

Clinical Policy: 25-hydroxyvitamin D Testing in Children and Adolescents

Reference Number: CP.MP.157

Effective Date: 12/17

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[Revision Log](#)

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Description

A global consensus statement recommends against universal screening for vitamin D deficiency in healthy children as there is insufficient evidence that the potential benefits of testing outweigh the potential harms.²

Policy/Criteria

- I. It is the policy of health plans affiliated with Centene Corporation[®] that 25-hydroxyvitamin D testing in healthy, including obese but otherwise healthy, children (age ≥ 1 and ≤ 18) is **not medically necessary** because these tests have not been demonstrated to have a clear clinical benefit.

Background

Measurement of 25-OH-D (25-hydroxyvitamin D) concentration is the appropriate screening test for vitamin D deficiency, as opposed to 1,25-OH₂-D, which has little to no predictive value related to bone health.⁶ However, there is lack of agreement concerning the best type of assay to conduct when measuring 25-hydroxyvitamin D.⁴ Furthermore, there is substantial controversy concerning cutoff levels to define vitamin D deficiency, as the evidence is inconsistent regarding optimal levels of vitamin D.¹

Prevalence of vitamin D deficiency in children (defined in the study as levels < 20 ng/mL) is estimated to be about 14%, although estimates range from 14% to 37%.^{3,6} Rates of deficiency vary among certain populations, with increased risk among black and Hispanic teenagers, as well as overweight and obese children and adolescents.⁶ Reduced serum vitamin D in overweight and obese children and adolescents reflects sequestration in adipose tissue, but little is known about the significance of low serum vitamin D in this population.⁴

A global consensus of 33 experts, convened at the request of the European Society for Pediatric Endocrinology, reviewed the available literature on prevention and management of nutritional rickets, and determined that routine vitamin D screening is not recommended for healthy children.² They note the frequent coexistence of dietary calcium and vitamin D deficiency, which alters the threshold for development of rickets, and makes a single screening value impractical.² The global consensus panel advocates for identification and screening of groups at high risk for vitamin D deficiency based on clinical factors, as opposed to universal screening as public health policy.

The American Academy of Pediatrics (AAP) – Section on Endocrinology advises against ordering vitamin D concentrations routinely in otherwise healthy children, including children who are overweight or obese.⁵ The AAP's report on optimizing bone health recommends

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screening for vitamin D deficiency only in children and adolescents with conditions associated with reduced bone mass and/or recurrent low-impact fractures.⁶

For healthy children and adolescents who are not ingesting enough foods with vitamin D, the AAP recommends supplementation with vitamin D, as does the global consensus panel convened by the European Society for Pediatric Endocrinology.

Coding Implications

This clinical policy references Current Procedural Terminology (CPT®). CPT® is a registered trademark of the American Medical Association. All CPT codes and descriptions are copyrighted 2017, American Medical Association. All rights reserved. CPT codes and CPT descriptions are from the current manuals and those included herein are not intended to be all-inclusive and are included for informational purposes only. Codes referenced in this clinical policy are for informational purposes only. Inclusion or exclusion of any codes does not guarantee coverage. Providers should reference the most up-to-date sources of professional coding guidance prior to the submission of claims for reimbursement of covered services.

Table 1: CPT codes not medically necessary when billed with a corresponding ICD-10CM in Table 2

CPT® Codes	Description
82306	Vitamin D; 25 hydroxy, includes fraction(s), if performed

Table 2: ICD-10-CM diagnosis codes not medically necessary when billed with a corresponding CPT code in Table 1.

ICD-10-CM Code	Description
E66.01	Morbid (severe) obesity due to excess calories
E66.09	Other obesity due to excess calories
E66.1	Drug-induced obesity
E66.3	Overweight
E66.8	Other obesity
E66.9	Obesity, unspecified
Z00.00	Encounter for general adult medical examination without abnormal findings
Z00.129	Encounter for routine child health examination without abnormal findings
Z00.8	Encounter for other general examination
Z68.52	Body mass index (BMI) pediatric, 5 th percentile to less than 85 th percentile for age
Z68.53	BMI pediatric, 85 th percentile to less than 95 th percentile for age
Z68.54	BMI pediatric, greater than or equal to 95 th percentile for age

Reviews, Revisions, and Approvals	Date	Approval Date
Policy created	12/17	12/17

References

1. U.S. Preventive Services Task Force. Final Recommendation Statement: Vitamin D Deficiency: Screening. U.S. Preventive Services Task Force. December 2016.
2. Munns CF, Shaw N, Kiely M, Specker BL, Thacher TD, et al. Global Consensus Recommendation on Prevention and Management of Nutritional Rickets. *J Clin Endocrinol Metab.* 2016 Feb;101(2):394-415. Co-Published in *Horm Res Paediatr.* 2016;85(2):83-106.
3. Saintonge S, Bang H, Gerber LM. Implications of a new definition of vitamin D deficiency in a multiracial us adolescent population: the National Health and Nutrition Examination Survey III. *Pediatrics* 2009; 123:797.
4. Misra M. Vitamin D insufficiency and deficiency in children and adolescents. In: UpToDate, Waltham, MA. Accessed December 11, 2017.
5. Golden N, Abrams S, and the AAP Committee on Nutrition. Optimizing Bone Health in Children and Adolescents. *Pediatrics.* 2014 Oct; 134(4):e1229-43. doi: 10.1542/peds.2014-2173.
6. Turer CB, Lin H, Flores G. Prevalence of vitamin D deficiency among overweight and obese US children. *Pediatrics* December 2012;doi:10.1542/peds2012-1711.
7. Munns CF, Shaw N, Kiely M, Specker BL, Thacher TD, et al. Global Consensus Recommendation on Prevention and Management of Nutritional Rickets. *J Clin Endocrinol Metab.* 2016 Feb;101(2):394-415. Co-Published in *Horm Res Paediatr.* 2016;85(2):83-106.

Important Reminder

This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. The Health Plan makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. “Health Plan” means a health plan that has adopted this clinical policy and that is operated or administered, in whole or in part, by Centene Management Company, LLC, or any of such health plan’s affiliates, as applicable.

The purpose of this clinical policy is to provide a guide to medical necessity, which is a component of the guidelines used to assist in making coverage decisions and administering benefits. It does not constitute a contract or guarantee regarding payment or results. Coverage decisions and the administration of benefits are subject to all terms, conditions, exclusions and limitations of the coverage documents (e.g., evidence of coverage, certificate of coverage, policy, contract of insurance, etc.), as well as to state and federal requirements and applicable Health Plan-level administrative policies and procedures.

This clinical policy is effective as of the date determined by the Health Plan. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a

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discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall govern. The Health Plan retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

This clinical policy does not constitute medical advice, medical treatment or medical care. It is not intended to dictate to providers how to practice medicine. Providers are expected to exercise professional medical judgment in providing the most appropriate care, and are solely responsible for the medical advice and treatment of members. This clinical policy is not intended to recommend treatment for members. Members should consult with their treating physician in connection with diagnosis and treatment decisions.

Providers referred to in this clinical policy are independent contractors who exercise independent judgment and over whom the Health Plan has no control or right of control. Providers are not agents or employees of the Health Plan.

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Note: For Medicaid members, when state Medicaid coverage provisions conflict with the coverage provisions in this clinical policy, state Medicaid coverage provisions take precedence. Please refer to the state Medicaid manual for any coverage provisions pertaining to this clinical policy.

Note: For Medicare members, to ensure consistency with the Medicare National Coverage Determinations (NCD) and Local Coverage Determinations (LCD), all applicable NCDs, LCDs, and Medicare Coverage Articles should be reviewed prior to applying the criteria set forth in this clinical policy. Refer to the CMS website at <http://www.cms.gov> for additional information.

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