Pulmonary delivery of siRNA against acute lung injury/acute respiratory distress syndrome

Makhloufi Zoulikha\textsuperscript{a}, Qingqing Xia\textsuperscript{a}, George Frimpong Boafo\textsuperscript{a}, Marwa A. Sallam\textsuperscript{b}, Zhongjian Chen\textsuperscript{c}, Wei He\textsuperscript{a,c,*}

\textsuperscript{a}Department of Pharmaceutics, School of Pharmacy, China Pharmaceutical University, Nanjing 211198, China
\textsuperscript{b}Department of Industrial Pharmacy, Faculty of Pharmacy, Alexandria University, Alexandria 21521, Egypt
\textsuperscript{c}Shanghai Skin Disease Hospital, Tongji University School of Medicine, Shanghai 200443, China

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Abstract The use of small interfering RNAs (siRNAs) has been under investigation for the treatment of several unmet medical needs, including acute lung injury/acute respiratory distress syndrome (ALI/ARDS) wherein siRNA may be implemented to modify the expression of pro-inflammatory cytokines

KEY WORDS Pulmonary administration; Drug delivery;

Abbreviations: AAV, adeno-associated virus; Ago-2, argonaute 2; ALI/ARDS, acute lung injury/acute respiratory distress syndrome; AM, alveolar macrophage; ATI, alveolar cell type I; ATII, alveolar cell type II; AV, adenovirus; CFDA, China Food and Drug Administration; COPD, chronic obstructive pulmonary disease; CPP, cell-penetrating peptide; CS, cigarette smoke; CXCR4, C–C motif chemokine receptor type 4; DPI, dry powder inhaler; DAMPS, danger-associated molecular patterns; DC-Chol, 3\text{b}-(N\text{0},N\text{0}-dimethylethylene diamine)-carbamoyl cholesterol; DODMA, 1,2-dioleoyl-N,N-dimethyl-3-aminopropane; DOGS, dioctadecyl amido glycine spermine; DOPC, 1,2-dioleoyl-sn-glycero-3-phosphocholine; DOPE, 1,2-dioleoyl-L-\alpha-glycero-3-phosphatidylethanolamine; DOSPA, 2,3-dioleyloxy-N-[2-(sperminecarboxamido)ethyl]-N,N-dimethyl-1-propanaminium; DOTAP, 1,2-dioleoyl-3-trimethylammonium-propane; DOTMA, N-[1-(2,3-dioleoyloxy)propyl]-N,N,N-trimethylammonium; DPPC, 1,2-dipalmitoyl-sn-glycero-3-phosphocholine; dsDNA, double-stranded DNA; dsRNA, double-stranded RNA; EC, endothelial cell; EPC, egg phosphatidylcholine; EXOs, exosomes; FDA, US Food and Drug Administration; HALI, hyperoxic acute lung injury; Hem-CLP, hemorrhagic shock followed by cecal ligation and puncture septic challenge; HMGB1, high-mobility group box 1; HMVEC, human primary microvascular endothelial cell; HNPs, hybrid nanoparticles; ICAM-1, intercellular adhesion molecule-1; IFN, interferons; LPS, lipopolysaccharides; MEND, multifunctional envelope-type nano device; MIF, macrophage migration inhibitory factor; miRNA, microRNA; mRNA, messenger RNA; Myd88, myeloid differentiation primary response 88; N/P ratio, nitrogen/phosphate ratio; NETs, neutrophil extracellular traps; NF-\kappa B, nuclear factor kappa B; NPs, nanoparticles; PAI-1, plasminogen activator inhibitor-1; PAMAM, polyamidoamine; PAMPs, pathogen-associated molecular patterns; PD-1, programmed death ligand-1; PDGF\textsubscript{R\alpha}, platelet-derived growth factor receptor-\alpha; pDNA, plasmid DNA; PEEP, positive end-expiratory pressure; PEG, polyethylene glycol; PEL, polyethyleneimine; PF, pulmonary fibrosis; PFC, perfluorocarbon; PLGA, poly(D,L-lactic-co-glycolic acid); PMs, polymeric micelles; PRR, pattern recognition receptor; PS, pulmonary surfactant; RIP2, receptor-interacting protein 2; RISC, RNA-induced silencing complex; RNAi, RNA interference; ROS, reactive oxygen species; shRNA, short RNA; siRNA, small interfering RNA; SLN, solid lipid nanoparticle; SNALP, stable nucleic acid lipid particle; TGF-\beta, transforming growth factor-\beta; TLR, Toll-like receptor; TNF-\alpha, tumor necrosis factor-\alpha; VILI, ventilator-induced lung injury, Ventilator-Associated Lung Injury; *Corresponding author.

E-mail address: weihe@cpu.edu.cn (Wei He).

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1. Introduction

Acute lung injury (ALI) or its more severe clinical manifestation, acute respiratory distress syndrome (ARDS), is an acute inflammatory lung disease that leads to high rates of morbidity and mortality annually. Recently, ARDS has been divided into mild, moderate, and severe by the Berlin definition according to the ratio of the partial pressure of arterial oxygen to the fraction of inspired oxygen, and therefore excluded the term ALI, since it is analogous to mild ARDS. ALI/ARDS originates from a wide variety of direct (pulmonary) lung injuries such as pneumonia, or indirect (extrapulmonary) insults such as sepsis. Regardless of the differences between the pulmonary and extrapulmonary forms of ALI/ARDS, the terminal pathophysiological characteristics are remarkably similar, including dysfunction of the alveolar-capillary membrane, enhanced inflammation, and decreased alveolar fluid clearance, resulting in pulmonary edema, impaired gas exchange, and hypoxemia.

Despite the extensive studies and the multiple possible targets of ALI/ARDS, there are still no specific pharmacological treatments. To date, the best practice against ALI/ARDS remains protective ventilation, including low tidal volumes and high positive end-expiratory pressures (PEEP). ALI/ARDS originates from a wide variety of direct (pulmonary) lung injuries such as pneumonia, or indirect (extrapulmonary) insults such as sepsis. Regardless of the differences between the pulmonary and extrapulmonary forms of ALI/ARDS, the terminal pathophysiological characteristics are remarkably similar, including dysfunction of the alveolar-capillary membrane, enhanced inflammation, and decreased alveolar fluid clearance, resulting in pulmonary edema, impaired gas exchange, and hypoxemia.

Table 1

| Frequency | Direct (pulmonary) lung injury | Indirect (extrapulmonary) lung injury |
|----------|-------------------------------|-------------------------------------|
| Common   | Pneumonia                     | Non-pulmonary sepsis                |
|          | Aspiration of gastric contents | Major trauma                        |
|          | Lung contusion                | Transfusion of blood products       |
|          | Inhalation of toxic gases (NO₂, ammonia, phosgene) | Acute pancreatitis                  |
|          | Near drowning                 | Hemorrhagic shock or reperfusion injury |
|          | Embolism (fat embolism, amniotic fluid embolism) | Drug overdose (narcotics, sedatives, aspirin, thiazides) |
|          | Reperfusion edema following lung transplantation |                                      |

The characteristics, including clear anatomy, accessibility, relative immune privilege, and relatively low enzyme activity, make the lung a good target for local siRNA therapy. However, the translation of siRNA is restricted by the inefficient delivery of siRNA therapeutics to the target cells due to the properties of naked siRNA. Thus, this review will focus on the various delivery systems that can be used and the different barriers that need to be surmounted for the development of stable inhalable siRNA formulations for human use before siRNA therapeutics for ALI/ARDS become available in the clinic.
In this review, we will first introduce the pathophysiological mechanisms of ALI/ARDS as the first step to determine the possible targets for siRNA. The potential of siRNA therapeutic as a novel approach to combat ALI/ARDS is then emphasized. Then, we highlight the privileges of the pulmonary route for the treatment of ALI/ARDS and we discuss the different intracellular and extracellular barriers that siRNA or siRNA carriers have to overcome to reach their targets. Finally, the different delivery vectors that can be applied for siRNA pulmonary delivery in ALI/ARD are presented.

2. Pathogenesis of ALI

ALI/ARDS is divided into 3 phases: exudative, proliferative, and fibrotic. Initially, the injured lung goes by an acute/early or exudative phase characterized by damage of the alveolar barrier and pulmonary edema. Then, the exudative phase evolves into a proliferative or organizing phase after approximately 1–2 weeks to begin the process of lung repair, and finally, it may progress to an irreversible fibrotic phase in some patients.27

2.1. Exudative phase

This phase starts with the disruption of alveolar-capillary integrity by pulmonary or extrapulmonary injuries. Initially, resident alveolar macrophages (AMs) recognize nonendogenous pattern-associated molecular patterns (PAMPs), such as lipopolysaccharides (LPS) and lipoteichoic acid, or endogenous danger-associated molecular patterns (DAMPs), such as histones and high-mobility group box 1 (HMGB1), via pattern recognition receptors (PRRs) such as Toll-like receptors (TLRs). This leads to NF-κB-dependent polarization of AMs into M1-like macrophages, that release pro-inflammatory mediators and chemokines contributing to the excessive recruitment of neutrophils and monocytic cells.30–37

The aggregation of neutrophils within the pulmonary vasculature can also result from the complement activation and release of C5a. In the lung, activated neutrophils release numerous cytotoxic substances, including reactive oxygen species (ROS), various proinflammatory cytokines, granular enzymes, and neutrophil extracellular traps (NETs). In ALI/ARDS, AMs and neutrophils are the primary sources of proinflammatory cytokines, that release pro-inflammatory mediators and chemokines contributing to the excessive recruitment of neutrophils and monocytic cells.30–37

2.2. Proliferative phase

Overlapping with the exudative phase, the proliferative phase starts. In this phase, ATI cells proliferate and differentiate into ATI cells to replace injured and necrotic cells, leading to the reabsorption of alveolar edema and restoration of alveolar architecture and function.13,15,27,39 Besides the epithelial integrity, endothelial integrity must also be restored, but this process is less well understood. Interestingly, it has been suggested that bronchoalveolar stem cells also have a role in this resolution.41 Simultaneously, resolution of inflammation involves termination of pro-inflammatory signaling and clearance of apoptotic neutrophils.41 AMs shift from M1 to M2 macrophages able to produce anti-inflammatory cytokines such as IL-4 and IL-13. This results in a decrease of proinflammatory cytokines production and removal of neutrophils, necrotic cells, and debris from the alveolar space. Also, it has been reported that T-regulatory lymphocytes may suppress cytokines production and neutrophil cell death contributing to the resolution of lung inflammation.41,42

2.3. Fibrotic phase

Pulmonary fibrosis is a more frequent complication in ALI/ARDS of primary pulmonary origin, which may start in parallel with the previous phases in some patients. It is marked by extensive basement membrane damage, persistent edema, loss of elastic tissue, and obliteration of alveolar spaces with a dense irregular matrix.27,34,45 This is mainly due to the proliferation and accumulation of activated fibroblasts, secreting extracellular matrix proteins such as collagen, predominantly the fibrillar collagen type I, but also type III collagen, resulting in the development of interstitial and intra-alveolar fibrosis that may slowly resolve or cause irreversible lung destruction. The observed fibroblast activation may be regulated by growth factors such as TGF-β, coagulation cascade proteins, and proinflammatory cytokines.44

This phase does not happen in all ALI/ARDS patients but has been associated with mechanical, biochemical, and histological evidences of fibrosis with a doubling of lung collagen, progressive hypoxia, multiorgan failure, and increased mortality. (Fig. 1D)27,34,46

3. siRNAs as promising therapeutics for ALI/ARDS

RNA interference (RNAi) is a natural cellular process that silences gene expression by promoting the degradation of mRNA.44 RNAi mechanism was uncovered in Caenorhabditis elegans by Fire and Mello in 1998, but it was until 2006 when it gained momentum as they won the Nobel Prize in Physiology or Medicine.46 RNAi is a post-transcriptional gene regulation process wherein double-
stranded endogenous or exogenous noncoding RNA molecules, including small interfering RNAs (siRNA), microRNAs (miRNAs), short RNAs (shRNAs), or piwi-interacting RNAs (piRNAs), induce silencing of homologous messenger RNA (mRNA) in a sequence-specific way. Initially, the process starts by breaking down the double-stranded RNA (dsRNA) into smaller RNA duplexes, known as siRNA, by the enzyme Dicer. The resulting 21–23 nucleotide dsRNA, or the directly transfected synthetic siRNA, are then incorporated into the RNA-induced silencing complex (RISC). Subsequently, the endonuclease argonaute 2 (Ago-2) component of the RISC unwinds the duplex into two single-stranded RNAs, the “passenger or sense” that will be released, and the “guide or antisense” strand that will guide the complex RISC to the complementary target mRNA sequence. Finally, the complex cleaves the target mRNA, mediated by Ago-2, leading to inhibition of mRNA translation and the knockdown of the target gene.

Synthetic siRNAs are directly delivered to the cytosol, however, the plasmid-based shRNAs need to cross the cellular membrane to penetrate the nucleus where they are transcribed before being exported to the cytosol to be processed into active siRNAs, and has therefore attracted less attention as a pharmaceutical agent in gene therapy. Typically, siRNA can trigger specific gene silencing since the antisense strand only binds to mRNA that is perfectly complementary to it. By contrast, miRNA that has a stem-loop and whose biogenesis is initially started in the nucleus binds imperfectly to mRNA and can degrade multiple mRNA targets causing toxicity. Accordingly, RNAi therapy has unlimited choices of targets and it is possible to design effective RNAi drug targeting for any disease-promoting gene according to the mRNA sequence. siRNA is more favorable for therapeutic use over other gene therapies because it can be easily synthesized, does not require genome integration, and thereby avoids the gene therapy-related risks of mutation and teratogenicity, with no problem of permanent modification since siRNA therapy can be stopped and controlled at any time and stage of treatment. The delivery of siRNA is easier compared with shRNA plasmid or pDNA delivery since its site of action is the cytosol. Furthermore, siRNA has a small size, higher transfection efficacy, potency and specificity, fewer obstacles, and

Figure 1  Schematic illustration of normal alveolus (A), injured alveolus during the exudative (B), proliferative (C), and fibrotic (D) phases in ALI/ARDS.
lower immune response, making them the best fit for RNAi
therapeutics\textsuperscript{22,23}. Since the discovery of the Nobel
prize-winning mechanism of RNAi fifteen years ago\textsuperscript{46}, it has become a promising drug target
for treating a myriad of diseases, including genetic diseases,
autoimmune and inflammatory diseases, cancer, and viral infec-
tion\textsuperscript{53}. To date, three siRNA-based therapeutics are already in
the pharmaceutical market. The first is ONPATTRO\textsuperscript{®} (patisiran), a
lipid nanoparticle administered intravenously once every 3
weeks, approved by the US Food and Drug Administration (FDA)
in August 2018, to treat hereditary transthyretin amyloidosis
(hATTR). The second is GIVLAARI\textsuperscript{™} (givosiran) approved in
the USA in November 2019 to treat acute hepatic porphyria. The
third is Oxlumo\textsuperscript{™} (lumasiran) approved in November 2020 by the
EU and FDA to treat primary hyperoxaluria type 1. Both givosiran
and lumasiran are administered subcutaneously as N-acetylga-
lactosamine (GalNAc) conjugates\textsuperscript{54,55}. Beyond the 3 siRNAs-
based drugs already approved, there are another 7 in phase 3 tri-
als. For instance, nedosiran is developed to treat primary hyper-
oxaluria, vutrisiran for the treatment of hATTR like patisiran,
teprasiran for the prevention of acute kidney injury after transplant
or cardiovascular surgery, fitusiran, for hemophilia A and B,
tivanisiran to treat ocular pain and dry eye disease, and cosdosiran
to treat nonarteritic anterior ischemic optic neuropathy. Also,
siRNA therapy was expanded from orphan and rare diseases to
more prevalent conditions, such as inclisiran for hypercholester-
olemia\textsuperscript{56}. Most of these drug candidates are GalNAc conjugates,
except teprasiran, cosdosiran, and tivanisiran, which were not
delivered with a carrier or conjugated to a targeting ligand.

Notably, siRNAs have been previously reported to be an
effective nucleic acid-based therapy in respiratory diseases\textsuperscript{57–60},
including ALI/ARDS, by targeting the essential inflammatory
pathways involved in this disease\textsuperscript{51,58}. Much work has been
done to find potential therapeutic targets of ALI/ARDS, such as those
involved in epithelial-endothelial dysfunction, immune cell
recruitment, or fibroblast activation; to reinforce lung defense
mechanisms and enhance lung repair processes\textsuperscript{59,63}. Nevertheless,
these studies need to be accompanied by simultaneous research
about the best delivery strategies of siRNA therapy of ALI/ARDS
to achieve rapid successful outcomes.

4. Pulmonary delivery of siRNA for treatment of ALI/
ARDS: Advantages and barriers

4.1. Advantages of siRNA pulmonary delivery

The delivery of drugs to the lung can be achieved directly via the
pulmonary route or indirectly by systemic administration. How-
ever, nucleic acids can be degraded by serum nucleases in the
bloodstream, hurdling the systemic delivery for siRNA\textsuperscript{19}. In
contrast, the local administration of siRNAs offers several
important benefits over systemic delivery, such as minimized undesirable systemic effects, as well as reduced dose of siRNAs
due to the high bioavailability at the target site and the lower lung
nuclease activity improving siRNA stability and efficacy (Fig. 3)\textsuperscript{20,23,58,62,64}. Most importantly, the pulmonary route allows
direct access to lung EpiCs, key target cells for gene knockdown
in ALI/ARDS\textsuperscript{21}.

Although inhalation of siRNAs is clinically the most common
and non-invasive method, the majority of in vivo studies were
conducted intratracheally or intranasally due to their simplicity
and ease of administration to animals compared to the inhalation
formulations\textsuperscript{21,58,62}. Clinically, intranasal instillations are easy to
perform. However, the particles are vastly filtered out in the
human nasal cavity. Similarly, intratracheal administration is
relatively invasive and not suitable for human use\textsuperscript{57}.

It has been reported that the most appropriate method for
pulmonary administration of siRNA-based therapeutics is inhalation
using pressurized metered dose inhalers (pMDIs), requiring
aerosolization of the drug by suspension or dissolution in a pro-
pellant\textsuperscript{57}. The two other inhalation devices, nebulizers and dry
powder inhalers (DPIs) are less common owing to the higher
chance of siRNA degradation by the applied stress of nebulizers, and the difficulty to formulate siRNA-containing powders or preserving the integrity and bioactivity of siRNA, which hinders the use of DPIs\(^\text{x9}\). However, the use of liquid formulations is not suitable for long-term storage. Accordingly, DPIs seem to be the more promising option because they are more stable, and have a lower risk of contamination extending their shelf-life\(^\text{26,63,66}\).

### 4.2. Barriers of siRNA pulmonary delivery

Despite all the advantages of siRNA pulmonary delivery over systemic delivery, it still faces several barriers. This includes extracellular barriers that can be anatomical, physiological, metabolic, and immunological due to the inherent properties of the lung as well as intracellular barriers, related to the cellular uptake and endosomal escape\(^\text{69,70}\), all with considering the disease pathological changes. Overcoming these barriers is indispensable to develop a successful inhalable siRNA against ALI/ARDS\(^\text{61}\).

#### 4.2.1. Extracellular barriers

To reach its target, siRNA needs first to overcome the under-mentioned extracellular barriers.

##### 4.2.1.1. Anatomical barriers

The human respiratory tract includes 23 groups of branching airways. The first 16 generations from the nasal cavity to the bronchi constitute the region of the conducting airways responsible for the conduction of gases, and the next 17–23 generations extending from the bronchioles to alveoli compose the respiratory airways responsible for the gaseous exchange\(^\text{71}\). This branched structure of the airways with different diameters and lengths presents an initial hurdle that siRNA must pass along to reach the deep lung area\(^\text{57,66}\). Here, the site of deposition mainly depends on the size of particles. For deep penetration into the alveolar region, particles should have an optimal aerodynamic diameter between 1 and 5 \(\mu m\), smaller enough to not deposit in the upper airways and larger enough for not being exhaled during normal breathing\(^\text{52,72}\). With maximum deposition in alveoli region between 1 and 2 \(\mu m\). However, 100 \(\mu m\) particles have been stated to settle successfully into the alveolar space due to the Brownian diffusion mechanism\(^\text{77,82,66,73}\). To avoid the exhalation of nanoparticles and improve their lung deposition, the “Trojan microparticles” or nanoparticles-in-microparticles, have been proposed to deliver them as disso-ciable microparticles having desired aerodynamic range (1–5 \(\mu mol/L\) in size) but can disassemble into nanoparticles upon lung deposition. Simultaneously, this strategy provides the aerodynamic behavior of microparticles and the ability of nanoparticles to avoid clearance\(^\text{52,75}\). Accordingly, a previous study conducted in a rodent model using near-infrared (NIR) fluorescent NPs reported a negligible systemic absorption (less than 5% after 1 h) of particles bigger than 100 \(\mu m\) after intratracheal administration. This study showed also that particles smaller than 34 \(\mu m\) were rapidly removed to the lymph and those smaller than 6 \(\mu m\) were quickly excreted in the urine\(^\text{76}\). These findings support the use of 100–200 \(\mu m\) as the optimal nanoscale for pulmonary route\(^\text{74}\). Typically, the extrapulmonary distribution of NPs is neglected in this nanoscale range. However, it may be enhanced by the loss of alveolar-capillary barrier integrity associated with ALI/ARDS\(^\text{75}\).

##### 4.2.1.2. Physiological and metabolic barriers

After lung deposition, the next hurdle for siRNA pulmonary delivery to overcome is the clearance of particles. Clearance mechanisms depend on the site of deposition in the lung. The particles are mainly removed by the mucociliary clearance in the conducting airways and the AMs in the respiratory region\(^\text{75,77}\). The continuous turnover of mucus, the mucociliary clearance driven by the physical action of the ciliated EpiCs, and the possible interactions (electrostatic, hydrophilic, hydrophobic) with particles, constitute a major pulmonary delivery barrier\(^\text{75,76}\). The interaction of pulmonary delivered particles with mucus and their mucociliary clearance are both surface and particle-size-dependent\(^\text{75}\). It was found that particles larger than 500 \(\mu m\) were sterically immobilized. In contrast, 100–200 \(\mu m\) diameter particles rapidly penetrated the mucus of human respiratory systems\(^\text{79}\). Also, positively charged particles will be more trapped in the mucus rich in anionic mucin glycoproteins than the negatively charged particles. Yet, the commonly utilized siRNA delivery vehicles are polycationic which may hurdle their penetration to the mucus barrier, and therefore their therapeutic efficacy\(^\text{82}\). In contrast, neutral particles

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**Figure 3** Pulmonary versus systemic delivery of siRNA to the lung.
exhibited higher average transport rates\textsuperscript{75}. The hydrophilic and neutral surfaces can be achieved by coating particles with hydrophilic and neutrally charged polymers, such as polyethylene glycol (PEG), or charge reversible peptides. In addition to the glycosylated regions, the mucin peptides may also trap hydrophobic particulates (Fig. 4)\textsuperscript{72,78}. Favorably, an enhanced mucoadhesion could prolong the lung retention time of pulmonary delivered particles, leading to an improved and pharmacokinetic profile\textsuperscript{75}. The impaired mucociliary clearance in ALI/ARDS can further prolong this retention.

Besides the respiratory mucus, the alveolar surface is covered by pulmonary surfactant (PS), a complex mixture of lipids and proteins\textsuperscript{82}. PS can strongly interact with drugs hindering their effectiveness\textsuperscript{19,57}. Fortunately, this may not have a great impact on ALI/ARDS since surfactant production is disrupted. Formerly, PS has been considered as a siRNA delivery barrier to the respiratory target cells. Lately, a beneficial role of PS in local pulmonary siRNA delivery has been illustrated, hypothesizing that the lung surfactant proteins may act as an “endogenous” delivery vector\textsuperscript{82}.

On the other hand, the ALI/ARDS-increased alveolar fluid can hamper the drug molecules from reaching the epithelium. Moreover, the possible interaction of the cationic siRNA complexes with the anionic constituents of alveolar liquid, including phospholipids and proteins, especially in protein-rich edema fluid in ALI/ARDS constitutes another barrier that can hinder their diffusion. Also, the consolidated or collapsed alveoli further hinder the transducing of acutely injured lungs. Likewise, an associated cough can increase the removal of siRNA from the airways\textsuperscript{19,61}. Despite the lower metabolic capacity of the lung and its negligible nuclease activity, naked siRNA or siRNA carriers may undergo enzymatic degradation following pulmonary delivery, mediated by enzymes such as antitrypsin and proteases\textsuperscript{61,75}. 4.2.1.4. Therapeutic barriers. Mechanical ventilation (MV) largely impacts the aerosolization efficiency and deposition within the respiratory tracts of the pulmonary delivered siRNA/siRNA carriers\textsuperscript{83,84}. Harmfully, the presence of heat and humidity in the ventilator circuit can reduce the efficiency of drug delivery\textsuperscript{83}. In patients receiving mechanical ventilation, it is difficult to ensure that the aerosolized particles are efficiently delivered to distal airways\textsuperscript{85}. It has been reported that the aerosol generated by nebulizers in mechanically ventilated patients was mostly deposited in proximal airways with a lung deposition lower than 20% of the nominal dose\textsuperscript{4,86}. Thus, inhalation devices and administration procedures have to be improved for patients under MV\textsuperscript{84}. Positively, the time required to deliver a specified amount of drug to its target and the proportion of drugs delivered could be better managed by controlling the ventilator settings (tidal volume, airflow, inspiratory time/total duration of the breath)\textsuperscript{83,85}.

4.2.2. Intracellular barriers
After overcoming the aforementioned extracellular barriers, siRNA has to bypass a set of intracellular barriers to reach RNAi machinery in the cytoplasm.

4.2.2.1. Cellular uptake. The anionic cell membrane is another barrier to siRNA delivery. Naked siRNA does not easily cross the cell membrane due to its negative charge and

Figure 4 Strategies to overcome the mucosal barrier.
high molecular weight. Thus, effective delivery strategies may be used to facilitate siRNA cellular uptake. The major cell uptake pathway associated with the non-viral delivery of siRNA is endocytosis. Generally, the cationic siRNA-carrier interacts with anionic proteoglycans and enters the cells by endocytosis. Clathrin-dependent endocytosis is the best characterized and the most common route involved in the cellular uptake of siRNA during pulmonary delivery, particularly if the diameter is less than 200 nm. Particles under 150 nm can both avoid macrophage uptake and be efficiently endocytosed. Besides, caveolea-dependent endocytosis may also mediate siRNA cellular entry, especially for sizes greater than 500 nm.

4.2.2.2. Endosomal escape. After cellular entry, the entrapped siRNA/siRNA-carrier vesicles are transported in early endosomes that mature into late endosomes (pH 5–6), and then into lysosomes (pH ~4.5) containing nuclease and other degradative enzymes. Thereby, siRNAs must escape into the cytosol (pH ~7.4) at an early stage to avoid lysosomal degradation. The endosomal escape is the rate-limiting step for efficient siRNA delivery. Several strategies have been investigated to promote the endosomal escape, including the incorporation of polymers such as polyethyleneimine (PEI) due to their high buffering capacity causing an increased osmolarity and endosome disruption, fusogenic lipids such DOPE (1,2-dioleoyl-L-α-glycero-3-phosphatidylethanolamine) or pH-sensitive fusogenic peptides such as GALA (a repeat of glutamic acid-alanine-leucine-alanine, WEAAEAELAEALAEELAEELAA) owing to their ability to undergo conformational changes in low pH, causing a destabilization of endosomal membrane. Notably, it has been reported that even 0.1% endosomal escape in the cytosol of target cells may be enough for effective RNAi performance since siRNA can amplify itself within the cell. For caveolea-mediated endocytosis, vectors are entrapped in nonacidal vesicles, transported to the Golgi or endoplasmic reticulum, and in that way escaping the lysosomal degradation.

5. siRNA delivery systems for treatment of ALI/ARDS

Although the use of siRNAs seems to be a powerful approach against ALI/ARDS, its clinical application is still hindered by many obstacles as discussed earlier. Similar to other nucleic acid-based therapeutics, this includes off-target effects, triggering of interferon responses, poor stability, and inefficient in vivo delivery, which have been the main challenge to applying siRNA-based therapeutics in humans. The properties of suitable delivery systems for siRNAs as potential therapeutic agents include the ability to: (1) entrap siRNA into a stable particle, (2) effectively protect siRNA from enzymatic degradation, (3) avoid pulmonary extracellular barriers, (4) possess high transfection efficiency to facilitate its cellular uptake, (5) have good intracellular trafficking properties to promote the endosomal escape and the cytosolic release of siRNA. Also, an optimal siRNA delivery vector should be biodegradable, biocompatible, with negligible toxicity and immunogenicity, and should not cause inflammation. Also, it should be suitable for large-scale manufacturing (Fig. 5). Owing to the similarities between DNA and RNA, many vectors tested for DNA delivery have been explored for drug delivery of siRNA to the lung against ALI/ARDS (Table 2).

5.1. Viral vectors

Virus-based delivery vectors, such as adenoviruses, adeno-associated viruses, retroviruses, lentiviruses, were used to introduce genetic material into host cells after removing their virulence genes and inserting engineered genes into their genomes.

5.1.1. Adenovirus

Adenovirus (AV), a non-enveloped double-stranded DNA (dsDNA) virus, was approved for clinical trials in 1990 as the first viral gene delivery vector. Gendicine, which is an AV-based system loaded with mutated p53 gene, was approved by the CFDA in 2003 for treatment of head and neck squamous cell carcinoma as the first-ever gene therapy. AV is widely used in lung-targeted gene therapy. However, AV-mediated gene delivery is limited by AV-induced immunological responses, inflammation, and non-specificity of cell targeting. Further, AV-mediated gene transfer is relatively inefficient in the absence of epithelial damage. However, this is not a limitation in the case of ALI/ARDS which is characterized by epithelial barrier dysfunction.

5.1.2. Adeno-associated virus (AAV)

AAV is a small single-stranded DNA nonpathogenic parvovirus that is replication-defective and needs a helper virus to complete its lytic life cycle. AVVs are safer and less immunogenic compared to AV, which makes them an attractive candidate for gene delivery. However, they have lower transfection efficiency. The AAV serotypes have different tissue tropisms due to different capsid proteins. The serotype AAV6 has been reported to be the most effective vector in infecting pulmonary epithelium. Interestingly, AV capsid has been modified to improve both the safety and efficacy of these vectors for their application in gene therapy of several human diseases especially lung diseases. Recently, Duncan et al. highlighted the mucus-penetrating ability of AAV6 as a potential mechanism to achieve robust in vivo lung gene transfer and effective inhaled gene therapy. Successfully, AAV6 was used in a recent study to deliver miRNA-21-5p intending to inhibit apoptosis of ATII cells in hyperoxic acute lung injury (HALI) rats’ model. Encouragingly, two AAV-based therapies have been approved by the FDA, luxturna (AAV2-based drug) indicated for rare eye disease and zolgensma, (AAV9-based drug) for spinal muscular atrophy, in 2017 and 2019, respectively.

5.1.3. Retroviruses

Retroviruses are enveloped RNA viruses characterized by their ability to reverse transcribing single-stranded RNA into dsDNA, before its insertion into the genome. The non-specificity, the insertional mutagenesis, and the risk of tumorigenesis are their major drawbacks. The most common type of retroviruses is lentivirus. Lentiviral vectors have been reported to have lower genotoxicity risk compared to other retroviral vectors.

Generally, despite the high transduction efficiency of viral vectors, their clinical application is limited by cytotoxicity, high immunogenicity, high potential of harmful insertional mutagenesis in the host genome, and even carcinogenesis. Besides, viral delivery systems are more suitable for RNAi therapeutics that require transport into the nucleus such as miRNA and shRNA but not for siRNA that acts in the cytoplasm.
5.2. Non-viral vectors

Non-viral carriers have been developed to avoid the immune system stimulation associated with viral vectors. Despite their relative safety and low-cost production, these vectors generally do not possess the high level of tissue tropism and transfection efficiency of viral vectors. The non-viral vectors applied for the pulmonary delivery of siRNA include various delivery systems, such as lipid-based, polymer-based, peptide-based delivery systems, and hybrid nanoparticles, etc.

5.2.1. Lipid-based delivery vectors

Commonly, lipid-based systems are the most investigated vectors for siRNA delivery. These include liposomes, stable nucleic acid lipid particles (SNALPs), solid lipid nanoparticles (SLNs), pH-responsive lipids, and exosomes (Table 3 and Fig. 6).

5.2.1.1. Lipoplexes (cationic liposomes).

So far, liposomes are the only nanomedicine approved by the FDA for inhalation. Two types of liposomes could deliver siRNA to the lung. Cationic liposomes by complexation and neutral liposomes by inclusions of fusogenic lipids, such as DOPE in cationic liposomes, were implemented for siRNA transfection. Many commercially available transfection agents such as Lipofectamine (DOTMA/DOPE), or lipofectamine (DOSPA/DOPE) which are based on cationic liposomes, were implemented for siRNA transfection.

Despite the several advantages of cationic liposomes, including efficient in vitro transfection, high loading capacity, structural flexibility, excellent biocompatibility, biodegradability, and ease of large-scale production, their advancement is still hindered by some drawbacks. These include safety concerns, nonspecific tissue uptake, and premature siRNA release due to their interaction with the cell membrane and thereby reduce the transfection efficiency. The inclusion of fusogenic lipids, such as DOPE in cationic liposomes, is thought to increase the interactions between the liposomal and endosomal membranes facilitating the release of siRNA. The transfection efficiency can be further improved by optimizing the ratio of lipid:siRNA or N/P ratio (Nitrogen to phosphate ratio) to obtain a slightly positive charged lipoplex.

Various modifications have been made to protect cationic liposomes from non-specific interactions, reduce their immunological responses and cellular toxicity, and promote endosomal escape. This was achieved by the addition of PEG on their surface to form “stealth liposomes.” However, PEGylation may be more advantageous in the systemic delivery challenged with the reticuloendothelial clearance system than the pulmonary administration since it can hinder their interaction with the cell membrane, and thereby reduce the transfection efficiency. The inclusion of fusogenic lipids, such as DOPE in cationic liposomes, is thought to increase the interactions between the liposomal and endosomal membranes facilitating the release of siRNA.

The ideal delivery system for siRNA.

![Figure 5](image-url)
cationic lipids such as DOTAP are less toxic than the multivalent cationic lipids such as Lipofectamine®/C226, but also less efficient in packing and binding nucleic acids115,125,141. Recently, Xu et al.91 used both naked siRNA and liposomal encapsulated siRNA by intratracheal and i.v. delivery, respectively, to silence PD-L1 (Programmed death ligand-1) expression significantly up-regulated in pulmonary ECs and/or EpiCs after hemorrhagic shock followed by cecal ligation and puncture septic challenge (Hem-CLP)-induced ALI in mice model. Interestingly, treatment with naked siRNA mainly affected EpiCs but did not target ECs. In contrast, PD-L1 siRNA delivery by liposome mostly targeted ECs, but not EpiCs. Consequently, i.v. administration of liposomes loading the same siRNA attenuated the development of Hem-CLP induced ALI. These results were not observed when naked siRNA was given intratracheally, due to the key role of pulmonary ECs in mediating the PD-1: PD-L1 pathway mainly targeted by i.v. delivery of the liposomal form91. In another study conducted by Dong et al.92, Lipofectamine loaded with siRNA was administered intratracheally to suppress Receptor-interacting protein 2 (RIP2) upregulated in cigar ette smoke (CS)-induced ALI mouse model.

Rip2 knockdown successfully ameliorated CS-
induced inflammation compared with the control siRNA. Intravenously administered Lipofectamine siRNA was also used to achieve targeted protein and cell-specific knockdown in AMs in lung ischemia-reperfusion injury rat model. Toll-like receptor-4 (TLR4) siRNA was specifically taken up by AMs, but not by pulmonary artery ECs nor ATII cells. Consequently, TLR4 siRNA-specific knockdown could significantly ameliorate lung injury. Furthermore, neither siRNA nor the lipid vector administration was associated with interferon production.

5.2.1.2. Neutral liposomes. Neutral liposomes composed of neutral lipids, such as cholesterol and its derivatives, DOPC (1,2-dioleoyl-sn-glycero-3-phosphocholine), and DOPE; have also been tested for siRNA delivery by the encapsulation of siRNA into

Figure 6  Schematic representation of different lipid-based vectors used for siRNA delivery against ALI/ARDS.

Figure 7  Schematic illustration of cellular uptake and intracellular trafficking of lipoplexes.
the aqueous core of the liposome. The loss of net charge improves the safety profile of lipid vectors, but it hinders their interaction with siRNA resulting in a relatively low siRNA entrapment compared to their cationic counterparts (less than 10%). Hence, neutral lipids are often used as transfection enhancers with other vectors. Frequently, DOPE is used as a fusogenic lipid to mediate endosomal release due to its ability to shift to an inverted hexagonal phase at the acidic environment of endosomes supporting the fusion between the nanocarrier and the endosomal membrane.

5.2.1.3. Stable nucleic acid lipid particles (SNALPs). SNALPs were first developed in 2001 as a new generation of cationic liposomes. SNALPs are composed of a mixture of an ionizable positively charged phospholipid, such as 1,2-dioleoyloxy-N,N,N-trimethyl-3-aminopropane (DODMA) or 1,2-dioleyl-3-dimethylammonium-propane (DODAP), as well as a neutral helper lipid (including Chol and fusogenic lipids), to overcome the two siRNA intracellular barriers (cellular uptake and endosomal release), with a PEG-coating to prevent degradation and nonspecific interactions.

5.2.1.4. Solid lipid nanoparticles (SLNs). Besides the liposomal formulations, SLNs have been adapted for siRNA delivery to the lung to avoid stability issues associated with liposomes. SLNs are composed of a neutral hydrophobic core, where siRNA can be incorporated by the hydrophobic ion-pairing approach and coated with a lipid membrane. Usually, lipids such as fatty acids/esters/alcohols, triglycerides, waxes, and cholesterol, are used in formulating SLNs. Most recently, aerosolizable TNF-a siRNA-encapsulated SLNs were successfully prepared for potential pulmonary delivery, suggesting the possibility to deliver the siRNA-SLNs dry powder into the lung to silence the gene expression of a gene of interest in cells, such as AMs or lung EpiCs.

5.2.1.5. Multifunctional envelope-type nano device (MEND). MEND is a nanoparticle in which siRNA is entrapped in the inner phase of a lipid bilayer modified with functional molecules such as peptides. Also, pH-responsive lipids can be incorporated into the lipid membranes to control their intracellular trafficking. For example, a GALA-MEND has been recently designed and optimized by incorporating a pH-responsive lipid (YSK05) to deliver CD31 siRNA to the pulmonary endothelium based on GALA peptide targeting. The incorporated YSK05 enhanced the endosomal release of the optimized MEND resulting in an improved knockdown in the lung endothelium by comparison with DOTMA or DOTAP-MENDs when intravenously injected into mice. Furthermore, the incorporated helper lipids such as DOPE synergistically enhanced both the selectivity and efficiency of the designed MEND. Thus, GALA modified MEND may be a potential carrier of siRNA in curing various lung diseases including ALI/ARDS.

5.2.1.6. Exosomes (EXOs). EXOs are endogenous extracellular nanovesicles derived from the endosomal system with a particle size ranging from 30 to 150 nm. EXOs are released by various types of cells into the extracellular environment as nature’s nano delivery system of lipids, proteins, and nucleic acids, playing an important role in intercellular communication.

Interestingly, Zhang et al. used serum-derived exosomes as vehicles to deliver siRNA against Myd88 (involved in innate immune signaling) to the lung. siRNAs were first loaded into serum EXOs via modified calcium-mediated transfection and then were intratracheally instilled in LPS-induced ALI mouse model. Efficiently, the siRNA-carriers were internalized by AMs, achieved specific gene silencing, and modulated LPS-induced inflammation. More interestingly, serum-derived EXOs themselves were neither inflammmogenic nor immunogenic when delivered into the lung. Curiously, the intratracheally instilled EXOs were only taken up by AMs, but not by LPS-activated neutrophils. Accordingly, the delivery of EXO-based siRNA to other lung cells, such as EpiCs, requires the development of new methods to avoid their uptake by AMs. Another study suggests that exosomes of human induced pluripotent stem cells with a size of 122 nm isolated by differential centrifugation, could be used as a natural delivery system of ICAM-1 siRNA to inhibit the intercellular adhesion molecule (ICAM-1) protein expression in HMVECs (Human primary pulmonary microvascular endothelial cells). EXOs were successfully internalized by LPS activated HMVECs, leading to selective suppression of ICAM-1 surface expression in vitro which may alleviate ALI in vitro. Based on those findings, EXOs seem like an attractive candidate for siRNA delivery because of their natural potential to carry genetic material, their native biocompatibility with low inherent immunogenicity and toxicity in comparison with exogenous carriers. Furthermore, they can potentially evade the endosomal pathway and lysosomal degradation. However, their purification techniques are cumbersome and expensive for large-scale manufacturing. Polymer-based carriers can be separated into two categories: polycations and polymeric nanoparticles.

5.2.2. Polymer-based delivery vectors Polymer-based vectors are attractive candidates for siRNA delivery systems because of their facile synthesis and lower immunogenicity compared to liposomes. Generally, polymer-based carriers can be separated into two categories: polycations and polymeric nanoparticles.

5.2.2.1. Polycations. Similar to lipoplexes, polyplexes or dendriplexes are formed by electrostatic interactions between their cationic amine (N) groups and the anionic phosphate (P) groups of nucleic acids. They include synthetic polycations such as poly-ethyleneimine (PEI), polyamidoamine (PAMAM) dendrimers, and natural polycations such as chitosan. The transfection efficiency of cationic polymers is influenced by their structure, size, and surface charge. Also, the degree of complexation quantified by the N/P ratio affects the net charge, size, and stability of the delivery system. Typically, an excess of the polymer is needed to generate stable cationic vehicles and facilitate their internalization. However, an excessively high positive charge may cause cytotoxicity. Hence, an appropriate N/P ratio should be found for each system to assure the balance between transfection efficiency and cytotoxicity. The polycations internalized by the clathrin-dependent pathway use the “proton sponge” effect to escape from endosomes. The proton sponges cationic polymers, such as PEI and PAMAM, prevent acidification of endocytic vesicles owing to their high buffering capacity inciting the membranes H+-ATPases to actively translocate H+ from the cytoplasm into the endosome to maintain the desired pH. The increased protonation causes an influx of chloride and water into the endosome leading to disruption of the endosomal membrane, and eventually endosomal escape of the endocytosed polyplexes into the cytosol.

Although PEI has an excellent gene condensation ability, high buffering capacity leading to great endosome escape activity, and
high transfection efficiency\textsuperscript{153}, it suffers from molecular weight-dependent cytotoxicity and nonbiodegradability hindering its application as an effective delivery system\textsuperscript{154}. The positive charge of PEI has been incriminated in necrotic and apoptotic cell death both in vitro and in vivo\textsuperscript{155}. Also, it has been demonstrated that linear PEIs are less toxic than their branched counterparts\textsuperscript{156}.

5.2.2.1. Polyplexes. In vivo-jetPEI\textsuperscript{TM} is a linear PEI that has low toxicity and high efficiency for local gene and siRNA delivery compared to the other cationic lipids and polymers\textsuperscript{157}. Fu et al.\textsuperscript{158} used JetPEI polyplex as a transfecting agent to deliver paxillin siRNA into the mouse lung by intratracheal instillation to suppress LPS-induced paxillin accumulation which is hypothesized to mediate the associated endothelial hyperpermeability. Resultantly, in vivo knockdown of paxillin attenuated LPS-induced pulmonary endothelial hyperpermeability and injury. Further, other pulmonary cells might be affected by this knockdown\textsuperscript{159}.

In a previous study, dexamethasone-conjugated PEI (DEXA-PEI) was utilized as a siRNA vehicle into lung tissue for the knockdown of macrophage migration inhibitory factor (MIF) overexpressed in particularate matter (PM)-induced airway inflammation. The results showed that DEXA-PEI/MIF siRNA reduced the PM-induced MIF expression in PM-activated EpiCs compared to naked MIF siRNA after intratracheal application in mice. The maximized reduction of inflammation compared to dexamethasone (which itself, with or without scrambled siRNA, has an anti-inflammatory effect) suggested the synergistic effect between siRNA and the developed delivery system\textsuperscript{157}.

Fluorination may increase the stability of polycations, lower their toxicity, and improve cellular siRNA delivery through enhancement of cellular uptake and endosomal escape\textsuperscript{160}. Interestingly, perfluorocarbon (PFC), a biocompatible polymer widely used in pulmonary therapies due to its high oxygen dissolving capacity, has been demonstrated to attenuate ARDS in both humans and animals\textsuperscript{159}. For example, it has been demonstrated that lung delivery of vaporized PFC improved the gas exchange in a detergent-induced lung injury\textsuperscript{161}. Another study also stated that inhalation of PFC may protect from blast lung injury\textsuperscript{159}. Similarly, it was reported that i.v. administration of PEI emulsion significantly improved LPS-induced ALI in rat’s model\textsuperscript{162}. Therefore, PFC may represent an excellent candidate for siRNA delivery of ALI/ARDS. To achieve combined inhibition of CXCR4 (C→X→C motif chemokine receptor type 4) and PAI-1 (plasminogen activator-inhibitor-1) which are overexpressed in LPS-activated EpiCs\textsuperscript{159}, Wang et al.\textsuperscript{159} prepared PFC emulsion polyplexes containing a fluorinated polymeric CXCR4 antagonist for the delivery of siRNA against PAI-1. The resultant polyplex emulsions protected siRNA from degradation and boosted siRNA uptake and endosomal escape with no significant cytotoxicity observed. After intratracheal instillation, PFC emulsion polyplexes showed strong lung retention and improved therapeutic outcomes in LPS-induced ALI model in mice\textsuperscript{159}. However, some reports indicated that the presence of fluorine can negatively influence the pulmonary distribution of siRNA polyplexes. It was found that siRNA-nonfluorinated polyplexes were mainly accumulated in the lungs after i.v. delivery in contrast to their fluorinated counterparts that were majorly distributed to the liver. Beneficially, the fluorine modification did remarkably reduce PEI cytotoxicity without altering its binding affinity and improved the in vitro siRNA delivery potency\textsuperscript{162}.

5.2.2.1.2. Polymeric micelles (PMs). PMs are nanocarriers (usually <100 nm) formed of a hydrophilic polymeric shell, such as PEG, and a hydrophobic core, such as PEI\textsuperscript{163,164}. Their biocompatibility, low toxicity, and relatively high stability make them attractive delivery systems\textsuperscript{165}. Cationic micelle-forming block polymers can be used to trap negatively charged siRNAs in their core via electrostatic interaction\textsuperscript{166}. Interestingly, PMs can be engineered to achieve active targeting and improved cellular uptake by their conjugation with specific targeting ligands or stimuli-responsive components\textsuperscript{167}. However, they are limited by low loading efficiency and complexity in transport through the cell membrane\textsuperscript{168}.

Most recently, Hou J et al.\textsuperscript{168} developed PMs composed of branched PEI modified with PEGs and loaded with siRNAs against PTPN13 (protein tyrosine phosphatase-N13) and NADPH oxidase-4, which are promoters of pulmonary fibrosis (PF) that constitutes a severe complication in ALI/ARDS. The fibroblast-targeting was achieved by the anti-platelet-derived growth factor receptor-α (PDGFRα) antibody onto the carrier. The developed carrier achieved remarkable antibifibrotic effects and successfully suppressed the development of bleomycin-induced PF in murine model\textsuperscript{168}. The same PMs loaded with siRNA against runt-related transcription factor-1 and decorated with an anti-stem-cell antigen-1 antibody could successfully inhibit PF in mice by preventing the differentiation of lung resident mesenchymal stem cells into myofibroblast\textsuperscript{169}.

5.2.2.1.3. Dendriplexes. Polycationic dendrimers, characterized by their unique three-dimensional structure, can also be used to condense siRNA into dendriplexes\textsuperscript{170}. Cationic PAMAM and phosphorous dendrimers were widely tested for siRNA delivery owing to their positive charge, their high degree of surface functionality, their excellent buffering capacity, and high transfection efficiency\textsuperscript{149}.

PAMAM dendrimers were applied to locally deliver TNF-α siRNA to the lung. Compared with non-complexed siRNAs, the cationic dendriplexes with a size between 127 and 153 nm displayed a high siRNA condensation, low cytotoxicity, increased cellular uptake, and transfection performance, and efficiently silence TNF-α in vitro in LPS-activated macrophages. The intranasal pre-administration of PAMAM dendriplexes in a murine LPS-induced lung inflammation model resulted in improved performance 4 h after the LPS challenge. However, PAMAM dendriplexes showed lower efficacy than non-complexed TNF-α siRNA on the 3rd day, requiring more frequent administrations. This could be explained by the higher stability of the non-complexed siRNA and their deep penetration into the lung retard- ing their removal compared to dendriplexes that undergo fast mucociliary clearance\textsuperscript{160}. Interestingly, Agnoli et al.\textsuperscript{171} engineered inhalable dry powders of TNF-α siRNA-loaded G4 PAMAM dendriplexes embedded in microparticles formed of a saccharide matrix, aiming to avoid the unfavorable exhalation of nanoparticles and facilitate their transport. The designed system was well-tolerated and showed an enhanced cellular uptake of siRNA leading to efficient silencing of TNF-α in LPS-activated macrophages. Favorably, the suitability of this system for inhalation therapy was confirmed by the efficient knockdown and the preserved siRNA integrity upon spray drying in the reconstituted NPs\textsuperscript{171}.\textsuperscript{171}
Comparatively, it has been reported that cationic phosphorus dendrimers have better delivery efficiency of siRNA than PAMAM dendrimers\textsuperscript{172}. Further, these dendrimers themselves could possess a potent anti-inflammatory action\textsuperscript{173}. A previous study conducted by Bohr et al.\textsuperscript{97} inspected the potential of TNF-α siRNA cationic phosphorus dendrimers to modulate LPS-induced ALI both in vitro and in vivo. In their research, phosphorous third-generation (G3) dendrimers were functionalized with either cationic pyrollidinum (DP) or morpholinum (DM) terminal groups. Results showed a strong binding between siRNA and both DP and DM dendrimers, with stronger binding for DP. This was attributed to the different pKa values between DP (8.1) and DM (7.1) resulting in higher ionization of DP in the pH of complexation. However, DP dendriplexes exhibited higher cytotoxicity against macrophages. This could be partly related to their higher surface charge. Similarly, DP dendriplexes showed better cellular uptake and improved in vitro silencing efficiency of TNF-α in LPS-activated macrophages than DM dendriplexes and/or non-complexed siRNA. In correlation with in vitro results, the nasal administration of DP dendriplexes in LPS-induced ALI murine model strongly inhibited TNF-α expression, much higher than DM dendriplexes and free siRNA\textsuperscript{97}. Accordingly, in implementing dendriplexes for siRNA delivery, the dose should be carefully adjusted to compromise between achieving the highest efficacy and at the same time avoiding carrier-associated toxicity.

Poly-l-lysine (PLL) is a well-known cationic polymer that has been widely studied as a non-viral gene delivery vector owing to its strong positive charge\textsuperscript{74}. Recently, Yang J et al.\textsuperscript{30} used dendritic PLL to deliver TNF-α siRNA to the lung. Interestingly, they have used charge-reversible pro-peptide of RAGE-binding peptide modified with cis-acconitic anhydride for the surface-coating of the delivery nanocomplex and giving it a negative surface charge. This allowed an efficient mucosal penetration of the developed nanosystem after intratracheal delivery. Then, the positive charge was recovered in the mild acidity of the inflamed alveolar space (pH ≈ 6.8), allowing efficient transfection of AMs and TNF-α silencing\textsuperscript{30}.

5.2.2.2. Polymeric nanoparticles. The polyplex formation depends on electrostatic interactions and the anionic nature of the RNA payload. However, their superstructure may be compromised by the competitive interactions with other physiological poly-anions\textsuperscript{147}. For this reason, polymers capable of forming neutral or anionic NPs have been suggested. Unlike polycations, siRNA is either dispersed throughout the polymeric matrix or adsorbed on the NPs surface. Appropriate surfactants or cationic polymers such as PEI or chitosan can be added to the matrix\textsuperscript{174}. Practically, the delivery of siRNA by these materials is challenged by their relative low nucleic acid loading and transfection efficiency\textsuperscript{147,148}. NPs based on hydrophobic polymers such as PLGA [poly(\textit{D,L}-lactic-co-glycolic acid)] have attracted more attention to deliver siRNA\textsuperscript{52,57,175}. The endosomal escape of PLGA NPs is due to the surface charge reversal caused by the pH change from anionic (in the physiological and alkaline pH) to cationic (in the endosome/lysosome acidic pH)\textsuperscript{52,174}. After the endosomal escape, PLGA nanoparticles may remain in the cytosol for more than 14 days, allowing a slow release of the payload\textsuperscript{174}. Further, combination with cationic polymers or cationic lipids can improve its transfection efficiency and intracellular trafficking\textsuperscript{176}. Frede et al.\textsuperscript{177} designed a multi-shell nanoparticle (size of 145 nm and zeta potential of + 23 mV), consisting of a calcium phosphate core, coated with siRNAs, encapsulated within PLGA, and finally coated with a final outer layer of PEI, as a vehicle for local treatment of pulmonary inflammatory disorders. Successfully, multiple nasal instillations of the calcium phosphate PLGA NPs loaded with siRNAs against different pro-inflammatory mediators (CCL-2, IP-10, and IFN-γ) to mice suffering from TH1 cell-mediated lung inflammation showed fruitful internalization (mainly endocytosed by macrophages and dendritic cells and less by EpcCs and lymphocytes), protected siRNA molecules, and decreased gene expression, thereby mitigated the inflammatory responses in the lung\textsuperscript{177}.

5.2.3. Peptide-based delivery vectors

Peptides are other possible carriers that can be applied for siRNA delivery because of their stability, low cytotoxicity and immunogenicity, biocompatibility, and biodegradability\textsuperscript{179}. Peptides can be used alone or as an assistant component to other systems, such as fusogenic peptides used to bypass the endosomal entrapment barrier and cell-penetrating peptides (CPPs) used to facilitate the cellular entry\textsuperscript{180}.

5.2.3.1. Fusogenic peptides. Fusogenic peptides are short peptides that can promote endosomal release by increasing interactions with endosomal membrane\textsuperscript{180}. GALA is a synthetic pH-sensitive fusogenic peptide able to convert from a random coil structure at pH6 to an amphipathic a-helical structure able to insert into membranes and cause membrane leakage at pH6, promoting the endosomal escape\textsuperscript{181,182}. As abovementioned, GALA-modified MENd has been proposed as a siRNA carrier to take advantage of the dual role of GALA, enhancing the endosomal release, and targeting the lung endothelium\textsuperscript{129}.

5.2.3.2. Cell-penetrating peptides (CPPs). CPPs are short chains of about 30 or fewer amino acids. Based on differences in physicochemical characters, CPPs can be classified into three classes: cationic, hydrophobic, and amphipathic\textsuperscript{183–185}. Internalization of siRNA conjugated with CPPs can be mediated via endocytic or nonendocytic pathways depending on the particle size, peptide type, and siRNA loading method\textsuperscript{186}. However, some CPPs have limitations due to non-specificity and easy proteolysis\textsuperscript{187}. Moreover, CPP-mediated delivery of siRNAs is majorly challenged by endosomal entrapment\textsuperscript{188}. It has been reported that CPPs remain localized in the endosome and are further captured by lysosomes rather than undergoing endosomal escape\textsuperscript{188}. Therefore, various modifications have been used with CPPs to promote endosomal escape and improve RNAi efficiency. This includes the attachment of fusogenic peptides such as HA2 peptide derived from influenza virus hemagglutinin, fusogenic lipids such as DOPE, or the use of buffering agents such as chloroquine\textsuperscript{173}. Generally, CPPs can either vectorize siRNA by direct covalent conjugation through a linker or by non-covalent conjugation through electrostatic and hydrophobic interactions between CPPs and siRNA, typically forming nanoparticle structures\textsuperscript{184,189}. So far, no CPPs or CPPs/cargo complexes have been approved by the FDA\textsuperscript{185}.

5.2.3.2.1. Covalent CPPs delivery of siRNA. Besides the activation of innate immunity, the covalent conjugation of siRNA to CPPs can sterically hinder the interaction between siRNA and RISC reducing its efficiency. Additionally, conjugates with charged nucleic acids are unstable and difficult to purify\textsuperscript{187,190,191}. Hence, covalent modification is most suitable for charge-neutral oligonucleotides than charged molecules such as siRNA\textsuperscript{120}.

5.2.3.2.2. Non-covalent CPP delivery of siRNA. The formation of nanoparticle complexes between anionic nucleic acids...
and CPPs is mainly driven by electrostatic and/or hydrophobic interactions\(^{195,196,197}\). Nevertheless, CPPs have poor encapsulating efficiency, and the nanoparticles formed are more heterogeneous and less defined than their CPP/siRNA conjugate counterparts\(^{196,197}\). Further, CPPs have relatively low transfection efficiency requiring additional modifications to form efficient and stable CPP polyplexes because of their low molecular weight, weak electron charge density, inefficient gene complexation capability, limited capacity for endosomal escape, and rapid degradation under physiological conditions. Generally, CPPs are not used alone but as assistant components in lipopolymer and polymeric polyplexes\(^{178}\).

Interestingly, CPP can be used for the co-delivery of genes and drugs\(^{186}\). For example, R3V6, a cationic amphiphilic peptide was used as a carrier for the delivery of the combinatorial therapy of sphinogosine-1-phosphate lyase siRNA (siS1PLyase), reported to be involved in acute pulmonary inflammation, and recombinant HMGB1A, an antagonist of the pro-inflammatory cytokine HMGB1. The ternary NPs of siS1PLyase-HMGB1A-R3V6 has been designed to test the synergistic effect of siS1PLyase-HMGB1A against LPS-induced ALI model\(^{98,192}\). Remarkably, the ternary NPs and siRNA/R3V6 nanocomplexes displayed higher stability compared with naked siRNA or siRNA/PEI, confirming that R3V6 could protect siRNA from nucleases and proteases. The designed nanocomplex exhibited similar delivery efficiency into the lung EpiCs compared to siRNA-HMGB1A-PEI and siRNA-HMGB1A-Lipofectamine. Moreover, it was shown to be less toxic than both of them with higher anti-inflammatory potency against LPS-activated macrophages. Consistently, the intratracheal administration of siS1PLyase-HMGB1A-R3V6 NPs efficiently reduced the S1Plyase level, the inflammatory response, and apoptosis in lung tissue. These findings confirmed the advantages of the peptide R3V6 over other types of carriers in terms of toxicity and combined delivery of siS1Plyase and HMGB1A\(^{188}\).

Later, Ge et al.\(^{99}\) developed guanidinated and fluorinated bifunctional polypeptides for the lung delivery of TNF-\(\alpha\) siRNA into AMs aiming to simultaneously enhance siTNF-\(\alpha\) polyplexes cell membrane penetration by the guanidinium groups, and enable their transmucous penetration by fluorocarbon modification. Resultantly, the cellular uptake of fluorinated siTNF-\(\alpha\)-loaded cationic polypeptides by macrophages was more effective than free siRNA, nonfluorinated, and PEI-siRNA polyplexes. Remarkably, the fluorine modification of the cationic polypeptides potentiated the mucus permeation by 240 folds. Compatibly, the TNF-\(\alpha\) knockdown of all the fluorinated polyplexes in LPS-activated macrophages was more effective than the nonfluorinated and PEI-siRNA polyplexes. This was confirmed in vivo by a potent TNF-\(\alpha\) silencing, reduced pro-inflammatory cytokines, recovered pulmonary ventilation function, and alleviated tissue damage in the LPS-ALI mice model after intratracheal administration\(^{197}\).

5.2.4. Hybrid nanoparticles (HNPs)

Hybrid lipid-polymer NPs combine both the natural biocompatibility and low immunogenicity of lipids, and the high binding affinity for RNA of polymeric nanocarriers\(^{201,198}\). In this respect, Angelo et al.\(^{194}\) designed hybrid lipid-polymer nanoparticles (size of 150 nm, and zeta potential close to \(-25\) mV) as a pulmonary delivery carrier of siRNA to the human airway EpiCs. They used PLGA and dipalmitoylphosphatidylcholine (DPPC), with or without PEI as the third component in their formulations\(^{195}\). DPPC, the principal lipid constituent of pulmonary surfactant, was expected to moderate HNPs interactions in the airways\(^{195}\), while the inclusion of PEI would likely enhance siRNA entrapment inside the HNPs. Resultantly, the optimized siRNA-loaded HNPs displayed an optimal in vitro aerodynamic behavior after administration with a vibrating mesh nebulizer. They exhibited good stability in artificial mucus, excellent penetration inside extracellular lung barriers, and efficient internalization inside the airway EpiCs without inducing any proinflammatory or cytotoxic effects\(^{196}\).

Recently, Dormenval et al.\(^{196}\) aimed to design physiochemically stable inhalable solid dosage forms for siRNA delivery suitable for large-scale production. Therefore, they prepared spray-dried TNF-\(\alpha\) siRNA-loaded lipid-polymer hybrid nanoparticles (LPNs) supposed to treat chronic obstructive pulmonary disease (COPD)-associated inflammation. The prepared TNF-\(\alpha\) siRNA-loaded LPNs (size of 197.1 nm, PDI of 0.098, and zeta potential of 18.1 mV) were maintained in the same range till 10-fold scaled up formulations. Favorably, the spray drying neither affected the siRNA integrity, nor the in vitro release and thereby efficient gene silencing was maintained\(^{196}\).

Kusumoto et al.\(^{197}\) designed a GALA-modified MEND constituted of an anionic core of siRNA-PEI coated with a cationic liposomal envelope of DOTMA, EPC (egg phosphatidylcholine), and cholesterol. They aimed to study the impact of GALA surface modification which was supposed to enhance the endosomal release and target the pulmonary ECs. GALA/MEND loaded with siRNA against CD31 could efficiently silence the endothelial marker gene CD31 compared to the unmodified MEND indicating the high potency of this system to deliver siRNA to the lung endothelium, one of the promising targets to treat ALI/ARDS\(^{197}\).

Interestingly, Merckx et al.\(^{198}\) designed a hybrid nanoparticle composed of siRNA-loaded cationic dextran nanogel core coated with a shell of the hydrophobic surfactant protein B (SP-B) and incorporated in a phospholipid mixture of DOPC and eggPG (LL-\(\alpha\)-phosphatidylglycerol), or DPPC and eggPG. The proteolipid coated siRNA-loaded nanogels were then administered by tracheal aspiration to silence TNF-\(\alpha\) in LPS-induced ALI murine model. Remarkably, the incorporation of SP-B model in the DPPC:eggPG coated formulation improved cytosolic siRNA delivery to the respiratory immune cells leading to a significant gene silencing compared to other nanocomposites. However, the administration of these nanocomposites to the naïve mice increased the inflammatory markers\(^{199}\). Most recently, the same group has demonstrated that the proteolipid coated siRNA-loaded nanogels can be freeze-dried and then reconstituted before nebulization for inhalation therapy without influencing their integrity or efficiency\(^{198}\).

6. Conclusions and outlook

The field of siRNA therapy is attracting great interest due to its potential therapeutic to silence any gene of interest including the deadliest diseases, such as ALI/ARDS. Despite the extensive studies and the multiple possible targets of ALI/ARDS, there are still no specific pharmacological treatments. To date, the best practice against ALI/ARDS remains protective ventilation that can aggravate lung damage or even cause lung injury. Therefore, gene silencing via siRNA can be implemented to open up new treatment opportunities for ALI/ARDS. Many preclinical reports are supporting the suitability of siRNA-based therapies for ALI/ARDS by recovering the integrity of lung epithelial and/or
endothelial barriers or reinforcing the pulmonary defense mechanisms. However, the majority of these studies focus only on the target of siRNA instead of looking into the delivery perspective. The pulmonary route seems to be the best choice to deliver siRNA into the lung effectively and safely. Despite the advantages of naked siRNA delivery, namely its simplicity and reduced toxicity, the results obtained from the local administration of free siRNA were controversial between different studies. Multiple designed vectors for gene delivery are used in several diseases. However, only some of them have been tested for siRNA delivery against ALI/ARDS. Most of the relevant studies used the available commercial transfection agents such as Lipofectamine, known for their inherent toxicity and instability. Hence, to be feasible and viable in the clinic, the development of an efficient and safe delivery system suitable for the pulmonary delivery of siRNA against ALI/ARDS seems promising. This has been proofed by a lot of recent studies that demonstrated the possibility to maintain the integrity of siRNA while formulating inhalable forms. 

Both viral and non-viral vector systems have been designed or adapted to protect siRNA and deliver it to its cytosolic target after bypassing the different extracellular and intracellular barriers. Given the inherent immunogenicity of viral vectors, non-viral vectors are increasingly used to deliver nucleic acids. Although non-viral delivery systems are less efficient than viral vectors, they are less immunogenic, less expensive, and easier to produce in large amounts. Exosomes have also been proposed to deliver siRNA into the lung as nature’s nano delivery system. However, exosome-mediated delivery is mainly hindered by purification techniques. Comparatively, liposomal delivery systems showed good transfection efficiency, polymer-based nanocarriers displayed sustained release profiles and higher cellular internalization potential, and peptide vectors exhibited enhanced endosomal escape. Interestingly, hybrid nanocarriers seem to be the best delivery system since it allows to combine the advantages of these different carriers into one single vector. Nevertheless, sophisticated approaches usually have characterization and manufacturing scale-up difficulties that may hinder their clinical translation.

Strategies such as the use of a cocktail of siRNAs silencing different mRNAs might be used in combination for targeting different pathways to provide a synergistic silencing effect given the complex pathological mechanisms and pathways involved in ALI/ARDS. Also, the use of targeting peptides to maximize uptake of vectors into alveolar cells or the codelivery of siRNA with other therapeutic agents may further improve the therapeutic outcomes. Innovative nebulization technologies are also needed to improve the delivery of siRNA/carryer to the distal lung while maintaining its integrity. Besides the delivery issues, there still exist many limitations to overcome. These include safety issues, manufacturing scale-up, the development of inhalation devices that can both maintain siRNA integrity and be suitable for patients under mechanical ventilation. Further, the extrapolation of results from animal studies to human applications is another bottleneck regarding the big differences in the airways between rodents and humans. Various injury models should be combined to create a similar human ALI/ARDS model such as cecal ligation and puncture followed by hemorrhage rather than LPS injury alone. Moreover, in some studies, siRNA treatment was given preventively or in the early phase of the disease which is not practical in the clinical application where the diagnosis and treatment of ARDS are delayed. Frequently, the studies assessed only the inflammatory mediators’ levels without histopathological investigations which is an important point to consider especially when using nanoparticles since the inhalation of NPs itself can cause inflammation. Additionally, the fibroproliferation process may present another promising target for siRNA therapy besides the inflammatory process. Remarkably, most of the studies that investigated the efficiency of delivery vectors in ALI/ARDS used limited targets such as anti-TNF-α siRNA. Similarly, studies that investigated the potential therapeutic targets for siRNA in ALI/ARDS did not focus on the delivery strategies. However, for successful outcomes, both the targets and the delivery strategies should be considered while developing a siRNA therapy for ALI/ARDS. Despite the encouraging preclinical results, the authorities do not list any clinical trial on siRNA delivery against ALI/ARDS. Therefore, more work is needed to develop the appropriate carrier delivering siRNA to the right target in the suitable device to see the first inhalable siRNA-based therapy against ALI/ARDS in the near future.

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Author contributions

Wei He conceived and designed the work. Makhloufi Zoulilka, Qingqing Xiao, and George Frimpong Boafo co-wrote the paper. Wei He, Zhongjian Chen, and Marwa A. Sallam commented and corrected the paper. All of the authors have read and approved the final manuscript.

Conflicts of interest

All the authors declare that this article content has no conflict of interest.

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