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Cost-effectiveness analysis of Chinese patent medicines for the treatment of postmenopausal osteoporosis in China

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Introduction: Evidence indicates that Chinese patent medicines can significantly increase bone mass in patients with osteoporosis and alleviate symptoms associated with low bone density. Although the therapeutic effects of these two drugs have been compared both directly and indirectly, no economic-related studies currently exist. Therefore, this study aims to assess the cost-effectiveness of Xianling Gubao Capsules compared to Jintiang Capsules and non-treatment for postmenopausal osteoporosis from the perspective of Chinese healthcare providers.

Methods: A Markov microsimulation model was employed to estimate the cost-effectiveness of the Xianling Gubao capsule and the Jintiang capsule in a hypothetical cohort of postmenopausal osteoporotic women aged 55 to 74 years with no prior history of fractures, over a treatment period of 6 months. Model parameters, including transition probabilities and costs, were derived from Chinese sources. Efficacy data for the treatments were obtained from two network meta-analyses. Outcomes were expressed as incremental costs per quality-adjusted life-year (QALY) gained. Sensitivity analyses were performed to ensure the robustness of the findings, with a cost-effectiveness threshold established at three times the Gross Domestic Product (GDP) per capita in China (\$38,223) per QALY.

Result: Compared to the control group that did not receive drug treatment, the preventive therapy using Chinese patent medicine significantly increased bone mineral density and reduced the probability of fractures across all age groups in the intervention group. The incremental cost-effectiveness ratios (ICERs) for the Jintiang capsule compared to the Xianling Gubao capsule ranged from \$11,955 per QALY at age 55 to \$9,711 per QALY at age 74, indicating that the cost-effectiveness of the Jintiang capsule improved consistently with age. Sensitivity analyses confirmed the robustness of the results across all parameter variations, with the annual cost of the Jintiang capsule identified as the most sensitive factor.

Conclusion: From the perspective of Chinese healthcare providers, preventive therapy using Chinese patent medicine, when compared to a control group that did not receive drug treatment, resulted in increased bone mineral density and a reduced probability of fractures across all age levels in the intervention group. Additionally, the Jintiang capsule appears to be a cost-effective treatment option for postmenopausal women with osteoporosis.

KEYWORDS

osteoporosis, Chinese patent medicines, postmenopausal, cost-effectiveness, Xianling Gubao capsules, Jintiang capsules

Introduction

Osteoporosis is a chronic, progressive skeletal disorder characterized by reduced bone mass, deterioration of bone microstructure, and increased fragility, which collectively elevate the risk of fragility fractures that can severely compromise patients' quality of life (1). Projections indicate that by 2040, nearly 319 million individuals worldwide will be at risk of osteoporotic fractures, with 55% of these cases expected to occur in Asia (2). According to the national census data from the National Bureau of Statistics of China at the end of 2021, the population aged 60 and above in China was 267.36 million, accounting for 18.9% of the total population. Among this group, the population aged 65 and above was 200.56 million, representing 14.2% (3). The Diagnosis and Treatment Guidelines for Primary Osteoporosis (2022) in China indicate that the prevalence of osteoporosis among women aged 50 and above is 32.1%, which is six times higher than that of men in the same age group (4). Furthermore, the prevalence of osteoporosis significantly increases among females aged 60 and above (1). Osteoporosis can lead to various types of fractures. A study conducted in 2015 estimated that the medical expenses for major osteoporotic fractures in China would reach as high as 11 billion, 20 billion, and 25 billion USD in 2015, 2035, and 2050, respectively (5). Therefore, it is essential to identify safe, effective, and economical treatment options.

Traditional osteoporosis medications include bisphosphonates, parathyroid hormone, selective estrogen receptor modulators, calcium supplements, estrogen replacement therapy, and calcilytics (6–8). While these drugs have demonstrated varying degrees of efficacy in the treatment of osteoporosis, patient compliance remains unsatisfactory due to the occurrence of adverse reactions (9–11). Hormone replacement therapy (HRT), a common treatment for osteoporosis, has been associated with an increased risk of cardiovascular disease and breast cancer (12). Additionally, long-term calcium intake, a routine preventive measure for osteoporosis, has also been linked to a heightened risk of myocardial infarction (13). Given the chronic nature of osteoporosis, it is crucial to balance the associated risks and benefits (14). In China, Chinese patent medicines (CPMs) are ready-made medications formulated in specific dosage forms based on prescriptions or standards guided by the principles of traditional Chinese medicine (15, 16). These medicines are widely utilized in the treatment of osteoporosis (15, 17). Evidence indicates that CPMs can significantly enhance bone mass in patients with osteoporosis and alleviate symptoms associated with low bone mass. The Xianling Gubao Capsule and Jintiang Capsule are two CPMs primarily recommended by various treatment guidelines (1, 18). Although there have been both direct and indirect comparisons of the therapeutic effects of these two medications, no economic evaluations have been conducted (19). Therefore, this study aims to compare the cost-effectiveness of Xianling Gubao Capsules and Jintiang Capsules, as well as non-treatment options for postmenopausal osteoporosis, from the perspective of Chinese healthcare providers, thereby addressing a significant gap in the existing literature.

Methods

Study design

This study utilized a 100,000 hypothetical individuals Markov microsimulation model to estimate the cost-effectiveness of Xianling Gubao Capsules and Jintiang Capsules in Chinese postmenopausal women, compared to no intervention. Each cycle lasts 1 year, during which each participant may experience a hip fracture, clinical vertebral fracture, or other types of fractures. Adopting the perspective of Chinese healthcare providers and extending the analysis to a lifetime horizon, the states are continuously updated until the patient's death. A consistent discount rate of 3% was applied to both costs and health outcomes to account for time preference. The analysis was conducted using TreeAge Pro (Healthcare Version) 2022, in accordance with the Consolidated Health Economic Reporting Standards (CHEERS), as detailed in [Supplementary Table 1 \(20\)](#).

The model meticulously simulated a cohort of Chinese postmenopausal women with no prior history of fragility fractures across various age groups: 55–59, 60–64, 65–69, and 70–74 years. The base case focused on individuals aged 55–59 years. In our model construction and parameterization, age-specific mean and standard deviation (SD) data of BMD shown in [Table 1](#) were derived from Wang et al. (21). The simulation incorporated a normal distribution derived from the mean and standard deviation (SD) values reported by Wang et al., based on the statistical assumption of a large-sample distribution. An initial BMD value was sampled from this normal distribution at the commencement of the Markov microsimulation and subsequently assigned randomly to individual participants. In accordance with established guidelines (1, 18), patients in all groups were administered calcium and activated vitamin D. The cohort was then assigned to receive either Xianling Gubao treatment (Xianling Gubao capsules 1.5 g b.i.d. for 6 months) or Jintiang treatment (Jintiang capsules 1.2 g t.i.d. for 6 months), while the no-intervention group was designated as the status quo.

Model structure

[Figure 1](#) illustrates the structure of the Markov model, which includes the states of no fracture, simple fracture, complex

TABLE 1 Mean BMDs, SDs of the femoral neck in different initial medicated ages of female (g/cm²).

| Age (years) | Number | Mean | SD | 95%CI |
|-------------|--------|------|------|--------------|
| 55–59 | 3,152 | 0.73 | 0.13 | (0.72, 0.73) |
| 60–64 | 3,155 | 0.66 | 0.12 | (0.65, 0.66) |
| 65–69 | 925 | 0.58 | 0.11 | (0.58, 0.59) |
| 70–74 | 137 | 0.55 | 0.13 | (0.53, 0.58) |

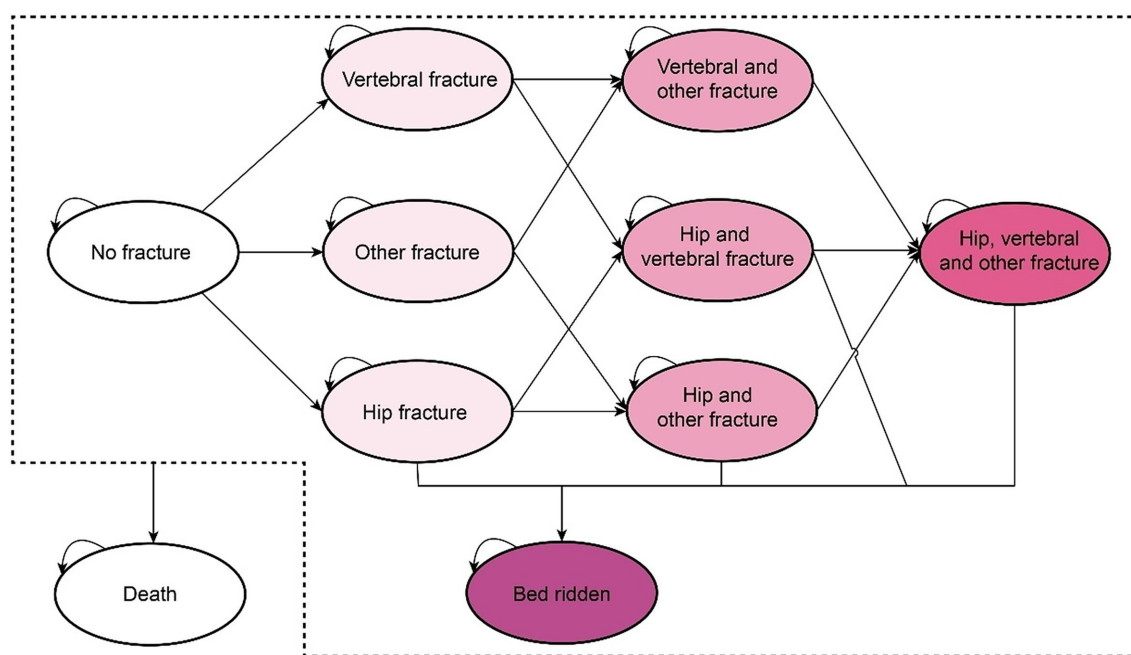


FIGURE 1

The Markov model structure for the disease progression of postmenopausal women with osteoporosis (omitting the arrow pointing to the death state).

fracture, bedridden due to hip fracture, and death from fracture or other causes. The simple fracture state indicates that individuals had experienced any hip, vertebral, or other types of fractures. The complex fracture state was defined as the occurrence of multiple simple fracture events. It was assumed that a certain proportion of individuals diagnosed with hip fractures may transition to the bedridden state without experiencing any additional fractures.

All patients entered the model in a “no fracture” healthy state. During each cycle, patients could experience a fracture, remain healthy, or die. Patients in the “fracture” state could either remain in the same fracture state if a refracture occurred, transition to another fracture state if a new fracture occurred, or move to the corresponding “post-fracture” state. Among these states, only patients with hip fractures may enter a bedridden state. For instance, patients who have experienced a vertebral fracture might subsequently experience another vertebral fracture or a hip fracture. Fractured patients could not return to the “no fracture” healthy state and would remain in the “post vertebral fracture” or “post hip fracture” state unless another fracture occurs or they entered the bedridden state. Ultimately, all patients were subject to the risk of death, and upon decease, they were transferred to the terminal death state.

Model parameters

All model parameters were sourced from China whenever possible to ensure their relevance to the local healthcare environment. In the absence of local data, information was synthesized from published literature through systematic literature searches. The input data utilized in the model is presented in the following section (Table 2).

Transition probabilities

Fracture risks

The transition probability of the fracture state was calculated based on age-specific and BMD-specific incidence rates of fragility fractures (Supplementary Table 2). Equations for the incidence of hip, vertebral, and other fractures associated with age and BMD were derived from published epidemiological data on the Chinese population (22–27). The fitted algorithms were evaluated using the R-squared statistic and adjusted for clinical plausibility. The probability of fracture was further modified based on the relative risk associated with a history of previous fractures, as individuals who have experienced any osteoporotic fracture are at an increased risk of subsequent fracture events. These values were obtained from a meta-analysis (28). Additionally, the probability of a bedridden state resulting from a hip fracture was extracted from a prior study conducted in Japan (29).

Mortality

Baseline age-specific mortality rates in the general population were extracted by sex and age from the seventh national population census in China (30). Excess mortality rates attributed to hip fractures were derived from published literature on Chinese women by multiplying the age-dependent risk ratio of mortality following a hip fracture (31).

Treatment

The efficacy data for Xianling Gubao capsules and Jintiang capsules were derived from a network meta-analysis of 22 randomized controlled trials that compared the effectiveness of various anti-osteoporotic agents (32). This study represents the largest analysis to date concerning the two aforementioned drugs and included a total of 2,016 postmenopausal women diagnosed with primary osteoporosis. Compared to placebo, Xianling Gubao capsules significantly enhanced

TABLE 2 Estimates of parameters used in the model.

| Parameter | Base-case value | Range | Distribution | Source |
|--|-----------------|-------------|--------------|--------|
| Relative risk of fracture in individuals with osteoporosis | | | | |
| RR of hip fractures with complications | | | | |
| History of previous fractures | 1.97 | 1.12–3.48 | Log-normal | (29) |
| RR of vertebral fractures with complications | | | | |
| History of previous fractures | 1.91 | 1.50–2.43 | Log-normal | (29) |
| RR of other fractures with complications | | | | |
| History of previous fractures | 1.91 | 1.50–2.43 | Log-normal | (29) |
| Probability of bedridden after hip fracture | 0.136 | 0.095–0.177 | Fixed | (32) |
| The therapeutic effect of drugs | | | | |
| Xianling Gubao capsule | | | | (59) |
| Increase in bone density (g/cm ²) | 0.05 | 0.01–0.08 | Beta | |
| Treatment time (year) | 0.5 | | Fixed | |
| Jintiange capsule | | | | (59) |
| Increase in bone density (g/cm ²) | 0.11 | 0.03–0.19 | Beta | |
| Treatment time (year) | 0.5 | | Fixed | |
| Cost (in 2022 China) | | | | |
| Annual Cost of Medication (US \$) | | | | |
| Xianling Gubao capsule | \$93.24 | ±20% | Triangular | (36) |
| Jintiange capsule | \$574.82 | ±20% | Triangular | |
| Annual medical expenses (US \$) | \$439.78 | ±20% | Triangular | |
| Cost of fracture treatment (US\$) | | | | |
| Hip fracture | \$7,379.82 | ±20% | Triangular | (60) |
| Vertebral fracture | \$1,361.12 | ±20% | Triangular | |
| Other fracture | \$1,758.30 | ±20% | Triangular | |
| Annual bedridden care expenses (US\$) | \$4,948.90 | ±20% | Triangular | |
| Health utility (QALY) | | | | |
| Age-Related Baseline Utility | | | | (41) |
| 55–59 | 0.88 | 0.862–0.897 | Beta | |
| 60–65 | 0.869 | 0.852–0.885 | Beta | |
| 66–70 | 0.827 | 0.802–0.851 | Beta | |
| 71–74 | 0.808 | 0.770–0.846 | Beta | |
| Disutility resulting from hip fractures | | | | (42) |
| First year | ×0.776 | 0.720–0.844 | Beta | |
| subsequent years | ×0.855 | 0.800–0.909 | Beta | |
| Disutility resulting from vertebral fracture | | | | |
| First year | ×0.724 | 0.667–0.779 | Beta | |
| subsequent years | ×0.868 | 0.827–0.922 | Beta | |
| Disutility resulting from other fracture | | | | (39) |
| First year | ×0.910 | 0.880–0.940 | Beta | |
| subsequent years | ×1 | | | |
| Utility of Bedridden State | 0.192 | | Fixed | (41) |
| Discount rate | | | | |
| Cost | 0.03 | 0–0.05 | | (35) |
| QALYs | 0.03 | 0–0.05 | | (35) |

lumbar and femoral neck BMD (Mean Difference, MD = 0.13, 95% CI [0.03, 0.22], MD = 0.17, 95% CI [0.06, 0.29]). In contrast, Jintiange capsules significantly improved femoral BMD compared to placebo (MD = 0.11, 95% CI [0.03–0.19]).

In addition, treatment-related adverse events were excluded from the model because previous studies did not find any

statistically significant differences between patients treated with Xianling Gubao capsules and those treated with Jintiange capsules (33, 34). Therefore, we assumed that treatment-related adverse events had a negligible impact on the costs and outcomes for patients receiving either Xianling Gubao capsules or Jintiange capsules.

Health resource use and costs

According to the Chinese Pharmacoeconomic Guidelines (35), the cost evaluation was conducted from the perspective of Chinese healthcare providers. Direct medical costs encompass the expenses associated with treatment regimens resulting from fracture events, direct medical expenses for each health state, and other medical expenditures. The cost data for this section were primarily obtained from a multi-center survey in China (36). The costs of Xianling Gubao capsules and Jintiang capsules were calculated based on the market share of generic drugs and their branded counterparts in China, utilizing official databases from China's National Medical Products Administration (NMPA) (37). The estimated annual cost of Xianling Gubao capsules was USD 93.24 (3 capsules, twice a day), while the cost for Jintiang capsules was USD 574.82 (3 capsules, three times a day). All related costs were adjusted to 2022 Chinese Yuan (CNY) using the Consumer Price Index (CPI). For reference, the average exchange rate in 2022 was USD 1 = CNY 6.7321.

Utilities

Baseline utility data were extracted from published literature to provide reference value for the decision analysis model (38). The utility values were derived from a Chinese large population using EQ-5D. The disutility multiplier associated with post-fracture in the first and subsequent years was derived from meta-analysis (39, 40). Utility for bed-ridden state was collected from a study of Chinese patients provided with nursing care (41).

Statistical analysis

In base case analysis, using first-order Monte Carlo simulation, total costs and QALYs for each treatment with Xianling Gubao capsules, Jintiang capsules, and no treatment were estimated at different starting ages of 55, 60, 65, and 70 years. To estimate the cost-effectiveness of Xianling Gubao capsules and Jintiang capsules, an incremental cost-effectiveness ratio (ICER) was calculated by dividing an incremental cost by an incremental quality-adjusted life-year (QALY) to obtain the cost per QALY gained. To explore key drivers of parameters, deterministic sensitivity analyses were conducted, and parameters assessed and their ranges are shown in Table 2. Probabilistic sensitivity analyses were conducted by a second-order Monte Carlo simulation with 1,000 iterations and selecting the assigned parameters distributed randomly (shown in Table 2). Following the analyses, cost-effectiveness acceptability curves were illustrated to determine the probability of being cost-effective for each strategy based on an assumed willingness-to-pay (WTP) threshold of three-time GDP per capita in China (\$38,223) per QALY gained.

Results

Model validation

This model was validated by comparing the age-specific incidence of hip fractures and clinical vertebral fractures per year with estimates from published epidemiological surveys (shown in Figure 2) (23, 42).

Base-case analyses

Table 3 presents the costs, QALYs, and ICERs for Xianling Gubao or Jintiang compared with no treatment at the starting ages of 55, 60, 65, and 70 years. Compared with the control group without drug treatment, the preventive treatment with Chinese patent medicine increased bone mineral density and reduced fracture probability at all age levels in the intervention group. Model simulation results based on females aged 55–59 showed that the use of Xianling Gubao capsules reduced hip fracture incidence by 8.20% and vertebral fracture incidence by 12.03%, with an increase in total per capita cost of \$119.21 and an increase in quality-adjusted life years (QALYs) by 0.0379. The use of Jintiang capsule reduced hip fracture rates by 18.33% and vertebral fracture rates by 30.05%, with an increase in total per capita cost of \$956.38 and an increase in QALYs by 0.08. Under the willingness-to-pay (WTP) threshold of three times China's per capita GDP, the use of traditional Chinese medicine for preventive treatment had a cost-effective advantage. Moreover, compared with the Xianling Gubao capsule, the Jintiang capsule was cost-effective (ICER: \$11,955/QALY).

For the population aged 60–74, the use of traditional Chinese medicine for preventive treatment still had a cost-effective advantage. Compared with the Xianling Gubao capsule, the Jintiang capsule was cost-effective (ICER: \$9858–10731/QALY) under the condition of WTP being 3 times China's per capita GDP.

One-way sensitivity analyses

One-way sensitivity analyses comparing the Xianling Gubao capsule with the Jintiang capsule or no treatment indicated that the ICERs were most sensitive to the discount rate, the loss of utility due to fractures, the first-year BMD of the study population, and drug acquisition costs (Figure 3). The graph demonstrates that the cost-effectiveness results of CPM treatment are relatively stable and unaffected by changes in these parameters. It is cost-effective to initiate preventive treatment with CPMs for individuals with below-average BMD who experience a significant impact on their quality of life due to fractures.

Probabilistic sensitivity analyses

At a WTP threshold of \$3,000 (approximately 0.24 times GDP per capita), the use of CPMs for preventive treatment in the female population aged 55–59 was deemed cost-effective. When the WTP exceeded \$24,000 (about 1.88 times per capita GDP), the economic benefits of using the Jintiang capsule became more pronounced compared to the Xianling Gubao capsule. Under the WTP conditions established in the study, the probabilities of the Xianling Gubao capsule and the Jintiang capsule being cost-effective were 31 and 49%, respectively (Figure 4).

Discussion

Previous systematic reviews had indicated a relative scarcity of cost-effectiveness research concerning CPMs (43). In this study, we constructed a Markov model based on life status to simulate the

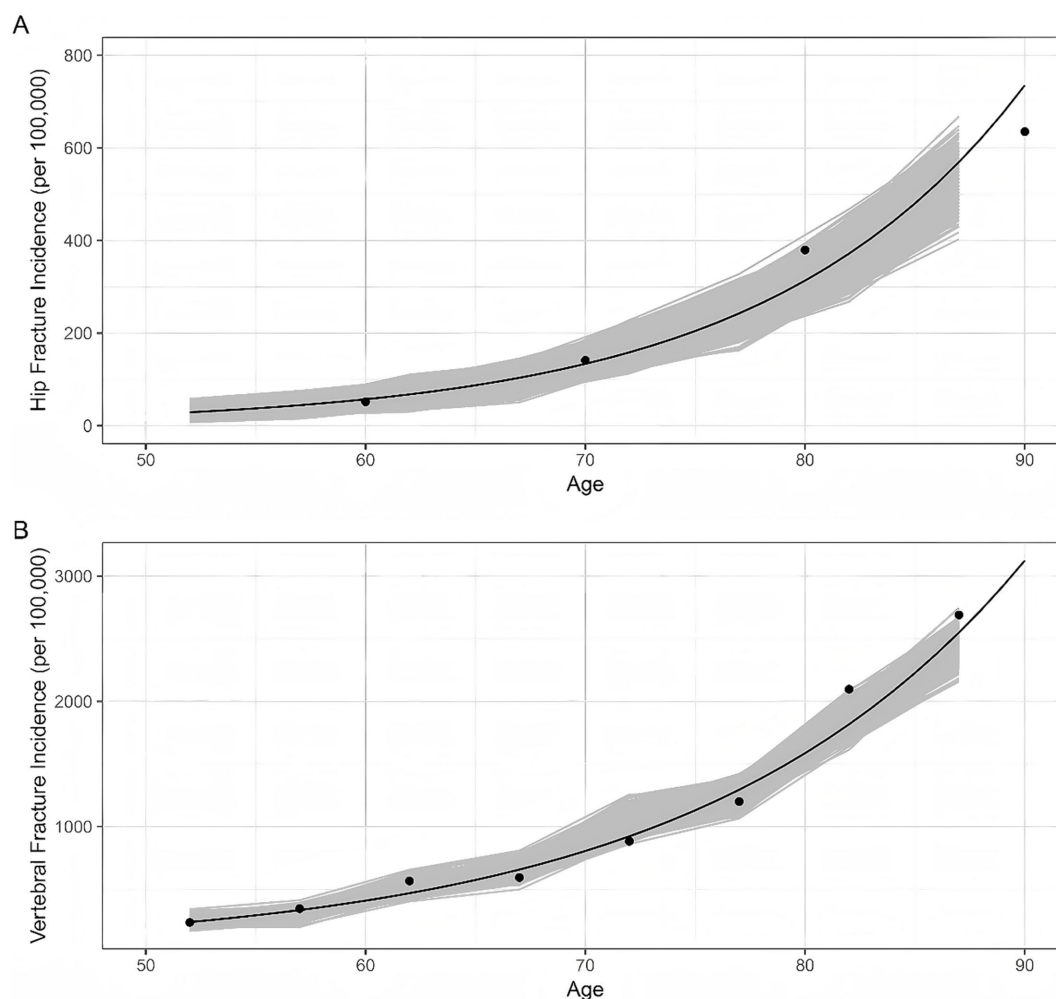


FIGURE 2

Model validation results. Dots represent the data reported by epidemiological surveys in China. Lines represent the trend curves fitted by the data above-mentioned. Shaded areas represent the model outputs of 1,000 simulations. (A) Hip fracture incidence; (B) Vertebral fracture incidence.

long-term effects of CPMs on the treatment outcomes of menopausal women with osteoporosis. Our base analysis suggests that, at a WTP threshold of \$3,000 per QALY, the use of CPMs for preventive treatment in postmenopausal women aged 55 to 59 was cost-effective. Furthermore, at a WTP threshold equivalent to three times China's per capita GDP, the combined use of Jintiang capsules was cost-effective across all age groups. This study provided reference values for future long-term economic evaluations of CPMs for osteoporosis in postmenopausal women.

The primary treatment approaches for bone loss diseases such as osteoporosis are HRT and bisphosphonates. Continuous HRT was associated with a high risk of breast cancer and endometrial cancer, as well as coronary artery issues and other cardiac disorders, while bisphosphonates could lead to osteonecrosis of the jawbone and skeletal system (44–46). Due to these adverse effects, the clinical use of HRT and bisphosphonates is limited. Therefore, new treatment strategies were needed to develop osteoporosis treatments that are less likely to cause adverse reactions to some extent (47). CPMs had gained popularity due to their minimal adverse effects while effectively treating various ailments. Traditional Chinese Medicine (TCM) had been utilized to address a range of orthopedic conditions,

particularly osteoporosis, fractures, and rheumatism, with notable success (48, 49). Xianling Gubao and Jintiang treatments played significant roles in managing osteoporosis with CPMs, making them the preferred choice for treating postmenopausal osteoporosis according to clinical application guidelines for Chinese patent medicines. Xianling Gubao capsules consist entirely of CPMs (50), which help regulate the balance of serum calcium and phosphorus deposition, enhance levels of vitamin D3 and alkaline phosphatase (ALP), and improve bone mineral density through the synergistic effects of various natural herbs targeting multiple pathways (51, 52). The primary component of Jintiang capsules was artificial tiger bone powder. The physical and chemical properties, as well as the pharmacological effects were consistent. Researchers studying the serum of patients with osteoporosis found that Jintiang capsules inhibited the κ B inhibitor signaling pathway by downregulating the overexpression of osteopontin, ultimately reducing MMP-3 expression. This action may strengthen the kidneys and bones, alleviate inflammation and pain, and combat osteoporosis (53). Furthermore, recent studies have identified Jintiang capsules as the first CPM that could effectively improve primary osteoporosis and enhance muscle strength (54). While the therapeutic potential of

TABLE 3 Cost-effectiveness of Xianling Gubao compared with Jintiange or no treatment at the starting ages of 55, 60, 65, and 70 years.

| Treatment Strategy | Probability of patients experiencing a fracture within the study period | | Cost, \$ | QALYs | ICER, \$/QALY |
|--------------------------|---|--------------------|----------|---------|---------------|
| | Hip fracture | Vertebral fracture | | | |
| 55–59 years | | | | | |
| No Treatment | 6.71% | 21.03% | 7174.94 | 13.0513 | Ref |
| Xianling Gubao Treatment | 6.16% | 18.50% | 7294.16 | 13.0892 | 3,147 |
| Jintiange Treatment | 5.48% | 15.47% | 8131.32 | 13.1313 | 11,955 |
| 60–64 years | | | | | |
| No Treatment | 7.30% | 23.58% | 6791.35 | 11.4976 | Ref |
| Xianling Gubao Treatment | 6.90% | 20.81% | 6883.21 | 11.5373 | 2,313 |
| Jintiange Treatment | 5.94% | 17.40% | 7681.96 | 11.5879 | 9,858 |
| 65–69 years | | | | | |
| No Treatment | 6.60% | 20.40% | 6075.22 | 9.9942 | Ref |
| Xianling Gubao Treatment | 6.09% | 18.04% | 6184.11 | 10.052 | 1886 |
| Jintiange Treatment | 5.29% | 14.91% | 6978.64 | 10.0784 | 10,731 |
| 70–74 years | | | | | |
| No Treatment | 7.46% | 23.00% | 5572.26 | 8.0271 | Ref |
| Xianling Gubao Treatment | 6.49% | 20.63% | 5646.1 | 8.0875 | 1,221 |
| Jintiange Treatment | 5.86% | 17.16% | 6393.42 | 8.1116 | 9,711 |

CPMs in managing osteoporosis is evident, there is a lack of economic studies to support their widespread adoption. Therefore, further economic evaluations are necessary to inform clinical and policy decisions regarding the integration of CPMs into standard osteoporosis treatment protocols.

The choice of research perspective in economic evaluation determines the measurement range of cost. Our results demonstrated that the use of CPMs preventive treatment appears to be a cost-effective treatment option for postmenopausal osteoporotic women at the starting age of 55 from the perspective of Chinese healthcare providers. Our results also revealed that the cost-effectiveness of Jintiange Treatment improved with an increase in the starting age. Our findings were consistent with previous economic evaluations in which the Jintiange treatment was generally cost-effective compared with the Xianling Gubao treatment (19, 32, 55). We did not find any studies that were different from this conclusion, which may be caused by the lack of relevant economic studies. However, pharmacoeconomic assessments were based on data from clinical trials or real-world data, and further CPMs clinical trials and real-world studies were recommended.

Currently, the pharmacoeconomic studies on the treatment of osteoporosis in menopausal women published both at home and abroad mainly concentrate on RANKL inhibitors, bisphosphonates, and other Western medicine preparations. There have been relatively few studies on CPMs for treating osteoporosis in menopausal women. Only Lai Fuchong et al. (34) carried out an economic analysis of Xianling Gubao capsules and Jintiange capsules in the treatment of type-I osteoporosis. The results indicated that the cost-effectiveness ratio of alendronate combined with Xianling Gubao capsule was the lowest, followed by alendronate combined with Jintiange capsule. Nevertheless, the limitations of this study include the lack of ICERs, which are essential for definitively determining the most cost-effective treatment strategy. Furthermore, the data were gathered from a restricted patient sample across two hospitals, without taking into account potential confounding factors. This may have an impact on

the generalizability of the results. The current study aims to fill this gap by presenting a comparative economic assessment of Xianling Gubao Capsules and Jintiange Capsules in postmenopausal women of various age groups at the start of treatment. Such an analysis is crucial for promoting the rational clinical use of CPMs in osteoporosis management, providing valuable perspectives for healthcare providers and policymakers when considering the incorporation of these treatments into standard care guidelines.

Although there were studies on the synergistic effect of CPM combined with conventional Western medicine in the treatment of osteoporosis (56), unfortunately, we had not found any direct comparative studies on the treatment of osteoporosis in postmenopausal women with CPM and Western medicine, including safety, efficacy and economy. Although no direct research evidence comparing CPM and Western medicine in the treatment of osteoporosis in postmenopausal women has been found so far, our team had fully realized the importance of this research gap for clinical practice and health policy-making. Currently, we were actively promoting the preparatory work for subsequent observational and real-world studies to provide solid methodological support for future research design. We were confident that through standardized research implementation and rigorous data analysis, we will gradually fill this research gap and provide more comprehensive evidence-based basis for the optimization of treatment decisions for osteoporosis in postmenopausal women.

As with any modeling research, our analyses had certain limitations. First, due to the absence of epidemiological data fully consistent with the study population, the efficacy parameter of the model transition probability involved a series of assumptions and corrections. This might deviate to some extent from the actual disease outcomes. Second, owing to the lack of relevant research, our data on the efficacy of drug treatment were solely obtained from a network meta-analysis. Although this was the largest study related to both drugs, the data dated back to 2016, and the study endpoint was bone mineral density (BMD), which

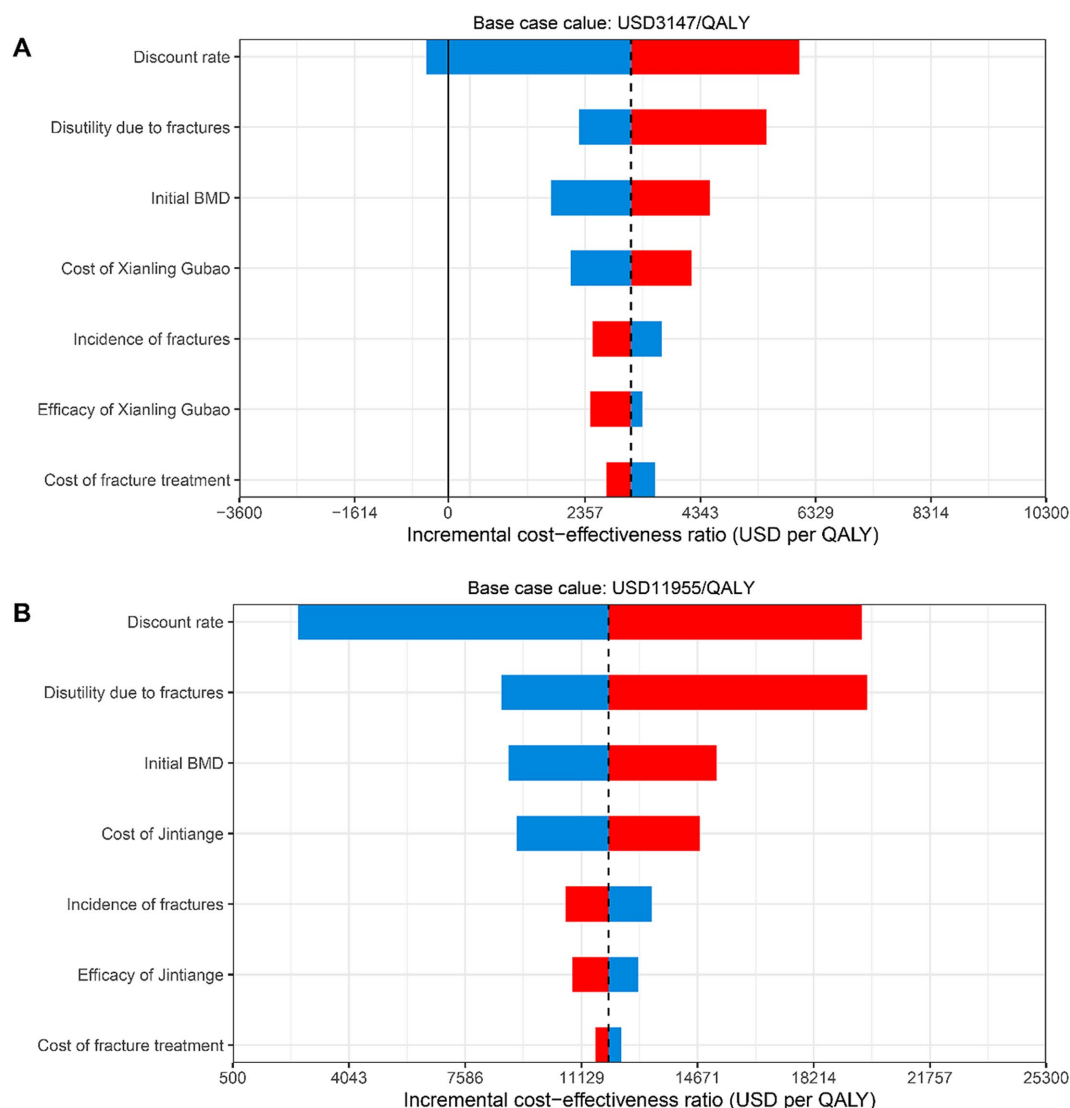


FIGURE 3

Tornado diagrams for one-way sensitivity analyses on the incremental cost-effectiveness ratio of Xianling Gubao capsule compared with Jintiang capsule or no treatment. Blue and red represent the ICER results in lower limit and upper limit values of parameters used, respectively. (A) Xianling Gubao; (B) Jintiang.

was used to predict the incidence of fractures subsequently. This could introduce bias to our results. Third, to maintain model parsimony, we did not incorporate adverse events. However, serious adverse events caused by Xianling Gubao capsules and Jin Tiange in the treatment of osteoporosis are considered rare (33, 34). Thus, they were unlikely to affect the results of our cost-effectiveness analyses. Similarly, this was a commonly adopted assumption in previous economic evaluations (57, 58). The impact of treatment-related adverse events on costs and outcomes can be regarded as minimal. Nevertheless, if data on adverse events become available, they should be integrated into the model.

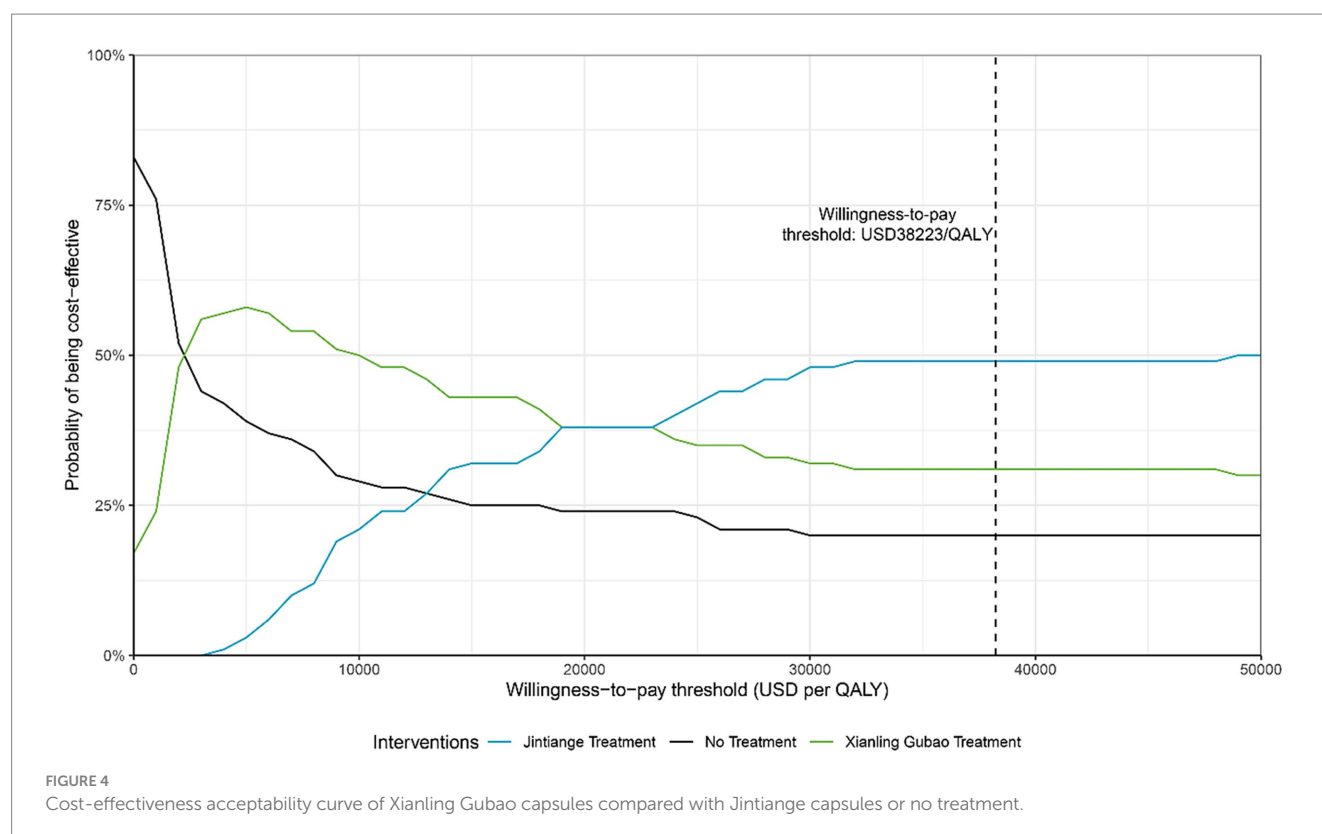
Conclusion

From the perspective of the Chinese healthcare providers, compared with the control group without drug therapy, the

preventive treatment with Chinese patent medicine increased bone mineral density and reduced fracture probability at all age levels in the intervention group, and Jintiang capsule appears to be a cost-effective treatment choice for postmenopausal osteoporotic women. This study provides valuable information to both clinical practitioners and decision-makers in ensuring the rational use of Chinese patent medicine, especially in the face of the growing clinical and economic burden of osteoporotic fractures in China.

Data availability statement

The original contributions presented in the study are included in the article/Supplementary material, further inquiries can be directed to the corresponding author.



Author contributions

CW: Writing – original draft, Writing – review & editing. XL: Data curation, Writing – original draft. JL: Investigation, Writing – review & editing. YZ: Methodology, Writing – review & editing. RY: Writing – review & editing, Conceptualization, Supervision.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

Generative AI statement

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Supplementary material

The Supplementary material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fpubh.2025.1596676/full#supplementary-material>

References

- Chinese Society of Osteoporosis and Bone Mineral Research. Guidelines for the diagnosis and treatment of primary osteoporosis (2022). *Chin Gen Pract.* (2023) 26:1671–91. doi: 10.12114/j.issn.1007-9572.2023.0121
- Oden A, McCloskey EV, Kanis JA, Harvey NC, Johansson H. Burden of high fracture probability worldwide: secular increases 2010–2040. *Osteoporos Int.* (2015) 26:2243–8. doi: 10.1007/s00198-015-3154-6
- National Bureau of Statistics. (2021). Bulletin of the seventh national population census. Available online: https://www.stats.gov.cn/zt_18555/zdtjgz/zgrkpc/dqcrkpc/ggl/202302/t0230215_1903997. (accessed on 2024-11-20).
- Ruihua D, Xiaoming X, Jixiang Z, Ning L. Best evidence summary of the exercise management in women with postmenopausal osteoporosis. *Chin Nurs Res.* (2022) 36:640–4. doi: 10.12102/j.issn.1009-6493.2022.04.013
- Si L, Winzenberg TM, Jiang Q, Chen M, Palmer AJ. Projection of osteoporosis-related fractures and costs in China: 2010–2050. *Osteoporos Int.* (2015) 26:1929–37. doi: 10.1007/s00198-015-3093-2
- Burns RB, Rosen H, Berry S, Smetana GW. How would you manage this patient with osteoporosis? Grand rounds discussion from Beth Israel Deaconess Medical Center. *Ann Intern Med.* (2018) 168:801–8. doi: 10.7326/M18-0950
- Chang B, Quan Q, Li Y, Qiu H, Peng J, Gu Y. Treatment of osteoporosis, with a focus on 2 monoclonal antibodies. *Med Sci Monit.* (2018) 24:8758–66. doi: 10.12659/MSM.912309
- Vidal M, Thibodaux RJ, Neira LFV, Messina OD. Osteoporosis: a clinical and pharmacological update. *Clin Rheumatol.* (2019) 38:385–95. doi: 10.1007/s10067-018-4370-1
- Netelenbos JC, Geusens PP, Ypma G, Buijs SJ. Adherence and profile of non-persistence in patients treated for osteoporosis—a large-scale, long-term retrospective study in the Netherlands. *Osteoporos Int.* (2011) 22:1537–46. doi: 10.1007/s00198-010-1372-5
- Hilgsmann M, Bours SP, Boonen A. A review of patient preferences for osteoporosis drug treatment. *Curr Rheumatol Rep.* (2015) 17:61. doi: 10.1007/s11926-015-0533-0
- Tadrous M, Mamdani MM, Juurlink DN, Krahn MD, Lévesque LE, Cadarette SM. Comparative gastrointestinal safety of bisphosphonates in primary osteoporosis: a network meta-analysis-reply to Pazianas and Abrahamsen. *Osteoporos Int.* (2014) 25:2671–2. doi: 10.1007/s00198-014-2789-z
- Rossouw JE, Anderson GL, Prentice RL, LaCroix AZ, Kooperberg C, Stefanick ML, et al. Risks and benefits of estrogen plus progestin in healthy postmenopausal women: principal results from the Women's Health Initiative randomized controlled trial. *JAMA.* (2002) 288:321–33. doi: 10.1001/jama.288.3.321
- Cipriani C, Pepe J, Minisola S, Lewiecki EM. Adverse effects of media reports on the treatment of osteoporosis. *J Endocrinol Invest.* (2018) 41:1359–64. doi: 10.1007/s40618-018-0898-9
- Reginster JY, Pelousse F, Bruyere O. Safety concerns with the long-term management of osteoporosis. *Expert Opin Drug Saf.* (2013) 12:507–22. doi: 10.1517/14740338.2013.793669
- Cui X, Wang S, Cao H, Guo H, Li Y, Xu F, et al. A review: the bioactivities and pharmacological applications of Polygonatum sibiricum polysaccharides. *Molecules.* (2018) 23:23. doi: 10.3390/molecules23051170
- Zhang ND, Han T, Huang BK, Rahman K, Jiang YP, Xu HT, et al. Traditional Chinese medicine formulas for the treatment of osteoporosis: implication for antiosteoporotic drug discovery. *J Ethnopharmacol.* (2016) 189:61–80. doi: 10.1016/j.jep.2016.05.025
- Zhao H, Zhao N, Zheng P, Xu X, Liu M, Luo D, et al. Prevention and treatment of osteoporosis using Chinese medicinal plants: special emphasis on mechanisms of immune modulation. *J Immunol Res.* (2018) 2018:1–11. doi: 10.1155/2018/6345857
- National Administration of Traditional Chinese Medicine (NATCM). Clinical application guidance for treating osteoporosis by Chinese patent medicine(2021). *Chin J Integr Tradit West Med.* (2022) 42:393–404. doi: 10.7661/j.cjim.20220204.063
- Zhao J, Zeng L, Wu M, Huang H, Liang G, Yang W, et al. Efficacy of Chinese patent medicine for primary osteoporosis: a network meta-analysis. *Complement Ther Clin Pract.* (2021) 44:101419. doi: 10.1016/j.ctcp.2021.101419
- Husereau D, Drummond M, Petrou S, Carswell C, Moher D, Greenberg D, et al. Consolidated health economic evaluation reporting standards (CHEERS) statement. *Int J Technol Assess Health Care.* (2013) 29:117–22. doi: 10.1017/S0266462313000160
- Wang L, Yu W, Yin X, Cui L, Tang S, Jiang N, et al. Prevalence of osteoporosis and fracture in China: the China osteoporosis prevalence study. *JAMA Netw Open.* (2021) 4:e2121106. doi: 10.1001/jamanetworkopen.2021.21106
- Zhang CG, Feng JN, Wang SF, Gao P, Xu L, Zhu JX, et al. Incidence of and trends in hip fracture among adults in urban China: a nationwide retrospective cohort study. *PLoS Med.* (2020) 17:e1003180. doi: 10.1371/journal.pmed.1003180
- Bow CH, Cheung E, Cheung CL, Xiao SM, Loong C, Soong C, et al. Ethnic difference of clinical vertebral fracture risk. *Osteoporos Int.* (2012) 23:879–85. doi: 10.1007/s00198-011-1627-9
- Lau EM, Woo J, Leung PC, Swaminathan R. Low bone mineral density, grip strength and skinfold thickness are important risk factors for hip fracture in Hong Kong Chinese. *Osteoporos Int.* (1993) 3:66–70. doi: 10.1007/BF01623375
- Kwok AW, Gong JS, Wang YX, Leung JC, Kwok T, Griffith JF, et al. Prevalence and risk factors of radiographic vertebral fractures in elderly Chinese men and women: results of Mr. OS (Hong Kong) and Ms. OS (Hong Kong) studies. *Osteoporos Int.* (2013) 24:877–85. doi: 10.1007/s00198-012-2040-8
- Tsang SW, Kung AW, Kanis JA, Johansson H, Oden A. Ten-year fracture probability in Hong Kong southern Chinese according to age and BMD femoral neck T-scores. *Osteoporos Int.* (2009) 20:1939–45. doi: 10.1007/s00198-009-0906-1
- Kwok TCY, Su Y, Khoo CC, Leung J, Kwok A, Orwoll E, et al. Predictors of non-vertebral fracture in older Chinese males and females: Mr. OS and Ms. OS (Hong Kong). *J Bone Miner Metab.* (2017) 35:330–7. doi: 10.1007/s00774-016-0761-z
- Kanis JA, Johnell O, De Laet C, Johansson H, Oden A, Delmas P, et al. A meta-analysis of previous fracture and subsequent fracture risk. *Bone.* (2004) 35:375–82. doi: 10.1016/j.bone.2004.03.024
- Yoshimura M, Moriwaki K, Noto S, Takiguchi T. A model-based cost-effectiveness analysis of osteoporosis screening and treatment strategy for postmenopausal Japanese women. *Osteoporos Int.* (2017) 28:643–52. doi: 10.1007/s00198-016-3782-5
- Office of the Leading Group of the State Council for the Seventh National Population Census. China population census yearbook (2020). Available online at: <http://www.stats.gov.cn/sj/pcsj/rkpc/7rp/zk/indexch.htm> (accessed on 2024-11-5).
- Koh GCH, Tai BC, Ang LW, Heng D, Yuan JM, Koh WP. All-cause and cause-specific mortality after hip fracture among Chinese women and men the Singapore Chinese health study. *Osteoporos Int.* (2013) 24:1981–9. doi: 10.1007/s00198-012-2183-7
- Luo MH, Zhao JL, Xu NJ, Xiao X, Feng WX, Li ZP, et al. Comparative efficacy of Xianling Gubao capsules in improving bone mineral density in postmenopausal osteoporosis: a network meta-analysis. *Front Endocrinol.* (2022) 13:839885. doi: 10.3389/fendo.2022.839885
- Zhongzhou Z, Dongsheng X, Xiujun J, Yongsheng S. Network meta-analysis of six Chinese patent medicines in treatment of knee osteoarthritis. *Pharm Clin Res.* (2024) 32:338–42. doi: 10.13664/j.cnki.pcr.2024.04.023
- Fuchong L. Clinical study of Xianlinggubao and Jintianghe capsules in treating type I Osteoporosischinese abstract. *Master.* (2020). doi: 10.27460/d.cnki.gzyzc.2020.000079
- Gueno L, Shanlian H, JiuHong W. China guidelines for pharmacoeconomic evaluations. *China J Pharm Econ.* (2020) 3–9. <https://www.cpa.org.cn/cpadmn/attached/file/20201203/1606977380634185.pdf>
- Qu B, Ma Y, Yan M, Wu HH, Fan L, Liao DF, et al. The economic burden of fracture patients with osteoporosis in western China. *Osteoporos Int.* (2014) 25:1853–60. doi: 10.1007/s00198-014-2699-0
- National Medical Products Administration. Available online at: <https://www.nmpa.gov.cn/yaopin/index.html> (accessed on 2024-10-30).
- Si L, Shi L, Chen MS, Palmer AJ. Establishing benchmark EQ-5D-3L population health state utilities and identifying their correlates in Gansu Province, China. *Qual Life Res.* (2017) 26:3049–58. doi: 10.1007/s11136-017-1614-5
- Hilgsmann M, Ethgen O, Richy F, Reginster JY. Utility values associated with osteoporotic fracture: a systematic review of the literature. *Calcif Tissue Int.* (2008) 82:288–92. doi: 10.1007/s00223-008-9117-6
- Si L, Winzenberg TM, de Graaff B, Palmer AJ. A systematic review and meta-analysis of utility-based quality of life for osteoporosis-related conditions. *Osteoporos Int.* (2014) 25:1987–97. doi: 10.1007/s00198-014-2636-2
- Liu N, Zeng LX, Li Z, Wang JE. Health-related quality of life and long-term care needs among elderly individuals living alone: a cross-sectional study in rural areas of Shaanxi Province, China. *Bmc Public Health.* (2013) 13:13. doi: 10.1186/1471-2458-13-313
- Zheng XQ, Xu L, Huang J, Zhang CG, Yuan WQ, Sun CG, et al. Incidence and cost of vertebral fracture in urban China: a 5-year population-based cohort study. *Int J Surg.* (2023) 109:1910–8. doi: 10.1097/JIS.0000000000000411
- Zhang F, Kong LL, Zhang YY, Li SC. Evaluation of impact on health-related quality of life and cost effectiveness of traditional Chinese medicine: a systematic review of randomized clinical trials. *J Altern Complement Med.* (2012) 18:1108–20. doi: 10.1089/acm.2011.0315
- Dinger J, Bardenheuer K, Heinemann K. Drospirenone plus estradiol and the risk of serious cardiovascular events in postmenopausal women. *Climacteric.* (2016) 19:349–56. doi: 10.1080/13697137.2016.1183624
- He JB, Chen MH, Lin DK. New insights into the tonifying kidney-yin herbs and formulas for the treatment of osteoporosis. *Arch Osteoporos.* (2017) 12:2–13. doi: 10.1007/s11657-016-0301-4
- Spivakovsky S. Treatment for bisphosphonate-related osteonecrosis of the jaw. *Evid Based Dent.* (2017) 18:56. doi: 10.1038/sj.ebd.6401243
- Lungu AE, Lazar MA, Tanea A, Rotaru H, Roman RC, Badea ME. Observational study of the bisphosphonate-related osteonecrosis of jaws. *Clujul Med.* (2018) 91:209–15. doi: 10.15386/cjmed-838

48. Mukwaya E, Xu F, Wong MS, Zhang Y. Chinese herbal medicine for bone health. *Pharm Biol.* (2014) 52:1223–8. doi: 10.3109/13880209.2014.884606
49. Suvarna V, Sarkar M, Chaubey P, Khan T, Sherje A, Patel K, et al. Bone health and natural products- an insight. *Front Pharmacol.* (2018) 9:981. doi: 10.3389/fphar.2018.00981
50. Wang X, He Y, Guo B, Tsang MC, Tu F, Dai Y, et al. In vivo screening for anti-osteoporotic fraction from extract of herbal formula Xianlinggubao in ovariectomized mice. *PLoS One.* (2015) 10:e0118184. doi: 10.1371/journal.pone.0118184
51. Zihui Z, Qiong L, Wenying L, Baige Z, Lin M, Zinan Z. Systematic review of efficacy of Xianling Gubao capsules combined with Methorexatein the treatment of rheumatoid arthritis with Osteoporosis. *Evaluation Anal Drug-Use in Hospitals China.* (2021) 21:329–32. doi: 10.14009/j.issn.1672-2124.2021.03.017
52. Yuan L, Wu R, Liang Z, Wang H. Meta-analysis of xianling gubao capsule in the treatment of osteoporotic vertebral compression fracture. *Chin J Mod Appl Pharm.* (2023) 40:830–838. doi: 10.13748/j.cnki.issn1007-7693.20214277
53. Z Y, Xu W, J J, Z Y, W S, X Y. Systemic review of Jintiang capsules in treatment of postmenopausal osteoporosis. *Zhongguo Zhong Yao Za Zhi.* (2019) 44:186–92. doi: 10.19540/j.cnki.cjcm.20180709.007
54. Liang H, Wang O, Cheng Z, Xia P, Wang L, Shen J, et al. Jintiang combined with alfacalcidol improves muscle strength and balance in primary osteoporosis: a randomized, double-blind, double-dummy, positive-controlled, multicenter clinical trial. *J Orthopaedic Translation.* (2022) 35:53–61. doi: 10.1016/j.jot.2022.05.002
55. Nan P, Shanshan L, Jiangxia y, Haijing G. Economic evaluation of Jintiang capsule in the treatment of postmenopausal osteoporosis. *Chin J New Drugs.* (2024) 33:1403–8. doi: 10.3969/j.issn.1003-3734.2024.13.015
56. Jin H, Huang C, Zhang Y, Dong Y, Xiong Q, Wang D, et al. Meta-analysis of the synergistic effect of traditional Chinese medicine compounds combined with conventional Western medicine in the treatment of osteoporosis. *Front Endocrinol (Lausanne).* (2025) 16:1606753. doi: 10.3389/fendo.2025.1606753
57. Darba J, Kaskens L, Sorio Vilela F, Lothgren M. Cost-utility of denosumab for the treatment of postmenopausal osteoporosis in Spain. *Clinicoecon Outcomes Res.* (2015) 7:105–17. doi: 10.2147/CEOR.S78349
58. Hiligsmann M, Reginster JY. Cost effectiveness of denosumab compared with oral bisphosphonates in the treatment of post-menopausal osteoporotic women in Belgium. *PharmacoEconomics.* (2011) 29:895–911. doi: 10.2165/11539980-000000000-00000
59. You R, Mori T, Ke L, Wan Y, Zhang Y, Luo F, et al. Which injected antiosteoporotic medication is worth paying for? A cost-effectiveness analysis of teriparatide, zoledronate, ibandronate, and denosumab for postmenopausal osteoporotic women in China. *Menopause (New York, NY).* (2021) 29:210–8. doi: 10.1097/gme.0000000000001911
60. Chinese National Bureau of Statistics. (2023). Consumer Price Index 2022(2023-01-09). Available online at: https://www.sg.gov.cn/sgtjjdcd/gkmlpt/content/2/2354/mpost_2354542.html#7996 (Accessed on 2024-10-09).