Animal heat activated cancer therapy by a traditional catalyst TiO$_2$-Pd/graphene composites

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Cancer therapy is one of the most important challenges in clinical medicine. So far, different methods have been developed for cancer therapy, such as radiation therapy, surgery, chemotherapy, and photodynamic therapy. Here we propose a new concept for cancer therapy, i.e., killing the cancer cells simply via reactive oxygen species (ROS) generated by TiO$_2$-Pd/graphene composites. Activated by animal heat of 37 °C, the electrons in the valence band can be excited to the conduction band of TiO$_2$ via the energy levels of Pd species and graphene, generating ROS without light irradiation or electric excitation. The tumors in BALB/c mice are successfully regressed at animal heat without any other external conditions, such as radiation, UV, visible and IR irradiation. Our results suggest that the design of animal heat activated cancer therapy is a feasible concept for practical applications of cancer treatments.

Cancer can be usually treated by surgery, chemotherapy, radiation therapy, hormonal therapy, targeted therapy (including immunotherapy such as monoclonal antibody therapy) and synthetic lethality$^{1-7}$. The choice of therapy depends upon the location and grade of the tumor and the stage of the disease, as well as the general state of the patient (performance status). However, there are still serious side effects for these cancer treatment methods, which may cause damage on human bodies. Therefore, designing and developing new concept about the complete removal of the cancer without damage to the rest of the body is still the ideal goal of treatment.

During the last decades, many photosensitizers have been employed extensively for photodynamic therapy$^{8-10}$. It is known that photodynamic catalyst can generate charge carriers and then produce reactive oxygen species (ROS) on surface under irradiation. These ROS can damage biological DNA and proteins, damaging and killing the cancer cells eventually$^{3,6}$. In addition, photothermal treatment uses near-infrared absorbing nanoparticles to generate heat, resulting in thermal ablation to kill cancer cells. CuS@MSN based theranostic nanoparticles have been designed and synthesized for tumor vasculature targeting and photothermal therapy$^2$. The Au–Cu nanocrystals exhibit a notable photothermal effect to kill cancer cells irradiated by a 808 nm laser$^{11}$. However, irradiation necessarily used for photodynamic and photothermal cancer treatment is difficult to be applied in vivo system. Therefore, it is of great importance to design and develop new cancer therapy concepts without any additional irradiation.

Herein, we introduced a new concept, animal heat activated cancer therapy with a traditional catalyst. The ROS can be generated by TiO$_2$-Pd/graphene without any external conditions, which could kill the cancer cells and regress the tumors in BALB/c mice under animal heat (37 °C), implying the feasibility of brand new concept for the cancer treatment. This concept may possess potential advantages for the practical cancer therapy, such as less body damage, selectivity on killing cancer cells and normal body cells.

Results and discussion

Raman spectra, XPS and HRTEM are applied to investigate the structure of TiO$_2$-Pd/graphene. As shown in Fig. 1A, the Raman peaks of anatase at 144 cm$^{-1}$ (E$_g$), 194 cm$^{-1}$ (E$_g$), 396 cm$^{-1}$ (B$_1g$), 516 cm$^{-1}$ (A$_1g$ and B$_1g$), and 638 cm$^{-1}$ (E$_g$) are observed in TiO$_2$ and TiO$_2$-Pd samples. For TiO$_2$/graphene and TiO$_2$-Pd/Graphene, besides the Raman peaks of anatase, the peaks of reduced graphene at 1315 and 1585 cm$^{-1}$ are detected$^{12,13}$. Similar

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results are obtained from the XRD patterns (Fig. S1). No other XRD peaks, such as PdO can be observed. The cell volume and lattice parameters of TiO2-Pd x% and TiO2-Pd/Graphene x% derived from XRD data remain almost unchanged (Fig. S2 and Table S1) compared with TiO2. This suggests that Pd ions are not doped into TiO2 lattice in substitutional or interstitial mode, as the ionic radius of Pd2+ (86 pm) is much larger than that of Ti4+ ions (68 pm). Therefore, it can be deduced that the Pd ions might exist on the surface of TiO2. Moreover, according to the XPS results, the Cl 2p3/2 peaks (198.5 eV) for TiO2-Pd/Graphene (Fig. S3A) locates between that of TiCl4 (198.2 eV) and PdCl2 (198.9 eV), ascribed to the surface O–Pd–Cl structure.14. Two pairs of doublet Pd 3d peaks can be observed for TiO2-Pd/Graphene (Fig. 1C). One Pd3d5/2 peak at 337.7 eV is attributed to –O–Pd–Cl structure (i.e., one Pd2+ ion is linked with one Cl– ion and one unsaturated oxygen ion) on the surface. The other peak at 336.2 eV for Pd 3d5/2 is ascribed to –O–Pd–O– species on the surface (i.e., one Pd2+ ion is linked with two unsaturated oxygen ions, which has been confirmed by our previous work).14. The molar ratio of Pd/Ti for TiO2-Pd and TiO2-Pd/Graphene is 4.19%/100% and 4.14%/100%, respectively. Moreover, the C1s peak at 284.3 eV (Fig. S3B) is ascribed to the graphene. HRTEM image of TiO2-Pd/Graphene (Fig. 1B) confirms TiO2-Pd nanoparticles are attached on the surface of graphene. In addition, the peak of 1216 cm−1 in the FT-IR spectra of TiO2-Pd/Graphene is ascribed to vibration of C–O bond (Fig. S5). The XPS Ti 2p spectra (Fig. S24 and S25) also confirms the graphene and TiO2-Pd are connected via the Ti–O–C bonds.

The surviving fraction of the A549 cells was measured by standard 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyl-tetrazolium bromide (MTT) assay to evaluate the ability of killing cancer cells for all samples at 37 °C, as shown in Fig. 2A. The surviving fraction of the A549 cells is about 96% for TiO2, 85% for graphene, 71% for TiO2/graphene, 53% for TiO2-Pd and 16% for TiO2-Pd/Graphene, respectively, when the concentration is 25 μg/ml after 16 h of incubation at 37 °C. Moreover, it is found that the surviving fraction of cancer cells (16%) for TiO2-Pd/Graphene is about 1/5 times as that for pure TiO2. With the increase of sample concentration, the performance of killing cancer cells falls in order of TiO2 < graphene < TiO2/graphene < TiO2-Pd < TiO2-Pd/Graphene. Furthermore, we have also checked the surviving fraction of cancer cells treated with TiO2-Pd x% and TiO2-Pd/Graphene x% and the surviving fraction is the lowest for TiO2-Pd1.5% and TiO2-Pd/graphene40% (Fig. S6). Furthermore, the increased Pd content (> 1.5%) and graphene content (> 40%) would inhibit the performance of killing cancer cells. Thus, the introduced Pd species and graphene into TiO2 system is the important key factor for the improved ability of killing cancer cells. The optical microscopy images of A549 cancer cells in the presence of samples (100 μg/mL) are shown in Fig. S9. Most of the A549 cells survive well in the blank experiment (without the addition of any samples) (Fig. S9a). The TiO2 can hardly influence the surviving of the cancer cells (Fig. S9b). It is found that the relative amount of healthy cancer cells decreased for TiO2/graphene (Fig. S9c) and TiO2-Pd (Fig. S9d). As we expected, seldom healthy cancers cell can be observed in Fig. S9e for TiO2-Pd/Graphene samples, suggesting its high efficiency on killing cancer cells. These results are in good agreement with MTT results in Fig. 2A.

The effect of temperature on the surviving fraction of cancer cells treated with various concentration of TiO2-Pd/Graphene for 4 h is evaluated in Fig. 2B. The surviving fraction of cancer cells for TiO2-Pd/Graphene decreased with the increased system temperature. When the concentration of TiO2-Pd/Graphene is about 25 μg/ml, the surviving fraction of A549 cells are approximately 80%, 70%, 60%, 55%, and 50% at 4, 12, 25, 37 and 39 °C, respectively. Similar results are observed in the synovial cells (Fig. S7). This indicates the system
temperature has great influence on the activity of killing cancer cells for TiO$_2$-Pd/graphene, and TiO$_2$-Pd/graphene can effectively inhibit the growth of cells at animal heat condition (37 °C).

Fig. S8 shows the relationship between the surviving fraction of the different (A549 and smooth muscle cells) cells and concentration of TiO$_2$-Pd/graphene. It is found that only about 10% of cancer cells survived and 40% of smooth muscle cells survived at 37 °C for 16 h with a concentration of 25 μg/mL. This suggests a possible selectivity on cancer therapy for TiO$_2$-Pd/graphene. It is expected that further investigation should be carried out by combining other techniques, such as targeted therapy, tumor injection, magnetic control to avoid the side effects on normal cells.

For in vivo animal heat activated therapy study, we employed subcutaneous 4T1 xenograft model in BALB/c mice to study the efficacy of animal heat activated cancer therapy (37 °C) after treatment by PBS (phosphate buffer saline), TiO$_2$, TiO$_2$-Pd and TiO$_2$-Pd/graphene. Twelve tumor-bearing mice were randomly and evenly divided into four groups, which were intratumorally injected with PBS, TiO$_2$, TiO$_2$-Pd and TiO$_2$-Pd/graphene, respectively. Figure 2C shows the thorough regression of tumor volume is observed only in the group with intratumoral injection of TiO$_2$-Pd/graphene, while the volume of tumor increases for the group with PBS and TiO$_2$ and remains unchanged for the group with TiO$_2$-Pd. For the group with intratumoral injections of TiO$_2$-Pd/graphene, the average relative tumor volume decreased to approximately 50% and for the group with intratumoral injections of TiO$_2$-Pd, the average relative tumor volume remains almost unchanged. Figure 2D displays the representative photographs of tumors on mice that the intratumorally injected with PBS control and TiO$_2$-Pd/graphene. The growth of tumor is inhibited when injected with TiO$_2$-Pd/graphene. These experiment results indicate that TiO$_2$-Pd/graphene is an effective functional material to inhibit cancer growth under animal heat (37 °C) without external irradiation or heating.

To get the physical insight of the band structure and density of states for TiO$_2$-Pd/graphene, the theoretical calculation results of pure TiO$_2$, TiO$_2$-Pd and graphene are shown in Fig. S12–S15. Pure TiO$_2$ indicates an indirect band gap of 2.46 eV (Fig. S12). The zero point corresponds to highest level that electrons occupy. For TiO$_2$-Pd, some new quasi-continuous energy levels in the band gap (Fig. S13 and S14) are attributed to the contribution of –O–Pd–Cl and O–Pd–O species. Meanwhile, it is obvious that the states of Pd 4d for –O–Pd–Cl and O–Pd–O species plays a great role to make the slightly connection between the conduction band and valence band. For TiO$_2$-Pd (Fig. S13 and S14), the density of states for the conduction band and valence band overlapped with that for the energy levels of Pd species (Pd 4d), implying that the electrons in the valence band can transfer to the energy levels of Pd species (Pd 4d) or even conduction band excited by external energy. Moreover, the graphene (Fig. S15) exhibit as a zero-gap semiconductor, implying electrons can transfer freely in the energy band of graphene. These theory calculation results imply that the electrons can easily transfer between TiO$_2$-Pd and graphene when TiO$_2$-Pd/Graphene are excited by external energy such as animal heat. These thermal excited electrons and holes can transfer to the surface to further generate ROS. This theoretical calculation can be further demonstrated by the UV – Vis absorption spectra and XPS valence band spectra (Fig. S16–S18).
In Fig. S16, according to the above theory calculation, the stronger absorption in visible region for TiO₂-Pd is caused by the electron transition from valence band to the quasi-continuous energy levels of Pd species (Pd 4d) or from the quasi-continuous energy levels of Pd species (Pd 4d) to the conduction band. Moreover, for TiO₂/graphene and TiO₂-Pd/graphene, the strong absorption in visible region is due to the continuous energy band structure of graphene.

For XPS valence band spectra (Fig. S18; the work function of instrument: 4.10 eV), the valence band top for pure TiO₂ is 2.75 eV (2.35 eV vs NHE), and the highest occupied state level for graphene is about 2.29 eV (1.89 eV vs NHE). For TiO₂-Pd, the valence band top is almost the same as that of TiO₂, and a small hump from 2.75 to 0.10 eV (2.35 to −0.3 eV vs NHE) is attributed to the energy level of Pd species (Pd 4d) occupied by electrons. For TiO₂-Pd/graphene, the highest energy levels that electrons occupy extend to −0.6 V (−1.0 eV vs NHE), owing to the synergetic effect of graphene and TiO₂-Pd. These results imply the electrons in the valence band can transfer to the energy levels of Pd species (Pd 4d) or even the conduction band of TiO₂ through the quasi-continuous energy levels freely and continuous energy band structure of graphene excited by the external energy. According to the results, the schematic band structure is determined for TiO₂-Pd/graphene, as shown in Fig. 3E.

The Raman spectra, cyclic voltammetry curves and fluorescence spectra of the DCF are applied to investigate the relationship between the generation of ROS and temperature. The Raman spectra of TiO₂-Pd and TiO₂-Pd/graphene (Fig. S19) indicates the peak intensity was enhanced with the increase of temperature, suggesting an enhanced vibrating energy of lattice, benefiting the electrons’ transfer to surface of sample.

The cyclic voltammetry curves of pure TiO₂, graphene, TiO₂/graphene, TiO₂-Pd and TiO₂-Pd/graphene at 37 °C (in Fig. 3A) indicate the ability to give and receive electrons increases when Pd species and/or graphene are introduced into TiO₂ system. For the cyclic voltammetry curves of the TiO₂-Pd/graphene at different temperatures, the redox peaks of TiO₂-Pd/graphene are improved with the increase of temperature (Fig. 3B). This suggests that TiO₂-Pd/Graphene may exhibit enhanced ability to give and receive electrons under thermal excitation.

The amounts of ROS (OH, O₂⁻, etc.) generated by TiO₂, graphene, TiO₂/graphene, TiO₂-Pd and TiO₂-Pd/graphene at 37 °C were evaluated by the photoluminescence (PL) intensity of the DCF (Fig. 3C). No PL peak is detected in the blank experiment. For TiO₂ and graphene, a weak PL peak is found, suggesting a small amount of ROS formed. Moreover, for TiO₂/graphene and TiO₂-Pd, the PL intensity further increases, suggesting more ROS can be generated than TiO₂. The PL intensity for TiO₂-Pd/graphene is the strongest among all the samples, indicating TiO₂-Pd/graphene is most effective to generate ROS among all the samples at 37 °C. Figure 3D shows generation amount of ROS for TiO₂-Pd/graphene at different temperatures. It is noted that the PL intensity significantly increases with the increase of temperature, indicating the increased temperature is benefit for the generation of reactive oxidation species. This also confirms that the electrons and holes in TiO₂ can be excited by heat and transfer to surface to form ROS.

Figure 3. Cyclic voltammetry of all samples at 37 °C (A) and TiO₂-Pd/graphene at different temperature (B). Verification of ROS generation ability for all samples at 37 °C (C) and TiO₂-Pd/Graphene at different temperatures (D). Schematic band structure of TiO₂-Pd/graphene composite (E).
Based on the aforementioned discussion, the mechanism of generating ROS (such as ·OH, O$_2^-$) and kill the cancer cells for all samples can be explained using the schematic band structure shown in Fig. 3E. The electrons in the conduction band (~0.45 eV, vs NHE) excited by animal heat (37 °C), whose potential is more negative than that of O$_2$/O$_2^-$ (~0.33 eV, vs NHE), can transfer to the surface and directly captured by the adsorbed O$_2$ molecules on the surface to form O$_2^-$ active species as shown in Eq. (1). The holes in the valence band (~2.65 eV, vs NHE), whose potential is more positive than H$_2$O /H$^+$ (0.82 V, vs NHE), are captured by surface absorbed H$_2$O molecules to form O$_2$ and H$^+$, which can further react with thermal electrons to produce the hydroxyl free radical OH· as shown in Eqs. (2), (3) and (4).

$$O_2 + e^- = O_2^- \quad (1)$$
$$E_{\text{redox}}^0 = -0.33 \text{ V vs NHE}$$

$$2H_2O + 4h^+ \rightarrow O_2 + 4H^+ \quad E_{\text{redox}}^0 = 0.82 \text{ V vs NHE}$$

$$O_2 + 2H^+ + 2e^- = H_2O_2 \quad (2)$$

$$H_2O_2 + e^- = OH^- + HO^- \quad (3)$$

$$OH^- + h^+ = HO \quad (4)$$

For pure TiO$_2$, electrons can hardly be excited to the conduction band by the animal heat (37 °C), leading to a poor killing cancer cell activity, because of large band gap (3.0 eV). For TiO$_2$-Pd, a small amount of electrons can be excited by the animal heat and transfer to the conduction band through the quasi-continuous energy levels of Pd species, leaving holes in the valence band of TiO$_2$. For TiO$_2$/graphene, the electrons could also be excited by the animal heat and transfer to the conduction band of TiO$_2$ through the continuous energy levels of graphene, leaving holes in the valence band of TiO$_2$. The electrons and holes would react with adsorbed O$_2$ and H$_2$O molecules on surface to form a small amount of ROS, which would kill the cancer cells directly. For TiO$_2$-Pd/graphene, the electrons in the valence band can accept the energy from the animal heat and transfer to the conduction band through the quasi-continuous energy levels of Pd species and the continuous energy levels of graphene, generating more charge carriers than TiO$_2$-Pd and TiO$_2$/graphene. Therefore, more ROS can be produced and the TiO$_2$-Pd/graphene exhibited the better capability of killing cancer cells than the other samples.

In summary, we have demonstrated a new concept for developing high efficient animal heat activated cancer treatment for TiO$_2$-Pd/graphene. The electrons and holes can be excited through the energy levels of Pd species and graphene at animal heat, generating ROS which can further kill the cancer cells. This may afford a feasible and efficient approach for cancer therapy application, without any other external conditions such as radiation, UV, visible and IR irradiation that may cause serious body damage.

**Methods**

**Catalyst preparation.** Synthesis of Pd-TiO$_2$. All chemicals used were of analytical grade and the water was deionized water (>18.2 MΩ cm). At room temperature, certain volume of PdCl$_2$ (0.1036 mol/L) solution was mixed with 40 mL of ethanol. Then 1 mL of HCl solution (12 mol/L) and 12 mL of Ti(OC$_4$H$_9$)$_4$ was added dropwise into the mixture under vigorous stirring. The mixture was stirred until the formation of TiO$_2$ gel, followed by being aged for 24 h. The obtained gels were dried at 373 K for 12 h and annealed at 723 K in a muffle for 2.5 h. The resultant samples were denoted as Pd-TiO$_2$. Pure TiO$_2$ was prepared using the same procedure, while without the addition of PdCl$_2$ solution. Unless stated otherwise, the nominal molar ratio of Pd$^{2+}$ to Ti$^{4+}$ is fixed at 1.5% in the precursor and. For comparison, other molar ratios were also used for Pd$^{2+}$ to Ti$^{4+}$ (such as 0.5%, 1.0%, 2.0%, 2.5% and 3.0%).

Synthesis of graphene oxide (GO). GO was prepared from crystalline flake graphite powder according to the modified method reported by Hummers and Offeman. In brief, 10 g of graphite powder and 5 g of NaNO$_3$ were added into 230 mL of cooled (273 K) concentrated H$_2$SO$_4$. Then, 30 g of KMnO$_4$ was added gradually with continuous stirring and cooling, and the temperature of the mixture was maintained below 293 K. After the ice bath was removed, the mixture was stirred at 308 K for 30 min. 460 mL of distilled water was added slowly to cause an increase in temperature to 371 K, and the mixture was maintained at that temperature for 15 min. The reaction was terminated by addition of 1.4 L of distilled water followed by 25 mL of 30% H$_2$O$_2$ aqueous solution. The solid product was separated by centrifugation and washed repeatedly with 5% HCl solution (2L) and deionized water until sulfate anion could not be detected with BaCl$_2$. The resultant solid was dried in vacuum at 323 K to obtain GO.

Preparation of TiO$_2$-Pd/graphene. GO was first dissolved in deionized water by ultrasonic treatment for 20 min. Then, Pd-TiO$_2$ was added into the GO colloidal solution and the mixture was ultrasonic for another 1 h to obtain a homogeneous suspension. The resultant composite was collected by drying at 333 K and then
triturated to powder in an agate mortar. Finally, the powder was calcined at 300 °C for 2 h under Ar atmosphere. The resulting products were TiO2-Pd/graphene.

Characterizations. The XRD patterns were acquired using a Rigaku D/max 2500 X-ray diffraction spectrometer (Cu Kα, λ = 1.54056 Å). The average crystallite size was calculated according to the Scherrer formula (D = k λ/β cosθ). Raman spectra were taken on a Renishaw inVia Raman microscope by using the 785 nm line of a Renishaw HPN IR785 semiconductor laser. The Fourier Transform Infra-Red (FT-IR) spectra were recorded for KBr disks containing the powder sample with an FT-IR spectrometer (MAGNA 560). The BET surface areas of the samples were determined by nitrogen adsorption−desorption isotherm measurements at 77 K (Micrometrics Automatic Surface Area Analyzer Gemini 2360, Shimadzu). XPS measurements were carried out by using a Thermo ESCALAB 250 spectrometer with an Al Kα monochromator source and all the binding energies were calibrated to the adventitious C1s peak at 284.8 eV. Diffuse reflectance UV−visible (UV−Vis) absorption spectra were recorded on a UV−Vis spectrometer (U-4100, Hitachi). ROS in the presence of samples was qualitatively detected by the H2DCF-DA assay. Photoluminescence (PL) spectra were acquired by using the 325 nm line of a nanosecond Nd: YAG laser (NL303G) as excitation source. The experimental setup consists of a spectrometer (Spex 1702), a photomultiplier tube (PMT, Hamamatsu R943), a lock-in amplifier, and a computer for data processing. The cyclic voltammetry curves were measured using an electrochemical workstation (CorrTest, Wuhan, Inc.) in a conventional three-electrode cell at different temperature. The samples/ITO is used as working electrode, Pt is used as counter electrode and the saturated calomel electrode (SCE) was used as reference electrode.

The generation ability of reactive oxygen species (ROS) for thermal catalysts can be estimated by measuring the fluorescent intensity of 2',7'-dichlorofluorescein (DCF). The 2',7'-dichlorodihydrofluorescein (DCFH, non-fluorescent) can rapidly react with ROS in the thermal catalysis system to form 2',7'-dichlorofluorescein (DCF, fluorescent). By measuring the fluorescent intensity of DCF, the generation ability of ROS can be determined for thermal-catalyst. Experimental process is as follow: 5 mg of catalysts were added into 5 ml centrifuge tube, then 1 ml working solution (975 μl of diluents and 25 μl staining fluid containing DCFH) was added. The mixture was shock and heated at different temperature for 30 min (room temperature, 65 °C). Then, the mixture was centrifuged and the supernatant was taken to detect the fluorescent intensity. The exciting wavelength for excitation was 491 nm.

Apoptosis assay. Human Lung Carcinoma cells (A549), smooth muscle cells and synovial cells were cultured in RPMI 1640 medium in 96-well plates, containing 10% fetal calf serum (FCS) in a humidified incubator with an atmosphere of 5% CO2 in air at 37 °C. The cell density was 2 × 104 cells per well. After being seeded for 24 h, the media were replaced by culture media containing a series of TiO2, Pd-TiO2 and TiO2-Pd/Graphene nanoparticles with increasing concentrations in RPMI 1640 medium and then the plates were placed into the humidified incubator. After another 16 h for the interaction between the cancer-cells and the nanocomposite particles, cell viabilities were measured by the standard MTT assay, a colorimetric assay based on the ability of viable cells to reduce 3-[4,5-dimethylthiazol-2-yl]-2,5-diphenyltetrazolium bromide. The survival rate and the error bar are shown in Fig. 2B (the case of incubating with nanocrystals only).

In vivo thermal therapy. BALB/c mice bearing A549 lung carcinoma tumors were intratumorally injected with TiO2, Pd-TiO2 or graphene/Pd-TiO2 (80 ul of 4 mg/ml solution for each mouse), respectively. The images were taken by an high definition camera. The tumor sizes were measured by a caliper every other day and calculated as volume = (tumor length) × (tumor width)2/2. Relative tumor volumes were calculated as V/V0 (V0 is the tumor volume when the treatment was initiated). The animal protocol in this study conformed to the Guide for the Care and Use of Laboratory Animals (the Guide, NRC 2011), and it was also approved by the Institutional Animal Care and Use Committee at Nankai University (Approval ID 201009080081).

Supporting information. Thermal catalytic activity, XRD, XPS Cl 2p, PDOS, Absorption spectra, XPS valence band spectra, Raman spectra, PL spectra and thermal excited PL spectra.

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References
1. Ancona, A. et al. Lipid-coated zinc oxide nanoparticles as innovative ROS-generators for photodynamic therapy in cancer cells. Nanomaterials https://doi.org/10.3390/nano8030143 (2018).
2. Chen, F. et al. In vivo tumor vasculature targeting of CuS@MSN based theranostic nanomedicine. ACS Nano 9, 3926–3934. https://doi.org/10.1021/acsnano.5b04341 (2015).
3. Duan, D. B. et al. Activating TiO2 nanoparticles: gallium-68 serves as a high-yield photon emitter for cerenkov-induced photodynamic therapy. ACS Appl. Mater. Interfaces 10, 5278–5286. https://doi.org/10.1021/acsami.7b17902 (2018).
4. Han, Y. et al. Theranostic micelles based on upconversion nanoparticles for dual-modal imaging and photodynamic therapy in hepatocellular carcinoma. Nanoscale 10, 6511–6523. https://doi.org/10.1039/c7nr09717d (2018).
5. Huang, C.-X. et al. Controlled synthesis of upconverting nanoparticles/CuS yolk-shell nanoparticles for in vitro synergetic photothermal and photodynamic therapy of cancer cells. J. Mater. Chem. B 5, 9487–9496. https://doi.org/10.1039/c7tb02733h (2017).
6. Seo, J. W. et al. Development of water-soluble single-crystalline TiO2 nanoparticles for photocatalytic cancer-cell treatment. Small 3, 850–853. https://doi.org/10.1002/smll.200600488 (2007).
7. Wu, M.-X. & Yang, Y.-W. Metal–organic framework (MOF)-based drug/cargo delivery and cancer therapy. *Adv. Mater.* [https://doi.org/10.1002/adma.201606134](https://doi.org/10.1002/adma.201606134) (2017).
8. Dou, Q. Q., Rengaramchandran, A., Selvan, S. T., Paulmurgan, R. & Zhang, Y. Core-shell upconversion nanoparticle-semiconductor heterostructures for photodynamic therapy. *Sci. Rep.* [https://doi.org/10.1038/srep08252](https://doi.org/10.1038/srep08252) (2015).
9. Lucky, S. S., Soo, K. C. & Zhang, Y. Nanoparticles in photodynamic therapy. *Chem. Rev.* 115, 1990–2042. [https://doi.org/10.1021/cr5004198](https://doi.org/10.1021/cr5004198) (2015).
10. Lismont, M., Dreesen, L. & Wuttke, S. Metal–organic framework nanoparticles in photodynamic therapy: current status and perspectives. *Adv. Funct. Mater.* [https://doi.org/10.1002/adfm.201606314](https://doi.org/10.1002/adfm.201606314) (2017).

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Author contributions

Y.C. conceived the project and designed the experiments. Y.Y., P.J. and Y.Y. performed the samples preparation, structural characterization, in vivo thermal therapy and apoptosis assay. H.L. and L.Z. performed the theory calculation. S.J. and W.Y. helped and analyzed the paper. Y.Y. and Y.C. wrote this paper.

Competing interests

The authors declare no competing interests.

Additional information

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