

1 **Clinical Practice Guideline:** **Bone Density Screening – Peripheral**

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3 **Date of Implementation:** **July 13, 2006**

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5 **Effective Date:** **January 29, 2026**

6
7 **Product:** **Specialty**

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9
10 **GUIDELINES**

11 American Specialty Health – Specialty (ASH) considers Peripheral Bone Density
12 Screening medically necessary when following United States Preventive Services Task
13 Force (USPSTF) guidelines.

14
15 The current US Preventive Services Task Force (USPSTF) recommends screening for
16 osteoporosis with bone measurement testing to prevent osteoporotic fractures in women
17 65 years and older. The USPSTF recommends screening for osteoporosis with bone
18 measurement testing to prevent osteoporotic fractures in postmenopausal women younger
19 than 65 years who are at increased risk of osteoporosis, as determined by a formal clinical
20 risk assessment tool. (June 2018). Risk factor determination should be performed using the
21 FRAX tool or another valid and reliable tool. The USPSTF concludes that the current
22 evidence is insufficient to assess the balance of benefits and harms of screening for
23 osteoporosis to prevent osteoporotic fractures in men. The most commonly used bone
24 measurement test used to screen for osteoporosis is central dual-energy x-ray
25 absorptiometry (DXA); other screening tests include peripheral DXA and quantitative
26 ultrasound (QUS). Central DXA measures bone mineral density (BMD) at the hip and
27 lumbar spine. Most treatment guidelines recommend using BMD, as measured by central
28 DXA, to define osteoporosis and the treatment threshold to prevent osteoporotic fractures.

29
30 **DESCRIPTION/BACKGROUND**

31 Several tools are available to assess osteoporosis risk: the Simple Calculated Osteoporosis
32 Risk Estimation (SCORE; Merck), Osteoporosis Risk Assessment Instrument (ORAI),
33 Osteoporosis Index of Risk (OSIRIS), and the Osteoporosis Self-Assessment Tool (OST).
34 These tools seem to perform similarly and are moderately accurate at predicting
35 osteoporosis. The FRAX tool (University of Sheffield), which assesses a person’s 10-year
36 risk of fracture, is also a commonly used tool. The FRAX tool includes questions about
37 previous DXA results but does not require this information to estimate fracture risk.
38 Because the benefits of treatment are greater in persons at higher risk of fracture, one
39 approach is to perform bone measurement testing in postmenopausal women younger than
40 65 years who have a 10-year FRAX risk of major osteoporotic fracture (MOF) (without
41 DXA) greater than that of a 65-year-old white woman without major risk factors. Bone
42 density measurement is performed as a screen for conditions such as osteopenia and

1 osteoporosis, bone weakening conditions due to bone resorption occurring at a faster rate
2 than bone formation. This change in bone density can be due to aging or disease processes
3 and is related to a multitude of factors, including hormonal changes, calcium consumption,
4 diet, and level of physical activity. Having osteopenia and/or osteoporosis is a risk factor
5 for fracture, and because these disease processes begin weakening bones long before
6 fractures occur, early screening for, and treatment of, decreased bone density can be useful
7 for preventing fractures. Studies have shown that screening those at risk for osteoporosis
8 can reduce the risk of fractures associated with falls or other injuries. The most commonly
9 used bone measurement test used to screen for osteoporosis is central DXA; other
10 screening tests include peripheral DXA and quantitative ultrasound (QUS). All the
11 osteoporosis drug therapy studies reviewed by the USPSTF used central DXA to determine
12 eligibility for study enrollment. Peripheral DXA measures BMD at the lower forearm and
13 heel. Quantitative ultrasound also evaluates peripheral sites and has similar accuracy in
14 predicting fracture risk as DXA, while avoiding the risk of radiation exposure; however, it
15 does not measure BMD (June 2018).

16
17 For peripheral bone density measurement, there are 3 different types of scans that can be
18 performed to test bone density: photon absorptiometry, peripheral dual energy x-ray
19 absorptiometry, and ultrasound.

20
21 Photon absorptiometry uses low doses of radiation but is very slow compared to all other
22 bone density tests using radiation. Although very popular in the past, this method is no
23 longer as commonly used. The radioactive source gradually decays and must be replaced
24 over time. It is also not as accurate as other tests using radiation such as dual energy x-ray
25 absorptiometry (DXA).

26
27 A modified version of the DXA scan is called peripheral dual energy x-ray absorptiometry
28 (P-DXA). This uses the x-ray technique of DXA but only measures density in the limbs
29 such as the wrist or the heel. It uses low doses of radiation and is faster than traditional
30 DXA.

31
32 Ultrasound uses sound waves to determine bone mineral density (BMD) for heel scan
33 screenings. Ultrasound is rapid and does not use radiation. This technique is generally used
34 as a prescreening tool for bone mineral density. If evidence of bone loss is detected, the
35 patient is generally referred for a more comprehensive scan of the hip and spine using
36 DXA. The most commonly used type of ultrasound for a heel scan is quantitative
37 ultrasound, and there are numerous devices using slightly varying techniques designed for
38 this type of ultrasound. Quantitative ultrasound works by evaluating two measures,
39 broadband ultrasound attenuation (BUA) and speed of sound (SOS). SOS is a measurement
40 of how quickly sound travels through the bone, while BUA is a measure of how much
41 sound is absorbed by the bone.

1 The advantage of these devices is the ability to bring bone density screening assessments
2 to a large portion of the population who otherwise would not be able to have testing. These
3 machines cost considerably less than those evaluating the hip and spine. However, it is
4 important to note that density changes in the heel and wrist occur much slower than those
5 in the hip or spine. The heel may be normal in bone density even when sites such as the
6 hip or spine are already significantly abnormal. The rate of false negative findings is,
7 however, low enough to support the use of these techniques as a screening procedure.

8
9 There are inherent risks in any procedure that involves radiation such as the photon
10 absorptiometry and x-ray, and as such these should be used only after the benefits and risks
11 have been assessed.

12 13 **EVIDENCE REVIEW**

14 **DXA**

15 Bone measurement testing with central DXA is the most commonly used and studied
16 method for the diagnosis of osteoporosis. Central DXA uses radiation to measure BMD at
17 central bone sites (hip and lumbar spine), which is the established standard for diagnosis
18 of osteoporosis and for guiding decisions about treatment. DXA can also be used at
19 peripheral bone sites (such as the lower forearm and heel) to identify persons with low
20 bone mass; however, most treatment guidelines recommend follow-up with central DXA
21 before initiating treatment for osteoporosis. Screening with peripheral DXA and other
22 imaging techniques may help increase access to screening in geographic locations (e.g.,
23 rural areas) where machines that perform central DXA may not be available. The USPSTF
24 identified 2 studies ($n = 712$) that reported on the accuracy of peripheral DXA at the
25 calcaneus to identify osteoporosis; compared with central DXA, the area under the curve
26 (AUC) ranged from 0.67 to 0.80 in women with a mean age of 61 years.

27 28 **QUS**

29 Quantitative ultrasound is another imaging technique used at peripheral bone sites (most
30 commonly the calcaneus), and it does not require radiation exposure. Compared with
31 central DXA, the AUC for QUS measured at the calcaneus in women ranged from 0.69 to
32 0.90, with a pooled estimate of 0.77 (95% CI, 0.72-0.81; 7 studies; $n = 1,969$). In men, the
33 AUC ranged from 0.70 to 0.93, with a pooled estimate of 0.80 (95% CI, 0.67-0.94; 3
34 studies; $n = 5,142$). However, QUS does not measure BMD, that is the current diagnostic
35 criteria for osteoporosis. In addition, drug therapy trials for osteoporosis treatment
36 generally use central DXA measurement of BMD as criteria for inclusion of study
37 populations. Thus, before QUS results could be routinely used to initiate treatment without
38 any further DXA measurement, a method for converting or adapting QUS results to the
39 DXA scale needs to be developed. Chou et al. (2014) demonstrated, “in a multiracial
40 referral population heel BMD predicts central osteoporosis and prevalent vertebral
41 fractures equally well in African American as in Caucasian women and may be better than
42 central BMD in assessing fragility in glucocorticoid users.” These studies indicate that

1 quantitative ultrasound is an effective and safe prescreening tool for bone mineral density
 2 that is quick and involves no radiation. Peripheral DXA was found to be a useful
 3 measurement of bone density but does involve the use of radiation, and as such should be
 4 used with care after the benefits and risks have been considered. Hashimi and Elfandi
 5 (2016) aimed to find out whether heel ultrasound is as good as central bone densitometry
 6 scanning in diagnosing osteoporosis in patients who are at high risk of osteoporosis. The
 7 recruited patients attended for a DEXA scan of the left hip and lumbar spine. All subjects
 8 had an ultrasound of the left heel using the quantitative heel ultrasound machine. The
 9 results of DEXA scan were blinded from the results of ultrasound and vice versa. The
 10 sensitivity and specificity of the ultrasound heel test to predict osteoporosis were 53%
 11 (95%CI: 29-77) and 86% (95%CI: 75-96) respectively. Specificity for predicting bone
 12 mineral density (BMD)-defined osteoporosis was high (86%), but sensitivity was low
 13 (53%). Authors concluded that heel ultrasound result in the osteoporotic range was highly
 14 predictive of BMD-defined osteoporosis. A positive ultrasound heel test in high-risk
 15 patients is more useful in ruling in osteoporosis than a negative test to rule out osteoporosis.

17 **PRACTITIONER SCOPE AND TRAINING**

18 Practitioners should practice only in the areas in which they are competent based on their
 19 education training and experience. Levels of education, experience, and proficiency may
 20 vary among individual practitioners. It is ethically and legally incumbent on a practitioner
 21 to determine where they have the knowledge and skills necessary to perform such services.

22
 23 It is best practice for the practitioner to appropriately render services to a patient only if
 24 they are trained, equally skilled, and adequately competent to deliver a service compared
 25 to others trained to perform the same procedure. If the service would be most competently
 26 delivered by another health care practitioner who has more skill and expert training, it
 27 would be best practice to refer the patient to the more expert practitioner.

28
 29 Best practice can be defined as a clinical, scientific, or professional technique, method, or
 30 process that is typically evidence-based and consensus driven and is recognized by a
 31 majority of professionals in a particular field as more effective at delivering a particular
 32 outcome than any other practice (Joint Commission International Accreditation Standards
 33 for Hospitals, 2020).

34
 35 Depending on the practitioner's scope of practice, training, and experience, a member's
 36 condition and/or symptoms during examination or the course of treatment may indicate the
 37 need for referral to another practitioner or even emergency care. In such cases it is prudent
 38 for the practitioner to refer the member for appropriate co-management (e.g., to their
 39 primary care physician) or if immediate emergency care is warranted, to contact 911 as
 40 appropriate. See the *Managing Medical Emergencies (CPG 159 – S)* clinical practice
 41 guideline for information.

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