Targeted Delivery of Auristatin-Modified Toxins to Pancreatic Cancer Using Aptamers

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Pancreatic cancer is one of the most lethal malignancies. Treatment with the first-line agent, gemcitabine, is often unsuccessful because it, like other traditional chemotherapeutic agents, is non-specific, resulting in off-target effects that necessitate administration of subcurative doses. Alternatively, monomethyl auristatin E (MMAE) and monomethyl auristatin F (MMAF) are highly toxic small molecules that require ligand-targeted delivery. MMAE has already received FDA approval as a component of an anti-CD30 antibody-drug conjugate, brentuximab vedotin. However, in contrast to antibodies, aptamers have distinct advantages. They are chemicals, which allows them to be produced synthetically and facilitates the rapid development of diagnostics and therapeutics with clinical applicability. In addition, their small size allows for enhanced tissue distribution and rapid systemic clearance. Here, we assayed the toxicity of MMAE and MMAF conjugated to an anti-transferrin receptor aptamer, Waz, and an anti-epidermal growth factor receptor aptamer, E07, on the pancreatic cancer cell lines Panc-1, MIA PaCa-2, and BxPC3. In vitro, our results indicate that these aptamers are a viable option for the targeted delivery of toxic payloads to pancreatic cancer cells.

INTRODUCTION

Pancreatic cancer is the 4th leading cause of cancer death.1 Pancreatic ductal adenocarcinoma (PDAC) is by far the most common subtype of pancreatic cancer, representing 90% of cases. A minority of patients (10%–15%) present with disease that is amenable to surgery, the only treatment that offers any hope of cure. However, even with adjuvant therapy (i.e., chemotherapy), 80% of these patients experience a relapse within 2 years of surgery.2

Most patients present with either locoregional disease that precludes surgery because of vascular invasion or advanced-stage disease with distant metastases. For patients with advanced disease, current therapy relies on gemcitabine as the first-line treatment, yet the benefit is modest.3 Overall, this treatment regimen results in a 5 year survival rate of only 8%.1 Numerous studies using novel cytotoxic agents and novel combinations of existing chemotherapies have failed to improve survival.4 Disappointingly, PDAC is one of the few cancers with recent increases in incidence or mortality.1 Because of the extremely aggressive nature of PDAC, in particular its propensity for distant metastasis and broad chemoresistance, there is an obvious need for the development of new therapies.

Advances in the field of antibody-drug conjugates (ADCs) have demonstrated the power of targeting potent cytotoxic drugs directly to cell surface receptors. The targeted delivery of small molecule cytotoxics, such as calicheamicin, has been replaced with newer classes of drugs, the maytansine and auristatin derivatives, both of which work by inhibiting microtubule assembly.5–7 Due to the extreme potency of ADCs, achieving a therapeutic effect requires less than 1% of an injected dose to localize to tumors.8 The development of emtansine, a maytansine derivative, led to U.S. Food and Drug Administration (FDA) approval of ado-trastuzumab emtansine (Kadcyla), an anti-HER2 ADC for the treatment of metastatic breast cancer.9 In addition, in July 2011, brentuximab vedotin (Adcentris), an anti-CD30 monomethyl auristatin E (MMAE) ADC, was approved by the FDA for the treatment of Hodgkin’s lymphoma and systemic anaplastic large-cell lymphoma.10 In the case of PDAC, MMAE has also shown promise when conjugated to an anti-tissue factor antibody11 and an anti-mesothelin antibody.12 It has also shown efficacy as a sensitizer for ionizing radiation therapy.13

Targeting PDAC requires knowledge of cell-specific surface receptors, and although no magic bullets are known to home exclusively to PDAC, the epidermal growth factor receptor (EGFR) and the transferrin receptor (TfR, CD71) have demonstrated high levels of expression on PDAC.14,15 In addition, EGFR overexpression is associated with poor prognosis and disease progression,16 while TfR overexpression supports malignant growth by modulating mitochondrial respiration and reactive oxygen species production.15,17 Both receptors are internalized by clathrin-mediated endocytosis, suggesting a potential mechanism for delivery of toxins.

Work in the field of targeted delivery has focused largely on the use of antibodies; however, challenges in the field remain (reviewed in Donaghy8). Aptamers are single-stranded oligonucleotides selected to bind a target with high affinity and specificity. Their ability to be chemically synthesized facilitates toxin conjugation, and their small size may be optimal for tumor penetration. An anti-TfR aptamer, Waz, and an anti-EGFR aptamer, E07, have already been selected...
and validated\textsuperscript{18,19} Furthermore, E07 has been used to deliver gemcitabine to the EGFR-expressing PDAC cell line MIA PaCa-2 to induce cell death\textsuperscript{20}

Previously, our lab demonstrated the ability to generate aptamer-toxin conjugates using a recombinant variant of the ribosomal toxin gelonin and the minimized anti-prostate-specific membrane antigen (PSMA) aptamer, A9.min\textsuperscript{21} Here, we assessed the ability to generate small molecule aptamer-toxin conjugates. Using thiol-reactive variants of the auristatin toxins MMAE and monomethyl auristatin F (MMAF), we generated aptamer-toxin conjugates using both the anti-EGFR aptamer, E07, and the anti-TfR aptamer, Waz, and tested their cytotoxicity on three distinct PDAC cell lines: Panc-1, MIA PaCa-2, and BxPC3. Our analysis takes advantage of the varying levels of expression of EGFR and TfR on these cell lines, as well as fundamental differences in the behavior of MMAE and MMAF.

RESULTS

Validation of Target Expression and Aptamer Binding on PDAC Cell Lines

The PDAC cell lines Panc-1, MIA PaCa-2, and BxPC3 are all reported to express TfR and EGFR\textsuperscript{15,22,23} To confirm this, we screened these cell lines for target receptor expression by flow cytometry using commercially available antibodies. Cells were incubated with phycoerythrin (PE)-labeled TfR and EGFR antibodies. After incubation, cells were washed and read by flow cytometry. As seen in Figure 1, all three PDAC cell lines express both TfR and EGFR, with Panc-1 and MIA PaCa-2 cells demonstrating higher expression levels than BxPC3 cells. The dose-response curves were used to determine the apparent dissociation constant ($K_D$) and maximal binding ($B_{max}$) for each aptamer on each cell line (Table 1).

Synthesis and Characterization of Aptamer-Auristatin Conjugates

We generated aptamer-auristatin conjugates from the same thiol-modified aptamers used to generate the preceding dye-labeled conjugates. Conjugates were made using one of two thiol-reactive toxin variants: the membrane-permeable auristatin derivative MMAE bearing a cathepsin-cleavable valine-citrulline linker (MC-VC-PAB-MMAE) or the membrane-impermeable auristatin derivative MMAF (MC-MMAF) (Figure 3A). Reduced aptamers were incubated with a 3-fold molar excess of either MC-VC-PAB-MMAE or MC-MMAF. The progress of the reaction was monitored by reverse-phase high-performance liquid chromatography (HPLC) and routinely proceeded to 100% (Figure 3B). Conjugates were subsequently desalted, and their identities were confirmed by mass spectrometry (Figure 3C).

Because of the potential for the presence of the drug to interfere with aptamer binding function, we performed a competition binding experiment to ensure that drug conjugation did not adversely affect
aptamer function. DyLight 650-labeled aptamers (Waz or E07) were incubated with increasing concentrations of MMAE-, MMAF-, or Alexa Fluor 488 (AF488)-labeled aptamer. As shown in Figure 3D, both the drug conjugates and the dye conjugate demonstrate a similar ability to compete with DyLight 650 for binding, indicating that aptamer function is not adversely affected by drug conjugation.

Critical to the function of the targeted toxin is its ability to separate from the targeting ligand upon uptake and entry into the cytoplasm. The valine-citrulline dipeptide in MC-VC-PAB-MMAE renders it susceptible to degradation by cathepsin B in the acidic environment of the endosome. MC-MMAF, however, contains no such linkage. To validate this chemistry while conjugated to an aptamer, we incubated MMAE- and MMAF-conjugated aptamers both with cathepsin B in acidic buffer and without cathepsin B in neutral buffer. As shown in Figures 4A and S1, free aptamer can be readily distinguished from both the MMAE and the MMAF conjugates on a 20% denaturing (7 M urea) polyacrylamide gel. Only aptamer-MMAE conjugates, not MMAF conjugates, incubated with cathepsin B under acidic conditions demonstrated an increase in electrophoretic mobility, indicating cleavage. No cleavage of MMAE was observed without cathepsin B at neutral pH.

Finally, although cathepsin B is known to be ubiquitous, with varied expression levels and distribution throughout the endosomes and lysosomes of most cells,24 we performed a co-localization study using AF488-Waz and Magic Red, an activity-based probe that detects the presence of cathepsin B in cells. Following cellular uptake, the aptamer (Figure 4B, green) demonstrated characteristic punctate staining, consistent with localization to endosomal and lysosomal compartments that co-localized well with that of cathepsin B staining (Figure 4B, red).

**Evaluation of Aptamer-Auristatin Conjugate Toxicity**

We assessed the efficacy of our aptamer-targeted toxins on three PDAC cell lines. Cells were incubated with conjugates for 4 days, at which time toxin-containing media were replenished with fresh media. After an additional 2 days, toxicity was determined. All assays were performed using both free toxin and a nontargeting control-toxin conjugate (C36-MMAE/F).

Conjugates of Waz and E07 bearing the cathepsin-cleavable, membrane-permeable MC-VC-PAB-MMAE both resulted in improvement in cytotoxicity on the PDAC cell lines when compared to
Figure 3. Synthesis, Purification, and Characterization of Aptamer Auristatin Conjugates

(A) Chemical structure of MMAE and MMAF used in our studies. MMAE includes the cleavable linker, valine-citrulline, and the self-immolative spacer, p-amino-benzylcarbonyl (PABC). (B) HPLC traces of Waz during reduction and conjugation. (Top panel) Analytical analysis of purified, reduced Waz. (2nd panel) Analysis of Waz MMAE.

|          | Predicted MW | Measured MW |
|----------|--------------|-------------|
| C36      | 12085.2      | 12084.3     |
| C36 MMAE | 13400.8      | 13401.3     |
| C36 MMAF | 13099.3      | 13099.6     |
| Waz      | 15933.4      | 15931.3     |
| Waz MMAE | 17249.0      | 17248.5     |
| Waz MMAF | 16857.5      | 16856.1     |
| E07      | 15686.4      | 15685.7     |
| E07 MMAE | 17002.0      | 17002.1     |
| E07 MMAF | 16610.6      | 16610.4     |

(D) Relative Fluorescence E07 DyLight650 vs Waz Dylight650

(legend continued on next page)
toxicity of a nontargeting control, C36-MMAE (Figure 5). Conjugates targeting TfR (Waz-MMAE) demonstrated the greatest increase in cytotoxicity when compared to the negative control. On all three cell lines, Waz-MMAE conjugates displayed half-maximal inhibitory concentration (IC50) values of 35 nM, while conjugates targeting EGFR proved somewhat less toxic and demonstrated a more varied response: IC50 of 55 nM on PANC-1 cells and ~120 nM on MIA PaCa-2 and BxPC3 cells (Table 3). Consistent with its ability to enter cells on its own, free MMAE yielded the lowest observed IC50 values across all three cell lines.

Similar experiments were performed with conjugates generated using the non-membrane-permeable MMAF-toxin conjugates. As shown in Figure 5 and summarized in Table 3, results with MMAF were somewhat more varied across cell types. Targeting TfR proved the most toxic: the IC50 on MIA PaCa-2 cells was relatively low MMAE-conjugated Waz. (3rd panel) Analysis of MMAF-conjugated Waz. (Bottom panel) Analysis of DyLight 650-conjugated Waz. A shift in the observable product, absence of a second peak, and superimposition of the 260 and 650 traces for DyLight 650 indicate >99% conjugation. Reduction, conjugation, and removal of excess dye or toxin were performed as described in Materials and Methods. (C) Comparison of predicted and measured molecular weights of aptamers and aptamer-toxin conjugates. Mass spectrometry was used to measure the molecular weight of aptamers and aptamer-toxin conjugates. (D) Competition analysis of dye and toxin (MMAE or MMAF)-labeled aptamers. Constructs tested are as indicated. Assay was performed as described in Materials and Methods.

DISCUSSION

Aptamer-toxin conjugates have the potential to improve the therapeutic index of chemotherapeutics by targeting toxins to cancer cells while bypassing healthy cells. Their small size abets tissue penetration, and their rapid clearance may improve off-target effects, a

![Figure 4. Further Characterization of Aptamers and Aptamer Conjugates](image)

(A) Cleavage of aptamer-toxin conjugates in the presence of cathepsin B. Lane 1: aptamer reduced with 10 mM TCEP at 70°C for 10 min. Lane 2: aptamer-MMAE conjugate incubated in PBS (pH 7). Lane 3: aptamer-MMAE conjugate incubated in PBS (pH 4), with 0.1 mg/mL of cathepsin B. Lane 4: aptamer-MMAF conjugate incubated in PBS (pH 7). Lane 5: aptamer-MMAF conjugate incubated in PBS (pH 4), with 0.1 mg/mL of cathepsin B. Samples in lanes 2–5 were incubated for 24 hr at 37°C. Arrows indicate reduced, unconjugated aptamer. (B) Colocalization of Waz-AF488 (Waz; green) and cathepsin B (CB; red). (51 nM), while the IC50s on Panc-1 and BxPC3 cells were somewhat higher (190 and 450 nM, respectively). Targeting EGFR provided the greatest toxicity on MIA PaCa-2 cells (220 nM) and had relatively less effect on BxPC3 cells (~1,000 nM) and Panc-1 cells (>1,000 nM). Consistent with its inability to cross the extracellular membrane, MMAF alone was significantly less toxic (~10-fold) than MMAE. Conjugation of MMAF to our nontargeting control aptamer rendered it even less toxic (>1,000 nM on Panc-1 and BxPC3 cells and ~950 nM on MIA PaCa-2 cells).

To ensure the mechanism of action of both Waz and E07 conjugates, we performed toxicity assays on B16 cells, a murine cell line for which neither Waz nor E07 demonstrate any significant binding (Figure S2). Consistent with these data, no significant toxicity (IC50 >1,000 nM) was observed using these constructs on these cells, a level similar to that seen using our non-targeted control, C36 (Table 3; Figure S3).

Finally, to ensure that the observed differences in cell death were not a result of conjugate degradation and drug release over the course of the assay, we performed a stability assay using fluorescently labeled aptamers in culture media. Little difference in stability was observed (Figure S4).
potential consequence of the long half-lives of antibodies.\textsuperscript{25,26} Perhaps more importantly, as chemicals, aptamers may simplify drug development, removing the biological component (i.e., the antibody) and providing a facile means for site-specific conjugation.

Previously, we streamlined conjugation of an aptamer to a protein toxin by chemically synthesizing a minimized aptamer bearing a 5′ thiol and generating an activated pyridyl disulfide for subsequent toxin conjugation.\textsuperscript{21} Here, we exploited the maleimide functional groups on commercially available MMAE and MMAF derivatives to further simplify the process of conjugating aptamers to these small molecule toxins. Unlike our previous work, these aptamer-small molecule toxin conjugates are chemicals, not biologics.

Based on the observation that many PDACs overexpress TfR and EGFR, we assessed the ability to use anti-TfR and anti-EGFR
cytoplasm is critical for function, limits on the rate of release from them more or less toxic. Because migration of the toxin to the cytoplasm is critical for function, limits on the rate of release from the aptamer or the endosome will affect the observed toxicity. For example, in the case of the protein toxin, gelonin, endosomal escape is the rate-limiting step in ligand-targeted delivery, not ligand binding or conjugate internalization.27

In the case of the MC-VC-PAB-MMAE conjugates, in which the MMAE toxin is membrane permeable following cleavage, the rate-limiting step may be processing within the endosome. In the case of membrane-impermeable MMAF, following uptake, toxicity is likely a consequence of both aptamer degradation and endosomal escape, processes that are likely different for different cells lines and different uptake pathways.

Although the IC_{50} of aptamer-MMAE conjugates were consistently lower than those of aptamer-MMAF conjugates, killing by the non-targeting control C36-MMAE was significant, resulting in a relatively low therapeutic index for these conjugates on PDAC cells even though killing of off-target B16 cells was negligible (Figure 5; Figure S3; Table 3). Aptamer-MMAF conjugates, however, while less toxic, demonstrated a much larger therapeutic index on PDAC cells and again, negligible killing on B16 cells. MMAE is a more toxic molecule than MMAF,28 which may partially explain this result. In addition, once released, MMAE can permeate the cellular membrane to kill other cells (i.e., the bystander effect). In vivo, the bystander effect can help kill surrounding cells in the tumor.29,30 However, in vitro, it may have inadvertently lowered the IC_{50} of C36-MMAE.

We acknowledge that this work only examines two variants of the auristatins. When we initiated this work, MMAE was only commercially available conjugated to the cathepsin B-cleavable valine-citrulline dipeptide linker and MMAF was only supplied conjugated to a non-cleavable linker. This had the effect of coupling cell permeability (MMAE is cell permeable; MMAF is not) with linker cleavability. MMAE and MMAF are now commercially available with a variety of linkers: non-cleavable MMAE and cleavable MMAF, as well as substitutions for the valine-citrulline linkage to other dipeptide linkages28 and substitutions for the maleimidocaproyl to a bromoacacetamidecaproyl.25 These substitutions have been shown to directly affect the therapeutic index of antibody-auristatin conjugates.25,28 Moving forward, we aim to explore the effect of these new linker chemistries on the toxicity of aptamer-drug conjugates. We hypothesize that MMAF conjugated to a cathepsin B-cleavable linker holds particular promise. Though MMAF is less toxic than MMAE, its inability to cross the lipid bilayer into other cells, coupled with its higher aqueous solubility,31 should result in an improved therapeutic index in vitro if endosomal escape can be facilitated by a cleavable linker. However, the behavior of different toxin variants on different cell lines appears idiosyncratic and likely must be determined empirically.25,31,32

Finally, it is worth comparing our work to a publication by Yoon et al.33 In that work, the authors took advantage of another important aspect of aptamers: the ability to use them agonistically to target-specific cell types without prior knowledge of specific surface receptors. The authors adapted a previously reported anti-PDAC aptamer,

### Table 2. Apparent K_{D} and B_{max} of Aptamers on PDAC Cell Lines

| Cell Line | Apparent K_{D} (nM) | B_{max} (Relative Units) |
|-----------|---------------------|-------------------------|
| Panc-1    | 170 ± 59            | 51 ± 22                 |
| E07       | 51 ± 22             | 10 ± 1.2                |
| MIA PaCa-2| 160 ± 42            | 26 ± 20                 |
| Waz        | 14 ± 1.3            | 4.9 ± 0.79              |
| BxPC3     | 110 ± 66            | 67 ± 16                 |
| E07       | 12 ± 2.4            | 16 ± 1.0                |
| B16       | >1,000              | >1,000                  |

Aptamer dose-response curves were used to determine the apparent K_{D} for Waz and E07 on Panc-1, MIA PaCa-2, BxPC3, and B16 cells. Apparent K_{D} values are shown in nanomolars, B_{max} values are shown in relative units, and both were calculated based on a plot of the median fluorescence normalized to the fluorescence of C36 to account for nonspecific uptake. Results are based on a minimum of three independent experiments.
All other synthesis reagents were purchased from Glen Research. CO2 and 99% humidity. Panc-1, MIA PaCa-2, and B16 cells were aptamer stability for MMAF conjugates. Future work should attempt to improve narrow therapeutic window. The largest therapeutic window was MMAE conjugates were correspondingly low, suggesting a rather anti-transferrin aptamer, Waz, and the anti-EGFR aptamer, E07, synthesized and prove toxic to PDAC cells. Here, we assayed the conjugated to MMAE and MMAF on three PDAC cell lines. Our results cannot be chosen based on binding alone. While aptamer-MMAE combinations. While the literature has given considerable attention to P19, and demonstrated its ability to target and deliver the auristatin MMAE and a derivative of maytansine (DM1). The toxins were attached to a short oligonucleotide and then to the aptamer via hybridization, unlike the direct conjugation strategy we employed here. In that study, dose-dependent killing with the targeted toxin was also observed. Altogether, these results bode well for the development aptamer-small molecule toxin conjugations for the treatment of PDAC, as well as other cancers.

In summary, aptamer-auristatin conjugates are quickly and reliably synthesized and prove toxic to PDAC cells. Here, we assayed the anti-transferrin aptamer, Waz, and the anti-EGFR aptamer, E07, conjugated to MMAE and MMAF on three PDAC cell lines. Our results show that the best aptamer for targeted delivery of a toxin to a cell line cannot be chosen based on binding alone. While aptamer-MMAE conjugates displayed low IC50 values across all three cell lines, C36-MMAE conjugates were correspondingly low, suggesting a rather narrow therapeutic window. The largest therapeutic window was found for MMAF conjugates. Future work should attempt to improve aptamer stability for in vivo use and explore different linker and toxin combinations. While the literature has given considerable attention to ADCs for the treatment of PDAC, our data suggest that aptamer-drug conjugates are a viable alternative with many potential advantages.

**MATERIALS AND METHODS**

**Cells and Media**
Cells were purchased from ATCC and maintained at 37°C with 5% CO2 and 99% humidity. Panc-1, MIA PaCa-2, and B16 cells were maintained in DMEM supplemented with 10% fetal bovine serum (FBS). BxPC3 cells were maintained in RPMI supplemented with 10% FBS.

**Aptamer Synthesis and Labeling**
Aptamers were synthesized in house via solid-phase synthesis on an Expedite 8909 DNA synthesizer (Biolytic Lab Performance, Fremont, CA). Phosphoramidite monomers ([2’-hydroxyl A/G, 2’-fluoro-modified C/U] were purchased from ChemGenes (Wilmington, MA). All other synthesis reagents were purchased from Glen Research (Sterling, VA). Aptamers were synthesized on an inverted dT or fluorescein (FAM) controlled pore glass (CPG) column (where indicated bearing a 5’ thiol modification using a thiol modifier, C6 S-5 phosphoramidite. The final 5’ dimethoxytrityl group was left on to facilitate purification. Aptamer sequences can be found in Table 2.

Table 3. IC50 of Aptamer-Toxin Conjugates on PDAC Cell Lines

|                  | Toxic (nM) | C36 (nM) | Waz (nM) | E07 (nM) |
|------------------|-----------|----------|----------|----------|
| Panc-1 MMAE      | 11 (8.9–13)| 110 (76–148)| 35 (31–41)| 25 (44–69) |
| MMAF >1,000      |           | 190 (130–280) |        |       |
| MIA PaCa-2 MMAE  | 7.5 (6.5–8.7)| 290 (250–340)| 37 (33–41)| 130 (90–190)|
| MMAF >1,000      |           |           |         |         |
| BxPC3 MMAE       | 16 (15–18)| 220 (200–240)| 39 (37–41)| 110 (100–120) |
| MMAF >1,000      |           |           |         |         |
| B16 MMAE         | 56 (46–69)| >1,000   | >1,000   | >1,000   |
| MMAF >1,000      |           |           |         |         |

The IC50 values of aptamer-toxin conjugates, nontargeting control toxin conjugates, and toxin alone on PDAC cell lines are reported in nanomolars, including the 95% confidence interval. Results are based on three discrete measurements at each concentration.

In that study, dose-dependent killing with the targeted toxin exceeded to 100%. Excess dye and toxin were removed by buffer exchanging into PBS± using a 10 K concentrator (Millipore, Ireland). Reduced aptamers were incubated for 1 hr at room temperature with a 3- to 5-fold molar excess of mal- 

eimide-labeled DyLight 650 (Thermo Fisher Scientific, Waltham, MA), AF488 (Life Technologies, Carlsbad, CA), MC-VC-PAB-MMAE (Concordis Biosystems, San Diego, CA), or MC-MMAF (Concordis Biosystems). Conjugation was confirmed via HPLC and routinely proceeded to 100%. Excess dye and toxin were removed by buffer exchanging into PBS± using a 10 K concentrator (Millipore).

Aptamers in 0.1 M TEAA were reduced with 10 mM Tris (2-chloroethyl) phosphate (TCEP) by heating to 70°C for 10 min and cooling for 1 hr at room temperature. Reduction was confirmed by HPLC. TCEP was removed by buffer exchanging into PBS± using a 10 K concentrator (Millipore, Ireland). Reduced aptamers were incubated for 1 hr at room temperature with a 3- to 5-fold molar excess of maleimide-labeled DyLight 650 (Thermo Fisher Scientific, Waltham, MA), AF488 (Life Technologies, Carlsbad, CA), MC-VC-PAB-MMAE (Concordis Biosystems, San Diego, CA), or MC-MMAF (Concordis Biosystems). Conjugation was confirmed via HPLC and routinely proceeded to 100%. Excess dye and toxin were removed by buffer exchanging into PBS± using a 10 K concentrator (Millipore).

The identity of the labeled aptamers was confirmed by liquid chromatography–mass spectrometry (LC-MS), using a Waters Acquity ultra-performance liquid chromatography (UPLC) system coupled to a Waters Synapt G2 quadrupole-time-of-flight hybrid mass spectrometer. For analysis, aptamers were resuspended at 200 μM in PBS± using a 10 K concentrator. All samples were desalted using a linear gradient of MeOH in 400 mM hexafluoroisopropanol (HFIP)/8 mM triethylamine (TEA) using a Waters XBridge C18 2.1 × 50 mm column. The resulting mass spectra were deconvoluted using MagTran.34

**Antibody and Aptamer Binding Assays**
To determine antibody binding, cells were washed in PBS± and lifted with 0.5% trypsin, and 50,000 per well were stained with PE-labeled anti-TIR antibody (Catalog No. 2270530, Sony Biotechnology, San
distinguish live from dead cells. Cells were then washed twice with FACS buffer, pelleted in a swinging-bucket centrifuge at 300 g for 5 min, and resuspended in FACS buffer with 100 ng/mL of bisbenzimide to distinguish live from dead cells. Cells were read on an iCyt Eclipse EC800 flow cytometer, and data were analyzed using FlowJo.

To determine aptamer binding, cells were seeded at 50,000 cells per well in a 96 well tissue culture plate. The following day, 90 μL of fresh media containing 1.0 mg/mL of DNA, sodium salt, and salmon testes (single-stranded DNA [ssDNA]; Millipore, Billerica, MA) was added to the cells and allowed to incubate for 1 hr. DyLight 650-labeled aptamers were prepared as 10 × stocks in PBS, heated to 70°C for 3 min, and allowed to renature at room temperature for 15 min. Aptamers were then added to cells and incubated for 1 hr at 37°C. Media were removed, and cells were washed twice with 100 μL of PBS and lifted with 0.5% trypsin. FACS buffer was added to inactivated the trypsin, and cells were pelleted by centrifugation and resuspended in FACS buffer with bisbenzimide. Cells were read on an iCyt Eclipse EC800 flow cytometer, and data were analyzed using FlowJo.

Aptamer Competition Assays
MIA PaCa-2 cells were washed in PBS and lifted in 0.5% trypsin, and 50,000 per well were suspended in FACS buffer with 1.0 mg/mL of ssDNA for 30 min. DyLight 650-, AF488-, MMAE-, and MMAF-conjugated aptamers were prepared as 20 × stocks in PBS, heated to 70°C for 3 min, and allowed to renature at room temperature for 15 min. To confirm that conjugating to DyLight 650, MMAE, or MMAF did not perturb aptamer binding, DyLight 650-labeled aptamer was mixed with the same aptamer labeled with AF488, MMAE, or MMAF and then added to cells. The final concentration on cells of DyLight 650-labeled aptamer was held constant at 100 nM, while the concentration of AF488-, MMAE-, and MMAF-labeled aptamer ranged from 10 to 1,000 nM. Aptamers were incubated with cells for 30 min, after which they were pelleted in a swinging-bucket centrifuge at 300 × g for 5 min, washed twice in FACS buffer, and resuspended in FACS buffer with 100 ng/mL of bisbenzimide to distinguish live from dead cells.

Cathepsin B Assay
Assay for cathepsin B cleavage was modified from Koga et al. 11 50 ng of FAM-labeled aptamer-toxin conjugates were incubated for 24 hr at 37°C in either PBS (pH 7) or PBS (pH 4) with 0.1 mg/mL of cathepsin B (Athens Research & Technology, Athens, GA). Conjugates were run on a 20% polyacrylamide denaturing gel and imaged using a Storm 840 imaging system (GE Healthcare, Marlborough, MA).

Microscopy
Cells were seeded at 50,000 cells per well in an 8 well Lab-Tek chambered cover glass (Thermo Fisher Scientific). The following day, 180 μL of fresh media containing 1.0 mg/mL of DNA, sodium salt, and salmon testes (ssDNA; Millipore, Billerica, MA) was added to the cells and allowed to incubate for 1 hr. Aptamers were added to cells at a final concentration of 500 μM, and Magic Red cathepsin B assay (ImmunoChemistry Technologies, Bloomington, MN) was added to cells at the manufacturer’s recommended concentration. After 1 hr, cells were washed twice with PBS and imaged at 60× using a Nikon Eclipse Ti 90.

Evaluation of Aptamer-Auristatin Conjugate Toxicity
Cells were seeded at 4,000 cells per well in a 96 well tissue culture plate. The following day, cells were blocked for 1 hr with 90 μL of fresh media containing 1.0 mg/mL of ssDNA and 1% penicillin-streptomycin. Aptamer-toxin conjugates were prepared as 10 × stocks in Dulbecco PBS (DPBS), heated to 70°C for 3 min, and renatured at room temperature for 15 min. MMAE and MMAF (MedChem Express, Princeton, NJ) were prepared as 5,000 × stocks in DMSO and then diluted to 10 × in DPBS. Aptamer-toxin conjugates and toxins alone were added, and cells were incubated at 37°C with 5% CO2 and 99% humidity. After 4 days, toxin-containing media were replaced by fresh media. After 2 additional days, cell viability was assayed with AlamarBlue (Invitrogen, Carlsbad, CA). Half-maximal inhibitory concentration (IC50) was determined using the variable slope model in GraphPad Prism 7 software.

SUPPLEMENTAL INFORMATION
Supplemental Information includes Supplemental Materials and Methods and four figures and can be found with this article online at https://doi.org/10.1016/j.omtn.2017.11.013.

AUTHOR CONTRIBUTIONS
C.K. conducted all experiments. Methods and approaches were conceived by C.K. and M.L.

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