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PRESENTATION

- Post-menopausal women
- Pre-menopausal on tamoxifen or GnRH
- Women on aromatase inhibitors

- Baseline BMD
- 25-OH vitamin D¹
- Evaluate for new low impact fracture

25-OH Vitamin D normal (≥ 30 ng/mL) and BMD normal (T-score ≥ -1.0) and no new low impact fracture

25-OH Vitamin D abnormal (< 30 ng/mL)

BMD abnormal³(T-score < -1.0 to -2.4) and any vitamin D level and no new low impact fracture

BMD abnormal³(T-score ≤ -2.5) and any vitamin D level⁵ and no new low impact fracture

Any BMD with New low impact fracture

TREATMENT

- Repeat tests in 2 years **and**
- Reinforce universal recommendations²

A

- Ergocalciferol 50,000 IU once a week for 8 weeks, then continue once a month, **or**
- Over the counter vitamin D3 1,000-2,000 IU daily **and**
- Recheck vitamin D, calcium, and albumin on the next visit **and**
- Reinforce universal recommendations²

- Reinforce universal recommendations² **and**
- Repeat DXA every 1-2 years **and**
- Consider medical therapy or referral to bone health specialist based on risk factors (assess by FRAX⁴)

- Start bisphosphonates:
 - Alendronate 70 mg po weekly, **or**
 - Risedronate 35 mg po weekly or 150 mg po monthly, **or**
 - Ibandronate 150 mg po monthly or 3 mg IV every 3 months, **or**
 - Zoledronic acid 5 mg IV once a year (use institutional order set) **or**
- Start denosumab at 60 mg subcutaneously every 6 months (use institutional order set) **or**
- Refer to bone health specialist **and**
- Reinforce universal recommendations²
 - If bone loss risks have changed significantly or major therapeutic intervention has been undertaken, obtain a 12 month follow up DXA

- Start universal recommendations² **and**
- Refer to Bone Health Specialist

GnRH = Gonadotropin-releasing hormone
 BMD = Bone Mineral Density
 DXA = Dual-energy X-ray Absorptiometry
 IU = International Units

¹ 25-hydroxyvitamin D, also known as 25-hydroxycholecalciferol, calcidiol or abbreviated as 25-OH Vitamin D, the main vitamin D metabolite circulating in plasma

² Universal recommendations:

- Elemental calcium 1,000 – 1,200 mg/day from all sources
- Avoid tobacco (see [Tobacco Cessation algorithm](#))
- Vitamin D 800 – 1,000 IU/day
- Limit alcohol
- Weight-bearing/muscle - strengthening exercises (see [Physical Activity algorithm](#))
- Limit caffeine

³ Abnormal BMD: Osteopenia, T-score between -1.0 and -2.4; Osteoporosis, T-score ≤ -2.5

⁴ FRAX[®] - Fracture Risk Assessment Tool at www.shef.ac.uk/frac

⁵ If vitamin D level is < 26 ng/mL, replenish with supplementation prior to initiating medical therapy for osteoporosis. See Box A for recommendation on vitamin D repletion.

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SUGGESTED READINGS

- Black, D. M., Bauer, D. C., Schwartz, A. V., Cummings, S. R., & Rosen, C. J. (2012). Continuing bisphosphonate treatment for osteoporosis - for whom and for how long? *New England Journal of Medicine*, 366(22), 2051-2053. doi:10.1056/NEJMp1202623
- Centre for Metabolic Bone Diseases, University of Sheffield. (n.d.). *FRAX® Fracture risk assessment tool. Calculation tool*. Retrieved from <https://www.sheffield.ac.uk/FRAX/tool.aspx>
- Coleman, R. E., Wright, J., Houston, S., Agrawal, R., Purohit, O. P.-K., Hayward, L., . . . Marshall, H. (2012). Randomized trial of marker-directed versus standard schedule zoledronic acid for bone metastases from breast cancer. *Journal of Clinical Oncology*, 30(15_suppl), 511. doi:10.1200/jco.2012.30.15_suppl.511
- Gralow, J. R., Biermann, J. S., Farooki, A., Fournier, M. N., Gagel, R. F., Kumar, R., . . . Van Poznak, C. (2013). NCCN task force report: Bone health in cancer care. *Journal of the National Comprehensive Cancer Network*, 11(suppl_3), S1-S51. doi:10.6004/jnccn.2013.0215
- Institute for Clinical Systems Improvement (ICSI). (n.d.). *Guidelines: Osteoporosis*. Retrieved from: <https://www.icsi.org/guideline/osteoporosis/osteoporosis/>
- National Comprehensive Cancer Network. (2020). *Breast Cancer*. (NCCN Guideline Version 6.2020). Retrieved from https://www.nccn.org/professionals/physician_gls/pdf/breast.pdf
- National Osteoporosis Foundation. (2015). *NOF's clinician's guide to prevention and treatment of osteoporosis*. Retrieved from <https://my.nof.org/bone-source/education/clinicians-guide-to-the-prevention-and-treatment-of-osteoporosis>
- The DIPART (Vitamin D Individual Patient Analysis of Randomized Trials) Group. (2010). Patient level pooled analysis of 68 500 patients from seven major vitamin D fracture trials in US and Europe. *BMJ*, 340(7738), b5463. doi:10.1136/bmj.b5463
- U.S. Preventive Services Task Force. (2011). Screening for osteoporosis: U.S. preventive services task force recommendation statement. *Annals of Internal Medicine*, 154(5), 356-364. doi:10.7326/0003-4819-154-5-201103010-00307

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DEVELOPMENT CREDITS

This survivorship algorithm is based on majority expert opinion of the Breast Survivorship work group at the University of Texas MD Anderson Cancer Center. It was developed using a multidisciplinary approach that included input from the following:

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