors) may reflect broader structural or healthcare system influences.

Therefore, these findings should be interpreted cautiously and primarily as indicators of patterns within osteoporosis care rather than direct targets for intervention.

4.3 Limitations and strengths

This systematic review has several limitations. First, the review was restricted to English-language publications, which may introduce language bias. This is particularly relevant in a global context, as non-English studies may capture underrepresented healthcare settings and system-level barriers. Consequently, the findings may not fully reflect the global diversity of osteoporosis care and may have limited generalisability. Second, no additional data were sought from the original study authors, which may have limited the completeness of extracted data and the risk of bias assessment. Although the risk of bias was systematically assessed, its impact on the interpretation of findings should be considered. Studies rated as having a higher risk of bias were retained to ensure a comprehensive overview, but were interpreted with caution. Most determinants identified in fair- and poor-quality studies were also reported in studies of good methodological quality, including male sex, ethnicity, comorbidity burden, prior osteoporosis diagnosis or bone mineral density testing, treatment costs, and the lack of routine fracture risk assessment. In contrast, several healthcare professional and organisational factors, including limited familiarity with osteoporosis treatment, knowledge gaps, workload concerns, perceptions regarding responsibility for osteoporosis management, referral and prescription practices, and constraints related to specialised treatment plans, were identified primarily in studies with a higher risk of bias (53, 57, 60, 62) and should therefore be interpreted with greater caution. Nevertheless, these factors may represent important hypotheses for future research.

The observational design of included studies, variability in study quality, heterogeneity in outcome definitions, population characteristics and follow-up time limit the strength of inference.

However, the review has notable strengths. It employed a comprehensive search strategy across five major databases, ensuring broad coverage. It includes a diverse, multinational dataset comprising 43 original research studies (20-62) from a wide range of healthcare systems and populations, thereby improving the findings' generalizability. The categorisation of factors into patient, physician, and system-level domains allowed for a structured analysis of the multifactorial nature of osteoporosis under-treatment. RoB was systematically assessed using validated tools (17-19), adding methodological rigour.

4.4 Contributions to the field

This review synthesises evidence on gaps in osteoporosis treatment across multiple healthcare systems, providing a comprehensive overview of the determinants of undertreatment. By structuring these into patient-, physician-, and system-level domains, it highlights the multifactorial nature of the problem.

It also demonstrates substantial international variability in post-fracture care, reflecting differences in healthcare organisation, clinical practice, and access to fracture prevention services. Physician-level barriers, such as uncertainty about treatment initiation and inconsistent care pathways, represent actionable targets for intervention. Finally, the review identifies key research gaps, including the optimisation of care for men, improved risk stratification, and the development of scalable post-fracture care models.

5 CONCLUSIONS

This systematic review confirms that substantial osteoporosis treatment gaps persist globally, particularly following fragility fractures. These gaps are driven by a combination of patient-, physician-, and system-level factors, including demographic disparities, comorbidity burden, socio-economic disadvantage, and healthcare system limitations.

Marked international variability in post-fracture treatment reflects differences in healthcare organisation, clinical guidelines, and access to diagnostic and treatment services. Physician-related barriers, including concerns about treatment safety, limited confidence in osteoporosis management, and fragmented care pathways, further contribute to undertreatment.

Future research should focus on optimising management for high-risk groups, such as men, improving risk-stratification tools, and developing scalable models of integrated post-fracture care to reduce global disparities in treatment.

ACKNOWLEDGEMENT

We thank the Public Health Department at the University of Ljubljana Medical Faculty for invaluable advice and guidance.

CONFLICTS OF INTEREST

The authors declare that no conflicts of interest exist.

FUNDING

The study did not receive any additional funding.

ETHICAL APPROVAL

Ethical approval was not required for this study, as it is a systematic review of previously published data.

DATA AVAILABILITY STATEMENT

All data and materials supporting the results of this study are presented in the main text, with any additional information available from the corresponding author upon reasonable request.

AI USAGE STATEMENT

During the preparation of this manuscript, the authors used the GPT language model to check grammar and spelling, refine wording, and assist with reference formatting. The authors subsequently reviewed and edited all content and accept full responsibility for the final version of the publication.

ORCID

Kaja Gril Rogina:
<https://orcid.org/0000-0002-3195-7652>

Luka Petravič:
<https://orcid.org/0000-0001-7087-7316>

Christos Oikonomidis:
<https://orcid.org/0009-0006-4591-5097>

REFERENCES

1. Kanis JA, Norton N, Harvey NC, Jacobson T, Johansson H, Lorentzon M, et al. SCOPE 2021: A new scorecard for osteoporosis in Europe. *Arch Osteoporos*. 2021 Dec;16(1):82. doi: 10.1007/s11657-020-00871-9.
2. World Health Organization. Fragility fractures [Internet]. 2024 [cited 2025 Apr 22]. Available at: <https://www.who.int/news-room/fact-sheets/detail/fragility-fractures>
3. Kates SL, Miclau T. The fracture liaison service: Its history, current state, how it works, and future directions. *OTA International*. 2022 Jun;5(3S):e192. doi: 10.1097/OI9.0000000000000192.
4. Ayub N, Faraj M, Ghatan S, Reijers JAA, Napoli N, Oei L. The treatment gap in osteoporosis. *J Clin Med*. 2021 Jul 5;10(13):3002. doi: 10.3390/jcm10133002.
5. Giangregorio L, Papaioannou A, Cranney A, Zytaruk N, Adachi JD. Fragility fractures and the osteoporosis care gap: An international phenomenon. *Semin Arthritis Rheum*. 2006 Apr;35(5):293-305. doi: 10.1016/j.semarthrit.2005.11.001.
6. Choudhry NK, Winkelmayer WC. Medication adherence after myocardial infarction: A long way left to go. *J Gen Intern Med*. 2008 Feb;23(2):216-8. doi: 10.1007/s11606-007-0478-8.
7. Page MJ, McKenzie JE, Bossuyt PM, Boutron I, Hoffmann TC, Mulrow CD, et al. The PRISMA 2020 statement: An updated guideline for reporting systematic reviews. *BMJ*. 2021 Mar 29;372:n71. doi: 10.1136/bmj.n71.
8. National Institute for Health and Care Research. PROSPERO [Internet]. 2019 [cited 2025 Apr 22]. Available at: <https://www.crd.york.ac.uk/prospéro>
9. National Library of Medicine. PubMed [Internet]. 2025 [cited 2025 Apr 22]. Available at: <https://pubmed.ncbi.nlm.nih.gov>
10. Scopus. Scopus preview [Internet]. 2025 [cited 2025 Apr 22]. Available at: <https://www.scopus.com>
11. Clarivate. Web of Science [Internet]. 2024 [cited 2025 Apr 22]. Available at: <https://clarivate.com/webofscience>
12. ScienceDirect: Science, health and medical journals, full text articles and books [Internet]. 2025 [cited 2025 Apr 22]. Available at: <https://www.sciencedirect.com>
13. Europe PMC. [Internet]. 2016 [cited 2025 Apr 22]. Available at: <https://europepmc.org>
14. Systematic Review Accelerator. SR-Accelerator [Internet]. 2024 [cited 2025 Apr 22]. Available at: <https://sr-accelerator.com>
15. Systematic Review Accelerator: Deduplicator tool. SR-Accelerator [Internet]. 2024 [cited 2025 Apr 22]. Available at: <https://sr-accelerator.com/#/deduplicator>
16. Microsoft Excel. Microsoft Excel, spreadsheet software [Internet]. 2024 [cited 2025 Apr 22]. Available at: <https://www.microsoft.com/en-us/microsoft-365/excel>
17. Ottawa Health Research Institute. The Newcastle-Ottawa Scale (NOS) for assessing the quality of nonrandomised studies in meta-analyses [Internet]. 2021 [cited 2025 Apr 22]. Available at: https://www.ohri.ca/programs/clinical_epidemiology/oxford.asp
18. Ottawa Health Research Institute. Newcastle-Ottawa Scale adapted for cross-sectional studies [Internet]. 2022 [cited 2025 Apr 22]. Available at: https://repository.up.ac.za/bitstream/handle/2263/84405/Nyawo_SystematicSuppl1_2022.pdf?sequence=2
19. CASP. Critical appraisal checklists [Internet]. 2018 [cited 2025 Apr 22]. Available at: <https://casp-uk.net/casp-tools-checklists/>
20. Bliuc D, Tran T, Chen W, Alarkawi D, Alajlouni DA, Blyth F, et al. The association between multimorbidity and osteoporosis investigation and treatment in high-risk fracture patients in Australia: A prospective cohort study. *PLoS Med*. 2023 Jan;20(1):e1004142. doi: 10.1371/journal.pmed.1004142.
21. Cheah MH, Lai PSM, Khor HM, Ratnasingam J, Chandrasekaran CSK, Singh S, et al. Fragility fracture care gap at a tertiary teaching hospital in Malaysia. *Arch Osteoporos*. 2023 May;18(1):63. doi:10.1007/s11657-023-01256-4.
22. Fujii T, Mori T, Komiyama J, Kuroda N, Tamiya N. Factors associated with non-initiation of osteoporosis pharmacotherapy after hip fracture: Analysis of claims data in Japan. *Arch Osteoporos*. 2023 Jul;18(1); doi:10.1007/s11657-023-01314-x.
23. Mahaisavariya C, Vanitcharoenkul E, Kitcharanant N, Chotiarnwong P, Unnanuntana A. Exploring the osteoporosis treatment gap after fragility hip fracture at a Tertiary University Medical Center in Thailand. *BMC Geriatr*. 2023 Feb 3;23(1):70. doi:10.1186/s12877-023-03778-5.
24. Hoit G, Whelan D, Atrey A, Ravi B, Ryan G, Bogoch E, et al. Association of age, sex and race with prescription of anti-osteoporosis medications following low-energy hip fracture in a retrospective registry cohort. *PLoS One*. 2022 Dec 1;17(12):e0278368. doi:10.1371/journal.pone.0278368.
25. Wu Y, Chen C, Chang C, Li C, Wang R, Sun Z, et al. Association between outpatient visits and initiating medication among elderly patients after an osteoporotic vertebral fracture. *J Clin Med*. 2022 Sep;11(17):5035. doi:10.3390/jcm11175035.
26. Nakatoh S, Fujimori K, Ishii S, Tamaki J, Okimoto N, Ogawa S, et al. Insufficient increase in bone mineral density testing rates and pharmacotherapy after hip and vertebral fracture: Analysis of the National Database of Health Insurance Claims and Specific Health Checkups of Japan. *Arch Osteoporos*. 2021 Dec;16(1); doi:10.1007/s11657-021-00992-9.

27. Skjødt M, Ernst M, Khalid S, Libanati C, Cooper C, Delmestri A, et al.. The treatment gap after major osteoporotic fractures in Denmark 2005-2014: A combined analysis including both prescription-based and hospital-administered anti-osteoporosis medications. 2021 Oct;32(10). doi: 10.1007/s00198-021-05890-x.
28. Walsh ME, Nerdrum M, Fahey T, Moriarty F. Factors associated with initiation of bone-health medication among older adults in primary care in Ireland. *Age Ageing*. 2021 Sep;50(5):1539-1546. doi:10.1093/ageing/afab033.
29. Iconaru L, Smeys C, Baleanu F, Kinnard V, Moreau M, Cappelle S, et al. Osteoporosis treatment gap in a prospective cohort of volunteer women. *Osteoporos Int*. 2020 Jul;31(7):1535-1543. doi:10.1007/s00198-020-05339-7.
30. Skjødt M, Khalid S, Ernst M, Rubin K, Martinez-Laguna D, Delmestri A, et al. Secular trends in the initiation of therapy in secondary fracture prevention in Europe: A multi-national cohort study including data from Denmark, Catalonia, and the United Kingdom. *Osteoporos Int*. 2020 Aug;31(8):1535-1544. doi:10.1007/s00198-020-05358-4.
31. Bougioukli S, Kollia P, Koromila T, Varitimidis S, Hantes M, Karachalios T, et al. Failure in diagnosis and under-treatment of osteoporosis in elderly patients with fragility fractures. *J Bone Miner Metab*. 2019 Mar;37(2):327-335. doi: 10.1007/s00774-018-0923-2.
32. Gani L, Reddy SK, Alsuwaigh R, Khoo J, King TFF. High prevalence of missed opportunities for secondary fracture prevention in a regional general hospital setting in Singapore. *Arch Osteoporos*. 2017 Dec;12(1):60. doi: 10.1007/s11657-017-0356-x.
33. Keshishian A, Boytsov N, Burge R, Krohn K, Lombard L, Zhang X, et al. Examining the treatment gap and risk of subsequent fractures among females with a fragility fracture in the US Medicare population. *Osteoporos Int*. 2017 Aug;28(8):2361-2369. doi: 10.1007/s00198-017-4072-6.
34. Sattari M, Cauley J, Garvan C, Johnson K, LaMonte M, Li W, et al. Osteoporosis in the Women's Health Initiative: Another treatment gap? *Am J Med*. 2017 Aug;130(8):935-942. doi: 10.1016/j.amjmed.2017.02.042.
35. Wang CY, Fu SH, Yang RS, Shen LJ, Wu FLL, Hsiao FY. Age- and gender-specific epidemiology, treatment patterns, and economic burden of osteoporosis and associated fracture in Taiwan between 2009 and 2013. *Arch Osteoporos*. 2017 Oct;12(1):35. doi: 10.1007/s11657-017-0385-5.
36. Yusuf AA, Matlon TJ, Grauer A, Barron R, Chandler D, Peng Y. Utilization of osteoporosis medication after a fragility fracture among elderly Medicare beneficiaries. *Arch Osteoporos*. 2016 Dec;11(1):25. doi: 10.1007/s11657-016-0285-0.
37. Roux S, Beaulieu M, Beaulieu MC, Cabana F, Boire G. Priming primary care physicians to treat osteoporosis after a fragility fracture: An integrated multidisciplinary approach. *J Rheumatol*. 2013 May;40(5):751-759. doi: 10.3899/jrheum.120908.
38. Siris E, Modi A, Tang J, Sen S. Substantial under-treatment among women diagnosed with osteoporosis in a US managed-care population: A retrospective analysis. *Curr Med Res Opin*. 2013 Jan;30(1):97-104. doi: 10.1185/03007995.2013.851074.
39. Leslie W, Brennan S, Prior H, Lix L, Metge C, Elias B. The post-fracture care gap among Canadian First Nations peoples: A retrospective cohort study. *Osteoporos Int*. 2012 Mar;23(3):985-991. doi: 10.1007/s00198-011-1880-y.
40. Leslie W, Giangregorio L, Yogendran M, Azimae M, Morin S, Metge C, et al. A population-based analysis of the post-fracture care gap 1996-2008: The situation is not improving. *Osteoporos Int*. 2011 May;23(5):771-779. doi: 10.1007/s00198-011-1630-1.
41. Bessette L, Jean S, Davison K, Roy S, Ste-Marie L, Brown J. Factors influencing the treatment of osteoporosis following fragility fracture. *Osteoporos Int*. 2009 Nov;20(11):1873-1880. doi: 10.1007/s00198-009-0898-x.
42. Haaland DA, Cohen DR, Kennedy CC, Khalidi NA, Adachi JD, Papaioannou A. Closing the osteoporosis care gap: Increased osteoporosis awareness among geriatrics and rehabilitation teams. *BMC Geriatr*. 2009 Jul;9:28. doi: 10.1186/1471-2318-9-28.
43. Papaioannou A, Kennedy C, Ioannidis G, Gao Y, Sawka A, Goltzman D, et al. The osteoporosis care gap in men with fragility fractures: The Canadian Multicentre Osteoporosis Study. *Osteoporos Int*. 2008 Apr;19(4):425-430. doi: 10.1007/s00198-007-0483-0.
44. El Miedany Y, El Gaafary M, Gadallah N, Mahran S, Fathi N, Abu Zaid MH, et al. Osteoporosis treatment gap in patients at risk of fracture in Egypt: A multi-center, cross-sectional observational study. *Arch Osteoporos*. 2023 May;18(1):26. doi: 10.1007/s11657-023-01252-8.
45. Martínez-Laguna D, Carbonell C, Bastida JC, González M, Micó-Pérez RM, Vargas F, et al. Prevalence and treatment of fragility fractures in Spanish primary care: PREFRAOS study. *Arch Osteoporos*. 2022 Jul;17(1):93. doi: 10.1007/s11657-022-01124-7.
46. Yavropoulou MP, Makras P, Athanasakis K, Bournia VK, Mathioudakis K, Tsolakidis A, et al. Prevalence and patterns of anti-osteoporotic drug use based on 2019 real-world nationwide data in Greece. *Arch Osteoporos*. 2022 Jun;17(1):94. doi: 10.1007/s11657-022-01126-5.
47. Malle O, Borgström F, Fahrleitner-Pammer A, Svedbom A, Dimai SV, Dimai HP. Mind the gap: Incidence of osteoporosis treatment after an osteoporotic fracture - results of the Austrian branch of the International Costs and Utilities Related to Osteoporotic Fractures Study (ICUROS). *Bone*. 2021 Jan 1;142:115539. doi: 10.1016/j.bone.2019.115071.
48. McCloskey E, Rath J, Heijmans S, Blagden M, Cortet B, Czerwiński E, et al. The osteoporosis treatment gap in patients at risk of fracture in European primary care: A multi-country cross-sectional observational study. *Osteoporos Int*. 2021 Feb;32(2):251-259. doi: 10.1007/s00198-020-05557-z.
49. Roh Y, Lee E, Ahn J, Kim H, Gong H, Baek K, et al. Factors affecting willingness to get assessed and treated for osteoporosis. *Osteoporos Int*. 2019 Jul;30(7):1373-1379. doi: 10.1007/s00198-019-04952-5.
50. van der Velde R, Wyers C, Teesselink E, Geusens P, van den Bergh J, de Vries F, et al. Trends in oral anti-osteoporosis drug prescription in the United Kingdom between 1990 and 2012: Variation by age, sex, geographic location and ethnicity. *Bone*. 2017 Jan;94:54-60. doi: 10.1016/j.bone.2016.10.013.
51. Cunningham TD, Di Pace BS, Ullal J. Osteoporosis treatment disparities: A 6-year aggregate analysis from national survey data. *Osteoporos Int*. 2014;25(9):2119-2125. doi: 10.1007/s00198-014-2747-9.
52. Curtis JR, McClure LA, Delzell E, Howard VJ, Orwoll E, Saag KG, et al. Population-based fracture risk assessment and osteoporosis treatment disparities by race and gender. *J Gen Intern Med*. 2009;24(8):942-947. doi: 10.1007/s11606-009-1031-8.
53. Barton DW, Griffin DC, Carmouche JJ. Orthopedic surgeons' views on the osteoporosis care gap and potential solutions: Survey results. *J Orthop Surg Res*. 2019 Mar;14(1):110. doi: 10.1186/s13018-019-1103-3.
54. Beshyah SA, Al-Saleh Y, El-Hajj Fuleihan G. Management of osteoporosis in the Middle East and North Africa: A survey of physicians' perceptions and practices. *Arch Osteoporos*. 2019 Jun;14(1):38. doi: 10.1007/s11657-019-0609-y.
55. Dewan N, MacDermid JC, MacIntyre NJ, Grewal R. Therapist's practice patterns for subsequent fall/osteoporotic fracture prevention for patients with a distal radius fracture. *J Hand Ther*. 2019 Oct 1;32(4):665-672. doi: 10.1016/j.jht.2018.03.001.
56. Mendis A, Ganda K, Seibel M. Barriers to secondary fracture prevention in primary care. *Osteoporos Int*. 2017 Oct;28(10):2707-2714. doi: 10.1007/s00198-017-4131-z.
57. Duyvendak M, Nauntun M, van Roon E, Brouwers JRB. Doctors' beliefs and knowledge on corticosteroid-induced osteoporosis: Identifying barriers to improve prevention. *Int J Clin Pharm*. 2011 Jun;36(3):423-430. doi: 10.1111/j.1365-2710.2010.01185.x.
58. Feldstein AC, Nichols G, Orwoll E, Elmer PJ, Smith DH, Herson M, et al. The near absence of osteoporosis treatment in older men with fractures. *Osteoporos Int*. 2005 Aug;16(8):949-955. doi: 10.1007/s00198-005-1950-0.
59. Bennett MJ, Center JR, Perry L. Exploring barriers and opportunities to improve osteoporosis care across the acute-to-primary care interface: A qualitative study. *Osteoporos Int*. 2023;34(7):1459-1467. doi: 10.1007/s00198-023-06748-0.

60. Papaioannou A, McCloskey E, Bell A, Ngui D, Mehan U, Tan M, et al. Use of an electronic medical record dashboard to identify gaps in osteoporosis care. *Arch Osteoporos*. 2021;16(1):49. doi: 10.1007/s11657-021-00919-4.
61. Leslie W, Edwards B, Al-Azazi S, Yan L, Lix L, Czapkowski P, et al. Cancer patients with fractures are rarely assessed or treated for osteoporosis: A population-based study. *Osteoporos Int*. 2020 Feb;32(2):329-336. doi: 10.1007/s00198-020-05596-6.
62. Cheung M, Ho A, Wong S. Post-fracture care gap: A retrospective population-based analysis of Hong Kong from 2009 to 2012. *Hong Kong Med J*. 2018 Dec;24(6):553-561. doi: 10.12809/hkmj187227.
63. Wang M, Seibel MJ. Secondary fracture prevention in primary care: A narrative review. *Osteoporos Int*. 2024 Aug;35(8):1359-1376. doi: 10.1007/s00198-024-07036-1.
64. Giangregorio L, Papaioannou A, Cranney A, Zytaruk N, Adachi JD. Fragility fractures and the osteoporosis care gap: An international phenomenon. *Semin Arthritis Rheum*. 2006 Apr;35(5):293-305. doi: 10.1016/j.semarthrit.2005.11.001.
65. Little EA, Eccles MP. A systematic review of the effectiveness of interventions to improve post-fracture investigation and management of patients at risk of osteoporosis. *Implement Sci*. 2010 Dec;5(1):80. doi: 10.1186/1748-5908-5-80.
66. Vescini F, Chiodini I, Falchetti A, Palermo A, Salcuni AS, Bonadonna S, et al. Management of osteoporosis in men: A narrative review. *Int J Mol Sci*. 2021 Dec 20;22(24):13640. doi: 10.3390/ijms222413640.
67. Phelan JC, Link BG, Tehranifar P. Social conditions as fundamental causes of health inequalities: Theory, evidence, and policy implications. *J Health Soc Behav*. 2010 Mar;51(1_suppl):S28-S40. doi: 10.1177/0022146510383498.
68. Job C, Adenipekun B, Cleves A, Gill P, Samuriwo R. Health professionals' implicit bias of patients with low socioeconomic status (SES) and its effects on clinical decision-making: A scoping review. *BMJ Open*. 2024 Jul;14(7):e081723. doi: 10.1136/bmjopen-2023-081723.
69. Šter M, Švab I, Petek D, Živčec G. Adherence to osteoporosis treatment with once-monthly bisphosphonate. *Zdr Varst*. 2011;50(1): 45-54. doi: 10.2478/v10152-010-0027-4.