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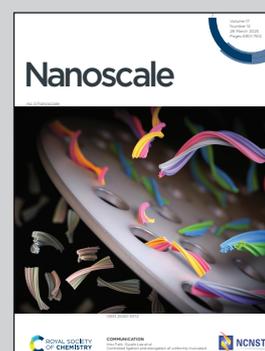
Enhanced antifungal activity of siRNA-loaded anionic liposomes against the human pathogenic fungus *Aspergillus fumigatus*

This study reports the development of siRNA-loaded anionic liposomes, co-encapsulating low-dose amphotericin B, to enhance siRNA penetration through the fungal cell wall of *Aspergillus fumigatus*. Targeting mRNAs of three key genes, these liposomes achieve a clear fungal growth inhibitory effect, demonstrating for the first time the antifungal potential of siRNA against human fungal pathogens.

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In this study, we explored the potential of anionic LIP for delivering siRNA into *A. fumigatus*, inspired by the use of anionic LIP in AmBisome® for delivering amphotericin B, a conventional antifungal drug.¹⁶

Results and discussion

We formulated anionic LIP with the same components as AmBisome®, including DSPG (1,2-distearoyl-*sn*-glycero-3-phosphoglycerol), HSPC (hydrogenated soybean phosphatidylcholine), and cholesterol (Chol; Fig. S1, ESI†) in a molar ratio of 0.8 : 2 : 1. We synthesized LIP *via* the “thin film hydration” method,^{17,18} which involved dissolving lipids in an organic solvent, forming a thin film, rehydrating it to create multilamellar LIP structures, and then converting them to unilamellar structures *via* ultrasound and extrusion (Fig. S2, ESI†).^{18,19} Dynamic light scattering analysis showed that five extrusion cycles through a 100 nm polycarbonate membrane produced LIP with a size of 143 ± 2 nm and a Zetasizer count rate within the recommended range.²⁰ Additional extrusion cycles reduced the size (137 ± 3 nm after 11 cycles and 130 ± 2 nm after 21 cycles) without affecting polydispersity or count rate, leading us to select a maximum of 21 cycles for monodisperse LIP (Table S1, ESI†).

To visualize the LIP, we synthesized fluorescently labeled liposomes (RhoB-LIP) by incorporating 2 mol% rhodamine DHPE (rhodamine B 1,2-dihexadecanoyl-*sn*-glycero-3-phosphoethanolamine, triethylammonium salt; Fig. S1†),^{21–24} which showed no differences in particle size or zeta-potential compared to unlabeled LIP (Fig. S3, ESI†). We co-incubated RhoB-LIP with *A. fumigatus* strain A1160p+ (MFIG001)²⁵ in *Aspergillus* minimal medium (AMM)²⁶ for 16 h at 37 °C. RhoB-LIP accumulated around the hyphal cell wall, but did not enter the fungal cells (Fig. S4, ESI†). This suggested that electrostatic interactions between the positively charged hyphal cell wall¹³ and the anionic LIP were insufficient for LIP entry. This observation aligns with previous findings where anionic LIP were trapped in the cell wall of another human fungal pathogen, *Candida albicans*.²⁷ Interestingly, in this previous study, anionic LIP loaded with amphotericin B (AmBisome®) crossed the cell wall and reached the cell membrane due to amphotericin B (AmB)'s high binding affinity for ergosterol, a critical component of the fungal cell membrane. Specifically, while anionic LIP were confined to the junction between the outer and inner cell wall layers, AmBisome® penetrated the inner β -glucan-chitin layer and reached the cell membrane. Anionic LIP loaded with both AmB and gold particles, which do not deform during passage, also successfully reached the cell membrane. In contrast, gold particles alone could not diffuse through the outer and inner cell wall layers, suggesting that AmB-loaded anionic LIP can displace the polysaccharide chains of the cell wall. Brief exposure to AmBisome® increased the cell wall's relative porosity five-fold without significantly affecting fungal cell viability. Notably, in ergosterol-deficient *Candida* mutants, AmBisome® were

trapped at the junction of the outer and inner cell wall layers, similar to anionic LIP, emphasizing the crucial role of the AmB–ergosterol interaction in this process.²⁷ Inspired by these previous findings, we explored incorporating AmB as an adjuvant into anionic LIP to potentially enable siRNA delivery into fungal cells.

To load siRNA into LIP, we incorporated it into the aqueous rehydration solution, as siRNA is hydrophilic. However, the negative charges of siRNA and phospholipids can cause electrostatic repulsion, impeding efficient encapsulation. We used protamine, a protein rich in arginine residues, which confers a high positive charge, to complex with siRNA and neutralize its negative charge.^{28,29} Protamine sulfate, which is FDA-approved and widely used in medicine, is also a component of neutral protamine Hagedorn (NPH) insulin for long-acting delivery. Its extensive use in diabetic patients has demonstrated minimal toxicity and immunogenicity,³⁰ making it a safe choice for our drug delivery system design. Moreover, protamine sulfate has shown no inhibitory effect on the growth of molds, including *Aspergillus*.³¹ Protamine and siRNA were co-incubated at an N/P ratio of 1 : 1 (N = positively charged amino acids, P = negatively charged phosphate groups on siRNA) before loading into LIP (Fig. S2†). Although free protamine from incomplete incorporation in the siRNA–protamine complex may cause liposome aggregation,³² our experiments with fluorescently labeled non-targeting siRNA (DY547-siRNA; Table S2, ESI†) showed no noticeable changes in LIP characteristics, including size and zeta-potential (Table S3, ESI†). In fact, protamine complexation improved siRNA loading across various batches, compared to those without protamine, resulting in more consistent and higher loading percentages (Table S3†). The concentration of loaded siRNA was quantified using a calibration curve of free DY547-siRNA (Fig. S5, ESI†). Thus, we included protamine as a standard component in the LIP formulation.

Integrating AmB as an adjuvant in siRNA-loaded anionic LIP aimed to keep the AmB content minimal to avoid inhibiting fungal growth. More specifically, binding of AmB with ergosterol in the fungal cell membrane can lead to membrane disruption and ultimately cell death.^{33,34} While leveraging AmB's ability to enhance siRNA penetration into fungal cells, it is crucial to minimize its potential adverse effects on membrane integrity, particularly to avoid off-target interactions of AmB with cholesterol in mammalian cell membranes.³⁵ We added AmB to the initial organic solution in a molar ratio of 0.8 : 2 : 1 : 0.01 (DSPG : HSPC : Chol : AmB). The concentration of encapsulated AmB was determined using high-performance liquid chromatography (HPLC) according to calibration curves of free AmB (Fig. S6, ESI†). To assess AmB's impact on the interaction between siRNA-loaded LIP and *A. fumigatus*, we used DY547-siRNA for visualization. After 16 h of co-incubation with *A. fumigatus*, weak signals were detected on the fungal cell wall with LIP loaded solely with siRNA (LIP \times DY547-siRNA). In contrast, siRNA signals were observed inside the fungal hyphae when LIP were co-loaded with AmB and siRNA (LIP \times AmB \times DY547-siRNA; Fig. S7, ESI†). Importantly,



microscopy revealed that hyphal growth remained largely unaffected by AmB (final concentration: $0.43 \mu\text{g mL}^{-1}$). These findings indicate that the concentration of AmB in this formulation effectively facilitated siRNA delivery but was likely too low to inhibit fungal growth, explaining the lack of visible antifungal effects. In comparison to previous approaches, cationic multilamellar liposomes used for siRNA encapsulation showed no accumulation of siRNA on the hyphal cell wall of *A. fumigatus*,¹⁴ whereas our anionic LIP formulation (LIP \times siRNA) achieved slight accumulation on the hyphal cell wall. Furthermore, using the cationic cell penetrating peptide PAF26 for siRNA conjugation allowed a small amount of PAF26 to penetrate the fungal hyphae, while siRNA was only observed on the fungal cell wall, co-localized with PAF26, indicating that PAF26 cannot translocate siRNA into *A. fumigatus*.¹⁴ In contrast, our LIP \times AmB \times siRNA formulation successfully delivered siRNA into the fungal hyphae, demonstrating the potential of anionic LIP, coupled with low-concentration AmB as an adjuvant, to facilitate siRNA entry into fungal cells.

Encouraged by these results with non-targeting siRNA, we explored functional siRNAs targeting three transcription factor-encoding genes in *A. fumigatus*: *hapB*, *hapX*, and *sreA*. The *hapB* gene encodes a subunit of the CCAAT-binding transcriptional regulatory complex (CBC), regulating over a third of the genome.³⁶ Deletion of *hapB* results in severe growth reduction³⁷ and attenuated virulence.²⁶ The *hapX* gene cooperates with the CBC for adaptation to low iron environments, and its deletion also reduces virulence.^{26,38} While the *sreA* gene is involved in iron regulation³⁹ and is dispensable for virulence, the double deletion of *sreA* and *hapX* is lethal to the fungus.³⁸ Additionally, double deletions of *sreA* with *hapB* or *hapX* are lethal in *Aspergillus nidulans*, closely related to *A. fumigatus*.⁴⁰ These findings led us to target both individual and combined gene silencing for antifungal effects *via* siRNA treatment.

In our initial screening, individual candidate siRNAs were co-incubated with germinated spores of *A. fumigatus* for 24 h at 37 °C in AMM, followed by RNA isolation and quantitative real-time PCR (qRT-PCR) assessment of target mRNA levels, using primers listed in Table S4 (ESI†). We selected siRNAs for each target gene (*hapB*-1, *hapX*-3, and *sreA*-1; Table S2†) that achieved 40–50% mRNA knockdown (Fig. S8A, ESI†) but did not visibly inhibit fungal growth. Given the role of these genes in iron regulation, we tested the effect of switching from iron-replete to iron-depleted conditions after 18 h. This led to a transient and ambiguous delay in fungal growth on day 2 for siRNA-treated samples (Fig. S8B†), confirming the siRNAs' functionality, but without clear growth inhibition.

Next, we co-loaded functional siRNA with AmB into anionic LIP (LIP \times AmB \times siRNA) and co-incubated siRNA targeting *hapB* (*hapB*-1) with germinated spores of *A. fumigatus* for 18 h, followed by a medium shift to AMM-Fe. On day 2, we observed no visible hyphal growth in samples treated with LIP \times AmB \times *hapB*-1, while those treated with scrambled siRNA (LIP \times AmB \times *hapB*-S) or LIP \times AmB showed comparable growth to PBS-treated controls. By day 3, sporulation (generation of green

spores) was evident in controls but not in treated samples, although hyphal growth resumed. By day 4, growth differences disappeared (Fig. 1A). We then tested combinations of individually encapsulated siRNAs (Fig. 1B), finding that dual siRNA treatment extended inhibition to 4 days, and triple siRNA treatment to 5 days. In contrast, scrambled siRNAs or LIP \times AmB showed growth similar to controls. These results suggest that targeting *hapB*, *hapX*, and *sreA* simultaneously is more effective than targeting *hapB* alone or in combination with *hapX* or *sreA*.

Building on these encouraging findings, we tested whether a second dose of siRNAs could prolong antifungal effects. Upon shifting to AMM-Fe, we included a second dose of siRNAs. Two doses of *hapB*-1 with two scrambled siRNAs extended antifungal effects to 5 days, while two doses of all three functional siRNAs achieved effects lasting at least 10 days (Fig. 1C), underscoring the antifungal efficacy of repeated dosing.

In initial experiments, growth differences were noted between samples and controls even before shifting to AMM-Fe. The medium change, involving a centrifugation step, likely introduced bias: shorter hyphae might be removed from samples, while longer hyphae tend to adhere more strongly to the well bottom in controls. Additionally, the medium change could also account for the temporary variations in growth inhibition observed among triplicates (*e.g.*, Fig. 1C, day 6, in the group treated with LIP \times AmB \times *hapB*-1/*hapX*-S/*sreA*-S), as it is difficult to completely prevent the removal of short hyphae from each well, leading to some degree of variation among the triplicates. To minimize bias and variation, we conducted subsequent experiments solely in AMM, despite AMM-Fe better mimicking the *in vivo* iron-restricted conditions encountered by *A. fumigatus*.⁴¹

To better monitor the antifungal effects of siRNAs, we measured the optical density of fungal cultures at 600 nm (OD_{600}).⁴² Following siRNA treatment, OD_{600} at 37 °C was monitored for at least 45 h (Fig. 2A). Controls treated with PBS, LIP \times AmB, and scrambled siRNAs showed detectable fungal growth at 15 h, reaching saturation (OD_{600} above 1.0) between 40–45 h. Treatment with *hapB*-1 and two scrambled siRNAs delayed growth detection to 35 h, while three functional siRNAs extended this to 52 h. Monitoring OD_{600} was extended to 60 h for these two cases (Fig. 2B): in *hapB*-1 and scrambled siRNA treatments, OD_{600} reached 1.0 at 60 h, but stayed below 0.2 with three functional siRNAs, indicating clear fungal growth inhibition even in iron-replete conditions.

In all experiments mentioned above, we had utilized the *A. fumigatus* strain A1160p+. To ensure broad applicability, we tested the efficacy of LIP \times AmB \times siRNA on three clinical isolates: Af293, D141, and ATCC46645, compared to A1160p+. These strains are widely used in research and show varied virulence.⁴³ After siRNA treatment, growth inhibition was consistent across all strains on day 2 (Fig. S9, ESI†). By day 3, while growth resumed in A1160p+ and Af293 targeting *hapB* alone, D141 and ATCC46645 showed minimal growth. Targeting *hapB*, *hapX*, and *sreA* uniformly suppressed hyphal formation



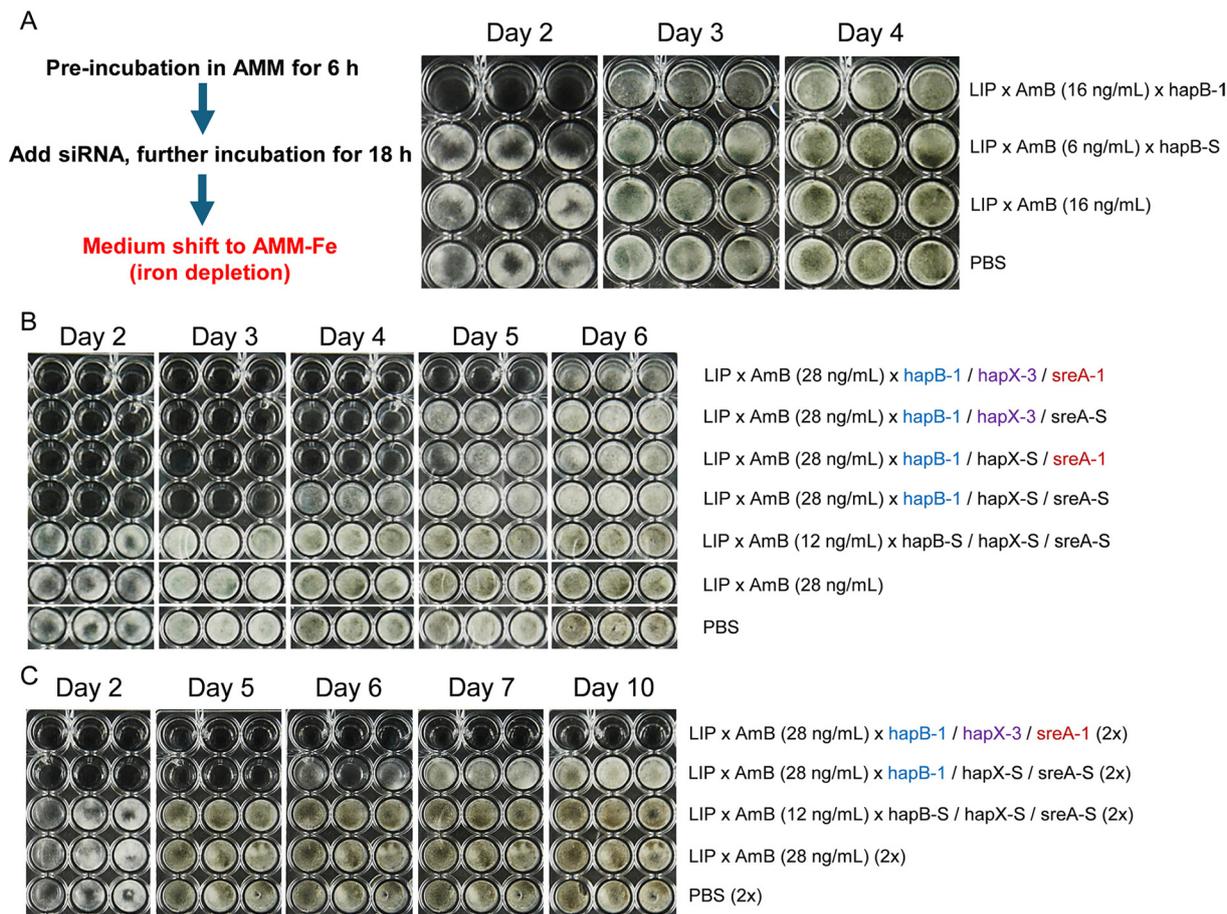


Fig. 1 Antifungal effect of liposomes co-loaded with amphotericin B and functional siRNA (LIP × AmB × siRNA) on *A. fumigatus* strain A1160p+. (A) After 6 h pre-incubation of fungal spores in AMM, LIP × AmB × *hapB-1* (final concentration: approximately 1.5 nM) was added for an 18 h co-incubation, followed by a medium shift to iron-depleted AMM (AMM-Fe). This delayed hyphal growth until day 2 and sporulation until day 3. Equivalent concentrations of scrambled control (LIP × AmB × *hapB-S*) and AmB (LIP × AmB) did not inhibit fungal growth. (B) After 6 h pre-incubation of fungal spores in AMM, LIP × AmB × siRNA formulations (final concentration: approximately 1.5 nM each, 4.5 nM in total) were added for an 18 h co-incubation, followed by a shift to AMM-Fe. Targeting *hapB*, *hapX*, and *sreA* simultaneously prolonged the antifungal effect, compared to targeting *hapB* alone, *hapB* + *hapX*, or *hapB* + *sreA*. (C) Following the same initial procedure, a second dose of siRNAs (final concentration: approximately 1.5 nM each, 4.5 nM in total) was included upon shifting to AMM-Fe. Two doses of siRNAs targeting *hapB*, *hapX*, and *sreA* showed an antifungal effect for at least 10 days.

across all strains, with the effect lasting until day 4 in D141 and ATCC46645 but resuming in A1160p+ and Af293. Overall, our findings confirmed that the LIP × AmB × functional siRNA formulations are effective across various *A. fumigatus* strains, albeit with some variation in the duration of RNA knockdown. This variation likely stems from differing growth rates of the tested strains, as evident on day 2 in the PBS-treated controls (Fig. S7†), where green spores appeared in the wells of A1160p+ and Af293, while only mycelial growth occurred in D141 and ATCC46645.

In summary, co-loading siRNA and AmB into anionic LIP effectively delivers siRNA into fungal cells. Minimal AmB facilitates LIP crossing the fungal cell wall, enabling siRNA entry. Targeting *hapB*, *hapX*, and *sreA* inhibited fungal growth across four *A. fumigatus* strains for 3–4 days under iron-replete conditions, reflecting the temporary nature of RNAi knockdown. This study represents the first *in vitro* demonstration of

siRNA's antifungal potential against *A. fumigatus*, highlighting RNAi-based gene silencing as a promising new strategy for antifungal therapy. Furthermore, our strategy of targeting multiple regulatory genes simultaneously has proven effective, particularly with combinations that have been shown to be lethal to fungi when deleted concurrently.

Further research should explore additional targets to extend the antifungal effect or achieve a fungicidal outcome through extensive gene expression perturbation. One promising avenue involves targeting the 18 transcription factor-encoding genes that play critical roles in *A. fumigatus* pathogenicity.⁴⁴ Our preliminary data suggest that combining multiple target genes could be the most effective strategy to enhance RNAi-based antifungal efficacy. Additionally, due to cytotoxicity concerns with AmBisome® to human cells,⁴⁵ and the potential off-target effects of siRNAs on human genes, which may not be fully mitigated by targeting fungal-specific genes or avoiding hom-



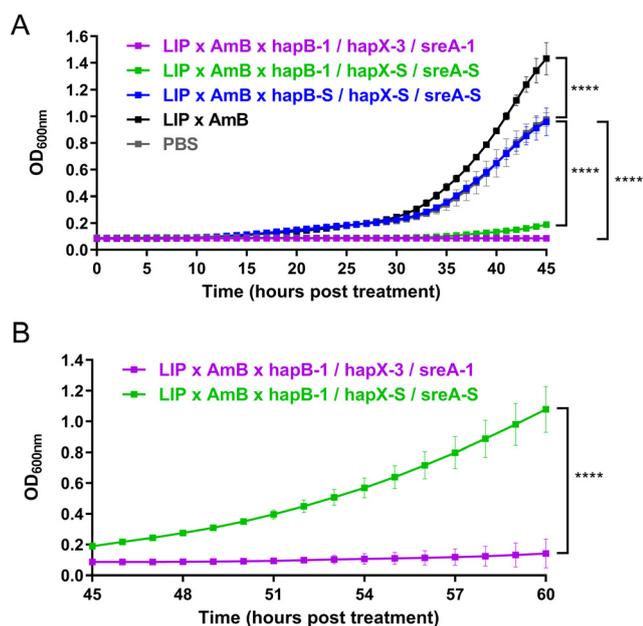


Fig. 2 Antifungal effect of LIP x AmB x siRNA under iron-replete conditions. (A) & (B) After 6 h pre-incubation of fungal spores in AMM, LIP x AmB x siRNA formulations (final concentration: approximately 1.5 nM each, 4.5 nM in total) were added. OD_{600nm} measurements were taken at 37 °C at 10 min intervals (in triplicates). (B) Due to observed delays in fungal growth, measurements were extended to 60 h to compare the effects of siRNAs targeting *hapB*, *hapX*, and *sreA* simultaneously vs. targeting *hapB* alone. A two-way ANOVA with multiple comparisons was performed at each time point against the PBS-treated control group in (A) and between the two groups in (B). Significant differences (****: $p < 0.0001$) are highlighted for the final time points (45 h in A and 60 h in B). No significant difference was found between the PBS group (grey) and the LIP x AmB x *hapB-S/hapX-S/sreA-S* group (blue) at 45 h ($p = 0.67$).

ologous mRNA regions, further optimization of our delivery system is essential. For instance, incorporating a fungal-specific ligand, such as dectin-1 (binds to β -glucans in the fungal cell wall),²¹ could reduce off-target effects of both AmB and siRNAs on host cells, thereby improving the safety profile of RNAi-based antifungal therapies. Moreover, the need for fungal-specific delivery systems extends beyond RNAi to other RNA knockdown strategies, such as antisense oligonucleotides (ASOs) that induce RNase H-dependent mRNA degradation,⁴⁶ and CRISPR-mediated interference (CRISPRi) utilizing dCas9 (a nuclease-deactivated form of CRISPR-associated protein 9) to block transcription by binding temporarily to the gene's regulatory region.⁴⁷ ASO strategies have successfully silenced mRNA in *C. albicans*; however, they exhibit toxicity to mammalian cells at concentrations exceeding 40 nM,⁴⁸ indicating unspecific uptake. Similarly, CRISPRi has been successful in gene knockdown in *C. albicans*, but is predominantly a tool for genetic analysis and requires lithium acetate-based transformation for fungal cell entry.⁴⁹ Given these challenges, our future research will focus on developing fungal-specific siRNA delivery strategies, which could also improve the clinical applicability of the other RNA-targeting approaches mentioned.

Conclusions

Co-loading siRNA and AmB into anionic LIP effectively delivers siRNA into fungal cells. Targeting key fungal regulatory genes with functional siRNAs achieves a clear growth inhibitory effect, demonstrating the potential of this novel antifungal strategy.

Data availability

The data supporting this article have been included as part of the ESI.†

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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