Ectopic expression of Klotho in fibroblast growth factor 23 (FGF23)-producing tumors that cause tumor-induced rickets/osteomalacia (TIO)

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ABSTRACT

Tumor-induced rickets/osteomalacia (TIO) is a rare paraneoplastic syndrome caused by tumors that ectopically express fibroblast growth factor 23 (FGF23). FGF23 is a bone-derived hormone that regulates serum phosphate concentrations. Patients with TIO develop hypophosphatemic rickets/osteomalacia due to FGF23 excess and suffer from symptoms such as leg deformities, bone pain, skeletal muscle myopathy, and multiple fractures/pseudofractures. Usually, successful surgical removal of the causative tumors normalizes serum FGF23 and phosphate concentrations in patients with TIO. Most FGF23-producing tumors associated with TIO are histologically called phosphaturic mesenchymal tumor, mixed connective tissue variant (PMTMCT). The precise mechanism by which these tumors ectopically overproduce FGF23 outside of bone is yet to be clarified. Therefore, we performed an RNA sequencing analysis of a PMTMCT that was found in the left parotid gland of a patient with TIO. Among the upregulated genes, we focused on Klotho, the protein product of which is a single pass transmembrane protein that works along with an FGF receptor 1c as a receptor complex for FGF23. Subsequent histological analysis confirmed the ectopic expression of Klotho in other PMTMCTs. From these results, we assume that the ectopic expression of Klotho in PMTMCTs enables a positive feedback loop in FGF23 production via the activation of FGF receptor 1c and exacerbates disease manifestations in TIO.

1. Introduction

Tumor-induced rickets/osteomalacia (TIO) is a rare paraneoplastic syndrome caused by tumors that overproduce fibroblast growth factor 23 (FGF23) (Shimada et al., 2001; Kumar, 2000). FGF23 is a bone-derived hormone that regulates serum phosphate concentrations (Fukumoto and Martin, 2009). FGF23 reduces phosphate reabsorption in the proximal renal tubules by downregulating sodium-phosphate cotransporters. FGF23 also decreases phosphate absorption in the intestine by reducing serum active vitamin D: 1,25-dihydroxyvitamin D [1,25(OH)2D] levels (Shimada et al., 2004). Therefore, chronic FGF23 excess leads to vitamin D-resistant hypophosphatemia. Adult patients with TIO suffer from symptoms such as bone pain, skeletal muscle myopathy, and multiple fractures/pseudofractures (Minisola et al., 2017). Additionally, growth retardation and leg deformities are common features of patients with childhood-onset TIO (Imel et al., 2006).

Historically, McCance reported a patient with osteomalacia with resistance to vitamin D in 1947, which is considered to be the first reported case of TIO (McCance, 1947). Weidner and Santa Cruz described 17 cases of mesenchymal tumors that caused osteomalacia or rickets in 1987 (Weidner and Santa Cruz, 1987). Later, it was revealed that FGF23 excess is responsible for the pathogenesis of TIO (Shimada et al., 2001; Kumar, 2000). Moreover, FGF23 excess also leads to several congenital hypophosphatemic diseases, such as autosomal dominant hypophosphatemic rickets (ADHR; OMIM #193100) (ADHR Consortium, 2000) and X-linked hypophosphatemia (XLH; OMIM #307800) (The HYP Consortium, 1995).
Table 1
Patient characteristics.

| Case | Age/sex | Location of the tumor | Serum parameters<sup>b</sup> | TmP/GFR<sup>c</sup> |
|------|---------|-----------------------|------------------------------|---------------------|
|      |         |                       | Calcium (8.8–10.1 mg/dL)    | (2.3–4.3 mg/dL)     |
|      |         |                       | Phosphate (2.7–4.6 mg/dL)   |                     |
|      |         |                       | Creatinine (M, 0.65–1.07 mg/dL; F, 0.46–0.79 mg/dL) | (106–332 U/L)       |
|      |         |                       | ALP (106–332 U/L)           | 10–65 pg/mL         |
|      |         |                       | iPTH (10–65 pg/mL)          | FGFR23 (10–50 pg/mL) |
|      |         |                       | 1,25(OH)<sub>2</sub>D (20–60 pg/mL) |                     |
| 1    | 77M     | Left parotid gland    | 8.8                          | 3.3                 | 0.72                  | 515                    | 80         | 187  | 70.7  | 1.7 |
| 2    | 53F     | Right external ear canal | 8.4                          | 2.7                 | 0.36                  | 1880                   | 47         | 464  | 36.4  | 2.0 |
| 3    | 51M     | Left sole             | 8.6                          | 2.0                 | 1.07                  | 1765                   | 22         | 893  | 52.7  | 0.9 |
| 4    | 36F     | Left nasal cavity     | 8.9                          | 2.3                 | 0.52                  | 270                    | 45         | 142  | 19.1  | 1.6 |
| 5    | 64F     | Right iliac bone      | 8.5                          | 2.2                 | 0.46                  | 832                    | 77         | 117  | 26.7  | 1.7 |
| 6    | 69M     | Right thyroid muscle  | 9.0                          | 2.2                 | 0.84                  | 292                    | 34         | 488  | 22.8  | 1.8 |
| 7    | 53M     | Right femur           | 9.3                          | 2.8                 | 1.16                  | 478                    | 220        | 1475 | 5.1   | 1.2 |
| 8    | 41F     | Left maxillary sinus  | 8.6                          | 2.3                 | 0.66                  | 610                    | 83         | 423  | 65.5  | 1.2 |
| 9    | 59M     | Left femur            | 9.0                          | 2.0                 | 0.57                  | 1242                   | 88         | 100  | 116.0 | 1.1 |
| 10   | 69M     | Right femur           | 8.2                          | 2.3                 | 0.61                  | 834                    | 37         | 3280 | 35.9  | 1.4 |
| 11   | 45M     | Right femoral neck    | 8.7                          | 1.5                 | 0.73                  | 679                    | 82         | 166  | 18.3  | 1.2 |
| 12   | 63M     | First lumbar vertebral body | 9.0                          | 2.5                 | 0.94                  | 404                    | 71         | 5211 | 47.3  | 1.5 |
| 13   | 59M     | Palate                | 8.2                          | 1.4                 | 0.65                  | 712                    | 140        | 94   | 53.0  | 0.8 |
|      |         | Average ± SD (all)    | 8.8 ± 0.4                    | 2.3 ± 0.5           | 0.71 ± 0.23           | 809 ± 519              | 79 ± 52    | 1003 ± 1543 | 438 ± 29.1 | 1.4 ± 0.4 |
|      |         | Average ± SD (Men, n = 9) | 8.8 ± 0.4                    | 2.2 ± 0.6           | 0.81 ± 0.21<sup>c</sup> | 769 ± 467              | 86 ± 62    | 1322 ± 1785 | 46.9 ± 32.9 | 1.3 ± 0.3 |
|      |         | Average ± SD (Women, n = 4) | 8.6 ± 0.2                    | 2.4 ± 0.2           | 0.5 ± 0.13<sup>c</sup> | 898 ± 694              | 63 ± 20    | 267 ± 182 | 36.9 ± 20.3 | 1.6 ± 0.3 |

<sup>a</sup> Age at the operation.
<sup>b</sup> Data prior to the operation, reference range for each parameter is given in brackets.
<sup>c</sup> Significant difference between male and female patients.
Most causative tumors for TIO are benign and located in bone or soft tissue (Folpe et al., 2004), except for a few cases of TIO caused by malignancies such as colorectal, ovarian, prostate, and thyroid cancers (Lin et al., 2014; Leaf et al., 2013; Mak et al., 2012; Abate et al., 2016). These mesenchymal tumors are histologically termed phosphaturic mesenchymal tumor, mixed connective tissue variant (PMTMCT). The typical features of PMTMCT are the dense proliferation of spindle-shaped cells, myxoid areas, chondroid or osteoid-matrix containing flocculent calcification, osteoclast-like giant cells, and blood vessels (Folpe et al., 2004). The majority of PMTMCTs are histologically benign, but a relatively small number of malignant PMTMCTs with distant metastases have been reported (Morimoto et al., 2014; Nair et al., 2017; Yavropoulou et al., 2018).

FGF23 is physiologically produced in osteocytes/osteoblasts and binds to a complex of FGF receptor 1c (FGFR1c) and Klotho (referred to as Klotho in this manuscript) to serve its function (Urakawa et al., 2006; Kurosu et al., 2006). Klotho is a single-pass transmembrane protein that shares sequence homology with family I beta-glycosidases (Matsumura et al., 1998). In human, the expression of Klotho is restricted to specific organs such as the kidney, prostate, placenta, and parathyroid glands. Considering the ubiquitous expression of FGFR1c, it is likely that Klotho enables the site-specific action of FGF23 (Urakawa et al., 2006). Klotho-deficient mice show FGF23 resistance with a 2000-fold increase in serum FGF23 concentrations compared to those in wild-type mice (Kuro-o et al., 1997). Moreover, because of the loss of FGF23 function, Klotho-deficient mice show similar phenotypes to those of Fgfr23-null mice, such as ectopic calcification, elevated serum 1,25(OH)2D concentrations, and hyperphosphatemia (Kurosu et al., 2006). Therefore, it is widely accepted that Klotho plays an essential role in FGF23 signaling.

The regulatory mechanism of FGF23 production in osteocytes/osteoblasts is not fully understood. Animal and human studies have shown that specific conditions, such as treatment with active vitamin D (Kolek et al., 2005; Yamamoto et al., 2010; Hertel et al., 2013; Sprague et al., 2015), phosphate overload (Hori et al., 2016; Perwad et al., 2005; Antoniucci et al., 2006; Ferrari et al., 2005), and uremia (Stubbs et al., 2007), upregulate FGF23 transcription in bone and elevate circulating FGF23 concentrations. Additionally, previous studies of inherited FGF23-related hypophosphatemic diseases have shown the involvement of several gene products in FGF23 transcription. For example, inactivating mutations in the phosphate-regulating endopeptidase homolog, X-linked (PHEX) gene lead to the upregulation of FGF23 production in patients with XLH, which is the most prevalent form of genetic FGF23-related hypophosphatemic rickets (Liu et al., 2003).

So far, the precise mechanism of ectopic FGF23 production in tumors that cause TIO is unknown. Recent studies have shown the presence of FNF1-FGRF1 or FNF1-FGF1 fusion genes in several tumors that are causative for TIO. However, the function of these fusion gene products remains unknown (Lee et al., 2015; Lee et al., 2016). Therefore, to examine the regulatory mechanism of FGF23 production in TIO-related tumors, we performed an RNA sequencing analysis using the parotid gland from a patient with TIO. Subsequent investigation confirmed the ectopic expression of Klotho in FGF23-producing tumors, which suggests the involvement of the pathogenesis of TIO.

2. Subjects and methods

2.1. Subjects

We performed RNA sequencing analysis using RNA extracted from a tumor and the adjacent normal tissue in the parotid gland of a patient with TIO (Takashi et al., 2017) (case #1). The brief medical history of the patient is as follows: the patient was a 77-year-old man who had been suffering from progressive pain in the back and the hip joints for eight years. Laboratory data showing hypophosphatemia and elevated serum FGF23 concentrations led to the diagnosis of FGF23-related hypophosphatemic osteomalacia. An 18F-FDG-PET/CT scan revealed a tumor in the left parotid gland. Higher FGF23 concentration in the left external jugular vein compared to other sites indicated that the tumor overproduced FGF23. When the tumor was surgically removed, the patient’s serum phosphate concentration returned to the normal range. The tumor was histologically diagnosed as a PMTMCT with positive FGF23 staining.

We subsequently enrolled 12 patients with TIO whose formalinfixed, paraffin-embedded (FFPE) tumor samples were available (cases #2–#13). Table 1 summarized the characteristics of patients and the location of the tumors. All the patients were treated with a combination of alfacalcidol and oral phosphate salts. Biochemical parameters including serum albumin, calcium, phosphate, alkaline phosphatase (ALP), creatinine (Cr), intact parathyroid hormone (iPTH), 1,25(OH)2D3, and urine phosphate and creatinine were collected from the medical record. These parameters were measured before the surgery. Serum FGF23 levels were evaluated using an FGF23 ELISA KIT (Kainos, Tokyo, Japan) that detects only full-length FGF23 (Endo et al., 2008). Although the reference range for serum FGF23 is 10–50 pg/mL for healthy subjects without chronic kidney disease, FGF23 levels more than 30 pg/mL are considered abnormally high in patients with hypophosphatemia and leads to the diagnosis of FGF23-related hypophosphatemia (Endo et al., 2008). Serum calcium concentrations were corrected by albumin if albumin was < 4.0 mg/dL. The reference ranges for adults are 8.8–10.1 mg/dL for serum corrected calcium, 2.7–4.6 mg/dL for serum phosphate, 106–332 U/L for serum ALP, 10–65 pg/mL for serum iPTH, 20–60 pg/mL for serum 1,25(OH)2D3, and 0.65–1.07 mg/dL (men) and 0.46–0.79 mg/dL (women) for serum creatinine. The tubular maximum reabsorption of phosphate per unit of glomerular filtrate (TmP/GFR) was calculated using serum phosphate, serum creatinine, urine phosphate, and urine creatinine concentrations. The reference range for TmP/GFR is 2.3–4.3 mg/dL.

This study was approved by the institutional review board of the University of Tokyo. We obtained written informed consent for the RNA sequencing and opt-out consent for both the immunohistochemical and RT-PCR analyses of tumor samples.

2.2. RNA sequencing of the parotid tumor

Total RNA was extracted from the fresh frozen parotid gland tissue and the adjacent normal parotid gland tissue using the NucleoSpin® RNA kit (Macherey-Nagel, Düren, Germany).

Paired-end sequencing libraries were constructed using the TruSeq RNA library Prep Kit v2 (Illumina, San Diego, CA). We sequenced the libraries on a MiSeq system (Illumina, San Diego, CA) using the 300-cycles MiSeq Reagent Kits v2 (Illumina, San Diego, CA). Data were analyzed on a CLC genomic workbench v8 (CLC bio Japan, Tokyo, Japan).

For RT-PCR, total RNA was reverse transcribed to cDNA using the PrimeScript® RT Master Mix (Takara Bio, Shiga, Japan). We amplified 100–300 base products of FGF23, Klotho, FGFR1c, and GAPDH cDNA using the SapphireAmp® Fast PCR Master Mix (Takara Bio, Shiga, Japan). The primer sets were as follows: 5′-TCTTTGTGTTTGACCCA CCT-3′ and 5′-CACCCTGAAACCATCCATG-3′ for Klotho, 5′-ATGCCTG AGTATAACACC-3′ and 5′-TCCAGAACGTTCAACACT-3′ for FGFR1c, and 5′-GCGACCCGTCAAAGGTG-3′ and 5′-CAGCATGCC CCCATGTA-3′ for GAPDH. We purchased the primer sets for FGF23 from TaKaRa Bio (HA181049). The primer sets designed to detect fusion genes were as follows: 5′-TCGTGGCCGCAGAGGAGTAC-3′ and 5′-GCTTCAATGCCATCTGTGGTCAAGT-3′ for the FNF1-FGRF1 fusion gene and 5′-TCGGGCCGCCGATACAGAGAAT-3′ and 5′-GCTTTTCCGGCGACTGATC-3′ for the FNF1-FGF1 fusion gene. PCR conditions were as follows: 1 min at 94 °C, followed by 35 cycles of 5 s at 98 °C, 5 s at 58 °C, and 5 s at 72 °C, with a final extension for 3 min at 72 °C. PCR products were electrophoresed on a 1% agarose gel containing ethidium bromide.
and visualized with UV light.

2.3. Immunohistochemistry of the tumor samples

Immunohistochemical (IHC) staining was performed using the Ventana BenchMark automated immunostainer (Ventana benchmark; Ventana Medical Systems Inc., Tucson, AZ) according to the manufacturer’s instructions. The primary antibodies used in this study were an anti-human Klotho monoclonal antibody that detects 55–261 amino acids of human Klotho (KM2076, Trans Genic Inc., Kobe, Japan) and an anti-phospho-Erk1/2 antibody that recognizes phospho-p44/42 MAPK (#4370, Cell Signaling Technology Japan, Tokyo, Japan).

2.4. Analysis of serum Klotho concentrations in the patients

A human soluble alpha-Klotho assay kit (IBL, Gunma, Japan) was used to measure the serum Klotho concentrations in 11 out of 13 patients.

2.5. Statistical analysis

The relationships between each biochemical parameter of the patients and the relationships between serum FGF23 and serum Klotho concentrations were analyzed using JMP® Pro14.0 (SAS Institute Japan Ltd., Tokyo, Japan). p values < 0.05 were considered statistically significant.

3. Results

3.1. Patient characteristics

We analyzed nine male patients and four female patients with TIO in our study. Even though all the patients were treated with a combination of alfacalcidol and oral phosphate salts, seven and ten of 13 patients showed hypocalcemia and hypophosphatemia, respectively (Table 1). Serum ALP levels were elevated in 10 patients. Eight patients showed secondary hyperparathyroidism. TmP/GFR levels were lower than the reference range in all the patients, suggesting that there was an increase in renal phosphate wasting as a result of FGF23 excess. In fact, there was a significant positive correlation between serum phosphate levels and TmP/GFR levels (r = 0.6179, p = 0.0244). Additionally, there was a significant negative correlation between serum corrected calcium and ALP (r = −0.4018, p = 0.0322). We did not find any significant correlation between serum FGF23 and other parameters. At the same time, there was no significant difference in parameters between male and female patients other than serum creatinine levels (Table 1).

3.2. RNA sequencing analysis of the tumor in the parotid gland

We detected several upregulated and downregulated genes in the parotid gland tumor compared to those in the adjacent normal tissue. Table 2 shows a list of representative upregulated genes in the tumor. We could not identify any significant fusion genes that involve FGFR1 or FGF1. The RT-PCR analysis confirmed the presence of FGF23, Klotho, and FGFR1c mRNA in the tumor (Fig. 1).

3.3. Immunohistochemistry of the causative tumors in TIO

Among 13 tumors from patients with TIO, nine tumors were positive for IHC staining for Klotho (Table 3). In these tumors, spindle-shaped tumor cells, which are considered the principal source of FGF23 in TIO, were positive for Klotho staining. Fig. 2 shows the images of tumor cells positive for Klotho staining (Fig. 2b–j) and those negative for Klotho staining (Fig. 2k and l), along with a positive control of renal tubular cells (a). Additionally, 11 of 13 tumors were positive for phospho-Erk1/

Table 2

| Name   | RPKM* | Fold change |
|--------|-------|-------------|
| Control | Tumor |
| DMP1   | 0.9545| 2411.07     |
| FGFI   | 0.6298| 67.6582     |
| FGFR2  | 0.1497| 207.605     |
| MEPE   | 5.2285| 7159.17     |
| SOST   | 2.4226| 3257.41     |
| SFRP4  | 1.5490| 1976.49     |
| Klotho | 0.4545| 446.978     |
| SPP1   | 12.9661| 11,899.7   |
| ACP5   | 0.3211| 236.662     |
| PHEX   | 0.1478| 104.634     |
| MMP9   | 2.9010| 1659.15     |

Table 3

| Case | Klotho | p-Erk |
|------|--------|-------|
|      | IHC | RT-PCR | IHC |
| 1    | +   | +      | +   |
| 2    | +   | N.E.   | +   |
| 3    | +   | +      | +   |
| 4    | +   | N.E.   | +   |
| 5    | +   | +      | +   |
| 6    | −   | N.E.   | +   |
| 7    | +   | N.E.   | +   |
| 8    | −   | N.E.   | +   |
| 9    | +   | N.E.   | N.E.|
| 10   | +   | N.E.   | N.E.|
| 11   | −   | N.E.   | N.E.|
| 12   | −   | N.E.   | N.E.|
| 13   | +   | N.E.   | N.E.|

FGFR1, fibroblast growth factor receptor 1; FGF23, fibroblast growth factor 23; MEPE, matrix extracellular phosphoglycoprotein; SOST, sclerostin; SFRP4, secreted frizzled-related protein 4; SPP1, secreted phosphoprotein 1; ACP5, acid phosphatase 5, tartrate resistant; PHEX, phosphate-regulating endopeptidase homolog, X-linked; MMP9, matrix metalloproteinase 9.

*I RPKM, reads per kilobase of exon per million mapped reads.
patients were between 94 and 5211 pg/mL. There was no significant correlation between serum Klotho and serum FGF23 concentrations (r = 0.2332, p = 0.4902) (Fig. 4).

4. Discussion

In this study, we detected the ectopic expression of Klotho mRNA and Klotho protein in the FGF23-producing tumors that cause TIO. First, the result of RNA sequencing analysis showed the upregulation of Klotho mRNA in a tumor in a parotid gland compared to the adjacent normal tissue. Second, IHC analysis of the FFPE samples revealed positive Klotho staining in nine of 13 FGF23-producing tumors. A previous report by Yavropoulou et al. showed the expression of Klotho mRNA in an FGF23-producing tumor in the periphery of the fibula (Yavropoulou et al., 2015). In our study, we confirmed both the expression of Klotho mRNA and Klotho protein in the majority of tumors in various locations. When limited organs such as renal tubular cells express Klotho under physiological conditions, we hypothesize that the ectopic expression of Klotho in these mesenchymal tumors may be involved in the pathogenesis of TIO. Although the exact cause and the meaning of Klotho expression in PMTMCs are not yet determined, we want to propose several hypotheses.

First, we conclude that the ectopic expression of Klotho in the tumor cells may lead to the activation of FGFRs. There is growing evidence that FGFRs not only serve as a receptor for FGF23 but also regulate FGF23 production. Wohrle et al. (2011) have shown that FGF9 induces Fgf23 expression in the rat osteosarcoma cell line UMR-106 cells in vitro, which suggests that FGF23 is a target of FGFR signaling in bone. Osteoglophonic dysplasia (OGD; OMIM #166250) is caused by activating mutations in the FGFR1 gene (White et al., 2005). It is reported that some patients with OGD develop FGF23-related hypophosphatemia in addition to various skeletal complications, such as craniosynostosis, dwarfism, and characteristic facial features (White et al., 2005). Conversely, conditional deletion of Fgfr1 in osteocytes from Hyp mice, which is a murine model of XLH, has resulted in reduced circulating FGF23 (Xiao et al., 2014). In addition, pharmacological inhibition of FGFRs results in transient repression of FGF23 mRNA in bone and a decrease in serum FGF23 concentrations in Hyp mice (Xiao et al., 2014), which is in turn followed by a compensatory increase in serum FGF23 levels after long-term therapy (Wohrle et al., 2013). Therefore, we hypothesize that the ectopic expression of Klotho in PMTMCs helps in the activation of the FGFR signaling pathway and creates a local positive feedback loop for FGF23 production (Fig. 5). In other words, ectopically expressed Klotho enables the autocrine and paracrine effects of FGF23. The positive staining with phospho-Erk1/2 (Fig. 3) is in line with the activation of the FGFR signaling pathway in FGF23-producing tumor cells.

Second, we conclude that the expression of Klotho in PMTMCs may reflect the osteoblastic differentiation of tumor cells. PMTMCs are often present in the bone or the soft tissue adjacent to the bone, which makes it difficult to interpret the results of genetic analysis of such tumors. In our study, however, we used a tumor in the parotid gland, which is not mineralized in physiological conditions, to rule out the possibility of contamination by calcified tissues. Nonetheless, the result of RNA sequencing analysis showed the upregulation of genes that are related to osteoblasts and osteocytes (Table 2). Moreover, matrix metalloproteinase 9 (MMP9) and acid phosphatase 5, tartrate resistant (ACP5), which are unique to osteoclasts and macrophages, were also upregulated. Since it has been shown that bone-forming cells such as osteoblasts and osteocytes express low amounts of Klotho (Raimann et al., 2013; Rhee et al., 2011; Komaba et al., 2017), the Klotho expression in tumor cells may reflect the differentiation of mesenchymal...
stem cells to osteoblastic lineage cells. Therefore, we hypothesize that a bone-like microenvironment is created in PMTMCs. This bone-like microenvironment could augment FGF23 production in tumor cells.

Klotho, which is also referred to as $\alpha$Klotho, is considered vital in the regulation of phosphate homeostasis, while $\beta$Klotho is required for FGF19 and FGF21 signaling (Kurosu and Kuro, 2009). There are different isoforms of Klotho protein: membrane-bound Klotho and soluble Klotho. Membrane-bound Klotho has a large extracellular domain that is subjected to ectodomain shedding and is released into the extracellular space as soluble Klotho (Matsumura et al., 1998). Several reports have suggested the involvement of soluble Klotho in the regulation of FGF23 production. Brownstein et al. have reported a patient with FGF23-related hypophosphatemic rickets who harbored a translocation of a chromosome adjacent to the Klotho gene and elevated serum Klotho concentrations (Brownstein et al., 2008). Additionally, treatment with an adeno-associated virus that produced Klotho resulted in increased circulating serum Klotho and caused FGF23-related hypophosphatemia in mice (Smith et al., 2012). However, in our study, no correlation was found between serum Klotho and FGF23 concentrations. Therefore, we hypothesize that membrane-bound Klotho rather than soluble Klotho is involved in the mechanism of FGF23 production.

Fig. 3. Phospho-Erk staining of FGF23-producing tumors.
Tumor cells were positive for phospho-Erk1/2 staining in 12 of 13 patients (b, case #1; c, case #2; d, case #3; e, case #4; f, case #5; g, case #6; h, case #7; i, case #8; j, case #9; k, case #12; l, case #13) as well as the positive control of the thyroid follicular cells (a). Original magnification for all photomicrographs is × 400.

Fig. 4. Relationships between serum FGF23 and Klotho concentrations in patients with TIO.
There was no significant correlation between serum FGF23 and Klotho concentrations in patients with TIO (n = 11).

Fig. 5. Autocrine/paracrine effects of FGF23 in FGF23-producing tumor cells.
Membrane-bound Klotho enables the autocrine and paracrine effects of FGF23. FGF23 that is secreted from tumor cells binds to a receptor complex of Klotho and FGFR1c and activates the FGFR signaling pathway to enhance the production of FGF23 in tumor cells. The positive feedback loop in the production of FGF23 and a bone-like microenvironment in PMTMCs may exacerbate the disease manifestations in patients with TIO. MAPK, Mitogen-activated Protein Kinase.
methylation of a CpG island in its promoter region appears to be responsible for the tissue-specific expression of the Klotho gene (Xu and Sun, 2015). We have conducted a methylation analysis using DNA extracted from tumors to see whether methylation status correlates with Klotho expression in FGF23-producing tumors. However, we found no correlation between promoter methylation status and the amount of Klotho mRNA expression (data not shown). Therefore, we conclude that DNA methylation does not principally regulate Klotho expression in FGF23-producing mesenchymal tumors.

The limitations of this study are as follows: First, as we used FFPE tissues instead of fresh frozen tissues for IHC analysis; the deterioration of samples over the years might have resulted in negative Klotho staining in some cases. Second, we could not determine whether the expression of Klotho and the presence of the FN1-FGFR1 or FN1-FGF1 fusion genes are mutually exclusive or not in FGF23-producing tumors. We did not detect these fusion genes for the four patients (cases #1, #3, #5, and #11) whose tumor RNAs were available. The presumed function of the protein products of the FN1-FGFR1 and FN1-FGF1 fusion genes is the activation of FGFR1 (Lee et al., 2016). Therefore, the inhibition of the protein products of the FN1-FGFR1 might explain the positive staining with p-Erk in Klotho negative samples. We will continue to study the mechanism of MAPK activation in PMTMCs by using fresh tumor samples in the future.

In conclusion, we found the ectopic expression of Klotho in FGF23-producing tumors. Although the precise regulatory mechanism of Klotho expression in PMTMCs is not clear, we hypothesize that Klotho helps to create a local positive feedback loop in the production of FGF23 through the activation of FGFR1 and exacerbates disease manifestations in patients with TIO. From a clinical perspective, the probable involvement of the FGR signaling pathway in the pathogenesis of TIO justifies the application of FGR inhibitors in patients with refractory TIO. Although the complete resection of tumors is always the optimal treatment for TIO, the results of our study suggest that patients without surgical indication suffering from unresectable, residual, or metastatic lesions may benefit from FGR inhibitors.

Conflict of interest

Dr. Kinoshita has received grants, KAKENHI 15K19528 and 17K16161, from Japan Society for the Promotion of Sciences (JSPS), during the conduct of the study; Dr. Takashi reports grants from JSPS, outside the submitted work; Dr. Ito reports grants from JSPS and research funding from Kyoto Hakko Kirin, outside the submitted work; Dr. Ikegawa has nothing to disclose; Dr. Mano has nothing to disclose; Dr. Ushiku has nothing to disclose; Dr. Fukayama has nothing to disclose; Dr. Nangaku reports advisory fees or research funding from Chugai Pharmaceutical, Taisho Pharmaceutical, Ono Pharmaceutical, and Kyowa Hakko Kirin.

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