

Rational design of cationic lipids for siRNA delivery

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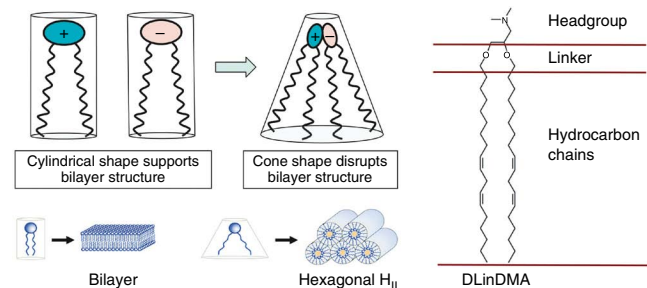
We adopted a rational approach to design cationic lipids for use in formulations to deliver small interfering RNA (siRNA). Starting with the ionizable cationic lipid 1,2-dilinoleyloxy-3-dimethylaminopropane (DLinDMA), a key lipid component of stable nucleic acid lipid particles (SNALP) as a benchmark, we used the proposed *in vivo* mechanism of action of ionizable cationic lipids to guide the design of DLinDMA-based lipids with superior delivery capacity. The best-performing lipid recovered after screening (DLin-KC2-DMA) was formulated and characterized in SNALP and demonstrated to have *in vivo* activity at siRNA doses as low as 0.01 mg/kg in rodents and 0.1 mg/kg in nonhuman primates. To our knowledge, this represents a substantial improvement over previous reports of *in vivo* endogenous hepatic gene silencing.

A key challenge in realizing the full potential of RNA interference (RNAi) therapeutics is the efficient delivery of siRNA, the molecules that mediate RNAi. The physicochemical characteristics of siRNA—high molecular weight, anionic charge and hydrophilicity—prevent passive diffusion across the plasma membrane of most cell types. Therefore, delivery mechanisms are required that allow siRNA to enter cells, avoid endolysosomal compartmentalization and localize in the cytoplasm where it can be loaded into the RNA-induced

silencing complex. To date, formulation in lipid nanoparticles (LNPs) represents one of the most widely used strategies for *in vivo* delivery of siRNA^{1,2}. LNPs represent a class of particles comprised of different lipid compositions and ratios as well as different sizes and structures formed by different methods. A family of LNPs, SNALP^{3–6}, is characterized by very high siRNA-encapsulation efficiency and small, uniformly sized particles, enabled by a controlled step-wise dilution methodology. LNPs, including SNALP, have been successfully used to silence therapeutically relevant genes in nonhuman primates^{6–8} and are currently being evaluated in several clinical trials.

An empirical, combinatorial chemistry-based approach recently identified novel materials for use in LNP systems⁷. A key feature of this approach was the development of a one-step synthetic strategy that allowed the rapid generation of a diverse library of ~1,200 compounds. This library was then screened for novel materials capable of mediating efficient delivery of siRNA *in vitro* and *in vivo*. Here, we instead used a medicinal chemistry (that is, structure-activity relationship) approach, guided by the putative *in vivo* mechanism of action of ionizable cationic lipids, for rational lipid design. Specifically, we hypothesized that after endocytosis, the cationic lipid interacts with naturally occurring anionic phospholipids in the endosomal membrane, forming ion pairs that adopt nonbilayer structures and disrupt membranes (Fig. 1)^{9–12}. We previously advanced the concept

Figure 1 Proposed mechanism of action for membrane disruptive effects of cationic lipids and structural diagram of DLinDMA divided into headgroup, linker and hydrocarbon chain domains. In isolation, cationic lipids and endosomal membrane anionic lipids such as phosphatidylserine adopt a cylindrical molecular shape, which is compatible with packing in a bilayer configuration. However, when cationic and anionic lipids are mixed together, they combine to form ion pairs where the cross-sectional area of the combined headgroup is less than that of the sum of individual headgroup areas in isolation. The ion pair therefore adopts a molecular 'cone' shape, which promotes the formation of inverted, nonbilayer phases such as the hexagonal H_{II} phase illustrated. Inverted phases do not support bilayer structure and are associated with membrane fusion and membrane disruption^{9,21}.



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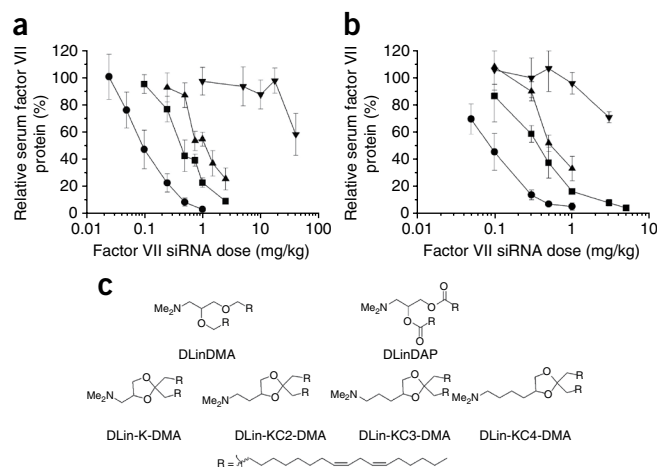


Figure 2 *In vivo* evaluation of novel cationic lipids. (a) Silencing activity of DLinDAP (▼), DLinDMA (▲), DLin-K-DMA (■) and DLin-KC2-DMA (●) screening formulations in the mouse Factor VII model. All LNP-siRNA systems were prepared using the preformed vesicle (PFV) method and were composed of ionizable cationic lipid, DSPC, cholesterol and PEG-lipid (40:10:40:10 mol/mol) with a Factor VII siRNA/total lipid ratio of ~0.05 (wt/wt). Data points are expressed as a percentage of PBS control animals and represent group mean ($n = 5$) \pm s.d., and all formulations were compared within the same study. (b) Influence of headgroup extensions on the activity of DLin-K-DMA. DLin-K-DMA (■) had additional methylene groups added between the DMA headgroup and the ketal ring linker to generate DLin-KC2-DMA (●), DLin-KC3-DMA (▲) and DLin-KC4-DMA (▼). The activity of PFV formulations of each lipid was assessed in the mouse Factor VII model. Data points are expressed as a percentage of PBS control animals and represent group mean ($n = 4$) \pm s.d. (c) Chemical structures of novel cationic lipids.

of using ionizable cationic lipids with pK_a s < pH 7.0 to efficiently formulate nucleic acids at low pH and maintaining a neutral or low cationic surface charge density at pH 7.4 (ref. 13). This strategy should provide better control of the circulation properties of these systems and reduce nonspecific disruption of plasma membranes. As positive charge density is minimal in the blood but increases substantially in the acidic environment of the endosome, this should activate the membrane-destabilizing property of the LNP. Although these attributes may account for the activity of these systems upon internalization by hepatocytes, they do not necessarily explain the high levels of hepatic biodistribution observed for many LNPs, including SNALP. Although these LNPs do not specifically include a targeting ligand to direct them to hepatocytes after systemic administration, it is possible that these LNPs associate with one or more proteins in plasma that may promote hepatocyte endocytosis.

The ionizable cationic lipid DLinDMA has proven to be highly effective in SNALP, has been extensively tested in rodents and non-human primates, and is now being evaluated in human clinical trials. Therefore, we selected it as the starting point for the design and synthesis of novel lipids. We chose the mouse Factor VII model⁷, as the primary *in vivo* screening system to assess functional LNP-mediated delivery to hepatocytes. Briefly, C57BL/6 mice received a single dose of LNP-formulated Factor VII siRNA through bolus tail vein injection and serum was collected from animals 24 h after administration to analyze Factor VII protein level. The initial screening of LNP-siRNA systems was conducted using LNPs prepared by a preformed vesicle method¹⁴ and composed of ionizable cationic lipid, distearoylphosphatidylcholine (DSPC), cholesterol and PEG-lipid (40:10:40:10 mol/mol), with a Factor VII siRNA/total lipid ratio of ~0.05 (wt/wt). Although not a bilayer-destabilizing lipid, a small amount of phosphatidylcholine was incorporated into the LNP to help stabilize the LNP both during formulation and while it was in circulation. A short acyl chain PEG-lipid was incorporated into the LNP to control particle size during formulation, but is designed to leave the LNP rapidly upon intravenous injection. As our goal was to identify novel ionizable cationic lipids for use in LNPs, we aimed to minimize other effects by using a single robust composition and set of formulation conditions suitable for all novel lipids tested. The preformed vesicle method employing the composition listed above provides a convenient platform for such testing, but uses a different formulation process, a different lipid composition and a different siRNA/lipid ratio than SNALP. The structure of DLinDMA can be divided into three main regions: the hydrocarbon chains, the linker

investigate the impact of increasing the number of *cis* double bonds in the hydrocarbon chains found the linoleyl lipid containing two double bonds per hydrocarbon chain (DLinDMA) to be optimal¹⁵. We therefore maintained the linoleyl hydrocarbon chains present in DLinDMA as an element in our lipid design, and focused on optimizing the linker and headgroup moieties.

The linker region in a bilayer structure resides at the membrane interface, an area of transition between the hydrophobic membrane core and hydrophilic headgroup surface. Our approach to linker modification of DLinDMA involved introducing groups expected to exhibit different rates of chemical or enzymatic stability and to span a range of hydrophilicity. A variety of these rationally designed lipids were made, characterized and tested (Supplementary Syntheses 1 and Supplementary Table 1). LNPs based on the ester-containing lipid DLinDAP showed substantially reduced *in vivo* activity compared to LNPs based on the alkoxy-containing lipid DLinDMA (Fig. 2). Further, LNPs based on DLin-2-DMAP, a lipid with one alkoxy linkage and one ester linkage, yielded activity intermediate between DLinDAP- and DLinDMA-based LNPs (Supplementary Table 1). Although it is uncertain why the ester-containing lipids are considerably less active *in vivo*, we speculate that the diester lipid (DLinDAP) is relatively inactive because it is more readily hydrolyzed *in vivo* than the alkoxy analog (DLinDMA), and therefore, unable to either protect the siRNA adequately before release from the endosome and/or survive long enough in the endosome to disrupt the membrane. These hypotheses are being investigated. LNPs based on lipids containing carbamate or thioether linkages also resulted in dramatically reduced *in vivo* activity. Interestingly, the introduction of a ketal ring linker into DLinDMA resulted in LNPs that were ~2.5-fold more potent in reducing serum Factor VII protein levels relative to the DLinDMA benchmark, with an ED₅₀ (that is, dose to achieve 50% gene silencing) of ~0.4 mg/kg versus 1 mg/kg, respectively (Fig. 2).

Given the importance of positive charge in the mechanism-of-action hypothesis guiding the lipid design, the effects of structural changes in the amine-based headgroup were investigated in the context of DLin-K-DMA as the new benchmark lipid. A series of headgroup modifications were made, characterized and tested to explore the effects of size, acid-dissociation constant and number of ionizable groups (Supplementary Syntheses 2 and Supplementary Table 2). Piperazino, morpholino, trimethylamino or bis-dimethylamino modifications tested were not better than the benchmark dimethylamino headgroup of DLin-K-DMA. As an additional parameter, the distance between the dimethylamino group and the dioxolane linker was varied

Table 1 Biophysical parameters and *in vivo* activities of LNPs containing novel lipids

Cationic lipid	Apparent lipid pK _a ^a	L _{II} to H _{II} phase transition temperature (°C) ^b	<i>In vivo</i> ED ₅₀ (mg/kg)
DLinDMA	6.8 ± 0.10	27	~1
DLinDAP	6.2 ± 0.05	26	40–50
DLin-K-DMA	5.9 ± 0.03	19	~0.4
DLin-KC2-DMA	6.7 ± 0.08	20	~0.1
DLin-KC3-DMA	7.2 ± 0.05	18	~0.6
DLin-KC4-DMA	7.3 ± 0.07	18	>3

^apK_a values ± s.d. (n = 3 to 9). ^bL_{II} to H_{II} phase transition was measured at pH 4.8 in equimolar mixtures with DSPS, using differential scanning calorimetric, repeat scans reproducible to within 0.1 °C.

affect both the pK_a of the amine headgroup as well as the distance and flexibility of the charge presentation relative to the lipid bilayer interface. Inserting a single additional methylene group into the headgroup (DLin-KC2-DMA) produced a dramatic increase in potency relative to DLin-K-DMA. The ED₅₀ for this lipid was ~0.1 mg/kg, making it fourfold more potent than DLin-K-DMA and tenfold more potent than the DLinDMA benchmark when compared head-to-head in the Factor VII model (Fig. 2a). Further extension of the tether with additional methylene groups, however, substantially decreased activity, with an ED₅₀ of ~0.6 mg/kg for DLin-KC3-DMA and >3 mg/kg for DLin-KC4-DMA (Fig. 2b).

As changes in lipid design and chemistry may affect the pharmacokinetics, target tissue accumulation and intracellular delivery of LNP formulations, we investigated the relative importance of these parameters on LNP activity at an early stage in this research program. Several of the novel lipids were incorporated into LNP-siRNA formulations containing cyanine dye (Cy3)-labeled siRNA. Plasma, liver and spleen levels of siRNA were determined at 0.5 and 3 h after injection at siRNA doses of 5 mg/kg, and the results are presented in Supplementary Table 3. In general, formulations that were the most active in the mouse Factor VII screens achieved the highest liver levels of siRNA at 0.5 h; however, delivery of siRNA to the target tissue was not the primary factor responsible for activity. This is supported by the observations that most formulations accumulated in the liver and spleen quite quickly and that some formulations with similar liver levels of siRNA had large differences in activity. Moreover, plasma pharmacokinetics alone did not predict activity. For example, although DLin-KC2-DMA and DLinDMA had virtually indistinguishable blood pharmacokinetic profiles in mice (data not shown), the activity of DLin-KC2-DMA in LNPs is approximately tenfold greater than the

same formulation with DLinDMA. Taken together, these results led us to conclude that rapid target tissue accumulation was important, but not sufficient, for activity. Moreover, other parameters were more critical for maximizing the activity of LNP-siRNA formulations.

Two important parameters underlying lipid design for SNALP-mediated delivery are the pK_a of the ionizable cationic lipid and the abilities of these lipids, when protonated, to induce a nonbilayer (hexagonal H_{II}) phase structure when mixed with anionic lipids. The pK_a of the ionizable cationic lipid determines the surface charge on the LNP under different pH conditions. The charge state at physiologic pH (e.g., in circulation) can influence plasma protein adsorption, blood clearance and tissue distribution behavior¹⁶, whereas the charge state at acidic pH (e.g., in endosomes) can influence the ability of the LNP to combine with endogenous anionic lipids to form endosomolytic nonbilayer structures⁹. Consequently, the ability of these lipids to induce H_{II} phase structure in mixtures with anionic lipids is a measure of their bilayer-destabilizing capacity and relative endosomolytic potential.

The fluorescent probe 2-(*p*-toluidino)-6-naphthalene sulfonic acid (TNS), which exhibits increased fluorescence in a hydrophobic environment, can be used to assess surface charge on lipid bilayers. Titrations of surface charge as a function of pH can then be used to determine the apparent pK_a of the lipid in the bilayer (hereafter referred to as pK_a) of constituent lipids¹⁷. Using this approach, the pK_a values for LNPs containing DLinDAP, DLinDMA, DLin-K-DMA, DLin-KC2-DMA, DLin-KC3-DMA and DLin-KC4-DMA were determined (Table 1). The relative ability of the protonated form of the ionizable cationic lipids to induce H_{II} phase structure in anionic lipids was ascertained by measuring the bilayer-to-hexagonal H_{II} transition temperature (T_{BH}) in equimolar mixtures with distearylphosphatidylserine (DSPS) at pH 4.8, using ³¹P NMR¹⁸ and differential scanning calorimetric analyses¹⁹. Both techniques gave similar results.

The data presented in Table 1 indicate that the highly active lipid DLin-KC2-DMA has pK_a and T_{BH} values that are theoretically favorable for use in siRNA delivery systems. The pK_a of 6.4 indicates that LNPs based on DLin-KC2-DMA have limited surface charge in circulation, but will become positively charged in endosomes. Further, the T_{BH} for DLin-KC2-DMA is 7 °C lower than that for DLinDMA, suggesting that this lipid has improved capacity for destabilizing bilayers. However, the data also demonstrate that pK_a and T_{BH} do not fully account for the *in vivo* activity of lipids used in LNPs. For example, although DLin-KC3-DMA and DLin-KC4-DMA have identical pK_a and T_{BH} values, DLin-KC4-DMA requires a more than fivefold higher dose to achieve the same activity *in vivo*. Moreover,

Figure 3 Efficacy of KC2-SNALP in rodents and nonhuman primates. (a) Improved efficacy of KC2-SNALP relative to the initial screening formulation tested in mice. The *in vivo* efficacy of KC2-SNALP (○) was compared to that of the unoptimized DLin-KC2-DMA screening (that is, PFV) formulation (●) in the mouse Factor VII model. Data points are expressed as a percentage of PBS control animals and represent group mean (n = 5) ± s.d. (b) Efficacy of KC2-SNALP in nonhuman primates. Cynomolgus monkeys (n = 3 per group) received a total dose of either 0.03, 0.1, 0.3 or 1 mg/kg siTTR, or 1 mg/kg siApoB formulated in KC2-SNALP or PBS as 15-min intravenous infusions (5 ml/kg) through the cephalic vein. Animals were euthanized 48 h after administration. *TTR* mRNA levels relative to *GAPDH* mRNA levels were determined in

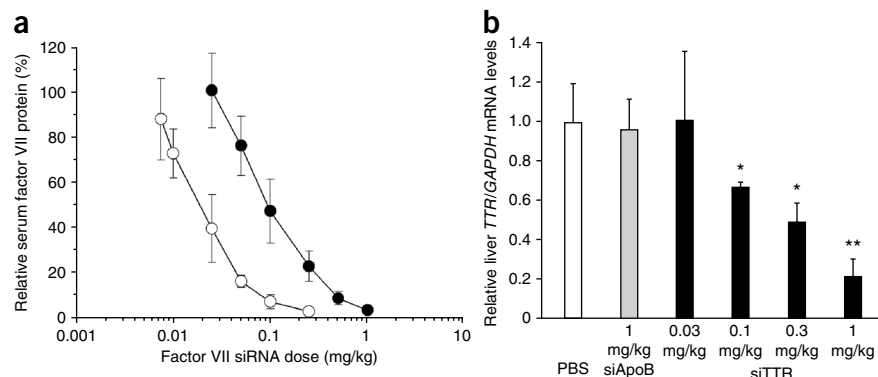


Table 2 Clinical chemistry and hematology parameters for KC2-SNALP-treated rats

Vehicle	siRNA dose (mg/kg) ^a	ALT (U/L)	AST (U/L)	Total Bilirubin (mg/dl)	BUN (mg/dl)	RBC (× 10 ⁶ /μl)	Hemoglobin (g/dl)	WBC (× 10 ³ /μl)	PLT (× 10 ³ /μl)
PBS		56 ± 16	109 ± 31	2 ± 0	4.8 ± 0.8	5.5 ± 0.3	11.3 ± 0.4	11 ± 3	1,166 ± 177
KC2-SNALP	1	58 ± 22	100 ± 14	2 ± 0	4.4 ± 0.6	5.6 ± 0.2	11.6 ± 0.6	13 ± 2	1,000 ± 272
KC2-SNALP	2	73 ± 9	81 ± 10	2.2 ± 0.4	4.3 ± 0.6	5.9 ± 0.3	11.6 ± 0.3	13 ± 4	1,271 ± 269
KC2-SNALP	3	87 ± 19	100 ± 30	2 ± 0	5.0 ± 0.8	6.0 ± 0.2	11.9 ± 0.4	15 ± 2	958 ± 241

^aNontargeting, luciferase siRNA. Sprague-Dawley rats ($n = 5$) received 15-min intravenous infusions of KC2-SNALP formulated siRNA at different dose levels. Blood samples were taken 24 h after administration. ALT, alanine aminotransferase; AST, aspartate aminotransferase; BUN, blood urea nitrogen; RBC, red blood cells; WBC, white blood cells; PLT, platelets.

DLin-KC2-DMA and DLin-KC4-DMA, which have very similar pK_a and T_{BH} values, exhibit a >30-fold difference in *in vivo* activity. This result suggests that other parameters, such as the distance and flexibility of the charged group relative to the lipid bilayer interface, may also be important. Thus, although the biophysical parameters of pK_a and T_{BH} are useful for guiding lipid design, the results presented in **Table 1** support the strategy of testing variants of lead lipids, even ones with very similar pK_a and T_{BH} values.

The lipid composition chosen for the initial formulation and screening of novel ionizable cationic lipids (cationic lipid/DSPC/cholesterol/PEG-lipid = 40:10:40:10 mol/mol, siRNA/total lipid ~ 0.05 wt/wt) was useful for determining the rank-order potency of novel lipids, but is not necessarily optimal for *in vivo* delivery. In addition, the *in vivo* activity of resultant LNP-siRNA formulations is affected by the formulation process employed and the resulting particle structure. Improvements in activity were possible with the preformed vesicle process by modifying and optimizing lipid ratios and formulation conditions (results not shown). However, we chose to further validate DLin-KC2-DMA activity specifically in the context of the SNALP platform, currently the most advanced LNP formulation for delivery of siRNA *in vivo*. We therefore tested *in vivo* a version of SNALP (termed KC2-SNALP), which uses less PEG lipid than reported previously⁶ and in which DLinDMA was replaced with DLin-KC2-DMA. The incorporation of DLin-KC2-DMA into SNALP led to a marked improvement in potency in the mouse Factor VII model; the measured ED_{50} decreased from ~0.1 mg/kg for the unoptimized screening formulation to ~0.02 mg/kg for the KC2-SNALP formulation (**Fig. 3a**). KC2-SNALP also exhibited similar potency in rats (data not shown). Furthermore, after a single administration in rats, KC2-SNALP-mediated gene silencing was found to persist for over 10 d (**Supplementary Fig. 1**).

In addition to efficacy, tolerability is another critical attribute of a suitable LNP-siRNA delivery system for human use. We therefore studied the single-dose tolerability of KC2-SNALP in rats—a popular rodent model for assessing the toxicology of siRNA and nucleic acid-based therapeutics. As doses near the efficacious dose level were found to be very well tolerated (data not shown), single-dose escalation studies were conducted starting at doses ~50-fold higher (1 mg/kg) than the observed ED_{50} of the formulation. To understand formulation toxicity in the absence of any toxicity or pharmacologic effects resulting from target silencing, we conducted the experiments using a nontargeting control siRNA sequence directed against luciferase. KC2-SNALP containing luciferase siRNA was prepared in the exact same manner as that containing Factor VII siRNA, and the resultant size, lipid composition and entrapped siRNA/lipid ratio were similar. Clinical signs were observed daily and body weights, serum chemistry and hematology parameters were measured 72 h after dosing. KC2-SNALP was very well tolerated at the high dose levels examined (relative to the observed ED_{50} dose) with no dose-dependent, clinically significant changes in key serum chemistry or hematology parameters (**Table 2**).

Given the promising activity and safety profile observed in rodents, studies were initiated in nonhuman primates to investigate the translation of DLin-KC2-DMA activity in higher species. For these studies, we chose to target transthyretin (*TTR*), a hepatic gene of high therapeutic interest²⁰. *TTR* is a serum protein synthesized primarily in the liver, and although amyloidogenic *TTR* mutations are rare, they are endemic to certain populations and can affect the peripheral nerves, leading to familial amyloidotic polyneuropathy, and the heart, leading to familial amyloid cardiomyopathy. Currently, the only disease-modifying therapy is liver transplantation. We treated cynomolgus monkeys with a single 15-min intravenous infusion of KC2-SNALP-formulated siTTR at siRNA doses of 0.03, 0.1, 0.3 and 1 mg/kg. Control animals received a single 15-min intravenous infusion of PBS or KC2-SNALP-formulated ApoB siRNA at a dose of 1 mg/kg. Tissues were harvested 48 h after administration and liver mRNA levels of *TTR* were determined. A clear dose response was obtained with an apparent ED_{50} of ~0.3 mg/kg (**Fig. 3b**). A toxicological analysis indicated that the treatment was well tolerated at the dose levels tested, with no treatment-related changes in animal appearance or behavior. No dose-dependent, clinically significant alterations in key clinical chemistry or hematological parameters were observed (**Supplementary Table 4**).

In summary, we applied a rational approach to the design of novel cationic lipids, which were screened for use in LNP-based siRNA delivery systems. Lipid structure was divided into three main functional elements: alkyl chain, linker and headgroup. With DLinDMA as a starting point, the effect of each of these elements was investigated in a systematic fashion, by holding the other two constant. First, the alkyl chains were established, then linker was varied and, finally, different headgroup structures were explored. Using this approach, important structure-activity considerations for ionizable cationic lipids were described and lipids with improved activity relative to the DLinDMA benchmark were identified. A SNALP formulation of the best-performing lipid (DLin-KC2-DMA) was well-tolerated in both rodent and nonhuman primates and exhibited *in vivo* activity at siRNA doses as low as 0.01 mg/kg in rodents, as well as silencing of a therapeutically significant gene (*TTR*) in nonhuman primates. Although the scope of the current work has been limited to hepatic delivery *in vivo*, the *TTR* silencing achieved in this work (ED_{50} ~ 0.3 mg/kg) represents a substantial improvement in activity relative to previous reports of LNP-siRNA mediated silencing in nonhuman primates.

METHODS

Methods and any associated references are available in the online version of the paper at <http://www.nature.com/naturebiotechnology/>.

Note: Supplementary information is available on the Nature Biotechnology website.

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AUTHOR CONTRIBUTIONS

J.C., M.A.C., P.R.C., T.D.M., M.J.H. and K.F.W. designed and advised on novel lipids. J.C., K.F.W. and M.S. synthesized novel lipids. M.J.H., T.D.M., J.C., K.F.W., M.M., K.G.R., M.A.M., M.T. and M.J. analyzed and interpreted lipid data. T.D.M., M.J.H. and M.A.T. co-directed novel lipid synthesis and screening program. S.C.S. designed and directed rodent *in vivo* studies. S.C.S., S.K.K., A.A., B.L.M., I.M., A.P.S., Y.K.T., R.A., T.B., D.W.Y.S. generated rodent *in vivo* data, including Factor VII and tolerability analyses. L.N., V.K., T.B., R.A., Q.C. and D.W.Y.S. developed novel siRNAs targeting *TTR*. R.A. and A.A. designed and directed NHP *in vivo* studies. S.C.S., S.K.K., A.A., B.L.M., I.M., A.P.S., Y.K.T., R.A., T.B., D.W.Y.S., S.A.B., J.Q., J.R.D. and A.D.F. analyzed and interpreted *in vivo* data. B.L.M., K.L., A.P.S., S.K.K., S.C.S. and E.J.C. generated and characterized preformed vesicle formulations with novel lipids. D.S. and C.K.C. developed methods and designed and conducted HPLC lipid analyses of preformed vesicle formulations. E.Y. and L.B.J. prepared SNALP formulations. P.R.C. directed biophysical studies and advised on methods. A.P.S., I.M.H., S.D. and K.W. performed biophysical characterization studies (pK_a , NMR, differential scanning calorimetric) of novel lipids and formulations. M.J.H., P.R.C., T.D.M., A.P.S., I.M.H. and K.F.W. analyzed biophysical data. S.C.S., M.J.H., A.A. and P.R.C. co-wrote the manuscript. T.D.M., M.M., M.A.M., M.A.T. and A.D.F. reviewed and edited the manuscript. S.C.S., M.J.H., A.A., P.R.C., I.M. and A.D.F. were responsible for approval of the final draft.

COMPETING INTERESTS STATEMENT

The authors declare competing financial interests: details accompany the full-text HTML version of the paper at <http://www.nature.com/naturebiotechnology/>.

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- de Fougerolles, A.R. Delivery vehicles for small interfering RNA *in vivo*. *Hum. Gene Ther.* **19**, 125–132 (2008).
- Whitehead, K.A., Langer, R. & Anderson, D.G. Knocking down barriers: advances in siRNA delivery. *Nat. Rev. Drug Discov.* **8**, 129–138 (2009).
- Judge, A.D. *et al.* Confirming the RNAi-mediated mechanism of action of siRNA-based cancer therapeutics in mice. *J. Clin. Invest.* **119**, 661–673 (2009).

- Judge, A.D. *et al.* Sequence-dependent stimulation of the mammalian innate immune response by synthetic siRNA. *Nat. Biotechnol.* **23**, 457–462 (2005).
- Morrissey, D.V. *et al.* Potent and persistent *in vivo* anti-HBV activity of chemically modified siRNAs. *Nat. Biotechnol.* **23**, 1002–1007 (2005).
- Zimmermann, T.S. *et al.* RNAi-mediated gene silencing in non-human primates. *Nature* **441**, 111–114 (2006).
- Akinc, A. *et al.* A combinatorial library of lipid-like materials for delivery of RNAi therapeutics. *Nat. Biotechnol.* **26**, 561–569 (2008).
- Frank-Kamenetsky, M. *et al.* Therapeutic RNAi targeting PCSK9 acutely lowers plasma cholesterol in rodents and LDL cholesterol in nonhuman primates. *Proc. Natl. Acad. Sci. USA* **105**, 11915–11920 (2008).
- Hafez, I.M., Maurer, N. & Cullis, P.R. On the mechanism whereby cationic lipids promote intracellular delivery of polynucleic acids. *Gene Ther.* **8**, 1188–1196 (2001).
- Xu, Y. & Szoka, F.C. Jr. Mechanism of DNA release from cationic liposome/DNA complexes used in cell transfection. *Biochemistry* **35**, 5616–5623 (1996).
- Zelphati, O. & Szoka, F.C. Jr. Mechanism of oligonucleotide release from cationic liposomes. *Proc. Natl. Acad. Sci. USA* **93**, 11493–11498 (1996).
- Torchilin, V.P. Recent approaches to intracellular delivery of drugs and DNA and organelle targeting. *Annu. Rev. Biomed. Eng.* **8**, 343–375 (2006).
- Semple, S.C. *et al.* Efficient encapsulation of antisense oligonucleotides in lipid vesicles using ionizable aminolipids: formation of novel small multilamellar vesicle structures. *Biochim. Biophys. Acta* **1510**, 152–166 (2001).
- Maurer, N. *et al.* Spontaneous entrapment of polynucleotides upon electrostatic interaction with ethanol-destabilized cationic liposomes. *Biophys. J.* **80**, 2310–2326 (2001).
- Heyes, J., Palmer, L., Bremner, K. & MacLachlan, I. Cationic lipid saturation influences intracellular delivery of encapsulated nucleic acids. *J. Control. Release* **107**, 276–287 (2005).
- Semple, S.C., Chonn, A. & Cullis, P.R. Interactions of liposomes and lipid-based carrier systems with blood proteins: Relation to clearance behaviour *in vivo*. *Adv. Drug Deliv. Rev.* **32**, 3–17 (1998).
- Bailey, A.L. & Cullis, P.R. Modulation of membrane fusion by asymmetric transbilayer distributions of amino lipids. *Biochemistry* **33**, 12573–12580 (1994).
- Cullis, P.R. & de Kruijff, B. The polymorphic phase behaviour of phosphatidyl-ethanolamines of natural and synthetic origin. A ^{31}P NMR study. *Biochim. Biophys. Acta* **513**, 31–42 (1978).
- Epand, R.M., Robinson, K.S., Andrews, M.E. & Epand, R.F. Dependence of the bilayer to hexagonal phase transition on amphiphile chain length. *Biochemistry* **28**, 9398–9402 (1989).
- Sekijima, Y., Kelly, J.W. & Ikeda, S. Pathogenesis of and therapeutic strategies to ameliorate the transthyretin amyloidoses. *Curr. Pharm. Des.* **14**, 3219–3230 (2008).
- Cullis, P.R., Hope, M.J. & Tilcock, C.P. Lipid polymorphism and the roles of lipids in membranes. *Chem. Phys. Lipids* **40**, 127–144 (1986).

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