Abstract. The aim of this study was to retrospectively determine the accuracy and feasibility of using ¹⁹⁹mTc-dextran (DX) lymphoscintigraphy for the localization of sentinel lymph nodes (SLNs) in breast cancer patients. The relevant factors affecting lymphoscintigraphy were also investigated. In this study, 235 breast cancer patients underwent ¹⁹⁹mTc-DX lymphoscintigraphic imaging and examination by a γ-probe method in combination with blue dye staining to detect SLNs. The detection results were considered in combination with rapid frozen pathology results to determine whether SLN metastasis was positive or negative. SLNs were identified in 191 patients by γ-probe detection among the 202 patients that tested positive by lymphoscintigraphic imaging, a coincidence rate of 94.6%. This suggested that lymph node metastasis had occurred and could be detected using lymphoscintigraphy. The axillary status of the breast cancer patients was also predicted using lymphoscintigraphy and the false-negative rate, sensitivity, specificity and positive predictive value were 13.3% (4/30), 90.7% (39/43), 23.4% (45/192) and 13.5% (21/155), respectively. The age of the patient, menstrual status, tumor location, tumor size, pathological type, preoperative biopsy and neoadjuvant chemotherapy were unrelated to the success of lymphoscintigraphy (P>0.05).

¹⁹⁹mTc-dextran lymphoscintigraphy can detect sentinel lymph node in breast cancer patients

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Introduction

Breast cancer is one of the most common cancers in females. The incidence of breast cancer ranks among the top two cancers in Chinese females, and is a serious threat to health (1). Lymphatic metastasis often occurs at the early stage of breast cancer, which is one of the prognosis factors and is key to determining the clinical stage and for guiding the selection of breast cancer treatment (2). Axillary lymph node dissection (ALND) in combination with pharmacology examination (¹⁹⁹mTc-dextran lymphoscintigraphy) has been considered to be the most accurate method for evaluating lymphatic metastasis; however, ALND usually leads to a series of short- or long-term complications, such as wound infection, hematoma formation, pain and limitation of shoulder activity (3). Moreover, ALND is not significant in the diagnosis of early-stage breast cancer patients who are axillary lymph node-negative, but may seriously affect the patient's quality of life (4). In the 1990s, the concept of sentinel lymph nodes (SLNs) was introduced into clinical practice. A number of studies have indicated that sentinel lymph node biopsy (SLNB) can predict axillary lymph node metastasis accurately (5,6).

In the treatment of breast cancer, SLNB is quite significant for the reduction of the upper extremity complications of patients and for the prediction of axillary lymph node state, and has gradually become an integral part of the comprehensive treatment of breast cancer. The identification and location of SLNs are the key findings of successful SLNB. Currently, the methods used for the identification and location of SLN include lymphoscintigraphy, blue dye methods, or a combination of the two methods.

This study retrospectively evaluated the accuracy of ¹⁹⁹mTc-dextran (DX) lymphoscintigraphy for the identification of SLN location in 235 consecutive cases of breast cancer in female patients, and analyzed relevant factors affecting the success of imaging.

Materials and methods

Patients. In this study, 235 consecutive cases of breast cancer in female patients diagnosed at the Affiliated Cancer Hospital of Guangxi Medical University (Nanning, China) from January
2009 to December 2012 were collected as the experimental subjects. All patients received lymphoscintigraphy prior to radical mastectomy at the Department of Nuclear Medicine of the Affiliated Cancer Hospital of Guangxi Medical University. The case inclusion criteria included the conditions as follows: i) female patients, ii) preoperative fine needle aspiration or biopsy and intraoperative frozen pathology verified breast cancer and iii) clinical stage T1-T2 phase patients. The case exclusion criteria were as follows (cases with any of the following were excluded from this study): i) patients receiving ipsilateral auxiliary trauma or previous surgery, ii) patients receiving ipsilateral breast cancer surgery, iii) pregnant or lactating patients and iv) patients with short-term relapse after radical mastectomy. Patients provided signed informed consent. Prior written and informed consent was obtained from every patient and the study was approved by the ethics review board of the Affiliated Tumor Hospital of Guangxi Medical University.

Equipment and materials. $^{99m}$Tc-DX and lyophilized dextran conjugate were provided by Beijing Senke Pharmaceutical Co., Ltd. (Beijing, China). The radiochemical purity by paper chromatography analysis was >90% and the marking rate was >95%, with a particle size of 50-200 nm. A molybdenum-technetium generator was provided by Beijing Atom Hi-Tech Co., Ltd. (Beijing, China). Single-photon emission computed tomography (SPECT) was achieved using a dual-head Discovery VH SPECT scanner purchased from GE Healthcare, Pittsburgh, PA, USA, which was configured with a low-energy high-resolution collimator. The sensitive ray energy range of the Europrobe hand-held $\gamma$ detector (Eurorad, Eckolsheim, France) was 100-1,000 keV, and 1% methylene blue (MB) was provided by Beijing Yongkang Pharmaceutical Factory (Beijing, China).

Preoperative lymphoscintigraphy for SLN location. Within 14-17 h prior to conducting the surgery, $^{99m}$Tc-DX 37-74 MBq (1-2 mCi)/0.5-1.0 ml, was injected at sites including the subcutaneous area the tumor site, the mammary areola and the area surrounding the biopsy residual cavity. Lymphoscintigraphy was performed following the injection at 15 min, 30 min, 1 h and 2 h, and at 1 h prior to surgery the next day. The SPECT acquisition conditions were set as matrix: 256x256; 1-fold magnification; peak energy, 140 keV; window width, 20%; and frame count, 5x10^3. A freshly prepared $^{99m}$TcO$_4$ point source was used as the location source, which included an active point source and four fixed-point sources located at the supraclavicular fossa, xiphoid, and bilateral areola. A collective focus of radioactive density, with the exception of that at the injection sites, was considered as positive lymph node imaging (Fig. 1A). Active point sources between the probe and imaging area helped to determine the surface location of the collective focus of density and were marked on the body surface (Fig. 1B).

Intraoperative blue dye methods and ‘hot spot’ detection. As shown in Fig. 1C, during the surgery, the patient was maintained in a supine position and 1% MB was administered by subcutaneous injection. After 5 min, SLNB was performed along the blue-stained lymph vessels to detect the stained lymph nodes. At the same time, the handheld $\gamma$ detector was used to detect hot nodules, which were those having a count 10-fold higher than the basal count. The counts of cold nodules were 10% of those of the hot nodules, and the counts of warm nodules were between those of the hot and cold nodules (Fig. 1D and E).

SLN treatment and axillary treatment methods. All the blue-stained lymph nodes and warm nodes resected during the surgery were considered as SLNs and were sent for rapid frozen section pathological biopsy. If the results suggested that SLN metastasis had occurred, routine ALND was carried out. If no metastasis was identified, the surgeon decided whether the patient could be treated with ALND. However, the lymph node detection was considered to have failed if no blue-stained lymph nodes or hot nodules were detected, and if this occurred, the patients were treated with ALND.

Evaluation criteria. The location accuracy of lymphoscintigraphy was evaluated on the basis of the criteria for evaluating SLNB (7). The following formulae were applied: i) $\gamma$-probe detection rate (%) = (SLN-positive cases/SLN cases detected) x100; ii) sensitivity = (imaging-positive cases/SLN metastasis cases); iii) false negative rate (%) = (imaging-false negative cases/SLN metastasis cases) x 100; iv) specificity (%) = (imaging-true negative cases/SLN-negative cases + SLN-false positive cases) x100; and v) positive predictive value = imaging-true positive cases/(SLN-true positive cases + SLN-false positive cases).

Statistical analysis. SPSS statistical software, version 15.0 (SPSS, Inc., Chicago, IL, USA) was used to analyze the data, which were compared using the $\chi^2$ test. Differences were considered to be statistically significant when P<0.05.

Results

Lymphoscintigraphy. The clinical characteristics of the patients are shown in Table I. The ages of the enrolled patients ranged from 24 to 77 years old, with a median age of 45 years. There were 63 cases of patients who were menopausal, and 172 cases that were premenopausal. There were 140 patients that had tumors located in the upper outer quadrant, while the remaining 95 cases had tumors in other quadrants. A tumor size ≤2 cm was found in 123 cases, and there were 112 cases with tumors >2 cm but ≤5 cm. Moreover, there were 16 cases of ductal carcinomas in situ, 185 cases of invasive ductal carcinomas, 12 cases of invasive lobular carcinoma and 22 cases of other types, including squamous cell carcinoma and mucous adenocarcinoma.

In this study, 202 patients among the 235 patients showed positive results in lymphoscintigraphy imaging. The detection rate was 86.0% (202/235). Moreover, 191 cases of these 202 patients were detected to have ‘hot nodules’ or ‘warm nodules’. In addition, there were 11 cases of patients in which no SLNs were detected by lymphoscintigraphy prior to surgery or by $\gamma$-probe methods during the surgery, but in which SLNs were detected using blue dye methods. The detecting coincidence rate was 94.6% (191/202) for lymphoscintigraphy in combination with the $\gamma$-probe method. The successfulness of
Figure 1. Examination of a breast cancer patient
(A) The left axillary lymph node was shown to be positive by lymphoscintigraphy. Arrow indicates the subcutaneous tracer injection site outside the left areola ($^{99}$mTc-DX, 37-74MBq, (1-2mCi)/0.5-1.0 ml was injected). 
(B) The node location was marked on the body surface after imaging. Arrow indicates the subcutaneous tracer injection site on the edge of the right areola. 
(C) Blue stained lymph nodes were investigated by lymphoscintigraphy. Arrow indicates the detected blue-stained lymph nodes. 
(D) Whether the blue-stained lymph nodes were hot nodes was determined. Arrow indicates $\gamma$ detector probe. 
(E) Probe following the removal of the lymph node. Arrow indicates the surgical site detected again by $\gamma$-detector probe.

Table I. Clinical characteristics of patients and the associated lymphoscintigraphy results.

| Clinical characteristics | Lymphoscintigraphy positive, n (%) | Lymphoscintigraphy negative, n (%) | $\chi^2$ | P-value |
|--------------------------|-----------------------------------|-----------------------------------|--------|---------|
| Patient age (years)      |                                   |                                   |        |         |
| ≤30                      | 5 (83.3)                          | 1 (16.7)                          | 0.054  | 0.973   |
| >30, ≤50                 | 133 (85.8)                        | 22 (14.2)                         |        |         |
| >50                      | 64 (86.5)                         | 10 (13.5)                         |        |         |
| Menstrual status         |                                   |                                   |        |         |
| Menopausal               | 52 (82.5)                         | 11 (17.5)                         | 0.491  | 0.483   |
| Premenopausal            | 150 (87.2)                        | 22 (12.8)                         |        |         |
| Tumor size               |                                   |                                   |        |         |
| T1: ≤2 cm                | 108 (87.8)                        | 15 (12.2)                         | 0.730  | 0.393   |
| T2: >2 cm, ≤5 cm         | 94 (83.9)                         | 18 (16.1)                         |        |         |
| Tumor location           |                                   |                                   |        |         |
| Upper outer quadrant     | 116 (82.6)                        | 24 (17.4)                         | 2.158  | 0.141   |
| Other quadrants          | 86 (90.5)                         | 9 (9.5)                           |        |         |
| Preoperative biopsy      |                                   |                                   |        |         |
| Yes                      | 136 (83.4)                        | 27 (16.6)                         | 2.162  | 0.141   |
| No                       | 66 (91.7)                         | 6 (8.3)                           |        |         |
| Neoadjuvant chemotherapy |                                   |                                   |        |         |
| Yes                      | 19 (95.0)                         | 1 (5.0)                           | 0.775  | 0.378   |
| No                       | 183 (85.1)                        | 32 (14.9)                         |        |         |
| Type of tumor            |                                   |                                   |        |         |
| Intraductal carcinoma    | 14 (87.5)                         | 2 (12.5)                          | 2.178  | 0.536   |
| Invasive ductal carcinoma| 157 (84.5)                        | 28 (15.5)                         |        |         |
| Invasive lobular carcinoma| 12 (100)                          | 0 (0)                             |        |         |
| Others                   | 19 (86.4)                         | 3 (13.6)                          |        |         |
lymphoscintigraphy was identified to have no association with the age of the patient, menstrual status, tumor location, tumor size, pathological type, preoperative biopsy and neoadjuvant chemotherapy (P>0.05; Table I).

Comparison of lymphoscintigraphy results with lymph node pathology results. A comparison of the results showed that there was one patient for which lymphoscintigraphic imaging gave a positive result prior to surgery, but in which blue stained lymph nodes or ‘hot spots’ were not observed during the surgery, while the pathology results following ALND found that lymph node metastasis had occurred (1/11). Moreover, there were 11 patients for whom negative results were obtained by lymphoscintigraphy prior to surgery, but blue-stained lymph nodes were detected during surgery. Intraoperative frozen section pathology and postoperative routine pathology confirmed that metastasis to the blue-stained lymph node had not occurred in these 11 patients, nor to the lymph nodes obtained from ALND. A further five patients (5/33) received negative lymphoscintigraphy imaging results and were not found to have SLNs during the surgery; however, following ALND, pathology results indicated that axillary lymph node metastasis had occurred.

Lymph node pathology results and axillary treatment. Pathology results demonstrated that there were 43 cases of SLN-positive patients and 159 cases of SLN-negative patients, with 17 cases that were axillary node-positive and 215 cases that were axillary node-negative. A total of 55 patients received only SLN resection, and the intraoperative frozen pathology biopsy showed that all these 55 cases were SLN-negative. Thus, they were not treated with ALND. A total of 180 cases received ALND, and pathology results suggested that 23 of them were SLN-positive but axillary node-negative.

Evaluation of the positioning accuracy of lymphoscintigraphy. Based on the evaluation criteria, the SLNB results obtained using $^{99m}$Tc-DX lymphoscintigraphy were analyzed. The detection rate of the $\gamma$ probe in patients that were positive by lymphoscintigraphy was 94.6% (191/202), and the false negative rate for predicting axillary status was 13.3% (4/30). Moreover, the sensitivity was 90.7% (39/43), the specificity was 23.4% (45/192) and the positive predictive value was 13.5% (21/155). These results suggest that the use of the $\gamma$-probe method in lymphoscintigraphy can localize SLN accurately for SLNB, but is not suitable for use in the determination or prediction of axillary lymph node metastasis.

Discussion

In the present study, lymphoscintigraphic imaging in combination with $\gamma$-probe analysis had an SLN detection rate of 94.6% (191/202), and the false negative rate for predicting axillary status was 13.3% (4/30). Moreover, the sensitivity was 90.7% (39/43), the specificity was 23.4% (45/192), and the positive predictive value was 13.5% (21/155). Due to the role of macrophage phagocytosis in lymph nodes, lymphoscintigraphic tracers were retained within the SLNs to enable the radiographic imaging. The lymph node contents as well as the distribution, shape, size and functional status of lymphatic vessels may be observed by lymphoscintigraphic imaging (8-11). Thus, this method can diagnose the lymph node metastasis of malignant tumors, and also determine pathological changes in the lymphatic system caused by a benign condition (12-14).

The detection rate of lymphoscintigraphy is ~90-97%, which is similar to that reported in the majority of literature (15,16), indicating that the use of a $\gamma$ probe in lymphoscintigraphy may provide accurate SLN localization for SLNB. However, the SLN imaging of the patients conducted using lymphoscintigraphy is not fully developed. Statistical analysis demonstrated that the success of lymphoscintigraphy had no correlation with patient age, menstrual status, tumor location, tumor size, pathological type, preoperative biopsy and neoadjuvant chemotherapy (P>0.05). However, relevant factors include the injection site and the degree of lymph node invasion (17,18).

The enrolled patients were all treated with the same radioactive tracers with the same volume of injection dose and the same imaging conditions so that the success rate of lymphoscintigraphy was less affected by these factors (19-21). $^{99m}$Tc-DX lymphoscintigraphy is able to accurately locate breast SLNs to guide breast SLN location with the use of a $\gamma$ probe and SLNB (22-24). However, individualized treatment for patients should be considered in order to improve the success rate of imaging and better guide SLNB.

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References

1. Mohan S and Ponnusankar S: Newer Therapies for the Treatment of Metastatic Breast Cancer: a Clinical Update. Indian J Pharm Sci 75: 251-251, 2013.
2. Lucci A, Hall CS, Lodhi AK, et al: Circulating tumour cells in non-metastatic breast cancer: a prospective study. Lancet Oncol 13: 688-695, 2012.
3. Valente SA, Levine GM, Silverstein MJ, et al: Accuracy of predicting axillary lymph node positivity by physical examination, mammography, ultrasonography, and magnetic resonance imaging. Ann Surg Oncol 19: 1825-1830, 2012.
4. Scarrano AM, Eiada R, Jacks LM, Kulkarni SR and Crystal P: Accuracy of unenhanced MR imaging in the detection of axillary lymph node metastasis: study of reproducibility and reliability. Radiology 262: 425-434, 2012.
5. Hung T, Piris A, Lobo A, et al: Sentinel lymph node metastasis is not predictive of poor outcome in patients with problematic nodal melanocytic tumors. Hum Pathol 44: 87-94, 2013.
6. McMasters M, Tuttle M, Carlson J, et al: Sentinel lymph node biopsy for breast cancer: A suitable alternative to routine axillary dissection in multi-institutional practice when optimal technique is used. J Clin Oncol 18: 2560-2566, 2000.
7. Clough KB, Nasr R, Nos C, Vieira M, Inguenault C and Poulet B: New anatomical classification of the axilla with implications for sentinel node biopsy. Br J Surg 97: 1659-1665, 2010.
8. Schwartz GF, Giuliano AE and Veronesi U: Consensus Conference Committee: Proceeding of the consensus conference of the role of sentinel lymph node biopsy in carcinoma or the breast April 19-22, 2001, Philadelphia, PA, USA. Breast J 8: 124-138, 2002.
9. Krag DN, Anderon SJ, Julian TB, et al: Sentinel-lymph-node resection compared with conventional axillary-lymph-node dissection in clinically node-negative patients with breast cancer: overall survival findings from the NSABP B-32 randomised phase 3 trial. Lancet Oncol 11: 927-933, 2011.
10. Goyal A, Newcombe RG, Chhabra A and Mansel RE; ALMANAC Trialists Group. Factors affecting failed localisation and false-negative rates of sentinel node biopsy in breast cancer—results of the ALMANAC validation phase. Breast Cancer Res Treat 99: 203-208, 2006.
11. Mátrai Z, Tóth L, Saeki T, et al: The potential role of SPECT/CT in the preoperative detection of sentinel lymph nodes in breast cancer. Orv Hetil 152: 678-688, 2011.
12. Pesek S, Ashikaga T, Krag LE and Krag D: The false-negative rate of sentinel node biopsy in patients with breast cancer: a meta-analysis. World J Surg 36: 2239-2251, 2012.
13. Shimazu K, Tamaki Y, Taguchi T, Takamura Y and Noguchi S: Comparison between periareolar and peritumoral injection of radiotracer for sentinel lymph node biopsy in patients with breast cancer. Surgery 131: 277-286, 2002.
14. Borgstein PJ, Meijer S, Pijpers RJ and van Diest PJ: Functional lymphatic anatomy for sentinel node biopsy in breast cancer: echoes from the past and the periareolar blue method. Ann Surg 232: 81-89, 2000.
15. Ogasawara Y, Yoshitomi S, Sato S, and Doihara H: Clinical significance of preoperative lymphoscintigraphy for sentinel lymph node biopsy in breast cancer. J Surg Res 148: 191-196, 2008.
16. Sabaté-Llobera A, Benítez-Segura A, Marí A, et al: Lymphoscintigraphy in oral squamous cell carcinoma sentinel node biopsy and its role in the surgical planning. Clin Nucl Med 39: e142-e145, 2014.
17. Noguchi M: Sentinel lymph node biopsy as an alternative to routine axillary lymph node dissection in breast cancer patients. J Surg Oncol 76: 144-156, 2001.
18. Uren RF, Howman-Giles R, Renwick SB and Gillett D: Lymphatic mapping of the breast: locating the sentinel lymph nodes. World J Surg 25: 789-793, 2001.
19. Kraft O and Havel M: Sentinel lymph nodes and planar scintigraphy and SPECT/CT in various types of tumours. Estimation of some factors influencing detection success. Nucl Med Rev Cent East Eur 16: 17-25, 2013.
20. Martínez-Rodríguez I, De Arcocha Torres M, Banzo I, et al: Evaluation of the contribution of the dynamic phase of lymphoscintigraphy to the detection of sentinel lymph node in breast cancer. Q J Nucl Med Mol Imaging 57: 296-300, 2013.
21. Brouwer OR, Vermeer L, van der Ploeg IM, et al: Lymphoscintigraphy and SPECT/CT in multicentric and multifocal breast cancer: does each tumour have a separate drainage pattern? Results of a Dutch multicentre study (MULTISENT). Eur J Nucl Med Mol Imaging 39: 1137-1145, 2012.
22. Nelson KP, Choudhury KR, Coleman RE, Shipes SW, Siler WL, Hubble WL and Wong TZ: Does the preparation and utilization of 99mTc-sulfur colloid affect the outcomes of breast lymphoscintigraphy? J Nucl Med Technol 41: 92-98, 2013.
23. Yeung HW, Cody III HS, Turlakow A, et al: Lymphoscintigraphy and sentinel node localization in breast cancer patients: A comparison between 1-day and 2-day protocols. J Nucl Med 42: 420-423, 2001.
24. Tanis PJ, van Sandick JW, Nieweg OE, Valdés Olmos RA, Rutgers EJ, Hoenfagel CA, and Kroon BB: The hidden sentinel node in breast cancer. Eur J Nucl Med 29: 305-310, 2002.