Pharmacology of drugs used in calcium & vitamin D disorders





Objectives

By the end of lecture, the students will be able to :

Recognize the common drugs used in calcium & vitamin D disorders

Classify them according to sources & Pharmacological effects

Detail the pharmacology of each drug, regarding, Mechanism, clinical utility in affecting calcium & vitamin D



BONE

Is a dynamic organ undergoes continuous remodeling process involving resorption of old bone by osteoclast & formation of new bone by osteoblast.

The dominant site of calcium storage in the body is bone, which contains nearly 99.9% of body calcium.

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The principal factors involved in calcium metabolism & bone remodeling are :
Parathyroid hormone (PTH)
Teriparatide
vitamin D
calcitonin

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PTH and vitamin D play central roles in the regulation of bone metabolism

The target tissues of Vit D and calcium

Bone (Absorption and resportion)

kidney (Reabsorption)

Intestine.(Calcium absorption)

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Parathyroid Hormone

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Is released from the parathyroid gland in

response to low plasma Ca²⁺ level

Effects of of PTH

BONE

Mobilization of Ca²⁺ and PO₄³⁻ from bone

In response to hypocalcemia , PTH stimulates osteoclast cells to increase the outward flux of calcium to restore serum calcium level.



↑ Ca²⁺ reabsorption

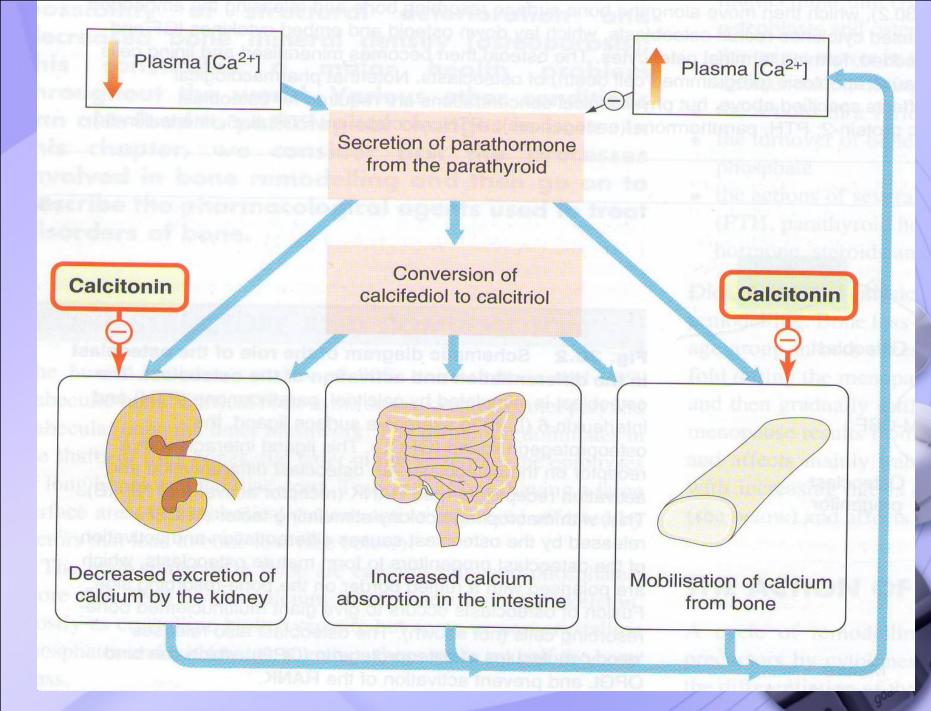
▲ formation of calcitriol which is the active form of vitamin D

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GIT

↑ absorption of Ca²⁺



Daily, intermittent administration of PTH for 1 to 2 hours / day leads to a net stimulation of bone formation.

Continuous exposure to elevated PTH leads to bone resorption and risk of fracture.

RESPONSE TO PTH

PTH

Intermittent

↑ osteoblast number/function

bone formation

bone mass/strength

Continuous

↑ osteoclast

↑ bone resorption

↑ serum Ca⁺

Clinical uses

Treatment of severe osteoporosis

Resistance cases failed to response to other medications

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Teriparatide

Synthetic polypeptide form of PTH (PTH analogue).

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Given, once / daily /subcutaneous injection

Therapeutic uses of Teriparatide

should not be used routinely due to carcinogenic effects. **Use in severs osteoporosis or patients not** responding to other drugs. **For treatment of osteoporosis in people** who have a risk of getting fracture increase bone mass & strength) Good for postmenopausal osteoporosis.

Side effects

Carcinogenic effect (osteosarcoma)

Diarrhea, heart burn, nauseaheadache, leg cramps

Hypotension when standing.

Elevated serum calcium can occur in some cases can lead to kidney stones

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Contraindications

Should not be used by people with increased risk for bone tumors (osteosarcoma) including : **People with paget's disease of bone People who had radiation treatment** involving bones **Children not recommended**

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Vitamin D

Cholecalciferol (vitamin D3) skin
Ergocalciferol (vitamin D2).plants
Vitamin D2 is the prescription form of vitamin D & is also used as food additive (milk).

Vitamin D3 is usually for vitamin D- fortified milk & foods & also available in drug combination products.

Vit D2 and Vit D3 have equal biological activities

Sources of Vitamin D

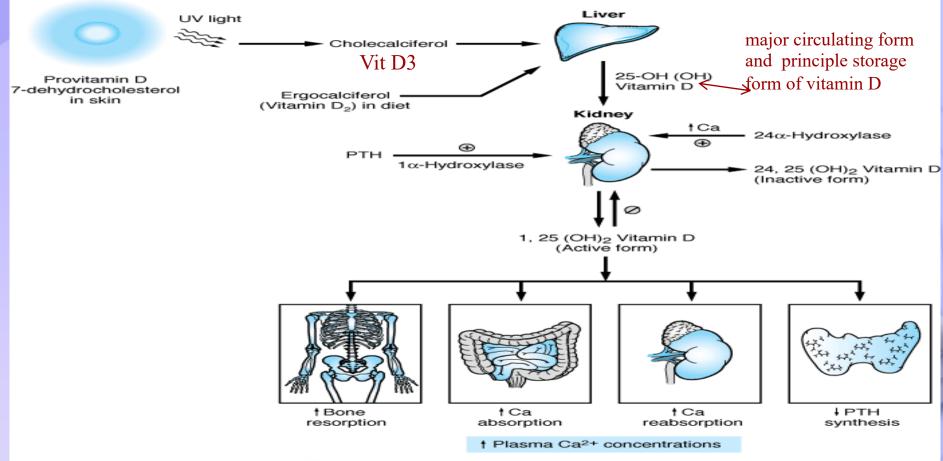


milk egg yolk fish oils

Note both D2 and D3 travel to the liver and then converted to active from in the kidneys see next slide

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Calcium and Vitamin D



Source: Molina PE: Endocrine Physiology, 3rd Edition: http://www.accessmedicine.com

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Vitamin D increases bone resorption, increases Ca^{2+} absorption from the intestine, increases renal Ca^{2+} reabsorption, and decreases the production of PTH by the parathyroid glands. The overall effect of vitamin D is to increase plasma Ca^{2+} concentrations.

Sunshine: Cholecalciferol (D3))

Generated in the skin from 7dehydrocholesterol by the action of ultraviolet radiation (sunshine).

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Vitamin D Metabolism

The initial transformation of D3 occurs in **liver** to (calcifediol) the main storage from of Vit D in our body.

In the **kidney** : **parathyroid hormone** stimulates the formation of active form of vitamin D (**calcitriol**)

Effects of Vitamin D

Bone : Activation of osteoblast cells

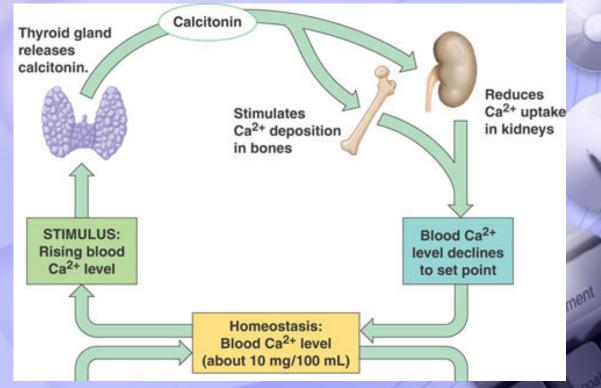
Kidney : Increased reabsorption of Ca²⁺ & PO₄.

GIT : Increased absorption of Ca²⁺

Calcitonin

produced by the parafollicular cells (C cells) of the thyroid gland. It is released when there is an elevated level of Ca²⁺ in the blood.

Calcitonin does not appear to be critical for the regulation of calcium homeostasis even if thyroid gland is removed



Effects of calcitonin

Bone : Decrease bone resorption by inhibiting osteoclast activity

Kidney : Decreases reabsorption of Ca²⁺ & PO₄, thus increasing their excretion

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Clinical uses of Calcitonin (it has lower efficacy compared to other drugs) **Osteoporosis (major indication;** alternative to other drugs) Hypercalcemia (short-term treatment of hypercalcemia of malignancy), Pagets disease **Routes of administration** S.C, Nasal spray

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Adverse effects

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Nausea
local inflammation (injection)
Flushing of face & hands
Nasal irritation

Thank You.

