Endoscopic Fluorescence-Guided Surgery for Sinonasal Cancer Using an Antibody-Dye Conjugate

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Objective: Endoscopic resection of sinonasal squamous cell carcinoma has become the standard of care, but challenges remain in obtaining clear resection margins. The current study evaluated the feasibility of endoscopic fluorescence-guided surgery (FGS) to improve surgical resection in a human sinus surgical model.

Methods: A fluorescence endoscope optimized for near-infrared (NIR) fluorescence detection was evaluated in a phantom study. Various endoscope diameters (4 and 10 mm) and viewing angles (0, 30, and 45 degrees) were evaluated to determine the feasibility of the system for IRDye800CW detection at various working distances (1–5 cm). Endoscopic FGS was then validated in a three-dimensional human sinus surgical model to which squamous cell tumors derived from mice were inserted. Mice had received intravenous panitumumab-IRDye800CW and upon fluorescence-guided tumor resection, mean fluorescence intensity (MFI) and tumor-to-background ratio (TBR) were calculated in situ and ex vivo settings.

Results: A significantly higher fluorescence intensity was found when using the 10-mm diameter endoscope compared to the 4mm diameter endoscope (P < .001). No significant difference in MFI was found among the viewing angles of the 4-mm diameter endoscope. Using the human sinus model, the highest MFI and TBR were obtained at a 1-cm working distance compared to longer working distances.

Conclusion: We demonstrate that clinically acceptable TBRs were obtained with several working distances to discriminate tumor tissue from adjacent normal tissue in a human sinus model, and that endoscopic FGS may have great potential in identifying residual tumor tissue regions during surgery.

Key Words: Fluorescence-guided surgery, endoscope, near-infrared, sinonasal cancer, squamous cell carcinoma.

INTRODUCTION

SINonasal squamous cell carcinoma (SNSCC) is the most common histology in malignant sinonasal tumors.1,2 Surgical resection is considered the standard of care treatment for SNSCC, and complete resection with tumor-free margins provides optimal survival.3 Recent studies have demonstrated the usefulness and feasibility of an endoscopic approach for malignant sinonasal tumors with acceptable survival outcomes and low rate of surgical complications.4–6 However, while endoscopy allows for minimally invasive surgical and direct visualization of the surgical field, clear resection margins can be difficult to ascertain due to a narrow field of view and the inability to palpate tumor edges. Consequently, rates of positive margins for endoscopic surgery for SNSCC are high (22–30%), which can result in increased local recurrence and decreased survival.7,8

Fluorescence-guided surgery (FGS) is increasingly being studied as a guide to differentiate tumor tissue from normal tissue in an intraoperative setting.9 Although several agents are under active clinical study,10–13 we investigated the anti-epidermal growth factor receptor (EGFR) antibody-dye conjugate panitumumab-IRDye800CW, which has demonstrated a high sensitivity and specificity of squamous cell carcinoma (SCC) detection in patients with head and neck cancer.14–18 We hypothesize that translating this
technique to endoscopic surgery may improve discrimination of tumor tissue from adjacent normal tissue. This would ultimately allow for more precise visualization of the tumor margins and reduce incomplete tumor resection. Thus, the objective of this study was to evaluate the potential clinical utility of the endoscopic sinonasal FGS technique in a novel human sinus surgical model and to determine the optimal surgical conditions for fluorescence detection.

MATERIALS AND METHODS

Endoscopic Fluorescence Imaging System

The endoscopic near-infrared (NIR) fluorescence imaging system included a D-Light P Xenon light source with ICG filters and an Image1 S H3-Z FI camera. Other components on the device were the Image1 S connect platform, the Image1 S H3-link, a Storz Aida HD Connect, and a fiber optic light cable connecting the light source and rigid Hopkins telescopes (KARL STORZ SE & Co KG, Tuttlingen, Germany). Using a foot-switch, the system allows a switchover between fluorescence and standard white light mode.

Shown in Figure 1 are the endoscopes that were evaluated. Three 4-mm diameter, 18-cm long endoscopes (0, 30, and 45 degrees) and two 10-mm diameter, 31-cm long, endoscopes (0 and 30 degrees) were evaluated. All of the endoscopes had built-in filters for NIR light detection, with all detected fluorescence signal being displayed on-screen as a blue signal overlaid onto a bright field image. Images taken with the endoscopic imaging device were compared with a closed-field fluorescence imaging-device (Pearl Trilogy Imaging System; LI-COR Biosciences Inc., Lincoln, NE, USA). The closed-field device shields itself from ambient light, allowing for consistent fluorescence imaging and served as a “gold-standard” fluorescence imaging device in this study, in a manner similar to as previously reported.18

Sensitivity to IRDye800CW

To determine the sensitivity of the device to NIR fluorescence detection of panitumumab-IRDye800CW, a 20-step 1:2 dilution range, ranging 4.58 mM to 3.94 pM, was generated of IRDye800CW carboxylate (LI-COR Biosciences Inc.) in water. For reference, the dilution range was first measured in the closed-field imaging device. Subsequently, the three rigid 4-mm diameter endoscopes (0, 30, and 45 degrees; Fig. 1) and two 10-mm diameter endoscopes (0 and 30 degrees) were used to measure fluorescence signals for all concentrations of dye to determine, quantitatively and qualitatively, the sensitivity of the device, in a manner similar to as previously reported.19 To determine the optimal working distance of the endoscope (ie, the working distance at which the highest sensitivity was reached) the experiment was performed at working distances ranging 1–5 cm.

Endoscopic Sinus Surgery Model

Nude (nu/nu) female mice, aged 6–8 weeks (Charles River Laboratories, Wilmington, MA, USA), were obtained and housed in accordance with our Institutional Animal Care and Use Committee (IACUC) guidelines. All experiments were conducted after review and approval of the institution’s IACUC guidelines. To generate the orthotopic head and neck squamous cell carcinoma (HNSCC) model, mice (n = 4) were injected with SCC-1 cells (0.5 × 10⁶; in 25 µL PBS; ATCC, Manassas, VA) in the floor-of...
mouth using a 30-G insulin syringe. Tumor growth was followed for 2–3 weeks by direct visual inspection and palpation, after which the mice were systemically injected with 100 μg of GMP-produced panitumumab-IRDye800CW.14 Mice were sacrificed 48 hours post-injection after which the skin overlying the floor of the mouth was removed and imaging in the closed-field device was performed to verify presence of panitumumab-IRDye800CW.

In order to simulate the anatomic and light conditions of SNSCC resection, sacrificed mice were placed in an anatomically accurate three-dimensional sinus model (sinus model otorhino neuro trainer; Pro Delphus, Pernambuco, Brazil).20 An overview of the experimental setup is given in Figure 2. Tumors were then resected under guidance of the endoscopic imaging using the 0 degree, 4-mm diameter, endoscope whereby various working distances (1, 2, and 3 cm) were assessed. Upon completion of resection, a second closed-field fluorescence image was acquired to verify the complete removal of the tumor tissue.

To determine the smallest detectable size of tumor tissue, excised tumor tissue and normal muscle were divided into parts, varying in weight from 0.2 to 20 mg (Fig. 5). Each piece was imaged using the fluorescence endoscopic imaging device, and for verification, each piece was also imaged in the closed-field device.

**Histopathology**

During resection of the tumor tissue from the mice, representative primary tumor tissue, tumor-to-normal margin tissue and wound-bed tissue was collected, formalin-fixed and paraffin embedded, after which blank slides (4-μm thickness) were obtained for a hematoxylin and eosin (H&E) stain. Subsequent immunohistochemistry was also conducted to assess for epidermal growth factor (EGFR) expression using an anti-EGFR antibody (clone EP38Y, Thermo Fisher Scientific, Waltham, MA, USA).14

**Fluorescence Microscopy**

For fluorescence microscopy, selected tissue slides were deparaffinized, and the nuclei were counterstained with 4',6-diamidino-2-phenylindole (DAPI, Prolong Diamond, Thermo Fisher Scientific). Stained slides were dried overnight, in the dark, and imaged using a custom setup inverted digital fluorescence microscope (DM6B, Leica Biosystems, Wetzlar, Germany) equipped with a highly sensitive Leica DFC9000GT camera (4.2-M pixel sCMOS camera), a metal halide LED light source (X-Cite 200DC, Excelitas Technologies, Waltham, MA, USA) for DAPI imaging, and a xenon arc lamp LB-LS/30 (Sutter Instrument, Novato, CA, USA) for NIR imaging of IRDye800CW. Image acquisition and processing was done using LAS X software (Leica Biosystems).

**Image Quantification and Data Analysis**

Endoscopic images that were acquired before, during, and after tumor resection were loaded into ImageJ (National Institutes of Health, Bethesda, MD, USA) to quantitatively assess the fluorescence signal in situ and ex vivo. Because the endoscopic imaging system presents the fluorescence signal as an overlay image whereby the fluorescence signal (in blue) is overlaid on the bright field image, acquired images were split into RGB color channels in ImageJ. Subsequently, the blue channel of each image was analyzed in ImageJ as being the “raw” fluorescence signal. Reference (“gold standard”) images acquired with the closed-field fluorescence-imaging device were analyzed using Image Studios software (LI-COR Biosciences Inc.). For fluorescence signal quantification in both ImageJ and Image Studio, regions of interest (ROIs) were drawn around the area of interest and mean fluorescence intensities (MFIs, a.u.) were calculated. Tumor-to-background ratios (TBRs) were calculated by dividing tumor MFI by the normal tissue MFI.

**Statistical Analysis**

Descriptive statistics was performed using GraphPad Prism software (Version 6.0c; GraphPad Software, La Jolla, CA, USA). Data is presented as means with standard deviations for continuous variables. To compare the endoscope variants (diameter and
Fig. 3. Fluorescence signals and tumor-to-background ratio in situ and ex vivo setting measured at three working distances (1–3 cm). Images taken by the endoscopic fluorescence imaging system of in situ (A) and ex vivo (B) normal muscle and tumor tissue at three working distances (1–3 cm). [Color figure can be viewed in the online issue, which is available at www.laryngoscope.com.]

Fig. 4. Fluorescence-guided resection of tumor. Using the 4mm diameter endoscope (0 degrees), fluorescence and bright field images were acquired prior to resection, after initial resection, and after re-resection of residual tumor. Histopathology confirmed presence of EGFR in the excised tumor tissue, but not in the normal tissue. Via fluorescence microscopy presence of panitumumab-IRDye800CW was confirmed in the tumor tissue. [Color figure can be viewed in the online issue, which is available at www.laryngoscope.com.]
degree) in the phantom study, Wilcoxon matched-pairs signed rank test or Friedman test were used. Mann–Whitney U test was used for comparison of the tumor MFIs and normal tissue MFIs. Exponential regression analysis was performed on data comparing the two imaging systems. A P-value of .05 or less was considered statistically significant.

**RESULTS**

**Ex-vivo Fluorescence Endoscopy Validation**

Three 4-mm diameter endoscopes of varying angles of view (0, 30, and 45 degrees) were used to image the dilution range of 20 concentrations of IRDye800CW at five working distances ranging 1–5 cm to evaluate the sensitivity of the endoscopes. For the 4-mm diameter endoscope (0 degree), the highest and most reliable fluorescence signals were obtained at the 2-cm working distance (Fig. 1A); therefore, we determined 2 cm as the optimal working distance for our phantom study. In terms of the degree of 4mm diameter endoscopes, no differences were found in MFI when changing the angle of the endoscope (0, 30, and 45 degree; P = .17, Fig. 1B). Visually, at a working distance of 2 cm, the lowest concentration of dye that produced a visible fluorescence signal in the raw images was 698 nM (Fig. 1C). Fluorescence imaging data obtained with the endoscopic device strongly correlated to that of the closed-field imaging device (R² = 0.99, Fig. 1C), which indicated the robustness of the endoscopic fluorescence imaging system. When comparing between 4-mm and 10-mm diameter endoscopes, a significantly higher MFI was found when using the 10-mm diameter endoscope (P < .001, Supplementary Fig. 1).

**Fluorescence Endoscopy Identifies Residual Tumor Deposits**

Using the human sinus model, the potential value of endoscopic FGS for tumor visualization and margin assessment for sinonasal surgery was assessed. Following tumor visualization (Fig. 4), an incomplete tumor resection was performed to evaluate whether or not the imaging device was sensitive enough to pick up residual tumor. As can be seen in Figure 4, residual tumor deposits could be clearly visualized and subsequently removed. In further assessment of excised tumor pieces, the smallest piece of tumor tissue (0.6 mg) had a visible fluorescence signal and could be discriminated from normal tissue (Fig. 5A). A significantly higher MFI was found for tumor tissue when compared to normal tissue (mean: 40.0 vs. 15.6 a.u., P < .05, Fig. 5b), demonstrating this technique could clearly discriminate the tumor tissue from normal tissue even though the tumor deposits were very small. Imaging results with the endoscopic device were corroborated by the closed-field imaging device. Histology, including immunohistochemistry of EGFR, confirmed the presence of tumor at areas that were positive for fluorescence (Fig. 4).

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Fig. 5. Panitumumab-IRDye800CW uptake in tumor and normal tissue. A) Tumor and normal muscle were serially bisected into pieces, with the smallest piece being 0.6 mg, and the fluorescence signal was imaged using the 4-mm diameter endoscope (0 degrees). The mean fluorescence signal (MFI) was plotted as a function of weight (mg) of both tumor (red) and normal (muscle; blue) tissue samples. B) Upon normalizing of the fluorescence signal, by dividing the measured MFI by the tissue weight, panitumumab-IRDye800CW distribution in tumor tissue was found to be significantly higher in tumor tissue compared to normal (muscle) tissue. [Color figure can be viewed in the online issue, which is available at www.laryngoscope.com.]
Discission

For sinonasal cancers, complete resection with clear margins significantly impacts oncologic outcomes in patients treated with minimally invasive endoscopic resection.22,23 Despite the fact that intraoperative frozen margins are reliably interpreted for sinonasal tumors,24 a clear demarcation of tumor from normal tissue or inflamed polyoid tissues is difficult within the sinonasal cavity. In the current study we demonstrate that endoscopic fluorescence imaging can be sensitive and specific to detect the tumor after systemic administration of a NIR tumor-targeting agent. Moreover, clinically relevant TBRs were obtained with several working distances to discriminate the tumor from adjacent normal tissue in a human sinus model, highlighting the potential of this technique to identify residual tumor regions during surgery. Results of the current study are in line with previous studies in open-field FGS approaches for identification of tumor in an in vivo and an ex vivo setting.15,17

The current study demonstrates that endoscopic imaging systems with NIR fluorescence-detection designed for ICG can be repurposed for use with IRDye800CW. Despite the consistency of the TBR at greater working distances in our model, the fluorescent signal (measured in MFI) was negatively correlated with working distance. The optimal working distance is thought to lie within 1–3 cm in a clinical setting; fluorescence imaging becomes brighter as the working distance is reduced and vice versa. However, there is an increase in light reflection off of tissue at shorter working distances, which may ultimately obscure tumor visualization in a surgical setting.

Endoscopes used for sinus surgery or endonasal skull base surgery generally have an outer diameter of 4 mm, whereas endoscopes with 10-mm diameter are common in the laparoscopic surgeries. Our results demonstrate the applicability of the FGS technique beyond open-field and laparoscopic applications (eg, in the abdominal cavity). Although the fluorescence signal obtained from 4-mm diameter endoscopes was less than the signal from 10-mm diameter endoscopes in our phantom experiments, a fluorescent signal was less than the signal from 10-mm diameter endoscopes in our phantom experiments, a fluorescent signal brighter as the working distance is reduced and vice versa. However, there is an increase in light reflection off of tissue at shorter working distances, which may ultimately obscure tumor visualization in a surgical setting.

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Clinically, endoscopic FGS has emerged as a promising technique to improve surgical precision during tumor resection. Utilizing NIR fluorescence imaging using ICG with laparoscopic and robot-assisted surgery have improved oncologic outcomes reported in procedures such as hepatectomy,25 nephrectomy,24 hysterectomy,25 and gastrectomy.25 We and others have reported the feasibility and potential clinical benefits of FGS using antibody dye–based imaging for patients with solid tumors.10,11,13–18,27–31 Combining FGS with the advantages of a minimally invasive, endoscopic approach may hold the potential to reduce morbidity while ensuring adequate oncologic outcomes. However, this has yet to be further investigated and validated in large prospective trials.

Conclusion

We demonstrate that endoscopic imaging of a tumor-specific NIR imaging agent can discern tumor tissue from normal tissue in a preclinical model under a range of conditions with a clinically relevant TBR at working distances from 1–3 cm.

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