

Barium titanate core – gold shell nanoparticles for hyperthermia treatments

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Abstract: The development of new tools and devices to aid in treating cancer is a hot topic in biomedical research. The practice of using heat (hyperthermia) to treat cancerous lesions has a long history dating back to ancient Greece. With deeper knowledge of the factors that cause cancer and the transmissive window of cells and tissues in the near-infrared region of the electromagnetic spectrum, hyperthermia applications have been able to incorporate the use of lasers. Photothermal therapy has been introduced as a selective and noninvasive treatment for cancer, in which exogenous photothermal agents are exploited to achieve the selective destruction of cancer cells. In this manuscript, we propose applications of barium titanate core–gold shell nanoparticles for hyperthermia treatment against cancer cells. We explored the effect of increasing concentrations of these nanoshells (0–100 µg/mL) on human neuroblastoma SH-SY5Y cells, testing the internalization and intrinsic toxicity and validating the hyperthermic functionality of the particles through near infrared (NIR) laser-induced thermoablation experiments. No significant changes were observed in cell viability up to nanoparticle concentrations of 50 µg/mL. Experiments upon stimulation with an NIR laser revealed the ability of the nanoshells to destroy human neuroblastoma cells. On the basis of these findings, barium titanate core–gold shell nanoparticles resulted in being suitable for hyperthermia treatment, and our results represent a promising first step for subsequent investigations on their applicability in clinical practice.

Keywords: barium titanate nanoparticles, gold nanoshells, NIR stimulation, human neuroblastoma

Introduction

Hyperthermia therapy, which entails the application of heat to treat disease conditions, has a long history dating back to ancient times. Hyperthermia is more precisely described as heating tissue to a temperature of 42°C–47°C for some tens of minutes,¹ leading to irreversible damage in the case of cancerous cells. At this temperature range, because of their physiological characteristics, tumor cells may undergo either necrotic or apoptotic death.^{2–6} Yet, hyperthermia has been synergistic with conventional cancer therapy methods to attain more effective results.^{3,7}

The main technological challenge of nanoscience and nanotechnology in recent years has concerned adding selectivity to the therapy, which has led to the idea of using exogenous heating agents. Since initial introduction of laser to hyperthermia, recent investigations in nanotechnology have generally focused on developing nanostructures with special optical properties that are appropriate for laser-induced hyperthermia, specifically metal nanostructures that demonstrate strong optical coupling of their plasmons with the laser light.^{8–13} The wavelength at which the surface plasmons resonate

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with the incident light (surface plasmon resonance), as well as the relative contribution of absorption or scattering to the total extinction spectra of the nanoparticles, strongly depends on the shape, size, type of metal, and surrounding dielectric environment,^{13–19} as well as on the particle orientation and interparticle interactions.²⁰ An increase in the size of the nanoparticles results in an increase of the extinction cross-section and the relative contribution of scattering to the total optical extinction.¹⁸ Plasmonic nanoparticles of a size much smaller than the wavelength of the light are better absorbers, and therefore prior candidates for localized photothermal heating, which is critical to invasive cancer treatment and tumor removal.^{21–23} These particles own a far-field extinction spectrum dominated by the dipole term from the series expansion of harmonic function of Mie theory.²⁴ Larger nanoparticles instead demonstrate larger scattering cross-sections²⁵ that can be exploited for contrast enhancement in bioimaging.^{26–29} The resonant excitation of larger particles though, when of 100 nm or over in size, shows significant contributions from higher order multipole modes that create a broader far-field extinction spectra. The combination of the absorption and scattering properties, which depends on the size of the particle, allows nanoparticle-based integrated diagnostic imaging and therapy.^{30–32}

During recent years, various nanostructures different in morphology, surface chemistry characteristics, chemical and physical properties, and therefore different optical features are intensively being investigated, each of which demonstrate their own appealing contribution to near-infrared (NIR) photo-assisted hyperthermia therapy and diagnosis of cancer, as well as to other scientific fields. Where diagnosis is concerned, with the intention to reach the molecular level, a great number of recent studies has been concentrated on surface-enhanced spectroscopy methods, such as surface-enhanced Raman spectroscopy (SERS). This requires particles or structures that possess the so called hot spots to enhance the near electrical field and to respectively increase the output signal of the targeted species. Various nanoparticles and structures are being introduced and studied for this aim.^{33,34} Other studies, on the other hand, show results of second harmonic generation (SHG) response from plasmonic nanoshell structures³⁵ and its dependence on the core dielectric constant as well as on the refractive index of the surrounding medium. Despite the fact that the main interest with plasmonic nanostructures is more concentrated on cancer therapy and diagnosis, these particles have had a significant influence on other fields of science as well.³⁶ Numerous nanostructures different in

shape, material, and structure, and therefore in chemical, physical, and optical properties have been and are being introduced and exploited for various applications. Among the variety of nanoparticles developed so far, nanoshells, introduced for the first time by Halas et al, present a new family of materials with tunable plasmon band, throughout the visible and into the infrared region of the electromagnetic spectrum.^{37–40} Being a hybrid structure, a variety of physical, chemical, and optical properties can be reached through a combination of different materials. These novel features can include as an instance the improved photocatalytic property of ZnO nanospheres upon being composed with a noble metal into nanoshell structure⁴¹ and many other examples in this regard. Therefore, nanoshells are still a subject of intense study either from a physical or chemical point of view.⁴² In the case of nanoshells, it is possible to modulate the relative contribution of the absorption or scattering of the particle by manipulating only the core/shell ratio and keeping the particle size unchanged.^{43,44} Nanoshells, as nanoscale heat sources, are currently being exploited in photothermal cell death and tumor destruction *in vitro*,^{45–47} with some other minor potential applications such as exogenous absorbers for tissue welding.^{48,49} In addition to being used as heating sources, these particles are either exploited as contrast agents, optically adjusted across a broad range of wavelengths in comparison with conventional molecular contrast agents.⁵⁰ Moreover, these particles are biocompatible and non-photobleaching,⁵¹ and have been employed in various imaging modalities.^{51,52} To date, the nanoshell-assisted photothermal approach has been demonstrated to be effective in *in-vitro* and some *in-vivo* experiments on breast, brain,^{9,53} and other tumor cell lines.

Following this line of research and our preliminary results on the synthesis of barium titanate core–gold shell ($\text{BaTiO}_3@$ goldshell) nanoparticles,⁵⁴ in this manuscript we present an optimized synthesis of $\text{BaTiO}_3@$ goldshell nanoparticles and demonstrate their potential ability for application in photothermal therapy. Recent results on the application of piezoelectric materials in biology, in particular barium titanate,⁵⁵ motivated us to select this material as the core of our nanoshells to further study the potential characteristics resulting from merging piezoelectric and plasmonic properties. Furthermore, we intend to study more the SHG properties of barium titanate nanoparticles and the transformation of their second harmonic nonlinear response upon accommodating in a nanoshell structure. As described in the results and discussion sections, experiments conducted on neuroblastoma cell lines demonstrate that $\text{BaTiO}_3@$

goldshell nanoparticles are effective heat-inducing agents. Furthermore, our studies show reasonable cytocompatibility of the particles and suggest their potential application for photothermal therapy.

Material and methods

Barium titanate (BaTiO_3) nanoparticles, 300 nm, 99.9% purity, were obtained from Nanostructured and Amorphous Materials Inc. (Houston, TX, USA). Hydrogen peroxide (H_2O_2) 30% analytical grade, and sodium hydroxide (NaOH) were purchased from Thermo Fisher Scientific (Waltham, MA, USA). Ammonia solution, 30% (NH_4OH), was purchased from Carlo Erba Reagenti SPA (Arese, Milano, Italy). (3-aminopropyl) triethoxysilane (APTES), 99.9% hydrogen tetrachloroaurate trihydrate ($\text{HAuCl}_4 \cdot 3\text{H}_2\text{O}$), 80% tetrakis(hydroxymethyl)phosphonium chloride (THPC), potassium carbonate (99.9%) and sodium borohydride (NaBH_4) were purchased from Sigma-Aldrich (St Louis, MO, USA). N_2 -saturated distilled water was used in all steps. Dulbecco modified Eagle's medium/Ham's F12 (DMEM/HF12), fetal bovine serum (FBS), trypsin, and phosphate buffered solution (PBS) were purchased from Euroclone S.p.A. (Pero, Milano, Italy) the LIVE/DEAD[®] viability assay was purchased from Life Technologies (Carlsbad, CA, USA); the WST-1 assay was purchased from Vinci Biochem Srl (Vinci, Firenze, Italy).

Preparation of gold nanoshells

Functionalization of BaTiO_3 nanoparticles

The functionalization of BaTiO_3 nanoparticles was carried out following our previous approach,⁵⁴ with some modifications. Briefly, 120 mg of BaTiO_3 nanoparticles were added to 20 mL of 1 M nitric acid in a plastic tube. The mixture was sonicated for 2 hours and left to stand overnight. The particles were washed with water until the solution reached approximately neutral pH and were then collected as a pellet by centrifugation at 5000 rpm for 7 minutes. The final pellets were resuspended in water. Next, the particles were treated with 50 mL of hydrogen peroxide at 90°C for 4 hours, providing the surface with the hydroxyl groups required for further functionalization with an organosilane.⁵⁶ The particles were then centrifuged under the same conditions as above and resuspended in 3 mL of water. The obtained BaTiO_3 core nanoparticles were reacted with 140 μL of APTES in the presence of 250 μL of ammonium solution in 46 mL of anhydrous ethanol, thus providing amine groups on the surface for gold seeding in the subsequent step. The mixture was stirred for 3 hours at room temperature and for 2.5 hours at 78°C, cooled and washed three times

with ethanol to remove the reactants and then resuspended in ethanol for storage at 4°C.

Gold nanoparticles

Aqueous solutions of colloidal gold nanoparticles with 1–2 nm diameters were prepared following the method proposed by Duff et al^{57,58} by reducing tetrachloroauric acid with THPC. The solution was gently stirred for 30 minutes and then kept in the refrigerator at 4°C in the dark for 4 weeks before being used in the following step.

Attachment of colloidal gold seeds to amine-functionalized BaTiO_3 nanoparticles

Core BaTiO_3 nanoparticles were seeded with the gold nanoparticles that were prepared in the previous step. Ten milliliters of the prepared colloidal gold nanoparticles was mixed with 1 mL of amine-functionalized BaTiO_3 nanoparticles in a tube and left to stir gently for 2 hours. The seeded gold nanoparticles on the barium titanate surface promote shell formation in the following step. The mixture was centrifuged three times at 5000 rpm. The supernatant was decanted, and the reddish pellet was redispersed and sonicated in 3 mL of distilled water.

Shell formation process

To grow a gold shell on the core nanoparticles, a growth solution (gold hydroxide solution) containing gold reducible salt was prepared.^{59,60} Briefly, to a flask containing a 0.16 mmol solution of potassium carbonate in 27.6 mL of distilled water, a 412.5 μL aliquot of a 25 mM $\text{HAuCl}_4 \cdot 3\text{H}_2\text{O}$ solution was added. The color of the solution gradually changed from transparent yellow to colorless over approximately 30 minutes. The resulting solution was left to stand overnight in the dark. Then, 26.7 μL of the original gold-seeded barium titanate nanoparticles, diluted in 2.5 mL of water, was mixed under vigorous stirring with a 1.9 mL aliquot of gold hydroxide solution. After 10 minutes, 1 mL of ice-cold freshly prepared 0.06 M of water solution of NaBH_4 was added to the mixture and stirred for 5 minutes. The solution changed color from opaque light violet to transparent blue within 5 minutes. The resulting gold nanoshells (NS) were centrifuged three times and redispersed in 1 mL of water for use. A simplified scheme of gold shell formation structure is reported in Figure 1A.

The prepared particles were further studied for their optical and thermal features. The morphology of the gold coverage was examined by a dual beam (FIB/SEM) microscope, Helios (Hillsboro, OR, USA) NanoLab 600i,

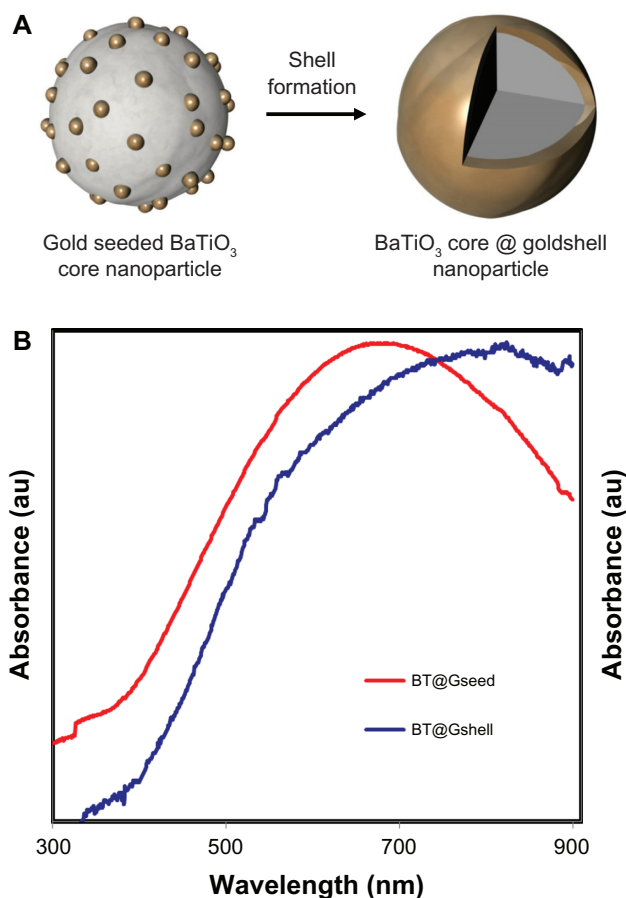


Figure 1 (A) Simplified scheme of gold-shell formation and structure of nanoparticles. (B) UV/Vis/NIR absorption spectrum of gold-seeded BaTiO₃ nanoparticles (red line) and BaTiO₃@gold-shell nanoparticles (blue line).

Abbreviations: BT@Gseed, gold-seeded BaTiO₃ nanoparticles; BT@Gshell, BaTiO₃@gold-shell nanoparticles; NP, nanoparticles; UV/Vis/NIR, ultraviolet-visible-near-infrared.

FEI. Samples were prepared by placing a drop of diluted nanoparticle solution on a silica wafer substrate and allowing the substrate to air-dry before observation.

The NIR absorption properties of the nanoparticles were studied by collecting the absorption spectra of their dispersion in water using a LAMBDA 45 PerkinElmer (Waltham, MA, USA) UV/Vis/NIR scanning spectrophotometer over a wavelength range from 400 to 900 nm. Once it was verified that the sample had the appropriate NIR peak resonance (~800 nm), more nanoparticles were produced for further biological experiments.

Cytocompatibility evaluation and in-vitro testing

Human neuroblastoma SH-SY5Y cells (ATCC CRL-2266) were cultured in DMEM/HF12 (1:1) with 10% fetal calf serum, 100 IU/mL penicillin, 100 µg/mL streptomycin, and 2 mM L-glutamine. Cells were maintained at 37°C in

a saturated humidity atmosphere containing 95% air/5% CO₂.

For viability testing, the SH-SY5Y cells were seeded in 96-well plates (5000 cells per well) and incubated with increasing concentrations of nanoshells (0, 10, 20, 50, and 100 µg/mL). At 24 and 72 hours from the beginning of treatment, cell metabolism was assessed with the WST-1 assay (2-(4-iodophenyl)-3-(4-nitrophenyl)-5-(2,4-disulfophenyl)-2H-tetrazolium monosodium salt, provided in a pre-mixed electro-coupling solution). Cell cultures were treated with 100 µL of culture medium with 10 µL of the pre-mixed solution added for 2 hours, and the absorbance was read at 450 nm with a microplate reader (Victor3; PerkinElmer). A further, qualitative evaluation of nanoshell cytocompatibility was performed with the LIVE/DEAD viability/cytotoxicity kit after 72 hours of treatment. The kit contains calcein AM (4 mM in anhydrous DMSO) and ethidium homodimer-1 (EthD-1, 2 mM in DMSO/H₂O 1:4 (v/v)) and allows for the discrimination between live cells (green-fluorescent stained by calcein) and dead cells (red-fluorescent stained by EthD-1). Cultures were rinsed with PBS, treated for 10 minutes at 37°C with 2 µM calcein AM and 4 µM EthD-1 in DMEM, and finally observed with an inverted fluorescence microscope (TE2000U, Nikon, Tokyo, Japan) equipped with a cooled CCD camera (DS-5MC USB, Nikon). Same experiments were also carried out on differentiating SH-SY5Y cells, cultured for 3 days in a low-serum medium (1%) supplemented with all-trans retinoic acid (20 µM) to evaluate cytocompatibility on a model of neuronal cells.

Nanoparticle internalization has been investigated in cells treated for 24 hours with 50 µg/mL of nanoshells by multimodal microscopy with an in-plane resolution of approximately 300 nm and a resolution of 1 µm along the optical axis. Coherent anti-Stokes Raman scattering (CARS) tuned to the CH₂ bond Raman shift (2850 cm⁻¹) has been exploited to image individual cells. The sum-frequency signals generated by the nanoshells at approximately 450 nm by the combination of 800 nm pump-and-probe pulses with 1045 nm Stokes photons allowed for the positions of the nanoshells to be determined.

In-vitro thermoablation experiments were performed after incubating the cells for 6 hours with 50 µg/mL of nanoshells. Thereafter, the cultures underwent the LIVE/DEAD assay procedure and underwent laser stimulation using a 785 nm laser diode (OBIS 785 LX, Coherent, Santa Clara, CA, USA) overlapping with the wavelength at the maximum surface plasmon resonance (SPR) absorption of the nanoparticles. The nominal power of laser was

50 mW, and the measured power at the sample surface was approximately 10 mW, focused on a spot of approximately 0.04 mm² for a power density of 250 mW/mm². The laser was integrated into the fluorescence microscope, thus offering the advantage of real-time tracking of cell damage. Experiments were performed by applying the laser light for 3 minutes. Control experiments were conducted using the same laser light source to stimulate non-nanoparticle treated cultures.

Statistical analysis

Analysis of the data was performed by an analysis of variance (ANOVA) followed by Student's *t*-test to test for significance, which was set at 5%. The WST-1 tests were performed in six replicates; all the remaining assays were performed in triplicate if not otherwise specified. In all cases, three independent experiments were conducted. The results are presented as the mean value \pm standard deviation.

Results and discussion

Nanoshells and their optical properties

Nanoshells with a core diameter of 300 nm and an approximate shell thickness of 20 nm were designed to have SPR peak absorption in the NIR region. As seen in Figure 1B, a significant spectral red shift with respect to the absorption spectra of the gold-seeded particles (red line) can be interpreted as an increase in the gold coverage, indicating shell formation (blue line). Figure 1B illustrates the maximum absorption at 820 nm for the final nanoparticles, where the large bandwidth of the peak is associated with the size of the core.¹⁷ According to the literature, very small, fresh THPC gold nanoparticles do not establish plasmonic features;^{61,62} nevertheless, aging the gold colloid for a period of 3–4 weeks causes an increase in the size of the particles. The attachment of gold seeds to BaTiO₃ cores was achieved through the existing electrostatic attraction between the positively charged amino-functionalized core particles (through APTES moieties) and the negatively charged gold nanoparticles.⁶³ In general, in an aqueous medium, the measured zeta potential of colloidal particles is strongly affected by the pH of the solution. This fact was taken into account to obtain uniform seeding.

Figure 2A shows a number of partially covered core particles with gold seeds having significant coverage uniformity at this intermediate step of shell growth. Consistent with previous observations,^{63,64} these nanoparticles serve as nucleus sites for the growth of a continuous gold shell by reducing additional gold precursors in their proximity. The reduction

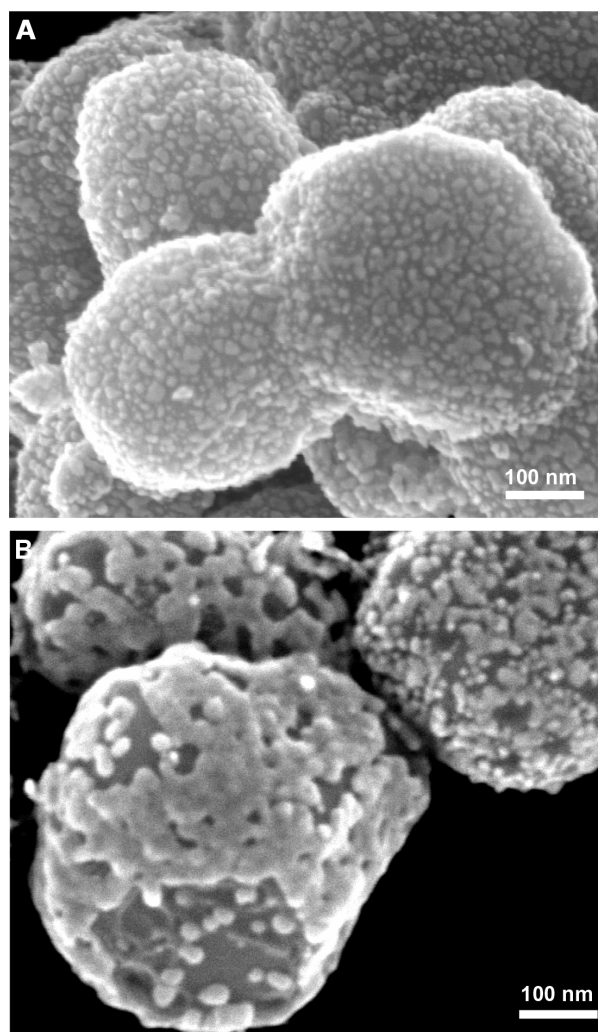


Figure 2 Scanning electron microscopy image of BaTiO₃ nanoparticles (A) seeded with gold and (B) shell formation.

of the gold in the last step allows the seed nanoparticles on the surface to grow and merge, thereby causing a systematic red shift of the SPR into the NIR (820 nm) region.^{39,63} The coalescence and merging of individual and separated gold nanoparticles into a shell are shown in Figure 2B, confirming shell formation, in this case with a smooth-looking shell. A scanning electron microscopy image and energy dispersive X-ray analysis on a single nanoshell showing its elementary composition is also reported (see Figure S1). As the dual beam images demonstrate, the core particles are rather spheroid with about 300 nm diameter and approximately 20 nm shell thickness.

To attain the most uniform shell layer, the surface of the core nanoparticles must be pre-treated with acid and then hydrogen peroxide, because of the essential role of having homogenous silanization and therefore seeding. When BaTiO₃ nanoparticles are exposed to acidic solution, the

BaCO₃ impurities present on the particle surface dissolve in water, leaving the particle surface similar in structure to a classic metal oxide lattice (eg, TiO₂). The pH of acidic solution affects the amount of Ba²⁺ leaching from the surface of the particle.⁶⁵ Placing the core particles in contact with an aqueous solution of hydrogen peroxide in optimized conditions provides the surface of the particles with an adequate number of hydroxyl groups that are essential for the attachment of APTES molecules. However, in the optimized conditions, H₂O₂ removes the adsorbed carbonate from the surface of the nanoparticle in a manner comparable to what occurs in a weak aqueous acetic acid solution.⁶⁶ As a consequence, when the particles were placed in H₂O₂ immediately after treatment with acid (nitric acid in our case), skipping the washing step or failing to properly wash the particles to reach neutral pH, the resulting mixture appeared yellowish with large clod-like aggregations of particles that were not easily broken. On the basis of these observations, the combination of acid and H₂O₂ causes an excessive loss of barium ions that leads to the degradation of the particle surface and its structure.

Nanoshell cytocompatibility and internalization

Cell viability was investigated with the LIVE/DEAD viability/cytotoxicity kit. The results on the compatibility of nanoshells revealed no evidence of cell membrane damage after the cells were treated with concentrations up to 50 µg/mL (Figure 3A) for 72 hours of incubation, demonstrating viabilities comparable to the control cultures (less than 5% of red, ie, necrotic, cells). At 100 µg/mL, however, a consistent decrement of cell density and increased cell death were observed.

These results were quantitatively confirmed by the WST-1 assay, which demonstrated excellent metabolic activity in cells treated with concentrations of nanoshells up to 50 µg/mL (Figure 3B) and no statistically significant decrement of viability compared with the controls after both 24 and 72 hours. After 72 hours, a highly tangible decrease of cell metabolism was observed in cells treated with 100 µg/mL of nanoshells (about 40%, $P < 0.05$). Analogous results have been achieved on differentiating SH-SY5Y cells, as a model of human neurons. No significant variation in metabolic activity, viability, and differentiation status was highlighted in cells treated with 0–50 µg/mL of nanoshells (see Figure S2).

Figure 4 shows results of internalization assessment: a composite image is reported, where the green channel

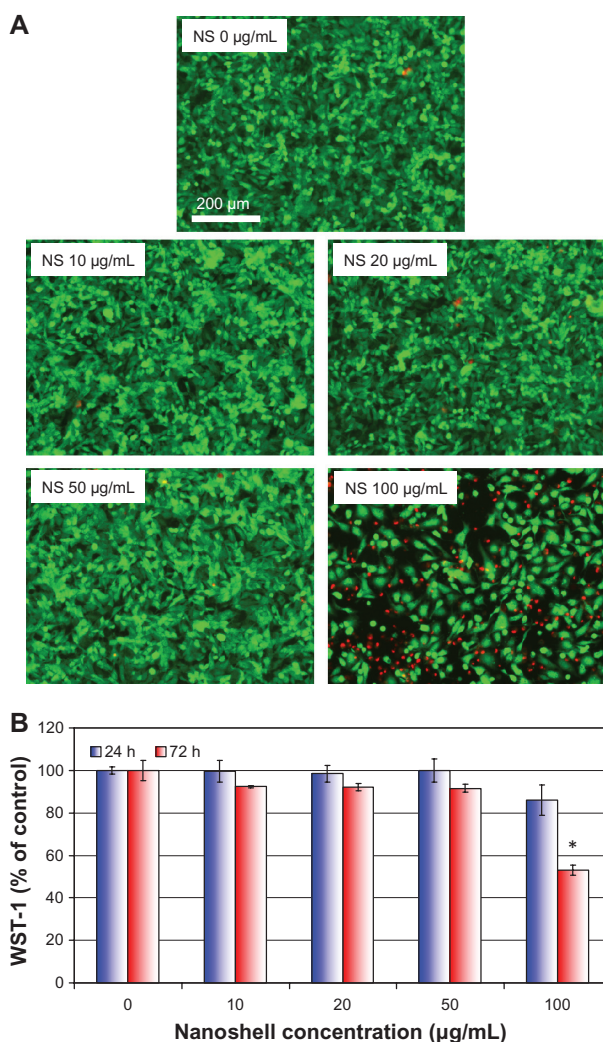


Figure 3 Cytocompatibility results: (A) LIVE/DEAD[®] assay and (B) WST-1 assay at 24 and 72 hours; * denotes $P < 0.05$.

maps the intensity of the CARS signal, and the red channel indicates the intensity of the sum-frequency signals. The dark spots are the shadows of nanoshells below or above the imaging plane. Example of CARS Z-stack acquisition is also reported (see Figure S3).

Considering all the collected data, an operative dose of 50 µg/mL was thus used for the following experiments.

Photothermal ablation

Gold nanoshells have been demonstrated to be suitable heating agents for photothermal ablation of cancerous cells.⁶⁴ Halas and co-workers demonstrated that gold nanoshells can be used for plasmonic photothermal therapy in the NIR region with two different modes for targeting the cells: passive^{25,50} and active.²⁹ In addition to the two modes, another approach for targeting tumor cells is the enhanced permeability and retention (EPR) effect. According to the EPR effect, the

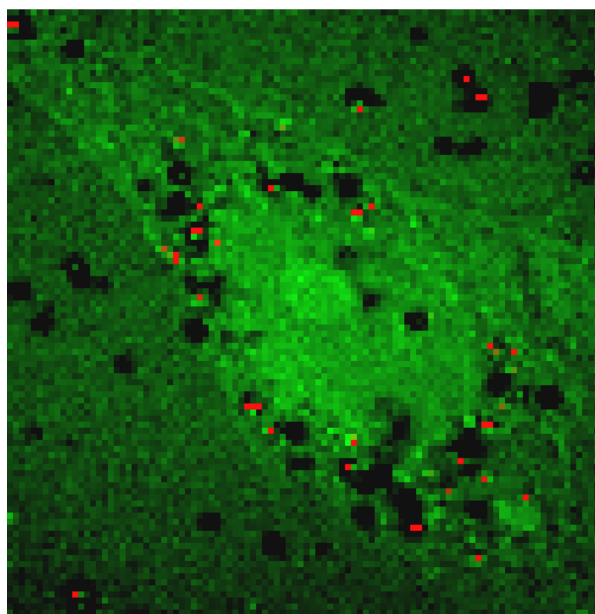


Figure 4 Multimodal image of an individual cell. The green channel maps the intensity of the coherent anti-Stokes Raman scattering signals from the CH_2 bonds; the red channel indicates the sum-frequency signals from the nanoshells. The field is $50 \mu\text{m} \times 50 \mu\text{m}$.

concentration of nanoparticles that accumulate in tumor tissues is greater than that in normal tissues or organs.^{67,68} Since it was discovered, the EPR effect has been recognized as the standard criterion for designing new anticancer agents.⁶⁹ It is assumed that based on the EPR effect, small nanoparticles ($<400 \text{ nm}$), including nanoshells, selectively accumulate in the tumor tissue by the passive extravasation associated with the leaky tumor vascularity.^{28,30,69} In theory, upon accumulation within tumors, nanoshells interfere in cell damage and aid in cancer/tumor treatment by absorbing NIR light and transforming it into heat as a result of their plasmonic characteristics. When using nanoshells that are designed to be scatterers (higher scattering cross-sections), cancerous cells could be detected even at an early stage. Here, we provide an in-vitro demonstration of the thermal ablation of neuroblastoma cell-lines using nanoshells designed to absorb NIR light.

Figure 5 shows cells incubated with nanoshells and stained with the LIVE/DEAD stain before and after treatment with the NIR laser. The ability of nanoshells to cause cell death under NIR light irradiation is demonstrated in the fluorescence images. Surviving and dead cells are shown in green and red, respectively. A bright field image of the sample and an image of the spot of laser exposure are also included in Figure 5. The left column illustrates cells that were incubated with nanoshells prior to exposure to the laser light in the bright field mode (Figure 5A) and in the

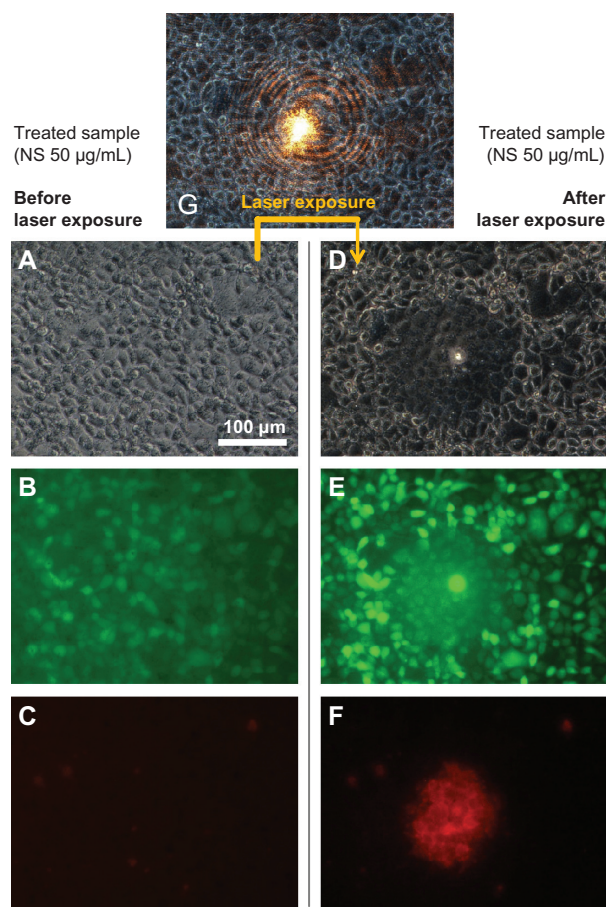


Figure 5 Photo-induced thermal ablation of neuroblastoma cells incubated with nanoshells prior to (left main column) and after (right main column) exposure to laser light. Cells are labeled with calcein (live cells in green) and with EthD-1 (dead cells in red).

fluorescence mode, indicating live (green) cells (Figure 5B) and dead (red) cells (Figure 5C), respectively. After exposure to laser light, significant cell death occurred, as shown in Figure 5E. No live cells remained within the spot of laser exposure or its vicinity, which is evident by the hole-like spot in the center of the green fluorescent background. The red spherical stain in Figure 5F corresponds to this mentioned zone and shows the dead cells within the border of the spot of laser exposure. Figure 5D shows a more tangible image of dead cells in the spot of laser exposure. Further control experiments to verify the influence of the laser alone were performed on cells not incubated with the nanoshells prior to exposure to laser light. No cell death is evident in the results of these experiments; cell damage is not observed in either of the control cases, confirming that highly effective, localized, and selective photothermal therapy would result from combining nanoshells and laser light at the appropriate wavelength for SPR absorption of the particles. Images for negative control on cells nonin-

cubated with nanoshells and treated with laser are shown in Figure S4.

Conclusion

Previous work by our group investigated BaTiO₃ nanoparticles and their potential applications in biology.⁷⁰ We recently reported our preliminary research on the development of BaTiO₃@goldshell nanoparticles.⁵⁶ In the present study, we demonstrated how the surface-plasmon-based-photo-absorption characteristics of these particles can effectively induce cell death selectively within the spot of laser exposure. The nanoshells were prepared through an accurate process combining molecular self-assembly and colloidal growth chemistry⁷¹ and showed suitable cytocompatibility at concentrations up to 50 µg/mL. The prepared nanoparticles were passively accumulated in cells and irradiated by an NIR laser. Localized photo-induced cell damage resulted from the studies carried out on neuroblastoma cell-lines. Future studies will be focused on addressing targeted delivery of nanoshells to specific cells and/or tissues, and on synergetic imaging and therapy of tumor cells. Thanks to their high second-order nonlinear optical susceptibility, BaTiO₃ nanoparticles are in fact considered good candidates to be exploited in SHG techniques. BaTiO₃ nanoparticles show substantial advantages over other SHG labeling probes, such as their ideal signal transfer,^{73,74} that make them suitable for use in in-vivo imaging. The simultaneous exploitation of piezoelectricity, SHG, and plasmon resonance, render therefore BaTiO₃@goldshell nanoparticles a promising theragnostic device in cancer research.

Executive summary

1. In this paper, we describe the synthesis and characterization of BaTiO₃@goldshell nanoparticles.
2. An optimal dose range of BNNTs (0–50 µg/mL) was identified for their in-vitro administration to cell cultures.
3. Nanoparticle internalization by neuroblastoma cells was verified through CARS.
4. The optical response of the prepared nanoshells was exploited to achieve cell destruction through hyperthermia following NIR laser stimulation.

Disclosure

The authors report no conflicts of interest in this work.

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Supplementary figures

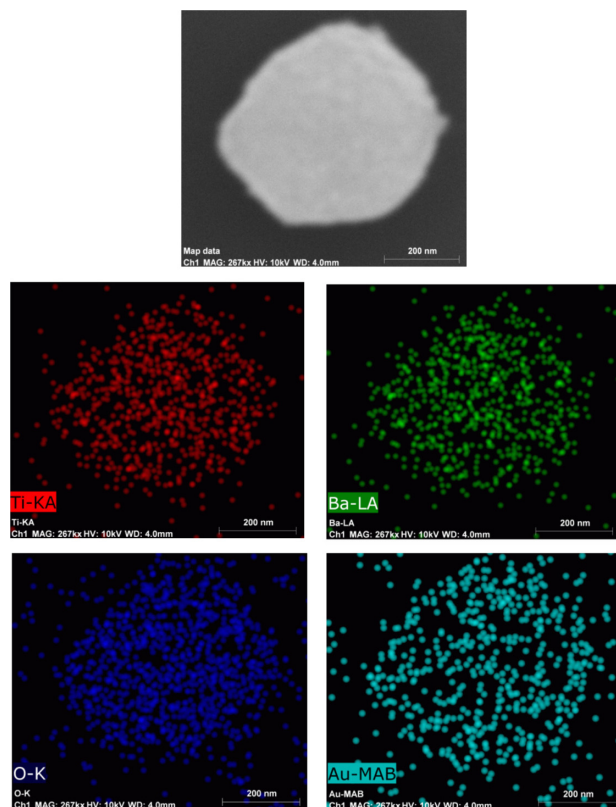


Figure S1 SEM image and EDX analysis on a single nanoshell, showing its composition in Ti, Ba, O and Au.

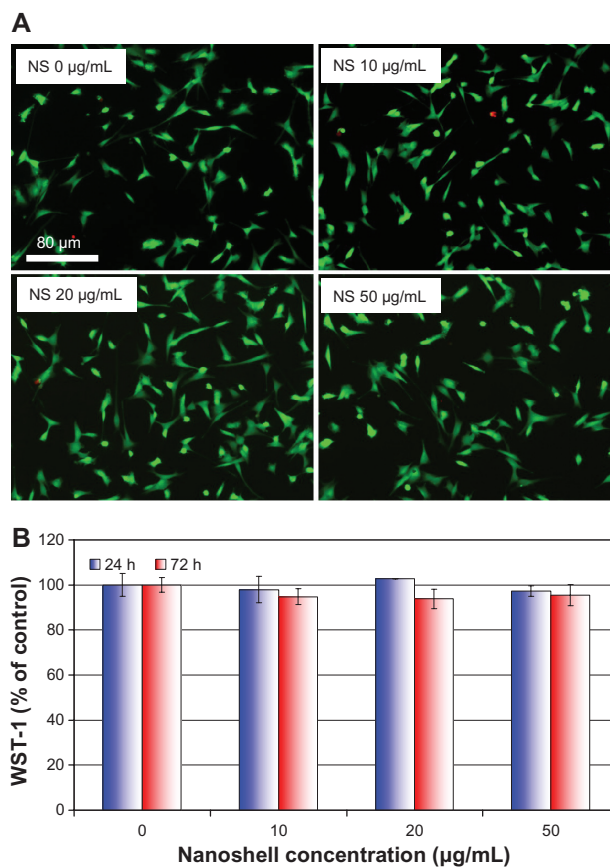


Figure S2 Cytocompatibility results on differentiating SH-SY5Y cells: the Live/Dead assay (A) and WST-1 assay at 24 and 72 h (B).

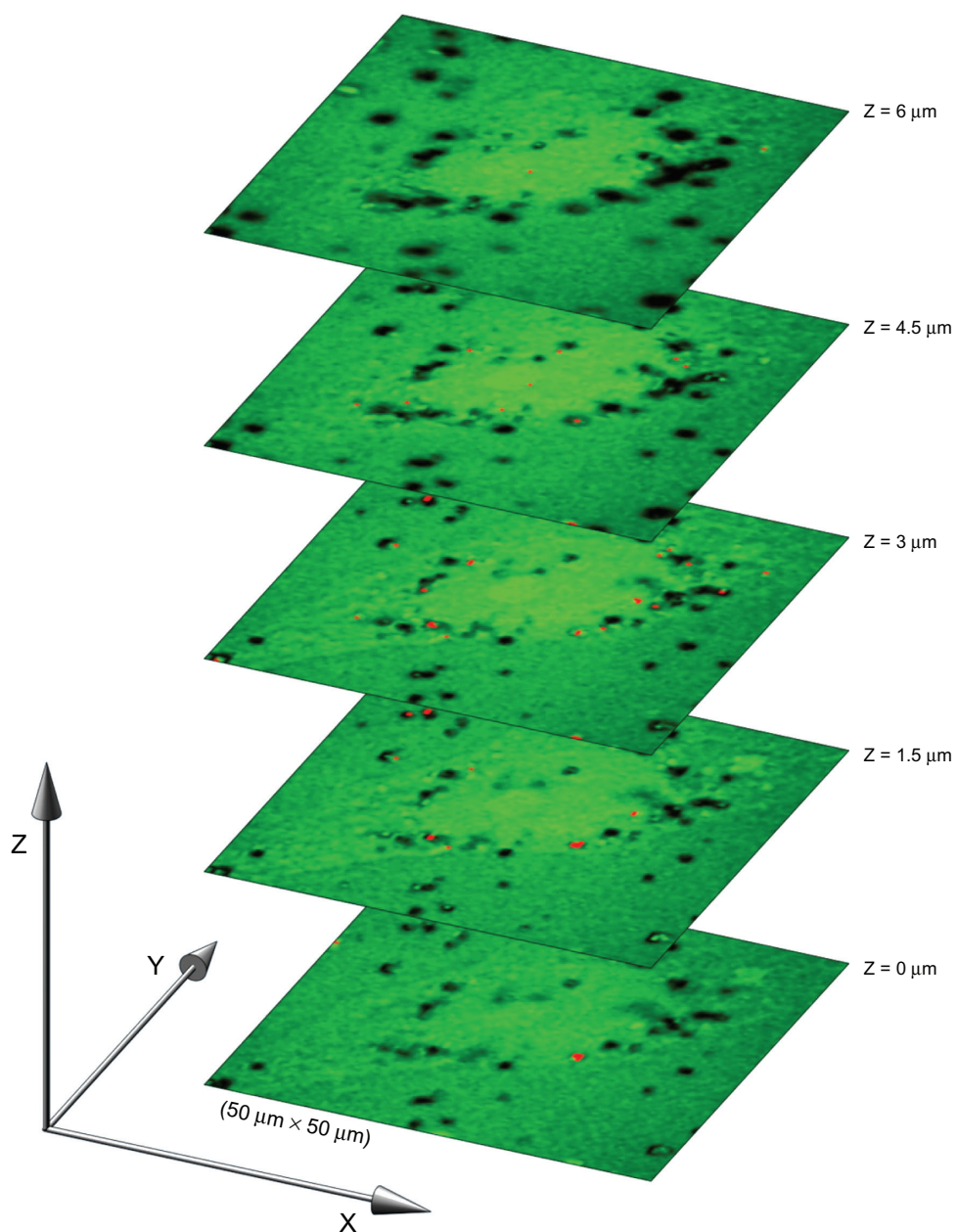


Figure S3 Example of CARS Z-stack acquisition of single cell including nanoshells. The green channel maps the intensity of the CARS signals from the CH₂ bonds; the red channel labels the nanoshells. The field is 50 μm × 50 μm.

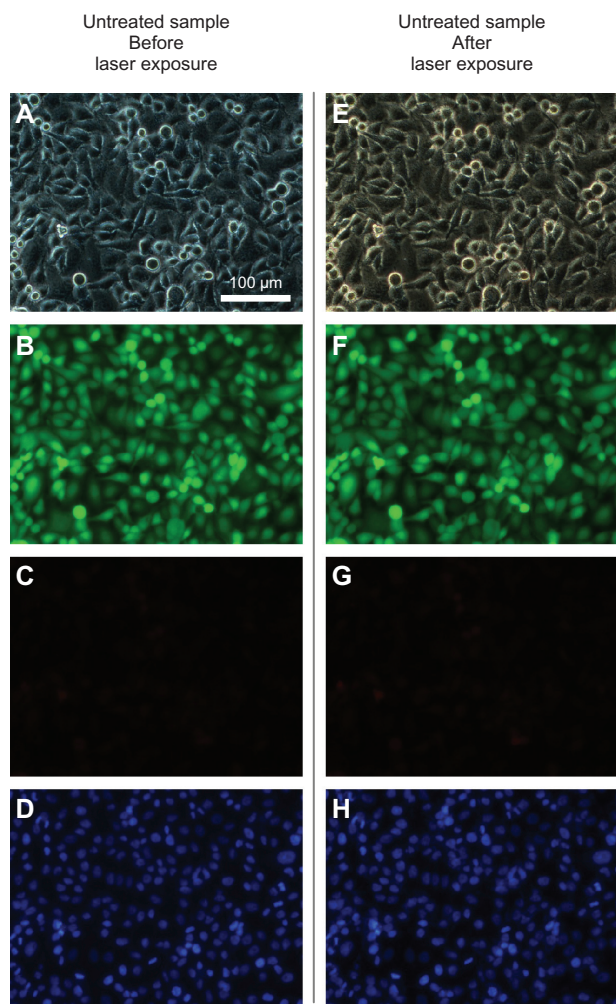


Figure S4 Neuroblastoma cells incubated without nanoshells before (left column, **A–D**) and after (right column, **E–H**) exposure to laser light. Cells are shown in bright field (**A** and **E**), labeled with calcein (in green, **B** and **F**), with EthD-1 (dead cells in red, **C** and **G**), and with nuclei labeled with Hoechst (in blue, **D** and **H**).

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