

Cross-Linked Forms of the Isolated N-Terminal Domain of the Lethal Factor Are Potent Inhibitors of Anthrax Toxin[∇]

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The proteins that comprise anthrax toxin self-assemble at the mammalian cell surface into a series of toxic complexes, each containing a heptameric form of protective antigen (PA) plus up to a total of three molecules of the enzymatic moieties of the toxin (lethal factor [LF] and edema factor [EF]). These complexes are trafficked to the endosome, where the PA heptamer forms a pore in the membrane under the influence of low pH, and bound LF and EF unfold and translocate through the pore to the cytosol. To explore the hypothesis that the PA pore can translocate multiple, cross-linked polypeptides simultaneously, we cross-linked LF_N, the N-terminal domain of LF, via an introduced cysteine at its N or C terminus and characterized the products. Both dimers and trimers of LF_N retained the ability to bind to PA pores and block ion conductance, but they were unable to translocate across the membrane, even at high voltages or with a transmembrane pH gradient. The multimers were remarkably potent inhibitors of toxin action in mammalian cells (20- to 50-fold more potent than monomeric LF_N) and in a zebrafish model system. These findings show that the PA pore cannot translocate multimeric, cross-linked polypeptides and demonstrate a new approach to generating potent inhibitors of anthrax toxin.

Bacillus anthracis causes pathology in infected human or animal hosts in part through the concerted action of three proteins, collectively termed anthrax toxin. The toxin consists of two enzymatic moieties, termed lethal factor (LF) and edema factor (EF), and a transport protein, termed protective antigen (PA), that delivers both LF and EF to the cytosol. LF is a 90-kDa zinc-dependent metalloprotease that cleaves mitogen-activated protein kinase kinases (6, 20, 24), and EF is an 89-kDa calmodulin-dependent adenylate cyclase (12). The intracellular actions of these enzymes impair the functions of various cells and can lead to the death of infected hosts.

Delivery of LF and EF to the cytosol begins with binding of PA (83 kDa) to a receptor. Two receptors have been identified: ANTXR1 (for anthrax toxin receptor 1; also known as ATR/TEM8) and ANTXR2 (for anthrax toxin receptor 2; also known as CMG2) (3, 23). Receptor-bound PA is proteolytically processed by furin or a furin-like protease (19), resulting in the removal of a 20-kDa fragment (PA20) from the N terminus. The remaining, receptor-bound fragment (PA63, 63 kDa) spontaneously oligomerizes, forming a ring-shaped heptamer, called the prepore, which is capable of binding up to three molecules of LF and/or EF with high affinity (17, 18). The resulting toxic complexes are internalized, and the low pH within the endosome promotes a conformational change in the prepore moiety that allows it to insert into endosomal mem-

branes and form a pore. The conformational transition of the prepore to the pore depends on the association of the 2β2-2β3 loops of the seven PA63 subunits to form a membrane-spanning, 14-stranded β barrel (1, 21, 22).

Recent evidence shows that the pore plays an active, chaperone-like role in the translocation of LF and EF across membranes (8, 15). Translocation requires unfolding of the enzymatic factors (26), and there is evidence that the pH gradient across the endosomal membrane drives the translocation process (7). The seven Phe-427 residues of PA63 form what we have termed the Phe clamp, a structure in the pore lumen that is believed to interact directly with the translocating polypeptide chain to promote its passage across the membrane (8).

The PA binding domain of LF, termed LF_N, corresponds to the N-terminal 263 amino acids of LF. LF_N binds to the prepore with high affinity (K_D of ~1 nM), and, when isolated as a discrete protein, this domain alone can be shown to translocate through the pore (28). Also, some fusion proteins containing LF_N fused to heterologous proteins are able to undergo PA-dependent translocation into cells (16, 26) or across planar lipid bilayers (9, 10, 14, 28). The crystallographic structure of LF shows LF_N to be a discrete helix-rich domain with a disordered N-terminal region that is essential for translocation. The disordered region, corresponding to the first ~30 amino acids and densely populated with acidic and basic residues, is believed to enter the pore and initiate N-terminal-to-C-terminal translocation of LF and EF across the membrane.

Although much has been learned in recent years about the translocation of anthrax toxin, many questions remain unanswered. Here we have addressed the question of whether multimeric, cross-linked polypeptides can be translocated through the heptameric PA63 pore simultaneously. The finding that cross-linked forms of the N-terminal domain of LF do not

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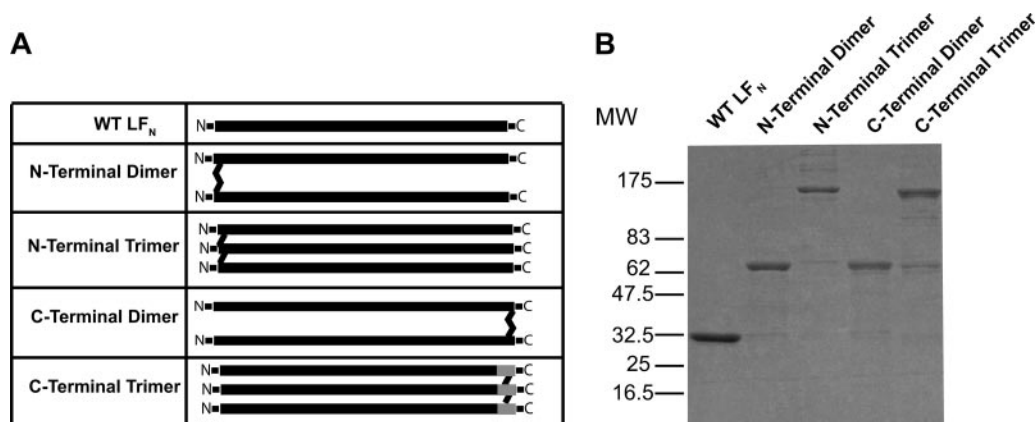


FIG. 1. Cross-linking of LF_N. (A) Schematic representation of constructs used in this study. The LF_N used contained an extra three amino acids at the N terminus due to the thrombin recognition site. (B) Cross-linked proteins were purified as described in Materials and Methods, analyzed by sodium dodecyl sulfate-polyacrylamide gel electrophoresis, and stained with Coomassie brilliant blue. WT, wild type; MW, molecular weight.

translocate led to the discovery that such multimers are potent inhibitors of anthrax toxin, both in cultured mammalian cells and in a vertebrate animal model.

MATERIALS AND METHODS

Plasmid construction. The expression construct pET15b-LF_N has been described previously (9). The mutations A1C and R263C were introduced into LF_N using QuikChange site-directed mutagenesis (Stratagene). The expression construct pET15b-LF_N-(Gly₄Ser)₃-Cys was made by amplifying LF_N by PCR using a primer that contained an in-frame addition of (Gly₄Ser)₃ repeats followed by a cysteine residue and stop codon. The DNA sequence encoding a fusion protein between LF_N and the catalytic domain of diphtheria toxin (DTA) was introduced into pET15b (Novagen) using 5' NcoI and 3' BamHI sites. A small polylinker containing the restriction sites XhoI, SacI, and NotI and an additional base to maintain the reading frame was introduced between the two protein domains. All constructs were verified by DNA sequencing.

Preparation of proteins. Recombinant LF_N and PA were purified as previously described (9, 27). PA was activated and heptamerized as previously described (5). Recombinant LF_N-DTA was expressed in BL21(DE3) *Escherichia coli* cells (Novagen). Growth and expression of LF_N-DTA were carried out in a 5-liter Bioflo 110 fermentor (New Brunswick Scientific). Cells were grown at 37°C in ECPM1 medium (2) to optical density at 600 nm of ~5, sparged with a 60% air-O₂ mixture, and induced at 30°C with 1 mM IPTG (isopropyl-β-D-thiogalactopyranoside). Bacteria were harvested and lysed with a French press and sonication in 20 mM Tris (pH 8.5)–1 M NaCl. The lysate was centrifuged at 30,000 × g, and the supernatant was applied to an Ni²⁺-chelated Sepharose column (Amersham Pharmacia). LF_N-DTA was eluted from the column using a 35 mM to 385 mM imidazole gradient. The His₆ tag was removed from LF_N-DTA using bovine α-thrombin (Enzyme Research Laboratories) overnight at room temperature. Thrombin was removed from cleaved LF_N-DTA using a Q-Sepharose anion-exchange column (Amersham Pharmacia).

Cross-linking. N-terminal LF_N dimers and trimers (using A1C LF_N) and C-terminal LF_N dimers (using R263C LF_N) and trimers [using LF_N-(Gly₄Ser)₃-Cys] were formed as follows. Protein was kept at 450 μM in 20 mM Tris (pH 8.5), 150 mM NaCl, and 500 μM tris-(2-carboxyethyl)phosphine. The pH of the solution was lowered to 7.5, and cross-linker (bismaleimidoethane for dimer formation and tris-[2-maleimidoethyl]amine for trimer formation [both from Pierce]) was added to protein at a 1:10 molar ratio. The reaction mixture was incubated at 4°C for 30 min. The addition of cross-linker to protein at a 1:10 molar ratio and incubation at 4°C for 30 min were repeated five times. Finally, the molar ratio of cross-linker to protein was raised to 1:1, and the reaction mixture was incubated overnight at 4°C. Dimeric and trimeric proteins were separated from monomeric protein using size exclusion chromatography over an S-200 column (Amersham Pharmacia).

Planar lipid bilayer translocation assay. Translocation of LF_N variants across black lipid membranes was measured as previously described (28). Briefly, a lipid bilayer of diphytanoylphosphatidylcholine was formed across a 0.2-mm-diameter

hole of a Delrin cup. The membrane separated two compartments containing 10 mM MES (morpholineethanesulfonic acid), 10 mM KH₂PO₄, 10 mM sodium oxalate, 100 mM KCl, and 1 mM EDTA (pH 5.5). Once membranes were formed, heptameric PA was added to the *cis* compartment, and a steady-state current level was reached (at a transmembrane voltage of +20 mV). LF_N was added to the *cis* compartment to a final concentration of 10 nM, and blockage of PA channels was measured by monitoring the decrease in current. The *cis* compartment was slowly perfused with a two-syringe system to remove unbound LF_N (Hamilton). The fraction of channels blocked by LF_N at any given time was calculated using the equation $F_b = I_t/I_o$, where F_b is the fraction of channels blocked, I_t is the current at any given time, and I_o is the steady-state current of open channels. Data were plotted either as the fraction of channels unblocked as a function of time or as the maximum percentage of channels blocked.

Translocation was initiated by increasing the voltage to +50 mV and was monitored by current increase across the membrane. The fraction of channels unblocked at any given time was calculated using the equation $F_u = (I_t + I_c)/(I_o + I_c)$, where F_u is the fraction of unblocked channels, I_t is the current at any given time, I_c is the current of blocked channels after addition of LF_N, and I_o is the steady-state current of open channels. Data were plotted either as the fraction of channels unblocked as a function of time or as the maximum percentage of channels unblocked.

Cell culture. CHO-K1 cells (ATCC CCL-61) were grown in Ham's F-12 medium supplemented with 10% fetal calf serum, 500 units/ml penicillin G, and 500 units/ml streptomycin sulfate. Cells were maintained as monolayers and grown in a humidified atmosphere at 5% CO₂.

Inhibition of protein synthesis. A protein synthesis assay was used to measure the ability of LF_N variants to inhibit the translocation of LF_N-DTA into cells (16). CHO-K1 cells (5×10^4 cells) were incubated with 100 pM PA83, 100 pM LF_N-DTA, and various concentrations of either monomeric or multimeric LF_N for 4 h at 37°C. Cells were then incubated with leucine-deficient medium supplemented with 1 μCi of [³H]leucine/ml for 1 h at 37°C and washed with phosphate-buffered saline. Cells were incubated twice for 10 min each with cold 5% trichloroacetic acid. The protein precipitate was solubilized in 100 μl 0.2 M KOH and neutralized with an equal volume of 0.1 M HCl. Tritiated protein was determined by scintillation counting. Fifty percent inhibitory concentrations (IC₅₀s) were calculated by fitting data to the equation $y = a + (b - a)/[1 + (x/c)^d]$, where a is the y minimum, b is the y maximum, c is the IC₅₀, and d is the slope.

Zebrafish studies. All zebrafish protocols were approved by the Institutional Animal Care and Use Committee of Children's Hospital, Boston. Breeding fish were maintained at 28.5°C on a 14-h light/10-h dark cycle. Embryos were collected by natural spawning, and raised in 10% Hanks saline at 32°C. Wild-type fish of the AB strain or a transgenic zebrafish line in which endothelial cells were labeled with enhanced green fluorescent protein (EGFP) [*Tg(fli1:EGFP)^{z1}*] (11) were used in these experiments. Microinjections were carried out as described by Weinstein et al. (25), with modifications described by Chan and Serluca (4). Combinations of PA, LF_N-DTA, LF_N, N-terminal LF_N dimer, and N-terminal LF_N trimer were prepared before injection. Phenol red (0.05%) was added under

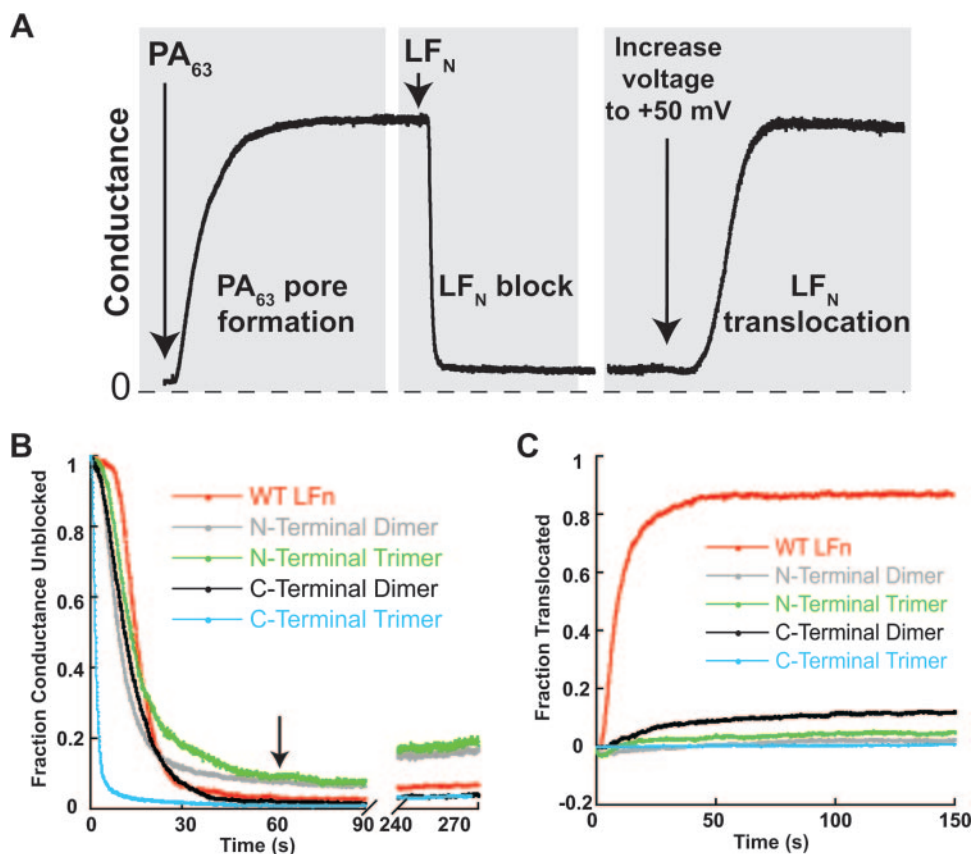


FIG. 2. Cross-linked LF_N cannot translocate across artificial membranes. (A) Schematic representation of translocation as assessed in planar lipid bilayers. Binding and translocation of LF_N were monitored by conductance. Pores initially form in membranes after addition of heptameric PA₆₃. Addition of LF_N rapidly lowers conductance, mimicking closed channels. Translocation, which is initiated by increasing the voltage to +50 mV, promotes passage of LF_N to the *trans* side of the membrane. (Modified from reference 13 with permission of the publisher.) (B) PA pores were formed in membranes at a transmembrane voltage of +20 mV and incubated in the presence of 10 nM monomeric or multimeric LF_N constructs. Ion conductance was measured during LF_N incubation and perfusion of the *cis* chamber. (C) PA channels blocked with monomeric or multimeric LF_N were subjected to an increase in transmembrane voltage to +50 mV, and ion conductance was monitored until a steady state was reached. WT, wild type.

each condition for visibility during injection, and samples were loaded into microinjection needles. Volumes of 30 nl or less were delivered into the anterior cardinal veins of embryos anesthetized with tricaine (Sigma) at 48 h postfertilization, resulting in injection of ~50 fmol PA, 11.5 fmol LF_N-DTA, and various amounts of inhibitor ($n = 20$ per treatment condition). After injection, embryos were transferred into fresh medium for recovery. Subjects were scored for attenuation of toxin effects under a dissecting scope at 20 h postinjection (hpi) to allow visualization of EGFP-labeled vasculature. Statistical analysis was conducted with Student's *t* test using SigmaStat 3.0 software.

RESULTS

Synthesis and characterization of LF_N multimers. Multimeric forms of LF_N were constructed by introducing Cys residues at either the N terminus (A1C) or the C terminus [R263C, LF_N-(Gly₄Ser)₃-Cys] of LF_N (Fig. 1A). Dimers cross-linked at the N or C terminus were prepared by incubating the A1C and R263C preparation, respectively, with the cysteine-reactive bifunctional cross-linker bismaleimidoethane. The corresponding trimers were generated by incubating the A1C and LF_N-(Gly₄Ser)₃-Cys preparations with the trifunctional cross-linker tris-[2-maleimidoethyl]amine. Both cross-linking reagents react with thiols to form nonreducible thioether link-

ages. We were successful in obtaining C-terminal trimers with the LF_N-(Gly₄Ser)₃-Cys preparation (Fig. 1B) but, for unknown reasons, not with R263C.

The multimers were purified by size exclusion chromatography and tested for the ability to undergo PA-dependent translocation across planar phospholipid bilayers, according to the protocol illustrated in Fig. 2A (28). PA₆₃ prepore was added to one compartment (defined as the *cis* compartment) of a planar lipid bilayer apparatus containing 100 mM KCl buffered at pH 5.5, and pore formation was monitored at a transmembrane voltage of +20 mV. After the conductance plateaued, the *cis* compartment was perfused to remove residual PA, and LF_N was added, causing rapid blockage of current as the protein bound to the pore. Following a second perfusion step to remove residual free LF_N, we increased the voltage to +50 mV to induce translocation of bound LF_N to the *trans* compartment. Translocation was manifested by the increase in conductance as LF_N exited the pore into the *trans* compartment.

As shown in Fig. 2B, both dimeric and trimeric forms of LF_N were effective conductance blockers, demonstrating that they retained the ability to bind to the LF/EF sites of the pore.

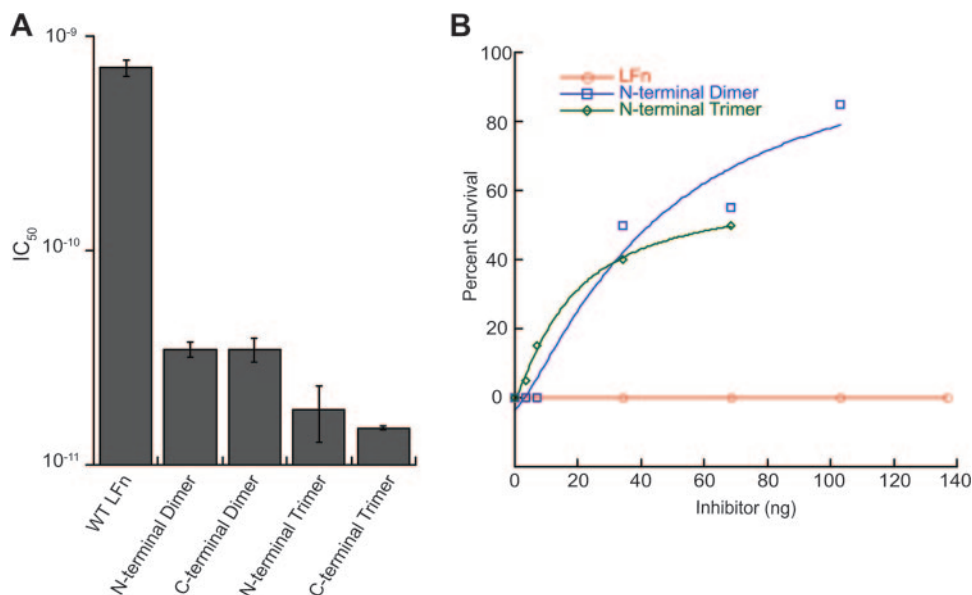


FIG. 3. Multimeric LF_N is a potent inhibitor of toxin action in cells and in zebrafish embryos. (A) CHO-K1 cells were incubated in the presence of PA, LF_N-DTA, and various concentrations of inhibitor. Toxin translocation was monitored through the incorporation of [³H]leucine into newly synthesized proteins, and IC₅₀s were calculated as described in Materials and Methods. (B) Zebrafish embryos were microinjected with PA, LF_N-DTA, and various concentrations of inhibitor ($n = 20$ for each data point). Protection as described in the text was scored as percent embryo survival. Inhibition experiments using trimer and dimer doses of 68.5 ng were repeated two times, in which significant increases in the survival of embryos treated with each multimer were observed (trimer, $P < 0.001$; dimer, $P < 0.001$). Statistics were completed using Student's t test. WT, wild type. Error bars indicate standard deviations.

Blockage by the monomer or the C-terminally cross-linked (C-linked) forms was close to 100%, whereas blockage by N-terminally cross-linked (N-linked) multimers was only ~90%. Also, blockage by the multimers was more rapid than that by the monomer, with the C-linked trimer showing remarkably fast kinetics. Two other C-linked trimers, prepared from LF_N-(Gly₄Ser)₂-Cys or LF_N-(Gly₄Ser)₄-Cys, were found to show the same fast kinetics of blockage as trimer prepared from LF_N-(Gly₄Ser)₃-Cys (data not shown). Finally, blockage by the N-linked forms was more readily lost during perfusion than that of the monomer or the C-linked forms (Fig. 2B, see 240- to 270-s time points).

Consistent with earlier findings, raising the voltage to +50 mV induced bound, monomeric LF_N to translocate through the pore (28). Translocation was rapid, with a half-time of several seconds, and conductance was restored to ~80% of the original value within a minute. However, none of the LF_N multimers was able to translocate to a significant degree, regardless of the site of cross-linking (Fig. 2C). Furthermore, translocation did not occur even at a high transmembrane voltage (+100 mV) or when we applied a transmembrane proton gradient by raising the pH of the *trans* compartment to neutrality (data not shown).

Multimeric forms of LF_N are potent inhibitors of toxin action on cells. The fact that multimeric forms of LF_N were able to bind to the PA pore but did not translocate suggested that they might function as potent inhibitors of toxin action. Such forms would be expected to bind to LF/EF binding sites on the PA heptamer more strongly than monomeric LF_N and could potentially form dead-end translocation complexes with the pore, effectively inactivating it.

We therefore measured the ability of the multimers to inhibit the PA-dependent action of LF_N-DTA on CHO-K1 cells. LF_N-DTA is translocated in a manner analogous to that of LF and EF (16), and, once in the cytosol, it catalyzes the ADP-ribosylation of EF-2, causing inhibition of protein synthesis. The inhibition of incorporation of radiolabeled leucine into newly synthesized proteins may therefore be used as an index of toxin action. As illustrated in Fig. 3A, the LF_N multimers gave IC₅₀s of between 15 and 35 pM, representing a 20- to 50-fold increase in inhibitory activity relative to monomeric LF_N (IC₅₀, ~700 pM). The two trimers were more potent than the dimers.

Multimers inhibit toxin action in zebrafish. To corroborate and extend the results from cell culture, we tested the ability of multimeric LF_N to inhibit LF_N-DTA-induced toxicity in a zebrafish embryonic vascular model. Coinjection of LF_N-DTA and PA induced rapid cell death in zebrafish embryos as early as 2 hpi, with tissues becoming necrotic by 8 hpi. When injected singly, neither LF_N-DTA nor PA produced observable effects at the latest time point, 20 hpi. Death (0% survival) was scored as severe necrosis and a complete lack of circulation at 20 hpi. Survival was defined as minimal to no tissue necrosis and retention of aortic blood flow. By these criteria, multimeric LF_N protected against LF_N-DTA effects, whereas monomeric LF_N did not (Fig. 3B).

Since the vasculature is directly exposed to injected toxin, we examined toxin effects on the endothelium using a transgenic zebrafish line in which endothelial cells expressed EGFP under the control of a *fli-1* promoter [*Tg(fli1:EGFP)^{z1}*]. Various concentrations of monomeric or multimeric LF_N were combined with LF_N-DTA and PA and

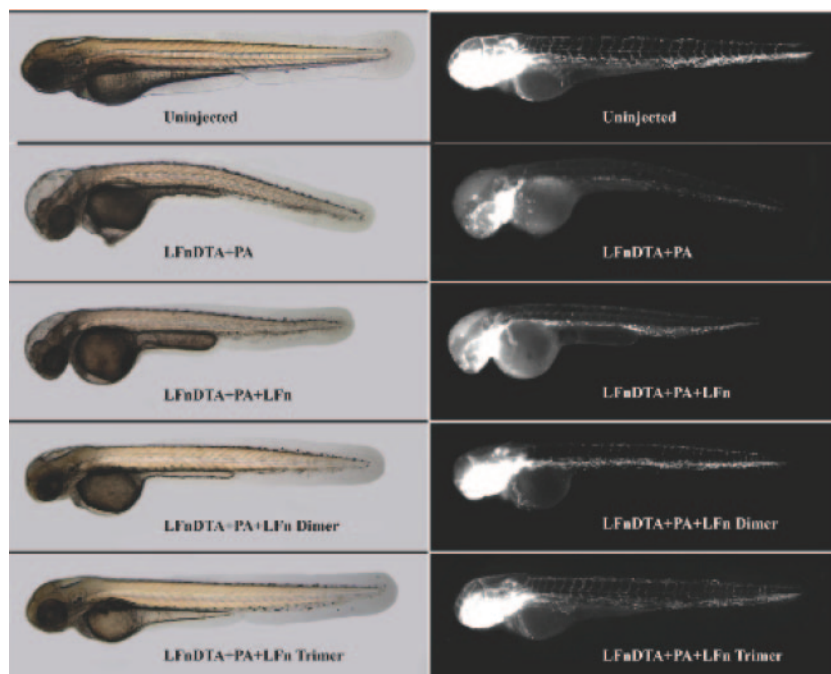


FIG. 4. Multimeric LF_N protects against LF_N -DTA effects on zebrafish vasculature. Zebrafish embryos expressing EGFP under the control of a *fli-1* promoter [$Tg(fli:EGFP)^{f1}$] (11) were injected with PA, LF_N -DTA, and/or 68.5 ng of inhibitor as indicated (representative embryos are depicted; $n = 20$). Photographs of the same embryos were taken for each pair of phase-contrast (left panels) and fluorescence (right panels) micrographs. Loss of fluorescence in LF_N -DTA- and PA-injected panels indicates endothelial cell death.

tested in comparison with controls. At 20 hpi, phenotypic changes were scored by overall morphology using phase-contrast microscopy, and changes in endothelial cells were scored under fluorescence (Fig. 4). A titratable increase in survival was observed in the presence of multimeric LF_N but not monomeric LF_N (Fig. 3B). Indeed, in the presence of multimeric LF_N , the vasculature of toxin-injected embryos closely resembled that of uninjected control fish, especially in the presence of the LF_N trimer (Fig. 4). Thus, multimerization of LF_N increased its potency as a toxin inhibitor, not only in cell culture but also in a vertebrate animal model.

DISCUSSION

Our findings show that dimers and trimers of LF_N formed by chemically cross-linking the N or C terminus do not translocate and, as a result, are potent blockers of toxin action. All of the multimers retained the ability to bind to the PA63 pore in planar phospholipid bilayers and to block ion conductance by the pore. Multimers cross-linked at the N terminus were less effective in blocking conductance (~90% inhibition) than those cross-linked at the C terminus (~98% inhibition). The C-linked multimers had free, unconstrained N termini that were apparently able to enter the pore and interact with the Phe clamp to form a tight seal against ion passage, as shown earlier with native LF_N (8). Cross-linking of the N termini must prevent formation of this seal or make the seal less tight.

The effect of cross-linking of LF_N on its affinity for the pore could not be easily predicted, because of potentially opposing effects of increased valence, which should strengthen the in-

teraction, and structural perturbation, which would most likely weaken it. We did not measure affinity directly, but the demonstration that conductance blockage by the multimers was complete within a minute and was not rapidly relieved upon perfusion indicates that cross-linking did not cause a major decrease in affinity. Conductance blockage was relieved during perfusion to a greater degree by the N-linked multimers than by the C-linked forms, suggesting that altering the interaction of the N termini with the Phe clamp diminishes affinity. These data support the notion that interaction of the N terminus of LF_N with the Phe clamp increases affinity of the protein for the pore.

Despite the fact that all of the multimers were able to bind to the pore and block its conductance, none of them translocated across the membrane under conditions known to promote translocation of monomers. There are at least two possible explanations for the inability to translocate. (i) Translocation might be restricted by the pore's physical dimensions, most likely its diameter at the narrowest point. There is clear evidence that the tertiary structure of a protein must be disrupted for translocation to occur (26). Consistent with this, the lumen of the transmembrane 14-strand β barrel of the pore has been estimated to be on the order of 15 Å, enabling it to accommodate a single α helix or an extended polypeptide but not a folded domain (7). Even narrower points in the translocation pathway may exist elsewhere in the pore, perhaps at the Phe clamp. Whether two or more extended polypeptides could be accommodated in the pore's β barrel or by the Phe clamp is difficult to predict. (ii) Translocation might be restricted by the pore's inability to pass certain chemical configurations generated by cross-

linking. Little is known about the ability of the pore to pass nonpeptidic chemical structures, and it is possible that it might not accommodate chemical configurations generated by reaction of the cross-linking agents with the Cys residues introduced into LF_N.

The mechanism by which translocation is blocked for the N-linked multimers may differ from that for the C-linked multimers, and indeed, more than one mechanism may contribute to the inactivity of either class of multimers. Our finding that N-linked forms of LF_N apparently did not interact with the Phe clamp to form a tight seal against ion flow suggests that the force-generating proton gradient across the membrane may be dissipated locally. Because the N termini of the C-linked multimers are unencumbered, one can envision that the N terminus of an LF_N domain would enter the pore and interact with the Phe clamp to form a tight seal to initiate threading of that domain through the pore. Complete translocation could be limited by inability of the Phe clamp or the transmembrane β barrel to accommodate the cross-linked intersection with the partner chain(s) or to translocate the partner chain in a C- to N-terminal direction, leading ultimately to a dead-end complex. Thus, our findings are consistent with the hypothesis that the PA₆₃ pore can translocate no more than a single polypeptide at once, but why cross-linked forms of LF_N do not translocate remains uncertain.

The ability of LF_N multimers to bind to the pore strongly but not translocate through it makes them potent inhibitors of toxin action. The dimers were \sim 20-fold more potent inhibitors than monomeric LF_N in a mammalian cell culture-based assay, and the trimers were even more potent, with the C-linked trimer showing the highest potency (\sim 50-fold greater than that of the monomer). The higher activity of the trimers may be related to greater avidity for the pore, based on valence. Two of the multimers were tested in a zebrafish model and shown to be potent inhibitors of the chimeric enzymatic effector LF_N-DTA when coinjected together with PA. Less than 100 ng of either a dimer or trimer raised zebrafish survival to $>$ 50% under conditions where 140 ng of monomeric LF_N caused no increase in survival. Also, the multimers blocked morphological changes induced in the zebrafish by the chimeric toxin.

Our data demonstrate successful protection against toxin action *in vivo* by multimeric LF_N. The need for an effective means to inhibit anthrax toxin action in unimmunized individuals infected with *B. anthracis* is well recognized, and our results offer a new way in which a potent anthrax toxin inhibitor can be generated by chemically modifying a component of the toxin. In addition, cross-linked forms of LF_N might also elicit toxin-neutralizing antibodies against LF, giving these forms dual protective functions.

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