

Persistence with osteoporosis medication treatments among Chinese patients with osteoporosis: a retrospective cohort study

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INTRODUCTION

- Osteoporosis, a chronic progressive disease, affects 32.0% of individuals aged 65 and older, including 51.6% of women and 10.7% of men in China.
- Inadequate treatment continuation increases the risk of fracture and mortality, and increased healthcare costs.
- However, the persistence of medication treatment among patients with osteoporosis in China remains unknown.

AIMS

- This study aimed to evaluate treatment persistence among Chinese patients with osteoporosis.

METHODS

STUDY DESIGN

- The study was a retrospective, observational cohort study utilizing electronic medical records (EMRs) obtained from a provincial healthcare database in Eastern China.

POPULATION

- Inclusion criteria: Patients diagnosed with osteoporosis and initiating anti-osteoporosis medications between January 2019 and October 2024.
- Exclusion criteria: Patients had prior osteoporosis medication use, more than one index medication, a diagnosis of Paget's disease or malignancy, missing demographic information, or a follow-up duration of less than six months.

OUTCOMES

- Persistence was defined as the time from therapy initiation to discontinuation, allowing a 60-day permissible gap, which was evaluated across treatment windows (naïve vs. non-naïve) and by index medication class.

STATISTICAL ANALYSIS

- Kaplan–Meier and log-rank tests: assess persistence.
- Cox proportional hazards models: identify predictors of non-persistence, adjusting for demographic and clinical factors.

RESULTS

- A total of 232,755 patients with osteoporosis were identified. The median follow-up time was 434 days (IQR: 125-908). The mean age was 65.2 years (SD = 13.5), and 72.9% were female.
- The 12-month persistence of denosumab (37.8%) was higher than other treatments, compared with PTHa (parathyroid hormone analogue) (28.6%), parenteral bisphosphonates (20.7%), oral bisphosphonates (7.9%), menatetrenone (7.8%), active vitamin D analogs (7.5%), MHT (menopausal hormone therapy) (2.7%), TCM (traditional Chinese medicine) and calcitonin (2.1%), and SERMs (selective estrogen receptor modulators) (1.8%) (Table 1).

Table 1 persistence of each drug at 12months

| Drugs | Individuals of Persistence | Individuals of Non-Persistence | % (95% CI) |
|----------------------------|----------------------------|--------------------------------|------------------|
| Oral bisphosphonates | 1,302 | 22,004 | 7.9 (7.6-8.3) |
| Parenteral bisphosphonates | 832 | 3,221 | 20.7 (19.5-22.0) |
| Denosumab | 2,378 | 8,090 | 37.8 (36.9-38.8) |
| Calcitonin | 153 | 9,722 | 2.1 (1.8-2.4) |
| MHT | 2 | 109 | 2.7 (0.9-8.2) |
| SERMs | 2 | 108 | 1.8 (0.5-7.2) |
| PTHa | 1 | 27 | 28.6 (16.7-48.9) |
| Active vitamin D analogs | 9,060 | 141,210 | 7.5 (7.4-7.6) |
| Menatetrenone | 112 | 2,485 | 7.8 (6.8-9.0) |
| TCM | 350 | 22,551 | 2.1 (1.9-2.3) |

- Persistence with oral medications was higher in non-naïve treatment windows than naïve treatment windows, which was particularly apparent for oral bisphosphonates (7.92% persistence at for naïve treatments vs 12.3% for non-naïve treatments). By contrast, persistence with parenteral bisphosphonates and denosumab was higher in naïve treatment windows than in non-naïve treatment windows (Figure 1).

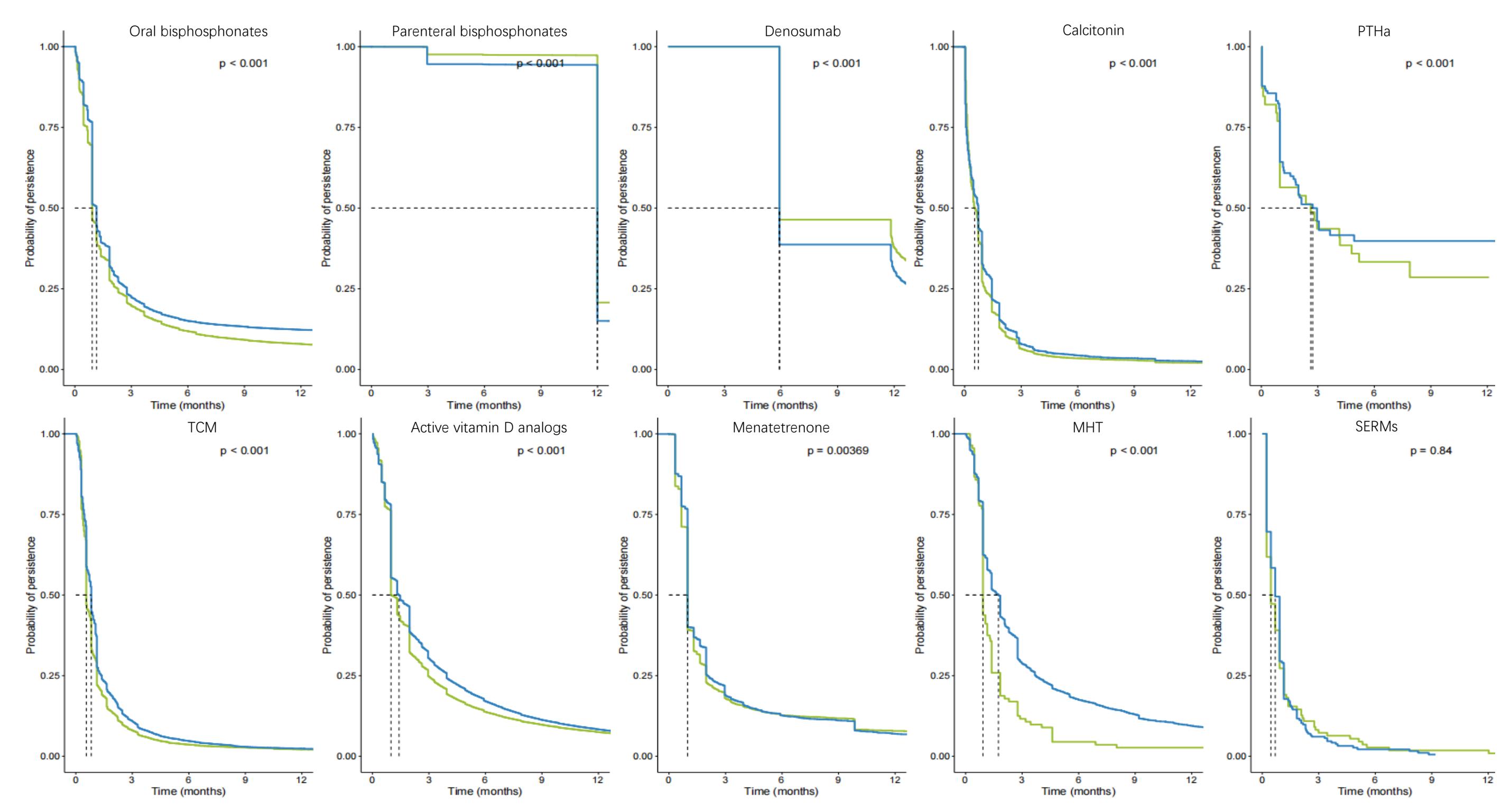


Figure 1 Kaplan–Meier analysis of discontinuation with medication class (naïve vs non-naïve treatments)

- Older age, female, osteoarthritis, and history of fractures were significantly associated with an increased risk of non-persistence.
- Patients with hypertension, diabetes, dyslipidemia, pneumonia, heart failure, ischemic stroke, rheumatoid arthritis, dementia, lupus, schizophrenia, or glaucoma were significantly less likely to discontinue treatment (Figure 2).

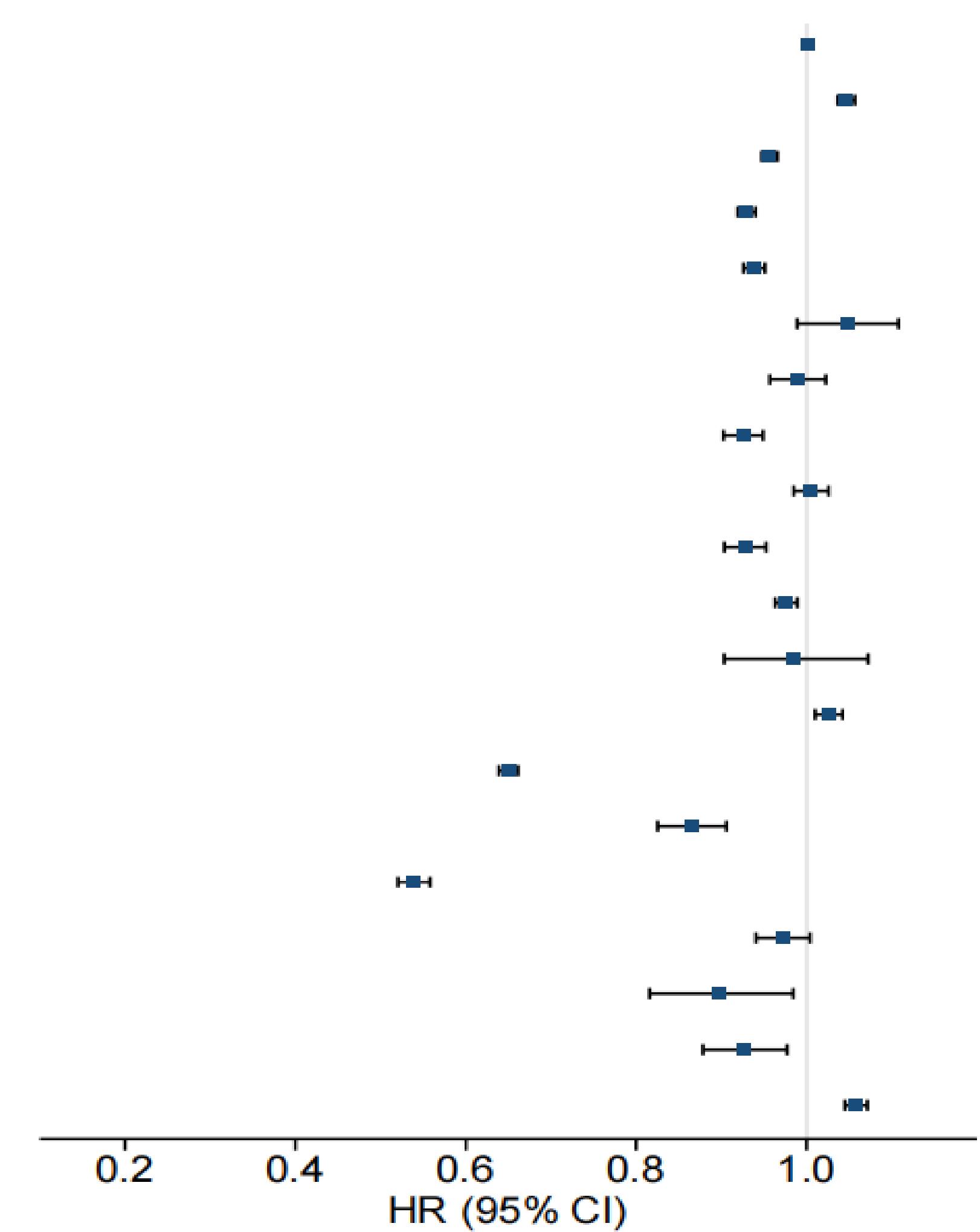


Figure 2 Hazards ratio and 95% confidence intervals for the associations between patient characteristics and non-persistence

CONCLUSIONS

Patients treated with denosumab exhibited higher treatment persistence compared to those receiving other AOMs, including oral bisphosphonates. Efforts to improve persistence are warranted, particularly among high-risk populations.

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