Can Ultrasound Alone Predict Papillary Thyroid Carcinoma with Desmoid-Type Fibromatosis? A Retrospective Analysis of 13 Cases, Focusing on the Stromal Area

Abstract

Purpose Papillary thyroid carcinoma with desmoid-type fibromatosis (PTC-DTF) is extremely rare. So far, only 4 cases describing the ultrasound findings of this variant have been reported. Here, we describe the ultrasound findings of 13 cases of PTC-DTF, focusing especially on the DTF area.

Materials and Methods We retrospectively analyzed the clinical reports, ultrasound reports, and ultrasound photographs obtained from medical records at Kuma Hospital.

Results The patients included 8 women and 5 men with a mean age of 47.9 years. The widest dimension of the nodules ranged from 16 to 79 mm (mean: 37.5 mm). The original ultrasound reports classified the nodules as either intermediate suspicion or high suspicion. A diagnosis of PTC was suspected in 12 nodules, and anaplastic carcinoma was suspected in 1 nodule. PTC-DTF presented with an irregularly shaped nodule (100 %), taller-than-wide sign (84.6 %), heterogeneous echo-genicity (100 %), no microcalcification (76.9 %), and no or mild flow signal on Doppler (75.0 %). The DTF area was identified in the ultrasound photographs of 8 nodules. DTF areas were generally heterogeneous (62.5 %) and more hypoechoic (71.4 %) than PTC areas. Microcalcification was not observed in the DTF areas. All of the DTF areas revealed no or mild flow signal. On ultrasound elastography, the DTF areas were not stiff, and they were more elastic than the PTC areas.

Conclusion It is difficult to predict PTC-DTF using ultrasound alone, and B-mode ultrasonography is more reliable than ultrasound elastography in the ultrasound diagnosis of malignant thyroid nodules.

Introduction

Papillary thyroid carcinoma (PTC) has many morphologic variants. PTC with extensive proliferation of fibroblasts and myofibroblasts in the stroma is extremely rare, and has been described as “PTC with nodular fasciitis-like stroma” [1–3]. Recently, Rebecchini et al. proposed that this variant should be renamed PTC with desmoid-type fibromatosis (DTF), because the mesenchymal component showed aberrant nuclear and cytoplasmic immunoreactivity for beta-catenin and harbored a heterozygous somatic activating mutation in the corresponding CTNNB1 gene [4]. Given its rarity, reports on PTC- DTF have been sporadic. To the best of our knowledge, only 4 cases describing ultrasound findings of PTC-DTF have been reported so far [4–6]. Here, we report the ultrasound findings of 13 PTC-DTF cases from a single institute, focusing especially on the DTF area. The aim of this study was to evaluate ultrasound features of PTC- DTF and correlate these features with the pathological findings.
Materials and Methods

Pathology reports of 10,659 cases of PTC resected at Kuma Hospital between January 2007 and December 2016 were reviewed. Of these, 13 cases (0.1 %) of PTC-DTF were ultimately included in this study. A diagnosis of PTC-DTF was made based on the presence of extensive fibroblast and myofibroblast proliferation in the stroma of PTC. PTC cases with desmoplastic changes, such as granulation tissue-like stroma, were excluded. DTF components accounted for 20% to 95% of the nodules. We then retrospectively analyzed the clinical and ultrasound reports with photographs obtained from medical records at Kuma Hospital. Ultrasound was performed using the APLIO 80 SSA-770 A (Toshiba Medical Systems Co., Ltd., Otawara, Japan) or APLIO 500 TUS-A500 (Toshiba) ultrasound machine with the PLT-805AT (Toshiba) or PLT-1005BT (Toshiba) probe. Real-time tissue elastography was performed in 2 cases using the EUB-7500 (Hitachi Medical Systems., Tokyo, Japan) ultrasound machine with the EUP-L74M (Hitachi) probe or APLIO 500 TUS-A500 ultrasound machine with the PLT-1005BT probe. Ultrasound reports were interpreted based on patterns proposed by the 2015 American Thyroid Association Management Guidelines [7]. For tissue stiffness, a 4-point scoring system proposed by Asteria et al. [8] was adopted. Statistical analysis was carried out with Fisher’s exact test or Student’s t-test. P-values < 0.05 were considered statistically significant.

Table 1

| Clinical and ultrasound findings of 13 cases of papillary thyroid carcinoma with desmoid-type fibromatosis. |
|---------------------------------------------------------------|
| **Female/male** | 8/5 |
| **Age [years] (mean)** | 19-77 (47.9) |
| **Preoperative serum thyroglobulin [ng/mL] (years)** | 7.0-444.1 (84.5) |
| **Antithyroglobulin antibody positive [ > 39.9 IU/mL]** | 1 (7.7 %) |
| **Original ultrasound report** |  |
| **Intermediate suspicion** | 2 (15.4 %) |
| **High suspicion** | 11 (84.6 %) |
| **Location [left/right/isthmus]** | 5/7/1 |
| **Maximum diameter [mm] (mean)** | 16-79 (37.5) |
| **A/T ratio** | 0.9-2.5 (1.6) |
| **A/T ratio>1.0** | 11 (84.6 %) |
| **Shape** |  |
| Round to oval | 0 (0 %) |
| Irregular | 13 (100 %) |
| Spiculated | 0 (0 %) |
| **Margin** |  |
| Well-defined | 7 (53.8 %) |
| Ill-defined | 6 (46.2 %) |
| Halo | 0 (0 %) |
| Lateral shadow | 3 (23.1 %) |
| **Echogenicity** |  |
| Homogeneous | 0 (0 %) |
| Heterogeneous | 13 (100 %) |
| Markedly hypoechoic | 5 (38.5 %) |
| Hypoechoic | 3 (23.1 %) |
| Isoechoic | 5 (38.5 %) |
| Hyperechoic | 0 (0 %) |
| **Calcification** |  |
| Macrocalcification (>2 mm) | 4 (30.8 %) |
| Microcalcification (<2 mm) | 3 (23.1 %) |
| Peripheral calcification | 1 (7.7 %) |
| Marked posterior shadowing | 1 (7.7 %) |
| **Vascularity** |  |
| No flow signal | 1 (8.3 %) |
| Mild flow signal | 8 (66.7 %) |
| Marked flow signal | 2 (16.7 %) |
| Perinodular flow signal | 2 (16.7 %) |
| **Acoustic posterior enhancement** | 6 (46.2 %) |
| **Extrathyroidal spread** | 5 (38.5 %) |
| **Suspected nodal metastases** |  |
| Central | 5 (38.5 %) |
| Lateral | 7 (53.8 %) |
| Data are expressed as number (%) or range (mean). A/T ratio, anteroposterior-to-transverse diameter ratio |

Results

Clinical and ultrasound findings of 13 patients with PTC-DTF are shown in Table 1. The patients included 8 women and 5 men with a mean age of 47.9 years (range 19–77 years). Cases with a history of ionizing radiation exposure or familial polyposis were excluded. Serum thyroglobulin levels ranged from 7.0 to 444.1 ng/mL (mean: 84.5 ng/mL) and were elevated (>46.5 ng/ml) in 6 patients (46.2 %). Anti-thyroglobulin antibody was positive (>39.9 IU/mL) in 1 patient (7.7 %).

Original ultrasound reports classified the nodules as either intermediate suspicion (2 nodules, 15.4 %) or high suspicion (11 nodules, 84.6 %). Among them, a diagnosis of PTC was suspected in 12 nodules, and anaplastic carcinoma was suspected in 1 nodule. There were no nodules with very low or low suspicion.

5 nodules were located in the left lobe, 7 in the right lobe, and 1 in the isthmus. The widest dimension of the nodules ranged from 16 to 79 mm (mean: 37.5 mm). The mean anteroposterior-to-transverse diameter (A/T) ratio was 1.6 (range: 0.9–2.5), and the taller-than-wide sign (A/T ratio > 1.0) was detected in 11 nodules (84.6 %) (Fig. 1). All of the nodules were irregularly shaped (Fig. 1), and no spiculated nodules were observed. The margins were either well-defined (7 nodules, 53.8 %) or ill-defined (6 nodules, 46.2 %). A thin, sonolucent rim surrounding the thyroid nodule (halo, marginal hypoechoic zone) was not observed in any case. Three nodules (23.1 %) exhibited a lateral shadow. Echogenicity was heterogeneous in all nodules (Fig. 2), and the predominant echogenicity was markedly hypoechoic in 5 nodules (38.5 %), hypoechoic in 3 nodules (23.1 %), and isoechoic in 5 nodules (38.5 %). No hyperechoic nodules were observed. Macrocacification (>2 mm) and punctate microcalcification (<2 mm) were observed in 4 nodules (30.8 %) and 3 nodules (23.1 %), respectively. One nodule (7.7 %) showed peripheral calcification (“eggshell” calcification), and 1 nodule (7.7 %) showed marked posterior shadowing. Twelve nodules were examined by Doppler ultrasound, 75.0 % of which showed no or a mild flow signal. Acoustic posterior enhancement was observed in 6 nodules (46.2 %), and 5 nodules (38.5 %) showed ex-
trathyroidal spread. Central and lateral nodal metastases were suspected in 5 nodules (38.5%) and 7 nodules (53.8%), respectively.

Based on the macro- and microscopic observations of the resected nodules, DTF and PTC areas were identified on the ultrasound images in 8 nodules and 7 nodules, respectively (Table 2). DTF areas were homogeneous in 3 nodules (37.5%) (Fig. 3a) and heterogeneous in 5 nodules (62.5%) (Fig. 4a). Compared with PTC areas, 2 DTF areas (28.6%) were more hyperechoic (Fig. 5a) and 5 DTF areas (71.4%) were more hypoechoic (Fig. 3a). Microcalcification was not observed in any DTF area. All of the DTF areas revealed no or a mild flow signal (Fig. 6a). Ultrasound elastography was performed in cases 7 and 13. In case 13, a large part of the DTF area that occupied the majority of the tumor was somewhat elastic (class 2) (Fig. 4c). In case 13, the DTF area was elastic (class 1), and the PTC area was somewhat elastic (class 2) (Fig. 7).

### Discussion

To the best of our knowledge, only 4 cases describing ultrasound findings of PTC-DTF have been reported so far [4–6]. In 2010, Lee et al. [5] first described the ultrasound findings of a PTC-DTF case in which the nodule showed homogeneous hypoechoicity with a circumscribed margin; color Doppler ultrasound showed minimal vascular flow in the nodule. The nodule in a case reported by Na et al. revealed a mixed echoic pattern composed of markedly and slightly hypoechoic portions [6]; the markedly hypoechoic portion showed increased blood flow. Rebecchini et al. described ultrasound findings of 2 PTC-DTF cases [4]; one of the nodules was partially cystic with sparse microcalcification and a central hyperechoic zone, and the other had an ill-defined border with no microcalcification. Thus, characteristic ultrasound findings of DTF components have not been well established.

In our study, PTC-DTF sonographically exhibited an irregularly shaped nodule with heterogeneous echogenicity and a taller-than-wide sign. The predominant echogenicity was markedly hypoechoic, hypoechoic, or isoechoic. The incidence of punctate microcalcification was low (23.1%) and 75.0% of the nodules showed no or a mild flow signal. The ultrasound findings of PTC-DTF did not appear to be

| Parameter                  | DTF area (n = 8) No. of nodules (%) | PTC area (n = 7) No. of nodules (%) |
|----------------------------|------------------------------------|------------------------------------|
| Echogenicity               |                                    |                                    |
| Homogeneous                | 3 (37.5%)                          | 0 (0%)                             |
| Heterogeneous              | 5 (62.5%)                          | 7 (100%)                           |
| Markedly hypoechoic        | 3 (37.5%)                          | 0 (0%)                             |
| Hypoechoic                 | 3 (37.5%)                          | 4 (57.1%)                          |
| Isoechoic                  | 2 (25.0%)                          | 3 (42.9%)                          |
| Hyperechoic a (n = 7)      | 2 (28.6%)                          |                                    |
| Hypoechoic b (n = 7)       | 5 (71.4%)                          |                                    |
| Calcification              |                                    |                                    |
| Macrocalcification (> 2 mm)| 1 (12.5%)                          | 1 (14.3%)                          |
| Microcalcification (< 2 mm)| 0 (0%)                             | 2 (28.6%)                          |
| Vascularity                |                                    |                                    |
| No flow signal             | 1 (12.5%)                          | 2 (28.6%)                          |
| Mild flow signal           | 7 (87.5%)                          | 3 (42.9%)                          |
| Marked flow signal         | 0 (0%)                             | 1 (14.3%)                          |
| Elastography## (n = 2)     |                                    |                                    |
| Class 1                    | 1 (50.0%)                          | 1 (100%)                           |
| Class 2                    | 1 (50.0%)                          |                                    |

Data are expressed as number (%). a; compared with PTC areas (n = 7)

DTF: desmoid-type fibromatosis; PTC: papillary thyroid carcinoma
Fig. 3 Papillary thyroid carcinoma with focal desmoid-type fibromatosis (DTF). The DTF area (•; areas surrounded by white dotted line) was homogeneous and more hypoechoic than the PTC area. (a; B-mode, longitudinal view, right; b; cut surface of resected thyroid).

Fig. 4 Papillary thyroid carcinoma with a massive desmoid-type fibromatosis (DTF) area. The echogenicity of DTF area (•; area surrounded by white dotted line) was heterogeneous and interpreted as being strain elastography class 2. (a; B-mode, longitudinal view, b; cut surface of resected thyroid, c; elastography).

Fig. 5 Papillary thyroid carcinoma (PTC) with a focal desmoid-type fibromatosis (DTF) area. The DTF area (•; areas surrounded by white dotted line) was more hyperechoic than the PTC area. Punctate microcalcification was seen in the PTC area. (a; B-mode, longitudinal view, b; cut surface of resected thyroid).

Fig. 6 Papillary thyroid carcinoma (PTC) with focal desmoid-type fibromatosis (DTF). The DTF area (•; areas surrounded by white dotted line) was homogeneous and more hyperechoic than the PTC area. No flow signal was seen. (a; B-mode, color Doppler, longitudinal view, b; cut surface of resected thyroid).
much different from those of conventional PTC. As PTC-DTF is composed of both PTC and DTF components, we believe that the 2 components should be evaluated separately.

As the name implies, DTF components of PTC-DTF are microscopically similar to DTF arising in the soft tissue [9]. According to ultrasound reports of soft tissue DTF [10]–[11], masses display heterogeneous or homogeneous echogenicity and are hypoechoic, isoechoic, or hyperechoic. Most of the cases showed posterior acoustic enhancement, substantial flow, and high resistive index values (>0.70). No calcifications or cystic areas were noted. In the present study, based on the macro- and microscopic findings, we could identify DTF areas on the ultrasound images in only 8 out of 13 nodules. DTF areas tended to be heterogeneous in echogenicity and more hypoechoic than PTC areas. Microcalcification was not observed. On Doppler ultrasound, the nodules revealed no or a mild flow signal, which is similar to soft tissue DTF.

Ultrasound elastography can be easily performed in the thyroid due to its superficial location [12, 13]. However, the assessment of deeply located nodules is difficult due to the depth-related decay of the stress [14]. Azizi et al. reported that thyroid nodule stiffness measured by elastography is an independent predictor of thyroid carcinoma with a positive predictive value that is greater than or equal to that of conventional ultrasonographic characteristics [15]. We hypothesized that the DTF area would be firm based on its composition of dense collagenous connective tissue. Contrary to our expectations, the DTF areas were not stiff, and in one case (case 7), the DTC area was more elastic than the PTC area. Thus, clinicians should be aware that PTC- DTF shows a benign pattern on ultrasound elastography. In conclusion, B-mode ultrasonography is more reliable than ultrasound elastography in the ultrasound diagnosis of malignant thyroid nodules.

Conclusion

This is the first description of ultrasound findings of PTC-DTF, focusing especially on the DTF area. The ultrasound findings of PTC-DTF include an irregularly shaped nodule, taller-than-wide sign, heterogeneous echogenicity, no microcalcification, and no or a mild flow signal. DTF areas tend to be heterogeneous in echogenicity and more hypoechoic than PTC areas. Despite the presence of abundant dense connective tissue, DTF areas show a benign pattern on ultrasound elastography. We therefore conclude that it is difficult to predict the DTF variant of PTC using ultrasound alone. Moreover, B-mode ultrasonography is more reliable than ultrasound elastography in the ultrasound diagnosis of malignant thyroid nodules. Further analysis with a greater number of cases is needed to clarify the ultrasound findings of DTF areas of PTC-DTF nodules.

Conflict of Interest

The authors declare no conflicts of interest.

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