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Note

siRNA LNCs – A novel platform of lipid nanocapsules for systemic siRNA administration

Stephanie David^{a,b,c}, Pauline Resnier^{a,b}, Alexis Guillot^{a,b}, Bruno Pitard^c, Jean-Pierre Benoit^{a,b}, Catherine Passirani^{a,b,*}

^aLUNAM Université, Université d'Angers, Angers, France

^bINSERM U1066 MINT, Micro et Nanomedecines biomimétiques, Angers, France

^cINSERM, UMR915, Université de Nantes, Nantes, France

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ABSTRACT

Several siRNA (small interfering RNA) therapeutics are undergoing clinical trials for cancer, respiratory diseases or macular degeneration, but most are administrated locally. In order to overcome the different barriers to attain an efficient siRNA action after systemic administration, nanocarriers able to carry and protect siRNA are awaited. With this aim, we developed a new platform of siRNA lipid nanocapsules (LNCs) using different cationic lipids, combining the properties of LNCs (siRNA protection and targeting) and lipoplexes (efficient siRNA delivery into the cell). The formulation was revealed to contain different compartments. A siRNA quantification method based on UV spectroscopy was developed to locate and quantify siRNA in each compartment. All in all, these novel siRNA LNCs presented sizes of about 55 nm with a neutral surface charge and siRNA encapsulation efficiencies up to 65% representing appropriate characteristics for systemic administration.

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1. Introduction

siRNA is rapidly broken down by nucleases and other blood components after intravenous administration. Moreover, between the administration and action site, different barriers have to be overcome. Therefore, different siRNA modifications and siRNA vectors have already been described in the literature, with a view to finding the “ideal” vector [1].

Lipid nanocapsules (LNCs), consisting of a lipid liquid core of triglycerides and a rigid shell of lecithin and polyethylene glycol, were developed in our laboratory [2]. The simple formulation process is based on phase inversions of an emulsion. These LNCs were recently modified to encapsulate DNA, complexed with cationic lipids, forming lipoplexes. These DNA LNCs were efficient for *in vitro* and *in vivo* transfection [3–5] in contrast to lipoplexes that were only efficient *in vitro*.

The association between LNCs and lipoplexes should combine LNC properties (nucleic acid protection, prolonged circulation time, possibility of active or passive targeting) with lipoplex properties (internalisation in cells, permitting nucleic acid action). With this in mind, the objective of this work was to develop siRNA LNCs allowing efficient systemic siRNA administration. In this study,

various ratios of different cationic lipids were used to formulate siRNA LNCs, and the size, charge and payload characteristics of these novel promising nanocarriers were assessed.

2. Materials and methods

2.1. siRNA LNC formulation

Basic lipid nanocapsules (LNCs) were formulated, as described before [2], by mixing 20% w/w Labrafac WL 1349 (caprylic-capric acid triglycerides, Gatefossé SA, Saint-Priest, France), 1.5% w/w Lipoid S75-3 (Lipoid GmbH, Ludwigshafen, Germany), 17% w/w Solutol HS 15 (BASF, Ludwigshafen, Germany), 1.8% w/w NaCl (Prolabo, Fontenay-sous-Bois, France) and 59.8% w/w water (obtained from a Milli-Q system, Millipore, Paris, France) together under magnetic stirring. Three temperature cycles between 60 and 95 °C were performed to obtain phase inversions of the emulsion obtained after mixing all the components. Then, a rapid cooling and dilution with ice-cooled water (1:1.4) at the phase inversion temperature (PIT) led to LNC formation.

To obtain siRNA LNCs, the water in the last step was replaced by lipoplexes which were prepared by adding equal volumes of siRNA (here a model siRNA targeted against PCSK9, (sense sequence: GGAAGAUCAUAAUGGACAGdTdT) Eurogenetec, Seraing, Belgium) and liposomes in a defined charge ratio of cationic lipid charge

* Corresponding author. Inserm U646, Université d'Angers, IBS-CHU, 4 rue Larrey, 49933 Angers Cedex 9, France. Tel.: +33 244 688534; fax: +33 244 688546.

E-mail address: catherine.passirani@univ-angers.fr (C. Passirani).

versus anionic siRNA charge. NaCl was added during preparation to obtain a final concentration of 0.15 M.

For liposome preparation, a cationic lipid DOSP (dioleylamine-succinyl paromomycin) (synthesis previously described in [6]), bis(guanidinium)-tris(2-aminoethyl)amine-cholesterol (BGTC) (synthesis previously described in [7]) or DOTAP (1,2-dioleyl-3-trimethylammoniumpropane) (Avanti® Polar Lipids Inc., Alabaster, AL, USA), solubilised in chloroform, was weighted in the ratio 1/1 (M/M), 3/2 (M/M) or 1/1 (M/M), respectively, with the neutral lipid 1,2-dioleyl-sn-glycero-3-phosphoethanolamine (DOPE) (Avanti® Polar Lipids Inc., Alabaster, AL, USA) to obtain a final concentration of 20 mM of cationic lipid charge, considering the number of lipid charges per molecule (4 for DOSP, 2 for BGTC and 1 for DOTAP). After chloroform evaporation under vacuum, deionised water was added to hydrate the lipid film overnight at 4 °C which was sonicated the next day.

DOSP micelles were prepared in the same way without the addition of DOPE.

3. Characterisation

3.1. Size and zeta potential measurements

Size and zeta potential of siRNA LNCs were measured using a Malvern Zetasizer® (Nano Series ZS, Malvern Instruments SA, Worcestershire, UK) at 25 °C, in triplicate, after dilution in a ratio of 1:100 with deionised water.

3.2. Agarose gel electrophoresis

To verify siRNA encapsulation in LNC, Triton® X100 (Sigma, Saint-Quentin Fallavier, France) was added to destroy LNCs. Samples were mixed with OrangeBlue loading dye (Promega, Madison, WI, USA) before deposition on 1% agarose gel containing ethidium bromide (Sigma, Saint-Quentin Fallavier, France) and migration at 100 V for 30 min.

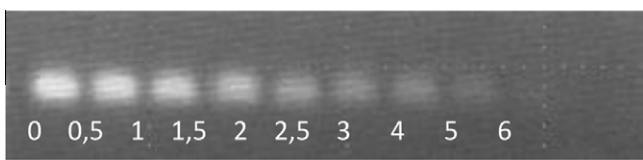
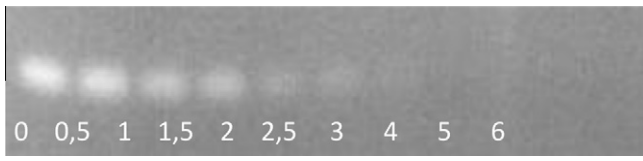


3.3. siRNA LNC purification

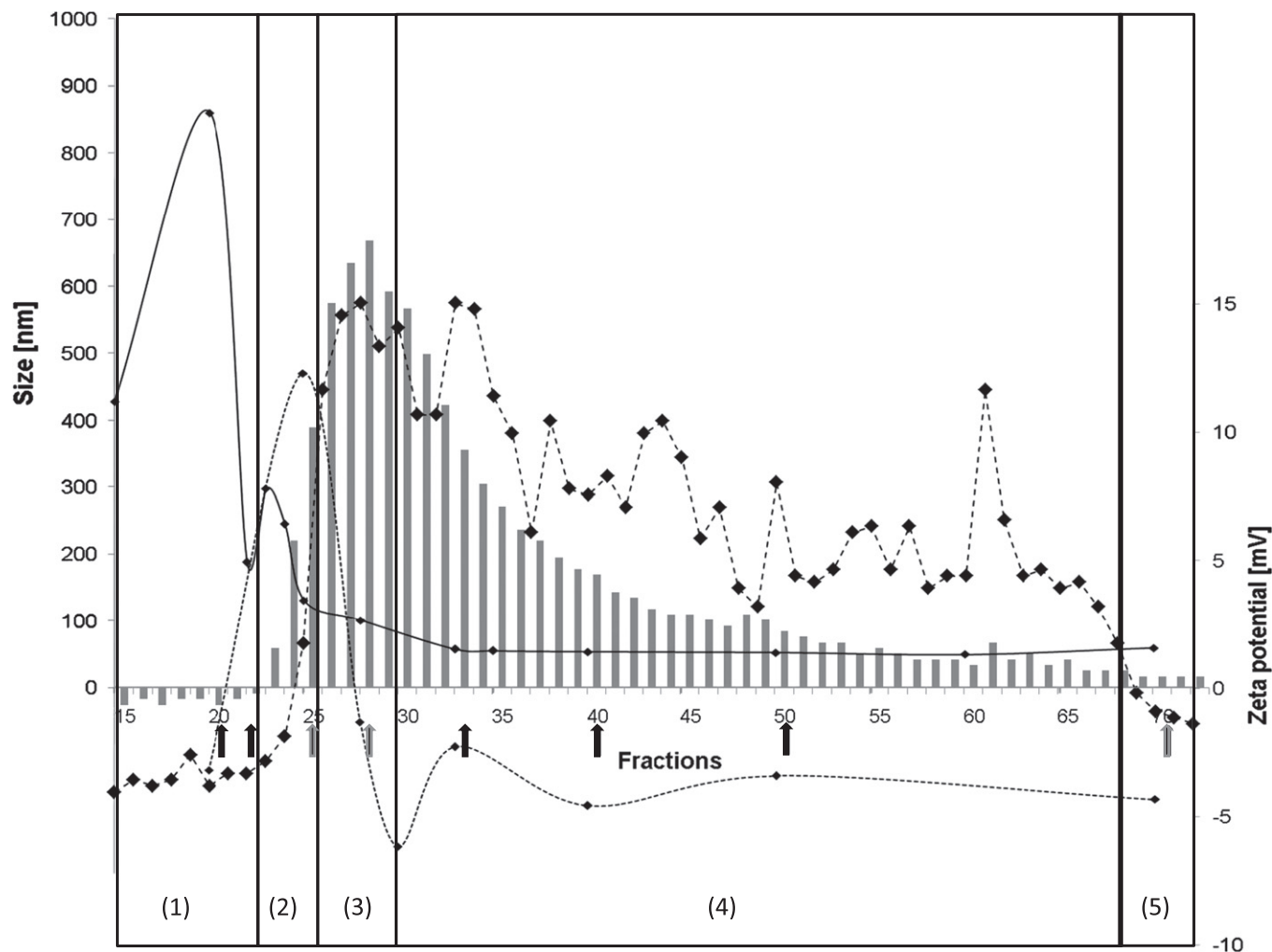
Five hundred microlitre of siRNA LNC formulation was deposited on a 1.5 × 40 cm Sepharose CL-4B column and were eluted with HEPES buffer (pH 7.4). Fractions of 500 µl or 1 ml were collected in glass tubes for further analysis; 100 µl of each fraction was used for turbidity measurements and analysed at 580 nm. PEG (polyethylene glycol) was quantified using 20 µl of each fraction, which was mixed with 5 µl KI/I₂ and 180 µl H₂O milli-Q before analysing at 492 nm using a Multiskan Ascent microplate reader (Thermo Fisher Scientific Cergy-Pontoise, France). Size and zeta potential measurements were taken as described above, and siRNA was evidenced using gel electrophoresis experiments.

3.4. siRNA quantification

One volume of the formulation was mixed with three volumes water (obtained from a Milli-Q-plus® system, Millipore, Paris, France), six volumes 1 M NaOH and two volumes chloroform, vortexed and immediately centrifuged for 15 min at 20,000 g and 4 °C. The aqueous phase, containing free siRNA and siRNA liberated from lipoplexes outside LNCs, was removed and analysed with a UV spectrophotometer (UVIKON 922, Kontron Instruments, Munich, Germany) at 260 nm. The volume removed for free siRNA quantification was replaced by ethanol. Then two volumes water and 10 volumes NaOH 1 M were added before vortexing and centrifuged a second time for 15 min at 20,000 g and 4 °C. The aqueous phase, containing the liberated siRNA from lipoplexes inside siRNA LNCs, was removed and analysed as previously mentioned at 260 nm. To analyse free and encapsulated siRNA quantity in siRNA LNCs, the same procedure was used using two volumes of formulation and replacing water by 1 M NaOH. The first aqueous phase contained free siRNA; the second aqueous phase contained siRNA encapsulated in siRNA LNCs. The siRNA quantity was calculated using a calibrating curve with different siRNA concentrations and compared to the total siRNA amount encapsulated in theory in siRNA LNCs. All samples were prepared in duplicate.

Table 1
Characteristics of siRNA lipoplexes used for siRNA LNC formulations.

Electrophoresis experiments	Lipid	CR	Mean diameter (nm)	Zone
	DOSP	2	1818	B
		2.5	1423	B
		3	1858	B
		4	1122	B
		5	605	C
	DOSP/DOPE	2	2229	B
		2.5	2026	B
		5	234	C
	BGTC/DOPE	2.5	1314	B
		3	770	B
		4	344	C
	DOTAP/DOPE	2.5	1110	B
		5	680	C



		(1) lipo	(2) lpx	(3) siLNC	(4) LNC/PEG	(5) siRNA
Size	—	850 nm	200 - 300 nm	100 nm	50 nm	50 nm
Zeta potential	---	-	5 – 10 mV	-5 – 5 mV	-5 mV	-5 mV
Turbidity		-	+	+	+	-
PEG	--	-	-	+	+	-
siRNA	↑	-	+	+	-	+
Conclusion		empty liposomes	Lipoplexes	siRNA LNCs	empty LNCs/ free PEG	free siRNA

Fig. 1. Purification of DOTAP/DOPE/siRNA LNC formulation (CR 2.5) on Sepharose columns.

4. Results and discussion

4.1. Lipoplex characterisation

The colloidal stability of lipoplexes was determined at different charge ratios (CR = cationic lipid charge/siRNA charge (+/-)) for various cationic lipids (DOSP, DOSP/DOPE, BGTC/DOPE and DOTAP/DOPE) using size, fluorescence and electrophoresis analysis (Table 1). In the way as for DNA lipoplexes, three different zones of colloidal stability A, B and C could be determined with specific properties in each zone [8]. In summary, in zone A, the quantity of cationic lipid is not sufficient to complex all nucleic acids, so they

are still detectable. In this zone, lipoplexes possess a small size and a negative surface charge. In zone B, all nucleic acids are complexed and are no longer detectable. The surface charge is neutral, which leads to aggregation and an augmented size. This was the case for all lipoplexes tested at CR 2 and 2.5, for DOSP/siRNA lipoplexes at CR 3 and 4 and for BGTC/DOPE/siRNA lipoplexes at CR 3. They presented a size over the arbitrary value of 700 nm and only a slight fluorescence in electrophoresis experiments. In zone C, the quantity of cationic lipids is predominant, and thus, the surface charge is positive leading to repulsion and smaller lipoplexes, which was observed for all lipoplexes tested at the CR 5 and for BGTC/DOPE/siRNA lipoplexes at CR 4. To encapsulate lipoplexes in the

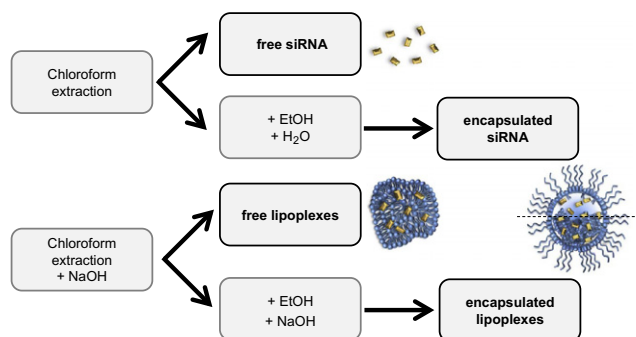


Fig. 2. Schematic representation of siRNA quantification. (For interpretation of the references to colour in this figure legend, the reader is referred to the web version of this article.)

liquid lipid core of LNCs, lipoplexes of zone B and C were chosen as, in zone B, the neutral surface charge, and the aggregation of lipoplexes suggests rather a lipophilic affinity; moreover, DNA lipoplexes of zone C were already encapsulated efficiently in LNCs.

5. siRNA LNC formulation and characterisation

Basic LNCs were developed by Heurtault et al. [2]. Their easy formulation process and their good characterisation make these nanocarriers attractive. To formulate them, the components were mixed together, and temperature cycles around the phase inversion temperature (PIT) were performed. In the last step, a large quantity of water was added at the PIT of the last cycle, resulting in the rapid cooling and dilution of the emulsion and the formation of LNCs. The addition of lipoplexes in the cooling water at the end of the formulation process contributed to siRNA LNC formulation and had the advantage of avoiding their exposure to high temperatures during the formulation.

siRNA encapsulation in LNCs was checked using electrophoresis experiments. As sometimes a low fluorescence band was visible, indicating an incomplete complexation or encapsulation, purification of siRNA LNCs DOTAP/DOPE CR 2.5 was performed on Sepharose columns. This also made it possible to determine whether siRNA was encapsulated in LNCs or simply complexed in lipoplexes. The collected fractions were then analysed by performing size and zeta potential measurements, PEG quantification, turbidity measurements and gel electrophoresis. The final result of these analyses (Fig. 1) suggested the presence of five different compartments in the formulation: (1) empty liposomes, (2) lipoplexes, (3) siRNA LNCs, (4) empty LNCs and (5) free siRNA. In the first com-

partment, empty liposomes were suspected as no PEG and no siRNAs were found and a size of about 850 nm was measured. In the second compartment, the size diminished, siRNAs were detected, the turbidity augmented and a slightly positive zeta potential was measured, indicating the presence of DOTAP/DOPE lipoplexes (CR 2.5). In the third compartment, PEG and siRNA were detected with a size of about 100 nm, and a near neutral zeta potential was measured, suggesting the presence of siRNA LNCs. In the fourth compartment, PEG but no siRNA was detected, and sizes about 50 nm and a slightly negative zeta potential were measured, indicating empty LNCs and/or free PEG. In the last compartment, no PEG was detected, but siRNAs were found, indicating free siRNA.

According to these results, our hypothesis is that siRNA can be situated in 4 different compartments: in the second compartment in lipoplexes outside LNCs; in the third compartment either (3a) complexed with cationic lipids in LNCs or (3b) associated with other components of the LNC formulation but dissociated from cationic lipids; or in the fifth compartment as free siRNA. To quantify and locate siRNA in the formulation, a quantification method using UV spectrometry analysis at 260 nm was developed (Fig. 2). Free siRNA was quantified using a simple chloroform extraction to separate other constituents of the formulation which interfered during UV spectroscopy analysis. siRNAs in lipoplexes outside LNCs were quantified in the same way, but NaOH was added to dissociate lipoplexes. siRNAs in LNCs were quantified using the “separated” formulations (without free siRNA and/or without lipoplexes) by addition of ethanol, to dissociate LNCs, and NaOH, to dissociate siRNAs from cationic lipids. To determine the siRNA quantity, calibrating curves for the different treatments were performed using siRNA quantities between 0 and the maximal theoretical charge in one volume siRNA LNC formulation ($c = 0.25$ g/l).

Afterwards, siRNA LNCs were formulated with different cationic lipids, and their size, zeta potential and encapsulation efficiency were determined (Table 2). The basic LNC formulation presented a size of 55 nm with limited distribution indicated by a very low polydispersity index (PDI) of 0.04 and a slightly negative zeta potential of about -5 mV. The zeta potential of formulations prepared by the novel process was comprised between -25 and $+25$ mV, and the encapsulation efficiencies in LNCs were situated between 4% and 65%. Both varied in relation to the cationic lipids used for lipoplex formation and the CR. The sizes were around 55 nm with a very low PDI with the exception of siRNA LNCs using DOTAP/DOPE/siRNA lipoplexes. Indeed, at CR 2.5, they showed two distinct populations of about 55 and 185 nm, and at CR 5, a size of 125 nm with the highest siRNA encapsulation efficiency at 65%. These lipids and CR were also previously used for DNA encapsula-

Table 2
Size, zeta potential measurements and encapsulation efficiency of siRNA LNC formulations.

LNC formulation	Size (nm)	PDI	Zeta potential (mV)	Encapsulation efficiency (%)				Total siRNA found (%)		
				siLNC (3a + 3b)						
Basic LNC formulation	55.5 ± 0.5	0.04 ± 0.01	-5.2 ± 0.5	Compartment 1						
LNC siRNA				(5) siRNA	(2) Lpx	(3a ²) siLNC	(3b ³) siLNC			
Cationic lipids	CR [±]									
BGTC/DOPE	2.5	55.3 ± 0.1	0.13 ± 0.03	1.5 ± 0.7	20 ± 0	66 ± 3	3 ± 1	10 ± 4	13 ± 3	108 ± 4
DOSP	2	51.6 ± 0.2	0.07 ± 0.00	-2.4 ± 0.7	2 ± 1	94 ± 5	0	4 ± 6	4 ± 6	107 ± 7
	2.5	60.4 ± 1.0	0.10 ± 0.02	-1.1 ± 0.1	8 ± 9	75 ± 18	13 ± 9	4 ± 1	17 ± 10	117 ± 14
	5	52.8 ± 0.0	0.08 ± 0.00	24.1 ± 1.7	20 ± 12	65 ± 13	11 ± 1	3 ± 0	14 ± 1	126 ± 8
				10.1 ± 3.2						
				-3.7 ± 4.3						
DOSP/DOPE	2	53.3 ± 0.2	0.03 ± 0.01	-25.7 ± 1.9	4 ± 1	80 ± 8	8 ± 12	7 ± 2	15 ± 10	75 ± 9
	2.5	59.7 ± 0.5	0.09 ± 0.01	-1.5 ± 0.4	2 ± 2	91 ± 0	2 ± 2	5 ± 0	7 ± 2	88 ± 2
	5	61.4 ± 0.3	0.15 ± 0.03	7.8 ± 0.9	12 ± 4	73 ± 1	11 ± 4	3 ± 0	15 ± 5	105 ± 6
DOTAP/DOPE	2.5	53.5 ± 6.8	0.29 ± 0.03	-0.3 ± 0.5	0	73 ± 19	11 ± 16	16 ± 3	27 ± 19	100 ± 5
		184.1 ± 28.4								
	5	124.8 ± 0.6	0.24 ± 0.00	1.6 ± 0.3	0	34 ± 1	20 ± 4	45 ± 5	65 ± 2	79 ± 7

tion in DNA LNCs forming efficient nanocarriers for *in vitro* [3] and *in vivo* [4] transfection. The size of DOTAP/DOPE siRNA LNCs (CR 5) was similar to that of other promising siRNA vectors (100–170 nm), with efficient transfection *in vivo* after systemic administration [9]. However, the first siRNA nanocarriers for systemic administration that are in phase I clinical trials (CALAA-01 [10]) have sizes of about 70 nm. This corresponds to siRNA LNC formulations containing the other lipids, but in contrast, they present only half or lower encapsulation efficiencies.

In summary, we developed a new platform of siRNA LNCs showing appropriate characteristics for systemic administration with good encapsulation efficiencies and/or small sizes and neutral surface charges. However, further investigations need now to be carried out to test their transfection efficiency *in vitro* and *in vivo* and to find a compromise between a small size and a high siRNA payload.

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