5-aminolevulinic acid for quantitative seek-and-treat of high-grade dysplasia in Barrett’s esophagus cellular models

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Abstract. High-grade dysplasia (HGD) in Barrett’s esophagus (BE) poses increased risk for developing esophageal adenocarcinoma. To date, early detection and treatment of HGD regions are still challenging due to the sampling error from tissue biopsy and relocation error during the treatment after histopathological analysis. In this study, CP-A (metaplasia) and CP-B (HGD) cell lines were used to investigate the “seek-and-treat” potential using 5-aminolevulinic acid-induced protoporphyrin IX (PpIX). The photodynamic therapy photosensitizer then provides both a phototoxic effect and additional image contrast for automatic detection and real-time laser treatment. Complementary to our studies on automatic classification, this work focused on characterizing subcellular irradiation and the potential phototoxicity on both metaplasia and HGD. The treatment results showed that the HGD cells are less viable than metaplastic cells due to more PpIX production at earlier times. Also, due to mitochondrial localization of PpIX, a better killing effect was achieved by involving mitochondria or whole cells compared with just nucleus irradiation in the detected region. With the additional toxicity given by PpIX and potential morphological/textural differences for pattern recognition, this cellular platform serves as a platform to further investigate real-time “seek-and-treat” strategies in three-dimensional models for improving early detection and treatment of BE.

Keywords: Barrett’s esophagus; photodynamic therapy; 5-aminolevulinic acid; PpIX; subcellular treatment.

1 Introduction

Barrett’s esophagus (BE) is a precancerous condition occurring in the lower esophagus, where the original squamous epithelium undergoes metaplastic transformation and is replaced by the intestinalized columnar cells. BE can progress through low-grade dysplasia to high-grade dysplasia (HGD), which significantly increases the risk of developing esophageal adenocarcinoma if early treatment is not provided. Although various interventions have been studied extensively, the various diagnostic techniques are still inadequate and suspected areas of HGD may be missed because of sampling errors for the initial biopsies or because of an inability to locate the same target areas on repeat endoscopy. This is a common problem because pathological analysis following four-quadrant biopsy is the current gold standard for evaluating suspected lesions. Because it can take several days to process, stain, and examine the biopsies, diagnosis and treatment are commonly separated into two steps, which leads to difficulties in accurate relocation of the lesions in separate procedures.

Optical biopsy is an emerging technique that takes advantages of the distinct tissue optical properties of targeted lesions to enable real-time diagnosis. In particular, the cellular-based optical biopsy retrieves subcellular Barrett’s-associated characteristics by use of point spectroscopy or endomicroscopy. However, there are substantial challenges in detecting dysplasia in BE with optical biopsy including poor signal-to-noise ratio, low specificity in fluorescence endoscopy imaging and significant patient-to-patient variability. To address these challenges, microscopic-based visualization of subcellular morphology with the addition of an exogenous contrast agent is desirable to enhance the contrast between normal, metaplastic, and HGD regions. Fluorescence of photosensitizers used in photodynamic therapy (PDT) has been reported to be beneficial in distinguishing drug–cellular interactions, giving PDT photosensitizers additional merits besides being therapeutic reagents. Using fluorescence from FDA-approved photosensitizers as a contrast agent becomes an attractive option for clinical translation of the techniques. In addition, the use of photosensitizers may provide additional treatment efficacy. In other words, integrated detection and treatment can be achieved using well-characterized light illumination immediately following a robust detection with quantitative feature extraction and classification. Palanca-Wessels et al. have developed metaplasia (CP-A) and HGD (CP-B, CP-C, and CP-D) cell lines from BE. The morphological features and cellular behaviors of these cell lines have been investigated along with normal squamous epithelial cells in both coculture and organotypic culture models. Therefore, these cell lines could potentially serve as a cellular-based platform to study the integrated early detection and treatment of BE.
In this study, we have focused on characterizing the subcellular irradiation efficacies in order to understand the potential protoporphyrin IX (PpIX) phototoxicity on both metaplasia and HGD cell lines after detection. The microscopic images obtained at subcellular resolution would be comparable with microendoscopic techniques in development. Among potential PDT “contrast agents,” 5-aminolevulinic acid (5-ALA) is an endogenous heme precursor that localizes well in mucosal tissues. It produces preferential accumulation of cytotoxic PpIX at the mitochondrial membranes, which can potentially lead to treatment selectivity; more importantly, PpIX provides the nucleus contrast for detection of HGD. To date, although 5-ALA has been studied extensively for treating BE in vivo, limited selectivity was noted for dysplastic cells. Our goal is to use the BE cellular platform to investigate the feasibility of integrated detection and treatment. The quantitative imaging analysis, feature extraction, and classification performance have been investigated in our group, followed by the characterization of localized, subcellular treatment efficacy in this study to examine the optimal irradiation scheme.

2 Materials and Methods

We characterized the dose response curves of two BE cell lines (CP-A and CP-B) to various doses of light fluence using one-photon and two-photon irradiations. Three irradiation regimes that target the whole cell, mitochondria, and cell nuclei were applied at each light dose in order to investigate the treatment efficacy as a result of targeted subcellular PDT. Experiments were conducted in the presence or absence of 5-ALA for positive and negative controls.

2.1 Cell Culture

CP-A (nondysplastic metaplasia, ATCC® CRL-4027™, American Type Culture Collection, Manassas, Virginia) and CP-B (HGD, ATCC® CRL-4028™) are both hTERT-immortalized epithelial cells derived from human esophagus. These well-characterized premalignant cell lines were selected as an in vitro platform to study subcellular treatment efficacy. Both cell lines were cultured under identical conditions using the same batch of MCDB 153 base medium (M7403, Sigma, St. Louis, Missouri), supplemented with 5% fetal bovine serum (16000-036, Gibco®, Life Technologies, Burlington, Ontario, Canada), 0.25 μg/ml Amphotericin B (A2942, Sigma), 1% penicillin/streptomycin (15140-122, Gibco®), 20 mg/ml bovine pituitary extract (P1476, Sigma), 100 mM 5-ALA dissolved in PBS (14190-144, Gibco®), 20 μg/ml Amphotericin B (A2942, Sigma), 1% penicillin/streptomycin (15140-122, Gibco®), 0.4 μg/ml hydrocortisone (H8888, Sigma), 140 μg/ml bovine pituitary extract (P1476, Sigma), 20 mg/L Adenine (A8626, Sigma), 0.1% insulin-transferrin-sodium selenite (I1884, Sigma), 20 ng/ml Recombinant EGF (E9644, Sigma), 4 mM glutamine (G7513, Sigma), and 1 mM cholora toxin (C8052, Sigma). Cells were maintained at 37°C in a water-jacketed CO2 incubator (Forma Series II, Thermo Fisher Scientific Inc., Waltham, Massachusetts) and passaged as recommended by ATCC. All experimental data were acquired using cells within 10 passages after cryopreservation to minimize the variability due to cell degradation.

2.2 Experimental Preparation

In this study, 5-ALA (A3785, Sigma) served as a fluorescent marker as well as a PDT photosensitizer. A stock solution of 100 mM 5-ALA dissolved in PBS (14190-144, Gibco®) was prepared and stored in 1 mL aliquots in the dark. Fresh photosensitizer-containing medium was prepared by further diluting the stock solution in the serum-free complete culture medium to a working concentration of 0.5 mM that provides sufficient contrast without yielding dark toxicity.

Cells were detached from a 25 cm² culture flask with 2.5× Trypsin-EDTA solutions (0.5%, 15400-054, Gibco®). A centrifugation process at 125g was used to resuspend cells in fresh trypsin-free culture medium. A total of 1 × 10⁵ cells were seeded on a gridded glass-bottom dish (P35G-2-14-CGRD, MatTek, Ashland, Massachusetts) 24 h before the experiment. The culture medium was then replaced by the 0.5-mM 5-ALA-containing medium for a specified incubation period before the experiment. Samples were rinsed with Hanks balanced salt solution (14025-092, Gibco®) supplemented with 20 mL of Hepes buffer (15630-106, Gibco®) before imaging and were immersed in 1 mL of Hanks-Hepes buffer during the imaging acquisition procedures.

2.3 Imaging Acquisition

The fluorescence emission of 5-ALA-induced PpIX was collected by a laser scanning confocal microscope (TCS SP5 and DMI 6000 B, Leica, Wetzlar, Germany) equipped with an argon-ion laser operating at 514 nm for one-photon irradiation and a femtosecond Ti:sapphire laser (Chameleon-Ultra, Coherent, Santa Clara, California) pulsing at 810 nm for two-photon excitation. The emission spectral band was set as 600 to 750 nm to maximize the detected PpIX intensity, in combination with a transmitted light detector to acquire bright field images.

2.4 Subcellular Treatment

Both cell lines were incubated with 5-ALA for 3 h before imaging and subcellular treatment. The light fluence levels used for imaging and dose response curves are summarized in Table 1, where the fluence was calculated based on the average power (P) measured at the sample plane, total exposure time (t), and the field of view (FOV); 246 × 246 μm². In order to understand the cytotoxic roles played by light fluence and PpIX and eventually to select an appropriate ablative approach, the dose response curves were compared between three treatment schemes selected manually based on the bright field and 5-ALA images using the built-in software of the microscope:

<table>
<thead>
<tr>
<th>One-photon excitation</th>
<th>Fluence (J/cm²)</th>
<th>Two-photon excitation</th>
<th>Fluence (J/cm²)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.15 × 6.4</td>
<td>0.002*</td>
<td>45 × 1.28</td>
<td>124*</td>
</tr>
<tr>
<td>28 × 5.12</td>
<td>0.24</td>
<td>45 × 5.12</td>
<td>495</td>
</tr>
<tr>
<td>70 × 10.24</td>
<td>1.18</td>
<td>45 × 25.6</td>
<td>2478</td>
</tr>
<tr>
<td>70 × 25.6</td>
<td>2.96</td>
<td>45 × 51.2</td>
<td>4955</td>
</tr>
<tr>
<td>70 × 51.2</td>
<td>5.92</td>
<td>45 × 102.4</td>
<td>9910</td>
</tr>
</tbody>
</table>

Note: Asterisks mean that the fluence used for imaging acquisition is 0.002 J/cm².
(1) Nucleus treatment: only cell nuclei were illuminated; (2) Mitochondrial treatment: only a portion of the perinuclear region highlighted by PpIX was irradiated, with the size of the irradiation field being similar to its cell nucleus; (3) Whole cell treatment: the light was applied through the whole FOV without any selection. The irradiated areas are demonstrated in Fig. 1. All treatment schemes were applied at the focal plane of the cells only, which focused on the center section of approximately 1 μm in thickness. To compare the treatment efficacy after various regimes, a cell viability dye, propidium iodide (PI) (P3566, Life Technologies), at a working concentration of 300 ng/mL was used to stain the nuclei of cells which had a compromised cell membrane at 2 h after the initial irradiation. The fluorescence intensity from PI was collected using the excitation light at 488 nm in combination with the emission range of 600 and 750 nm. Cell regions selected for photodynamic therapy (PDT) treatment were highlighted, including cell nuclei (solid line), partial mitochondria (dotted line), and the whole cell region that can be visualized in bright field images (dashed line).

3 Results

3.1 Fluorescence of Protoporphyrin IX

Sample PpIX fluorescence images of metaplastic (CP-A) and HGD (CP-B) cells are shown in Fig. 1. Differences in cell line morphology were observed when cell mitochondria were highlighted by PpIX; CP-A cells showed dense mitochondrial distribution close to the perinuclear region demonstrating a rounder shape, while the mitochondria of CP-B cells spread out through the cells demonstrating a more elongated morphology. Maximum PpIX synthesis typically occurs within 6 h of incubation.27 It is noted that PpIX intensity in CP-B at 3 h of incubation is comparable with that in CP-A at 6 h of incubation, indicating a faster PpIX synthesis in CP-B cells. Further analysis of the cellular fluorescence images identified a small set of morphological and textural features that differentiated metaplastic (CP-A) and dysplastic (CP-B) cells with a sensitivity of 95% and specificity of 87%, yielding an area under the curve of 0.95 for the receiver operating characteristics curve.31

3.2 Subcellular Irradiation

Light treatment was given at 3 h after 5-ALA incubation, where the metaplastic and HGD cells exhibited significantly different intracellular PpIX intensity31 (photon counts of 32 ± 3 in CP-B and 23 ± 2 in CP-A, p < 0.01) in our imaging analysis study (data not shown). We employed various regimes to investigate and characterize their treatment efficacy, which could be relevant when the cellular-based treatment or ablation is integrated with endomicroscopic technology. Figures 2–5 demonstrate the dose response curves of CP-B (HGD) and CP-A (metaplasia) cells to various laser irradiation doses using 514 nm (Figs. 2 and 3) or 810 nm (Figs. 4 and 5) irradiations and the cell survival rate associated with three subcellular treatment approaches.

Since both drug and light contribute to cell toxicity, the following comparisons of PDT treatment responses were made: (1) the dose responses of different cell lines, (2) the dose responses of subcellular treatment, (3) 5-ALA and control groups without drug administration, and (4) the effect of one-photon and two-photon illuminations.

When comparing the dose response between two different cell lines incubated with 5-ALA (Fig. 2), it was observed that the HGD cells (CP-B) exhibited a significantly lower survival rate than CP-A cells when the light fluence is greater than 2.96 J/cm² (p < 0.01); CP-B cells showed an approximately 20% reduction in viability comparing with CP-A cells when using either mitochondria or whole cell treatment. When a light fluence of 5.92 J/cm² was given, CP-B cells were observed to be 37% and 51% less viable than CP-A cells with the mitochondria and whole cell treatment, respectively. These facts suggested the dysplastic cell line was more sensitive to the treatment.
proportional cytotoxic contribution from the light irradiation and the photosensitizer will be discussed further when comparing 5-ALA groups with control groups.

Various extents of phototoxic effects from subcellular PDT treatment were also observed. In general, when cells were incubated with 5-ALA, whole cell and mitochondrial irradiations yielded better efficacy when compared with nucleus treatment. This trend is obvious when the light dose is higher than 2.96 J/cm² for CP-B (p < 0.01) and 5.92 J/cm² in CP-A (p < 0.01). These results indicate that the irradiation regime involving mitochondria (PpIX location) may enhance the treatment efficacy in HGD while keeping most of metaplasia cells unharmed. For example, when mitochondria of CP-B and CP-A were both irradiated with light at 5.92 J/cm², CP-B cells were observed to be less viable (p < 0.01). This selectivity may be further enhanced when integrated with quantitative feature extraction and detection.

The dose response curves were also collected using cells without 5-ALA administration (Fig. 3). The results observed from nonstained CP-A and CP-B cells indicated that both cell lines had equivalent responses to the photothermal and photochemical laser effects in no-drug controls. Both cell lines were observed to have greater than 80% of viability with good morphological appearance after being exposed to the high fluence of 5.92 J/cm², indicating that the light-induced photochemical and photothermal reactions at this level corresponded to approximately 20% cell death, with no difference observed between cell lines and subcellular regimes (p > 0.05). In other words, when comparing the results with 5-ALA groups, 20% of cell deaths caused by mitochondrial
and whole cell treatments was attributed to the laser effect, which agrees with the observed 20% cell deaths from the nucleus-only treatment in the 5-ALA group. The relationships between different treatment regimes are described and summarized in Table 2.

As demonstrated in Figs. 4 and 5, the dose response curves were also collected using two-photon irradiation. The relationships between different treatment regimes are described and summarized in Table 3. It was noted that the dose response curves were consistent with the results obtained from the one-photon treatment, where CP-B cells were more susceptible to PpIX phototoxicity. In addition, because the focal volume of two-photon irradiation is much smaller than one-photon irradiation, the difference in viability of CP-B caused by PpIX was observed to be smaller than it was with one-photon treatment, as summarized in items c and d of Table 2. This was consistent with one-photon treatments for 5-ALA-stained HGD showing better efficacy than two-photon treatment when mitochondria were irradiated, as observed in Figs. 2 and 4 (p < 0.05). CP-A cells showed a similar viability between one- and two-photon regimes, which could be attributable to less intracellular PpIX concentration at 3 h of incubation (as mentioned in Fig. 1) or more resistance to the photosensitizing toxicity.

### 4 Discussion

In this study, subcellular irradiation of two BE cell lines was investigated as a potential in vitro model for “seek-and-treat” for esophageal dysplasia. It has also been demonstrated that the intracellular PpIX could serve as a contrast agent to highlight both types of esophageal cells (Fig. 1), while acting as a photosensitizer for selective treatment (Figs. 2 and 4). Image processing and classification have been performed in the coculture model that showed 95% sensitivity and 87% specificity for differentiating HGD based on these images. The specific PDT efficacy achieved by subcellular irradiation was also characterized and correlated with the relative amount of intracellular PpIX activation and the light fluence applied to the cells. Using confocal fluorescence images of these cells, we found that both CP-B and CP-A cells achieved their maximum PpIX formation at mitochondria within 5 to 6 h, while the dysplastic cells (CP-B) demonstrated faster PpIX production than the metaplastic cells (CP-A), especially at 3 h of incubation (photon counts of $32 \pm 3$ in CP-B and $23 \pm 3$ in CP-A, p < 0.01). Treatment was given and characterized at an earlier time frame (3 h); therefore, even if a false positive of detection occurs, it may not be harmful for the falsely treated metaplastic cells.

Correlating the discrepancy of intracellular PpIX to treatment efficacy, CP-B cells were found to be more susceptible to PpIX activation when different light fluence and subcellular treatment regimes were given. In contrast, both nonstained cell lines were observed to demonstrate similar viability when the same fluence was applied. According to the results from Figs. 2–5, comparison between groups and the corresponding explanations were summarized in Table 2. It is clear that the viability reduction between the 5-ALA groups and control (nonstained) groups was solely due to the photosensitizing effects from PpIX. This conclusion is further supported when comparing the

### Table 2 The difference in cell viability between treatment regimes using one-photon PDT (5.92 J/cm²).

<table>
<thead>
<tr>
<th>Comparison</th>
<th>Viability</th>
<th>Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>CP-A to PpIX</td>
<td>CP-A&lt;sub&gt;PpIX-whole&lt;/sub&gt; 59%</td>
<td>−20% due to CP-A responses to PpIX&lt;sup&gt;a&lt;/sup&gt;</td>
</tr>
<tr>
<td></td>
<td>CP-A&lt;sub&gt;PpIX-mito&lt;/sub&gt; 74%</td>
<td>−3% (PpIX was partially activated)&lt;sup&gt;b&lt;/sup&gt;</td>
</tr>
<tr>
<td>CP-B to PpIX</td>
<td>CP-B&lt;sub&gt;PpIX-whole&lt;/sub&gt; 8%</td>
<td>−74% due to CP-B responses to PpIX&lt;sup&gt;c&lt;/sup&gt;</td>
</tr>
<tr>
<td></td>
<td>CP-B&lt;sub&gt;PpIX-mito&lt;/sub&gt; 37%</td>
<td>−35% (PpIX was partially activated)&lt;sup&gt;d&lt;/sup&gt;</td>
</tr>
<tr>
<td>Cell sensitivity</td>
<td>CP-B&lt;sub&gt;PpIX-whole&lt;/sub&gt; 8%</td>
<td>CP-B is more sensitive to treatment&lt;sup&gt;e&lt;/sup&gt;</td>
</tr>
<tr>
<td></td>
<td>CP-B&lt;sub&gt;PpIX-mito&lt;/sub&gt; 37%</td>
<td>CP-B is more sensitive to treatment&lt;sup&gt;f&lt;/sup&gt;</td>
</tr>
<tr>
<td>Cell to light fluence</td>
<td>CP-A&lt;sub&gt;control-whole&lt;/sub&gt; 75%</td>
<td>− Equivalence to laser fluence&lt;sup&gt;g&lt;/sup&gt;</td>
</tr>
<tr>
<td></td>
<td>CP-B&lt;sub&gt;control-whole&lt;/sub&gt; 82%</td>
<td>− Equivalence to laser fluence&lt;sup&gt;h&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

Note: Superscripts a–h correspond to the items discussed in the main text.
viability between the nucleus treatment and mitochondrial treatment because the nucleus treatment irradiated the part “without PpIX.” As shown in Table 2 from items a to d of one-photon irradiation, it was observed that the CP-B cells demonstrated 51% lower viability than CP-A cells after PpIX activation using whole cell irradiation. This was consistent with the comparison between 5-ALA incubated CP-A and CP-B cells with whole cell and mitochondrial irradiations (items e and f in Table 2). Although cell death was attributed to both photothermal and photochemical interactions, an additional 50% decrease in CP-B viability compared with CP-A cells may have been related to the discrepancy in intracellular PpIX intensity, where approximately 50% more PpIX intensity was observed in CP-B cells at 3 h of incubation, as mentioned in the previous paragraph. It should be noted that the difference in survival rate can also be affected by the intrinsic PDT resistance and sensitivity of CP-A and CP-B, respectively. However, CP-B viability is comparable with the previous 5-ALA-based cell study in a well-oxygenated environment. Furthermore, equivalent cell responses to laser irradiation were found to be consistent between group comparisons as shown in items g and h of Table 2. The nucleus treatment on 5-ALA and nonstained groups caused no difference in viability, and the whole cell treatment resulted in a lower survival rate due to additional energy deposition. Moreover, according to the treatment efficacy measured from the single-plane (thickness: one-photon ~0.8 μm; two-photon ~0.5 μm) subcellular treatment at the fluence range, it is straightforward to come up with an assumption that one-photon excitation demonstrated a better killing efficacy as more PpIX along the light path was excited to induce the phototoxic effect. However, it should be noted that the photosensitizing effect would not be proportional to the fluence if the fluence rate applied to the sample depleted the ground state photosensitizers. The fluence rates used in this study were up to 0.12 W/cm² in one-photon treatment, which have negligible depletion effects on ground state photosensitizers in either high or low oxygen conditions. On the other hand, two-photon excitation with the fluence rate of 96.8 W/cm² exceeded the reported limit, thus it was possible that the ground state photosensitizer was depleted during the treatment.

PI was used as a necrotic stain to quantify cell death a short period of time after irradiation (2 h). It has been reported that apoptosis could be the main biological effect induced by PDT, although this depends on the cell type, overall dose, etc.

### Table 3
The difference in cell viability between treatment regimes using two-photon PDT (9911 J/cm²).

<table>
<thead>
<tr>
<th>Comparison</th>
<th>Viability</th>
<th>Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>CP-A to PpIX</td>
<td>CP-A_PpIX-whole 54%</td>
<td>CP-A_control-whole 70%</td>
</tr>
<tr>
<td></td>
<td>CP-A_PpIX-mito 71%</td>
<td>CP-A_PpIX-nucleus 81%</td>
</tr>
<tr>
<td>CP-B to PpIX</td>
<td>CP-B_PpIX-whole 9%</td>
<td>CP-B_control-whole 65%</td>
</tr>
<tr>
<td></td>
<td>CP-B_PpIX-mito 58%</td>
<td>CP-B_PpIX-nucleus 87%</td>
</tr>
<tr>
<td>Cell sensitivity</td>
<td>CP-B_PpIX-whole 9%</td>
<td>CP-A_PpIX-whole 54%</td>
</tr>
<tr>
<td></td>
<td>CP-B_PpIX-mito 58%</td>
<td>CP-A_PpIX-mito 71%</td>
</tr>
<tr>
<td>Cell to light fluence</td>
<td>CP-A_PpIX-whole 81%</td>
<td>CP-A_PpIX-nucleus 84%</td>
</tr>
<tr>
<td></td>
<td>CP-B_PpIX-whole 70%</td>
<td>CP-A_PpIX-whole 66%</td>
</tr>
</tbody>
</table>

Note: Superscripts a–h correspond to the items discussed in the main text.

### Table 4
Scaling factors for whole cell and mitochondrial treatments.

<table>
<thead>
<tr>
<th>Groups</th>
<th>Fractional area (partial/total activation of PpIX)</th>
<th>Fractional changes in survival rate&lt;sup&gt;a&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td>One-photon excitation</td>
<td>45% ± 11%&lt;sup&gt;b&lt;/sup&gt;</td>
<td>47% ± 4%</td>
</tr>
<tr>
<td>Two-photon excitation</td>
<td>54% ± 11%</td>
<td></td>
</tr>
</tbody>
</table>

<sup>a</sup> Calculated using the ratios of item d to item c (Tables 2 and 3, values in “notes”).
<sup>b</sup> Calculation from a single plane. The one-photon excitation volume will be further affected by the sample thickness.
the cell morphological features shown in Fig. 6, cells tended to
demonstrate a necrotic cell death within 2 h after exposure to a
high fluence. This phenomenon could be the consequence of
PpIX-induced oxidative stress.40 This agreed with a previous
report that the oxidative stress-induced necrosis happens shortly
after irradiation; the affected cells change the morphology in less
than 1 h and go through necrosis within 2 h.41

5 Conclusion
In conclusion, the premalignant BE cell lines provide a platform
to investigate detection and treatment strategies using a PDT
photosensitizer. In addition to the distinct morphological pat-
terns revealed by PpIX fluorescence imaging, this study dem-
strated potential subcellular treatment strategies and their
corresponding efficacies for cellular-based PDT, which will be
an important component in real-time integrated detection and
treatment of BE using endomicroscopic technology. Our results
demonstrated that the HGD cells were less viable than metaplas-
tic cells after PDT due to increased accumulation of PpIX at
t earlier times. More importantly, whole cell and mitochondrial
treatments induced a comparable PDT effect, although the cell
viability is different due to the fractional changes of the irra-
diated PpIX area. In contrast, nucleus irradiation only exhibited
little phototoxicity. With the aid of automatic detection, this
indicates that the appropriately targeted irradiation schemes
directed at mitochondria may further enhance treatment selec-
tivity without introducing excessive phototoxicity for non-HGD
cells. Verifying these strategies in a coculture platform and three-
dimensional cell cultures will be the logical next step complemen-
tary to quantitative imaging processing and classification.

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The authors have declared that no competing financial inter-
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