Entrapment of Hydrophobic Drugs in Nanoparticle Monolayers with Efficient Release into Cancer Cells

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Drug delivery systems (DDSs) provide an important tool for increasing the efficacy of pharmaceuticals through improved pharmacokinetics and biodistribution. A wide variety of nanoscale materials such as liposomes, polymeric micelles, and dendrimers have been employed as drug carriers. Both covalent and noncovalent approaches can be applied to the conjugation of drugs into/onto these DDSs. Noncovalent approaches have the capability of employing active drugs, whereas covalent attachment generally requires chemical modification which can cause reduced efficiency of drug release or incomplete intracellular processing of a prodrug.

Recently, gold nanoparticle (AuNP) based drug/gene delivery systems have attracted attention due to their functional versatility, biocompatibility, and low toxicity. Recent studies have demonstrated a controlled release of payload by intracellular thiols. However, controlled dissociation of drugs in active form from covalent AuNP–drug conjugates remains a challenge for clinical applications.

Noncovalent incorporation of drugs into AuNP monolayers provides an alternative delivery strategy with the potential for avoiding drug release and prodrug processing issues. The structure of commonly used water-soluble AuNPs is similar to that of unimolecular micelles such as dendrimers, featuring a hydrophobic interior and a hydrophilic exterior. The alkanethiol monolayer of the nanoparticle coupled with the radial nature of the ligands creates “hydrophobic pockets” inside the monolayer of the AuNP where organic solutes can be partitioned, as demonstrated by Lucarini and Pasquato. We report here the use of these pockets to encapsulate drugs and deliver them with high efficiency to cells.

The biocompatible AuNPs used in this study feature two functional domains: a hydrophobic alkanethiol interior and a hydrophilic shell composed of a tetra(ethylene glycol) (TEG) unit terminated with a zwitterionic headgroup. Particles with this general structure have been shown to minimize nonspecific binding with biomacromolecules.

We chose three different hydrophobic guest compounds: 4,4-difluoro-4-bora-3a,4a-diaza-s-indacene (Bodipy) as a fluorescent probe and the highly hydrophobic therapeutics tamoxifen (TAF) and β-lapachone (LAP) as drugs (Figure 1). The nanoparticle–payload conjugates (AuNPZwit-Bodipy, TAF, and LAP) were prepared by a solvent displacement method. First, AuNPZwit (Au core: 2.5 ± 0.4 nm) and guest were dissolved in acetone/water, and the solvent slowly evaporated. The bulk of the excess guest precipitated out and was removed by filtration; the particles were further purified by multiple filtrations through a molecular weight cutoff filter until no free guest was observed, followed by dialysis against buffer. The number of entrapped guest molecules per particle was determined from 1H NMR spectra and NaCN-induced decomposition experiments (see Supporting Information) and varied depending on size, hydrophobicity (log P), and molecular structure of the hydrophobic molecules (Figure 1). The particle/guest complexes are stable in buffer for >1 month and to extended dialysis, a level of kinetic entrapment greater than that observed with dendrimers.

The ability of the delivery systems to release their payload was first explored in vitro using AuNPZwit-Bodipy in a two-phase dichloromethane (DCM)–water system, where the dye is quenched by the AuNP and photoluminescence (PL) was only observed upon...
dye release. In these studies, a rapid increase in PL intensity is observed along with transfer of Bodipy into the DCM layer (Figure 1c). Significantly, since no release is observed in monophasic aqueous conditions and no particle was observed in the DCM layer, payload release occurs interfacially.

Payload delivery to cells using AuNPZwit-Bodipy was determined by confocal laser scanning microscopy (CLSM) using human breast cancer (MCF-7) cells. Efficient delivery of the dye to the cytosol is observed after 2 h of incubation with AuNPZwit-Bodipy (Figure 2a–c). Cellular uptake of nanoparticles was studied using transmission electron microscopy (TEM) and inductively coupled plasma mass spectrometry (ICP-MS), using the analogous cationic particle/dye conjugate AuNPTTMABodipy as a positive control. Little cellular uptake of AuNPZwit was observed by either TEM (Figure 2d,e) or ICP-MS for AuNPZwit-Bodipy (31 ng/well at 4 h (Figures 2), 71 ng/well at 24 h Figure S3, corresponding to uptake of 0.06% and 0.14% of available particle, respectively), whereas substantial particle uptake was observed with AuNPTTMABodipy (1750 ng/well (4 h), 2150 ng/well (24 h)). Since no free dye was observed during the 24 h incubation of AuNPZwit-Bodipy in the medium or serum solution at 37 °C (Figure 1d), Bodipy delivery presumably occurs via a monolayer–membrane transfer process, consistent with our in vitro studies.16

Demonstration of drug delivery to MCF-7 cells through presumably the same mechanism was determined through cytotoxicity studies of free and encapsulated drugs using an Alamar blue assay (Figure 3). Notably, AuNPZwit itself was nontoxic at 30 μM. In contrast, IC50 values of 4 and 4.6 μM were observed using AuNPZwit-LAP and AuNPZwit-TAF, respectively. The delivery process was quite efficient, with the per drug molecule IC50 of AuNPZwit-TAF (46 μM) only 3-fold higher than that of TAF (16 μM), and with that of AuNPZwit-LAP (6.0 μM) essentially identical to that of LAP (5.2 μM).

In conclusion, we have demonstrated that hydrophobic dyes/drugs can be stably entrapped in a hydrophobic pocket of AuNPs and released into the cell by membrane-mediated diffusion without uptake of the carrier nanoparticle. Importantly, the small size of these nanocarriers coupled with their biocompatible surface functionality should provide long circulation lifetimes and preferential accumulation in tumor tissues by the enhanced permeability and retention (EPR) effect.17 Additionally, the noninteracting nature of their monolayer should make these systems highly amenable to targeting strategies. We are currently exploring these applications as well as the role of monolayer and guest structure in the encapsulation process.

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Supporting Information Available: Experimental procedures, synthesis of gold nanoparticles and complexes, and TEM of gold nanoparticles. This information is available free of charge via the Internet at http://pubs.acs.org.

References