

Liposomes and liposomal structures as tools Název: for labeling and screening of nucleic acids

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Datum: 19.4.2013

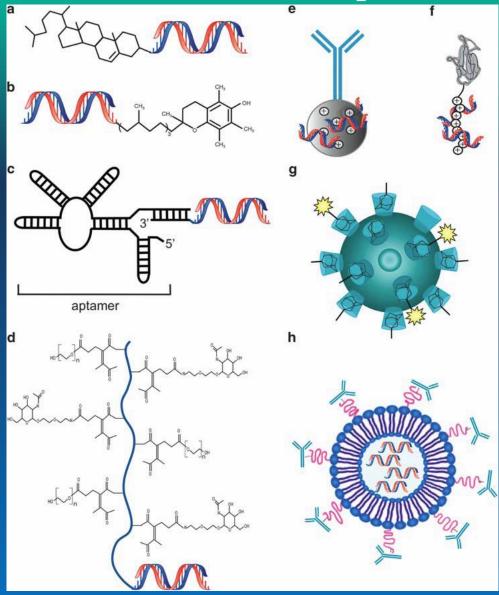
Reg.č.projektu: CZ.1.07/2.3.00/20.0148

Název projektu: Mezinárodní spolupráce v oblasti "in vivo" zobrazovacích technik

Outline:

- Liposomes, its structure, composition
- Preparation routes
- Liposome modification
- Cationic liposomes
- Conclusion

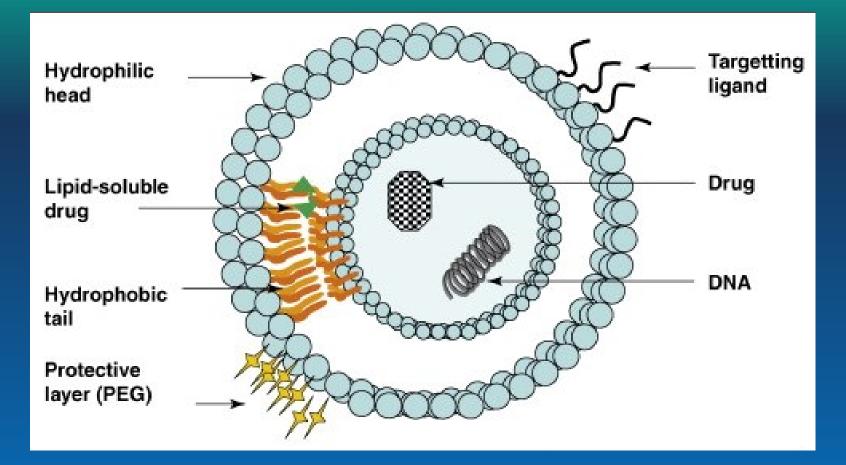
Nanotransporters of RNA



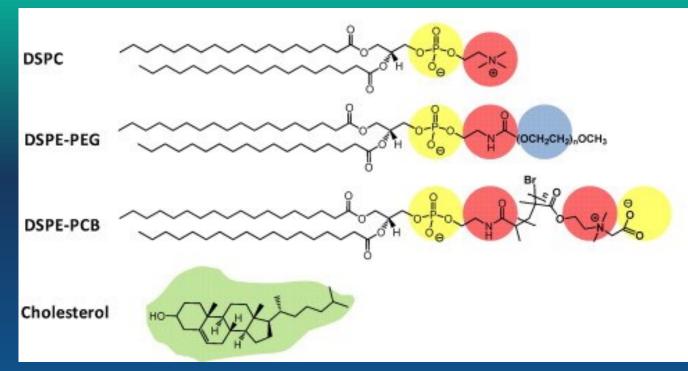
a) siRNAs that are directly conjugated to cholesterol b) small targeting molecules c) aptamer d) membrane-penetrating polymers linked to targeting small molecules e) complexed with fusion proteins composed of an antibody fragment or targeting peptide linked to an RNA-binding domain that is either protamine f) polyarginine g) encapsulated within nanoparticles h) liposomes

Peer D. Gene Therapy (2011) 18, 1127–1133

Liposome Nanotransporters



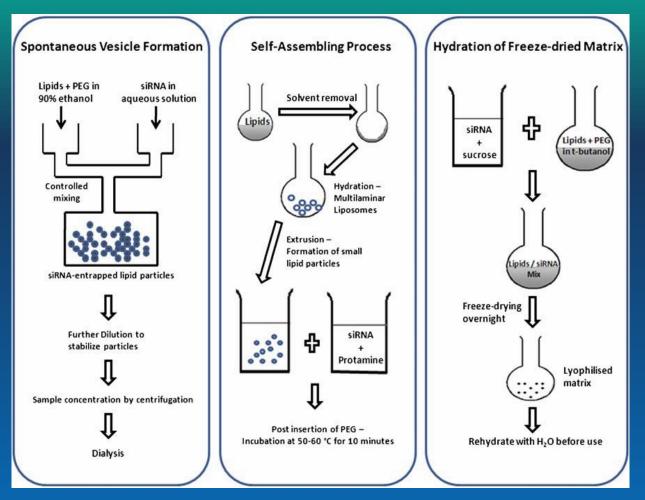
Liposome composition



- 1,2-distearoyl-sn-glycero-3-phosphatidylcholine (DSPC)
- 1,2-diarachidoyl-sn-glycero-3-phosphatidylcholine (DAPC)
- 1,2-dibehenoyl-sn-glycero-3-phosphatidylcholine (DBPC)
- 1,2-distearoyl-snglycero-3-phosphatidylethanolamine (DSPE)
- 1,2-dioleoyl-sn-glycero-3-phospho-rac-(1-glycerol) sodium salt (DOPC)

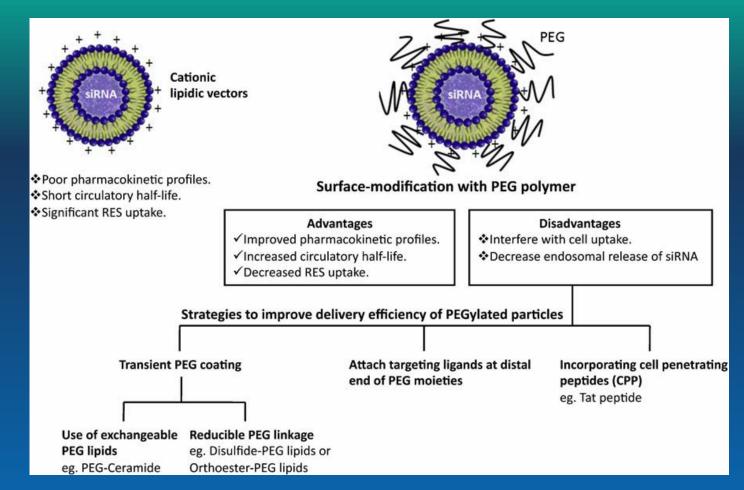
N. Dos Santos et al. / Journal of Controlled Release 105 (2005) 89-105

Lipidic Systems for In Vivo siRNA Delivery



 Formulation strategies for preparation of siRNA (small interfering) -loaded PEGylated lipid particles Sherry Y. Wu, A. J. McMillan The AAPS Journal, Vol. 11, No. 4, 2009

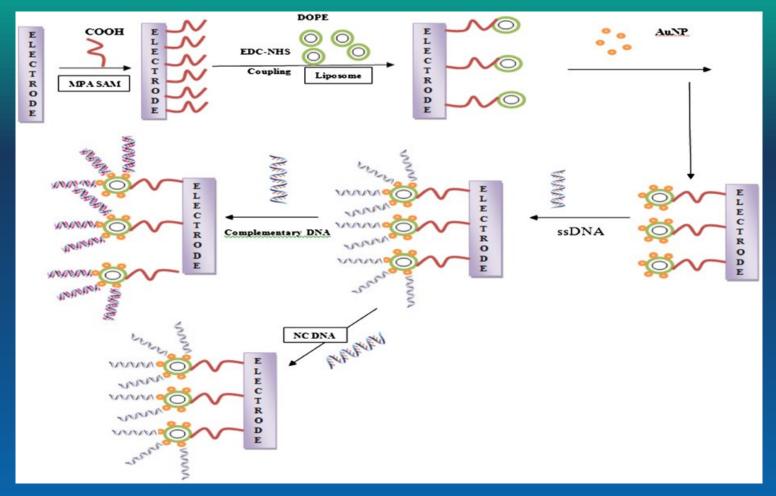
Lipidic Systems for In Vivo siRNA Delivery



• Strategies to enhance the delivery efficiency of PEGylated lipid particles

Sherry Y. Wu, A. J. McMillan The AAPS Journal, Vol. 11, No. 4, 2009

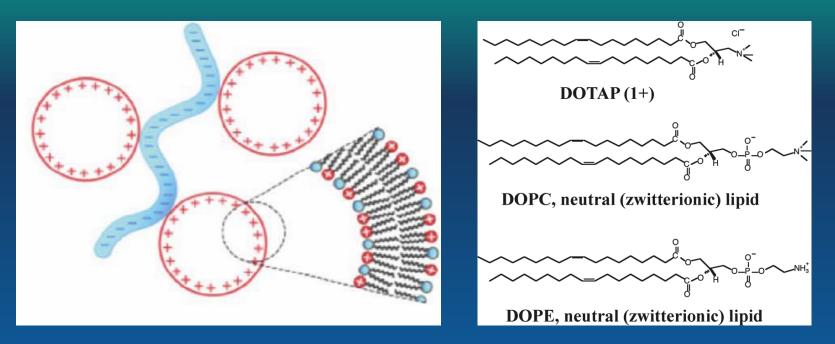
Gold surface supported spherical liposome



• Formation of spherical DOPE (1,2-dioleoyl-sn-glycero-3-phosphoethanolamine) AuNP nano-composite and DNA detection

M. Bhuvanaetal, Biosensors and Bioelectronics 41 (2013) 802

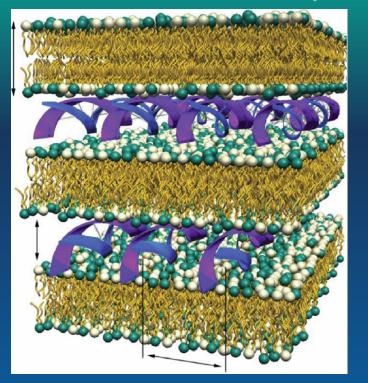
Cationic liposome-nucleic acid complexes: liquid crystal phases

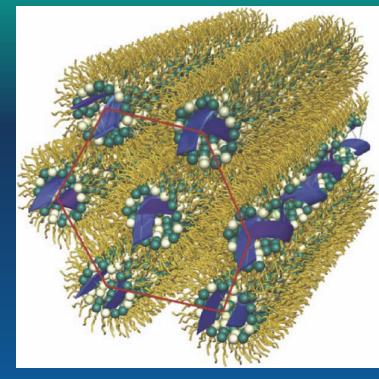


- Cationic liposome (CL) carriers of nucleic acids gene delivery
- Formation of complexes CL-nucleic acid complexes spontaneously assemble into novel liquid crystalline phases

C.R. Safinya et al., Liquid Crystals, 38:11-12, 1715 (2011)

Cationic liposome-nucleic acid complexes: liquid crystal phases

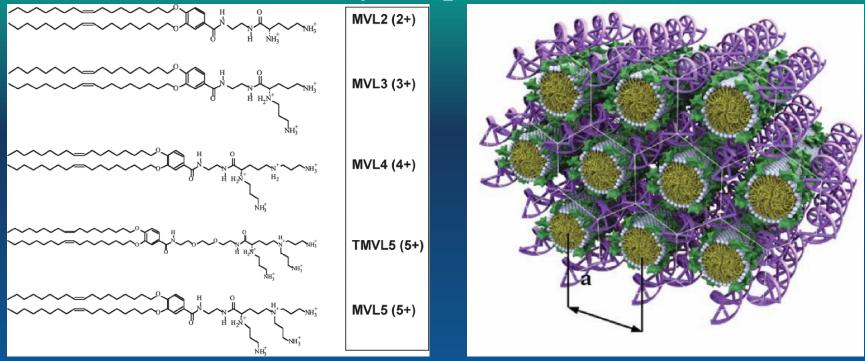




- Left: Schematic of lamellar phase with alternating lipid bilayer–DNA monolayer of cationic lipid–DNA (CL-DNA) complexes
- Right: Schematic of inverted hexagonal phase of cationic lipid–DNA (CL-DNA) complexes. In this phase, cylinders consisting of DNA coated with an inverse lipid monolayer are arranged on a hexagonal lattice.

C.R. Safinya et al., Liquid Crystals, 38:11-12, 1715 (2011)

Cationic liposome-nucleic acid complexes: liquid crystal phases



- Left: Chemical structure and maximum charge of custom-synthesised multivalent lipids (MVLs).
- Right: Schematic of CL-DNA complexes. In this phase, oppositely charged DNA chains surround cylindrical micelles arranged on a hexagonal lattice.

C.R. Safinya et al., Liquid Crystals, 38:11-12, 1715 (2011)

Conclusion

- Liposomes and their complexes can be exploited for delivery of RNA and DNA to cells.
- It can be a usefull tool for treatment of many diseases.
- There are still problems with gene therapy and many questions must be solved

Acknowledgments

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