**In-vitro Application of Doxorubicin Loaded Magnetoplasmonic Thermosensitive Liposomes for Laser Hyperthermia and Chemotherapy of Breast Cancer**

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**Abstract**

We describe doxorubicin loaded magnetoplasmonic thermosensitive liposomes (MPTL-DOX), which are designed to combine features of magnetic drug targeting and laser hyperthermia-triggered drug release. The synthesized magnetite/gold nanoshells are stabilized using polyvinyl pyrrolidone (PVP) with mean crystallite size of 15.8 ± 3.5 nm. The liposome formulation DPPC:cholesterol:DSPE-PEG2000 at 80:20:5 molar ratio shows DOX release of less than 5% at 37°C following 24 h incubation. MPTL-DOX shows encapsulation efficiencies of about 95% and 74% for DOX and magnetoplasmonic nanoshells (MPNS), respectively. The MPTL-DOX formulation displays a desired temperature sensitivity with 65% and 100% DOX release following laser irradiation and then 24 h incubation at 37°C, respectively. The rate of DOX release from liposome using this formulation is 0.09 which was obtained by heating to 43°C, and agrees well with the first kinetic model. A temperature rise between 4-12°C was achieved for MNS using 25 μg/ml and 300 μg/ml after 400 s respectively. For cytotoxicity measurement, one untreated (control) and two treatment groups are studied. The first treatment groups are: with MPNS only, with MPTL only, and laser irradiation only. The second treatment groups are: laser hyperthermia using MPTL, MPTL with magnetic field (MF), MPTL-DOX, and MPTL-DOX with MF. MPTL-DOX is targeted to breast cancer cell lines (MCF-7 cells) under a permanent magnetic field and exhibits a substantial increase in cytotoxicity and apoptotic effects. The results suggest that externally guided drug targeting can trigger drug release using an exogenous absorber in laser hyperthermia which can be used advantageously for thermo-chemotherapy of cancers.

**Keywords:** Magnetoplasmonic nanoshells; Doxorubicin; Thermosensitive liposome; Laser hyperthermia; MCF-7 breast cancer cells; Cytotoxicity

**Introduction**

Engineered nanomaterials hold great promise in drug delivery systems [1-3]. Novel magnetic nano-formulations such as liposomes, metallic/nonmetallic, and polymeric nanoparticles have increased the ability to deliver drugs for which conventional therapies have shown limited efficacy. When magnetoliposomes containing iron oxide nanoparticles are exposed to a magnetic field the liposomal temperature exceeds transition temperature and the contents are released [4]. Heating techniques may be improved by using a magnetic field localized to therapeutic site, for instance a tumor [5-7]. Hassannejad et al. have recently demonstrated that gold-coated superparamagnetic iron oxide nanoparticles (SPIONs) encapsulated by phospholipid liposomes can be used as exogenous absorber in laser thermal therapy [8]. Moreover, gold nanoparticles were conjugated with the anticancer drug Doxorubicin, for drug delivery to liver cancer recently [9]. When gold nanostructures are irradiated by laser, the absorbed optical energy raises their temperature, hence acting as a localized heat source that is quickly equalized within the nanoparticles. The heat is then transferred to the surrounding environment determined by their thermal relaxation time and laser pulse duration. Therefore, light-triggered thermal properties of liposomal Au nanostructures are different from the classical thermosensitive liposomes [10]. Au nanostructures absorb energy at a characteristic wavelength and exhibit surface plasmon resonance (SPR) [11,12]. Only part of the absorbed energy is emitted as photoluminescence while most of it is converted to heat [13]. Therefore, light-induced heating of liposomal Au nanostructures by UV, visible and near-infrared to release liposome-entrapped drugs has developed [14].

In addition to the required elements for triggered drug release, an optimized delivery system for cytotoxic drugs should have biophysical properties (e.g. stealth properties) to favour passive accumulation in tumors upon intravenous administration. This study is a novel approach towards implementing a PEG-shielded thermosensitive liposome encapsulating the DOX as a drug together with PVP-stabilized magneto-plasmonic nanoshells (MPNS), enabling magnetic drug targeting by static gradient magnetic fields and laser hyperthermia as a trigger for drug release, producing a synergistic cytotoxic effect. To the best of our knowledge, this combination of magnetic targeting/laser hyperthermia/drug release is novel and has not been addressed previously.

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Materials and Methods

Chemicals and reagents

All analytical reagents are used without further purification. Ferric chloride hexahydrate (FeCl₃·6H₂O, 99%), ferrous chloride tetrahydrate (FeCl₂·4H₂O, 99%), hydrochloric acid (HCl, 37%), sodium hydroxide, chloroform, formaldehyde solution (H₂CO, 37%), absolute ethanol, Triton X-100, and polyvidone25 are purchased from Merck (USA). Tetrakis (hydroxymethyl) phosphonium chloride (THPC) aqueous solution, Gold (III) chloride trihydrate (HAuCl₄·3H₂O, ≥94% Au basis), 3-amino propyltriethoxysilane (APTES), and Doxorubicin hydrochloride (DOX) are purchased from Sigma-Aldrich (USA). Dipalmitoylphosphatidylcholine (DPPC), cholesterol (Chol), and 1,2-distearyloxy-sn-glycero-3-phosphoethanolamine-N-[amino(polyethylene glycol)-2000] (ammonium salt) (DSPE-PEG2000) are purchased from Avanti Polar Lipids (USA). MCF-7 cell line is purchased from the Pasteur Institute (Iran). RPMI 1640 medium and fetal bovine serum (FBS) are purchased from GIBCO (USA). Annexin-V-FLUOS staining kit is purchased from Roche (Switzerland). Deionized water (18 MΩ·cm) is provided by a Milli-Q system and deoxygenated by vacuum for 1 hour prior to the use.

Synthesis of PVP- stabilized gold nanoshells

Gold coated superparamagnetic iron oxide nanoparticles (SPIONs) are fabricated by a multistep procedure as described earlier [15,16]. Briefly, SPIONs are synthesized by the coprecipitation method [17-19]. Then the surface of SPIONs is functionalized with APTES to generate an amine terminated surface [20]. A colloidal gold solution containing ~ 2 nm Au particles is prepared according to the Duff and Baker [21] method. After that, a continuous gold shell is grown around the SPIONs. In order to prepare PVP-coated gold nanoshells, in a typical procedure [8], the obtained nanoshell dispersion in water (2 ml) is centrifuged at 4000 g for 25 min. Then 1 ml supernatant is replaced by 1 ml PVP solution and the obtained mixture stirred for 24 hours at room temperature. After that, it is centrifuged twice under 4000 g for 25 min and washed with water in order to remove free PVP. As the hydrophilic head group (N-C=O) of PVP is attached to the surface of the particles, particle repulsion will occur due to steric interactions between polymers adsorbed on the particle’s surface.

Preparation of magnetoplasmonic liposomes

Liposomes are composed of DPPC:Chol:DSPE-PEG2000 at a molar ratio of 80:20:5, as suggested by Pradhan et al. [22]. A lipid mixture of chloroform stocks is prepared and dried at 42°C under nitrogen stream and further placed in a vacuum overnight. The lipid film is hydrated with a buffer consisting of PVP-coated gold nanoshell dispersed in 300 mM Citrate (pH 4.0) at 60°C. After hydration, 15 min of sonication (Tecnar 20, 190 W) is applied to break down any larger vesicles. Subsequently, the samples undergo 5 cycles of freeze-thaw including 10 min at -196°C, 10 min at 65°C and 30 s vortexing between cycles. Unencapsulated nanoparticles are removed by centrifuging at 4000 g for 5 min, after which the supernatant liposomal dispersion is centrifuged at 20000 g for 30 min to precipitate the liposomes encapsulating gold nanoshells. Liposomes are obtained by extruding the mixture 15 times with an extruder (Avanti Polar Lipids) at 55°C through polycarbonate membrane filters (Whatman PLC, Maidstone, Kent) with a pore size of 100 nm. Encapsulation efficiency of MPTL is calculated using ICP-MS results through the formula (Wₐ/Wᵢ) x 100%, where W_i is initial concentration of Fe (mg/ml) in hydration solution and W_a is encapsulated concentration of Fe (mg/ml) in the liposomes.

Doxorubicin loading into the magnetoplasmonic liposomes (MPTL-DOX)

Doxorubicin, a weak base cation with a pKa of 8.3, is encapsulated into the extruded liposomes using a pH-gradient loading protocol as described by Mayer et al., [23] with a slight modification: the exterior pH of the extruded liposomes is adjusted to 8 with sodium carbonate solution (500 mM) creating a pH gradient. The preheated liposomes are incubated with doxorubicin hydrochloride (DOX:lipid weight ratio of 1:10); with respect to the amount of original total lipid used for the liposome preparation at 60°C for 1 h. Unencapsulated DOX is removed by passing the liposome through a Sephadex-G50 (fine) column at a flow rate of 0.5 ml/min with HEPES buffered saline (10 mM HEPES pH 7.4 and 140 mM NaCl) as the eluent. The resulting liposomes (MPTL-DOX) are stored at 4°C until further use.

The DOX concentration of the liposomes is determined by a Cary Eclipse spectro-fluorometer equipped with ThermoScan Software (Varian, Palo Alto, CA) at 485 nm excitation and 590 nm emission wavelength. 20 μl DOX loaded liposomes are incubated in 1 ml HEPES buffered saline for up to 60 min and the fluorescence intensity is measured after various incubation times. Before measurement, the samples are allowed to cool down to room temperature. For quantification, a calibration curve is obtained with a dilution series of free doxorubicin in 1% Triton X-100 in HEPES buffered saline is used. Using the following formula, entrapment efficiency (‰) = (encapsulated drug in liposomes / amount of total drug) x 100% [24].

Cells and cell culture

The MCF-7 breast cancer cells are cultivated in RPMI 1640 medium supplemented with 10% FBS and maintained at 37°C in a 5% CO₂ incubator.

Cytotoxicity study

Cytotoxicity studies are done using MTT (3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide) tetrazolium reduction assay. As suggested by Kulshrestha et al. [25] 1×10⁴ cells per well are cultivated in 96-well plates overnight. The control group is untreated cells and the first MCF-7 treatment groups are: with magnetoplasmonic nanoshells only, with MPTL (at a lipid concentration of 10 mM) only, with laser irradiation only. For the control (untreated) group, the cells are washed with PBS, after that the fresh cell culture media is placed in an incubator for the next 24 h. For the MPNS and MPTL treatments, the cells are cultured in 200 μl medium containing nanoshells and MPTL (at lipid concentration of 10 mM), respectively. In order to study the effect of magnetic fields on nanoliposomes, during the first hour of incubation, the culture plates are positioned on the 96-well format magnetic plate (Chemicell-Germany) with a field gradient of 50-130 T/m. After 75 min, the cells are washed with PBS and incubated for another 24 h with fresh medium.

The second treatment groups are: laser hyperthermia mediated by MPTL only, MPTL with magnetic field (MF), MPTL-DOX only and MPTL-DOX with MF all at lipid concentration of 10 mM. In vitro hyperthermia of MCF-7 cells consisting of liposomes containing magnetoplasmonic nanoshells is done using Argon laser (Melles Griot /43 Series Ion Laser-USA) at 350 mW for 5 minutes with or without DOX. The thermometer is connected at the other end to a Windows-based laptop where the results are displayed on the screen. During the experiment the temperature is maintained at 37 ± 0.4°C recorded by a digital thermometer (Model No. HP-34420A).
In vitro evaluation of thermo-chemotherapy

In vitro evaluation of cellular uptake is based on 15 min incubation of cells in a solution containing Ca²⁺ ions and annexin V–FITC (at a final concentration of 1 µg/ml). The annexin V-binding buffer consists of 10 mM HEPES–NaOH, pH 7.4, 150 mM NaCl, 5 mM KCl, 1 mM MgCl₂, and 1.8 mM CaCl₂. The stained cells are mounted on slides and visualized under a fluorescence microscope. A two steps procedure of preparation and application of the multihybrid nanoliposomes is illustrated in Figure 1.

Characterization

Transmission electron microscopy (TEM) as performed using a CM 200 FEG STEM Philips-M.E.R.C operating at the voltage of 200 kV. Liposomes are observed by TEM following negative staining with uranyl acetate. Liposomes are diluted with distilled water and dropped on a PDL-coated copper grid. The excessive sample is removed with filter paper and air-dried for 1 min at room temperature. Subsequently, uranyl acetate solution (10 µl of 1%) is dropped onto the grid. After 1 min the excess staining solution is removed with filter paper and is allowed to dry in the air before introduction into the microscope. Fourier Transform Infrared Spectroscopy (FT-IR) spectra are recorded by TENSOR27 FT-IR spectrometer. The obtained nanoparticles are dried, mixed with KBr and compressed into a pellet. UV-VIS spectroscopy of nanoparticle suspensions is taken on a CARY100 UV-VIS spectrophotometer with a 10 mm optical path length quartz cuvette. The mean hydrodynamic radius and polydispersity of the liposomal vesicles are determined using Dynamic Laser Light Scattering (DLS). The scattered laser light intensity at 90° is measured by light scattering photometer using a wavelength of 488 nm (Brookhaven instrument Corporation, USA). Stability of magnetoliposomes formulation is evaluated by measuring the zeta potential. The zeta potential is measured as the particle electrophoretic mobility of charged, colloidal suspensions by means of laser microelectrophoresis in a thermostatted cell at room temperature (Brookhaven Instruments Corporation, USA). Every sample measurement is repeated 5 times. Encapsulation efficiency of gold nanoshells within the liposomes is determined by inductively coupled plasma mass spectrometry (ICP-MS). Samples for ICP-MS (VARIAN 735-ES) analysis are frozen, lyophilized, and dissolved in nitric acid hydrochloride, prepared by adding nitric acid (100 µL) and hydrochloric acid (300 µL of 37%) for 72 h to dissolve particles. Then, samples are diluted to 2 mL with HNO₃ (1.6 mL of 2%) and analyzed via ICP-MS against standards. Magnetization measurements are carried at 300 K in a magnetic field (H) of up to 20 kOe with a vibrating sample magnetometer (Meghnatis Daghigh Kavir Co. VSM/AGFM) that can measure magnetic moments as low as 10⁻³ emu. For the magnetization measurements, NPs are in dry powder form obtained by evaporating the water from the solution. The samples are dried from freeze dryer (Pishtaz Engineering Co. Model: FD-4).

Statistical analysis

All experiments are performed at least in triplicate. The value of particle size determined by TEM micrographs is the average of at least 50 measurements and reported as mean ± standard deviation. The SPSS 15.0 as used to perform the calculations. Paired-sample t-test is used to compare the difference between the control and treated groups. P values of <0.05 are considered statistically significant.

Results and Discussion

FTIR

Figure 2 shows the FT-IR spectra of bare SPION (a), amine-functionalized SPION (b), PVP (c) and PVP-covered gold nanoshell (d). The broad band at region of 3550-3200 cm⁻¹ can be attributed to the N-H stretching vibration and NH₂ bending mode of free NH₂ groups, respectively. Also, it can be seen that the characteristic bands of the Fe-O bond of amine-functionalized SPIONs shift to higher frequencies of 620 and 477 cm⁻¹ compared with that of bare SPIONs (at 579 and 444 cm⁻¹). These shifts have been attributed to the replacement of –H at the Fe-O-H groups on the surface of SPION by the more electronegative group of –Si(O-)₂⁻ which results in the enhancement of bond force constant for Fe-O bonds. The FT-IR spectrum of PVP (Figure 2c) mainly consists of two bands at 1282 and 1475 cm⁻¹ corresponding to the vibration of C-N and pyrrolidone ring. Similar bands are also seen in the FT-IR spectrum of PVP-coated gold nanoshells. However, comparing with the spectrum of PVP, the resonance peak of C-N, at 1282 cm⁻¹, was shifted to 1475 cm⁻¹ and the band of C=O, at 1664 cm⁻¹ shifted to 1647 cm⁻¹. The change of the spectrum indicates that the absorption of PVP on the gold shell is not on the basis of electrostatic attraction but gold atoms on the surface of nanoparticles would coordinate with N and O atoms.
have magnetization saturation (Ms) of about 46.94 emu/g. Absence of hysteresis loss with zero coercivity in the magnetization curve suggests the superparamagnetic nature of the particles. The Ms value of the resultant gold nanoshells decreases to 11.98 emu/g, when the gold is added. With the same magnetite core size and gold shell thickness of 3.5 nm, the Ms value for the synthesized nanoshells in this paper is larger than that reported by Xu et al. [28].

**TEM**

The physical size of particles at different steps of nanoshell fabrication is determined by bright field TEM. Based on TEM micrographs (Figure 4), the average diameter of SPIONs and SPION/gold nanoshells are 9.5 ± 1.4 and 15.8 ± 3.5 nm, respectively. TEM image confirms the formation of magnetoplasmonic liposome (Figure 5). The clear margin of each encapsulated nanoparticle indicates the steric hindrance of PVP-covered nanoshells. Based on TEM micrographs, the average diameter of SPION/gold nanoshells are 15.8 ± 3.5 nm. It is observed that MPNS are packed within the liposomes. The thickness of the lipid bilayer membrane surrounding the collection of nanoshells is 6.5 ± 1.6 nm measured by NIH Image J software (http://rsb.info.nih.gov/ij/), suggesting the formation of unilamellar liposomal structure. A value of 7-8 nm is also reported for lipid bilayer membrane [29].

**Encapsulation efficiency of nanoshells**

A magnetoplasmonic nanoshells (MPNS) encapsulation efficiency of 54% is obtained. Higher encapsulation of MPNS results in higher heating ability, therefore it increases the toxicity of the liposomes. In this aspect, this magnetoplasmonic liposomes is superior to the others reported in literature [22]. The concentration of nanoshells has

of PVP which is consistent with previously reported results [26]. FTIR confirms the presence of a PVP coating on the nanoshells.

**DLS**

The effect of surface coating on the stability of SPIONs is observable from DLS data. The volume-weighted hydrodynamic size measured by DLS for nanostructures are summarized in Table 1. The reduced hydrodynamic size of amine-functionalized SPIONs shows more stability of nanoparticle suspension after silanization reaction. This effect can be also seen after coating of the amine-functionalized SPIONs with a continuous thin layer of gold, which reduced the hydrodynamic size of gold nanoshells to 57 nm. After PVP coating of gold nanoshells the hydrodynamic size increased to 102 nm which is because the nanoshell’s surface functionalization with long chain PVP molecules. This result agrees well with the previously reported thickness in the case of PVP adsorbed on palladium nanoparticles [27]. The hydrodynamic diameter of the MPTL-DOX is 240 nm. The size distribution of this formulation is quite homogenous as polydispersity index is around 0.19.

**VSM**

Figure 3 shows the magnetization curve for magnetite core and gold nanoshells at ambient temperature. It indicates the superparamagnetic behaviour of nanostructures. It can be deduced that the magnetite NPs

![Figure 2: FT-IR spectra of bare SPION (a), amine-functionalized SPION (b), PVP (c) and PVP-covered SPION/gold nanoshells (d).](image-url)

![Figure 3: Variation of magnetization with applied magnetic field for bare magnetite core and magnetoplasmonic nanoshells.](image-url)
a significant effect on the amount of heat generated at a given power density. This effectively indicates that not only do the MPNSs play an important role in generating thermal effect, but also an enhanced absorption cross section for spherical metal nanoshells can be achieved at optimized value of concentration for a better laser-hyperthermia [16]. The absorption efficiency of our synthesized MPNS corresponds to 0.656 at SPR of 531 nm. Maximum value of the parameter DT/I shows the efficiency of transformation of absorbed optical energy by MPNS into the thermal energy, where T is the temperature and I is constant intensity of laser radiation during pulse duration. In our case, the nanoshell efficiency parameter equals 2×10^{-5} °C cm^2/W.

Furthermore, an experiment was set up to measure the temperature of the aqueous medium containing dispersed gold nanoshells and magnetoplasmionic liposomes where the the cuvette containing the sample was irradiated by laser at 17W/cm^2 for 400 s. A thermostt probe (Redfish Sensors Inc., Model QTGB-14 D3) with an accuracy of 0.15% was placed in the solution to monitor the temperature change where it was recorded by a thermometer (Model No. HP-34420A. The output of the thermometer was connected to a Windows-based laptop where the results were displayed. As it is shown in Figure 6, at low concentrations, the temperature of MPNS increases steadily and linearly up to 4°C in about 300 s where it reaches the turning point and saturates after 400 s. At 100 μg/ml the temperature reaches its maximum of 6.5°C but when the concentration was increased to 300 μg/ml, the temperatures increases linearly up to 8°C in about 120 s where it shows a turning point and at this stage the temperature rise gradually until it is maximized at 12°C after 400 s. Such a phenomenon can be due to the arrangement, density, and potential for aggregation of Au nanoshells. Therefore, it results in a stronger surface plasmon absorption of light and more efficient photothermal effects. All the experiments demonstrate that the temperature increases with increasing MPNS concentration and the exposure time. However, in the case of MPNS encapsulated liposomes, Figure 7, the maximum temperature obtained was about 6°C after 400 s which as expected is less than the case of without liposome. These results clearly confirm that the fabricated MNSs not only are suitable candidates for hyperthermia tumor therapy but even with liposome carriers the temperature increase is sufficient to cause the hyperthermic effect about 42-45°C. The question that how fast and what temperature is generally achievable, entirely depends on the MPNS concentration, the laser power and the irradiation time.

**Laser induced fluorescence spectroscopy (LIF)**

The fluorescence response of Dox was studied at room temperature using 488 nm wavelength of argon laser which is close to its excitation wavelength. The emission spectrum was recorded and processed by Spectrasuit software. The results of LIF experiment for Dox, TL, TL-Dox, and MPTL-Dox are shown in Figure 8 where a peak at 590 nm corresponding to Dox emission is observed. A small bump at 550 nm is thought to be due to Au nanoshell response.

Recent studies have indicated that AuNPs act as good quenchers...
of many fluorescence donors, due to the nanosurface energy transfer (NSET) effect [30]. Addition of gold nanoshells to TL-Dox result in a significant quenching of doxorubicin fluorescence (Figure 8d), demonstrating the presence of NSET between the doxorubicinyl groups and Au nanoshells.

**Encapsulation efficiency of DOX and drug release study under laser irradiation**

The drug encapsulation efficiency of 95% for DOX-loaded magnetoplasmonic liposome is calculated. The liposome formulation DPPC:cholesterol:DSPE-PEG2000 shows DOX release of less than 5% at 37°C following 24 h incubation. This particular lipid composition has Tm at around 41.3°C [22], as a thermosensitive formulation, so minimal leakage at 37°C and triggered DOX release at 43°C is expected. DOX release from magnetoplasmonic liposomes are 65% and 100% in 50% FBS following 5 min Argon ion laser (350 mW, l=514 nm) irradiation and 24 h incubation at 37°C, respectively. Laser induced hyperthermia raises the temperature from 37°C to 43°C which leads to the phase transition of the lipid combination causing massive release of the encapsulated drug. Most of the magnetic liposomes reported in earlier studies [4,31] have been composed of DPPC alone, which shows a high burst of drug release even at 37°C. By adding saturated lipids, the compressibility moduli of the liposome could be increased.

Furthermore, it has been established that drug release from low temperature sensitive liposomes occurs via grain boundary permeabilization when it is heated into the region of its phase transition temperature [5]. However, the grain boundary structures of the bilayer could bind proteins that are responsible for opsonization. Therefore, in order to reduce the toxicity (by avoiding the opsonization) a few mol% of DSPE-PEG2000 is added to cover the bilayer [5].

**Drug release from thermosensitive liposomes**

As reported by Pradhan et al., [22], phase transition temperature of the liposome prepared with formulation of DPPC:CHOL:DSPE-PEG2000 at the molar ratio of 80:20:5 is ~42°C. In this report, the drug release rate from similar lipid composition is measured as a function of time at 43°C. The release rate of thermosensitive liposome varies according to the composition of liposome, its preparation procedure, and heating temperature [32]. The relation between percentage release and exposure time is found to follow the first-order kinetics expressed as [33]

$$\% R(t) = R_c \left(1 - e^{-k_{rel}t}\right)$$  \hspace{1cm} (1)

where % R(t) is the percentage of drug released at exposure time t, k_{rel} is liposome release rate and R_c is the total percentage of drug released at a given heating temperature. It is assumed that at t=0 the temperature is 37°C. This equation is used to fit the experimental data obtained at 43°C [32]. From the best fitting curve (shown in Figure 9) obtained by using nonlinear least squares method, the release rate is found to be 0.07 for TL-Dox and 0.09 for MPTL-Dox. In addition, doxorubicin release from both of the formulations are assessed (Table 2).

**Cytotoxicity study**

Figure 10 shows the cytotoxic effect of laser, MPNS, and MPTL on MCF-7 cells. The untreated (control) and first treated (i.e. exposure of untreated cells to laser, MPNS, or and MPTL) groups are non-toxic. After being heated, the cells develop resistance to heat, which reduces the likelihood of being destroyed by direct thermal cytotoxic effects. Because Hyperthermia alters the cell walls by means of so-called heat shock proteins, cancer cells then react much more effectively to the radiation [34]. So before the laser hyperthermia experiment, the cells are normally preheated which in our case it was done by using a

Figure 8: LIF spectra (a) free Dox, (b) liposomes, (c) Dox loaded liposomes, (c) Dox and gold nanoshells co-loaded liposomes.
water bath (Lab Companion, USA). The heating effect of surrounding liquid medium on viability of MCF-7 and L-929 cells was assessed after being heated for 5 min at the temperature of 43°C. The cell viability is significantly (~%50) decreasing in L-929 cells, but as for MCF-7 cells, there is no significant difference between control and treated cells. The cytotoxic effects of laser hyperthermia and chemotherapy using the prepared formulation as single and combined modalities are illustrated in Figure 11. Data are presented as a percentage of the cell viability where viability of control group is taken as 100%.

Magnetic guidance is one of the strategies for the accumulation of nanoparticles. Therefore, when the magnetic field is applied, most of the MNS are accumulated at the bottom of the wells, where the cells are present. After that, based on the SPR absorption in gold nanoshells, an energy relaxation is followed through a non-radiative decay channels which results in an increase in kinetic energy, leading to overheating of the local environment around the light-absorbing species. Clearly, the hyperthermia effects are expected to be higher, where the applied magnetic field has accumulated of most of the nanoparticles around the cells. Because it increases the possibility of Au NPs aggregation, hence resulting in a stronger localized SPR and thus more efficient photothermal effects. Moreover, the combined laser hyperthermia treatment and magnetically targeting MPTL-DOX, at doxorubicin concentration of 0.5 µg/ml (IC-50 of MCF-7 cells) [35] and 0.1 µg/ml result in less than 1% and less than 15% cell viability (Figure 11), respectively. An analysis of these combined treatments using Valeriote's formula [36] shows that the magnetically targeted MPTL-DOX and laser hyperthermia treatments are synergistic in nature.

**Fluorescence microscopy**

Figure 12 shows that the internalized nanoliposomes release doxorubicin in the acidic organelles of MCF-7 cells, which would activate the fluorescence of doxorubicin quenched by Au nanoshells due to NSET. The fluorescence microscopy images of MPTL-DOX in MCF-7 cells obtained by a fluorescence microscope Eclipse 80i (Nikon, Japan). DOX-MPNS loaded liposomes are incubated with MCF-7 cells for about 75 min (with and w/o MF) and irradiated with Ar laser at 37°C. The image (a) shows the red fluorescence in the area or the nucleus of the MF applied to MPTL-DOX in MCF-7 cells. The nucleus of the cells in MPTL-DOX-w/o MF (d) does not show this fluorescence. This confirms that under magnetic field, the drug loaded NPs are internalized by the cells. It has been reported that chemosensitivity to DOX in MCF7 is due to the decreased level of DNA double-strand break repair proteins, Rad51 [37]. Aroui et al., have shown that the major mechanism of DOX activity is the inhibition of topoisomerase II and stabilization of a ternary drug-topoisomerase II (TOPO II)-DNA complex, causing DNA damage and induction of apoptosis [38]. Furthermore, Prasad et al., [39] have shown that hyperthermia causes apoptosis by irreversibly damaging the actin and tubulin structures of cells. Hyperthermia damages different cellular structures such as enzyme complexes required for DNA synthesis and
repair [40]. We believe that the presence of gold nanoshells enhances the laser hyperthermia effect in a controlled manner.

In addition, apoptosis is a mode of cell death that is accompanied by specific alterations to the plasma membrane. In Figure 12, membrane blebbing can be identified which is characteristic of apoptotic cells [41]. Redistribution of phosphatidylserine from the inner to the outer plasma membrane leaflet has become one of the most widely used markers for apoptotic cells in mammals. This is largely due to the availability of annexin V probe, as a sensitive phosphatidylserine-binding protein. The FITC-conjugated annexin V-binding assay provides a very specific, rapid and reliable technique to detect apoptosis. Analysis of fluorescence microscopy images is conducted to reveal the enhanced cell apoptosis caused by the magnetoplasmonic nanoshells and doxorubicin under combined chemotherapeutic and hyperthermia therapies. Therefore, this experiment indicates the doxorubicin loading into the liposomes in addition to indicating the potential application of this hybrid nanosystem in fluorescence imaging.

Conclusion

In the present study, fabrication of a new class of doxorubicin-loaded MPNS encapsulated thermosensitive liposomes is presented. The drug release behavior of the formulation after Ar laser irradiation and 24 h incubation at 37°C is studied. Moreover, the cytotoxic and apoptotic effects of combined laser hyperthermia and chemotherapy for MPTL-DOX are evaluated for MCF-7 cells. In our case, cytotoxicity and uptake of DOX are enhanced under magnetic field. We believe that preliminary studies suggest the potential application of MPTL-DOX as biomarker and for combined chemotherapy and laser hyperthermia. Further research is required for optimization of the effect of MPNSs size and laser wavelength for *in-vivo* higher penetration and thermal induction.

References


