The value of peri-lesional tissue stiffness and stiff rim sign in the differential diagnosis between benign and malignant breast lesions

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Abstract

Background: Shear wave elastography (SWE) is an important method in the diagnosis of breast lesions. The purpose of this study was to evaluate the value of tissue stiffness around breast lesion and stiff rim sign for the differentiation of benign and malignant lesions.

Methods: 192 patients (mean age, 44.6 ± 13.6 years) with 199 breast lesions proven by pathological examination underwent shear wave elastography (SWE). We first observed if there was a stiff rim sign. Then the shell around the breast lesion on SWE was automatically drawn by machine, with width of 1mm, 2mm and 3mm. Elasticity modulus of the lesion and surrounding tissue were recorded, including maximum elasticity (Emax), mean elasticity (Emean), minimum elasticity (Emin) and elasticity ratio (shell/lesion ratio). The optimal thresholds of elasticity modulus were calculated according to receiver operating characteristic (ROC) curve.

Results: There were 75 malignant lesions and 124 benign lesions. The average Emax, Emean of lesion and shell were significantly higher in the malignant group than in the benign group (P<0.05). The optimal cut-off value
of Emax for diagnosing malignant lesion was 101.7 Kpa, with a sensitivity of 66.3% and specificity of 87.9%. The optimal cut-off value of Emean was 29.1 Kpa, with a sensitivity of 65.3% and specificity of 79.8%. The stiff rim sign had a highest diagnostic performance for malignancy than other elastic parameters, with an accuracy of 88.4%. However, measuring peritumoral tissue stiffness can achieve a relatively high sensitivity, whereas specificity was not improved significantly.

**Conclusion** - The stiffness of tissue surrounding breast malignancy was significantly higher than benign lesion. Stiff rim sign has the potential to improve the diagnostic performance of breast lesions.

**Keywords** - Shear wave elastography, Stiff rim sign, Breast, Lesion

**Background**

Breast US elastography is a technique that can be used to assess the stiffness of lesions, which may be helpful for the differential diagnosis. Based on various literatures and evidence-based results, WFUMB Guidelines have recommended additional elastography to conventional US to improve the breast lesion characteristics [1]. Usually, there are two types of elastography methods, strain elastography (SE) or shear wave elastography (SWE) [2-3].

Compared with SE, SWE has the ability to measure lesion stiffness, so it is regard as a reproducible and quantitative means. Some studies have demonstrated that SWE showed high sensitivity and specificity for the characterization of breast lesions [4-6]. Besides the measurement of intra-lesional stiffness, there is a growing interest of measuring tissue stiffness around the breast lesion. Zhou et al pointed out the strain ratio of surrounding tissue and lesions may be helpful for breast lesions diagnosis [7]. Park et al found malignant breast lesion with a stiff rim were larger in size and associated with more aggressive pathologic subtypes [8]. The surrounding rim of higher stiffness which is characterized by desmoplastic reactions may be due to the infiltration of cancer cells into interstitial tissues or intraductal
components [9].

In our study, we used two methods to assess the stiffness of peri-lesional tissue. One was observing whether there was a stiff rim sign or not, the other was making a quantitative measurement of the stiffness around lesion (shell). Our aim was to find the value of peri-lesional stiffness in the differential diagnosis between benign and malignant breast lesions.

**Methods**

**Patients**

From June 2017 to December 2018, 192 patients (mean age 44.6 ± 13.6 years; age range, 18-73 years) with breast lesions were included in this study. The enrollment criteria were as follows: ⑴ no previous biopsy and treatment; ⑵ having complete clinical and imaging data; ⑶ lesions with qualified SW images. All patients were pathologically confirmed through US-guided core needle biopsy (CNB) or surgery. Informed consent was obtained from all patients and the study was approved by the Ethical Committee of our hospital.

**US and shear wave elastography**

Breast US examination was performed using DC-8 diagnostic US system (Mindray Medical International, Shenzhen, China) with a 5-12-MHz linear-array probes. Conventional US was first performed and images were classified according to Breast Imaging and Reporting and Data Systems (BI-RADS) lexicon including shape, margin, orientation, echo pattern, posterior acoustic features and the presence of calcification [10]. US BI-RADS grading results are judged only by ultrasound images, regardless of whether the patient receives mammography or not. After the examination of conventional US, a serious of elastography techniques were then preformed for the lesions by a specialist with at least five years experiences in breast imaging.

We switched ultrasound machine to the SWE mode. Patient was told to
hold breath for a few seconds and the probe was placed vertically on the skin surface with slight manual compression. Then the double dynamic mode was used to observe grey-scale image and SWE image. A square region of interest (ROI) was adjusted to include the whole lesion and adjacent breast tissue. Then SWE quality mode and SWE mode were displayed in order. High quality images were presented as uniform green without purple artifact. In the SWE mode, the stiffness of lesions was represented by color map from red (hard) to blue (soft). Two radiologists were blinded to pathological and clinical findings, observing and analyzing the image features. First, the stiff rim sign, defined as red or orange halo around the breast lesion, was observed (fig.1). Then we drew the border of lesion manually. Afterward, the shell just around the breast lesion was drawn, with width of 1mm (shell A), 2mm (shell B), and 3mm (shell C), respectively (fig.2). The maximum (Emax), mean (Emean), minimum (Emin) stiffness value in kPa of lesion and shell, and the elasticity ratio (shell/lesion ratio) were calculated automatically by the machine. All images were stored in the hard disk for further analysis.

Statistical analysis
We used SPSS 22.0 (SPSS, Chicago, IL) as the statistical software. SWE values were expressed as mean ± standard deviation. Quantitative data and Categorical data were analyzed using independent t-test and Chi-square test, respectively. Receiver operating characteristic (ROC) curve was constructed to determine the optimal cut-off values for Emax, Emin, Emean values and shell/lesion ratio. The sensitivity, specificity, accuracy and the area under the curve (AUC) of these parameters were also calculated. P<0.05 was considered statistically significant.

Results
Pathological diagnosis
Totally 192 patients with 199 breast lesions were enrolled in our study,
including 75 malignant lesions and 124 benign lesions. Of the malignant lesions, 61 were invasive ductal carcinoma (IDC), 8 were ductal carcinoma *in situ* (DCIS), 4 were mucinous carcinoma, 1 was invasive lobular carcinoma, and 1 was mixed metaplastic carcinoma. Of the benign lesions, 111 were fibroadenoma, 8 were intraductal papilloma, 3 were benign phyllode tumour, and 2 were inflammatory granuloma.

**Diagnostic performance of BI-RADS**

The mean diameter of lesions was $1.8 \pm 0.9$ cm. There was a significant difference of mean diameter between benign ($1.6 \pm 0.7$ cm) and malignant lesions ($2.2 \pm 1.1$ cm) ($P<0.05$). B mode BI-RADS categories were summarized in Table 1. For benign lesions, 58 were category 3, 50 were category 4a, 15 were category 4b, and 1 was category 4c. Whereas for malignant lesions, the lesion numbers classified as category 4a, 4b, 4c and 5 were 9, 13, 41, 12, respectively. If we set BI-RADS category 4a as the optimal cut-off, the sensitivity, specificity, and accuracy for diagnosing malignant lesions were 88.0%, 87.1%, and 75.1%.

**Diagnostic performance of SWE**

The Emax and Emean of lesions were significantly higher in the malignant group than in the benign group ($P<0.05$) (Table 1). According to the ROC analysis, the optimal cut-off was 101.7 Kpa for Emax showing a sensitivity of 65.3% and specificity of 87.9%. For Emean the optimal cut-off was 29.1 Kpa, with a sensitivity of 65.3% and specificity of 79.8%.

As shown in Table 1, Shell/lesion ratio was higher for malignant lesions than benign lesions, with significant difference ($P<0.05$). The Emax and Emean of shell A, shell B and shell C were also significantly higher in malignancies than in benign lesions ($P<0.05$). The sensitivity, specificity, and accuracy of the quantitative parameters concerning peri-lesional tissue stiffness for malignancy were shown in Table 2.
The number of lesions appearing stiff rim sign was 13 in benign group and 66 in malignant lesions, with significant difference (P<0.05). Stiff rim sign had a highest diagnostic performance for malignancy than other elastic modulus, with an accuracy of 88.4% (Fig.3-5). BI-RADS in combination with stiff rim sign can increase diagnostic sensitivity to 92.0% and accuracy to 82.3%.

**Discussion**

It is well known that elastography is an important method in the differential diagnosis of breast lesions. Different from strain elastography, SWE use high intensity acoustic pulse for tissue compression, causing measurable transverse shear wave. A recent meta-analysis showed the sensitivity, specificity and diagnostic odds ratio of SWE in combination with conventional US were 0.877, 0.849, and 40.164 [11].

The representative technique of SWE is virtual touch tissue imaging quantification (VTIQ) [12,13]. Our previous research showed SWVmax value had relative high diagnostic performance with a sensitivity of 74.0% and specificity of 92.1%, while setting 5.37 m/s as the optimal cut-off value. However, one major limitation of VTIQ is the ROI placement may be subjective and show inter-observer variability. In the present study, focal region include all the breast lesion and surrounding tissue. The machine automatically calculates Emax, Emax, Emin of lesion and shell, with no need for selecting ROI. Therefore, it may be a more objective method than VTIQ.

Besides the analysis of intra-tumoral elasticity, some scholars have showed interest in the elasticity of peri-tumoral tissue. Zhou et al [7] may be the first researchers to evaluate the stiffness of peri-lesional tissue. They found there was a significant high positive correlation between the strain ratio of the lesion and the strain ratio of the surrounding tissue in the malignant group (r=0.740, P<0.001). In the study by Xiao et al [14], quantitative elastography features of the lesion rim can significantly improve diagnostic specificity compared with conventional B-mode US alone. The possible
reasons of stiff rim sign and high stiffness of surrounding tissue may be due to
the microscopic environmental changes in the peri-lesional stroma of
malignant breast lesion. As we know, the elasticity is related to the content of
collagen fiber [15]. After the increase of angiogenesis and microvessel density,
cancer cells infiltrate into the stroma and cause desmoplastic reaction [16].
Another hypothesis is that the stiff peri-pesional tissue may cause attenuation
of shear wave, resulting in low shear wave amplitude of internal lesions [8].

In our study, we used two methods to evaluate the surrounding tissue
stiffness. One was to quantitatively measure the shell and shell/tumor ratio,
the other was to identify whether a stiff rim sign existed or not. The result
showed stiff rim sign yielded a satisfactory sensitivity of 88.0% and specificity
of 89.5%, which is higher than the diagnostic performance of lesion elasticity.
However, the sensitivity of most quantitative measurement data was high but
specificity was relative low. The reason of low specificity may be due to the
tissue of shell we chose in our study. The surrounding tissue of breast lesion
was glandular tissue, which had higher stiffness than fatty tissue. Therefore,
some benign lesions may show high stiffness of shell and shell/tumor ratio.
This result was in accordance with the research of Zhou et al [7]. Another
possible influence factor was that we cannot draw the outline of the lesion
precisely. Sometimes the shell may include the breast lesion tissue, thus
resulting in relatively low specificity and accuracy. On the contrary, the
intra-lesional elasticity had a low sensitivity and relatively high specificity. We
found that the sensitivity of Emax and Emean of tumor is 65.3%. This meant
about 34.7% malignant lesions would be missed if we used elasticity of
tumour value alone. False negative rate may be due to the size and type of
lesion. It has been reported that DCIS, mucinous carcinoma or small breast
carcinoma had lower SWV [17].

The stiff rim sign showed a high diagnostic performance compared with
other elastography. However, 13 lesions (eleven fibroadenomas, one
intraductal papillomas and one mastitis) were considered false positive,
whereas nine lesions (six IDCs and 3 DCISs) were considered false negative. We must acknowledge that stiff rim sign is limited in its subjectivity and image quality. Sometimes the red or orange halo around the lesion was incomplete, which was also regard as stiff rim sign. This may reduce the diagnostic sensitivity and specificity.

In our study, we found the sensitivity of BI-RADS was 88%, and its specificity was 87.1%. The high sensitivity and specificity were similar to stiff rim sign, but also higher than quantitative parameters of SWE. The diagnostic performance of SWE was not as effective as we expected. Therefore, we hold the opinion that SWE can only provide additional information of tumor elasticity, and the final diagnosis should rely on the combination of elastography with the BI-RADS classification standard. Our study showed the addition of stiff rim sign to BI-RADS can increase the accuracy (from 75.1% to 82.3%) and sensitivity (from 88.0% to 92.0%). Only three malignancies would be missed if we used combined method. The three false negative lesions were IDCs smaller than 1 cm, which was in line with the result reported by Vinnicombe [18] and Chen [19]. Some malignancies may have benign characteristics when they were small.

There were some limitations in our study. First, the malignancy in this study was mostly invasive ductal carcinoma (61/75, 81%). Therefore, we did not further analyze the elastography of different subtype. Second, we should admit that stiff rim sign was subjective to some extent. Inter-observer variability was not taken into account and could impact our results. Third, it was a single-center study and a large number of cases would be needed.

**Conclusion**

The stiffness of tissue surrounding breast malignancy was significantly higher than benign lesion. BI-RADS in combination with stiff rim sign can have a satisfactory diagnostic performance.
Abbreviations
AUC: area under the curve; BI-RADS: Breast Imaging and Reporting and Data Systems; CNB: core needle biopsy; DCIS: ductal carcinoma \textit{in situ}; IDC: invasive ductal carcinoma; ROC: receiver operating characteristic; SE: Strain elastography; SWE: Shear wave elastography; VTIQ: virtual touch tissue imaging quantification

Declarations
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Authors’ contributions
Kong WT organized the data and wrote the paper. Wang Y, Zhou WJ, Zhang YD searched and analysed the data. Zhuang XM collected the clinical data. Wang WP and Wu M revised the paper. All authors read and approved the manuscript.

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Availability of data and materials
Please contact the author for further data.

Ethics approval and consent to participate
This study was approved by Drumtower Hospital Ethics committee of Nanjing University. The study had adhered to the law of China and the 2008 Helsinki Declaration. Signed written informed consent was obtained from all participants.
Consent for publication
Not applicable.

Competing interests
The authors declare that they have no conflicts of interest.

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**Figure Legends**

**Fig.1** The stiff rim sign. The breast lesion was surrounded by red and orange halo, which may be caused by desmoplastic reaction.

**Fig.2** The sketch map of the method concerning quantitative measurement of shell. After drawing the outline of breast lesion manually, we drew the shell around lesion, with width of 1mm (shell A), 2mm (shell B), and 3mm (shell C), respectively.

**Fig.3** A fibroadenoma in a 24-year-old woman. A regular, hypo-echogenic solid breast lesion was seen on B-mode US (a), which was classified as BI-RADS 3. CDFI showed abundant blood flow signal in the lesion (b). The lesion was soft represented as blue (c, right) with a satisfactory quality of elastographic image (c, left). Quantitative measurement showed the Emax, Emean of shell and shell/lesion ratio were 45.0 Kpa, 13.3 Kpa and 0.78 (d).

**Fig.4** A invasive ductal carcinoma in a 47-year woman. B-mode US showed an irregular and heterogeneous lesion with micro-calcification (a). There was a stiff rim sign in SWE mode (b). The Emax, Emean of shell and shell/lesion ratio were 155.8 Kpa, 61.5 Kpa and 1.63 (c).

**Fig.5** A invasive ductal carcinoma in a 55-year woman. On B-mode US, the
small breast lesion (9mm) showed malignant characteristics, which can be classified as BI-RADS-4c (a). However, the lesion was soft showing almost blue on SWE mode (b), with low Emax, Emean of shell and shell/lesion ratio of 40.2 Kpa, 20.2 Kpa and 1.03, respectively.