Quantitative study on appearance of microvessels in spectral endoscopic imaging

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Abstract. Increase in abnormal microvessels in the superficial mucosa is often relevant to diagnostic findings of neoplasia in digestive endoscopy; hence, observation of superficial vasculature is crucial for cancer diagnosis. To enhance the appearance of such vessels, several spectral endoscopic imaging techniques have been developed, such as narrow-band imaging and blue laser imaging. Both techniques exploit narrow-band blue light for the enhancement. The emergence of such spectral imaging techniques has increased the importance of understanding the relation of the light wavelength to the appearance of superficial vasculature, and thus a new method is desired for quantitative analysis of vessel visibility in relation to the actual structure in the tissue. Here, we developed microvessel-simulating phantoms that allowed quantitative evaluation of the appearance of 15-μm-thick vessels. We investigated the relation between the vascular contrast and light wavelength by the phantom measurements and also verified it in experiments with swine, where the endoscopically observed vascular contrast was investigated together with its real vascular depth and diameter obtained by microscopic observation of fluorescence-labeled vessels. Our study indicates that changing the spectral property even in the wavelength range of blue light may allow selective enhancement of the vascular depth for clinical use.

Keywords: narrow-band imaging; blue laser imaging; vascular contrast; endoscopy; tissue-simulating phantom; three-dimensional microvascular imaging.

1 Introduction

In digestive endoscopy, it is well known that the emergence of microvessels in the superficial layer of the mucosa is an important aspect of cancerous disease. Clinical studies have shown that the increased density and irregular morphological features of microvessels are associated with tumor progression and that the observation of microvessels is crucial for cancer diagnosis. Thus, spectral endoscopy techniques such as narrow-band imaging (NBI) and blue laser imaging (BLI) which provide better visibility of microvessels, have been developed and are expected to play a significant role in gastrointestinal cancer diagnosis.

Short-wavelength light sources are utilized in spectral endoscopy techniques. There are two important optical properties that allow us to obtain high-contrast images of superficial microvessels. First, the absorption coefficient (μα) of blood is large at short wavelengths [Fig. 10]. For example, the absorption coefficient of blood μα at 420 nm is more than six times greater than that at 540 nm. This large absorption of light leads to high vascular contrast even for thin vessels including microvessels. Second, the scattering coefficient of mucosa is large at short wavelengths. Figure 10 shows an example of the reduced scattering coefficient μs′ for human gastric mucosa. The μs′ of biological tissue typically increases monotonically as the wavelength becomes shorter, and the gradient tends to be larger in the range of blue light due to the effect of Rayleigh scattering.

Thus, short-wavelength light does not penetrate into the deep layer of the mucosa by scattering and can detect only the superficial microvessels.

The NBI system, utilizing a xenon lamp and a set of optical filters, provides narrow-band blue illumination (centered at 415 nm and with a bandwidth of ±15 nm) for superficial vessel detection and green illumination (centered at 540 nm and with a bandwidth of ±15 nm) for visualizing deeper vessels. The spectral characteristics of NBI are determined by the optical filters and cannot be changed during observations. The BLI system employs two lasers (sharply peaked in the ranges of 410 ± 10 and 450 ± 10 nm) for its narrow-band illumination. For the BLI system, the spectral characteristics can be changed in principle by controlling the intensity ratio of those two lasers.

The emergence of spectral endoscopy systems with multiple light sources has increased the importance of understanding the relationship between the light wavelength and visualization of the microvessels in the mucosa. While the theoretical analysis for evaluating the spectral images has become a critical issue to understand spectral images, the theory and the quantitative analysis methods remain to be established.

Gono et al. in their pioneering work measured a mucosa underside of a human tongue using five narrow-band illuminations and demonstrated that the superficial capillary patterns
were markedly improved by exploiting light of around 415 nm. Du Le et al.\textsuperscript{16,17} in a recent work attempted quantitative comprehensión of vascular contrast and light wavelength by Monte Carlo simulations and phantom measurements and indicated some key factors to explain the behavior of vascular contrast by NBI.

To make a quantitative evaluation of the appearance of superficial microvessels like those that emerge in the early cancerous tissue in relation to the wavelengths, however, these preceding studies have two problems: (1) in the observation of real human tissue by Gono et al., the actual depths and diameters of the observed vessels remained unclear; and (2) in the phantom experiment by Du Le et al., they did not investigate superficial thin vessels with a depth less than 100 \(\mu\)m and diameter less than 20 \(\mu\)m that are especially targeted for cancer diagnosis.

Here, we developed experimental techniques for phantom measurements that allowed us to quantitatively evaluate the contrast of vessels with controllable depth and diameter including superficial microvessels as thin as 15 \(\mu\)m, that were comparable in size with actual vessels that emerge in early cancer. We also performed a swine experiment, where the contrast of each vessel observed by an endoscope at four different wavelengths was investigated with its actual vascular depth, which were obtained by fluorescence labeling of the vessels and observing them with a confocal microscope. The details of these experiments are described and discussed.

### 2 Simulation Study and Phantom Experiment

To validate the dependence of blood vessel contrast on the light wavelength, we performed a phantom experiment observing microvessel-simulating phantoms and Monte Carlo simulations. Since we focused on the physical behavior of vascular contrast of superficial thin vessels in the mucosa, in this work we used a simplified tissue model where a single vessel was embedded in a homogeneous medium as described in this section.

#### 2.1 Methods for Phantom Experiment

##### 2.1.1 Microvessel-simulating phantoms

We produced microvessel-simulating phantoms by employing the technique for making a capillary pattern based on photolithography by Arai et al.\textsuperscript{18} This technique enables us to make a capillary within a polydimethylsiloxane (PDMS) resin base. The structure of the phantom is presented in Fig. 2. The depth and diameter of the capillary are controllable in the range of 15 to 100 \(\mu\)m. We defined the phantom size as 30\(\times\)30 \(\times\)4 mm\(^3\)

![Fig. 1 Absorption coefficient \(\mu_a\) of blood (a) and reduced scattering coefficient \(\mu'_s\) of mucosal tissue (b).](image-url)

The absorption coefficient \(\mu_a\) was calculated using the molar absorbance coefficient of \textit{oxy}- and \textit{deoxy}-hemoglobin, as in Ref.\textsuperscript{12} by considering the hemoglobin oxygen saturation (\(\text{StO}_2\)). The mucosal \(\mu'_s\) is for human gastric mucosa taken from Ref.\textsuperscript{13}.
The image acquisition system is presented in Fig. 3. The diffused light from the 445-nm laser was irradiated on the phantom with an incident angle of 45 deg. (Here, since the 445-nm laser was exploited merely as a high-power monochromatic light source, the wavelength of 445 nm was not essential for this experiment). The reflected light image was captured with an EMCCD camera (iXon3 888, Andor) with an attached zoom lens (VH-Z100UR, Keyence). We obtained an image resolution of $0.65 \mu m/pixel$, which was sufficient to resolve even the thinnest capillary of $15-\mu m$ diameter. The image-taking procedure consisted of reinjection of the tartrazine solution into the capillary, refocusing the camera, and image acquisition. To assess the reproducibility and error of the measured vascular contrast, we repeated the procedure for more than 10 times for each phantom.

### 2.1.3 Vascular contrast

We defined the vascular contrast as $C = \log_{10}(I_b/I_v)$, where $I_v$ is the signal level of the vessel area and $I_b$ is that of the surrounding mucosal area. We obtained the contrast value for each acquired phantom image. Since we repeated the image acquisition procedure for plural times as explained above, we calculated the average and standard deviation for all the obtained images.

### 2.1.4 Determination of optical properties

The optical properties of the phantom resin base were determined by the integrating sphere technique (Fig. 4). The total transmittance $T_t$ and diffuse reflectance $R_d$ of the 2-mm-thick resin base were measured using an integrating sphere (RT-060SF, Labsphere) and a spectrometer (PMA-12, Hamamatsu Photonics). The reduced scattering coefficient $\mu_s'$ was determined by an inverse Monte Carlo technique [1, 2]. Here, we assumed a homogeneous, 4-mm-thick tissue model possessing the optical parameters of $\mu_a$ and $\mu_s'$ and calculated its $T_t$ and $R_d$ using the software package Monte Carlo for multilayered (MCML) [3]. We performed the calculation for various $\mu_a$ and $\mu_s'$ and determined the optical parameters that reproduced the measured values with minimal errors.

### 2.1.5 Verification of phantom structure

To verify that each phantom was made with its intended structure, we cut the phantom vertically to the vessel after the measurement and acquired an image of the cross section with a confocal microscope (OPTELICS H1200, Lasertec). We obtained the actual depth and diameter of the vessel by converting the size on the cross-sectional image into the real size based on the spatial resolution of the image.

### 2.1.6 Validity check of phantom size

As explained in Sec. [1], we prepared phantoms of the single size of $30 \times 30 \times 4 \text{ mm}^3$. To check the validity of this size, we prepared phantoms of different thicknesses and sizes and investigated the thickness/size dependence of the vascular contrast.

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Table 1  
Vessel specifications investigated in the simulation study and the analysis of the swine vessel images. Types 1, 2, and 3 correspond to the superficial microvessels, moderately deep and thick vessels, and deep and thick vessels, respectively.

<table>
<thead>
<tr>
<th></th>
<th>Type 1</th>
<th>Type 2</th>
<th>Type 3</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Simulation</strong></td>
<td>Depth $d$ ($\mu m$)</td>
<td>50</td>
<td>150</td>
</tr>
<tr>
<td></td>
<td>Diameter $\phi$ ($\mu m$)</td>
<td>15</td>
<td>50</td>
</tr>
<tr>
<td><strong>Swine data analysis</strong></td>
<td>Depth $d$ ($\mu m$)</td>
<td>10 to 100</td>
<td>137 to 215</td>
</tr>
<tr>
<td></td>
<td>Diameter $\phi$ ($\mu m$)</td>
<td>7 to 30</td>
<td>37 to 54</td>
</tr>
</tbody>
</table>

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Fig. 3  
Image acquisition system for the phantom experiment: (a) the overall view and (b) schematic.

Fig. 4  
Integrating sphere system to measure the total transmittance (a) and diffuse reflectance (b) spectra. A collimated parallel beam (5 mm in diameter) of a halogen lamp was injected at the phantom resin base attached at the integrating sphere, and the transmitted/reflected light was sent to the spectrometer through an optical fiber. When we measured the reference standard spectra, we detached the sample from the sphere.
As for the thickness, we prepared phantoms of Type 2 vessel at 473 nm, whose thickness was varied in the range of 3 to 8 mm. Here, note that the larger the scattering coefficient of the phantom is, the less the phantom reflectance becomes dependent on the thickness. Since the scattering coefficient is the smallest at 473 nm in the range of 405 to 473 nm, the thickness dependence may be observed most easily at this wavelength if it exists. As for the horizontal size, we prepared phantoms of Type 2 vessel at 405 and 473 nm, whose width across the capillary was varied in the range of 8 to 30 mm.

2.2 Methods for Simulation Study

To investigate the variation in blood vessel contrast depending on the wavelength of illumination, we performed Monte Carlo simulations as follows. We modified the simulation software MCML to enable the simulation of light propagation in a tissue that possesses an internal blood vessel. Figure 5 shows the tissue model for this simulation. The structure of the tissue model can be defined by two parameters: \( d \), the vessel’s depth from the surface, and its diameter \( \phi \). We modeled three types of vessels, as shown in Table 1. The optical parameters of the tissue can be defined by three parameters: the absorption coefficient \( \mu_a \), scattering coefficient \( \mu_s \), and scattering anisotropy parameter \( g \). For the inner part of the vessel, we used the \( \mu_a \) of hemoglobin for its absorption, where we defined the hemoglobin concentration as 150 g/L and the oxygen saturation as 70%.

![Fig. 5 Tissue model for the simulation study. A blood vessel with square cross section is embedded in the mucosal tissue. The vessel diameter \( \phi \) is defined as the side length of the vessel cross section. The total thickness of the mucosa was set to 4 mm to match with the actual condition.](image)

A typical value for gastrointestinal mucosa The scattering coefficient within the vessel was set to zero to accord with the optical properties of the phantom vessels (described in Sec. 2.1.1). For the outer part of the vessel, we set \( \mu_a \) to zero, assuming there was no significant absorption outside of blood vessels, and used reduced scattering coefficient \( \mu_s' = \mu_s (1 - g) \). The values of \( \mu_s \) and \( \mu_s' \) used in this simulation are presented in Fig. 5. Photons were homogeneously injected into the tissue on an area of 15×15 mm² with incident angles in the range of 0 deg to 70 deg, and the emitted photons from the tissue surface were counted. The simulation was carried out for wavelengths in the range of 400 to 500 nm in 5-nm steps. We calculated two billion events for one simulating condition (tissue structure, optical parameters, and light wavelength).

2.3 Results

Figure 6 shows the simulated vessel contrast for each type of vessel and wavelength tested. Here, we focus on the wavelengths of 405, 420, 445, and 473 nm at which spectral images were obtained in the swine experiment (described in Sec. 3.1.1). This result suggests the following properties regarding the image contrast of vessels: (1) the shallow and thin vessel (50-μm deep and 15-μm thick) presents the highest contrast at 420 nm, but it also presents a high density even at 405 nm, comparable with the case at 420 nm. The detectable optical density is low at 445 and 473 nm; (2) the vessel of moderate depth and diameter (150-μm deep and 50-μm thick) presents almost the same densities at three wavelengths: 405, 420, and 445 nm; and (3) the deep and thick vessel (350-μm deep and 100-μm thick) presents a higher density at 445 nm than at 405 nm.

Next, we verified the agreement between the simulations and phantom measurements. The measured actual vessel depth and diameter, and the scattering coefficients determined by the integrating sphere and inverse Monte Carlo method, were in agreement with their targeted values within 10% accuracy. Figure 7 shows the reflectance profiles and vessel contrast values obtained from the phantom images compared with the simulated values. All the measurements agreed well with the simulation results. Although there are small differences in the absolute values (within roughly 0.07 optical densities), the simulation results reproduced the wavelength dependence of vascular contrast for each vessel type very well. This accuracy is sufficient to allow discussion of the differences in vessel optical densities at multiple wavelengths as estimated by the simulations.

![Fig. 6 Simulated vascular contrast for each type of vessel. The four wavelengths 405, 420, 445, and 473 nm at which we observed in the swine experiment are indicated by solid lines.](image)

3 In Vivo Imaging Study with Swine

3.1 Methods

3.1.1 Ethics statement

Prior to the study, animal protocol was reviewed and approved by the FUJIFILM Animal Experiment Committee and also by the Animal Care and Use Committee of Nihon Bioresearch Inc.
3.1.2 Pig preparation

We used a female miniature pig (weight 45.1 kg and age 19 months) from Nippon Institute for Biological Science. It was anesthetized by inhalation of a mixture of nitrous oxide and oxygen containing 0.5% to 1.5% isoflurane. All the following procedures were carried out under anesthesia. We opened the abdomen to expose the entire stomach.

3.1.3 Endoscopy

We prepared a special endoscope system for experimental use, equipped with laser diodes of 405, 420, 445, and 473 nm for illumination. This system enabled us to obtain and save images at a speed of \(30\ \text{frames/s}\). The illumination was subsequently switched to each of the four lasers synchronously with the video frames. The stomach mucosa was observed in the following manner. We inserted the endoscope into the stomach through the mouth and observed the gastric mucosa at the boundary of the cardiac and nonglandular area in magnifying mode. When we recorded the images, we kept the endoscope tip approximately 2 mm above the mucosa using the 2-mm-deep hood attached at the tip (Fig. 9). The observed area was marked by making spots using electric cautery. The outer side of the area was also marked by spotting with black ink.

3.1.4 Fluorescent labeling of gastric vessels

To create a three-dimensional (3-D) image of the blood vessels in the observed area by endoscopy, we fluorescently labeled the gastric blood vessels using tomato lectin \(Lycopersicon esculentum\) agglutinin (LEA).\(^{22-25}\) LEA can selectively label vascular endothelial cells. To label the swine gastric vessels, we developed a method to administrate LEA to gastric vessels only. The abdomen was opened to expose the whole stomach, and then a cannula was inserted into the left gastric artery for agent injection. The left gastric vein was clamped, and the left and right gastroepiploic arteries, right gastric artery and vein, and short gastric artery and vein were ligated. A cannula for drainage was inserted into the left gastric vein. Next, 12 mg of FITC-labeled LEA (FL-1171, Vector Laboratories) was injected into the left gastric artery. Approximately 5 min later, the left gastric vein was released, and then 500 mL of PBS containing 4% paraformaldehyde was administered through the injection.
cannula for perfusion fixation of all gastric tissues and also to flush away the residual LEA in the blood. After fixation, the pig was euthanized by exsanguination of the abdominal aorta. The stomach was removed and cut open from the greater curvature, and then the tissue of the observed area was extracted. The excised tissue was immersed in a sucrose solution overnight, and then 100-μm-thick cryostat sections were prepared. We observed the sections with a confocal laser microscope (LSM 780, Zeiss) at ×10 magnification, using an argon laser (488 nm) for FITC excitation. We obtained cross-sectional images at steps of 5.97 μm.

3.1.5 Vessel image analysis

We converted the set of sectional microscopic images into a 3-D image. This 3-D image enabled us to obtain the diameter and depth of each vessel on the endoscopic image in the following manner. Figure 10 shows the 3-D fluorescence image of the extracted swine gastric tissue obtained by confocal microscopy. We divided the depth of this image into three levels (Level 1: 0 to 200 μm, Level 2: 200 to 400 μm, and Level 3: 400 to 600 μm) and integrated the data of each level into a two-dimensional maximum intensity projection (MIP) image. We obtained cross-sectional images at steps of 5.97 μm.

Fig. 10 Three-dimensional (3-D) vascular image of the extracted swine gastric tissue obtained by confocal microscopy: (a) the 3-D vascular image data; (b) the MIP images (projected on the xy-plane) of all the depth levels; and (c)–(e) the MIP images of the three divided levels.

3.1.6 Results of matching microscopic and endoscopic images

We converted the set of sectional microscopic images into a 3-D image. This 3-D image enabled us to obtain the diameter and depth of each vessel on the endoscopic image in the following manner. Figure 11 shows the 3-D fluorescence image of the extracted swine gastric tissue obtained by confocal microscopy. We divided the depth of this image into three levels (Level 1: 0 to 200 μm, Level 2: 200 to 400 μm, and Level 3: 400 to 600 μm) and integrated the data of each level into a two-dimensional maximum intensity projection (MIP) image. We converted the set of sectional microscopic images into a 3-D image. This 3-D image enabled us to obtain the diameter and depth of each vessel on the endoscopic image in the following manner. Figure 11 shows the 3-D fluorescence image of the extracted swine gastric tissue obtained by confocal microscopy. We divided the depth of this image into three levels (Level 1: 0 to 200 μm, Level 2: 200 to 400 μm, and Level 3: 400 to 600 μm) and integrated the data of each level into a two-dimensional maximum intensity projection (MIP) image. We converted the set of sectional microscopic images into a 3-D image. This 3-D image enabled us to obtain the diameter and depth of each vessel on the endoscopic image in the following manner. Figure 11 shows the 3-D fluorescence image of the extracted swine gastric tissue obtained by confocal microscopy. We divided the depth of this image into three levels (Level 1: 0 to 200 μm, Level 2: 200 to 400 μm, and Level 3: 400 to 600 μm) and integrated the data of each level into a two-dimensional maximum intensity projection (MIP) image. We converted the set of sectional microscopic images into a 3-D image. This 3-D image enabled us to obtain the diameter and depth of each vessel on the endoscopic image in the following manner. Figure 11 shows the 3-D fluorescence image of the extracted swine gastric tissue obtained by confocal microscopy. We divided the depth of this image into three levels (Level 1: 0 to 200 μm, Level 2: 200 to 400 μm, and Level 3: 400 to 600 μm) and integrated the data of each level into a two-dimensional maximum intensity projection (MIP) image.
compared these MIP images and the endoscopic images [Fig. 11(a)], focusing on the vascular pattern, and identified the corresponding vessels to match these two types of images [Fig. 11(b)]. This matching enabled us to obtain the depth and diameter of each vessel that appeared on both images. We then selected three types of vessels whose depths and diameters are in the ranges as presented in Table 1. We calculated the vascular contrast values for these vessels.

3.2 Results

We classified the vessels that appeared in the endoscopic images by their depths and diameters into the three types presented in Table 1 and calculated the mean vascular contrasts for each type. The vascular contrasts were obtained for images at four wavelengths (405, 420, 445, and 473 nm). The results are presented in Fig. 12. The contrast of superficial microvessels is the highest at 420 nm and decreases in the order: 405, 445, and 473 nm. This relation between vascular contrast and wavelength is consistent with the simulation results of Type 1 vessels. The wavelength-dependent differences in the vascular appearance can also be identified in the endoscopic images (Fig. 13), where the vascular appearance drastically changes as the wavelength changes between 420 and 445 nm. This change can be attributed to the behavior of vascular contrast at 445 nm. Contrast for superficial microvessels at 445 nm is low compared with that at 420 nm, but becomes higher and comparable with 420 nm for deeper and thicker vessels (Fig. 12). This property suggests that changing the spectral information, even in the wavelength range of blue light, will permit one to select the type of vessels to enhance the image for clinical use. In particular, combining two spectral images, such as 405 or 420 and 445 nm, and exploiting the difference in the superficial microvascular contrast of these images may enable enhancement of highly superficial microvessels with a depth less than, e.g., 50 μm. This could be useful for some clinical applications. For example, Konda et al. demonstrated that the superficial vascular density in Barrett’s esophagus increased as the lesion progressed from benign to invasive cancer and that the most superficial (0 to 50 μm) vascular density presented a significantly larger increase than those at depths of 0 to 150 μm.

The results presented above are based on monochromatic light at 405, 420, and 445 nm. However, NBI uses blue light at 415 nm with the bandwidth of ±15 nm. To investigate the effect of this band spread, we recalculated the contrast–depth relation of microvessels (ϕ = 15 μm and d = 50 μm) for each wavelength by convolution of the simulated reflectance spectrum and the Gaussian functions with σ = 10 and 20 nm [Fig. 14(a)]. We also investigated in the case of a triangular filter function as shown in Fig. 14(b) to simulate the diminishing power as the wavelength approaches 400 nm due to the common

4 Discussion and Conclusion

A simulation study and phantom measurements for common tissue models were performed. The vascular contrasts were matched with high precision for two types of vessel with different optical parameters (Fig. 11). These results validate our simulation method using the current conditions, thus we can discuss the vascular contrast based on the simulation results.

As for superficial microvessels, the vascular contrast was the highest at a wavelength of 420 nm. This property agrees with the findings of Du Le et al. and can be ascribed to the optical properties of blood absorption, which peaks around 420 nm (Fig. 1). In addition to 420 nm, we also investigated vascular contrasts at 405 and 445 nm, wavelengths that were not studied in that publication. It should be noted that a small difference in wavelength around 420 nm leads to prominent differences in vascular contrast–depth relations (Fig. 11). These wavelength-dependent differences in the vascular appearance can also be identified in the endoscopic images (Fig. 13), where the vascular appearance drastically changes as the wavelength changes between 420 and 445 nm. This change can be attributed to the behavior of vascular contrast at 445 nm. Contrast for superficial microvessels at 445 nm is low compared with that at 420 nm, but becomes higher and comparable with 420 nm for deeper and thicker vessels (Fig. 12). This property suggests that changing the spectral information, even in the wavelength range of blue light, will permit one to select the type of vessels to enhance the image for clinical use. In particular, combining two spectral images, such as 405 or 420 and 445 nm, and exploiting the difference in the superficial microvascular contrast of these images may enable enhancement of highly superficial microvessels with a depth less than, e.g., 50 μm. This could be useful for some clinical applications. For example, Konda et al. demonstrated that the superficial vascular density in Barrett’s esophagus increased as the lesion progressed from benign to invasive cancer and that the most superficial (0 to 50 μm) vascular density presented a significantly larger increase than those at depths of 0 to 150 μm.

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spectral property of a xenon lamp used for endoscopy (Fig. 14). The results are presented in Fig. 15 with limiting the wavelength range to less than 450 nm.

![Fig. 14](image1.png)

**Fig. 14** Optical band-pass filter functions used for the simulation of band broadening: (a) Gaussian filters with $\sigma = 10$ and 20 nm. We calculated at $\lambda_0 = 405, 420, \text{ and } 445 \text{ nm}$; and (b) triangular filter simulating the typical xenon lamp spectrum (Fig. 15) with limiting the wavelength range to less than 450 nm.

![Fig. 15](image2.png)

**Fig. 15** Typical spectrum of a xenon lamp as a white light source used for endoscopy (in-house data by FUJIFILM). The light intensity rapidly decreases as the wavelength approaches 400 nm.

![Fig. 16](image3.png)

**Fig. 16** Simulated effect of band broadening on the vascular contrast ($\phi = 15 \mu m \text{ and } d = 50 \mu m$).

The results in Fig. 15 indicate that it is important to consider not only the peak wavelength, but also the bandwidth and its shape. Additionally, our study indicated that changing the spectral property even in the wavelength range of blue light may allow selective enhancement of the vascular depth for clinical use. To obtain high contrast of superficial microvessels, exploiting the large contribution of light in the range of 400 to 420 nm is essential and using longer wavelengths will enable enhancement of deeper and thicker vessels. Vessels in the tumorous tissue vary in their depths depending on the tumor type and its stage of progression. Thus, it is expected that the spectral imaging techniques for vascular enhancement will develop to the next stage where the spectral property of the illumination can be optimally changed in the target-lesion oriented manner.

**References**


Biographies of the authors are not available.