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## Imaging Tumor-Induced Sentinel Lymph Node Lymphangiogenesis with LyP-1 Peptide

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### Abstract

Lymphangiogenesis in tumor-draining lymph nodes (LNs) starts before the onset of metastasis and is associated with metastasis to distant LNs and organs. In this study, we aimed to visualize tumor induced lymphangiogenesis with a tumor lymphatics-specific peptide Lyp-1. The LyP-1 peptide was labeled with a near-infrared fluorophore (Cy5.5) for optical imaging. At days 3, 7, 14 and 21 after subcutaneous 4T1 tumor inoculation, Cy5.5-LyP-1 was administered through the middle phalanges of the upper extremities of the tumor-bearing mice. At 45 min and 24 h postinjection, brachial LN fluorescence imaging was performed. *Ex vivo* fluorescence images were acquired for quantitative analysis of the fluorescence intensity. Tumor induced lymphangiogenesis was confirmed by LYVE-1 immunostaining and increased size of tumor side brachial LNs. Cy5.5-LyP-1 staining in LNs co-localized with LYVE-1, suggesting lymphatics-specific binding of LyP-1 peptide. The brachial LNs were clearly visualized by optical imaging at both time points. The tumor side LNs showed significantly higher fluorescence intensities than the contralateral brachial LNs at days 7, 14, and 21, but not day 3 after tumor inoculation. At day 21 after tumor inoculation, the average signal of tumor-draining LNs was  $78.0 \pm 2.44$ ,  $24.3 \pm 5.43$ ,  $25.6 \pm 0.25$  ( $\times 10^3$  photon/cm<sup>2</sup>/s) using Cy5.5-LyP-1, Cy5.5-LyP-1 with blocking, and Cy5.5 only, respectively. Tumor-draining brachial LNs showed extensive growth of lymphatic sinuses throughout the cortex and medulla. Use of LyP-1 based imaging probes with optical imaging offers a useful tool for the study of tumor-induced lymphangiogenesis. LyP-1 may serve as a marker of lymphangiogenesis useful in detecting “high risk” lymph nodes before tumor metastasis and after micro-metastasis, as well as for screening potential anti-lymphatic therapies.

### Keywords

Lymphangiogenesis; lymph node; LyP-1 peptide; optical imaging

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## Introduction

Regional lymph node metastasis represents the first step of tumor dissemination for a variety of common human cancers, such as carcinomas of the breast, colon, and prostate as well as melanoma (Tammela and Alitalo 2010). One mechanism that promotes metastasis to regional lymph nodes is tumor-induced lymph node (LN) lymphangiogenesis, which usually precedes metastasis and leads to increased tumor spread to distal LNs and further to distal organs (Harrell et al. 2007; Ji 2009). It has been found that tumor-associated lymphangiogenesis correlates with metastasis to tumor-draining LNs in several mouse models (Mandriota et al. 2001; Stacker et al. 2001). The lymphangiogenesis of LNs has been shown to be associated with the presence of non-sentinel LN metastases in animal tumor models (Harrell et al. 2007; Hirakawa et al. 2007; Hirakawa et al. 2005; Qian et al. 2006) and human breast cancer and melanoma (Van den Eynden et al. 2007). The induction of sentinel LN lymphangiogenesis may be driven by VEGF-A and VEGF-C in transgenic mouse models (Hirakawa et al. 2007; Hirakawa et al. 2005), which suggests that tumor-secreted factors are also strong mediators for local LN lymphangiogenesis (Ji 2009) and that increased sentinel LN lymphangiogenesis is involved in lymphatic spread of tumor cells (Ishii et al. 2010). In mouse models of skin carcinogenesis in which VEGF-A or VEGF-C was overexpressed in the skin, lymphangiogenesis was found to occur not only at the tumor site but also in the tumor-draining LNs, and to correlate with metastasis to lymph nodes and beyond (Hirakawa et al. 2007).

Conventional *in vivo* imaging approaches for cancer metastases in patients have focused on the detection of the cancer cells themselves (Jaffer and Weissleder 2005; Sharma et al. 2008). These methods have limited sensitivity, because relatively large anatomical sizes are required for reliable detection (Winnard et al. 2008). In addition, current imaging approaches to evaluate lymph node metastasis mainly rely on the size and the shape of the involved lymph node (Qian et al. 2006). However, nodal metastases are often microscopic, so neither computed tomography (CT) nor standard magnetic resonance (MR) imaging can rule them out reliably (Eiber et al. 2010). Thus, the ability to detect LN lymphangiogenesis may serve as a novel way of predicting LN metastasis. Furthermore, imaging lymphangiogenesis *in vivo* will be of great help in understanding LN lymphangiogenesis, as well as in developing therapeutics to control metastatic tumor cell dissemination through the lymphatics.

Therefore, various groups have performed studies to develop lymphatic imaging probes that would exceed the capabilities of the established “blue dye” procedure, and to improve identification and mapping of lymph nodes, especially sentinel lymph nodes during surgery (Ballou et al. 2007; Kosaka et al. 2009; Ting et al. 2010). Targeted optical probes for lymphatic imaging have also been developed, using a near-infrared (NIR) fluorescent indotricarbocyanine (Cy7) dye, that are based on monoclonal antibody recognition of endothelial ligands for the lymphocyte homing receptor L-selectin, MECA-79. These new probes have provided specific lymph node NIR fluorescent signals detectable within minutes after tail vein injection, and stable imaging has persisted for more than 24 h (Licha et al. 2005). Recently, in mouse models of LN lymphangiogenesis, Mumprecht *et al.* (Mumprecht et al. 2010) found that systemically injected antibodies against lymphatic epitopes accumulated in the lymphatic vasculature in tissues and LNs. Using an  $^{124}\text{I}$ -labeled antibody against the lymphatic vessel endothelial hyaluronan receptor-1 (LYVE-1), they found that anti-LYVE-1 immuno-PET enabled visualization of lymphatic vessel expansion in LNs bearing metastases, even though the metastases were not detected by  $^{18}\text{F}$ -FDG PET.

Compared with antibodies, peptide-based imaging probes with much smaller molecular size would allow faster clearance and lower immunogenicity. With peptidic phage display

screening on MDA-MB-435 xenografts, Laakkonen *et al.* (Laakkonen et al. 2002) identified a cyclic 9-amino-acid peptide (CGNKRTRGC), LyP-1. They found that LyP-1 peptide targeted to lymphatic endothelial cells instead of blood vessel endothelial cells, suggesting that this peptide could be used as a tumor lymphatics specific marker. Intravenous injection of fluorescein-conjugated LyP-1 led to good accumulation in primary MDA-MB-435 xenografts and their metastases. In addition, systemic LyP-1 peptide treatment of xenografted tumor mice inhibited tumor growth (Laakkonen et al. 2004). In this study, we explored the feasibility of LyP-1 peptide probes for LN lymphangiogenesis imaging.

## Materials and Methods

### General

Cy5.5-*N*-hydroxysuccinimide ester (Cy5.5-NHS) was purchased from GE Healthcare (Piscataway, NJ). LyP-1 peptide was purchased from C.S. Bio (Menlo Park, CA). All other solvents and chemicals were purchased from Sigma-Aldrich (St. Louis, MO).

### Synthesis of Cy5.5-LyP-1 conjugate

LyP-1 peptide (1 mg, 1  $\mu$ mol) dissolved in 100  $\mu$ L of DMSO was mixed with Cy5.5-NHS (GE Healthcare, Piscataway, NJ) (1 mg, 0.89  $\mu$ mol) and diisopropylethylamine (DIEA, 2  $\mu$ L). The reaction mixture was stirred in darkness at room temperature for 1 h and then quenched by 10  $\mu$ L of acetic acid. The crude product was purified by reversed-phase high-performance liquid chromatography (RP-HPLC) on a C<sub>18</sub> Vydac protein and peptide column (218TP510; 5  $\mu$ m, 250  $\times$  10 mm). The fractions containing Cy5.5-LyP-1 were collected, lyophilized, and dissolved in saline at a final concentration of 1 mg/mL. For analytical HPLC, a C<sub>18</sub> Vydac 218TP54 column (5  $\mu$ m, 250  $\times$  4.6 mm) was used and the flow was set at 1 mL/min using a gradient system starting from 95% solvent A (0.1% trifluoroacetic acid (TFA) in water) and 5% solvent B (0.1% TFA in acetonitrile (CAN)) (0–2 min) and ramped to 35% solvent A and 65% solvent B at 32 min. The Cy5.5-LyP-1 peptide had a retention time of 16.05 min. The identity of Cy5.5-LyP-1 was confirmed by MALDI-TOF MS (*m/z*: observed 1,890.70 [M + H<sup>+</sup>], calculated 1,888.15) (Figure 1A). For serum stability assay, 0.5  $\mu$ g of Cy5.5-LyP-1 peptide was added to 100  $\mu$ L mouse serums and incubated at 37 °C for various periods of time up to 24 h and aliquots were taken and quenched by adding 100  $\mu$ L acetonitrile. Each aliquot was centrifuged at 5000 rpm for 5 min. The supernatant (10  $\mu$ L) was analyzed by LC-MS.

### Tumor model development

The 4T1 murine breast cancer cell line was derived from a spontaneously arising BALB/c mammary tumor (Aslakson and Miller 1992). The rapid and efficient metastasis to organs affected in human breast cancer makes the 4T1 model an excellent mouse model for the study of metastatic progression of breast cancer in humans. Moreover, the model is syngeneic in immunocompetent BALB/c mice, which can be used to study the role of the immune system in tumor growth and metastasis (Tao et al. 2008). The cell line was obtained from the American Type Culture Collection (ATCC) (Manassas, VA) and maintained at 37 °C in a humidified atmosphere containing 5% CO<sub>2</sub> in Dulbecco's Modification of Eagles Medium (DMEM) with 10% fetal bovine serum (FBS). The tumor model was developed in 4–6 week-old female BALB/C mice (Harlan Laboratories) by subcutaneous injection of 5  $\times$  10<sup>6</sup> of 4T1 cells to the right shoulder of each mouse. Tumor growth was followed by caliper measurements made perpendicular to the tumors. Animal procedures were performed according to a protocol approved by the National Institutes of Health Clinical Center Animal Care and Use Committee (NIH CC/ACUC).

### In vivo optical imaging with Cy5.5-LyP-1

At days 3, 7, 14 and 21 after inoculation with 4T1 cells, tumor-bearing BALB/C mice (n = 5/group), under isoflurane anesthesia, were injected with either Cy5.5-LyP-1 (0.8 nmol) or Cy5.5 (0.8 nmol) via the middle phalanges of the upper extremities of both sides. At 45 min after injection, *in vivo* fluorescence imaging was performed with a Maestro II small-animal *in vivo* imaging system (CRI, Woburn, MA). A filter set with 615–665 nm excitation and long-pass >600 nm emission was used to detect Cy5.5 fluorescent signal. The fluorescence images consisting of Cy5.5 and autofluorescence spectra were then unmixed, based on their spectral patterns, using the manufacturer's software (Maestro software, CRI). At 24 h post injection, the mice were euthanized and a dorsal skin flap was elevated to expose the brachial lymph nodes on both sides for reimaging of fluorescence. Then the LNs were removed, and *ex vivo* fluorescence images were obtained to measure the fluorescence intensity. To study the lymph node specificity of LyP-1 peptide, Cy5.5-LyP-1 was co-injected with 300 µg of un-conjugated LyP-1 peptide as blocking agent.

### Histological evaluation and immunofluorescence staining

At 24 h after Cy5.5-LyP-1 injection, brachial lymph nodes from both sides were removed and embedded in optimum cutting temperature (O.C.T.) formulation (Sakura Finetek, Torrance, CA). Then the lymph nodes were sectioned into 4 µm slices and observed with an Olympus IX81 fluorescence microscopy (Olympus Inc., Melville, NY) equipped with an excitation band-pass filter (530–585 nm) and emission band-pass filter (605–680 nm). The same lymph node sections were also stained against lymphatic endothelial marker LYVE-1. Rabbit anti-mouse LYVE-1 antibody (1:100; Abcam) and FITC-conjugated goat anti-rabbit secondary antibody (1:400; Jackson ImmunoResearch Laboratories) were used. H&E staining was also performed on lymph node sections.

### Measurement and statistical analysis

Fluorescence intensity was measured by drawing regions of interest (ROIs) over removed lymph nodes. Quantitative data were expressed as means ± SD. Means were compared using a Student's t test. P values <0.05 were considered statistically significant.

## Results

### Optical imaging of lymphangiogenesis

Cy5.5-LyP-1 peptide showed high serum stability. Even after 24 hrs incubation, we did not observe apparent degradation of the peptide as determined by LC-MS (Figure 1B). Then, we established the tumor model by subcutaneous injection of murine breast cancer 4T1 cells on the right shoulder of BALB/C mice. In this model, the lymph flow drains into the collateral brachial lymph node, i.e. tumor-draining LN. Compared with the contralateral brachial LN, the tumor-draining brachial LN was significantly enlarged at days 7, 14 and 21 after inoculation (Figure 2). At 45 min after Cy5.5-LyP-1 injection via the middle phalanges of both upper extremities, brachial LNs on both sides were detectable with *in vivo* optical imaging. At day 3 after tumor cell inoculation, we did not observe significant fluorescence intensity increase in the tumor-draining LNs over the contralateral brachial LN. From day 7, fluorescence intensity of the tumor-draining brachial LN was progressively elevated. Meanwhile, there was no apparent fluorescent signal change in the contralateral LN (Figure 3A & B). At the 24 h time point, the primary tumor was removed for optimal LN visualization. The tumor-draining LN still retained a high fluorescent signal after removal of the primary tumor, especially in mice at 21 days after tumor inoculation. No apparent changes were observed in the contralateral LNs (Figure 3C & D).

*Ex vivo* optical imaging of brachial LNs is shown in Figure 4. Both the size and the fluorescence intensity of the tumor-draining LNs increased along with the time after tumor inoculation from day 7 to day 21. The fluorescence intensities were  $0.024 \pm 0.002$ ,  $0.038 \pm 0.002$ ,  $0.048 \pm 0.002$  and  $0.106 \pm 0.002 \times 10^6$  photon/cm<sup>2</sup>/sec respectively at days 3, 7, 14 and 21 after tumor cell inoculation, which were 1.02, 1.63, 2.04, and 4.52-fold higher than in the contralateral LNs (Figure 4B).

In order to confirm the specificity of LyP-1 peptide for lymphoangiogenesis imaging, optical imaging with Cy5.5 dye only was performed using the same tumor model as described above. Cy5.5 dye also showed LN uptake and retention at 24 hr after local injection (Figure 5). However, Cy5.5-LyP-1 showed significantly higher LN uptake compared with Cy5.5 ( $P < 0.05$  at day 14 and  $P < 0.01$  at day 21). More importantly, average signal of Cy5.5-LyP-1 in the LNs also showed significant increase from day 14 to day 21 while there was no apparent change with Cy5.5 only along with time. We also performed optical imaging using Cy5.5-LyP-1 with the presence of extra amount of cold LyP-1 peptide. At day 21, the average signal in the LNs after blocking is similar to that of Cy5.5 only, which is significantly lower compared with unblocked Cy5.5-LyP-1 imaging (Figure 5).

### Histological analysis of tumor induced lymphangiogenesis

To further demonstrate the lymphatic targeting of LyP-1 peptide, the tumor-draining brachial lymph nodes were sectioned after *ex vivo* fluorescence imaging, and fluorescent signals from Cy5.5-LyP-1 were observed with microscopy. As shown in Figure 6, Cy5.5-LyP-1 had a vessel-oriented distribution, and the abundance increased along with time after tumor inoculation. Since LYVE-1 is one of the most specific and widely used lymphatic endothelial markers, we stained the LN sections with anti-LYVE-1 antibody and visualized the staining with an FITC-conjugated secondary antibody. At day 3 after tumor inoculation, the tumor-draining LN showed few lymphatic sinuses. At day 7, the tumor-draining LN showed sparse lymphatic sinuses restricted to the cortex. Lymphangiogenesis was consistently observed from day 14 to day 21. At day 21, extensive lymphangiogenesis throughout the cortex and medulla was observed.

Interestingly, Cy5.5-LyP-1 showed a distribution pattern in the tumor-draining LNs that was very similar to the staining pattern of anti-LYVE-1 within these same nodes, confirming the lymphatic endothelium's recognition of the LyP-1 peptide. We also noticed that Cy5.5-LyP-1 distribution and anti-LYVE-1 staining did not fully overlap, indicating that LyP-1 does not necessarily bind to LYVE-1.

### Discussion

Lymphangiogenesis involves new lymphatic budding, sprouting, and remodeling. Moreover, lymphangiogenesis requires lymphatic endothelial cell (LEC) differentiation to form mature lymphatic networks, and this complicated process is regulated by multiple growth factors, cytokines and chemokines (Ji 2009). Previous studies have demonstrated that lymphangiogenesis in the LNs in the postnatal period is considered to be a major contributor to tumor metastasis (Hirakawa et al. 2009; Ji 2006). Since sentinel LNs are the first destinations in the spreading of cancer in many malignancies, lymphangiogenesis in the sentinel LNs may actively promote lymphatic metastasis. In a B16F10 melanoma model, greatly increased and enlarged lymphatic sinuses were found to be distributed throughout the cortex and medulla in tumor-draining LNs, preceding tumor metastasis (Harrell et al. 2007). Consistent with their findings, we also identified apparent lymphangiogenesis in tumor-draining LNs in a subcutaneous 4T1 murine breast cancer model. The enlargement of tumor-draining LNs, but not of the contralateral LNs, was observed at day 7 after tumor cell inoculation. LYVE-1 is a hyaluronic acid receptor with sequence similarity to CD44

expressed on lymphatic endothelial cells (Banerji et al. 1999). Anti-LYVE-1 immunostaining confirmed that extensive lymphangiogenesis happened in the tumor-draining LN. At the early stage of tumor growth, the tumor-draining LNs showed lymphatic sinus expansion mainly in medullary sinuses, and the LN lymphangiogenesis was in the cortical sinuses. With time, enlarged lymphatic sinuses and lymphangiogenesis distributed throughout the cortex and medulla, even though we did not identify tumor metastases in the tumor-draining LNs by H & E staining at day 21 after tumor inoculation.

In clinical practice, lymphatic mapping employs radiocolloid solutions (Schirrmester et al. 2004) or vital dyes (Suzuki et al. 2009). These strategies work well to provide information about the localization and function of lymph nodes. However, the signals are transient and only provide nonspecific visualization of the lymphatic system. Recent developments in optical imaging in conjunction with various molecular markers offer help in better understanding of the mechanisms underlying lymphangiogenesis and lymphatic metastases of tumors. For example, fluorescently labeled monoclonal anti-mouse LYVE-1 antibody has been utilized to monitor cancer cell trafficking in the lymphatic system in real time (McElroy et al. 2009). The recent identification of relatively specific peptides for lymphatic endothelium allowed analyses of lymph node lymphangiogenesis as well as tumor lymphangiogenesis (Laakkonen et al. 2004; Laakkonen et al. 2002; Laakkonen et al. 2008). Intravenously injected FITC conjugated LyP-1 peptide was observed to home to tumor-associated lymphatic vessels and tumor cells in MDAMB-435 xenografts and some other tumors (Laakkonen et al. 2004; Laakkonen et al. 2002; Laakkonen et al. 2008). However, due to the poor tissue penetration of photons and high background of FITC, FITC-LyP-1 is not an ideal probe for lymphangiogenesis imaging *in vivo*.

After conjugation of the tumor lymphatic endothelium specific LyP-1 peptide with NIR fluorescent dye Cy5.5, we successfully visualized tumor-draining LN *in vivo* at different stages of tumor growth. The optical imaging revealed that tumor growth in the shoulder was associated with increased lymphangiogenesis in the brachial lymph node. From day 7 to day 21, the size of tumor-draining LNs did not change dramatically, while the fluorescent intensity kept increasing, indicating that the lymphatic vessel density increased along with time. This was confirmed by immunostaining with anti-LYVE-1 antibody (Figure 6). Moreover, at 24 hr after Cy5.5-LyP-1 injection, we still could identify much stronger fluorescent signals inside tumor-draining LNs than Cy5.5-LyP-1 blocked with cold compound and Cy5.5 dye only, confirming specific detection of lymphangiogenesis with this peptide probe. The results demonstrated that targeted binding of Cy5.5-LyP-1 facilitated visualization of the expansion of lymphatic network within the tumor-draining sentinel LNs, even before tumor metastasis occurred.

We also noticed that the fluorescent signal did not homogeneously distribute within the tumor-draining LNs at all stages. We speculated that this phenomenon resulted from the unique structure and pattern of lymph flow in the LNs. Within a LN, each afferent lymphatic collects lymph from a different drainage field, and each lobule is potentially exposed to different concentrations of cytokines and stimulators. Thus, lobules within the same lymph node may have different levels of immunological activity and the cortical, paracortical and medullary compartments composed of these lobules will not necessarily have a uniform appearance (Sainte-Marie et al. 1982; Willard-Mack 2006). In this study, variation in the stimulation received from 4T1 tumor was likely the reason that the lymphangiogenesis within the tumor-draining LNs varied in different lobules, as reflected by both optical imaging and fluorescent immunostaining.

## Conclusion

In summary, we successfully visualized lymphangiogenesis in tumor-draining LNs, along with tumor growth, using Cy5.5-labeled LyP-1 peptide. This new technology offers unique advantages over current technologies for *in vivo* prediction and detection of lymph node metastasis. It enables a strategy of demonstrating “high risk” lymph nodes before tumor metastasis or after micro-metastasis occurs. It should be useful in both development of anti-lymphatic therapeutics and advancement of our understanding of the interactions between primary tumors and tumor-draining lymph nodes.

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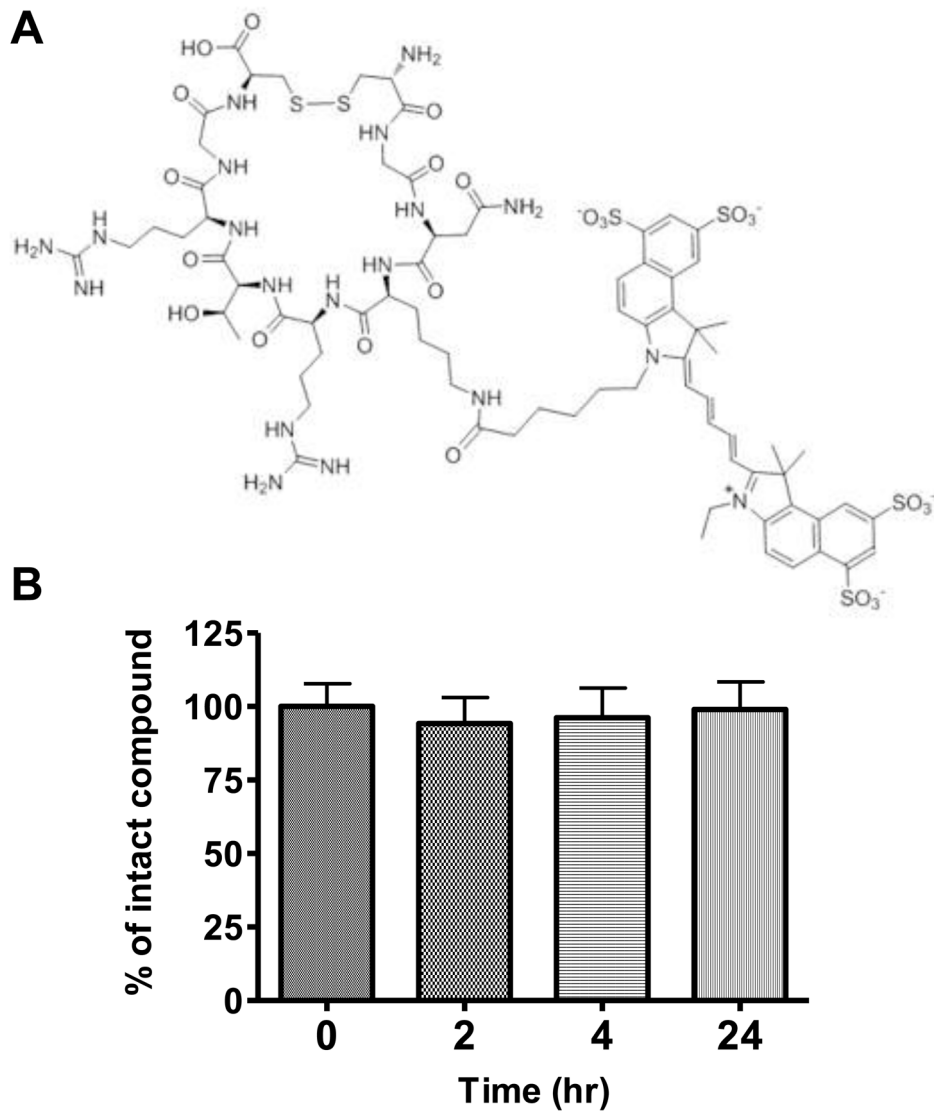
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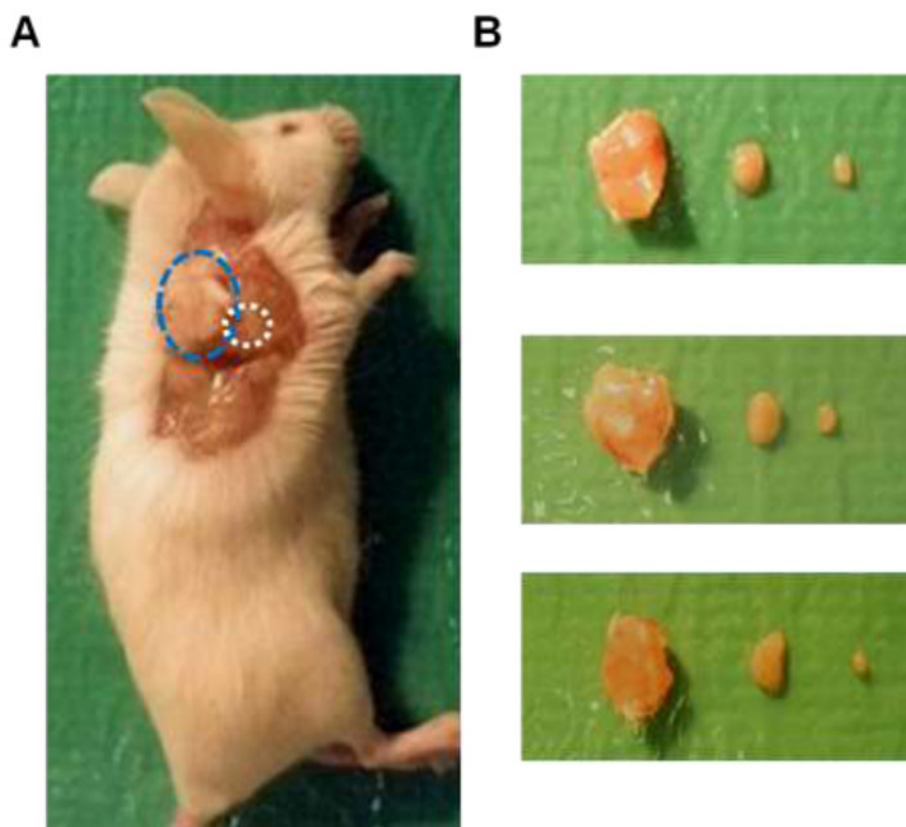
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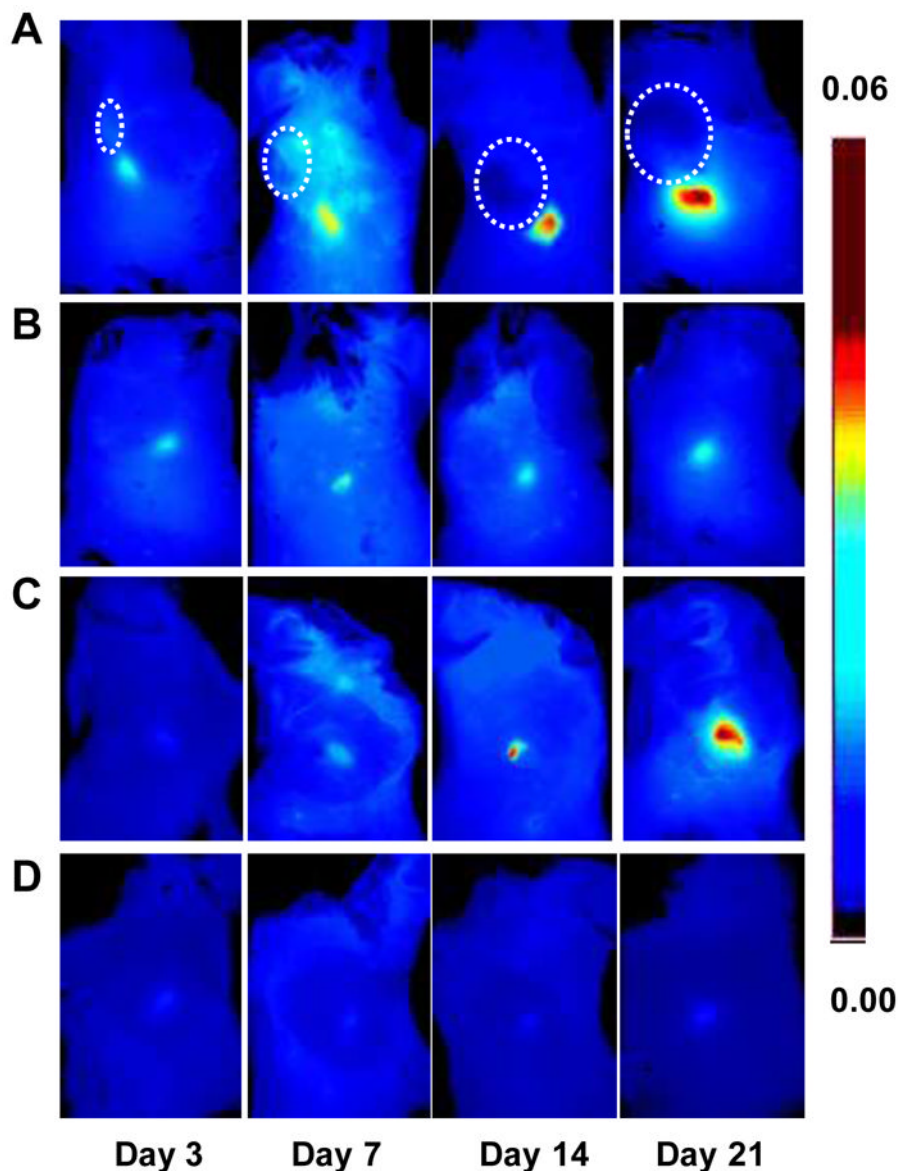
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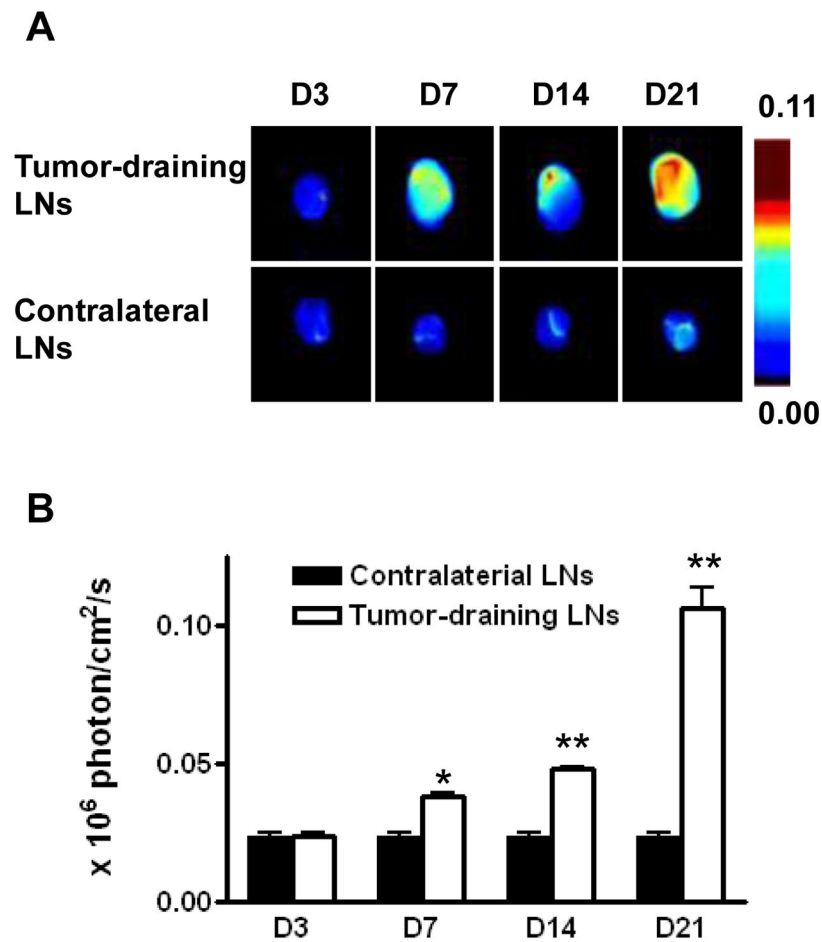
**Figure 1.** (A) Structure of Cy5.5-LyP-1. (B) Serum stability of Cy5.5-LyP-1 with LC-MS.



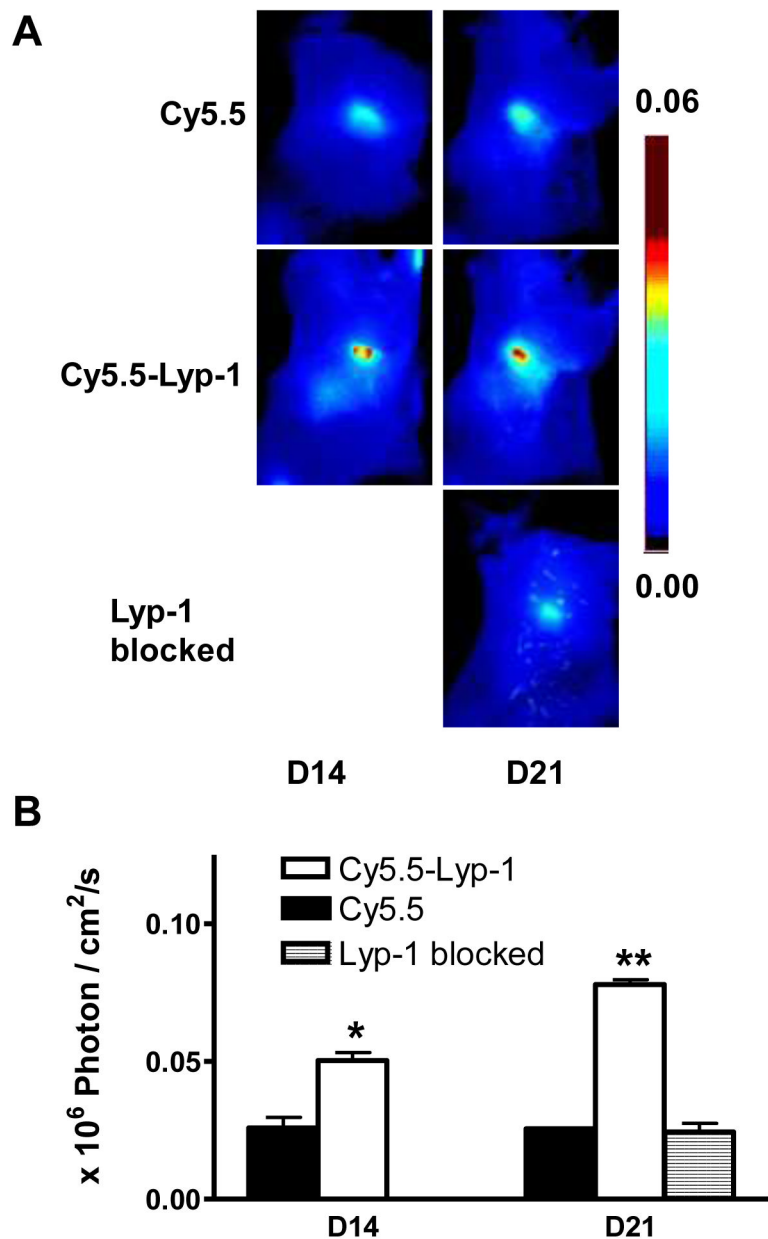
**Figure 2.**  
(A) Relative position of primary tumor (blue circle) and tumor-draining LN (white circle).  
(B) Photographs of three sets of tumors (left), tumor-draining lymph nodes (middle) and contralateral lymph nodes (right).



**Figure 3.** Optical imaging of tumor-induced lymphangiogenesis with Cy5.5-LyP-1. At days 3, 7, 14 and 21 after 4T1 cell inoculation, tumor-bearing BALB/C mice (n = 5/group), under isoflurane anesthesia, were injected with Cy5.5-LyP-1 via the middle phalanges of both upper extremities. At 45 min after injection, *in vivo* fluorescence imaging of both tumor-draining LNs (A) and contralateral LNs (B) was performed with a Maestro II small-animal *in vivo* imaging system. The fluorescence images consisting of Cy5.5 and autofluorescence spectra were then unmixed based on their spectral patterns with the manufacturer's software (Maestro software, CRI). The tumors are indicated by circles. (C and D) At 24h post injection, the mice were euthanized and a dorsal skin flap was elevated to expose the brachial lymph nodes on both sides for a repeat of fluorescence imaging (C. tumor-draining LNs; D. contralateral LNs).

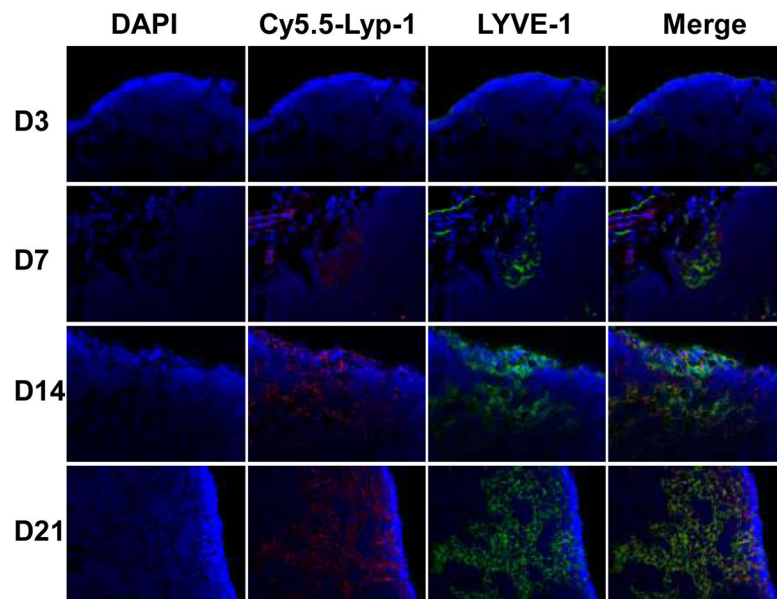


**Figure 4.** *Ex vivo* optical imaging of lymphangiogenesis. (A) Compared with the contralateral brachial LN, the tumor-draining brachial LN was significantly enlarged at day 7, 14 and 21 after tumor cell inoculation. (B) Measurement of Cy5.5-LyP-1 fluorescence intensity in LNs. Fluorescent signal increased dramatically from day 7 to day 21 in tumor-draining brachial LN (\* $P < 0.05$ , \*\* $P < 0.001$ ).



**Figure 5.**

(A) Optical imaging of tumor-induced lymphangiogenesis with Cy5.5-LyP-1 and Cy5.5. At days 14 and 21 after 4T1 cell inoculation, tumor-bearing BALB/C mice ( $n = 3/\text{group}$ ), under isoflurane anesthesia, were injected with either Cy5.5-LyP-1 or Cy5.5 via the middle phalanges of both upper extremities. At 24h post injection, the mice were euthanized and a dorsal skin flap was elevated to expose the tumor-draining brachial lymph nodes for fluorescence imaging. (B) Measurement of fluorescence intensity in LNs (\* $P < 0.05$ , \*\* $P < 0.01$ ).



**Figure 6.**

*Ex vivo* fluorescence staining of lymph node. At 24 h after Cy5.5-LyP-1 injection, brachial lymph nodes from both sides were removed and sectioned into 4  $\mu\text{m}$  slices and observed with an Olympus IX81 fluorescence microscopy. The same lymph node sections were also stained against LYVE-1, one of the most specific and widely used lymphatic endothelial markers. Nuclei were stained with DAPI.