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跳到**CASE REPORT** - **Tumor-induced osteomalacia** (TIO) is a rare paraneoplastic syndrome that presents with hypophosphatemia, myopathy, bone pain, and fractures. ... The systemic bone demineralization **caused** by renal phosphate wasting is often **due to** excessive production of FGF23. Abstract · DISCUSSION

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Name of Journal: *World Journal of Clinical Cases*

Manuscript NO: 46113

Manuscript Type: CASE REPORT

**Oncogenic osteomalacia caused by a phosphaturic mesenchymal tumor of
the femur: A case report**

Dong Tang, Xiao-Man Wang, Yong-Sheng Zhang, Xiao-Xiao Mi

Abstract

BACKGROUND

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Oncogenic Osteomalacia Caused by a Phosphaturic ...

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A previous report presented a young male patient with **oncogenic osteomalacia** caused by a tumor arising from the left side of the mandible, which was also found in the oral cavity 5). In the present case, however, the tumor is not believed to have originated in the mandible since it was clearly separated from the bone, although it destroyed some alveolar bones.

Cited by: 24

Author: In Myung Yang, Yong Koo Park, Yong Jun...

Publish Year: 1997

Phosphaturic mesenchymal tumor ... - PubMed Central (PMC)

<https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2783479>

Tumor-induced osteomalacia (TIO) is a rare paraneoplastic syndrome that results in renal phosphate wasting with hypophosphatemia. In most cases, the underlying cause of TIO is a small **mesenchymal** neoplasm that is often difficult to detect, resulting in delayed diagnosis. One such neoplasm is the ...

Cited by: 31

Author: Victoria L. Woo, Regina Landesberg, Erik...

Publish Year: 2009

Oncogenic osteomalacia caused by a phosphaturic ...

<https://thejns.org/spine/abstract/journals/j-neurosurg-spine/10/4/article-p329.xml> ▾

Phosphaturic mesenchymal tumors that cause the paraneoplastic syndrome known as **oncogenic osteomalacia** are rare. The authors report on the case of a 57-year-old man with a history of osteomalacia and in whom was diagnosed a thoracic spine tumor at the T-4 level.

A Rare Case of Phosphaturic Tumor/Oncogenic Osteomalacia ...

www.jocr.co.in/wp/2019/01/10/2250-0685-1304-fulltext ▾

Jan 10, 2019 · Keywords: Phosphaturic mesenchymal tumor, proximal femur, intralesional extended curettage Introduction Phosphaturic mesenchymal tumors (PMTs) are very rare tumors which are frequently associated with tumor-induced osteomalacia (TIO), a paraneoplastic syndrome that manifests as renal phosphate wasting [1].

Oncogenic osteomalacia associated with mesenchymal ...

<https://jmedicalcasereports.biomedcentral.com/articles/10.1186/1752-1947-6-181> ▾

Jul 02, 2012 · Tumor-induced osteomalacia is a paraneoplastic syndrome of hypophosphatemia. Osteomalacia causes multiple bone fractures and severe pain. We report the case of a 57-year-old Japanese man with tumor-induced osteomalacia associated with a middle cranial fossa bone tumor. The tumor was successfully resected by using a middle fossa epidural approach.

Cited by: 2

Author: Isao Chokyu, Kenichi Ishibashi, Takeo Go...

Publish Year: 2012

Ethmoid tumor and oncogenic osteomalacia: Case report ...

<https://www.sciencedirect.com/science/article/pii/S187972961830098X>

Oncogenic osteomalacia is a very rare disease usually caused by a phosphaturic mesenchymal tumor, particularly the “mixed connective tissue type”, secreting FGF-23 hormone. Objective The authors report a case of **ethmoid tumor** associated with **oncogenic osteomalacia** and discuss management