

DISCUSSIÓ

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Si el ió citrat no pot entrar al reticle cristal·lí de la hidroxapatita, perquè és més gros que el tamany del cristall ossi i el 90 % d'aquest citrat està a l'ossada (Taylor T.G. Biochim.Biophys. Acta, 148, (short commun), 1959) - (Brecevic L. i Furedi-Milhofer J. Calcif.Tiss.Int. 28:131, 1979), què hi fa ?.

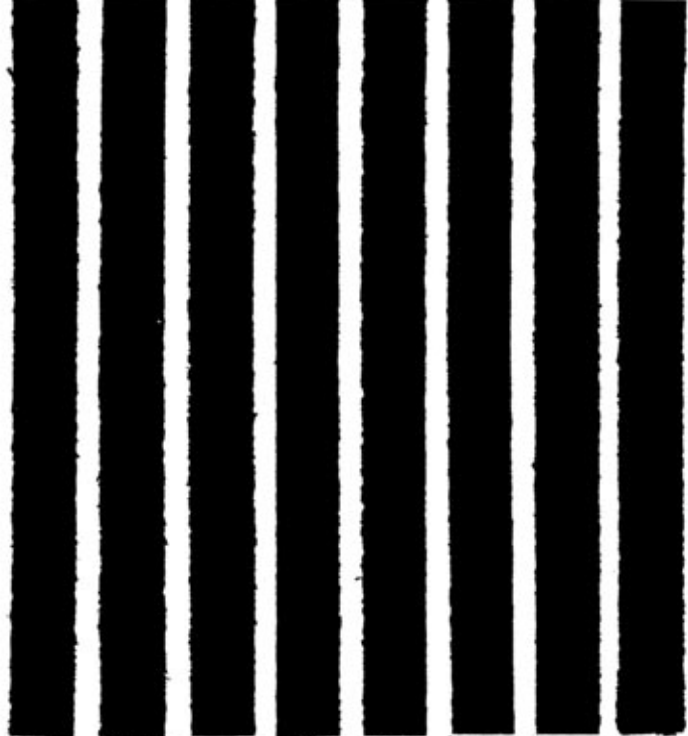
Si diverses malalties siguin infeccioses, metabòliques o genètiques i àdhuc la ingravitació, quecientment tenen una tendència específica a fer víctima l'os com a premonició de la seva actuació de conjunt (Un dels grans problemes de la pèrdua d'os és la seva universalitat, Garn S.M. i al. Fed.Proc. 26, 6:1729, 1967) - (La heterogeneïtat confusa de moltes malalties metabòliques de l'os prové de l'haver-hi molts camins per arribar a un resultat comú, Burr D.B. Am.J.Anat. 186:216, 1989), no és propici pensar que aquesta posició de conjunt que tendeix a la convergència en l'índole dels trastorns ossis, provocats per les situacions citades sigui regida, més endins, per un constituent - l'àcid cítric - que orienti la finalitat coincident donada la seva omnipresència a l'os ?.

I a més, per què no, sabent que la majoria de substàncies que custodien la modelació i la remodelació de l'os, directa o indirectament, demostren tenir un fort atraïment per l'àcid cítric que es palesa movent la seva quantia a l'organisme ?.

Si apleguem les tres interrogacions es pot córrer el risc de suposar que l'àcid cítric, ja que hi és i per no perdre peculiaritat, davant de la impossibilitat d'entrar a la intimitat de l'os opti per romandre al seu embolcall d'hidratació feït de conductor diferenciat que tria les substàncies que han de passar o sortir i a la inversa - rutina perenne - del cristall.

Perquè aquesta suggerència sigui consentida s'ha d'admetre que que pugui haver-hi uns intersticis habituals o circumstancials, sempre específics, que donin pas a les matèries d'intercanvi. Com afegiment, " tots els cristalls d'hidroxiapatita contenen un gran nombre de llocs vacants i intersticis pels ions " (Schottky) - i " els defectes de cristal·lització molt freqüents en les apatites fan que augmenti la solubilitat " (Arens i al. Caries 11:186, 1977) , finalment, " els ions citrat ataquen la superfície de l'os mineral, desplaçant simultàniament una quantitat equivalent de ions fosfat " (Pak i al. Calcif.Tiss Res. 4:69, 1967).

citrat sèrum



PTH

vitD

F.alc

GH

vitC

Tir

Estr

F

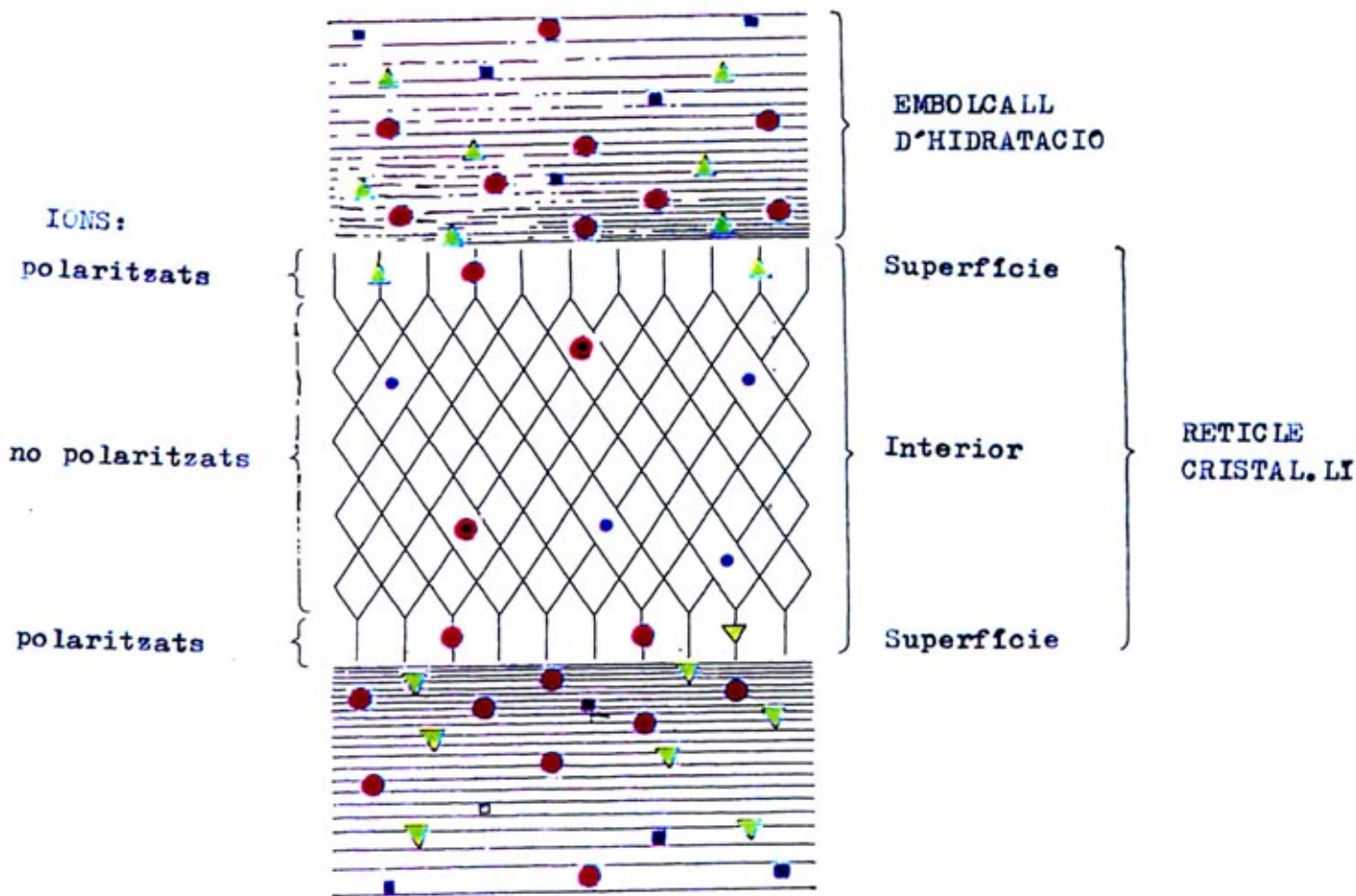
Cal

Cort

Gluc

Ins





- Llacunes del reticle
- Estronci⁺⁺ o Radi⁺⁺ a l'interior del reticle cristal.li, ocupant el lloc del calci
- ▲ Sodi⁺, citrat⁼, carbonat⁼, fluor⁻, a la superfície del cristall en substitució del Ca⁺ o del fosfat⁼ polaritzat
- Potasi⁺, Clor⁻, solament a l'embolcall d'hidratació

EPÍLEG.

" la disminució local del pH (pot ésser per certs efectes específics del ió citrat) seria suficient per a explicar la solubilització del mineral.

G. Vaes Symp.Int. sur l'Osteomalacie, pag.30, Tours (1.965).

CONCLUSIONS

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En els animals tractats s'evidencia:

- a) Un augment significatiu de l'àcid cítric sèric.
- b) Que les fosfatases alcalines disminueixen proporcionalment a l'augment de l'àcid cítric.
- c) Que l'anatomia estadística de les vísceres fetge, ronyó, melsa, cor i pulmó no presenta cap alteració a tenir en compte.
- d) Que l'anàlisi hemàtic, tret de les variacions de l'àcid cítric i de les fosfatases alcalines està dins dels marges tolerats.
- e) Que no hi ha desdibuixament dels mitocondris de cor.
- f) Un major creixement, en llargada, dels ossos llongs.
- g) Un remodelament subperiòstic aberrant.
- i) La massa òssia lamel·lar en franca remoció del mineral.

It seems:

Que la inhibició malònica respecte a la repercussió en el remodelament ossi ha seguit un curs, amb tots els alligonaments a fer-hi i les correccions necessàries, d'aproximació " lineal " al resultat buscat.

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