Correlation between Ultrasonographic Appearance of Papillary Thyroid Microcarcinoma and BRAF V600E Mutation

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The study was conducted to investigate the correlation between the ultrasonographic appearance of a thyroid nodule and the BRAF V600E mutation. Patients with thyroid nodules \((n = 186)\), for which BRAF V600E testing and cytopathology analysis were performed, and who underwent subsequent surgery for nodule resection were enrolled in this study. For each patient, color Doppler ultrasonography was performed to observe the variables of the nodules. The nodules were then characterized using the thyroid imaging reporting and data system classification TI-RADS. Furthermore, the ultrasonographic appearance of the control group, encompassing patients with nodular thyroid goiters, and the case group, encompassing patients with papillary thyroid microcarcinoma (PTMC), was statically analyzed. Similarly, a statistical analysis of the ultrasonographic appearance of the BRAF V600E-positive and BRAF V600E-negative subgroups was also performed. The accuracy was significantly different for the corresponding values when color Doppler ultrasonography, BRAF V600E testing, or cytopathology alone was used for diagnosis. There were significant differences in the ultrasonographic appearance variables between the control and case groups. Comparing with the BRAF V600E-negative subgroup of the case group, the ultrasonographic appearances of the BRAF V600E-positive subgroup showed less circumscribed and more irregularly shaped nodules, with significantly different aspect ratios of \(>1\). The combination of BRAF V600E testing and color Doppler ultrasonography or cytopathology improved the accuracy of the PTMC diagnose. We found that the ultrasonographic appearance of thyroid nodules was related to PTMC.

1. Introduction

Thyroid carcinoma is the most common malignant tumor of the endocrine system, and it has been increasingly studied by clinicians and researchers. In the past years, its incidence has gradually increased, with the fastest increase observed among all types of malignant tumors [1]. Specifically, in terms of pathological type of thyroid carcinoma, there has been a growing incidence of papillary carcinoma, while in terms of pathological stage microcarcinoma has exhibited the fastest increase in incidence [2, 3]. According to the latest SEER statistics, nearly 90% of thyroid carcinomas are papillary carcinomas [4], reinforcing the importance of studying papillary thyroid microcarcinoma (PTMC). In addition, microcarcinoma has a high cervical lymph node metastasis rate of 30–70% [5–7]. Some patients with microcarcinoma have local recurrences very soon after surgery or can even present initially with distant metastases to the lungs or bones [8, 9]. Therefore, the present study is aimed at assessing the risk of PTMC by exploring the ultrasonographic appearance of thyroid nodules.

Many thyroid carcinomas, mainly follicular cell-derived thyroid carcinomas, are caused by genetic mutations, through effects on molecular signaling pathways like MAPK and PI3K/AKT. The most commonly found mutated genes include \(BRAF, H-RAS, K-RAS, N-RAS,\) and \(PTEN\). The BRAF V600E mutation is the most common in papillary thyroid carcinoma (PTC) [10], corresponding to 28–83% of all gene mutations observed and accounting for about 90% of all \(BRAF\) mutations [11–13]. However, it has not
been found in normal thyroid tissues or benign lesions [14–18]. In addition, the BRAF V600E mutation is closely related to recurrence of papillary carcinomas and patient death. Therefore, the main objective of this study was to investigate PTMC and evaluate the correlation between its ultrasonographic appearance and the BRAF gene. According to the World Health Organization (WHO), a PTC with a maximum tumor diameter < 1.0 cm is considered a PTMC [13, 19, 20]. As defined by WHO in the 2004 edition of the Tumor Pathology and Genetics of Endocrine Organs, microcarcinoma is a subtype of thyroid carcinoma, and thyroid microcarcinoma is only referred to as a PTMC [21]. Therefore, all thyroid microcarcinomas examined in this study were PTMC.

Figure 1: Ultrasonography of thyroid nodules. (a, b) Ultrasonographic appearance of a thyroid nodule pathologically diagnosed as a nodular thyroid goiter showing hypoechogenicity, unclear borders. (c, d) Ultrasonographic appearance of a thyroid nodule pathologically diagnosed as PTMC and negative for the BRAF V600E mutation, showing hypoechogenicity and central blood flow. (e, f) Ultrasonographic appearance of a thyroid nodule pathologically diagnosed with PTMC and positive for the BRAF V600E mutation, showing hypoechogenicity, unclear borders, and aspect ratios > 1.
values were measured and the nodules were classification, and/or attenuation. Moreover, the elasticity modulus subjected to according to TI-RADS. Before surgery, each module was examined to determine if their aspect ratios were in the range of 0.5–1.0 cm, while nodules with diameters of less than 0.5 cm were excluded from this study. Additionally, the patients with two nodules were also excluded.

2. Material and Methods

2.1. Research Subjects. This study was approved by the Ethics Committee of China Medical University. Informed consent was obtained from all patients enrolled after full explanation of the purpose and nature of all procedures performed.

The study included 186 patients presenting thyroid nodules with a maximum diameter of <1.0 cm, for which preoperative BRAF V600E testing and cytopathology analysis were performed, and who underwent subsequent surgery to remove thyroid nodules at the Department of Thyroid Surgery of the First Affiliated Hospital of China Medical University from January 2017 to March 2018. None of the patients had surgical contraindications. The patients were aged 19–70 years, mean age of 38.6 ± 2.7 years, and included 143 females and 43 males. There were 100 pathologically diagnosed cases of PTMC, 80 cases of nodular goiters, and 6 cases of follicular thyroid carcinomas. All PTMC were solitary nodules. The diameters of PTMCs and nodular goiters were in the range of 0.5–1.0 cm, while nodules with diameters of less than 0.5 cm were excluded from this study. Additionally, the patients with two nodules were also excluded.

2.2. Procedures. Color Doppler ultrasound diagnostic apparatuses (Aixplorer, Supersonic Imagine, France; EPIQ7, Philips, USA) with probe frequencies of 5–12 MHz were used. The subjects were scanned in the supine position for thyroid gland to determine the size, number, location, internal echo, border, morphology, and blood flow of thyroid nodules; the nodules were examined to determine if their aspect ratios were >1 and if they had had cystic change, microcalcification, and/or attenuation. Moreover, the elasticity modulus values were measured and the nodules were classified according to TI-RADS. Before surgery, each module was subjected to fine needle aspiration for BRAF V600E testing and cytopathology analysis. The pathological diagnosis was conducted after nodular surgery. The nodule locations were classified as either in the central or peripheral regions, where the peripheral region included the isthmus and anterior, posterior, lateral, and medial margins.

Regarding the method to measure the elasticity modulus values, we used the Shear Wave Elastography (SWE) mode, in which the elastic range was set to 180 kPa, the diameter of the elasticity value measurement area was set to 5.0 mm, and the sampling frame was uniformly filled with color. During each measurement, patients were requested to hold their breath for 3–5 s, and, when stabilized, the images were stored as frames. Meanwhile, two-dimensional and elasticity images (with color coding in the range of 0–180 kPa) were displayed to determine the range and position of the region of interest (ROI) within the selected mass. Subsequently, the ROI elasticity modulus values were calculated, including the mean and minimum and maximum values in kPa. Each nodule was measured in triplicate and the results were averaged. Each patient was examined by the same methods, and the BRAF V600E testing was performed by the Amplification Refractory Mutation System (ARMS) method.

Patients with PTMC and nodular goiters were designated as the case and control groups, respectively. To diagnose thyroid nodules by either color Doppler ultrasonography, BRAF V600E testing, or cytopathology alone or by the latter two methods combined, we calculated the specificity, sensitivity, accuracy, and positive and negative predictive values; the accuracy was then statistically analyzed. Ultrasonographic appearance variables of the nodules (location, border, internal echo, morphology, microcalcification, aspect ratio > 1, attenuation, cystic change, blood flow, and elasticity modulus value) and patient age and gender were statistically compared between the two groups. The case group was further divided into two subgroups

| Classification | Pathology (number of cases) | Accuracy | P value* |
|----------------|-----------------------------|----------|----------|
| (1) Color Doppler ultrasonography | M 85 B 15 | 81.67% | >0.05 |
| | M 15 B 62 | | |
| (2) BRAF V600E testing | M 56 B 44 | 75.56% | >0.05 |
| | M 0 B 80 | | |
| (3) Cytopathology analysis | M 61 B 28 | 78.62% | >0.05 |
| | M 6 B 64 | | |
| Uncertain | M 11 B 10 | | |
| Combination of (1) and (2) | M 96 B 4 | 87.78% | <0.05 |
| | M 18 B 62 | | |
| Combination of (2) and (3) | M 83 B 13 | 88.55% | <0.05 |
| | M 6 B 64 | | |
| Uncertain | M 4 B 10 | | |

*P < 0.05: statistically different; P > 0.05: not statistically different; M: malignant nodule; B: benign nodule.
according to the BRAF V600E testing results (positive or negative). The ultrasonographic appearance variables of the nodules and patient age and gender in each subgroup were analyzed statistically to determine whether there was a correlation with BRAF V600E.

The thyroid nodules were diagnosed as benign and malignant using color Doppler ultrasonography based on the following criteria: the nodules of TI-RADS grades 3 and 4a were benign, while those of TI-RADS grades 4b, 4c, and 5 were malignant. The thyroid nodules were diagnosed as malignant or benign using BRAF V600E testing based on the following criteria: the nodules positive for BRAF V600E were malignant, while those that were negative were benign.

Each patient was examined using the same method, supervised by at least two physicians, both being associate professors, who reached an agreement through discussion in the case of controversy.

### Table 2: Correlation between the ultrasonographic appearances of PTMC and nodular thyroid goiter.

| Factor                  | Number of cases | Pathological expression | P value* |
|-------------------------|-----------------|-------------------------|----------|
|                         |                 | PTMC positive          | PTMC negative |   |
| Location                |                 | PTMC positive          | PTMC negative |   |
| Marginal region or near isthmus | 131             | 69                      | 62         | 0.194 |
| Central region          | 49              | 31                      | 18         |   |
| Gender                  |                 | PTMC positive          | PTMC negative |   |
| Female                  | 140             | 82                      | 58         | 0.128 |
| Male                    | 40              | 18                      | 22         |   |
| Age (years)             |                 | PTMC positive          | PTMC negative |   |
| ≥45                     | 97              | 51                      | 46         | 0.385 |
| <45                     | 83              | 49                      | 34         |   |
| Border echo             |                 | PTMC positive          | PTMC negative |   |
| Unclear                 | 96              | 68                      | 28         | <0.001* |
| Clear                   | 84              | 32                      | 52         |   |
| Internal echo           |                 | PTMC positive          | PTMC negative |   |
| Hypoechoic              | 152             | 92                      | 60         | 0.002* |
| Hyperechoic             | 28              | 8                       | 20         |   |
| Morphology              |                 | PTMC positive          | PTMC negative |   |
| Irregular               | 112             | 80                      | 32         | <0.001* |
| Regular                 | 68              | 20                      | 48         |   |
| Microlcalcification     |                 | PTMC positive          | PTMC negative |   |
| Yes                     | 136             | 84                      | 52         | 0.003* |
| No                      | 44              | 16                      | 28         |   |
| Aspect ratio            |                 | PTMC positive          | PTMC negative |   |
| >1                      | 56              | 36                      | 20         | 0.113 |
| <1                      | 124             | 64                      | 60         |   |
| Attenuation             |                 | PTMC positive          | PTMC negative |   |
| Yes                     | 36              | 12                      | 24         | 0.003* |
| No                      | 144             | 88                      | 56         |   |
| Cystic change           |                 | PTMC positive          | PTMC negative |   |
| No                      | 152             | 96                      | 56         | <0.001* |
| Yes                     | 28              | 4                       | 24         |   |
| Blood flow              |                 | PTMC positive          | PTMC negative |   |
| Central                 | 64              | 36                      | 28         |   |
| Marginal                | 116             | 64                      | 52         | 0.889 |
| Elasticity modulus value|                 | PTMC positive          | PTMC negative |   |
| >63 kPa                 | 89              | 60                      | 29         | 0.002* |
| <63 kPa                 | 91              | 40                      | 51         |   |

*P < 0.05: statistically different; P > 0.05: not statistically different.
specificity, sensitivity, accuracy, and positive and negative predictive values were calculated. Chi-square tests were used to compare the ultrasonographic appearance of thyroid nodules between the case and control groups and between the BRAF V600E-positive and BRAF V600E-negative subgroups of the case group. Univariate and multivariate analyses were performed by Cox proportional hazards regression models to estimate HR and 95% CI.

### 3. Results

The control group consisted of 80 patients, of whom 16 were diagnosed with thyroid nodules of TI-RADS grade 3, 46 of grade 4a, and 18 of grade 4b, according to the color Doppler ultrasonography. The BRAF V600E testing results of these thyroid nodules were all negative (Figures 1(a)–1(f)); and the cytopathology analysis diagnosed 6 cases of malignant nodules and 64 of benign, with 10 diagnostically uncertain cases. The case group consisted of 100 patients, of whom 15 were diagnosed with thyroid nodules of TI-RADS grade 4a, 67 of grade 4b, and 18 of grade 4c, according to the color Doppler ultrasonography. The BRAF V600E testing of the thyroid nodules showed positive results for 56 patients, while it was negative for 44 patients (Figures 1(a)–1(f)). The cytopathology analysis diagnosed 61 cases of malignant nodules and 28 of benign, with 11 diagnostically uncertain cases.

The color Doppler ultrasonography of the thyroid nodules alone exhibited specificity of 77.5%, sensitivity of 85%, accuracy of 81.67%, positive predictive value of 82.52%, and negative predictive value of 80.52%. Diagnosis of the nodules by BRAF V600E testing alone showed specificity of 100%, sensitivity of 56%, accuracy of 75.56%, positive predictive value of 100%, and negative predictive value of 64.52%. Diagnosis of the nodules by cytopathology analysis alone showed specificity of 91.43%, sensitivity of 68.54%, accuracy of 78.62%, positive predictive value of 90.04%, and negative predictive value of 69.57%. The combination of color Doppler ultrasonography and BRAF V600E testing showed specificity of 77.5%, sensitivity of 96%, accuracy of 87.78%, positive predictive value of 84.21%, and negative predictive value of 93.94%. The combination of BRAF V600E testing and cytopathology analysis showed specificity of 91.43%, sensitivity of 86.46%, accuracy of 88.55%, positive predictive value of 93.26%, and negative predictive value of 83.12%.

There was no significant difference \( (P > 0.05) \) in the accuracy of the PTMC diagnosis among color Doppler ultrasonography, BRAF V600E testing, and cytopathology analysis. However, each of the three methods, when used alone, was significantly different \( (P < 0.05) \) from their combinations in terms of accuracy of PTMC diagnosis, as shown in Table 1.

There was a statistically significant difference \( (P < 0.05) \) between the case and control groups for some of the Doppler ultrasonographic appearance variables, including unclear border, hypoechogenicity, irregular morphology, microcalcification, attenuation, cystic change, and elasticity modulus value. However, there was no statistical difference in location, gender, age, aspect ratio > 1, or blood flow type \( (P > 0.05) \) between the two groups, as shown in Table 2.

The univariate analysis showed that the PTMC was correlated with border echo, internal echo, morphology, microcalcification, aspect ratio, attenuation, cystic change, and elasticity modulus value. Further, the multivariate Cox regression analysis showed that the border echo (HR = 6.812, 95%CI = 2.897–16.016, \( P < 0.001 \)), internal echo (HR = 7.977, 95%CI = 3.211–19.817, \( P < 0.001 \)), morphology (HR = 6.154, 95%CI = 2.687–14.095, \( P < 0.001 \)), microcalcification (HR = 6.768, 95%CI = 2.835–16.159, \( P < 0.001 \)), attenuation (HR = 0.208, 95%CI = 0.076–0.572, \( P = 0.002 \)), cystic change (HR = 7.313, 95%CI = 2.521–21.211, \( P < 0.001 \)), and elasticity modulus value (HR = 3.074, 95%CI = 1.369–6.901, \( P = 0.006 \)) were independent predictors for diagnosis of PTMC (Table 3).

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### Table 3: Univariate and multivariate Cox regression of ultrasonographic appearances for PTMC.

| Univariate analysis | HR (95% CI) | P value | Multivariate analysis | HR (95% CI) | P value |
|---------------------|-------------|---------|-----------------------|-------------|---------|
| Location            | 0.617 (0.315–1.208) | 0.159 | —                     | —           | —       |
| Gender              | 1.728 (0.851–3.507)  | 0.130 | —                     | —           | —       |
| Age (years)         | 0.903 (0.499–1.635)  | 0.737 | —                     | —           | —       |
| Border echo         | 4.333 (2.313–8.118)  | <0.001* | 6.812 (2.897–16.016) | <0.001*     |         |
| Internal echo       | 2.697 (1.205–6.034)  | 0.016* | 7.977 (3.211–19.817) | <0.001*     |         |
| Morphology          | 4.500 (2.381–8.503)  | <0.001* | 6.154 (2.687–14.095) | <0.001*     |         |
| Microcalcification   | 2.629 (1.312–5.269)  | 0.006* | 6.768 (2.835–16.159) | <0.001*     |         |
| Aspect ratio        | 3.392 (1.417–8.121)  | 0.006* | 2.054 (0.862–4.894)  | 0.104       |         |
| Attenuation         | 0.412 (0.199–0.853)  | 0.017* | 0.208 (0.076–0.572)  | 0.002*      |         |
| Cystic change       | 4.929 (2.072–11.721) | <0.001* | 7.313 (2.521–21.211) | <0.001*     |         |
| Blood flow          | 1.091 (0.591–2.014)  | 0.781 | —                     | —           | —       |
| Elasticity modulus  | 2.638 (1.438–4.838)  | 0.002* | 3.074 (1.369–6.901)  | 0.006*      |         |

\( *P < 0.05 \): statistically different; \( P > 0.05 \): not statistically different.
There was a statistically significant difference ($P < 0.05$) between the BRAF V600E-positive and BRAF V600E-negative subgroups for some of the color Doppler ultrasonographic variables, namely, unclear border, irregular morphology, and aspect ratio $> 1$. However, there was no statistical difference ($P > 0.05$) for the other variables (location, gender, age, echo, microcalcification, attenuation, cystic change, blood flow type, and elasticity modulus value) between the two subgroups, as shown in Table 4.

The univariate analysis showed that the BRAF V600E mutation was correlated with border echo, morphology, aspect ratio, attenuation, and elasticity modulus value. Further, the multivariate Cox regression analysis showed that the border echo (HR = 17.889, 95%CI = 3.674 – 87.093, $P < 0.001$), morphology (HR = 0.07, 95%CI = 0.012 – 0.414, $P = 0.003$), and aspect ratio (HR = 23.476, 95%CI = 4.555 – 121.004, $P < 0.001$) were independent predictors for BRAF V600E mutation (Table 5).

### Table 4: Correlation between the ultrasonographic appearance of PTMC and the BRAF V600E mutation.

| Factor                   | Number of cases | BRAF V600E mutation | $P$ value* |
|--------------------------|-----------------|---------------------|------------|
|                          |                 | Positive | Negative |             |
| Location                 |                 |          |          |             |
| Marginal region or near isthmus | 68              | 38        | 30        | 0.972      |
| Central region           | 32              | 18        | 14        |            |
| Gender                   |                 |          |          |             |
| Female                   | 82              | 47        | 35        | 0.571      |
| Male                     | 18              | 9         | 9         |            |
| Age (years)              |                 |          |          |             |
| $\geq 45$                | 55              | 34        | 21        | 0.195      |
| $<45$                    | 45              | 22        | 23        |            |
| Border echo              |                 |          |          |             |
| Unclear                  | 70              | 49        | 21        | $<0.001^*$ |
| Clear                    | 30              | 7         | 23        |            |
| Internal echo            |                 |          |          |             |
| Hypoechoic               | 89              | 48        | 41        | 0.236      |
| Hyperechoic              | 11              | 8         | 3         |            |
| Morphology               |                 |          |          |             |
| Irregular                | 75              | 35        | 40        | 0.001*     |
| Regular                  | 25              | 21        | 4         |            |
| Microcalcification        |                 |          |          |             |
| Yes                      | 83              | 45        | 38        | 0.427      |
| No                       | 17              | 11        | 6         |            |
| Aspect ratio             |                 |          |          |             |
| $>1$                     | 36              | 30        | 6         | 0.001*     |
| $<1$                     | 64              | 26        | 38        |            |
| Attenuation              |                 |          |          |             |
| Yes                      | 14              | 11        | 3         | 0.067      |
| No                       | 86              | 45        | 41        |            |
| Cystic change            |                 |          |          |             |
| No                       | 92              | 49        | 43        |            |
| Yes                      | 8               | 7         | 1         | 0.061      |
| Blood flow               |                 |          |          |             |
| Central                  | 37              | 18        | 19        | 0.256      |
| Marginal                 | 63              | 38        | 25        |            |
| Elasticity modulus value |                 |          |          |             |
| $>63$ kPa                | 58              | 37        | 21        | 0.065      |
| $<63$ kPa                | 42              | 19        | 23        |            |

* $P < 0.05$: statistically different; $P > 0.05$: not statistically different.
4. Discussion

The progression of PTMC is rapid, and its incidence has significantly increased in recent years. Despite controversy over PTMC surgery, its metastasis to the central lymph nodes is indisputable, occurring at a high rate of 24–64%. Moreover, PTMC has high recurrence and mortality rates [22, 23]. Therefore, a high diagnostic accuracy is necessary in clinical settings, as surgery is the commonly accepted method of treating metastatic PMTC. However, currently, the diagnostic accuracy of PTMC is not high, and missed diagnoses and misdiagnoses occur frequently. At present, the main diagnostic method is color Doppler ultrasonography with an accuracy of 75–90% [24].

Recently, with the advances in molecular biology technology, rapid developments have occurred in the field of molecular research on thyroid carcinoma. As a research hotspot in recent years, the BRAF V600E mutation has become a new prospect for the early detection of PTMC. In fact, the BRAF V600E allele is an oncogene for thyroid carcinoma. However, the sensitivity of Braf V600E testing is very low (reaching only 45%), while its specificity is as high as 99.5% [13, 25–27]. Moreover, Li et al. [13, 28] reported that the Braf V600E mutation is closely related to extracapsular invasion of PTC, lymph node metastasis, and high TNM stage, leading to a higher capsular invasion rate [29]. PTCs with the BRAF V600E mutation are highly invasive and prone to infiltrate the tissues surrounding the thyroid gland, leading to late clinical staging [28–30] and worse prognosis [29, 31, 32]. Therefore, obtaining positive BRAF V600E testing results is a pivotal step for diagnosis. In this study, the BRAF V600E testing results and the ultrasonographic appearance of the thyroid nodules were combined to increase the accuracy of PTMC diagnosis, allowing the invasiveness of nodules to be inferred. With this combination, PTMC can be detected in a more reasonable manner to identify the nodules that are susceptible to metastasis.

In agreement with the literature [23, 25, 27], our findings revealed 100% specificity and 56% sensitivity for the BRAF V600E testing. This specificity rate means that patients positive for BRAF V600E are diagnosed with PTC without misdiagnoses. Therefore, positive results in BRAF V600E testing are of great value. However, the very low diagnostic sensitivity results in many missed diagnoses of PTC. Considering this, it is necessary to seek new markers to evaluate thyroid carcinoma. The statistical difference found between the combination of BRAF V600E testing with color Doppler ultrasonography or cytopathology and either method alone indicated that the combined diagnosis compensated for the limitations of the methods individually (i.e., the low sensitivities of BRAF V600E testing and cytopathology and the low specificity of color Doppler ultrasonography), thereby greatly enhancing the accuracy of PTMC detection and greatly reducing the proportion that cannot be determined by cytopathology analysis. For nodules that cannot be defined or tend to be benign by cytopathology analysis, once a positive result for BRAF V600E testing is determined, the nodule would be diagnosed as malignant. Therefore, BRAF V600E testing is a good auxiliary means for improving the diagnosis rate of thyroid nodules [34].

Statistically significant differences were observed in some of the ultrasonographic appearance variables (unclear border, hypoechoicogenicity, irregular morphology, microcalcification, attenuation, cystic change, and elasticity modulus value) between the control and case groups, indicating that these variables can be considered specific manifestations of thyroid malignant nodules, consistent with previously reported findings [33]. PTMC nests are not susceptible to liquefied necrosis, and cystic change is rarely seen, while they have a tendency to undergo longitudinal division. Therefore, irregular morphologies, unclear borders, and nodular attenuation are more likely to occur in microcarcinomas. However, there is no statistical difference in aspect ratios > 1, which is inconsistent with the research results of other scholars. The main reason is that the object of this

| Location | 0.985 (0.422–2.297) | 0.972 | 0.532 | 1.343 (0.483–3.733) | 0.572 | 0.196 | 1.693 (0.762–3.761) | 0.196 |
| Border echo | 7.667 (2.853–20.602) | <0.001* | 17.889 (3.674–87.093) | <0.001* |
| Internal echo | 0.439 (0.109–1.764) | 0.246 | — | — |
| Morphology | 0.167 (0.052–0.532) | 0.002* | 0.070 (0.012–0.414) | 0.003* |
| Microcalcification | 0.646 (0.218–1.910) | 0.430 | — | — |
| Aspect ratio | 7.308 (2.666–20.035) | <0.001* | 23.476 (4.555–121.004) | <0.001* |
| Attenuation | 6.668 (1.299–34.241) | 0.023* | 2.079 (0.345–12.541) | 0.425 |
| Cystic change | 0.163 (0.019–1.377) | 0.096 | — | — |
| Blood flow | 0.623 (0.275–1.413) | 0.258 | — | — |
| Elasticity modulus value | 2.513 (1.105–5.712) | 0.028* | 1.995 (0.228–17.475) | 0.533 |

*P < 0.05: statistically different; P > 0.05: not statistically different.
study is nodules with a maximum diameter of less than 1 cm, and the number of cases is relatively small. Among them, the proportion of malignant nodules with an aspect ratio greater than 1 is not significantly higher than that of benign nodules. In this study, the presence of unclear borders, irregular morphologies, and aspect ratios > 1 in the ultrasonographic appearance of the BRAF V600E-positive subgroup was significantly more common than in that of the negative group. This is also consistent with the findings of Skubisz et al. [34]. Malignant nodules with irregular morphologies are prone to mutations, meaning that it is necessary to pay attention to irregular morphologies when evaluating and classifying nodules. Moreover, unclear borders are usually accompanied by aspect ratios > 1. If such a nodule is suspected to be malignant, it should be surgically removed to prevent lymph node metastasis, because the probability that its oncogene test result will be positive is greatly increased, compared to more regular nodules. An explanation for the statistically significant difference in ultrasonographic appearance is that the cells of cancer nodules grow and are arranged in a longitudinal manner; in the early stages of thyroid carcinoma, the cancer cells in the anterior and posterior sides of the tumor undergo division, while those growing in other directions are at a relatively quiescent phase. This difference results in longer radial extension of the tumor in the anterior and posterior directions than in the right and left directions—a growth pattern totally contrasting with that of benign nodules [35]. In addition, the malignancy assessed by ultrasound was not obvious for six patients, but the BRAF V600E testing was positive, and surgery was not performed immediately. After one year, the nodule increased significantly and the anteroposterior diameter exceeded 1.0 cm, accompanied by lymph node metastasis. Therefore, BRAF V600E expression was positive and tumor invasiveness could be related [36, 37]. Further research is needed for future follow-up.

The diameter of the nodules selected in this study was between 0.5 and 1.0 cm, mainly because the study was primarily focused on papillary microcarcinomas (which have diameters < 1.0 cm). Additionally, the minimum diameter of the nodules investigated in the case group was 0.5 cm, primarily because examining smaller nodules would allow even smaller cancer nests to be used for BRAF V600E testing via fine needle aspiration, resulting in a higher false-negative rate.

Regarding the limitations of this study, as BRAF V600E testing in our hospital is performed relatively late and no large sample size was collected, a larger sample is needed to verify further our conclusions. Moreover, one patient with multiple nodules was excluded from the study and thus was not subjected to ultrasonographic analysis.

In conclusion, the combined application of BRAF V600E testing and color Doppler ultrasonography or cytopathology can improve the accuracy of PTMC diagnosis. Among the ultrasonographic observations of thyroid nodules, unclear borders, hypoechogenicity, irregular morphologies, microcalcification, attenuation, the absence of cystic change, and the elasticity modulus value can be used in combination as indicators to distinguish between benign and malignant nodules. For malignant nodules, ultrasonographic appearances with unclear borders, irregular morphologies, and aspect ratios > 1 were associated with positive results of BRAF V600E testing, suggesting that such nodules are prone to metastasis and are associated with poor prognosis. This correlative study of ultrasonographic appearances of thyroid nodules may represent a scientific foundation for artificial intelligence-based ultrasonographic diagnosis of thyroid carcinoma.

Data Availability
Data are available on request from the authors.

Conflicts 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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