UCLA MEDICAL	GROUP / Managed Care Operations	UCLA Health
DEPARTMENT:	Utilization Management	POLICY NUMBER: TBD
SECTION:	UM Program	Page 1 of 4
TITLE:	Bone Mineral Density Measurement (BMD)	ISSUE:
		EFFECTIVE:
Date Revised:	/02, 1/07,2/09, 5/10, 2/2012, 5/2016, 5/2017, 08/2019,8/2021	
APPROVED BY U	MC: 9/02, 1/07,2/09, 5/10, 3/2012, 3/26/2014, 5/6/16, 5/2017, 08/2019,	8/2021

BMD measurement is NEVER indicated unless the results will influence a treatment decision.

INDICATIONS FOR BMD MEASUREMENT:

- Bone mineral density (BMD) testing with DXA should be performed: NOF and USPSTF guideline: In women age 65 and older and men age 70 and older (USPSTF concludes that the current evidence is insufficient to assess the balance of benefits and harms of screening for osteoporosis to prevent osteoporotic fractures in men).
- NOF: In postmenopausal women and men above age 50–69, based on risk factor profile
- The USPSTF recommends screening for osteoporosis with bone measurement testing to prevent osteoporotic fractures in postmenopausal women younger than 65 years who are at increased risk of osteoporosis, as determined by a formal clinical risk assessment tool. Several tools are available to assess osteoporosis risk: the Simple Calculated Osteoporosis Risk Estimation (SCORE; Merck), Osteoporosis Risk

Assessment Instrument (ORAI), Osteoporosis Index of Risk (OSIRIS), and the Osteoporosis Self-Assessment Tool (OST). The FRAX tool (University of Sheffield), which assesses a person's 10-year risk of fracture, is also a commonly used tool • In postmenopausal women and men age 50 and older who have had an adult age fracture, to diagnose and determine degree of osteoporosis

- Vertebral imaging should be performed: In all women age 70 and older and all men age 80 and older if BMD T-score is ≤-1.0 at the spine, total hip, or femoral neck
- In women age 65 to 69 and men age 70 to 79 if BMD T-score is ≤-1.5 at the spine, total hip, or femoral neck
- In postmenopausal women and men age 50 and older with specific risk factors:
- Low-trauma fracture during adulthood (age 50 and older)
- Historical height loss (difference between the current height and peak height at age 20) of 1.5 in. or more (4 cm)

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- Prospective height loss (difference between the current height and a previously documented height measurement) of 0.8 in. or more (2 cm)
- Recent or ongoing long-term glucocorticoid treatment
- If bone density testing is not available, vertebral imaging may be considered based on age alone.
- Check for secondary causes of osteoporosis. Biochemical markers of bone turnover can aid in risk assessment and serve as an additional monitoring tool when treatment is initiated. **Monitoring patients**
- Perform BMD testing 1 to 2 years after initiating medical therapy for osteoporosis and every 2 years thereafter.
- More frequent BMD testing may be warranted in certain clinical situations.
- The interval between repeat BMD screenings may be longer for patients without major risk factors and who have an initial T-score in the normal or upper low bone mass range.
- Biochemical markers can be repeated to determine if treatment is producing expected effect.

FREQUENCY OF BMD MEASUREMENT: not more than every two years when medically indicated, unless the patient has a medical condition associated with accelerated bone loss. Patients with a normal scan should not have it repeated for five years, unless the patient has a medical condition associated with accelerated bone loss.

BMD measurement can be used to:

- Establish/confirm diagnosis of osteoporosis
- Predict future fracture risk
- Monitor changes in BMD due to medical conditions or therapy

BMD measurement is NEVER indicated unless the results will influence a treatment decision.

SUPPLEMENTAL INFORMATION

OSTEOPOROSIS: BMD values greater than 2.5 standard deviations below a non-hispanic white female reference group mean aged 20-29 years old (T score)

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OSTEOPENIA: BMD value between 1-2.5 standard deviations below a non-hispanic white female reference group mean-aged 20-29 years old (T score)

RISK FACTORS FOR OSTEOPOROTIC FRACTURE

NONMODIFIABLE

- Personal history of fracture as an adult
- History of osteoporosis in a first degree relative
- Caucasian race
- Advanced age
- Dementia
- Poor health/frailty

MODIFIABLE

- Current cigarette smoking
- · Low body weight
- Estrogen deficiency (early menopause (age <45) or bilateral ovariectomies, prolonged (>1 yr) premenopausal amenorrhea)
- Low calcium intake (lifelong)
- Alcoholism
- Impaired eyesight despite adequate correction
- Recurrent falls
- Inadequate physical activity
- Poor health/frailty

RISK FACTORS FOR OSTEOPOROSIS

Hypogonadism Amenorrhea/oligomenorrhea

Hyperprolactinemia Premature menopause

Endocrine and metabolic

disorders

Hyperparathyroidism or tumor secretion of PTH -rp

Untreated/inadequately treated hyperthyroidism

Cushing's syndrome Adrenal insufficiency

Acromegaly

Hypophosphatemia Hemochromatosis

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Porphyria

Gastrointestinal disorders Malabsorbtion syndromes

Chronic obstructive jaundice Primary biliary cirrhosis

Gastrectomy

Pancreatic insufficiency

Eating disorders Anorexia nervosa

Bulimia

Chronic anemia (sickle cell disease, thalassemia, hemophilia)

Cancers with diffuse bone involvement

Chronic disorders Rheumatoid arthritis

Ankylosing spondilytis Renal failure/hemodyalisis

Liver failure Amyloidosis

TPN or nutritional deficiencies Rickets, scurvy, etc.

Medications Glucocorticoids (three months or more of 5 mg prednisone /

equivalent daily) Anticonvulsants

Heparin

Thyroxin (excessive dose) Heparin and warfarin

Cyclosporine

GnRH analogues and antiestrogens

Depo-Provera (chronic)

Chemotherapy

Lithium Aluminum

Lifestyle factors Smoking

Excessive alcohol intake Excessive caffeine intake

Prolonged immobilization (three months or more)

REFERENCES

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US Preventive Services Task Force: Screening for Osteoporosis to Prevent Fractures US Preventive Services Task Force Recommendation Statement JAMA2531-2521:(24)319;2018.

The National Osteoporosis Foundation: Clinician's guide to Prevention and Treatment of Osteoporosis, <u>Osteoporos Int</u>. 2014; 25(10): 2359–2381.