

# Application of apolipoprotein E-modified liposomal nanoparticles as a carrier for delivering DNA and nucleic acid in the brain

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**Abstract:** An innovative drug delivery technology is urgently needed to satisfy unmet medical needs in treating various brain disorders. As a fundamental carrier for plasmid DNA or nucleic acids, we developed a liposomal nanoparticle (multifunctional envelope-type nano device [MEND]) containing a proton-ionizable amino lipid (YSK-MEND). Here we report on the impact of apolipoprotein E (ApoE) modification on the function of YSK-MEND in terms of targeting brain cells. The cellular uptake and function of YSK-MEND encapsulating short interference RNA or plasmid DNA were significantly improved as a result of ApoE modification in mouse neuron-derived cell lines (Neuro-2a and CAD). Intracerebroventricular administration of ApoE-modified YSK-MEND (ApoE/YSK-MEND) encapsulating plasmid DNA also resulted in higher transgene expression in comparison with YSK-MEND that was not modified with ApoE. Moreover, observation of fluorescence-labeled ApoE/YSK-MEND and expression of mCherry (fluorescence protein) derived from plasmid DNA indicated that this carrier might be useful for delivering and conferring transgene expression in neural stem cells and/or neural progenitor cells. Thus, this system may be a useful tool for the treatment of neurodegenerative disease.

**Keywords:** brain, DNA, oligonucleotide, delivery, nanoparticles

## Introduction

A technology for curing brain disorders, such as Alzheimer's disease and Parkinson's disease, constitutes an unmet medical need. Gene therapy or treatment with functional nucleic acid, ie, short interference RNA (siRNA), is an attractive method for meeting these needs. To realize these therapies, a nanosized carrier that is capable of delivering plasmid DNA and siRNA to brain parenchymal cells is essential. Topical administration into the brain is one of the rational approaches. Recently, a Phase I study of gene therapy for curing Parkinson's disease using the adeno-associated virus encoding L-amino acid decarboxylase is currently under way.<sup>1</sup> While the adeno-associated virus-based carrier is less immunogenic in comparison with other viral vectors,<sup>2</sup> an effective nonviral carrier would still be desirable for use as a safe gene carrier.

Previously, we developed a liposomal nanoparticle known as a multifunctional envelope-type nano device (MEND) for delivering plasmid DNA or siRNA to liver,<sup>3,4</sup> cancer,<sup>5,6</sup> lung,<sup>7</sup> and adipose tissue.<sup>8</sup> Further, to improve the in vivo kinetic and intracellular fate of MEND, neutral MEND (YSK-MEND) was prepared using a pH-sensitive cationic lipid with tertiary amines (YSK05) as a key component of the envelope structure.<sup>9</sup> Several previous studies revealed that the neutral lipid particles

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could accumulate effectively in the liver with the aid of serum apolipoprotein E (ApoE).<sup>10,11</sup> ApoE is a member of the soluble apolipoprotein family that plays a key role in the cell-to-cell delivery of lipoproteins via the low-density lipoprotein (LDL) receptor family or heparin sulfate proteoglycan.<sup>12</sup> Our recent findings demonstrated that preliminary incubation of YSK-MEND with recombinant ApoE stimulated cellular uptake of these particles in a cell line derived from mouse brain endothelial cells.<sup>13</sup> In that study, ApoE was spontaneously absorbed onto the surface of the MEND by simple incubation, and was then recognized, presumably via lipoprotein-related receptors.

Meanwhile, the physiological function of ApoE in the central nervous system is currently being clarified, especially in relation to the mechanism of Alzheimer's disease and lipid metabolism.<sup>14,15</sup> In the brain, ApoE is produced by glial cells, and then functions as a ligand for cellular uptake of the cholesterol-rich high-density-like lipoproteins, that target the LDL receptor family, including LDLR and LDLR-related protein (LRP)1, which is expressed on neuronal cells, glial cells, and astrocytes.<sup>14</sup> Therefore, ApoE/YSK-MEND was expected to accelerate its cellular uptake into brain parenchymal cells when administered in the brain. In this study, our first goal was to evaluate the *in vitro* function of ApoE/YSK-MEND in neuronal cells as a pDNA or siRNA carrier *in vitro*. The second goal was to evaluate the *in vivo* function of ApoE as a ligand for brain parenchymal cells via intracerebroventricular injection using plasmid DNA as the cargo.

## Materials and methods

### Chemicals

1,1'-dioctadecyl-3,3'-tetramethylindocarbocyanine perchlorate (DiD) was purchased from Invitrogen (Carlsbad, CA, USA). Hoechst 33342 was purchased from Dojindo Laboratories (Kumamoto, Japan). Recombinant human ApoE3 was purchased from BioVision Incorporated (Milpitas, CA, USA). 1-(monomethoxy polyethyleneglycol 2000)-2,3-dimyristoylglycerol (PEG-DMG) was purchased from NOF Corporation (Kanagawa, Japan). Cholesterol was purchased from Avanti Polar Lipids (Alabaster, AL, USA). For preparation of plasmid DNA encoding luciferase that is completely free from unmethylated CpG motifs [pCpGfree-Luc(0)], the multiple cloning site of pCpGfree-mcs (InvivoGen, San Diego, CA, USA) was preliminarily replaced with a new one (5'-Bgl II-Pvu II-Nco I-Sca I-Xba I-Nhe I-3') by ligating the hybridized oligonucleotide fragments to Bgl II/Nhe I digestion of pCpGfree-mcs (pCpGfree-NEWmcs). Thereafter, a fragment encoding CpG-free luciferase plus one CpG motif just below the stop codon; Luc (+1) was

obtained by Nco I/Nhe I digestion of pORF-Luc; Sh-CpG (InvivoGen), and then ligated to the Nco I/Nhe I digested site of pCpGfree-NEWmcs [pCpGfree-Luc (+1)]. Finally, the last remaining CpG motif was removed by inserting a CpG motif-free nucleotide fragment into a Dra III/Nhe I digested site of pCpGfree-Luc (+1), where we refer to it as a pCpGfree-Luc (0).

### Animals

ICR mice were purchased from Japan SLC Inc (Shizuoka, Japan). The experimental protocols were reviewed and approved by the Hokkaido University Animal Care Committee in accordance with the Guidelines for the Care and Use of Laboratory Animals.

### Cell culture

Neuro-2a cells derived from mouse neuroblastoma cells were maintained in Dulbecco's Modified Eagle's Medium-high glucose (Sigma-Aldrich, St Louis, MO, USA) supplemented with 10% fetal bovine serum, NaHCO<sub>3</sub> (3.7 g/L), antibiotic-antimycotic (Gibco, Thermo Fisher Scientific, Waltham, MA, USA), and L-glutamine (Gibco; final concentration 0.2 mM). CAD cells were maintained in Dulbecco's Modified Eagle's Medium/F-12 (Sigma-Aldrich) supplemented with 10% fetal bovine serum, penicillin (10,000 units/mL), and streptomycin (10 mg/mL). The cells were cultured under an atmosphere of 5% CO<sub>2</sub>/air at 37°C.

### ApoE/YSK-MEND formation

YSK-MEND composed of YSK05, cholesterol (70/30) plus 3 mol% of PEG-DMG was prepared by the tertiary butanol (t-BuOH) dilution procedure as described previously.<sup>9</sup> Briefly, 10 mM of total lipids were dissolved in a 90% t-BuOH solution. siRNA or plasmid DNA was dissolved in 20 mM citrate buffer (pH 4.0) and added dropwise into the lipid solution under vigorous mixing to avoid a low local concentration of t-BuOH. Subsequently, the solution was quickly diluted with citrate buffer to a final concentration of <20% t-BuOH. The t-BuOH was removed by ultrafiltration, followed by replacement with phosphate-buffered saline (pH 7.4). Finally, the solution was again ultrafiltered to concentrate the YSK-MEND. Recombinant ApoE was preliminarily dissolved in 5 mM phosphate buffer/0.5 mM dithiothreitol and stocked as 1 mg/mL ApoE solution. For *in vitro* study, ApoE/YSK-MEND was prepared by adding ApoE stock solution to YSK-MEND at the indicated concentration, and incubated at 4°C for 6 hours. For instance, to prepare the 1 µg/mL ApoE/YSK-MEND, 50 µL of YSK-MEND (0.1 mM of

lipid concentration) was added to 2.5  $\mu\text{L}$  of ApoE stock solution (1 mg/mL) and then incubated. For in vivo use, 192  $\mu\text{L}$  of YSK-MEND (0.2 ng/ $\mu\text{L}$  of plasmid DNA concentration) was added to 4.8  $\mu\text{L}$  of ApoE stock solution (1 mg/mL) and 43.2  $\mu\text{L}$  of phosphate-buffered saline. The final concentration of ApoE was adjusted to 20  $\mu\text{g}/\text{mL}$ . The average diameter and zeta potential of the ApoE/YSK-MEND were determined using a Zetasizer Nano ZS ZEN3600 (Malvern Instruments, Malvern, UK). DiD-labeled ApoE/YSK-MEND was prepared by adding DiD (0.5 mol% of total lipid) in the lipid composition.

### Quantification of cellular uptake of ApoE/YSK-MEND by flow cytometry

The cellular uptake of the ApoE/YSK-MEND was assessed by flow cytometry. Neuro-2a cells were seeded at a density of  $1 \times 10^5$  cells per well in six-well plates (Corning Inc, New York, NY, USA) in 2 mL of the culture medium at 37°C under an atmosphere with 5%  $\text{CO}_2$ . After 24 hours, the cells were incubated with DiD-labeled ApoE/YSK-MEND at a concentration of 1  $\mu\text{M}$  of total lipid in culture medium for 1 hour. The cells were washed with phosphate-buffered saline and then trypsinized in 1 mL of phosphate-buffered saline containing 0.5% bovine serum albumin and 2 mM ethylenediamine tetraacetic acid (MACS buffer). After filtration through nylon mesh to remove cell aggregates and dust, the cells were analyzed using a FACS Calibur™ flow cytometer (Becton Dickinson, Franklin Lakes, NJ, USA). Cellular uptake of DiD-labeled MENDs was expressed as the mean fluorescence intensity, calculated using CellQuest software (Becton Dickinson).

### Quantification of gene silencing effect of ApoE/YSK-MEND by quantitative realtime (RT)-PCR

Neuro-2a cells were seeded at a density of  $1 \times 10^5$  cells per well in six-well plates in 2 mL of culture medium at 37°C. After 24 hours, ApoE/YSK-MEND encapsulating anti- $\beta$ -site amyloid precursor protein cleaving enzyme 1 (BACE1) siRNA was transfected into these cells in 1 mL of culture medium at an siRNA concentration of 150 nM. The medium was replaced with fresh culture medium after 1 hour. Cells were harvested 24 hours after transfection. Total RNA

was isolated from cells by TRIzol reagent (Invitrogen) and reverse transcribed using a high capacity RNA-to-cDNA kit (Applied Biosystems, Waltham, MA, USA) according to the manufacturer's protocol. A quantitative polymerase chain reaction (PCR) analysis was performed using SYBR Green Master Mix (Applied Biosystems) and an Mx3500P real-time quantitative PCR system (Agilent Technologies, Santa Clara, CA, USA). All reactions were performed using appropriate primer sets for BACE1 and  $\beta$ -actin (Hokkaido System Science Co Ltd, Hokkaido, Japan). The sequences of the siRNA and primers are summarized in Table 1. Amplification was carried out by primary denaturing at 95°C (10 minutes), 40 cycles of denaturation at 95°C (15 seconds), and annealing at 60°C (1 minute). The  $\beta$ -actin endogenous gene was used for relative gene quantification. Specificity was preliminarily verified by melting curve analysis. The sequences of siRNA used in the present study are listed in Table 1.

### In vitro transfection studies

For transfection,  $8 \times 10^4$  Neuro-2a or CAD cells were cultured in a 24-well dish for 1 day. The cells were then washed with phosphate-buffered saline, and YSK-MEND and ApoE/YSK-MEND containing 0.2  $\mu\text{g}$  of plasmid DNA [pCpGfree-Luc (0)] were applied to the medium, and then was incubated for 24 hours. The cells were collected by centrifugation (2,000 g, 4°C) for 2 minutes, and then solubilized with 75  $\mu\text{L}$  of reporter lysis buffer (Promega, Madison, WI, USA). Luciferase activity in a 40  $\mu\text{L}$  aliquot of cell lysate was then measured using a luminometer (Luminescencer-PSN; ATTO Corporation, Tokyo, Japan). The amount of protein in the cell lysate was determined using a bicinchoninic acid protein assay kit (Pierce, Rockford, IL, USA). Luciferase activity is expressed as relative light units per mg of protein.

### Intracerebroventricular injection

ICR mice were anesthetized with isoflurane before and throughout treatment. ApoE/YSK-MEND was injected at 1 mm lateral and 0 mm caudal to bregma, and 3 mm deep from the dura using animal testing injection needles (two-step injection needle for tuberculin, Natsume Seisakusho Co., Ltd.) and a 50  $\mu\text{L}$  Hamilton syringe (Hamilton Company, Reno, NV, USA). The ApoE/YSK-MEND (1.6  $\mu\text{g}$  of total plasmid DNA)

**Table 1** Sequences of siRNA and primers used in the present study

| siRNA          | Sense                    | Antisense               |
|----------------|--------------------------|-------------------------|
| BACE1          | GAACCUAUGCGAUGCGAAUdTdT  | AUUCGCAUCGCAUAGGUUCdTdT |
| Primers        | Forward                  | Reverse                 |
| BACE1          | AGATCTCAAGATGGACTGCAAGGA | TCTTGGGCAAGCGAAGGTTG    |
| $\beta$ -actin | AGAGGGAAATCGTGCCTGAC     | CAATAGTGATGACCTGGCCGT   |

**Abbreviations:** BACE1,  $\beta$ -site amyloid precursor protein cleaving enzyme 1; siRNA, small interfering RNA.

was injected slowly over 20 seconds. The syringe was kept in place for 1 minute after injection to prevent backflow.

## Luciferase assay

ICR mice (aged 10–11 weeks, male) were administered ApoE/YSK-MEND encapsulating plasmid DNA encoding luciferase [pCpGfree-Luc (0)] by intracerebroventricular injection at a plasmid DNA dose of 1.6 µg per mouse. After 24 hours, the brains were harvested, separated into four sections, and frozen in liquid nitrogen. Each brain quarter was homogenized using Micro Smash MS-100R (Tomy Digital Microbiology Co Ltd, Tokyo, Japan) in 800 µL of lysis buffer (100 mM Tris-HCl, 2 mM ethylenediamine tetraacetic acid, 0.1% Triton X-100, pH 7.8), and the lysate was then centrifuged at 12,000 g for 10 minutes at 4°C. The four sections of supernatants individually prepared from each brain were mixed, and luciferase activity in a 20 µL aliquot of supernatant was then measured using a luminometer (Luminescencer-PSN). The amount of protein in the supernatant was determined using a bicinchoninic acid protein kit (Pierce). Luciferase activity is expressed as relative light units per mg of protein.

## Analysis of gene expression and distribution of ApoE/YSK-MEND after injection

ICR mice (aged 10–11 weeks, male) were treated with DiD-labeled ApoE/YSK-MEND encapsulating pmCherry-N1 (Clontech Laboratories Inc, Palo Alto, CA, USA), an mCherry-encoding plasmid DNA at a plasmid DNA dose of 1.6 µg per mouse by intracerebroventricular injection. At 24 hours post-injection, the mice were sacrificed and perfused intracardially with 4% paraformaldehyde in 0.1 M phosphate buffer. In order to observe the distribution of ApoE/YSK-MEND in brain tissue, the brain was excised and 100 µm thick coronal and horizontal brain sections were prepared using a DSK-100 Microslicer™ (Dosaka EM Co Ltd, Irvine, CA, USA). For immunohistochemistry, excised brain tissue was soaked in 4% paraformaldehyde overnight

and then cryoprotected in 20% sucrose in phosphate buffer for 1 day. Subsequently, the brain was frozen in powdered dry ice. Next, 40 µm thick coronal tissue sections were cut on a cryostat (CM1900; Leica Microsystems, Wetzlar, Germany). All slices were placed into 0.05% sodium azide in phosphate buffer and stored at 4°C until use.

## Immunohistochemistry

The sections were incubated with 0.3% Triton-X/phosphate-buffered saline at room temperature for 10 minutes, and blocked in 5% goat serum/Triton-X/phosphate-buffered saline (blocking buffer) for 1 hour. Primary antibodies against NeuN, GFAP, Iba1, and Sox2 were applied to tissue sections in blocking buffer at 4°C overnight. Information on the antigen, host species, source, and specificity is summarized in Table 2. The slices were rinsed three times in Triton-X/phosphate-buffered saline, followed by incubation with Alexa Fluor 488-labeled species-specific secondary antibodies (at a dilution of 1:500 in blocking buffer) at room temperature for 1 minute. Sections were rinsed three times in phosphate-buffered saline. Thereafter, the nuclei were stained with 40 µM of Hoechst 33,342. Sections were then coverslipped with Vectashield® hard set mounting medium (Vector Laboratories Inc, Burlingame, CA, USA). The stained samples were observed by confocal laser scanning microscopy (Nikon A1; Nikon Co Ltd, Tokyo, Japan).

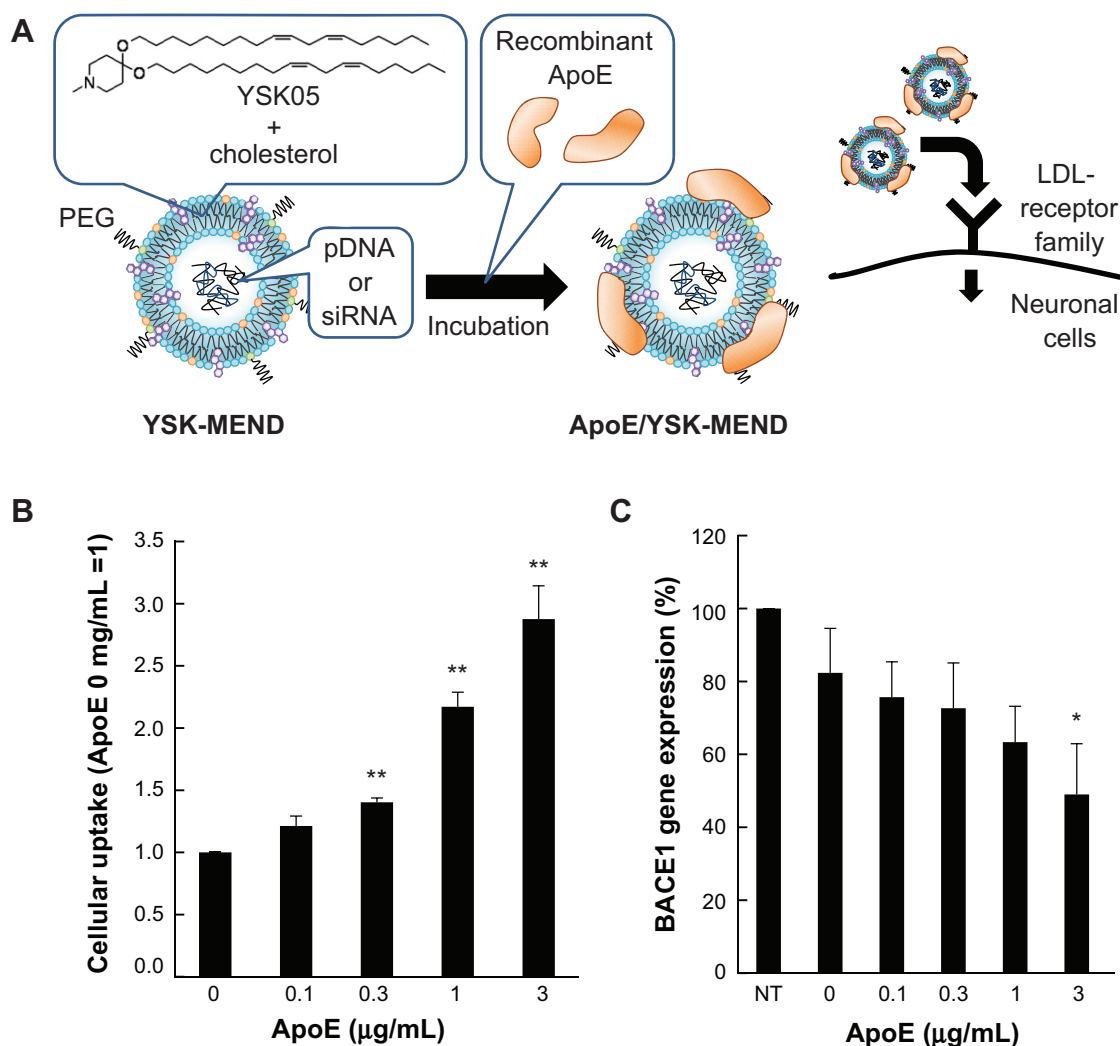
## Results and discussion

To validate the activity of ApoE as a cell targeting ligand in vitro, cellular uptake of YSK-MEND and ApoE/YSK-MEND encapsulating siRNA was first compared using a mouse neuron-derived cell line (Neuro-2a). To prepare ApoE/YSK-MEND, YSK-MEND was incubated with recombinant human ApoE3 at indicated concentrations for 6 hours (Figure 1A). The physicochemical properties of the ApoE/YSK-MEND encapsulating siRNA are shown in Table 3. The zeta potential tended to decrease depending on the ApoE modifications, while their sizes remained fairly constant. To evaluate the cellular uptake of MENDs, the particles were

**Table 2** Primary antibodies used in the present study

| Molecules | Host   | Clonality, isotype | Dilution ratio | Catalog number (supplier)                                |
|-----------|--------|--------------------|----------------|--|
| NeuN      | Mouse  | Monoclonal, IgG1   | 1:1,000        | MAB337<br>Chemicon International Inc., Temecula, CA, USA |
| GFAP      | Mouse  | Monoclonal, IgG1   | 1:600          | MAB360<br>Chemicon International Inc.                    |
| Iba1      | Rabbit | Polyclonal, IgG    | 1:1,000        | 019-19741<br>Wako Pure Chemicals, Osaka, Japan           |
| Sox2      | Rabbit | Polyclonal, IgG    | 1:1,000        | Ab97959<br>Abcam, Cambridge, MA, USA                     |

**Abbreviation:** Ig, immunoglobulin.



**Figure 1** Effect of ApoE modification on cellular uptake and gene knockdown efficacy of YSK-MEND in vitro.

**Notes:** (A) Schematic diagram illustrating the probable structure of ApoE/YSK-MEND, and a scheme for cellular uptake. (B) Neuro-2a cells were incubated with DiD-labeled ApoE/YSK-MEND encapsulating anti-BACE1 siRNA at a concentration of 1 µM of total lipids of MEND for 1 hour. Cellular uptake of ApoE/YSK-MEND was analyzed by flow cytometry. The data are represented as the mean ± standard deviation (n=3). Statistical analyses were performed by one-way analysis of variance, followed by Dunnett's test (\*\* $P < 0.01$ ) relative to ApoE-unmodified YSK-MEND. (C) Neuro-2a cells were treated with ApoE/YSK-MEND at a dose of 150 nM of total siRNA for 1 hour in the presence of serum. The gene silencing effect was then measured at 24 hours after transfection. The data are represented as the mean ± standard deviation (n=3). Statistical analyses were performed by one-way analysis of variance, followed by Dunnett's test (\* $P < 0.05$ , \*\* $P < 0.01$ ) relative to no treatment.

**Abbreviations:** DiD, 1,1'-dioctadecyl-3,3'-tetramethylindocarbocyanine perchlorate; LDL, low-density lipoprotein; ApoE, apolipoprotein E; BACE1,  $\beta$ -site amyloid precursor protein cleaving enzyme 1; pDNA, plasmid DNA; siRNA, small interfering RNA; PEG, polyethylene glycol; MEND, multifunctional envelope-type nano device; NT, no treatment.

labeled with DiD, a fluorescent marker. Cellular uptake was analyzed by flow cytometry. As shown in Figure 1B, cellular uptake was increased as a function of modified ApoE concentration. This result indicates that ApoE may well be useful as a ligand for targeting neuronal cells.

To evaluate the potential of ApoE/YSK-MEND as a carrier for nucleic acid delivery, Neuro-2a cells were incubated with

ApoE/YSK-MEND encapsulating siRNA, targeting BACE1, since knockdown of this gene is considered to be one of the strategies for curing Alzheimer's disease.<sup>16</sup> In parallel with enhanced cellular uptake (Figure 1B), the gene silencing effects increased with increasing ApoE concentration (Figure 1C).

To demonstrate the in vivo benefit of using ApoE as a ligand, YSK-MEND or ApoE/YSK-MEND was then

**Table 3** Physicochemical properties of siRNA-encapsulating YSK-MEND with or without ApoE modification

| ApoE (µg/mL)        | 0          | 0.1        | 0.3        | 1          | 3          |
|---------------------|------------|------------|------------|------------|------------|
| Size (nm)           | 103±22     | 104±19     | 105±19     | 105±17     | 105±19     |
| Zeta potential (mV) | -1.12±0.94 | -4.49±5.84 | -2.24±0.85 | -3.01±0.65 | -4.44±1.11 |

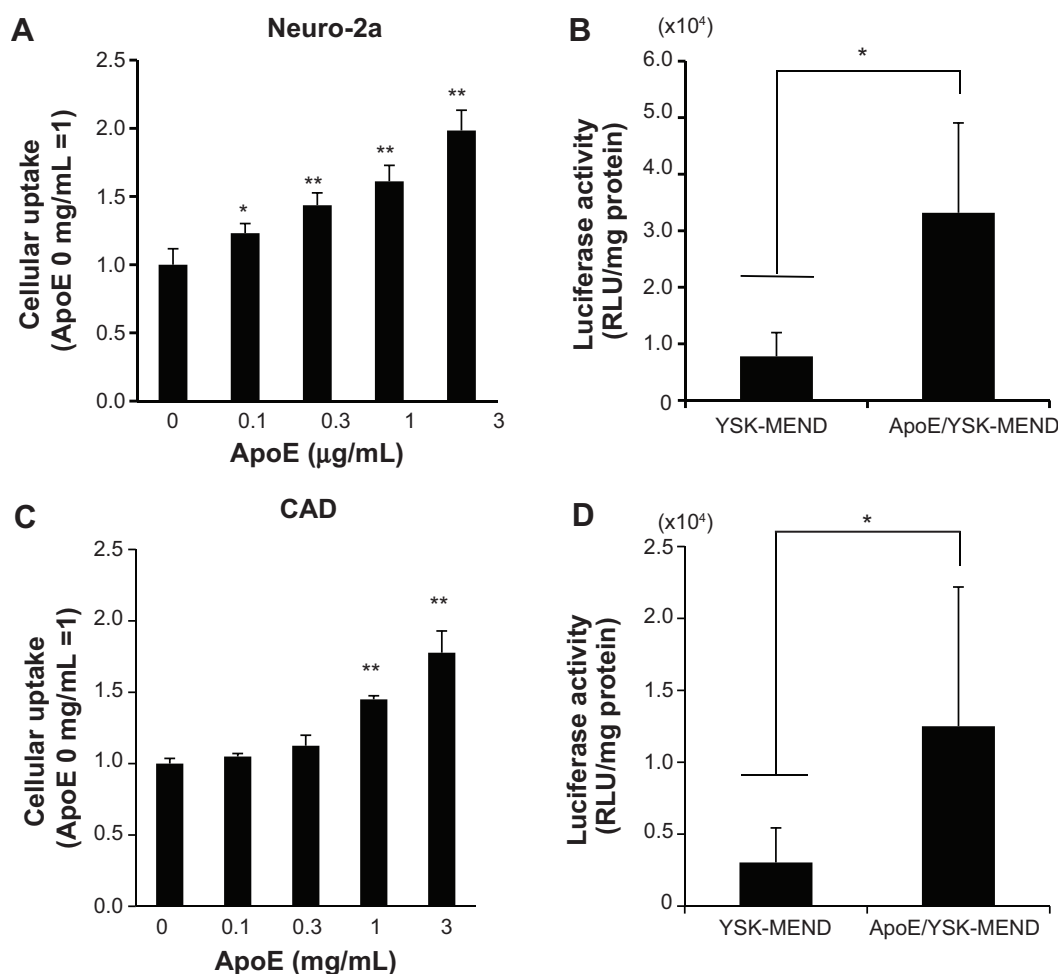
**Note:** Data are expressed as the mean ± standard deviation (n=3).

**Abbreviations:** ApoE, apolipoprotein E. siRNA, short interfering RNA; YSK-MEND, multifunctional envelope-type nano device prepared with YSK05.

administered to the mouse brain by intracerebroventricular injection. However, a recent study revealed that gene knockdown was achieved only in a narrow area within 1 mm of the site of injection when siRNA-encapsulating particles were administered.<sup>17</sup> Thus, in the *in vivo* studies, plasmid DNA encoding luciferase or mCherry instead of siRNA was used as a cargo to investigate the impact of ApoE modification on the function of YSK-MEND, since transgene expression can be quantitatively evaluated using whole brain tissue or visualized even when only a portion of the cells exhibited transgene expression. In this case, the plasmid DNA encoding luciferase was first encapsulated in YSK-MEND. For ApoE modification for *in vivo* use, YSK-MEND (6 mmol of total lipid in 240  $\mu$ L of phosphate-buffered saline) was incubated with recombinant ApoE3 at a concentration of 20  $\mu$ g/mL

overnight at 4°C. Prior to the *in vivo* study, the function of plasmid DNA-encapsulating ApoE/YSK-MEND was validated *in vitro*. In this study, another type of neuron-derived cell line (CAD), along with Neuro-2a, was used to address the generality of these phenomena. As shown in Figure 2A and C, cellular uptake of YSK-MEND was increased with increasing levels of ApoE modification in both Neuro-2a and CAD cells. Consistent with these data, uptake of ApoE/YSK-MEND (3  $\mu$ g/mL ApoE) was significantly higher than that of YSK-MEND (Figure 2B and D). Thus, the ApoE-dependent uptake machinery is available regardless of the type of cargo.

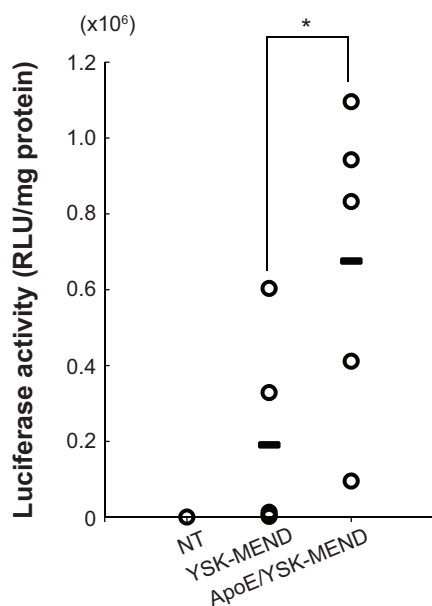
*In vivo* luciferase gene expression in whole brain was evaluated after 24 hours post-intracerebroventricular administration. Gene expression of YSK-MEND was significantly enhanced by modification with ApoE (Figure 3). Because of the



**Figure 2** Effect of ApoE-modification on cellular uptake and gene transfection activity of YSK-MEND *in vitro*.

**Notes:** Neuro-2a (A) and CAD (C) cells were incubated with DiD-labeled YSK-MEND or ApoE/YSK-MEND. Cellular uptake of ApoE/YSK-MEND was analyzed by flow cytometry. The data are represented as the mean  $\pm$  standard deviation ( $n=3$ ). Statistical analyses were performed by one-way analysis of variance, followed by Dunnett's test (\* $P<0.05$ , \*\* $P<0.01$ ) relative to ApoE-unmodified YSK-MEND. Neuro-2a (B) or CAD (D) cells were treated with YSK-MEND or ApoE/YSK-MEND (modified with 3  $\mu$ g/mL of ApoE) encapsulating pCpGfree-Luc (0) for 1 hour in the presence of serum. Gene transfection activity was then measured at 24 hours after transfection. The data are represented as the mean  $\pm$  standard deviation ( $n=3$ ). Statistical analyses were performed by one-way analysis of variance, followed by the Student's *t*-test (\* $P<0.05$ ).

**Abbreviations:** DiD, 1,1'-dioctadecyl-3,3'-tetramethylindocarbocyanine perchlorate; ApoE, apolipoprotein E; MEND, multifunctional envelope-type nano device; RLU, relative light units; YSK-MEND, multifunctional envelope-type nano device prepared with YSK05.



**Figure 3** Comparison of transgene expression of YSK-MEND and ApoE/YSK-MEND encapsulating plasmid DNA encoding luciferase after intracerebroventricular injection. ICR mice (aged 10–11 weeks, male) were administered ApoE/YSK-MEND at a dose of 1.6  $\mu$ g plasmid DNA/mouse by intracerebroventricular injection. After 24 hours, the mice were sacrificed and the luciferase activity was evaluated. Luciferase activity is expressed as relative light units per mg of protein. An open circle symbol and bars represent raw data for each mouse and the mean value, respectively ( $n=4$ ). Statistical analyses were performed using the unpaired t-test and the Mann–Whitney test. ( $*P<0.05$ ).

**Abbreviations:** ApoE, apolipoprotein E; MEND, multifunctional envelope-type nano device; NT, no treatment; RLU, relative light units; YSK-MEND, multifunctional envelope-type nano device prepared with YSK05.

existence of the blood–brain barrier, the protein concentration in cerebrospinal fluid is kept lower than that in serum. On the other hand, ApoE is abundantly supplied by astrocytes.<sup>18</sup> Ulrich et al reported that ApoE was also found in interstitial fluid at a concentration of 2  $\mu$ g/mL.<sup>19</sup> Thus, it is plausible that neutral liposomal particles, including YSK-MEND, might be inherently recognized by endogenous ApoE, and thereby gradually accumulated in brain cells. In fact, Rungta et al reported that neutralized lipid nanoparticles formed using an ionizable lipid successfully delivered siRNA to neurons when injected intracranially.<sup>17</sup> Uno et al also reported that  $\alpha$ -tocopherol-conjugated siRNA accumulated efficiently in neuronal cells after incorporation into high-density lipoprotein after intracerebroventricular injection.<sup>16</sup> Even in this situation, premodification of ApoE significantly enhanced gene expression in the brain. Thus, binding of ApoE to YSK-MEND is a rate-limiting process for cellular uptake.

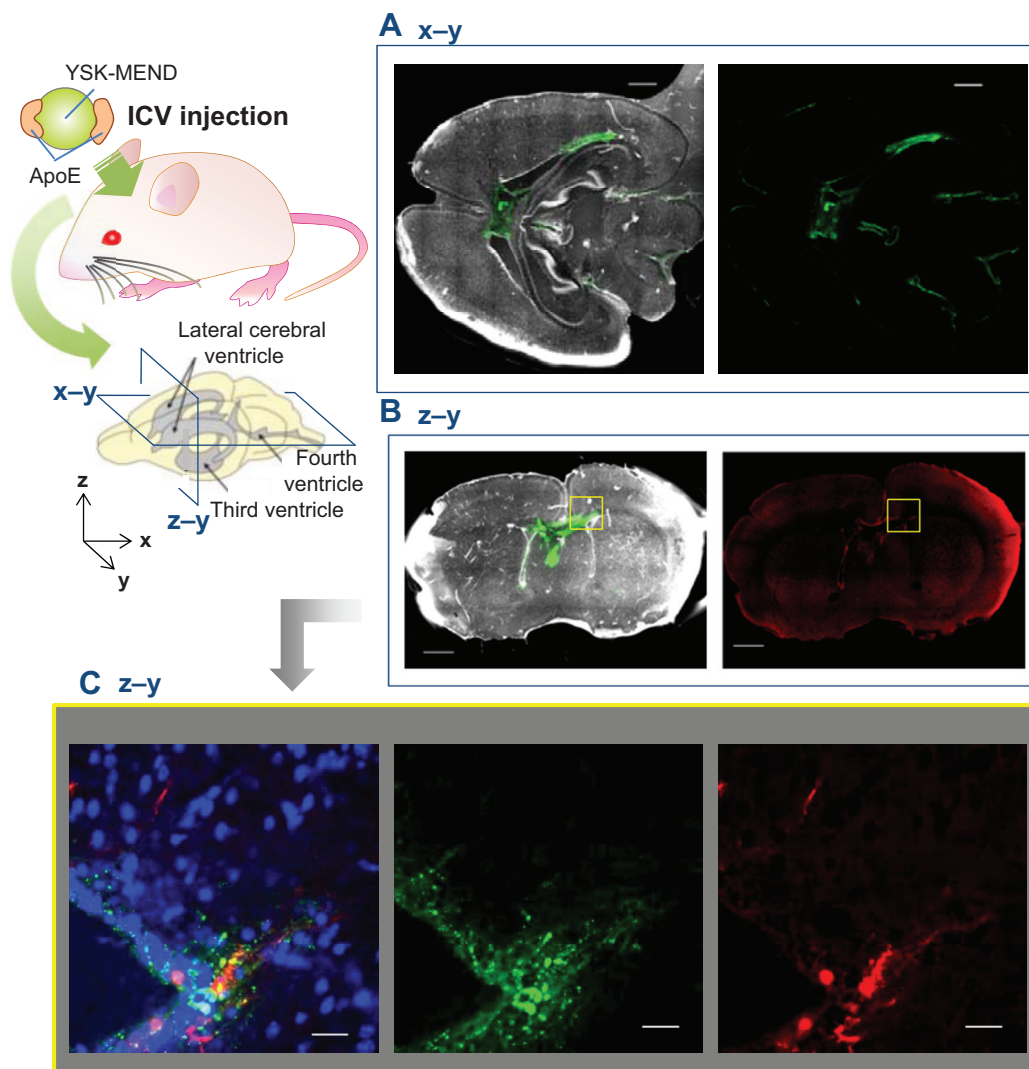
In order to identify the distribution of ApoE/YSK-MEND and the cells that are transfected, plasmid DNA encoding mCherry was encapsulated in DiD-labeled ApoE/YSK-MEND. Localization of ApoE/YSK-MEND and expression of mCherry are shown in Figure 4. Figure 4A and

B represent images in brain sections that were sliced along the coronal and horizontal planes, respectively. Figure 4C represents the magnified image from the rectangular area in the horizontal plane (Figure 4B). While ApoE/YSK-MEND was mainly observed in the paraventricular region, close to the site of administration, a fraction of the particles was detected on the surface of the cerebral cortex and cerebellum (Figure 4A). Therefore, ApoE/YSK-MEND could spread in the cerebrospinal fluid and become internalized in cells that are in direct contact with cerebrospinal fluid. The fluorescence derived from mCherry was detected in a part of the cells that take up ApoE/YSK-MEND, especially in the subventricular zone, just below the ependymal cell layer (Figure 4B and C).

The receptor for ApoE/YSK-MEND in the central nervous system remains to be elucidated. Deletion of LDLR or LRP1 in mice increases the concentration of ApoE in the central nervous system, so these two receptors play a key role in the metabolism and/or clearance of ApoE. Thus, it is most plausible that LDLR<sup>20</sup> or LRP1<sup>21</sup> contributed to cellular uptake of ApoE/YSK-MEND. The ApoE-dependent increase in cellular uptake and function of YSK-MEND in Neuro-2a in vitro also supports our hypothesis, since LDLR and/or LRP are also expressed in this cell line.<sup>22</sup>

To identify the type of cells transfected, we stained the brain sections with antibodies for typical brain cell markers. Expression of mCherry was found to be colocalized with Sox2- and/or GFAP-positive cells (Figures 5B and D), while colocalization was poorly observed in cells stained with NeuN and Iba1, markers for neuronal cells and microglia, respectively (Figure 5A and C). It has been reported that neural stem cells and neural progenitor cells are found in the subventricular zone.<sup>23</sup> Sox2 and GFAP are expressed at relatively high levels in these cells, and become negative after differentiation. Therefore, it is possible that mCherry is expressed in neural stem cells and neural progenitor cells.

Rungta et al and Uno et al reported that their ApoE-associated or high-density lipoprotein-associated carriers functioned in neural cells.<sup>16,17</sup> However, in our system, no gene expression was detected in neuronal cells. This discrepancy can be explained by differences in the cargo. In the previous studies, siRNA was delivered to the cytoplasm.<sup>24</sup> However, in this study, we delivered the necessary plasmid DNA to the nucleus.<sup>25,26</sup> The nuclear entry of plasmid DNA was severely limited by the double-layered nuclear envelope structure. Thus, non-dividing neuronal cells express hardly any transgenes, even



**Figure 4** Distribution of ApoE/YSK-MEND and mCherry expression in the brain after ICV administration of ApoE/YSK-MEND. ICR mice (10–11 weeks, male) were administered DiD-labeled ApoE/YSK-MEND encapsulating plasmid DNA encoding mCherry at a dose of 1.6  $\mu$ g plasmid DNA per mouse by intracerebroventricular injection. The frozen section was prepared 24 hours after administration and observed by confocal laser scanning microscopy. Nuclei were stained by Hoechst 33,342 ( $\lambda_{ex}/\lambda_{em}$  352 nm/461 nm; (A) and (B), white; (C), blue). DiD ( $\lambda_{ex}/\lambda_{em}$  644 nm/665 nm) was indicated by green. mCherry ( $\lambda_{ex}/\lambda_{em}$  587 nm/610 nm) was pseudocolored in red. (A) Coronal section, (B) horizontal section, and (C) magnified image of the region of interest surrounded by yellow rectangle: merged cells. Scale bars 1,000  $\mu$ m (A, B) and 25  $\mu$ m (C).

**Abbreviations:** DiD, 1,1'-dioctadecyl-3,3'-tetramethylindocarbocyanine perchlorate; ICV, intracerebroventricular; ApoE, apolipoprotein E; MEND, multifunctional envelope-type nano device; ex, excitation; em, emission; YSK-MEND, multifunctional envelope-type nano device prepared with YSK05.

when a particle can enter the cells. Unlike mature neurons, neural stem cells and neural progenitor cells actively undergo cell division, even in the adult brain.<sup>18</sup> Since the nuclear transfer of plasmid DNA and the resulting transgene expression was achieved in the mitosis phase when the nuclear membrane structure is temporarily diminished,<sup>27,28</sup> dividing neural stem cells and/or neural progenitor cells represent attractive targets for gene delivery.

## Conclusion

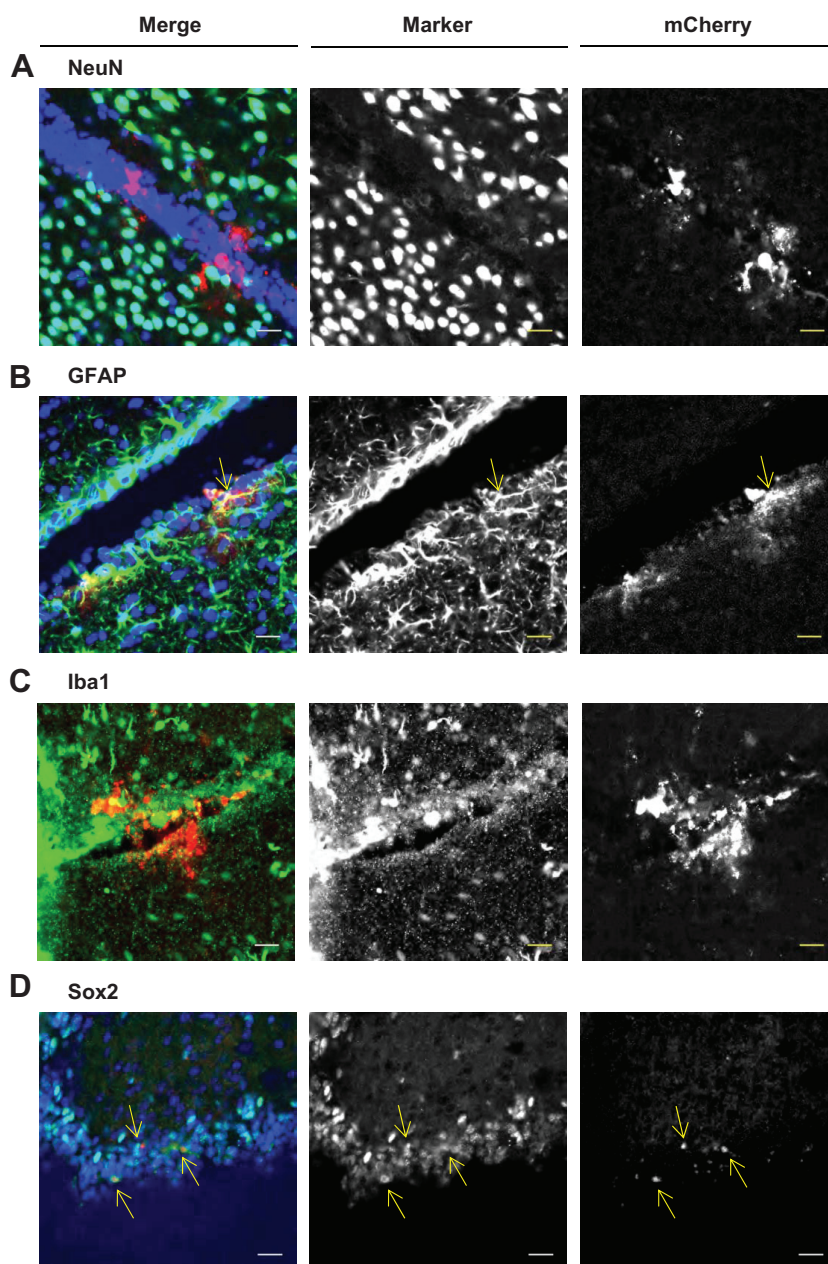
In conclusion, we demonstrated that ApoE/YSK-MEND may be useful as a nucleic acid delivery carrier when

introduced by means of intracerebroventricular injection. This nanotechnology is one of the potential solutions to unmet medical needs regarding development of a cure for neurodegenerative disease.

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**Figure 5** Identification of mCherry-expressing cells by immunostaining. ICR mice (aged 10–11 weeks, male) were administered with DiD-labeled ApoE/YSK-MEND at a dose of 1.6  $\mu$ g pDNA/mouse by intracerebroventricular injection. The frozen section was prepared 24 hours after administration and stained with antibodies which were typical cell markers for brain cells. **(A)** NeuN, **(B)** GFAP, **(C)** Iba1, and **(D)** Sox2, left panels: merged (blue; nuclei, green; immunostaining, red; mCherry), middle panels: immunostaining, right panels: mCherry. Yellow arrows indicated the mCherry/Sox2-positive cells. Scale bars represent 20  $\mu$ m.

**Abbreviations:** DiD, 1,1'-dioctadecyl-3,3'-tetramethylindocarbocyanine perchlorate; ApoE, apolipoprotein E; MEND, multifunctional envelope-type nanodevice.

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## Author contributions

Mina Tamaru, Taichi Nakatani, Yusuke Sato and Hiroto Hatakeyama acquired data. Kazuaki Kajimoto contributed to the analysis and interpretation of the results. Hidetaka Akita and Hideyoshi Harashima supervised and designed the experiments. Mina Tamaru and Hidetaka Akita wrote

the manuscript with critical input from the coauthors. All authors contributed to discussion of the results.

## Disclosure

The authors report no conflicts of interest in this work.

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