Nanoparticles for Gene Delivery

Joshua Lim

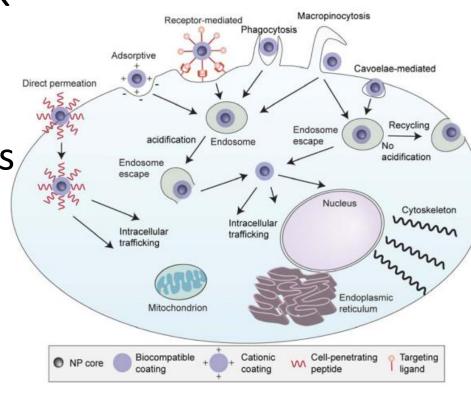


Project Overview

 Based on previous work with non-viral gene delivery method

 Iron oxide nanoparticles bring advantages

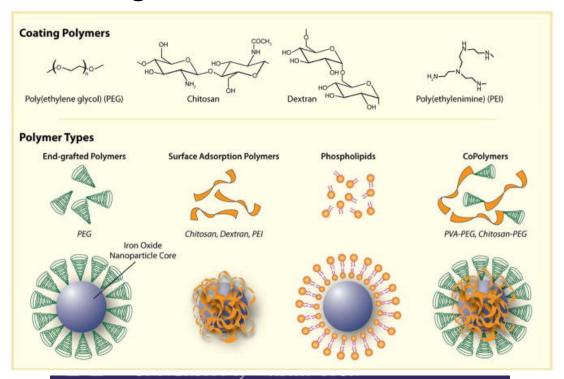
- Superparamagnetic
- Biodegradable
- Easily tunable



Iron Oxide Nanoparticles

Several components

- Iron oxide core
- Copolymer: Chitosan-PEG (polyethylene glycol)
- PEI for DNA binding-tried 1.2 kDa and 25 kDa PEI



NPBA-PEI

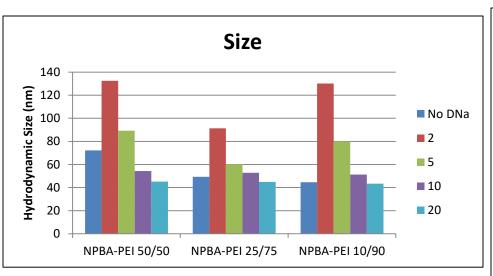
$$O_2N$$
 O_1
 O_2
 O_3
 O_4
 O_4
 O_4
 O_4

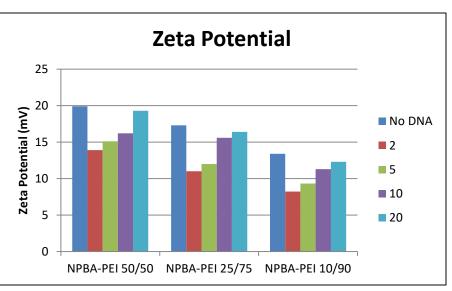
3,4-dihydroxy-6-nitrobenzaldehyde used to modify chitosan for higher affinity for iron oxide core

- Made batch of polymer, CPNBA-PEI, and incorporated into synthesis
- Tried different ratios of original polymer,
 CPBA, and new polymer, CPNBA-PEI

Characterization and DNA Binding

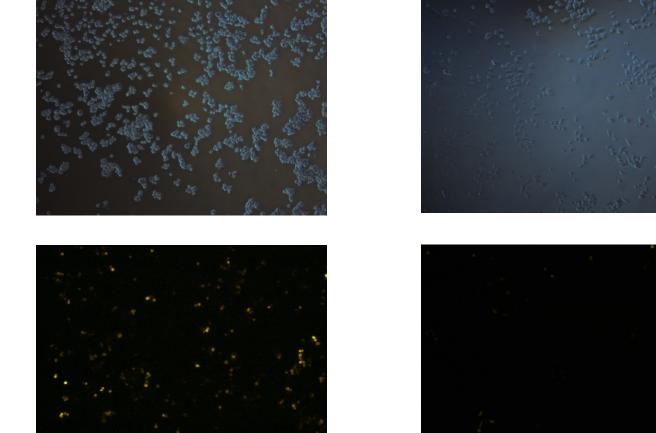
- Size and zeta potential measurements
- Bound DNA at four ratios of NP:DNA
- Red fluorescent protein DNA used





Cell Transfections

Worked best with NPBA-PEI 50/50 on SF767 cells



50/50 10:1 NP:DNA 1ug DNA

Lipofectamine control

Future Work

- Measure transfection efficiency
- Try optimizing targeting ligands for enhanced uptake in specific cells
- Try combination of 1.2 kDa and 25 kDa PEI

Acknowledgments

Neurological Surgery Summer Student Program

- Dr. Ellenbogen
- Jim Pridgeon
- Christina Buckman

Zhang Lab

- Prof. Miqin Zhang
- Prof. Forrest Kievit
- Zach Stephen, mentor