

Decoy exosomes as a novel biologic reagent to antagonize inflammation

This article was published in the following Dove Press journal:
International Journal of Nanomedicine

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Background: Exosomes are ubiquitous naturally secreted stable nanovesicles that can be engineered to target and deliver novel therapeutics to treat a host of human diseases.

Methods: We engineered the surfaces of cell-derived nanovesicles to act as decoys in the treatment of inflammation by antagonizing the major proinflammatory cytokine, tumor necrosis factor alpha (TNF α).

Results: Decoy exosomes were generated by displaying the TNF α binding domain of human TNF receptor-1 (hTNFR1) on the outer surface of exosomes using stably transfected HEK293 cells. We developed an efficient method to purify the engineered exosomes from conditioned medium based on sequential centrifugation, ultrafiltration, and precipitation. We characterized decoy exosomes using immune-quantification, nanoparticle tracking analysis, and confocal microscopy to confirm that they retain the correct orientation, size, and shape of naturally produced exosomes. We demonstrated the engineered decoy exosomes specifically antagonize activities of TNF α using an inflammatory reporter cell line.

Conclusions: Decoy exosomes produced in human cells serve as a novel biologic reagent for antagonizing inflammatory signaling mediated by TNF α .

Keywords: Decoy exosomes, Inflammation, CD63, Receptor, TNF α , Drug delivery

Introduction

Inflammation is a hallmark of many debilitating and life-threatening disorders, including inflammatory bowel disease,¹ rheumatoid arthritis,² psoriasis,³ atherosclerosis,⁴ Alzheimer's,^{5,6} sepsis,⁵ and cancer.^{7,8} Many of these disorders are chronic and impose a significant burden to family and society due to their high morbidity and mortality, and therapies are costly, both financially and for patient care.^{9–11} In the United States and Canada alone, it is estimated that more than 1.4 million people suffer from inflammatory bowel disease.¹² While many genetic and environmental factors play a prominent role in disease progression,¹³ the proinflammatory cytokine, TNF α , is recognized as the central player in eliciting inflammatory cascades.^{12,14,15} Protein therapeutics, such as anti-TNF α antibodies that selectively block the TNF α cascade, have become the mainstay therapy for the management of chronic inflammatory diseases.^{16–18} Despite the promise of the early studies, concerns about side effects of these protein drugs, including induction of anti-antibodies and immuno-suppression, may limit the applications to disease treatments.^{19–22} Additionally, the efficacy of antibody-based therapies is limited by their rapid clearance and poor penetration of tissue,²³ especially when compared to other biologics, such as exosomes.

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Thus, there is a pressing demand to introduce improved therapeutic agents to treat inflammatory diseases; to this end, we introduce a new class of decoy exosomes that antagonize the major proinflammatory cytokine, TNF α .

Exosomes are cell-derived nanovesicles that are being engineered by some laboratories to function as disease cell-targeting drug delivery vehicles. Exosomes can be engineered to display specific proteins on their external membrane, for example to target specific receptors on a tumor cell, and to display other functional proteins in their lumen.^{24–26} Circulating in blood and other body fluids, including synovial joints, engineered exosomes could be used to deliver large amounts of therapeutic reagents to targeted tissues. The intrinsic biocompatibility and high degree of tissue-penetration have led numerous groups to pursue the engineering of exosomes as drug-delivery vehicles to treat diverse human diseases.²⁷ For example, exosomes have been engineered to deliver siRNAs to neurons, microglia, and oligodendrocytes in the brain to affect specific gene knockdowns.²⁸ Exosomes bearing targeting and therapeutic proteins circulate in the body much longer than synthetic vehicles, and are found to be well suited to deliver anti-oncogenic KRAS small RNAs to deep-seated pancreatic cancer cells.²⁹ However, we are unaware of robust and high-fidelity strategies to engineer exosomes with protein therapeutics to treat inflammation. Here we show that engineered exosomes with TNF receptor (TNFR1) can be used to antagonize the proinflammatory cytokine TNF α . We further show the establishment in a proof-of-concept study that demonstrates the effectiveness of exosomes in delivering the “decoy-binding domain” of the TNF-receptor to a cell model of inflammation.

A vital component of this study is the application of an exosome surface display technology developed by our group to load diverse therapeutics and protein reporters to the inside or outside of the exosome membrane surface via genetic fusions.³⁰ Individually, we fuse target proteins to the outer and inner surface of exosomes using the endogenous transmembrane protein CD63, an abundant tetraspanin protein expressed on the surface of exosomes.^{31,32} In this study, we used tetraspanin CD63 to engineer exosomes bearing a surface-exposed TNFR1. Since these novel exosomes function as agonists in TNF-signaling of inflammation responses in cells, we refer to them as decoy exosomes. Because each decoy exosome displays multiple copies of TNFR1 on its surface, it is proposed to function as a biological sponge that absorbs

soluble TNF α in the affected tissue and thereby compromises TNF-mediated inflammation in the exosome-targeted tissue.

The goals of this study are two-fold: 1) to establish human cell lines in permanent culture that are genetically modified with our tetraspanin CD63 surface-display system to produce decoy exosomes bearing genetically transformed TNF α receptors, and 2) to demonstrate these decoy exosomes antagonize TNF α in a cellular model of inflammation. To achieve these goals, we designed and constructed a chimeric gene composed of three geometrically defined protein domains: the transmembrane CD63; the decoy receptor hTNFR1-EC on the outer surface; and the green fluorescent protein (GFP) on the inner surface of the exosomes. GFP functions as a fluorescent reporter of the expression of the chimeric gene and moreover enables imaging of the distribution of decoy exosomes in cell culture or tissue. Using a combination of genetic and cellular methods and analyses, we showed the transfection of HEK293 cells with the engineered constructs produce a fusion protein on the surface of exosomes with correct geometry. Moreover, we used fluorescence microscopy to show HEK293 cells produce large numbers of decoy exosomes that are released to the conditioned medium. Finally, we developed a protocol to purify decoy exosomes from conditioned medium in high-yield and showed that decoy exosomes exhibit a specific anti-TNF α activity on an inflammatory cell reporter system established by our group.³³

Because of their intrinsically high tissue-penetrating ability, we hypothesize decoy exosomes are more effective in delivering therapeutic (anti-inflammatory) cargo to target cells compared to the injection of protein-based therapeutics. Additionally, our work provides a proof-of-concept to develop agonist-binding engineered exosomes to treat other diseases, including cancer, cardiovascular, and neuronal disorders.

Material and methods

Materials

We purchased human recombinant TNF α from R&D Systems (Minneapolis, MN) and puromycin hydrochloride from ThermoFisher Scientific (Waltham, MA). We purchased the luciferase assay substrate and cell lysis reagents from Promega (Madison, WI). We purchased the organelle staining reagent LysoTracker Red DND-99, fusion genes for tracking cellular organelles of early (Rab5a-RFP) and late

(Rab7a-RFP) endosomes from ThermoFisher Scientific. Finally, we purchased the exosome marker (XPACK-RFP) from System Biosciences (SBI, Palo Alto, CA).

Design and construction of expression vectors

We constructed the fusion protein of CD63-GFP as reported.³⁰ We inserted either TNFR1-EC or RFP at the 2nd extracellular loop of CD63 using a fusion technology.³⁰ We confirmed the fidelity of all constructs by double-stranded DNA sequencing. Details of the coding sequences of the decoy chimeric protein appear in the Additional file 1 (Supplementary sequences). We have previously reported on the RFP-control chimaera.³⁰

Cell culture

We purchased the human embryonic kidney cells (HEK293) from Alstem (Richmond, California). Cells were cultured in high glucose Dulbecco's Modified Eagle Medium (DMEM) with 2mM glutamine (Gibco; MA, USA) supplemented with 10% fetal bovine serum (GE Healthcare Life Sciences; Issaquah, WA) and 100 U/mL penicillin/streptomycin (Gibco; MA, USA) at 37°C in 5% CO₂.

Transfection, drug selection, and establishment of stable cell lines

Stable cell lines expressing various CD63 fusion proteins were established as previously reported.³⁰ Briefly, we transfected HEK293 with plasmid DNA (1~2.5 µg/well) using either Lipofectamine (ThermoFisher Scientific, Waltham, MA) or FuGENE6 transfection reagent (Promega, Madison, WI) for 24~48 hrs. Next, we transferred the transfected HEK293 cells to a drug selection medium containing 5 µg/mL puromycin. The resultant cell lines stably expressed the fusion gene and remained GFP/RFP-positive for at least 50 days (equivalent to 25 passages). We maintained these stably transfected cells in medium with puromycin although before an experiment, we bathed cells in a puromycin-free medium for at least 2 passages. Stable reporter cell lines that detect TNF α signaling were also similarly established as previously reported.³³

Fluorescence-activated cell sorting (FACS)

Cultured HEK293 cells stably expressing the CD63-RFP-GFP or CD63-TNFR1-GFP fusion proteins were analyzed using an Accuri C6 Plus Flow Cytometer (BD Biosciences, San Jose, CA), as reported.³⁴ In brief, cells grown at

80~90% confluency on 3 cm dishes were collected and sorted using both FL1 and FL3A channels to quantify the levels of GFP and RFP intensity – we recorded over 10,000 events for each sample.

Exosome preparation

We prepared exosomes from the conditioned medium as previously reported.³⁴ Briefly, first we transferred the stably transfected cells to serum-free UltraCulture medium (Lonza, Allendale, NJ), and then we collected the conditioned medium, centrifuged to remove cell debris, and ultra-filtered to remove large extracellular vesicles (>200 nm) using a 0.22 µm filter. We mixed the filtered medium containing mainly exosomes with exosome-precipitation solution ExoQuick-TC (SBI, Palo Alto, CA). We determined the protein content of the resuspended pellet using a NanoDrop Lite (ThermoFisher Scientific, Waltham, MA). We have submitted all relevant data of our experiments to the EV-TRACK knowledgebase (EV-TRACK ID: EV190008) with an EV-Track score of 38%.³⁵

Nanoparticle analysis and characterization

Exosomes isolated from engineered cells were subjected to nanoparticle tracking analysis (NTA) using a NanoSight LM10 instrument (Malvern Instruments Ltd, Malven, UK) with a 405 nm and 60 mV laser source. Typically, we used 1 ml of a diluted exosome preparation for laser light scattering study usually making 3 recordings of 60 seconds per sample. We processed the data using the NTA software to determine the size distribution of the exosomes.

Dot-blot immuno-analysis

We used a dot-blot immuno-analysis to identify target proteins using premade dot-blot from SBI (Palo Alto, CA). Each blot contained 12 pre-printed spots and featured 8 antibodies for exosomal markers including CD63, CD81, ALIX, FLOT1, ICAM1, EpCam, ANXAS, and TSG101 as well as cytosolic GM130 to identify any cellular contamination. We used ~300 µg of exosome proteins in each assay according to the immuno-binding and detection protocol in the user manual.

Fluorescence and confocal microscopy

Images of live cells or fluorescently labeled exosomes were recorded using an Olympus fluorescence microscope (Waltham, MA) or Leica TCS SP8 confocal microscope

(Buffalo Grove, IL). To show the intracellular localization of fusion proteins, we recorded both fluorescent and phase contrast images from the same field. Any image modifications including brightness and contrast were applied to the entire image using Image J software.

Luciferase assay

Reporter luciferase assays were conducted using a luciferase assay kit according to the manufacturer's instructions (Promega, Madison, WI). First, we lysed the cells and then recorded the luciferase activity with a microplate reader (BMG LABTECH, Cary, NC) as previously reported.³⁶

Cell models for testing the antagonist activity of the decoy exosomes

We developed two cell models to assay for the antagonist activity of the decoy exosomes: 1) a cell co-culture model to assess the effect of newly released exosomes and 2) an exosome treatment model to assess the effect of purified exosomes. In the cell co-culture experiments, we combined 2.5×10^3 of inflammation reporter cells with 7.5×10^3 of either HEK293 cells producing the decoy exosomes (treatment group), RFP-exosome-control (mock control), or the parental HEK293 cells (no treatment control) in a 96-well plate. After 24 hrs of co-culture, we seeded cells in a serum-free Ultra-Culture either in the presence or absence of 1 ng/mL TNF α . After another 24 hr-incubation period, we first lysed the cells and then conducted a luciferase assay on the lysate. For the decoy exosome experiments, we seeded 1×10^4 of inflammation reporter cells in a 96-well plate overnight. Cells were then switched to serum-free Ultra-Culture in groups with either decoy exosomes, control exosomes, or no exosomes in either the presence or absence of 1 ng/mL TNF α . Cells without TNF α treatment were also included in the study to account for background signals from the reporter cells. After 24 hrs of treatment, we lysed the reporter cells and conducted the luciferase assays on the lysates as previously reported.³³

Data collection and statistical analysis

We used the Student's paired, two-tailed *t*-test to determine statistical significance of our studies, with *P*-values <0.05

being considered significant. We expressed the values for all measurements as mean \pm standard error.

Results and discussion

Design and construction of a fusion gene encoding the decoy receptor on the outer surface of exosomes

Unlike other nanoscale carriers such as liposomes, biopolymers, and gold nanoparticles,^{37,38} we produced exosomes within living human cells that harbor a repertoire of endogenous membrane proteins on their surface, including CD63, a hallmark biomarker of exosomes.^{39,40} We elected to engineer CD63 as a molecular scaffold for our exosome surface display technology – we have previously shown this approach allows one to project specific target-recognizing peptides on the outer and inner surface of the exosome membrane.³⁰ We introduced the extracellular TNF α binding domain of human TNF α receptor 1 (hTNFR1-EC) at the 2nd loop of CD63 in order to project the TNF α receptor on the external face of the exosome. The choice of insertion site of TNFR1 in the 2nd loop but not the 1st loop is due to two major considerations. First, the chosen site is between Alanine 133 and Serine 134, which is both on the outer surface of exosomes and near the most exposed region of CD63.³⁰ Second, the 1st loop is relatively small and tends to be covered by the larger 2nd loop, hence TNFR1 inserted at this loop may become sterically inaccessible to its ligand.^{39,40} To track and image the distributions of fusion proteins in cells and exosomes, we fused GFP to the C-terminus of CD63, which is located on the inner surface of exosomes (Figure 1A, upper panel). We also constructed an RFP fusion protein (replacing hTNFR1-EC) to produce the control group of fluorescently visible but inactive exosomes (Figure 1A, lower panel). To facilitate the establishment of stable cell lines that secrete these engineered exosomes, we cloned the coding sequence of each fusion into a vector that contained two expression cassettes: the CMV-driven fusion gene and the EF1 α -driven puromycin resistance gene for drug selection (Figure 1A). According to this design, the engineered exosomes would display either the hTNFR1-ED (decoy exosome) or RFP (control exosome) at the 2nd loop on the outer surface along with GFP on the inner surface of modified exosomes (Figure 1B). Thus, decoy cells would produce decoy exosomes with hTNFR1-ED and GFP, while control cells would produce control exosomes with

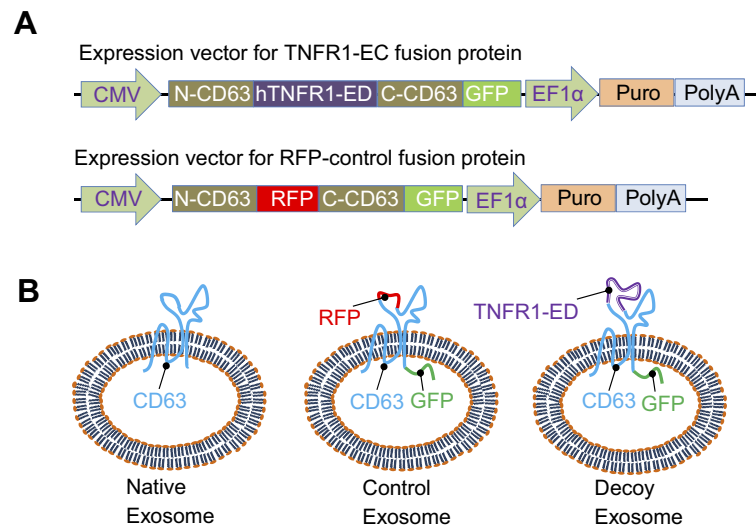


Figure 1 Surface display of extracellular domain (ED) on the outer surface of decoy exosome. **(A)** Schematic illustration of vectors expressing a decoy chimera containing the extracellular domain of human tumor necrosis factor receptor I (hTNFR1-ED, top) or a control RFP (bottom). **(B)** The strategy of presenting the decoy receptor on the outer surface of exosomes using a CD63 scaffold. CD63 (blue) anchors the membrane in an M-shaped topology with its two termini located within the lumen (native exosome, Left). The control exosomes have a C-terminal GFP with or without an RFP in the second loop of CD63 (RFP-control exosome, middle). The hTNFR1-ED (purple) is fused in the second loop on the outer surface of the exosome, while a reporter GFP (green) is attached to its C-terminus inside its lumen (decoy exosome, right). **Abbreviations:** CMV, cytomegalovirus promoter; N-CD63, N-terminus of CD63; hTNFR1-ED, the extracellular domain of the human TNFR1; C-CD63, C-terminus of CD63; EF-1 α , elongation factor-1 promoter; Puro, puromycin resistance gene; PolyA, polyadenylation signaling sequences.

no receptor but both RFP and GFP, whereas parental cells would neither express GFP nor RFP.

Establishment and characterization of stable cell lines that produce control or decoy exosomes

To permanently express the decoy proteins in human cells, we first established stable cell lines by transfecting HEK293 cells with our expression vector DNA followed by puromycin treatment for more than 40 days (Figure 2A). These stable cells consistently produce the decoy or control proteins, which may be integrated into pre-secreted exosomes in transfected cells.

Next, we characterized the expression level of each gene fusion in living HEK293 cells using FACS analysis. We analyzed the expression of CD63-RFP-GFP (control cells) or CD63-TNFR1-GFP (decoy cells) fusion proteins in stably transfected cultured cells using a BD Accuri C6 Plus Flow Cytometer. We also analyzed the control group, ie, the parental HEK cell in the same way. As expected, we recorded an increase in the intensity of GFP fluorescence in the decoy cells (Figure 2B, right), as compared to the parental control HEK cells (non-GFP, non-RFP) (Figure 2B, left). We also recorded significant increases in the intensities of RFP and GFP in mock engineered cells compared to the parental control HEK cells (Figure 2B, middle). These FACS results

showed we successfully generated stable cells expressing defined fusion genes. In the decoy group, ~64.7% of cells showed a dramatic increase in the level of GFP expression. In the RFP/GFP mock control group, ~49% of control cells expressed GFP and RFP. In contrast, almost all parental HEK293 cells were GFP- and RFP-negative. However, it is noteworthy that the distribution of fluorescence intensity was broad in the stably transfected cells (decoy and control), a common finding in studies that use the transfection protocol described above.

To establish whether the expressed fusion proteins were positioned on the exosomal membrane by CD63, we recorded fluorescent images of GFP and RFP in transfected cells. Confocal images revealed the distributions of both RFP (red) and GFP (green) in the cytosol of control cells identified as nodules (Figure 2C), and furthermore, we found significant co-localization of the emission signals, as evidenced by the strong yellow (merged) color in the overlay image (Figure 2C). In contrast, we found GFP positive nodules alone in the cytosol of decoy cells (Figure 2D), consistent with the finding of the FACS analysis detailed above (Figure 2B).

Exosomes are first generated within the endocytic compartments and then released to the extracellular environment. We conducted a series of co-localization studies to confirm our premise whether the punctuated nodule

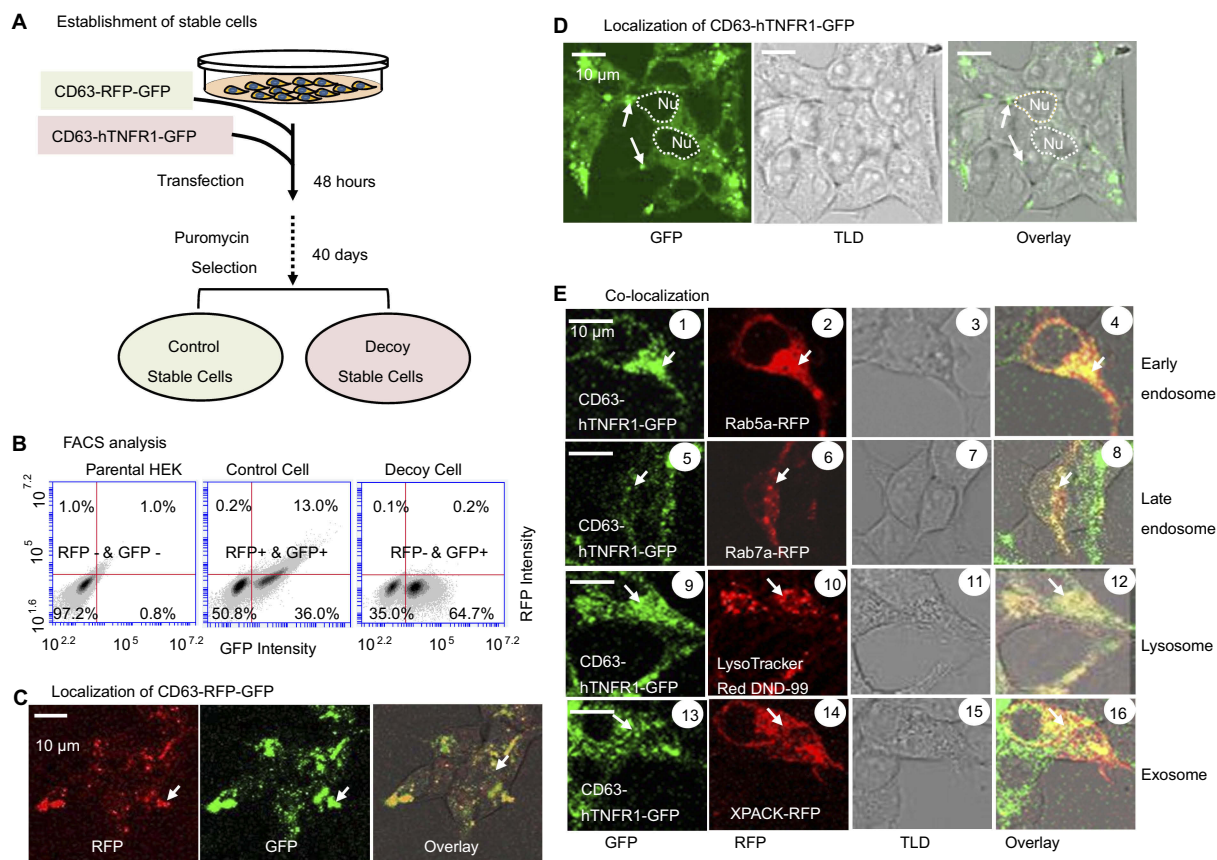


Figure 2 Establishment and characterization of stable source cell lines. **(A)** Scheme and timeline for the establishment of stable cell lines for production of either the CD63-RFP-GFP control exosomes or CD63-hTNFR1-GFP decoy exosomes. **(B)** FACS analysis of parental HEK293, CD63-RFP-GFP control, and CD63-hTNFR1-GFP stable cells. **(C, D)** Confocal images of the stable cell lines established by genetically transforming the parental HEK293 cells. Both GFP and RFP fusion proteins showed punctuated morphology in the cytoplasm for control CD63-RFP-GFP, while only GFP fusion proteins were expressed in CD63-hTNFR1-GFP cells. **(E)** Subcellular localization of the fusion proteins. Green fusion proteins were co-localized with red organelle markers for the early (1–4) or late endosomes (5–8), lysosomes (9–12), or exosomes (13–16) in the decoy exosome producing cells. Arrows indicate the organelle structures of endosome/lysosome/MVB. Scale bar: 10 μ m.

Abbreviation: TLD, transmitted light detector for light imaging.

structures identified in fluorescence images are endocytic compartments. First, we transfected stable decoy cells with a validated endosomal marker Rab5a-RFP, and then we recorded their intracellular localization expression using confocal microscopy followed by colocalization image analysis. As expected, we found the decoy cells expressed CD63-hTNFR1-GFP (Figure 2E, 1, 5) and the endosomal marker Rab5a-RFP (Figure 2E, 2, 6) after 2 days of the transfection. The signals of the two fluorescent proteins exhibited a remarkable degree of overlap in the early endosomes (Figure 2E, 4), and in late endosomes (Figure 2E, 8). Second, we transfected decoy cells with a validated lysosomal marker (LysoTracker Red DND-99) and observed an extensive overlap of green (CD63-hTNFR1-GFP) and red (LysoTracker Red) fluorescent signals – this overlap is apparent in the merged (yellow) images (Figure 2E, 1, 2). These confocal images confirm

the co-localization of CD63-hTNFR1-GFP in lysosomal compartments (Figure 2E, 1, 2), a finding that strongly indicates the expressed decoy receptors localize exclusively to exosomes. Finally, we transfected the stable decoy cells with a known exosome marker, XPACK. Similarly, we observed an extensive signal overlap between our fusion proteins (CD63-hTNFR1-GFP, green) and the exosome marker (XPACK-red) (Figure 2E, 1, 6), supporting the conclusion that our CD63-based fusion proteins target and subsequently anchored to the exosome.

Together, the results show stable decoy and control cell lines can produce either decoy receptors or control proteins. The findings of analysis of confocal fluorescence microscope images provide further evidence that CD63-based therapeutic proteins are incorporated into exosomes via the endogenous pathway that involves early and late endosomes as well as lysosomes.

Production and characterization of decoy exosomes

Next, we investigated whether decoy exosomes are released by cells into the culture medium. First, we isolated decoy exosomes from the conditioned medium of stable decoy cells through a combination of differential centrifugation, ultrafiltration, and precipitation, as reported previously (see exosome preparation flowchart, Figure 3A).³⁴ We employed nanoparticle tracking analysis (NTA) (Figure 3B, left panel) to record typical Brownian movement and size distributions of the purified exosomes. In summary, the mean size of 73–75 nm is consistent with that measured for natural exosomes (Figure 3B, right panel).³¹ To further confirm that our purified nanoparticles were exosomes, we performed an immune dot-blot analysis using a panel of 9 antibodies. The blot revealed positive stains for all 8 exosome markers (CD63, CD81, ALIX, FLOT1, ICAM1, EpCam, ANXAS, and TSG101), whereas the control cytosolic marker (GM130) was negative (Figure 3C). To qualitatively demonstrate that released decoy exosomes are loaded with CD63-TNFR1-GFP, we took confocal images from decoy and control exosome samples. The images showed strong GFP and/or RFP, indicating that the fusion proteins were targeted and anchored on the exosome (Figure 3D). Together, these

data demonstrate our stable decoy cells express, integrate and may target decoy receptors into exosomes. Importantly, the cells are found to release decoy exosomes into the surrounding culture medium. We further showed that these cells produce a sufficient number of exosomes to allow for characterizations.

Antagonistic activities of decoy exosomes

In an earlier study, we established a cell-based reporter system to precisely monitor inflammatory signaling mediated by TNF α (Additional File 2, Figure S1A)³³ – the assay can detect as little as 1 ng/ml of TNF α . Consistent with our previous observations,³³ we now show TNF α generates a robust yet unsaturated inflammatory response in our reporter cells (Additional File 2, Figure S1B). In this study, we quantified antagonistic activities of our decoy exosomes using 1 ng/mL of TNF α – this dosage is known to induce inflammatory responses both *in vitro* and *in vivo*.^{41,42} Our reporter cell models are similar to *in vitro* inflammatory assays, which have been used to study the acute response elicited by various inflammatory cytokines including TNF α as reported.⁴³

First, we tested the antagonistic activity of decoy exosomes using a co-culture experiment (Figure 4A). As demonstrated by the experiments detailed above, decoy cells express the hTNFR1-ED fusion protein in secreted exosomes. After

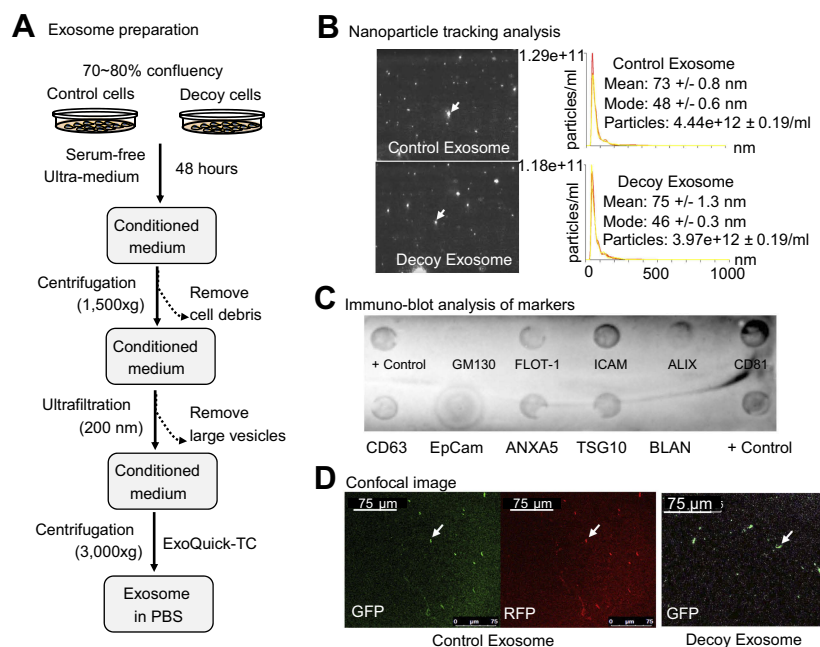


Figure 3 Exosome preparation and characterization. (A) A centrifuge-ultrafiltration-precipitation protocol used for exosome preparation. (B) Nanoparticle tracking analysis reveals exosomes in solution (left panel) along with particle size and distribution (right panel). (C) A dot-blot immuno-analysis was carried out using premade dot-blot to detect either exosomal makers, including CD63, CD81, ALIX, FLOT1, ICAM1, EpCam, ANXA5, and TSG10, or cytosolic protein GM130 for evaluating cellular contamination if any. (D) Confocal images of isolated control and decoy exosomes.

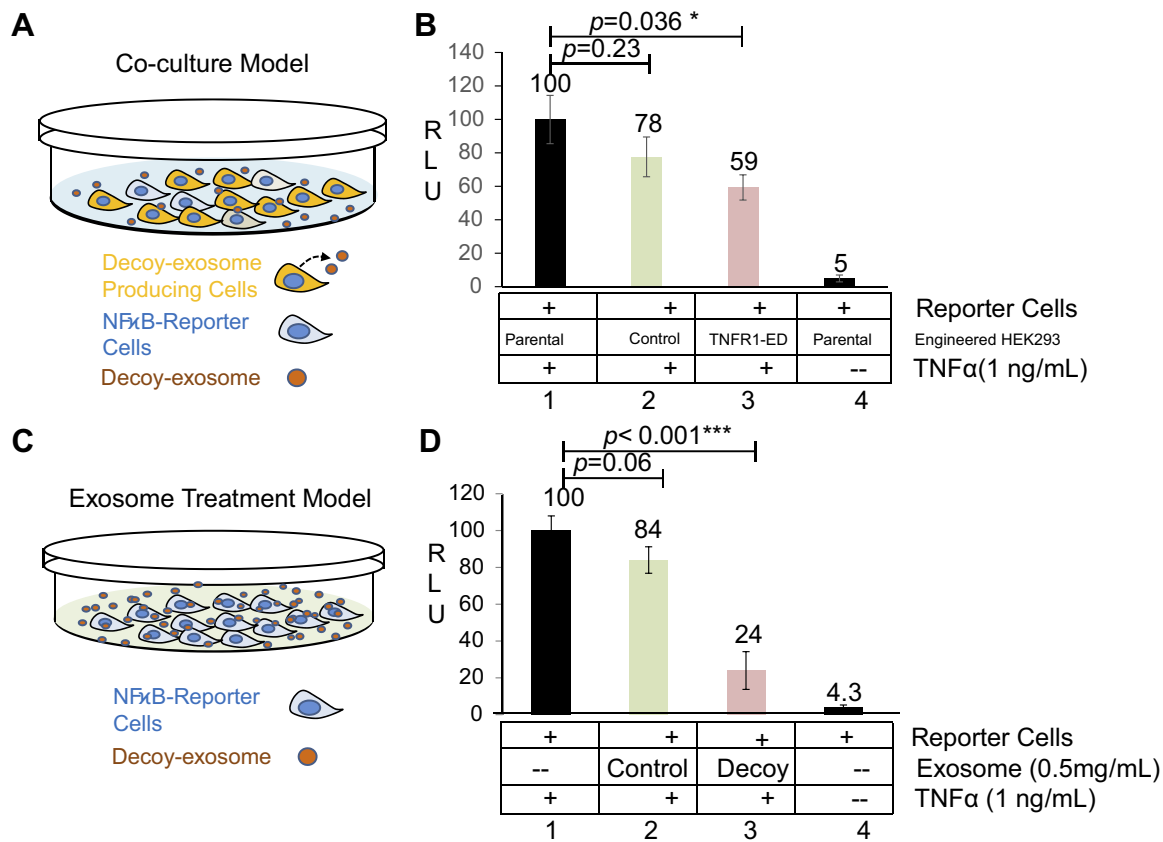


Figure 4 Efficacy of decoy exosomes on antagonizing TNFα activities in two cell models. **(A)** Co-culture experimental design for testing newly secreted exosomes and their antagonistic activities against TNFα. **(B)** Bar graph showing the specific antagonistic activities. Data are presented as mean ± SE, n=9 from 3 independent experiments. **(C)** Exosome treatment for testing the direct effect of decoy exosomes and their antagonistic activities against TNFα. **(D)** Bar graph showing the specific antagonistic activities against TNFα. Data are mean ± SE (n=6) from 2 independent experiments with two independent stable lines. The RLU was normalized to the reporter cells, which were the parental cells HEK293 cells without exosomes/engineered HEK293 cells. Column 2: Non-specific effect. Column 4: Background level activities. *Denotes P<0.05, while ***denotes P<0.001, using Student's t-test. **Abbreviation:** RLU, relative light units of luciferase activities.

mixing various source cells (parental, decoy, or mock control) with our established TNFα reporter cells for 24 hrs, we treated each co-cultured preparation with either TNFα (1 ng/mL) or vehicle control for an additional 24 hrs. Under this condition, each co-culture showed a significant increase in luciferase activity after TNFα treatment (Figure 4B, Column 1, 2 3 vs Column 4). Among them, the co-culture containing parental HEK293 cells showed a robust response to TNFα with a drastic increase in luciferase activity (~20-fold over vehicle control, Figure 4B, Column 1 vs Column 4). As expected, the luciferase activity decreased significantly in the co-culture containing decoy cells (~41% compared to parental control, P=0.036; Figure 4B, Column 1 vs Column 3), indicating an antagonistic effect of expressed TNFR1-ED towards the native TNFα receptor. In contrast, we only observed a minor decrease in luciferase activity in the co-culture containing mock control cells (~22% compared to parental control, P=0.23; Figure 4B, Column 1 vs Column 2). These data suggest decoy cells (TNFR1-ED) elicit a specific antagonism

towards the TNFα-receptors situated on cellular surfaces. Together, our results indicate that forced expression of TNFR1-ED antagonizes TNFα-mediated inflammatory signaling in HEK293 cells.

To further confirm that the observed antagonistic effects are solely due to the decoy exosomes, we treated the inflammation reporter cells directly for 24 hrs with purified exosomes prepared from various source cells (parental, decoy, and mock control), either in the presence or absence of TNFα (1 ng/mL) (Figure 4C). As expected, the reporter cells treated with parental exosomes had a robust inflammatory response to TNFα treatment, showing a marked increase (~23-fold over vehicle control) of luciferase activity (Figure 4D, Column 1 vs 4). In agreement with our co-culture results, the treatment of cells with decoy exosomes resulted in a significant decrease in luciferase activity compared to treatment with parental exosomes (~76%, P<0.001; Figure 4D, Column 1 vs 3). In contrast, treatment of cells with mock control exosomes induced a small and insignificant decrease of luciferase activity (~16%,

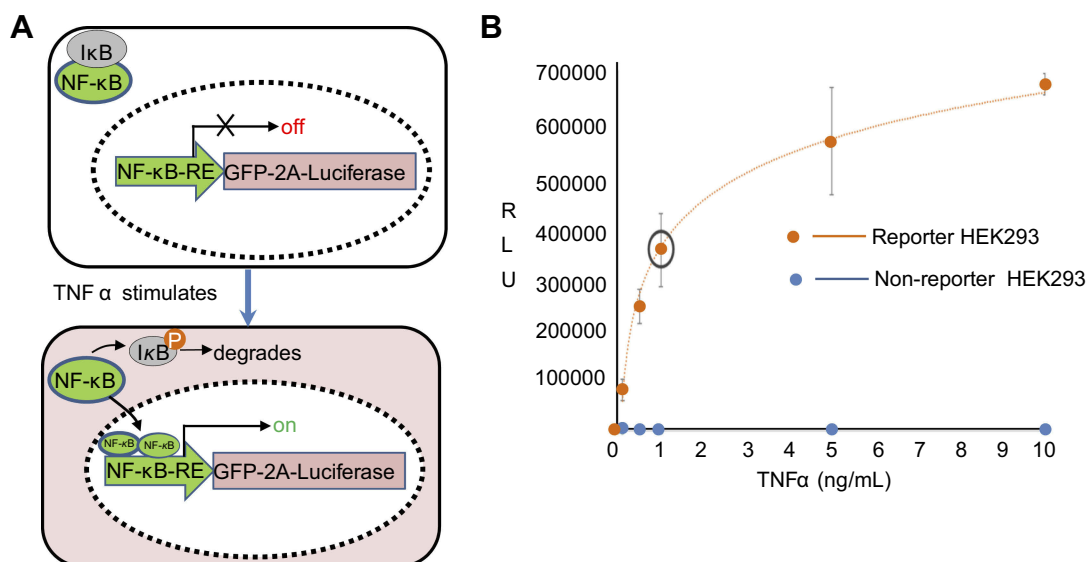


Figure S1 Reporter cell line and its dose-response to TNF α . (A) Molecular mechanism of the reporter line. A genetically encoded reporter circuit (NF- κ B-RE-driven GFP and fire luciferase gene with a self-splicing 2A peptide) was incorporated into the genome of HEK293 cells. In the absence of TNF α , the transcription factor NF- κ B remains in cytosol and associates with its inhibitor protein I κ B. Therefore, the reporter genes has little expression due to lack of NF- κ B binding to its response element (RE). In the presence of TNF α , TNF α binding to its receptor results in the phosphorylation of I κ B, which leads to its degradation and a release of NF- κ B. The freed NF- κ B then enters into nucleus and binds the RE in the promoter of reporter, resulting in an increase of luciferase activities. (B) A dose-response of TNF α . HEK293 cells were stimulated with incremental amount of TNF α for 24 hours. Reporter cells were then collected for the luciferase activities. A dose-dependent response to TNF α was evident (red line) as compared the non-reporter parental HEK293 (blue dots).

$P=0.06$; Figure 4D, Column 1 vs 2). Together, our results collectively demonstrate that decoy exosomes specifically antagonize TNF α , which indicates their potential as novel anti-inflammatory biologic therapeutic agents.

We initially tried two different concentrations of exosomes. At a lower concentration (0.1 mg/ml), the decoy exosomes produced ~24.6% inhibition of TNF α activity ($n=6$, $P=0.55$). At a higher concentration (0.5 mg/ml), our decoy exosomes produced ~76% inhibition of TNF α activity ($n=6$, $P<0.05$), indicating a high amount of decoy exosomes was required. Because decoy exosomes tended to aggregate if the concentration was beyond 0.5 mg/ml, we did not try a higher concentration to determine the maximum inhibition. It appears that the effective dosage to inhibit >75% of TNF α activity (exosomes, 0.5 mg/ml) is much higher than those of the chimeric bivalent inhibitor (TNFR1-Fc, ~10 ng/ml) or monoclonal antibodies (mAbs, 1~50 μ g/ml).^{44,45} However, one shall be careful not to overly interpret the significance of those *in vitro* results. Because the higher tissue-penetrating ability can only manifest in animals and patients, the advantage of this exosome-based method over other conventional decoy methods may not be fully appreciated until the decoy exosomes are finally tested in animal models.

The present study is focused on the earlier stage of development, including the development of an engineering strategy and proving in concept of decoy exosomes as a new

anti-TNF α reagent. A scale-up production of decoy exosomes from more desirable producing cells may be needed to obtain quality exosomes for animal studies to determine if decoy exosomes can be translated into a new class of clinical biologics for the treatment of rheumatic arthritis, psoriasis, and inflammatory bowel diseases.^{28,29,46}

Conclusion

We developed a molecular engineering approach to generate decoy exosomes that bind specifically to the inflammatory cytokine, TNF α . Our results show that these decoy exosomes antagonize TNF α -elicited signaling in cellular models of inflammation. In the future, our approach can be further exploited to display multiple receptors that are activated by other inflammatory cytokines, including interleukin-1, interleukin-6, and interleukin-12/23 – this feature, we argue, would elicit an even higher and more robust anti-inflammatory response.^{22,47,48} We also note that exosomes may exhibit a greater penetration of inflamed tissue compared to antibodies. Importantly, we anticipate extending the principle of decoy exosomes to different classes of membrane receptors to treat other diseases. For instance, decoy exosomes engineered to compete with the vascular endothelial growth factor (VEGF) could be used as a cancer therapy or for retinal diseases.^{49,50} In summary, our study demonstrates a new

avenue of therapeutics using decoy exosomes as a biological sponge to absorb detrimental factors in blood or tissues – decoy exosomes represent a novel class of biologics to treat human diseases, including inflammation, cancer, and cardiovascular disorders.

Data availability

All data generated or analyzed in this study are included in this published article and its Supplementary information files.

Acknowledgment

We thank Dr. Yan Jiang for critically reading and editing the manuscript. This work was supported by internal funds from the School of Engineering, Santa Clara University. GM acknowledges support from the Tsinghua-Berkeley Shenzhen Institute. The funding institute plays no role in the design of the study and collection, analysis, and interpretation of data.

Disclosure

The authors report no conflicts of interest in this work.

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Supplementary materials

Supplementary sequences

1. N-CD63-hTNFR1-EC-C-CD63-GFP coding sequences
2. N-CD63-hTNFR1-EC-C-CD63-GFP chimeric protein sequences

1. N-CD63-hTNFR1-EC-C-CD63-GFP coding sequences
 ATGGCGGTGGAAGGAGGAATGAAATGTGTGAAG
 TTCTTGCTCTACGTCCTCCTGCTGGCCTTTTTCGCGC
 TGTGCAGTGGGACTGATTGCCGTGGGTGTCGGGGC
 ACAGCTTGTCTGAGTCAGACCATAATCCAGGGGG
 CTACCCCTGGCTCTCTGTTGCCAGTGGTCATCATCG
 CAGTGGGTGTCTTCTTCTTCTGCTGGCTTTTGTGG
 GCTGCTGCGGGGCCTGCAAGGAGAACTATTGTCTT
 ATGATCACGTTTGCATCTTTCTGTCTCTTATCATGT
 TGGTGGAGGTGGCCGACGCCATTGCTGGCTATGTG
 TTTAGAGATAAGGTGATGTCAGAGTTTAATAACAAC
 TTCCGGCAGCAGATGGAGAATTACCCGAAAAACAA
 CCACACTGCTTTTGAATCTGGCATGGGCCTCTCCAC
 CGTGCCTGACCTGCTGCTGCCACTGGTGCTCCTGG
 AGCTGTTGGTGGGAATATACCCCTCAGGGGTATTG
 GACTGGTCCCTCACCTAGGGGACAGGGAGAAGAG
 AGATAGTGTGTGCCCAAGGAAAATATATCCACCC
 TCAAATAAATTCGATTTGCTGTACCAAGTGCCACAA
 AGGAACCTACTTGTACAATGACTGTCCAGGCCCGG-
 GGCAGGATACGGACTGCAGGGAGTGTGAGAGCGG
 CTCCTTACCGCTTCAGAAAACCACCTCAGACACT
 GCCTCAGCTGCTCAAATGCCGAAAGGAAATGGGT
 CAGGTGGAGATCTTCTTGTGCACAGTGGACCGGGA
 CACCGTGTGTGGCTGCAGGAAGAACCAGTACCGG
 CATTATTGGAGTGAAAACCTTTTCCAGTGTCTCAAT
 TGCAGCCTCTGCCTCAATGGGACCGTGCACCTCTC
 CTGCCAGGAGAAACAGAACACCGTGTGCACCTGC
 CATGCAGGTTTCTTTCTAAGAGAAAACGAGTGTGT
 CTCCTGTAGTAACTGTAAGAAAAGCCTGGAGTGCA
 CGAAGTTGTGCCTACCCAGATTGAGAATGTTAAG
 GGCAGTGGAGTACAGGCACCACAGGGCTCGATT
 TAAATTCGATCCTGGACAGGATGCAGGCAGATTTA
 AGTGCTGTGGGGCTGCTAACTACACAGATTGGGAG
 AAAATCCCTTCCATGTGCAAGAACCGAGTCCCGA
 CTCCTGCTGCATTAATGTTACTGTGGGCTGTGGGAT
 TAATTTCAACGAGAAGGCGATCCATAAGGAGGGCT
 GTGTGGAGAAGATTGGGGGCTGGCTGAGGAAAAA
 TGTGCTGGTGGTAGCTGCAGCAGCCCTTGAATTG
 CTTTTGTCGAGGTTTTGGGAATTGTCTTTGCCTGCT
 GCCTCGTGAAGAGTATCAGAAGTGGCTACGAGGTG
 ATGatgagagcagagagcggcctcccgcctgagatcgagtgccga

tcaccggcacctgaacggcgtggagttcagctggtggcggcgagagggca
 ccccaagcagggccgcatgaccaacaagatgaagagcaccaaaggcgcctg
 accttcagcccctacctgctgagccacgtgatggctacggcttctaccactcggc
 acctaccaccagcggctacgagaacctctctgcacgccatcaacaacggcggt
 acaccaacaccgcatcgagaagtagcaggacggcggcgtgctgcacgtgagct
 tcagctaccgctacgaggccggcggcgtgatcggcactcaagtggtgggac
 cggttccccgaggacagcgtgatcttcaccgacaagatcatccgagcaacgcc
 accgtggagcactgcacccatggcgataacgtgctggtgggagcttcgcc
 gcaccttcagcctcgcgcgacggcggctactacagcttctggtggacagccatg
 cactcaagagcggcctaccacccagcctctgcagaacggggcccatgttcg
 ccttcgccgctggaggagctgcacagcaacaccgagctggcagctgaggat
 accagcacgcttcaagaccccatcgccttcgagatcccgcgctcagtcgctc
 aattctgctgagcggcaccgcccggaccggctccaccggatctcgcATC-
 ATCATCATCATATAA

Abbreviations: N-CD63, N-terminus of CD63 coding sequences; C-CD63, C-terminus of CD63 coding sequences; hTNFR1-EC, human TNF α receptor 1 extracellular domain coding sequences; GFP, green fluorescent protein coding sequences

2. N-CD63-hTNFR1-ED-C-CD63-GFP chimeric protein sequences

MAVEGGMKCVKFLLYVLLLAFCACAVGLIAGV
 GAQLVLSQTIIQGATPGSLLPVVIIAVGVFLFLVAFV
 GCCGACKENYCLMITFAIFLSLIMLVEVAAIAGY
 VFRDKVMSEFNNNFRQQMENYPKNNHTAFESGM
 GLSTVPDLLLPLVLELLVGIYPSGVIGLVPHLGDR
 EKRDSVCPQKYYHPQNNISCTKCHKGTLYND
 CPGPGQDTCRECEGSFTASENHLRHCLSCSKCR
 KEMGQVEISSCTVDRDTCVCGCRKNQYRHYWSEN
 LFQCFNCSLCLNGTVHLSCQEKQNTVCTCHAGFF
 LRENECVSCSNCKKSLECTKLCLPQIENVKGTEDS
 GTTGLDLNSILDRMQADFKCCGAANYTDWEKIPS
 MSKNRVPDSCCINVTGCGINFNEKAHKEGCV
 KIGGWLKRNVLVAAAALGIAFVEVLGIVFACCL
 VKSIRSGYEVMMESDESGLPAMEIECRITGLNGV
 EFELVGGGEGTPKQGRMTNKMSTKGALTFSPYL
 LSHVMGYGFYHFGTYPSGYENPFLHAINNGGYTN
 TRIEKYEDGGVLHVSFSYRYEAGRVIKDFVVG
 GPFEDSVIFTDKIIRSNATVEHLHPMGDNVLVGSF
 ARTFSLRDGGYYSFVVDSHMHFKSAIHPSILQNG
 GPMFAFRVEELHSNTELGIVEYQHAFKTPIAFARSR-
 AQSSNSAVDGTAGPGSTGSRHHHHHH

Abbreviations: N-CD63, N-terminus of CD63; C-CD63, C-terminus of CD63; hTNFR1-EC, human TNF α receptor 1-extracellular domain; GFP, green fluorescent protein.

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