Parathyroid glands

(2)

Dr. Zaynab Alourfi PhD أ.م.د.زينب العرفي

وظيفة: ما هي المقاربة الصحيحة ؟

لمريض لديه بوال وسهاف وحصيات كلوية وعقدة مثبتة للسيستاميبي تحاليله المخبرية

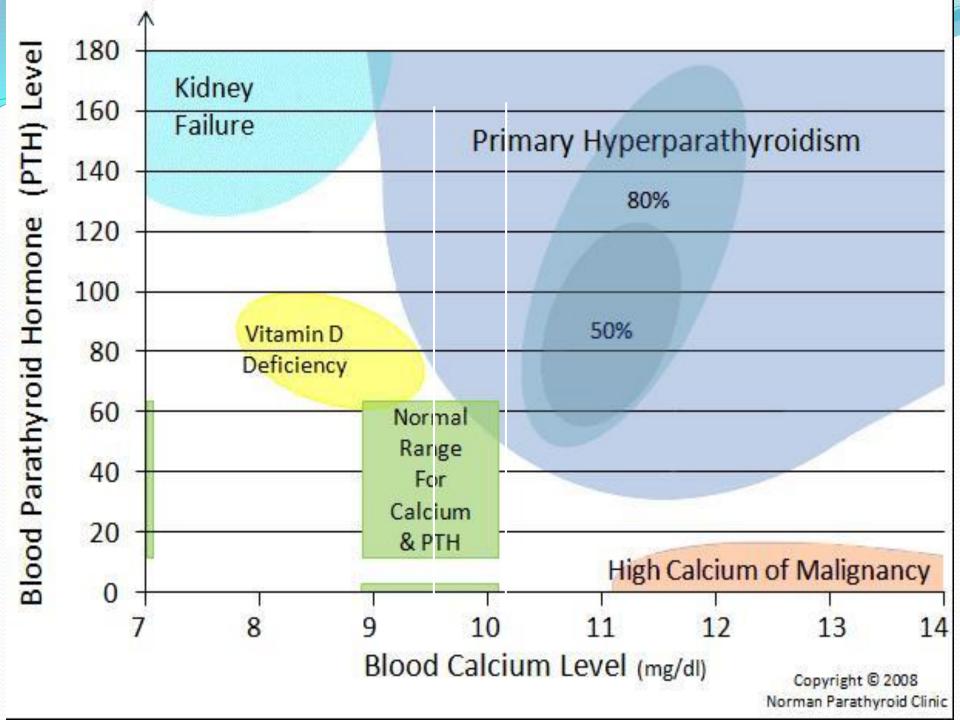
Ca = 9,5 mg/dl (normal: 8.5-10.5)

Albumin = 4 g/dl

PTH = 38 pg/ml (normal: 10 - 65)

تتوافق الحالة مع

Normocalcemic Primary hyperparathyroidism

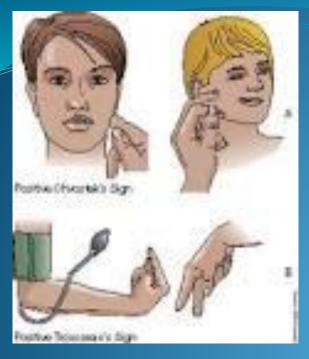


- نقاش حامي الوطيس بين وفاء عن فريق الطالبات وأسامة عن فريق الطلاب حول تحاليل لفادي (عمره ٢٥ سنة) أجراها من أجل التوظيف وهي كما يلي: السكر والكرياتنين والتعداد العام والصيغة طبيعية لكن الكالسيوم أقل من الطبيعي
 - الكالسيوم = λ, ξ ملغ/دل (ط λ, δ λ, δ) ما رأيك؟

الكالسيوم المعدّل

الكالسيوم المعدّل = الكالسيوم الحالي + 0.8 + 0.8 (4-الألبومين)

- أثناء تطبيق ما درسه طالب سنة رابعة عن علامات نقص الكلس جرب استقصاء علامة شفوستك على أخيه فكانت ايجابية اضطرب وسأل ما عليه أن يفعل؟
 - أ- يعاير الكالسيوم في الدم
 - ب- يعاير هرمون جارات الدرق
 - ت- يطلب ايكو درق
 - <u>-</u>_



Hypocalcemia







Trousseau's Signs

- Chvostek sign: Specificity is low (25 % of normal indiveduals have a mild Chvostek sign)
- Trosseau sign is more specific
 (1% to 4% of normal indiveduals have positive)

Causes of Hypocalcemia

Hypoparathyroidism

Surgical/ Idiopathic/ Neonatal

Familial/ Autoimmune

Deposition of metals (iron, copper,

aluminum)

Postradiation/ Infiltrative

Functional (in hypomagnesemia)

Resistance to PTH action

Pseudohypoparathyroidism

Renal insufficiency

Medications that block osteoclastic bone

resorption

Plicamycin, Calcitonin, Bisphosphonates

Failure to produce 1,25(OH)₂D normally

Vitamin D deficiency

Hereditary vitamin D-dependent rickets, type 1 (renal 25-OH-vitamin D 1alphahydroxylase deficiency)

Resistance to 1,25(OH)₂D action

Hereditary vitamin D-dependent rickets, type 2 (defective VDR)

Acute hyperphosphatemia

Crush injury with myonecrosis

Rapid tumor lysis

Parenteral phosphate administration

Excessive enteral phosphate

Oral (phosphate-containing antacids)

Phosphate-containing enemas

Acute pancreatitis

- Citrated blood transfusion
- Rapid, excessive skeletal mineralization

Hungry bones syndrome

Osteoblastic metastasis

Vitamin D therapy for vitamin D deficiency

Miss L is a 16 year old woman with no significant past medical history, who is brought to the ER by her mother after she noted her to be acting bizarrely for the past several weeks. Thought to be actively psychotic, a psychiatry consult is asked to see the patient, who recommends checking routine labs:

```
Sodium – 142 meq/L (136–146 meq/L)
Potassium – 4.1 meq/L (3.5–5.0 meq/L)
Magnesium – 2.3 mg/dL (1.5–2.3 mg/dL)
Calcium (total) – 6.9 mg/dL (8.7–10.2 mg/dL)
Phosphate – 4.4 mg/dL (2.5–4.3 mg/dL)
Albumin – 4.2 g/dL
Creatinine – 0.8 mg/dL
```

```
- ما هو التشخيص الأرجح مما يلي وما هي الخطوة التالية الدين المنطوق التالية المنطوق كانب عبد اللازمة كانب كانب المنطوق كانب اللازمة للتشخيص؟ - ما هي الاجراءات اللازمة للتشخيص؟
```

Pseudohypoparathyroidism (PHP)

1- **PHP type IB** is a disorder of isolated resistance to PTH: biochemical features of hypocalcemia, hyperphosphatemia, and secondary hyperparathyroidism.

2- <u>PHP type IA</u> has, in addition to biochemical features, a characteristic somatic phenotype known as **Albright hereditary osteodystrophy** (AHO).

Short stature, round face, short neck, obesity, brachydactyly (short digits), shortened metatarsals, subcutaneous ossifications, and often reduced intelligence. Because of shortening of the metacarpal bones—most often the fourth and fifth metacarpals—affected digits have a dimple, instead of a knuckle, when a fist is made. Primary hypothyroidism is frequently seen. Less commonly, these patients have abnormalities of reproductive function—oligomenorrhea in females and infertility in males due to primary hypogonadism.



Pseudo Pseudohypoparathyroidism (PPHP)

Certain individuals in families with PHP inherit the somatic phenotype of AHO without any disorder of calcium metabolism.



• بعد استئصال درق تام حدث لدى المريض نوبة تكزز، عولجت بالكالسيوم وريدياً وكذلك فموياً. تحسن المريض وطلب منك تخريجه. ما رأيك

Treatment of Hypocalcemia

Acute Hypocalcemia

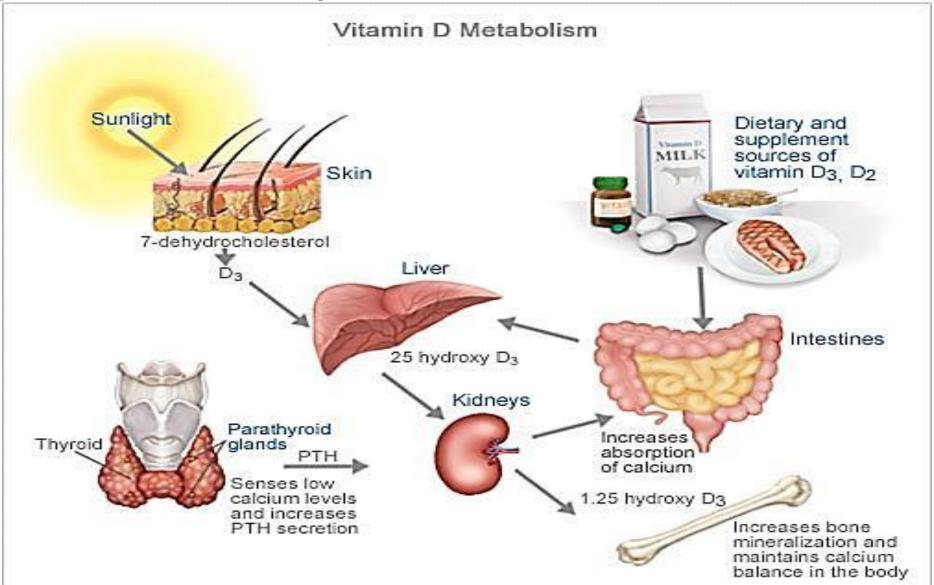
Patients with tetany should receive **intravenous calcium** as calcium chloride (272 mg calcium/10 mL), calcium gluconate (90 mg calcium/10 mL), or calcium gluceptate (90 mg calcium/10 mL). Approximately 200 mg of elemental calcium can be given over several minutes. The patient must be observed for stridor and the airway secured if necessary.

Oral calcium and a rapidly acting preparation of vitamin D should be started. If necessary, calcium can be infused in doses of 400 to 1000 mg/24 h until oral therapy has taken effect.

Intravenous calcium is irritating to the veins and is best infused into a large vein or through a central venous catheter.

Vitamin D Deficiency

Vitamin D synthesis and metabolism



Vitamin D sources:

• Natural sources of vitamin D :

Sun Exposure:

- Ultraviolet (UV) B (290–320) nanometers penetrates uncovered skin and converts cutaneous 7dehydrocholesterol to previtamin D₃
- The amount of sunlight needed to synthesize adequate amounts of vitamin D varies, depending upon the person's age, sun exposure, season, time of day, length of day, cloud cover, smog, skin melanin content, sunscreen and underlying medical problems.

 Vitamin D promotes <u>calcium</u> absorption in the gut and maintains adequate serum calcium and phosphate concentrations.

Needed for bone growth and bone remodeling.

Vitamin D has other roles in the body, including modulation of cell growth, neuromuscular and immune function, and reduction of inflammation.

Diagnosis of vitamin D deficiency

- A normal level of vitamin D is defined as a 25(OH)D concentration greater than 30 ng/mL (75 nmol/L).
- Vitamin D <u>insufficiency</u> is defined as a 25(OH)D concentration of 20 to 30 ng/mL (50 to 75 nmol/L).
- Vitamin D <u>deficiency</u> is defined as a 25(OH)D level less than 20 ng/mL (50 nmol/L).

Vitamin D deficiency

Causes:

- Lack of vitamin D in the diet, often in conjunction with inadequate sun exposure
- Inability to absorb vitamin D from the intestines
- Inability to process vitamin D due to kidney or liver disease

Groups at risk:

- Breastfed infants
- Older adults
- People with limited sun exposure
- People with inflammatory bowel disease and other conditions causing fat malabsorption
- People who are obese or who have undergone gastric bypass surgery
- Chronic kidney disease.
- Liver disease.

Recommended Dietary Allowances (RDAs) for Vitamin D [1] Age Male Female Pregnancy Lactation o-12 months* 400 IU 400 IU (10 mcg) (10 mcg) 600 IU 600 IU 1-13 years (15 mcg) (15 mcg) 600 IU 600 IU 600 IU 600 IU 14-18 years (15 mcg) (15 mcg) (15 mcg) (15 mcg) 600 IU 600 IU 600 IU 600 IU 19-50 years (15 mcg) (15 mcg) (15 mcg) (15 mcg) 600 IU 600 IU 51-70 years (15 mcg) (15 mcg) 800 IU 800 IU >70 years

Adequate Intake (AI): established when evidence is insufficient to develop an RDA and is set at a level assumed to ensure nutritional adequacy.

(20 mcg)

(20 mcg)

Tolerable Upper Intake Levels (ULs) for Vitamin D

Age	Male	Female	Pregnancy	Lactation
o-6 months	1,000 IU (25 mcg)	1,000 IU (25 mcg)		
7–12 months	1,500 IU (38 mcg)	1,500 IU (38 mcg)		
1-3 years	2,500 IU (63 mcg)	2,500 IU (63 mcg)		
4–8 years	3,000 IU (75 mcg)	3,000 IU (75 mcg)		
≥9 years	4,000 IU (100 mcg)	4,000 IU (100 mcg)	4,000 IU (100 mcg)	4,000 IU (100 mcg)

<u>Tolerable Upper Intake Level (UL):</u> maximum daily intake unlikely to cause adverse health effects

الوظيفة: ملء الجدول التالي

World Health Organization Definition of Bone Mass-Dual-energy X-ray			
absoeptiometry (DEXA)			
Normal bone mass	T score		
Low bone mass	T score		
Osteoporosis	T score		
Established	T score		
osteoporosis			

