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Intracellular Delivery of Functional Proteins with DNA–Protein Nanogels–Lipids Complex

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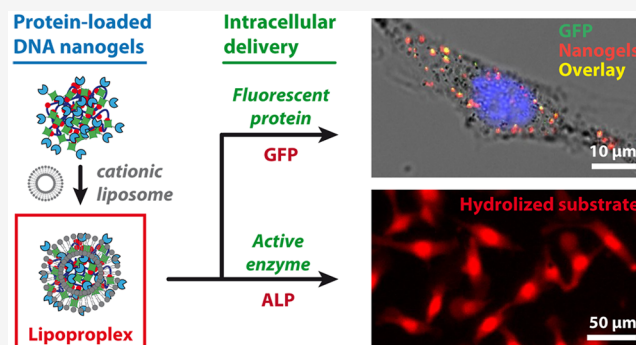
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ABSTRACT: Using functional proteins for therapeutic purposes due to their high selectivity and/or catalytic properties can enable the control of various cellular processes; however, the transport of active proteins inside living cells remains a major challenge. In contrast, intracellular delivery of nucleic acids has become a routine method for a number of applications in gene therapy, genome editing, or immunization. Here we report a functionalizable platform constituting of DNA–protein nanogel carriers cross-linked through streptavidin–biotin or streptactin–biotin interactions and demonstrate its applicability for intracellular delivery of active proteins. We show that the nanogels can be loaded with proteins bearing either biotin, streptavidin, or strep-tag, and the resulting functionalized nanogels can be delivered into living cells after complexation with cationic lipid vectors. We use this approach for delivery of alkaline phosphatase enzyme, which is shown to keep its catalytic activity after internalization by mouse melanoma B16 cells, as demonstrated by the DDAO–phosphate assay. The resulting functionalized nanogels have dimensions on the order of 100 nm, contain around 100 enzyme molecules, and are shown to be transfectable at low lipid concentrations (charge ratio $R_{\pm} = 0.75$). This ensures the low toxicity of our system, which in combination with high local enzyme concentration ($\sim 100 \mu\text{M}$) underlines potential interest of this nanoplatform for biomedical applications.



INTRODUCTION

Due to their highly specific biological activities and intrinsic biocompatibility, delivery of active proteins into cells represents great interest not only for fundamental research¹ and intracellular sensing,² but also for numerous therapeutic applications.^{3–5} However, despite their high potential, proteins suffer from chemical and thermal degradation, kidney clearance, and difficulty to penetrate through biological membranes.⁶ In particular, enzymes present a major limitation to be sensitive to both medium and the temperature, leading to the loss of their catalytic activity in inappropriate conditions thus limiting their applicability in both enzyme replacement therapy^{7,8} and enzyme prodrug therapy.^{9,10} Another aspect requiring consideration is the localization of proteins. Indeed, while most therapeutic proteins currently on the market have an extracellular localization, the intracellular transport of active proteins aiming cytosolic targets could strongly expand their exploitation for biomedical purposes.^{11,12} Yet the transport to the cytosol is still a major challenge, mainly due to physical impediment of the cell membrane and possible protein inactivation due to endocytosis.^{13–15}

Various vectorization systems have thus been proposed for protein protection and intracellular delivery, such as PEGyla-

tion, conjugation with cell-penetrating (CPP) or specifically engineered peptides, and formulations with liposomes or polymeric nanoparticles.^{5,16–18} DNA represents another promising protein carrier in nanomedicine,^{19,20} due to its intrinsic biocompatibility, programmability and transfectability. In fact, nucleic acids can be readily delivered into cells by using various nonviral cationic transfection agents, among which, cationic lipids represent the most involved vectors in clinical studies.²¹ In this view, recently, DNA has been bound to β -galactosidase and glucose oxidase in order to both protect and deliver them into the cells using lipofectamine as a transfection agent.²² In this case, individual enzyme molecules were covalently conjugated to several single-stranded DNA oligonucleotides. In another example, three luciferase molecules inside tubular DNA origami nanostructures were delivered into the cells via polyethylenimine transfection reagent.²³ A modular protein delivery system

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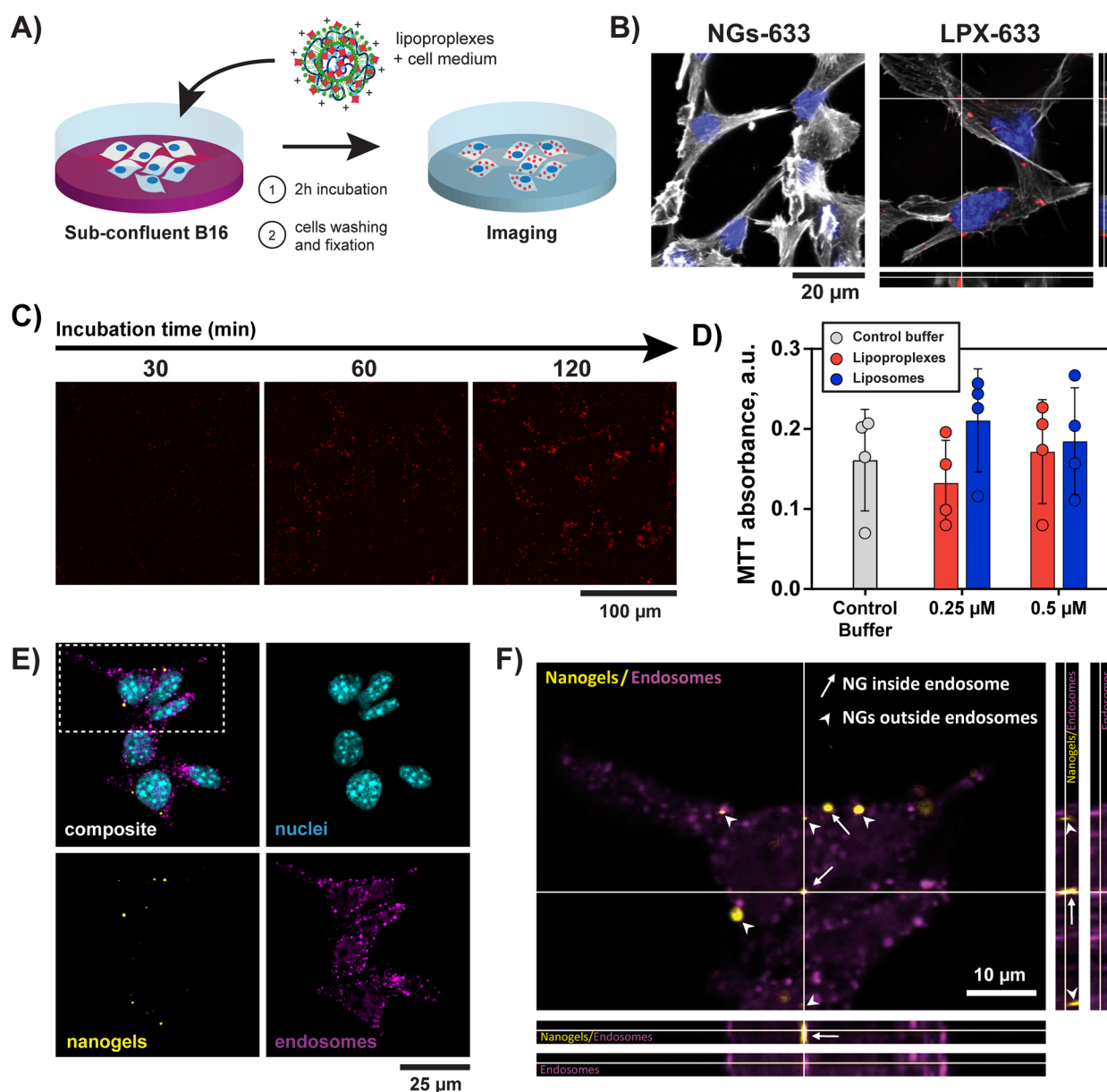


Figure 2. Internalization of DNA–protein nanogels mediated by cationic liposomes. (A) Scheme of experimental procedure for delivery of lipopropoxes into cells. (B) Confocal microscopy observation of B16 cells treated with AlexaFluor633-labeled nanogels in the absence (NG-633) and in the presence (LPX-633) of cationic lipids after 2h of incubation. Gray color corresponds to actin labeled with phalloidin ($\lambda_{\text{Ex}} = 488$ nm and $\lambda_{\text{Em}} = 495$ –550 nm), blue to the nucleus stained with DAPI ($\lambda_{\text{Ex}} = 405$ nm and $\lambda_{\text{Em}} = 415$ –450 nm), and red to nanogels containing AlexaFluor633-labeled streptavidin ($\lambda_{\text{Ex}} = 638$ nm and $\lambda_{\text{Em}} = 645$ –700 nm). Orthogonal views are given for the cells treated with LPX-633 (z -depth = 3 μm). (C) Widefield fluorescence microscopy images of B16 cells incubated with LPX-633 during different times ($\lambda_{\text{Ex}} = 578 \pm 11$ nm and $\lambda_{\text{Em}} = 641 \pm 38$ nm). (D) MTT cell viability assay. Formazan absorbance for B16 cells after 2 h of incubation with different concentrations of lipopropoxes or liposomes. Circles show data points, and bars show the mean values \pm sd. Four independent tests, statistic Kruskal–Wallis multiple comparison test: n.s. (E, F) Confocal microscopy analysis of B16 cell with EEA1 immunolabeling of endosomes after LPX-555 delivery. (E) Composite image and the separate channels of nanogels labeled with AlexaFluor555 (nanogels, colored in yellow $\lambda_{\text{Ex}} = 555$ nm and $\lambda_{\text{Em}} = 645$ –700 nm), EEA1 immunolabeling (endosomes, colored in magenta, $\lambda_{\text{Ex}} = 638$ nm and $\lambda_{\text{Em}} = 645$ –700 nm) and DAPI staining (nuclei, colored in blue, $\lambda_{\text{Ex}} = 405$ nm and $\lambda_{\text{Em}} = 415$ –450 nm). (F) Zoom on the selected region from (E) and orthogonal views along the white lines (z -depth = 2.29 μm). Arrows: nanogels colocalized with endosomes, arrowheads: nanogel not colocalized/close to endosomes.

than 1.5. We thus fixed the ratio $R_{\pm} = 1.5$ and analyzed the resulting LPX by flow cytometry, a method which enables codetection of different fluorophores within individual particles of nanometric size.³⁰ The resulting dot-plots for NGs-633 only, FITC-labeled liposomes only, and FITC-LPX-633 are given in the Figure 1B. For NGs and liposomes, the dot plots demonstrate a higher fluorescence signal for AlexaFluor633 (71.96%) and FITC (75.99%) signals, respectively. However,

when NGs and liposomes were mixed together, a large population of particles demonstrating the signal from both fluorophores (86.34%) was detected, indicating successful complexation. The observation that the mean FITC fluorescence in the LPX decreased, while the mean AlexaFluor633 fluorescence increased can be explained either by FRET energy transfer between the fluorophores, or by change of their fluorescent properties in lipidic microenvironment,^{31,32} both

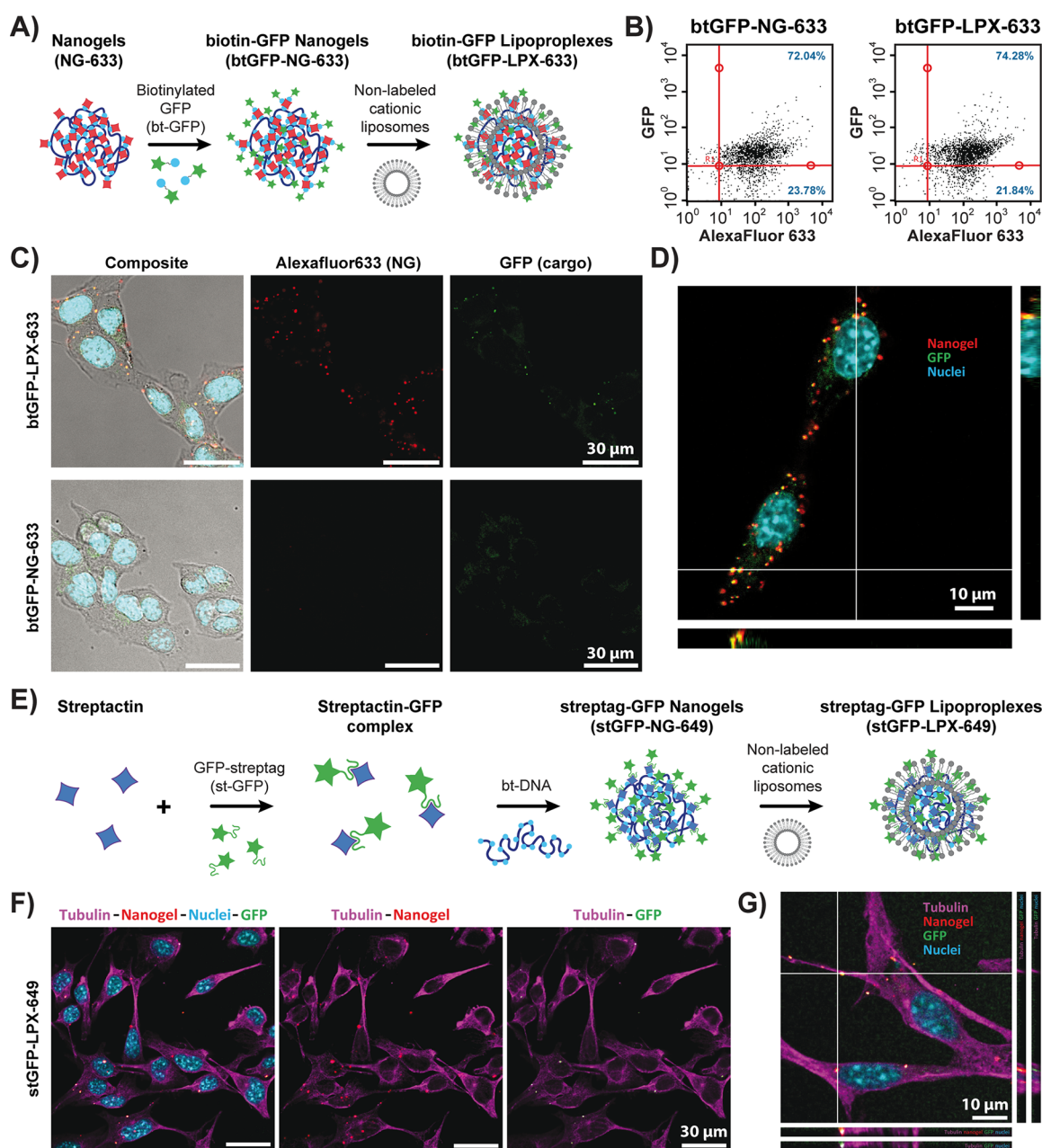


Figure 3. Delivery of lipoproplexes functionalized with GFP. (A) Scheme of postfunctionalization of NG-633 with biotinylated GFP (btGFP) and preparation of the corresponding lipoproplexes (btGFP-LPX-633). (B) Flow cytometry dot plots for btGFP-NG-633 (left) and btGFP-LPX-633 (right). The dot plots represent intensity of GFP fluorescence ($\lambda_{\text{Ex}} = 488 \text{ nm}$; $\lambda_{\text{Em}} = 525/30 \text{ nm}$) versus the fluorescence intensity of the NG-633 (AlexaFluor633, $\lambda_{\text{Ex}} = 640 \text{ nm}$; $\lambda_{\text{Em}} = 661/19 \text{ nm}$). Red lines separate the dot plots into 4 quadrants. Blue numbers give the percentage of observed events in the upper right and lower right quadrants. (C) Transmission and confocal microscopy images of B16 cells incubated either with btGFP-NG-633 (bottom) or with btGFP-LPX-633 (top). Cells are imaged in transmission light and overlaid with fluorescence signal from confocal observations. Green color corresponds to GFP cargo protein ($\lambda_{\text{Ex}} = 488 \text{ nm}$ and $\lambda_{\text{Em}} = 495\text{--}550 \text{ nm}$), cyan to the nucleus stained with DAPI ($\lambda_{\text{Ex}} = 405 \text{ nm}$ and $\lambda_{\text{Em}} = 415\text{--}450 \text{ nm}$), and red to nanogels containing AlexaFluor633-labeled streptavidin ($\lambda_{\text{Ex}} = 638 \text{ nm}$ and $\lambda_{\text{Em}} = 645\text{--}700 \text{ nm}$). (D) Higher magnification confocal microscopy observation of the sample from (C) with orthogonal views ($z\text{-depth} = 5.79 \mu\text{m}$). (E) Scheme of preparation of GFP-functionalized streptactin–DNA nanogels (stGFP-NG-649) based on streptag–streptactin binding and preparation of the corresponding lipoproplexes (stGFP-LPX-649). (F) Confocal microscopy observation of B16 cells incubated with stGFP-LPX-649. Green color corresponds to GFP cargo protein ($\lambda_{\text{Ex}} = 488 \text{ nm}$ and $\lambda_{\text{Em}} = 495\text{--}550 \text{ nm}$), cyan to the nucleus stained with DAPI ($\lambda_{\text{Ex}} = 405 \text{ nm}$ and $\lambda_{\text{Em}} = 415\text{--}450 \text{ nm}$), magenta to the immunolabeled tubulin with AlexaFluor555-labeled antibody ($\lambda_{\text{Ex}} = 552 \text{ nm}$ and $\lambda_{\text{Em}} = 584\text{--}613 \text{ nm}$), and red to nanogels containing DY649-labeled streptavidin ($\lambda_{\text{Ex}} = 638 \text{ nm}$ and $\lambda_{\text{Em}} = 645\text{--}700 \text{ nm}$). (G) Higher magnification confocal microscopy observation of the sample from (F) with orthogonal views ($z\text{-depth} = 2.65 \mu\text{m}$).

further confirming binding between DNA–protein nanogels and cationic lipids leading to the efficient formation of the lipoproplexes.

Delivery of Lipoproplexes into Cells. The possibility of delivery of the NGs-633 into living cells by their vectorization with cationic liposomes was then investigated with B16 mouse melanoma cells. The subconfluent cells were incubated with

LPX-633 (or control solution containing NGs-633 in absence of liposomes) in the presence of DMEM medium without serum for 2 h (Figure 2A), then fixed and stained for subsequent microscopy observations. No NGs were observed within the cells after treating them with NGs in the absence of liposomes (Figure 2B, left). In contrast, when the cells were treated with LPX-633, fluorescent signal of AlexaFluor633 was clearly observed as fluorescent spots within the cells. By performing confocal microscopy acquisition at different Z-planes, the presence of the NGs-633 in the same focal planes as cytoskeleton and nucleus was confirmed thus proving that the NGs could be internalized by the cells (Figure 2B, right). Liposomes were thus necessary to deliver the nanogels into the cells. These observations were also confirmed by wide field fluorescence microscopy observations (Figure S3), where all tested lipopropexes demonstrated corresponding fluorescence inside B16 cells, regardless of the fluorescent labeling. In contrast, only poor fluorescent signal was detected in B16 cells in the control samples where cells were treated with NG-633 in the absence of liposomes or with AlexaFluor633-streptavidin protein in the presence of liposomes. Contrary to several previous reports on intracellular delivery of DNA-based nanomaterials,^{33–35} we did not detect any internalization of our nanogels in cells in absence of the transfection agent. This difference in internalization could be due to a difference in rigidity between the highly structured tetrahedral DNA nanostructures having edges and corners³⁴ and our soft nanogels, as it has been demonstrated that the penetration through the membrane depends on the rigidity of the particle.³⁶ On the other hand, it has already been shown that the use of a transfection agent could increase the internalization of a DNA-based nanocages in cells.³⁵ Moreover, we propose that this agent can allow the nanogel to escape degradation in lysosomes. Indeed, Liang et al. showed that bare DNA tetrahedron, after endocytosis, ends up trapped in lysosomes,³⁴ whereas transfection agents, such as cationic lipids, are known to allow the exit of the endosomes.²⁸

The kinetics of lipopropex internalization by the cells was then analyzed. Lipopropex samples (or controls corresponding to NGs in absence of liposomes) were incubated with cells for 30, 60, or 120 min, before cell fixation and widefield fluorescence imaging (Figures 2C and Figure S4). We observed that nanogel internalization progressively increased upon incubation and reached a maximum after 2 h of incubation (Figure S5). The toxicity of the lipopropexes after internalization in B16 cells was then assessed with MTT assay (Figure 2D).³⁷ In this test, formazan absorbance at 570 nm is proportional to the number of viable cells, and no significant difference was observed between lipopropexes, cationic liposomes, and the control sample (buffer alone), indicating no toxicity for the B16 cells ($n = 4$, Kruskal-Wallis multiple comparison test). Moreover, the detection by confocal microscopy of the nanogels inside the cells 26 h after their internalization (Figure S6), further confirmed their biocompatibility and stability in the cellular microenvironment.

We then investigated the fate of the nanogels after their internalization by the cells. First, we studied colocalization of FITC-labeled lipids with NG-633 at different times after their internalization by the cells. We found that 30 min after delivery, fluorescent signal corresponding to FITC-lipid could be colocalized with the NG-633 fluorescent signal, suggesting that NG-633 and FITC-lipid were still associated. After longer period (5 and 26 h post incubation with FITC-LPX-633), only

fluorescent signal corresponding to NG-633 was observed inside cells (Figure S6). This suggests the dissociation of lipids from nanogels followed by the loss of fluorescent signal of FITC-liposome probably due to “dilution” of lipids in the cell lipidic environment following a probable lipid exchange occurring during FITC-LPX-633 cell internalization. Indeed, during internalization of nucleic acid vectorized by cationic liposomes, it is known that a lipid exchange occurs with endosomal membrane inducing disruption of endosomal membrane and delivery of nucleic acid in cytoplasm.²⁸ In order to provide evidence of endosomal escape of our DNA–protein NGs inside cells, we performed immunolabeling of endosomes and lysosomes after the delivery of lipopropexes. For this purpose, anti-EEA1 and anti-LAMP1 antibodies were used for the immunostaining of early endosomes and lysosomes, respectively. After 2 h of incubation of cells with fluorescent LPX (in this experiment, we used AlexaFluor555 streptavidin for nanogel formation, resulting in LPX-555 to allow multichannel observation of nanogels, cell nuclei and immunolabeling), we observed only rare events of colocalization of NG with EEA1 (Figure 2E and F). Interestingly, in most cases, the nanogels were found close to endosomal compartments. We thus hypothesize that this observation was performed after the endosomal escape. Concerning lysosomes, we observed no colocalization of NG with LAMP1 labeled compartments (Figure S7). The endosomal-lysosomal system is a dynamic process where intracellular membranous compartment interconvert from early endosomes, that mature to late endosomes and then fused into lysosomes.³⁸ As the lysosome appears to be the “last” step of this pathway, we can assume that if NGs are rarely observed inside early endosomes, it is expected that NGs inside lysosome may be more rare or even impossible to observe.

Intracellular Delivery of GFP-Loaded Lipopropexes.

We then explored the possibility of loading biotinylated cargo proteins into already formed NGs. Biotinylated green fluorescent protein (btGFP) was used as a model protein due to its fluorescent properties. The preparation of btGFP-loaded nanogels (btGFP-NGs) was conducted in two steps: (i) DNA–protein nanogels were first formed with AlexaFluor633-functionalized streptavidin in the conditions described above, and (ii) btGFP was then added and bound to remaining available biotin-binding streptavidin sites on the nanogels (Figure 3A). By AFM, the size of the resulting GFP-functionalized nanogels was measured to be 130 ± 51 nm (Figure S1B). The NGs were then characterized by flow cytometry (Figures 3B and Figure S8A). Colocalization of the btGFP with AlexaFluor633 fluorescence in the NGs demonstrates successful formation of GFP-functionalized NGs (btGFP-NGs-633), as 72.04% of fluorescent events were observed in the upper-right quadrant of the dot plot. The NGs were then complexed with nonfluorescent cationic liposomes in order to form GFP-functionalized lipopropexes, btGFP-LPX-633. Flow cytometry (Figure 3B right and Figure S8B) demonstrated that the formation of the lipopropexes did not interfere with GFP functionality, since the signal of functional GFP protein was still codetected with AlexaFluor633 from the nanogels after adding the liposomes, and the percentage of colocalized fluorescence events in the upper-right quadrant of the dot plot stayed quite unaffected (74.28%).

B16 cells were then incubated with the btGFP-LPX-633 and observed by confocal microscopy. Figure 3C shows the presence of fluorescent nanogels in the cells only when they have been complexed with the liposomes. Even though the fluorescence

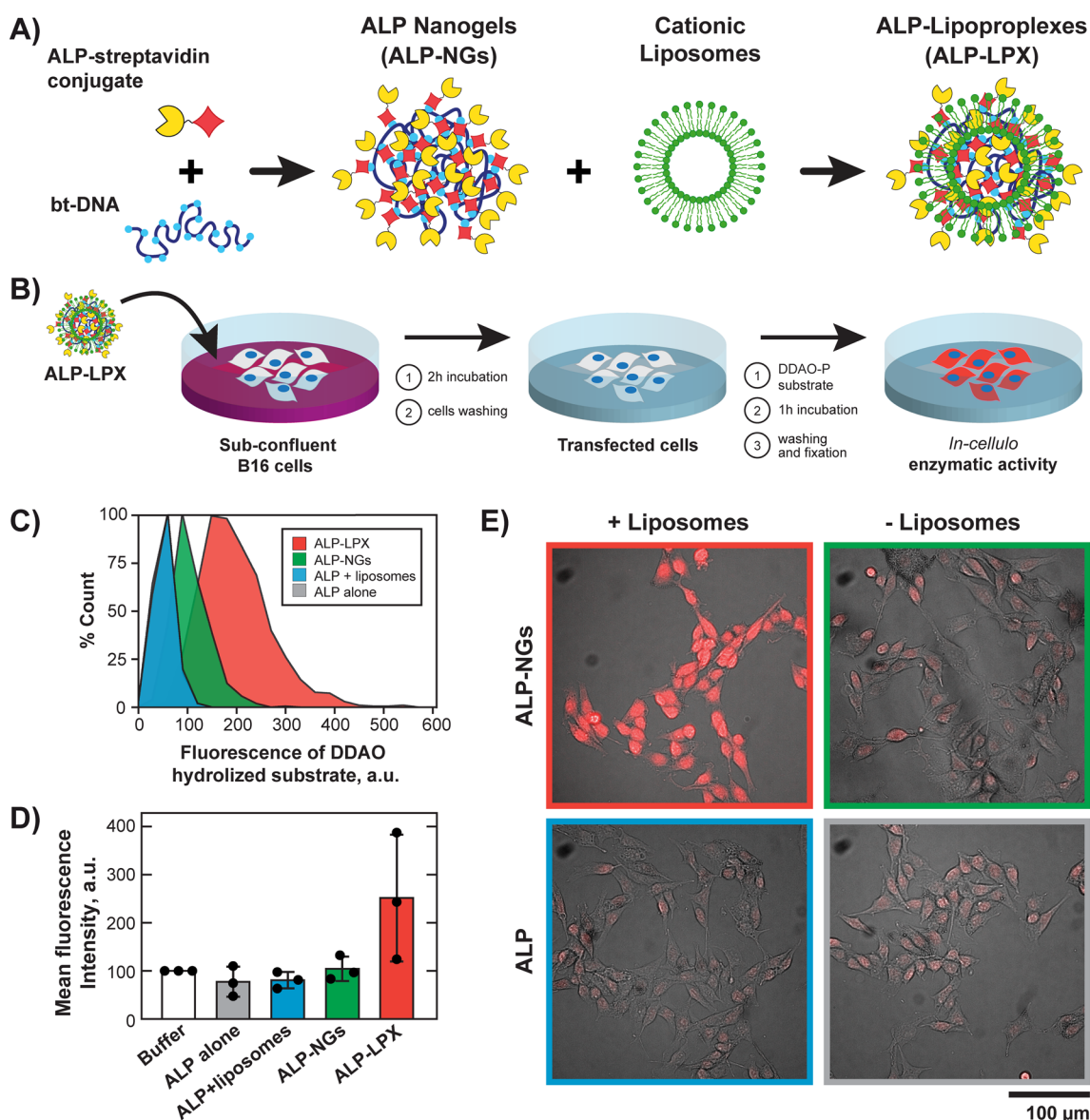


Figure 4. Delivery of ALP-functionalized lipoproplexes and enzymatic activity assay in the cells. (A) Preparation of ALP-loaded lipoproplexes (ALP-LPX). (B) Schematic representation of the experimental procedure for ALP delivery and measurement of its activity inside the cells. (C) Flow cytometry distribution of hydrolyzed DDAO substrate inside B16 cells treated with ALP-LPX and in control samples ($\lambda_{\text{Ex}} = 640 \text{ nm}$; $\lambda_{\text{Em}} = 661/19 \text{ nm}$). The distribution for ALP alone control is not visible, as it has practically the same shape as the one for ALP+liposomes control. (D) Normalized intracellular fluorescence intensities of hydrolyzed DDAO (mean \pm s.d.) detected by fluorescence microscopy. White bar shows intrinsic ALP activity (normalized to 100%) for cells incubated with the phosphate buffer (in the absence of ALP). Black circles show the data points. ($n = 3$ independent experiments, Kruskal–Wallis Test and Dunn’s multiple comparison test, $p = 0.0412$ ALP-LPX vs ALP+liposomes and $p = 0.0689$ for ALP-LPX vs ALP alone). (E) Example of combined transmission and fluorescence microscopy images of hydrolyzed DDAO substrate after treatment of B16 cells with enzyme alone (ALP) or enzyme loaded into DNA nanogels (ALP-NGs) in the presence and absence of liposomes.

signal of the GFP was weaker, at least $76 \pm 8\%$ (for over 400 events) of the nanogels demonstrated colocalization of the AlexaFluor633 red fluorescent signal from streptavidin with the green fluorescence from the GFP (see example of composite image and orthogonal views in Figure 3D), in agreement with the flow cytometry analysis. We typically observed 7.7 ± 4.5 internalized particles per cell (for 140 cells) in a single confocal plane, and around 50 internalized nanogels per cell have been observed in one particular z-stack of 7 confocal planes shown in the Figure S9. Biotinylated GFP protein can thus be successfully delivered into the cells within DNA-streptavidin complexed with cationic liposome vector.

To extend the applicability of our method, we devised another functionalization approach by using GFP fused to the twin-streptag oligopeptide sequence (stGFP).^{39,40} Twin-streptag is a 28 amino acid tag having 2 binding sequences for the biotin-binding pocket of streptavidin, with a strong affinity (in the range of pM) for the engineered variant of streptavidin, streptactin. We first found that streptactin labeled with DY649 dye was also able to cross-link the bt-DNA, but at higher protein concentrations (above 300 nM) than with the streptavidin–biotin system (Figure S10A). To functionalize the nanogels, we first obtained a stoichiometric complex of stGFP with streptactin and then used it for cross-linking of bt-DNA (Figure 3E and Figure S10B). We found that even more stGFP–streptactin

complex was necessary in this case to produce compact nanogels (around 1000 nM). We then formed the lipoproplexes with the stGFP-NG-649, incubated them with B16 cells, and imaged them by confocal microscopy (Figure 3F). Here again, fluorescent spots with colocalized fluorescence from streptactin (red) and GFP fluorescence (green) were found inside the cells when they were treated with stGFP-LPX-649 (Figure 3G). These results expand the functionalizability of DNA-streptavidin (or DNA-streptactin) nanogels and demonstrate that nanogel-based approach for intracellular protein delivery can also be applicable for cloned proteins bearing not biotin but specific oligopeptide tags.

Intracellular Delivery of Enzyme-Loaded Lipoproplexes. We finally investigated whether the lipoproplexes could be used as carriers for an active enzyme and if it would keep its catalytic activity after delivery into the cells. For this purpose, bt-DNA was directly cross-linked in one step with commercially available streptavidin conjugated to alkaline phosphatase (ALP) enzyme (Figure 4A)²⁶ to give functionalized nanogels (ALP-NGs). The formation of ALP-NGs was characterized by AFM imaging (Figures S11), and the concentration of streptavidin-ALP conjugate of 15 nM (for 1 μ M of bt-DNA in phosphate groups) was selected for further experiments, giving an apparent diameter of 133 ± 41 nm (Figure S1C). This ratio of streptavidin-ALP to bt-DNA corresponds to up to 100 enzymes per nanogel. The complexation of nonfluorescent ALP-NGs with FITC-labeled liposomes was then confirmed by flow cytometry (Figure S12). We observed that complexation with functionalized nanogels led to an increase of mean fluorescence signal distribution of liposomes from the charge ratio $R_{\pm} = 0.75$ (i.e., 0.5 μ M liposomes for 2 μ M bt-DNA). Such a low R_{\pm} ratio was thus sufficient for the formation of ALP-loaded lipoproplexes (ALP-LPX).

We then checked if the ALP enzyme kept its catalytic activity within the lipoproplexes *in vitro*. For the assay, 9H-(1,3-dichloro-9,9-dimethylacridin-2-one-7-yl) phosphate, diammonium salt (DDAO-P) fluorogenic substrate was used to follow the hydrolysis of its phosphate group by alkaline phosphatase via appearance of far-red fluorescence emission of the resulting DDAO. We found that incorporation of the ALP into the nanogels only slightly decreased the catalytic activity of the enzyme to 88% (Figure S13). Adding liposomes to the ALP alone did not affect its activity, whereas within ALP-loaded lipoproplexes, the enzyme conserved 79% of its initial catalytic activity, as compared to the activity of the native ALP-streptavidin conjugate. The ALP enzyme thus stayed available to the DDAO-P substrate and catalytically active even after loading into the DNA nanogels and their complexation with cationic lipids.

Finally, the delivery of ALP-functionalized lipoproplexes (ALP-LPX) into B16 cells was investigated (Figure 4B). The cells were treated with ALP-LPX for 2 h and washed to remove any remaining nanogels. The efficiency of delivery could not be assessed for the nonfluorescent NGs, instead we directly characterized the enzymatic activity of ALP inside the cells. For this purpose, DDAO-P substrate was added into the cell medium, and after 40 min of incubation the substrate was removed from the medium, cells were rinsed, and intracellular fluorescence of the hydrolyzed DDAO was analyzed either by flow cytometry after detaching (Trypsin-EDTA) and fixing the cells (Figure 4C), or by fluorescence microscopy after rinsing and fixing the cells onto culture plate (Figure 4D, E and Figures S14 and S15). We found that even in absence of added ALP, the

B16 cells were able to slightly hydrolyze the DDAO-P substrate giving a basal level of alkaline phosphatase activity.⁴¹ Flow cytometry showed no effect of adding enzyme alone and enzyme in the presence of liposomes on the basal fluorescence of the cells. The distribution of fluorescence of hydrolyzed substrate slightly increased when cells were incubated with ALP-functionalized nanogels (ALP-NG) without liposomes, but the increase in fluorescence was much stronger when the cells were incubated with ALP-LPX (Figure 4E). Figure S15 shows fluorescence distributions for 3 independent experiments, and Figure 4D gives the average fluorescence value of the triplicate ALP delivery experiments normalized by the basal level of ALP activity inside B16 cells treated with phosphate buffer only. We systematically observed that fluorescence intensity corresponding to the hydrolyzed substrate inside the cells treated with ALP-LPX, was higher than for the controls containing phosphate buffer only, free enzyme (free ALP), ALP with liposomes or ALP-functionalized nanogels without liposomes, with significant difference for ALP-LPX vs ALP+liposomes and ALP-LPX vs ALP alone (Kruskal–Wallis Test and Dunn’s multiple comparison test). In an additional control, we compared ALP enzymatic activity inside cells incubated with functionalized and nonfunctionalized LPX. Figure S16 demonstrates that no enhancement of ALP enzymatic activity after delivery of the enzyme-free LPX was observed, indicating that the increased enzymatic activity in ALP-LPX was due to the catalytic activity of delivered enzyme and not a result of increasing the basal expression of alkaline phosphatase due to the interaction of lipoproplexes with cellular environment. All these results show that the ALP has been efficiently delivered inside the cells in its active state via functionalized lipoproplexes. To the best of our knowledge, it is a first example of efficient delivery of a large local number of active enzymes.

DISCUSSION

We reported a new vector-carrier-cargo platform called lipoproplex, schematically represented in Figure S17, and demonstrated its ability to transport active proteins through the cell membrane. For this purpose, we used DNA nanogels cross-linked via streptavidin–biotin or streptactin–biotin interactions. This platform enables loading of the protein cargo into DNA carrier in several ways (i.e., using streptavidinated, biotinylated or strep-tagged cargo proteins) thus opening the possibility of large number of functionalities that can be potentially introduced into a single nanogel. We used cationic liposomes for the vectorization of DNA–protein NGs and named the resulting complex as “lipoproplex”, which evokes the complex between DNA and liposomes (i.e., lipoplex) but is intended to emphasize the importance of the protein component in our DNA–protein nanogels. Three-dimensional structure of the nanogels demands lower concentration of cationic liposomes for complexation (down to $R_{\pm} = 0.75$), which in combination with intrinsic biocompatibility of DNA nanoscaffold results in nontoxic protein delivery system as demonstrated by the MTT assay. For a proof of concept, here we investigated delivery of two proteins: biotinylated (or strep-tagged) fluorescent protein GFP and the enzyme alkaline phosphatase conjugated to streptavidin. In both cases, the complexation with cationic liposomes and subsequent internalization by the cells did not affect the respective properties of the proteins: GFP retained its fluorescence emission, whereas the ALP kept its efficiency to hydrolyze the DDAO-P substrate.

Both strategies thus proved to be adapted for the nanogels functionalization.

By considering that all streptavidin-ALP conjugates enter the nanogels and each NG consists of one DNA scaffold molecule, under our conditions (15 nM of strep-ALP for 1 μ M of bt-DNA in phosphates) each nanogel can contain up to 100 loaded enzyme molecules. By assuming the nanogels to have a spherical morphology and 130 nm diameter, the local concentration of the loaded protein can be evaluated to around 130 μ M. High number of potentially loadable proteins can be useful for delivery into cells of multifunctional nanogels containing a large number of different proteins, and the high local concentration of the loaded proteins is promising for biomedical applications. This approach can thus be of particular interest for codelivery of several active enzymes at controlled stoichiometry. Cotransfection of genes coding for different enzymes was suggested for cancer therapy^{42,43} and treatment of metabolic diseases.⁴⁴ But it was shown in some cases to result in the expression of only one or the other enzyme rather than the desired dual expression of both enzymes,⁴⁵ because of the competition for the use of intracellular resources available for gene expression.⁴⁶ Direct delivery of NGs containing defined enzymatic composition avoids the coexpression step and can thus overcome this issue. Moreover, high local enzyme concentrations within the nanogels can be of particular interest for creating or restoring complex metabolic pathways requiring the close spatial positioning of several catalytic processes within an enzymatic cascade.

CONCLUSIONS

In conclusion, we present a new DNA–protein nanogel platform for the transport of active loaded proteins into living cells. The loaded proteins have been efficiently internalized by the cells while keeping their specific properties, which underlines potential therapeutic interest of lipopropoxes for delivery and codelivery into the cells of active enzymes. We anticipate that DNA–protein nanogels could be employed as nanoscaffold for the transport of other biological and chemical entities, not only by incorporating them into the nanogel by streptavidin/streptactin-biotin binding, but also by orthogonal binding into DNA double helix (i.e., intercalative drugs, such as doxorubicin).²⁴ Moreover, the applicability of our systems to biotinylated, streptavidinated, or strep-tagged recombinant proteins, nanoparticles, aptameric sequences, or antibodies opens up the possibility of using nanogels as universal scaffolds with a high degree of functionalizability that not only could encapsulate a number of desired features in a single nanosized material but also could protect the incorporated entities from the biological microenvironment. All of these results underline the great potential of using DNA transfection agents for soft DNA nanomaterials delivery.

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/jacs.3c08000>.

Materials and methods, AFM images and frequency size distribution for different nanogels (Figure S1); fluorescent microscopy analysis of lipopropox aggregation (Figure S2); wide-field microscopy observation of B16 cells treated with different lipopropoxes and controls (Figure S3); kinetics of lipopropoxes internalization and

its analysis (Figures S4–S5); confocal microscopy images of B16 cells at different times after incubation with FITC-LPX-633 (Figure S6); LAMP1 immunolabeling on B16 cells after LPX-633 incubation (Figure S7); flow cytometry analysis of GFP-functionalized nanogels and lipopropoxes (Figure S8); z-stack projection of confocal planes for B16 cells treated with GFP-functionalized lipopropoxes (Figure S9); AFM observation of DNA–streptactin nanogels and their functionalization with st-GFP (Figure S10), AFM observation of formation of nanogels functionalized with streptavidin-ALP conjugate (Figure S11); flow cytometry analysis of formation of ALP-functionalized lipopropoxes (Figure S12); effect of incorporation into nanogels and lipopropoxes on the acellular enzymatic activity of ALP (Figure S13); fluorescence microscopy detection of intracellular catalytic activity of ALP after intracellular delivery of ALP-LPX (Figure S14); box plot fluorescence distributions for 3 individual experiments of ALP-LPX delivery (Figure S15); comparison of the ALP enzymatic activity of B16 treated with ALP-functionalized and nonfunctionalized lipopropoxes (Figure S16); and schematic representation of lipopropox-based proteins delivery platform (Figure S17) (PDF)

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Notes

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