

Vitamin D Practice Guidelines



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		Desirable 20 ng/ml IOM recommendations		Desirable 30 ng/ml ES Recommendations at risk for D deficiency			
Life stage group	AI	EAR	RDA	UL	Daily requirement		UL
Infants							
0 to 6 months	400 IU (10g)			1,000 IU(25 μg)) 400–1,000	IU	2,000 IU
6 to 12 months	400 IU (10 g)			1,500 IU(38 μg)) 400–1,000	IU	2,000 IU
Children							
1–3 yr		400 IU (10 μg)	600 IU(15 μg)	2,500 IU(63 μ	g) 600–1,000) IU	4,000 IU
4-8 yr		400 IU (10 μg)	600 IU(15 μg)	3,000 IU(75 μ	g) 600–1,000) IU	4,000 IU
9-13 yr		400 IU (10 μg)	600 IU(15 μg)	4,000 IU(100 µ	ug) 600–1,000) IU	4,000 IU
14- 18 yr		400 IU (10 μg)	600 IU(15 μg)	4,000 IU(100 µ	ug) 600–1,000	บเบ	4,000 IU

Institute of Medicine (IOM) Ross et al. J Clin Endocrinol Metab 96: 53–58, 2011 Endocrine Society (ES) Holick et al. J Clin Endocrinol Metab 96 (7):1-, 2011 Vitamin D Intakes Recommended by the IOM and the Endocrine Society (ES) Practice Guidelines Committee / Pregnancy and Lactation

	ION	/ recommend	ES recommendations for patients at risk for vitamin D deficiency		
Life stage group	EAR	RDA	UL	Daily	UL
Pregnancy	-	2			2
14-18 yr	400 IU (10 µg)	600 IU (15 μg)	4,000 IU (100 μg)	600–1,000 IU	4,000 IU
19- 30 yr	400 IU (10 µg)	600 IU (15 μg)	4,000 IU (100 µg)	1,500–2,000 IU	10,000 IU
31-50 yr	400 IU (10 µg)	600 IU (15 µg)	4,000 IU (100 µg)	1,500–2,000 IU	10,000 IU
Lactation	400 IU (10 µg)	600 IU (15 µg)	4,000 IU (100 µg)		
14-18 yr	400 IU (10 µg)	600 IU (15 μg)	4,000 IU (100 µg)	600–1,000 IU	4,000 IU
19- 30 yr	400 IU (10 µg)	600 IU (15 μg)	4,000 IU (100 µg)	1,500–2,000 IU	10,000 IU
31-50 yr	400 IU (10 µg)	600 IU (15 µg)	4,000 IU (100 µg)	1,500–2,000 IU	10,000 IU

Vitamin D intakes Recommended by the Practice Guidelines Committee / Adults



	ION	l recommenda	ES recommendations for patients at risk for vitamin D deficiency				
Life stage group	EAR	RDA	UL	Daily	UL		
Adults							
19-30 yr	400 IU (10 µg)	600 IU (15 µg)	4,000 IU (100 µg)	1500–2,000 IU	10,000 IU		
31-50 yr	400 IU (10 µg)	600 IU (15 µg)	4,000 IU (100 µg)	1,500–2,000 IU	10,000 IU		
51-70 yr	400 IU (10 µg)	600 IU (15 µg)	4,000 IU (100 µg)	1,500–2,000 IU	10,000 IU		
> 70 yr	400 IU (10 µg)	600 IU (20 µg)	4,000 IU (100 µg)	1,500–2,000 IU	10,000 IU		

Lebanese Osteoporosis Guidelines 2003 and 2007Universal Recommendations



- Maintain a physically active lifestyle with adequate exposure to sunlight
- Avoid smoking and high alcohol intakes
- Maintain dietary calcium intake around 1.5 gm of elemental calcium in PM estrogen deficient women or men >65 years and vitamin D intake of 600 to 800 IU/day
- Provide calcium and vitamin D supplementation in the elderly

Lebanese Osteoporosis Guidelines 2003 and 2007 Universal Recommendations

- Avoid a low weight <60 kg in men or 50 kg in women or a low Body Mass Index BMI<20 kg/m2.
- The prevention of osteoporosis begins with optimal bone mass acquisition during growth. Factors hindering bone mass acquisition, such as malnutrition and inadequate Calcium or Vitamin D intake, should be considered, identified and addressed during childhood.
- Address known factors that stimulate bone resorption or inhibit bone formation, including hypogonadism, primary hyperparathyroidism, hyperthyroidism and hypercortisolism.
- Develop fall prevention awareness and programs in the elderly.
- Hip protection and/or soft floor covering in elderly environment.

Lebanese Osteoporosis Guidelines 2012



- Elemental calcium at 1200 mg/day
 - Intake > 1500 mg/day has limited benefit and may increase risk of kidney stones or CV disease
- Vitamin D3 800-1000 IU/day
 - Many patients will need more to achieve serum 25-OH-D level of 30 ng/ml or higher
- Regular weight-bearing exercise
- Fall prevention
- Avoid tobacco use and excess alcohol intake

Lebanese Vitamin D Practice Guidelines 2013

- To most consistently improve clinical outcomes such as fracture risk, fall risk and minimize bone loss an optimal serum level of 25-hydroxyvitamin D is probably above 75 nmol/L (30 ng/ml)-Desirable Range 30-60 ng/ml
- For most Lebanese supplementation is needed to achieve this level.
- Exposure to natural sunlight, when used in moderation (avoiding sunburn), can contribute to vitamin D sufficiency in the summer in subjects who do seek such exposure
- The recommended vitamin D intake is 15–25 µg (600–1000 IU) daily for childrenadolescents and low-risk adults under 50 years of age, and 20–50 µg (1000–2000 IU) for high-risk and older adults, with potential for consideration of higher doses.
- Doses up to 50 μg (2000 IU) are safe and do not require monitoring, but if higher doses are sometimes needed, monitoring is appropriate.

Lebanese Vitamin D Practice Guidelines 2013



- Serum 25-hydroxyvitamin D should NOT be measured in routine practice
- It should be measured in following instances:
- High risk individuals: such as those with osteoporosis on pharmacologic therapy, with fractures
- Conditions known to affect vitamin D metabolism or action
 - Steroids, anticonvulsants, malabsorption, by-pass surgery, cirrhosis (need to order both 25-D and 1,25D level), patients with high PTH levels
- In individuals on doses > 2000 IU /day



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- Once a stable dose-level are reached routine monitoring of 25-OHD level is not recommended
- Recommend using a laboratory with rigorous QA measures (See technical recommendations and guide Dr Daher)
- Recommend against using 1,25 (OH)₂ D except in select conditions: such as disorders of vitamin D and PO4 metabolism.
- Research is needed on impact of low D levels on multiple non-classical outcomes

Hypovitaminosis D in EMR



- Hypovitaminosis D is strikingly common in "apparently healthy" individuals, lowest levels are in the Middle East-silent precursor of NCDs
 - Predictors age, gender, veiling, season, parity, SES
 - Genetic polymorphisms in metabolic pathway may contribute: CYP21R
- This has a negative impact on musculoskeletal health
 - Vit D status inversely correlates with PTH (R=-0.2 to-0.37) and directly with bone mass (R=0.2-0.35)-Elderly with OP have lower 25-OHD, and higher levels of PTH
 - 25-OHD level positively correlates with bone mass
 - RCT show that Ca/D (> 700IU/day) reduce falls and fracture risk.
 - Deleterious impact on maternal & neonatal health is anticipated but not established (*Morley et al. JCEM 2006, Javaid et al. Lancet 2006*)
- There may be an effect on non-classical outcomes need for RCT
 - Cardiovascular: for eg Pre-ecclampsia (Bodnar JCEM 2007)
 - Insulin resistance & DM including gestational diabetes: (Pittas Diabetes Care 2007)
 - Infections and auto-immune disorders
 - Cancer

Hypovitaminosis D in EMR



- Assay variation somewhat limits comparability across studies and is a major obstacle in advancing field-Need for QA programs
- Calcium intake does, and VDR polymorphisms may, modulate effect of hypovitaminosis D on major outcomes
- Recommendations in western populations need to be adjusted upwards in Eastern Mediterranean Region
- Evidence lacks to define optimal dose in:
 - pregnant and breast-feeding women, infants, pre-pubertal children and nonclassical outcomes worldwide and for the elderly in EMR.
- Meanwhile, suggest increments in recommended doses
 To achieve desirable 25-OHD level 25-30 ng/ml