INTRODUCTION

Osteoporosis in Myanmar is still a neglected disease entity in the national priority health problems. Due to the low level of awareness on osteoporosis and its consequent fragility fractures, limited availability and accessibility of standard diagnostic tools and medications, osteoporosis is mainly under diagnosed and under-treated in daily clinical practice. Rounds of discussions were held among relevant practitioners and professionals to identify national priorities for standard clinical practice guidelines. This Myanmar CPG on osteoporosis focuses on the assessment and management of post menopausal women and men over age 50 who are at high risk of fragility fractures and the integration of new tools for assessing the 10-year risk of fracture into overall management.

OBJECTIVES

To develop Myanmar Clinical Practice Guidelines (CPG) for health care providers that focus on prevention, diagnosis, and treatment aspects of osteoporosis and osteoporosis related fragility fractures among postmenopausal women and men over the age of 50.

(1) To develop guidelines on healthy lifestyle (namely nutrition and physical activity) for the prevention of osteoporosis in Myanmar context.

(2) To formulate cost effective ways to diagnose osteoporosis using bone densitometry technique, Osteoporosis Screening Tool for Asians (OSTA), spine radiographs, 10-year fracture risk assessment (FRAX) tool.
(3) To formulate evidence-based recommendations on the pharmacologic management of osteoporosis
(4) To develop the assessment plan for fragility fracture(s)
(5) To develop recommendations for the management of fragility fracture(s)

METHODOLOGY FOR THE DEVELOPMENT OF MYANMAR CPG FOR OSTEOPOROSIS

The Guidelines Committee was formed among the members of the Special Interest Group on Osteoporosis.

First of all, rounds of discussion were held among physicians, endocrinologists, orthopedic surgeons, gynecologists, radiologists, rheumatologists, physiatrists, general practitioners and allied health professionals to ascertain their view on osteoporosis and to identify priorities for the Myanmar CPG on osteoporosis.

Reviews of the literature were made according to these priorities to collect updated knowledge on diagnosis, prevention and treatment for osteoporosis and fragility fracture. Special references were made on those guidelines with reliable methodology and relevant epidemiological background to Myanmar (Canada, American, Asian, Philippine, Australia, Strong bone Asia). The Guideline Committee developed clinical Practice Guideline on osteoporosis with particular emphasis on the issues raised by discussion with Myanmar Health Care Providers.

The draft of the Myanmar CPG on Osteoporosis was discussed among the relevant healthcare providers on 2012 World Osteoporosis Day and resulted in the final CPG on osteoporosis.

SUMMARY OF RECOMMENDATIONS

Prevention
Non-pharmacological measures
1. Exercise
   (a) For those with or at risk for osteoporosis: appropriate resistance training and/or weight-bearing aerobic exercise.
   (b) For those with vertebral fractures: directed core stability exercises.
   (c) For those at risk of falls: exercises that focus on balance (e.g., Tai chi, balance and/or gait training).
   (d) For those in long-term care at high risk: use of hip protectors.
2. Smoking: Avoidance of tobacco usage is recommended
3. Alcohol: Avoidance of excessive consumption of alcohol is recommended (<3 units per day in men and <2 units per day in women)
4. Exercises: daily exercise of 20 – 30 minutes for 3 – 5 times a week such as regular walking, jogging, aerobic dancing and tai-Chi or Chi-gong

Vitamin D and Calcium
Calcium: 600–1200 mg oral daily should be prescribed.
Vitamin D:
For healthy adults at low risk of vitamin D deficiency, routine supplementation with 400–1000 IU (10–25 mg) vitamin D<sub>3</sub> daily is recommended.

For adults over age 50 at moderate risk of vitamin D deficiency, supplementation with 800–1000 IU (20–25 mg) vitamin D<sub>3</sub> daily is recommended.

Screening
Osteoporosis Screening Tool for Asians (OSTA) is recommended to identify the individual's risk for Osteoporosis where central dual x-ray absorptiometry (DXA) is unavailable.

![Figure 1. Osteoporosis self-assessment tool for Asians (OSTA)](image)

Recommendation
- High risk patients: to measure BMD, if possible, and consider drug treatment even if BMD is not available.
- Medium risk patients: to measure BMD and consider drug treatment if BMD is low.

Low risk patients: not to measure BMD unless other risk factors are present. Fracture Risk Assessment Tool (FRAX) tool should be used for all postmenopausal women with at least one WHO risk factor.

Diagnosis
Recommendation
It is recommended that diagnosis of osteoporosis be based on WHO diagnostic classification criteria for bone mass using the Dual Energy X-ray Absorptiometry (DEXA) as the gold standard.

According to the World Health Organization criteria, osteoporosis is diagnosed by bone mineral density (BMD) measurements.
Normal bone density: T-score greater than -1
Osteopenia (low bone mass): T-score between -1 and -2.5
Osteoporosis: T-score lower than -2.5

(Note: The T-score is the number of standard deviations of BMD below the young normal mean).

Areas for measuring BMD

DEXA is currently the 'gold standard' for the diagnosis of osteoporosis. BMD can be measured at the hip, the lumbar spine and the total body.

Ultrasound measurements are not recommended for diagnosing osteoporosis.

Indications for BMD measurements

The OSTA is applicable in Myanmar as the resources for DEXA are constrained.

1. DEXA measurements should be performed on the following subjects:
   1. Postmenopausal women 65 years or older, regardless of additional risk factors.
      This recommendation includes women 65 years or older, who have been taking osteoporosis therapy and who have not had a BMD test.
   2. Postmenopausal women younger than 65 years and with 1 or more of the following additional risk factors for osteoporosis:
      • Parental history of hip fracture;
      • Current cigarette smoking;
      • A body weight less than 57.2 kg for Caucasians; for Asian populations, a criteria of BMI <19
      • Use of (or plans to use) oral corticosteroids at 7.5mg/day for longer than 3 months;
      • Serious long-term conditions thought to increase fracture risk, such as hyperthyroidism or malabsorption.

Postmenopausal women who have had a fracture of any type as an adult after age 45 years

2. Biochemical markers of bone turnover
   Biochemical markers of bone turnover can be used for assessing adherence to, effectiveness of therapy and assessment of fracture risk if available.

   The following biochemical markers of bone turnover can be measured in serum and urine:
   Biochemical markers of bone formation (in serum)
   • Procollagen type I propeptides (PINP)
   • C and N-telopeptides of type I collagen cross-link (in serum and urine)

3. Other Tests
   The following are the routine investigations that should be performed in patients with osteoporosis: complete blood count and erythrocyte sedimentation rate, renal function tests, liver function tests, general biochemistry including serum calcium, phosphate, and alkaline phosphatase, thyroid function tests, testosterone level in men.

   Other special tests such as serum protein electrophoresis, parathyroid hormone and 25-hydroxy vitamin D are only indicated if the initial work-up is suggestive of a related disorder.

Treatment

Recommendations

Among those with BMD examination treatment should be started if:

a. Vertebral compression fracture/s confirmed through radiograph (clinical osteoporosis);
b. BMD T-score of ≤ -2.5;
c. BMD T-score between -1 and - 2.5 SD with any of the following:
   c1. History of previous fracture
   c2. Secondary causes associated with high risk for fracture
   c3. 10-year probability of hip fracture ≥ 3% or any major osteoporosis related fracture of ≥20% based on the FRAX estimates

   Among those without BMD measurement, it is suggested that treatment be started if patient:

   (i) belongs to the high risk category based on OSTA tool where central BMD cannot be done or not available
   (ii) has a 10-year probability of hip fracture ≥ 3% or any major osteoporosis related fracture of ≥ 20% based on the FRAX estimates

Specific therapies for osteoporosis

Recommended pharmacologic options

i. Alendronate is the first line option for all osteoporosis. The recommended dose is 70 mg once weekly or 10 mg daily for 5 years. It should be used with caution in renal impairment and history of oesophageal reflux/hiatus hernia. These agents should be taken on an empty stomach at least 30 minutes before breakfast.

ii. Zoledronic acid is recommended as a second line for those who have GI intolerance to oral bisphosphates. Recommended dosage is 5 mg IV infused over at least 15 minutes every 12 months.

iii. Raloxifene, selective estrogen receptor modulators (SERM) is recommended as 3rd line of medical treatment. It is effective for prevention and treatment
of vertebral fracture in post-menopausal women. Prescribed dosage is 60 mg once daily. It has not been shown to decrease the risk of hip fractures, but reduces the risk of vertebral fractures.

iv. Strontium ranelate has been shown to both increase osteoblastic bone formation and reduce osteoclastic bone resorption. Given at 2 grams daily orally, it may be used as first line therapy in high risk patient or in those intolerant of bisphosphonates as a second line therapy. It is indicated in post-menopausal osteoporosis for reduction of fracture risk in hip and vertebrae.

v. Calcitonin spray 200IU daily in alternating nostrils can be recommended in the initial first month after an acute vertebral compression fracture. It is not recommended for long term treatment. It has no effect on prevention of hip fracture. It is less effective for increasing bone density than estrogen or bisphosphates.

vi. Teriparatide (1-34 parathormone) is not available locally and currently needs prescription by a specialist in bone disease or an endocrinologist. It can be recommended in those who suffer further fractures despite treatment with other agents.

vii. Estradiol is not recommended for treatment of osteoporosis.

**Special Groups (Long-term Corticosteroid [CS] Therapy)**

It is reasonable to consider starting prophylactic therapy in patients on chronic steroids. Long term corticosteroid of ≥3 months at a prednisolone-equivalent dose ≥7.5 mg daily.

Alendronate is recommended for the treatment of steroid-induced osteoporosis.

- Measurement of BMD using DEXA is currently recommended for assessment of fracture risk in individuals treated with CS.
- Anti-resorptive therapy is recommended for subjects with a BMD T-score of -1.5 and in whom it is intended to continue therapy for at least 3 months; or subjects aged 65 years or over with a prior fragility fracture. Calcium and vitamin D supplementation are recommended for all CS patients.

**Osteoporosis Related Fragility Fractures**

The fracture during activity that would not normally injure young healthy bone (i.e., Fall from standing height or less) with trivial force usually accompanied with risk factors for osteoporosis or risk factors for fall or risk factors for fracture. (IOF)

**Recommendation**

- It is suggested that exercise be encouraged among both the housebound elderly and those in the community due to its benefit on balance and indirectly on fracture prevention.
- Other than exercise, provision of hip protectors can be considered to reduce incidence of hip fracture.
- It is recommended that high risk Postmenopausal Women (PMW) be given pharmacologic options (e.g., bisphosphonates, selective estrogen receptor modulator, hormonal replacement therapy, calcitonin, strontium ranelate to increase BMD or reduce fracture risks.)

**Clinical assessment strategies for fragility fracture**

The major risk factors for fragility fracture are prior fragility fracture; increasing age, low BMD, low body weight, family history of osteoporotic fracture, glucocorticoids use and current smoking.

There are three main assessment areas after fracture

(1) Bone morphology and bone mass density
(2) Patient’s underlying medical diseases
(3) Underlying etiology of osteoporosis and risk factors

**Methods of BMD**

**When to assess BMD**

DEXA is not practicable immediately after getting fragility fracture.

X-ray observation—osteopenia on x-ray implies significant bone loss with decreased opacity, thin cortices, wide canals, current fractures, healing fractures.

DEXA—current gold standard for BMD

Fragility fracture patient assessment in addition to routine pre-op or fracture evaluation:-

- Thorough history taking and physical examination relevant to fragility fracture should be done.

**Laboratory tests are recommended for assessment to exclude secondary osteoporosis**

**Policy for osteoporotic fracture management:** immediate hospitalization whenever feasible. The following objectives should be followed:

(1) Immediate comfort with pain relief and protection
(2) Setup the appropriate specific fracture care
(3) Minimize dependences and maximize the mobility
(4) Treat underlying osteoporosis and monitor
(5) Identification and prevention of risk factors
(6) Patient’s education

**Monitoring Plan after fracture fixation**

- Fracture stability follow up every three weeks
- Fracture union follow up at every six weeks
- Counseling for risk reduction immediately after discharge and at follow-up
CONCLUSION

This is the first published guideline that focused on both medical and surgical aspect of osteoporosis and fragility fracture. It is advisable for all stakeholders to use the guideline so as to be able to give optimal health care for individual afflicted with the condition. This guideline is meant to help clinicians to make appropriate decision in management of osteoporosis and its consequent fragility fracture. However, clinician's best clinical judgments are the most important assets in the management of these patients. Regular review of these guidelines will be made in the light of fresh and new evidence, which will come up in the medical literature in future.

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