Distant lung metastases caused by a histologically benign phosphaturic mesenchymal tumor

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Summary

Tumor-induced osteomalacia (TIO) is a rare form of hypophosphatemia usually caused by phosphaturic mesenchymal tumors (PMTs); the biologic behavior of PMTs is under investigation. Herein we present a case of TIO with a protracted course over 12 years leading to a fatal outcome. A 39-year-old man presented with weakness in 2004 and was found to have decreased serum phosphorus, phosphaturia and low levels of 1,25-dihydroxyvitamin D3. Four years later he developed a painful left calf mass. The lesion was resected, but recurred causing extreme pain and dysfunction. Radiological examination showed a large cluster of soft tissue tumors affecting all the muscle compartments of the calf and a smaller lesion inside the metaphysis of the tibia. Above-knee amputation was performed. Histological examination of all lesions showed a cellular spindle cell neoplasm with variously sized vessels, wide vessel-like spaces and scattered deposits of calcified extracellular material. The tumor infiltrated skeletal muscles, subcutaneous fat and the proximal end of the fibula. The tibial lesion had identical histology. Three years after the amputation the patient presented with cough and dyspnea. Radiological examination, followed by an open biopsy, showed that there were multiple metastatic nodules of PMTs in both lungs. Shortly after the diagnosis the patient died. This case illustrates that even benign cases of PMTs may lead to a fatal outcome and the classification of PMTs into benign and malignant should be reassessed in order to correspond to its biological behavior.

Learning points:

- PMTs, aside from having locally aggressive behavior, may metastasize and cause death
- PMTs may behave aggressively despite 'benign' histological findings
- Accurate diagnosis of tumor-induced osteomalacia and patient management require a multidisciplinary approach

Background

Tumor-induced osteomalacia (TIO) is a rare form of hypophosphatemia caused by phosphaturic mesenchymal tumors (PMTs) located in the bone or soft tissue. The diagnosis is usually based on the biochemical profile of

severely low levels of phosphate (PO_4) associated with phosphaturia and low or inappropriately normal levels of 1,25-OH-vitamin D. Increased levels of fibroblast growth factor 23 (FGF 23) that is excreted by the tumor is the

pathophysiological factor responsible for phosphaturia. Localization and resection of the tumor can cure completely osteomalacia. The majority of PMTs are benign, and malignancy or malignant transformation of PMTs with distant metastasis is very uncommon (1). We describe herein the follow-up of a previously reported case (2) with a histopathologically benign phosphaturic mesenchymal tumor that caused severe hypophosphatemia and multiple recurrences, and finally led to amputation of the affected limb. Although there was no evidence of histologic malignancy, 13 years after the initial diagnosis and 2 years after amputation the patient developed fatal lung metastases.

Case presentation

This is a follow-up presentation of a case that was part of our study registered on Clinical trials.gov (protocol number NCT01660308) and approved by the Scientific Review Board of AHEPA University Hospital (protocol number 51741).

Our patient presented with a medical history of TIO caused by a histopathologically benign PMTs with a locally aggressive behavior, which led to multiple recurrences (2).

On his first admission to our center the patient had severe hypophosphatemia (1.8 mg/dL, RR: 2.7–4.5 mg/dL) and elevated 24 h urine phosphate (1797.4 mg/24 h, RR: 400–1300 mg/24 h), elevated alkaline phosphatase (ALP) levels (300 IU/L, RR: 40–150 IU/L), elevated parathyroid hormone (PTH) (19.4 pmol/L, RR: 1.8–6.03 pmol/L), and serum Ca^{2+} levels (10.8 mg/dL, RR: 8.2–10.6 mg/dL), and low to normal 1,25 (OH)₂-vitamin D levels (18 pg/mL, RR: 18–24 pg/mL) and 25-OH-vitamin D (20.9 ng/mL, RR: 40–100 ng/mL), as previously described (2). The ratio of maximum rate of renal tubular reabsorption of phosphate to glomerular filtration rate (TmPO₄/GFR) as determined by the Walton and Bijvoet nomogram was noted to be low at 0.3 mmol/L (0.8–1.4) confirming the excessive loss of phosphate in the urine (2). There were no signs of glycosuria, aminoaciduria or proteinuria. In our initial approach of the patient, the medical team decided resection of the tumor mass which was located in the periphery of the head and the upper third of the left fibular diaphysis due to the patient's medical record of repeated recurrences and his biochemical profile of markedly decreased phosphate and 1,25 (OH)₂ vitamin D levels and significantly increased plasma FGF-23 levels. Radiological examination of the left limb showed that the mass of the calf consisted of multiple tumors and that

there was a separate lesion at the tibial metaphysis (2). The above-mentioned data combined with the clinical presentation of the patient with excruciating pain and limp dysfunction led to the decision of an aggressive operation of high amputation of the affected limb up to the height of the distal femur. At that time, we also performed a molecular analysis of biopsies taken from the tumor and from cells isolated from bone marrow. We recorded an over-expression of genes related to growth factor receptors (PDGFRA, PDGFRB and VEGFR3) that have been documented before in various types of cancers (3, 4), but not in histologically benign PMTs.

Two years after amputation, the patient had persistently low PO₄ levels (1.8 mg/dL), despite being supplemented with oral phosphate and calcitriol; FGF23 levels were also markedly increased (5630 pg/mL). The tubular reabsorption of phosphate (TRP, 75.97%), TmP/GFR (0.5/24 h), and urinary phosphorus (29.38 mmol/L), were indicative of hyperphosphaturia. Apart from general muscle weakness and cramps, attributed to chronic hypophosphatemia, he complained of persistent cough and shortness of breath that was aggravated in the last 4 months, restricting him from operating simple everyday tasks and deteriorating even more his quality of life.

Investigation

A chest X-ray revealed multiple pulmonary nodules with ground glass appearance in both lungs and a PET scan was performed to further evaluate the multiple pulmonary nodules (Fig. 1).

Open lung biopsy was performed. Histological examination of the excised lung nodule revealed metastatic PMTs (Fig. 2). The histologic features were similar to those of the primary tumor (2) and there was no mitotic activity in at least 40 high power fields ($\times 400$ magnification; Fig. 3). The cellular proliferation marker, Ki/67/MIB1 was positive in less than 5% of the neoplastic cells (Fig. 4). The neoplastic cells were positive for vimentin, and negative for desmin S 100 protein and CD34 antigen. The stroma had areas of hyalinization, ossification and hemosiderin deposits.

Outcome and follow-up

Six months after the diagnosis of lung metastases, the patient was admitted to the emergency department due to acute dyspnea and ultimately succumbed to respiratory failure 24 h later.

Figure 1
Axial PET-CT scan image showing numerous metabolically active pulmonary nodules.

Discussion

The histopathological diagnosis of PMTs is a challenge, considering the rarity of the tumor and its non-specific histological and immunohistochemical findings. As demonstrated by Folpe and *et al.* (5) most PMTs cases compose of small, bland, spindled cells that produce a distinctive 'smudgy' matrix, with a well-developed capillary network (often described as Hemangiopericytoma-like vessels) and possibly microcystic change. This matrix calcifies in an unusual flocculent or 'grungy' pattern. Mitotic activity is usually mild (0–1 mitoses per high power magnification field). Immunohistochemistry does not positively contribute

Figure 3
Lung metastasis of PMTs, H&E stain x400.

to the diagnosis of PMTs, other than the exclusion of other mesenchymal neoplasms. The exception to this is the immunohistochemical detection of FGF23 expression. Nevertheless, commercially available FGF23 antibodies lack sufficient specificity to reliably diagnose PMTs (6).

The rarity and the elusive histopathological presentation of PMTs have resulted in poor understanding of its biological behavior. According to the World Health Organization (WHO) classification, PMTs has two types: benign and malignant (4). While there are no standard distinctive histopathological criteria between the two forms, the latter is defined on the basis of 'frankly sarcomatous features' (high mitotic rate, prominent atypia and pleomorphism and necrosis). Most reported

Figure 2
Lung metastasis of PMTs with ossification, H&E stain x40.

Figure 4
Lung metastasis of PMTs, Ki67/MIB1 stain x400.

Table 1 Reported cases of PMTs with malignant behavior.

Publication	Number of cases	Site	Histopathological evidence of malignancy	Metastasis	Follow-up/outcome
Folpe <i>et al.</i> (5)	1 out of 32	Various	3 of 32 cases malignant PMTs		For those patients with available follow-up all were alive free of disease except for four patients with recurrent tumors. One of those developed lung and bone metastasis and was diagnosed with malignant PMTs
Morimoto <i>et al.</i> (10)	2	(1) Right pelvis (2) right pelvis (found at the age of 25 untreated until the age of 31)	(1) Malignant PMTs (2) first diagnosis: benign PMTs; metastasis diagnosis: malignant transformation	(1) Lung and bones (2) lung, bones, liver and elbow	(1) DOD from rapidly progressive lung metastases 32 months after second embolization (2) recurrence 2 years after surgery, metastasis, DOD due to respiratory failure (total clinical course: 26 years)
Qari <i>et al.</i> (8)	1	Mandible	No and Ki67 <3%	Lung and soft tissue	24 year-course, multiple metastases, AWD
Wasserman <i>et al.</i> (9)	2	Floor of mouth-nasal cavity	(1) Yes (2) yes	Lung-lung	(1) >20 year-course multiple recurrences (2) >10 year-course multifocal disease
Qju <i>et al.</i> (7)	1	Right leg	Yes (Ki67 >40%)	Lung	Recurrence after first surgery, AWNED 10 months after the last operation

AWD, alive with disease; AWNED, alive with no evidence of disease; DOD, dead of disease; PMTs, phosphaturic mesenchymal tumors.

cases of PMTs are of the benign type and have either a favourable outcome or recurred locally. Cases of PMTs with distant metastases are rare and usually have evident histological features of malignancy (5, 7, 8, 9, 10) (Table 1). To the best of our knowledge the current case is only the second report of PMTs with histologically benign features presenting with distant metastasis on long-term follow-up. In the other case, reported by Qari *et al.* (8), the patient presented with a 1.2cm destructive osteolytic lesion in the mandible and the biopsy showed a cellular spindle cell neoplasm with osteoid-like production, compatible with PMTs. After multiple local recurrences, the tumor eventually metastasized to the lungs as multiple, small foci of tumor. Seventeen years from original presentation, it was confirmed that the disease had become widespread with lesions in the soft tissue. Our case was diagnosed as benign PMTs based on the WHO classification criteria, due to lack of sarcomatous features in all examined specimens, including metastatic sites. The only finding that was suggestive of an aggressive behavior was over-expression of genes related to growth factor receptors (2). Our patient had multiple recurrences, distant metastases and eventually died of the disease, indicating that despite absence of histologic evidence, the tumor had malignant potential.

This case emphasizes the need for long-term follow-up for patients with PMTs.

Declaration of interest

The authors declare that there is no conflict of interest that could be perceived as prejudicing the impartiality of the research reported.

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Patient consent

Written informed consent from the patient was obtained before any analysis was performed.

Author contribution statement

M P Yavropoulou wrote the first draft. C Poullos and P Hytioglou performed the pathology; C Foroulis, S Tournis, I Kesisoglou, K Kotsa, John G Yovos and P Zebekakis, revised and approved the final draft.

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