

## RESEARCH ARTICLE

View Article Online  
View Journal | View IssueCite this: *Mater. Chem. Front.*,  
2019, 3, 127

# Enhanced efficacy of photothermal therapy by combining a semiconducting polymer with an inhibitor of a heat shock protein†

Tingting Sun,<sup>ab</sup> Xingxing Chen,<sup>ab</sup> Xin Wang,<sup>c</sup> Shi Liu,<sup>a</sup> Jun Liu<sup>id</sup><sup>a</sup> and Zhiqiang Xie<sup>id</sup><sup>\*a</sup>

Photothermal therapy (PTT) is a highly effective therapeutic modality in tumor therapy. Nevertheless, the PTT of cancer is also accompanied by thermoresistance of cells, which may alleviate the PTT efficacy or even lead to tumor recurrence. Heat shock protein 90 (HSP90) is closely related to the resistance, so a combination of inhibitors of HSP90 and photothermal agents could improve the PTT efficacy. In this work, gambogic acid (GA), an inhibitor of HSP90, is incorporated into a photothermal nanoagent, so as to reduce the influence of thermoresistance and maximize the PTT efficiency. Poly((2,5-diyl-2,3,5,6-tetrahydro-3,6-dioxo-pyrrolo(3,4-c)pyrrole-1,4-diyl)-alt-(2,2':5',2''-terthiophene-5,5''-diyl)) (PDPP3T) is selected as the robust photothermal material, and polymer-GA nanoparticles (PGNPs) incorporating both GA and PDPP3T are prepared. The PGNPs exhibit excellent photothermal activity with a high photothermal conversion efficiency of 36%, and great heating reproducibility and photoacoustic imaging performance. Besides, PGNPs are stable enough in water, in physiological conditions or under irradiation. The inhibition of HSP90 dramatically increases the apoptosis rate of cells, accordingly promoting the efficacy of PTT. Mice treated with PGNPs under 808 nm laser irradiation possess the most effective tumor inhibition, while no obvious systemic toxicity has been observed. Our study provides a valid approach to improve the PTT efficacy for more potent cancer therapy.

Received 12th September 2018,  
Accepted 19th November 2018

DOI: 10.1039/c8qm00459e

rsc.li/frontiers-materials

## Introduction

Photothermal therapy (PTT), as a non-invasive therapeutic technique, has been extensively studied in recent years because of its high efficiency and specificity in tumor destruction.<sup>1–12</sup> PTT is based on the photothermal conversion materials which convert the absorbed near-infrared (NIR, 700–1100 nm) light to heat in tumor tissues and cells, sequentially ablating the tumors.<sup>13–15</sup> NIR light is less harmful to normal tissues and possesses deeper penetration depth than ultraviolet or visible light.<sup>16</sup> An 808 nm laser is the most widely selected light source because of the low absorption of water in this region, which can significantly mitigate the heating effect of normal tissues.<sup>16–18</sup> In the last decade, a myriad of organic or inorganic nanomaterials

with strong absorbance in the NIR window have been extensively studied as photothermal agents (PTAs).<sup>6,11,19–35</sup> Among them, semiconducting polymer nanoparticles (SPNs) possess high photothermal ability to convert light into heat for PTT and photoacoustic (PA) imaging.<sup>36–49</sup> More recently, the photothermal features of SPNs have also been combined with other therapeutic methods, for instance, controlling the thermosensitive ion channels<sup>50</sup> or enzyme activity.<sup>10</sup>

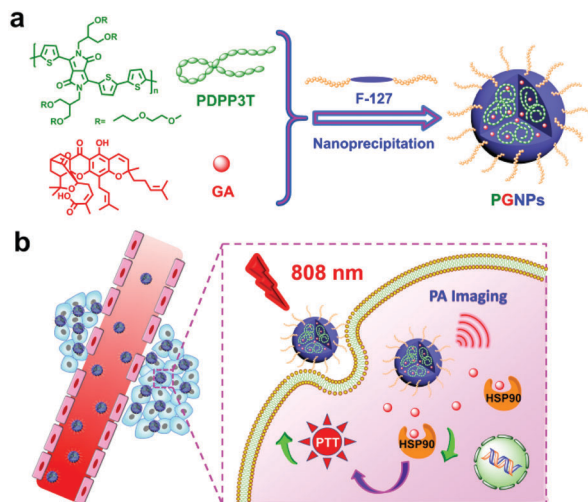
In order to ablate tumors thoroughly, harsh photothermal heating (high temperature over ~50 °C) is required to induce complete necrosis of cells.<sup>51–54</sup> However, due to inevitable heat diffusion, photothermal treatment at a high temperature may threaten the healthy tissues and cells nearby.<sup>55</sup> Normally, when the temperature reaches the “hyperthermia range” (42–47 °C), tumor cells die, probably due to the degeneration of proteins.<sup>51</sup> Nevertheless, the tumor cells are able to activate their cytoprotective pathways such as the induction of heat shock proteins (HSPs), which might rapidly repair thermal damage to proteins and lead to the thermotolerance of tumor cells upon laser irradiation.<sup>1,51,55–59</sup> For example, HSP90, as a molecular chaperone regulating the function and stability of several signaling proteins, is closely related to antiapoptosis and tumor growth.<sup>1,60</sup> It is reasonable that the inhibition of overexpressed HSP90 might

<sup>a</sup> State Key Laboratory of Polymer Physics and Chemistry, Changchun Institute of Applied Chemistry, Chinese Academy of Sciences, 5625 Renmin Street, Changchun, Jilin 130022, P. R. China. E-mail: xiez@ciac.ac.cn; Fax: +86 431 85262775; Tel: +86 431 85262775

<sup>b</sup> University of Chinese Academy of Sciences, Beijing 100049, P. R. China

<sup>c</sup> Department of Thyroid Surgery, The First Hospital of Jilin University, 71 Xinmin Street, Changchun, Jilin 130021, P. R. China

† Electronic supplementary information (ESI) available: Characterization, synthesis and additional figures. See DOI: 10.1039/c8qm00459e



**Scheme 1** Schematic illustration of (a) the preparation and (b) the enhanced PTT efficacy of PGNPs.

surmount tumor thermoresistance thus improving the PTT efficacy of SPNs.<sup>1,2,7,54,61–63</sup>

Herein, gambogic acid (GA), a natural product, was selected as the inhibitor for HSP90.<sup>64</sup> Poly((2,5-diyl-2,3,5,6-tetrahydro-3,6-dioxo-pyrrolo(3,4-c)pyrrole-1,4-diyl)-*alt*-(2,2':5',2''-terthiophene-5,5''-diyl)) (PDPP3T) with branched oligo(ethylene glycol) (OEG) as the side chains and strong absorption in the NIR region was employed as the photothermal material. PDPP3T structures have already been used for photovoltaic devices,<sup>65,66</sup> but their application in the PTT field has not been studied in detail yet. In this work, PDPP3T and GA were co-encapsulated in pluronic F-127 (F-127) to form nanoparticles (PGNPs), and the PTT potential of the PGNPs was demonstrated both *in vitro* and *in vivo* (Scheme 1). Nanoparticles without encapsulation of GA (PNPs) or PDPP3T (GNPs) were also prepared as controls. PGNPs which integrate PTA and HSP90 inhibitors could stressfully block the overexpressed HSP90 upon laser irradiation, thereby enhancing both early and late apoptosis of cells to ultimately improve the PTT efficiency.

## Experimental

### Materials and reagents

PDPP3T was synthesized according to previous methods (yield: 50.0%).<sup>66–68</sup> GA was purchased from Dalian Meilun Biotech Co., Ltd. F-127 was purchased from Shanghai yuanye Bio-Technology Co., Ltd. The cell viability (live-dead cell staining) assay kit and Annexin V-FITC/propidium iodide (PI) apoptosis detection kit were purchased from Jiangsu KeyGEN Biotechnology Co., Ltd. Anti-HSP90 antibody was purchased from Abcam Corporation. FITC-labeled Goat Anti-Mouse IgG (H + L) was purchased from Beyotime Biotechnology Co., Ltd. The DeadEnd™ Fluorometric TUNEL System was purchased from Promega Corporation. The other chemicals were used as obtained commercially.

### Preparation of PGNPs

For the preparation of PGNPs, PDPP3T, GA and F-127 were all dissolved in 2 mL of tetrahydrofuran (THF). Then the solution was quickly dispersed into 10 mL of deionized water under sonication and sonicated for 5 min. After THF was fully evaporated, it was dialysed for 24 h. For the preparation of Nile red (NR) labeled PGNPs (NR@PGNPs), NR, PDPP3T, GA and F-127 were all dissolved in the organic phase. PNPs and GNPs were prepared with the same method in the absence of GA or PDPP3T, respectively.

### Photothermal effects of PGNPs

PGNPs with different concentrations (0–5  $\mu\text{g mL}^{-1}$ ) were irradiated with an 808 nm laser (1.0  $\text{W cm}^{-2}$ ) for 10 min, and the temperature was recorded every 10 s. Besides, for a fixed concentration of PGNPs (5  $\mu\text{g mL}^{-1}$ ), the influence of different power densities (0.3–1.7  $\text{W cm}^{-2}$ ) was also recorded. The photothermal response of the PGNPs in water (5  $\mu\text{g mL}^{-1}$ ) was recorded with irradiation for 10 min and the temperature during the cooling down period was also measured to determine the photothermal conversion efficiency in accordance with previous methods.<sup>69,70</sup> The temperature variations of the PGNPs (5  $\mu\text{g mL}^{-1}$ ) in water over 6 cycles of heating and natural cooling were measured to investigate their photostability.

### Cellular uptake

For investigation of the cellular uptake of PGNPs by human liver hepatocellular carcinoma (HepG2) cells, NR@PGNPs were employed. Firstly, cells were seeded in 6-well culture plates with clean coverslips (1  $\times 10^5$  cells per well) for 24 h. The medium (Dulbecco's modified Eagle's medium, DMEM) was then replaced with NR@PGNPs in culture medium and the cells were incubated at 37 °C for 0.5 or 2 h. After the cells were washed with phosphate-buffered saline (PBS, pH 7.4) three times, they were fixed with 4% formaldehyde. Hoechst 33258 was used to stain the nuclei. The samples were examined *via* confocal laser scanning microscopy (CLSM) by using a Zeiss LSM 700. Flow cytometry (FCM) was also performed to quantify the cellular uptake.

### *In vitro* PTT efficacy

The cytotoxicity of the nanoparticles was measured *via* 3-(4,5-dimethylthiazol-2-yl)-2,5-diphenyltetrazolium bromide (MTT) assays. HepG2 cells were harvested in a logarithmic growth phase and seeded in 96-well plates (8  $\times 10^3$  cells per well) followed by incubation in DMEM for 24 h. After that, cells were treated with DMEM, PNPs, GNPs or PGNPs at various concentrations for 4 h at 37 °C. Then laser irradiation (808 nm, 1.0  $\text{W cm}^{-2}$ , 10 min) was performed. After incubation for another 20 h, 20  $\mu\text{L}$  of MTT solution (5  $\text{mg mL}^{-1}$  in PBS) was added and the plates were incubated at 37 °C for another 4 h, followed by removal of the culture medium containing MTT and addition of 150  $\mu\text{L}$  of dimethyl sulfoxide to each well. Finally, the absorbance of the formazan product at 490 nm was measured *via* a microplate reader after being shaken for 5 min.

## Cell apoptosis detection assays

Cells were seeded in 6-well culture plates and pretreated with the same conditions as in MTT assays. After treatment, the cells were harvested, washed and stained with the Annexin V-FITC/PI detection kit for about 20 min. Finally, the ratio analysis of apoptosis was determined *via* FCM.

## Live-dead cell staining

Cells were pretreated with the same conditions as in the MTT assays. After treatment, calcein-acetoxymethyl (calcein-AM)/PI were used to stain the cells for 30 min at room temperature. Finally, cell staining results were obtained by using a fluorescence microscope.

## Immunofluorescence staining

Cells were seeded in 6-well culture plates with clean coverslips and pretreated with the same conditions as in the MTT assays. After laser irradiation for 1 h, the cells were firstly stained with a mouse primary anti-HSP90 antibody and then Alexa Fluor 488-labeled goat anti-mouse secondary antibody. Finally, the cells were stained with Hoechst 33258 for CLSM.

## PA imaging

*In vitro* and *in vivo* PA imaging were all performed with the reported methods.<sup>71</sup> For *in vivo* PA imaging, tumor-bearing nude mice were injected with PGNPs (1.5 mg kg<sup>-1</sup>) intravenously, and PA images were acquired at 0, 1, 5, 7, 12 and 24 h. The images were analyzed using ViewMOST™ software. After injection for 24 h, the mice were sacrificed, and the major organs (heart, liver, spleen, lungs and kidneys) of the mice were harvested for PA imaging.

## *In vivo* antitumor activity and biosafety

Animal care and handling procedures were according to the guidelines of the Regional Ethics Committee for Animal Experiments.

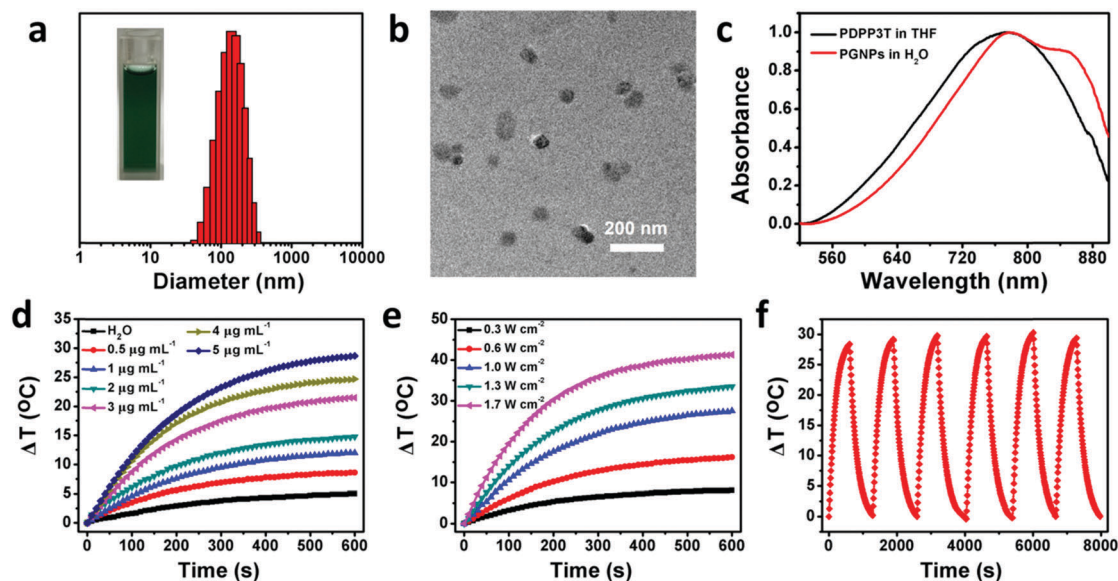
Female Kunming (KM) mice were obtained and maintained under required conditions. To evaluate antitumor effects, the animal model we used was subcutaneous hepatocarcinoma H22 tumor xenografts. The tumor-bearing mice were divided into eight groups (four in each group) randomly. They were intravenously injected with PBS, GNPs (0.15 mg kg<sup>-1</sup> of GA), PNP (1.5 mg kg<sup>-1</sup> of PDPP3T) or PGNPs (1.5 mg kg<sup>-1</sup> of PDPP3T and 0.15 mg kg<sup>-1</sup> of GA) respectively with or without laser irradiation. After different treatments, the tumor dimensions (length and width) were measured every 2 d. Tumor volumes were calculated *via* length × width<sup>2</sup>/2 (mm<sup>3</sup>). 10 d later, they were all sacrificed and the tumors and main organs were collected. The main organs (heart, liver, spleen, lungs and kidneys) and tumors were fixed in 4% paraformaldehyde solution and then embedded in paraffin, sliced and stained using haematoxylin and eosin (H&E) and TUNEL.

For biosafety analysis, 4 KM mice in each group were injected with PGNPs (1.5 mg kg<sup>-1</sup>) intravenously and then irradiated with an 808 nm laser. After 10 d, the liver/kidney function makers were obtained from an automatic biochemical analyzer. Besides, the complete blood of the mice was obtained for hematology analysis.

## Results and discussion

### Preparation and characterization of PGNPs

The synthetic routes of PDPP3T<sup>66</sup> and the chemical structure of GA are shown in Scheme S1 (ESI<sup>†</sup>). The chemical structure of PDPP3T was confirmed by proton nuclear magnetic resonance (<sup>1</sup>H NMR) and elemental analysis. The number-average molecular weight (M<sub>n</sub>) of PDPP3T is about 30 000 g mol<sup>-1</sup>. GA is not only an inhibitor of HSP90 but also a chemotherapeutic agent,<sup>72</sup> which can inhibit the proliferation of many kinds of



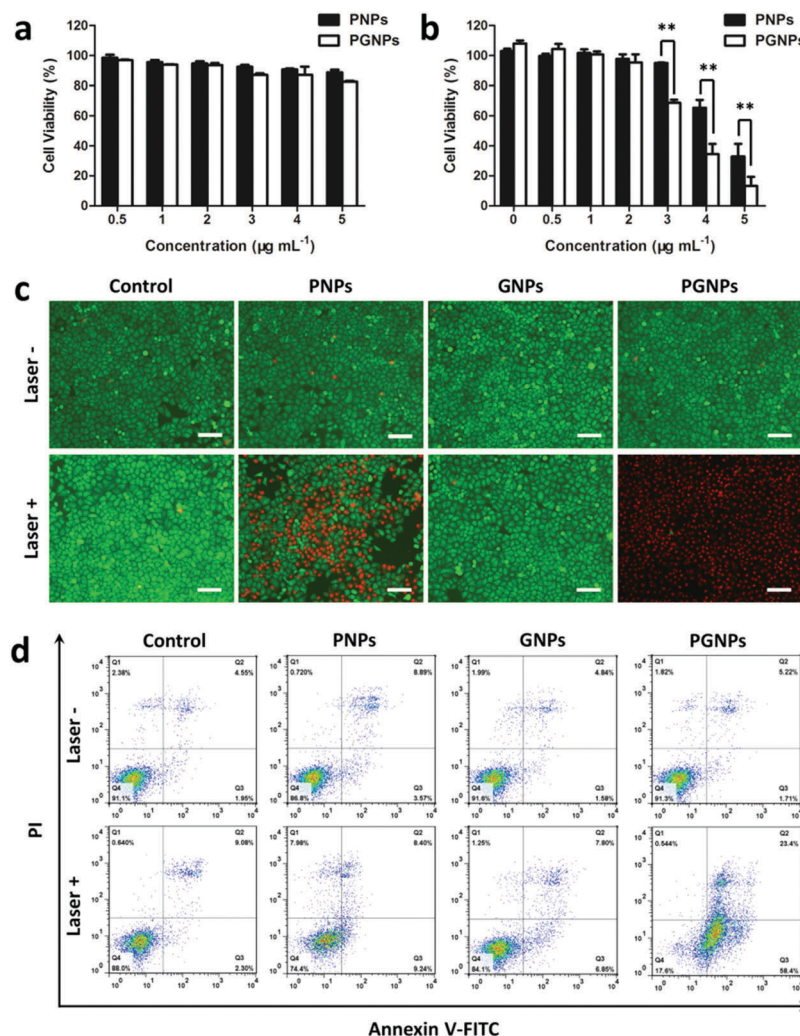
**Fig. 1** Basic characterizations and photothermal performance of PGNPs. (a) DLS result (Inset: Picture of PGNPs in water) and (b) TEM image of PGNPs. (c) Absorption spectra of PDPP3T in THF and PGNPs in water. Photothermal conversion behavior of PGNPs by varying (d) concentrations (0–5 μg mL<sup>-1</sup>) and (e) laser power densities (0.3–1.7 W cm<sup>-2</sup>). (f) Heating reproducibility of PGNPs (5 μg mL<sup>-1</sup>) in water over 6 cycles of heating-cooling.



malignant cells, including hepatoma,<sup>73</sup> lung cancer,<sup>74</sup> leukemia,<sup>75</sup> breast carcinoma<sup>76</sup> and gastric carcinoma.<sup>77</sup> The cytotoxicity of GA against HepG2 cells was first evaluated to determine the dosage of administration under the premise of ignoring its chemotherapeutic effect. As shown in Fig. S1 (ESI<sup>†</sup>), a maximum concentration of  $0.5 \mu\text{g mL}^{-1}$  would be applicable. Then, PGNPs were prepared *via* a nanoprecipitation method. After optimizing the conditions, the loading contents were 15 wt% of PDPP3T and 1.5 wt% of GA, accompanied by loading efficiencies of 91 and 73%, respectively. The contents of PDPP3T and GA in the PGNPs were determined from the standard curves (Fig. S2 and S3, ESI<sup>†</sup>). The obtained solution of PGNPs is in dark green, and the hydrodynamic diameter of the PGNPs is 134.9 nm with a polydispersity index (PDI) of 0.15 as determined by dynamic light scattering (DLS) (Fig. 1a). The nanoparticles in the transmission electron microscopy (TEM) image (Fig. 1b) exhibit a uniform spherical morphology with sizes

smaller than that measured by DLS, probably resulting from the volume shrinkage during the drying process of the TEM sample. PGNPs possess good stability both in water and PBS (pH 7.4) containing 10% fetal bovine serum (FBS) (Fig. S4, ESI<sup>†</sup>), which is of great importance for their storage and practical application. Besides, the hydrodynamic diameters of PNP and GNPs are 148.8 and 133.8 nm with PDI of 0.172 and 0.166, respectively (Fig. S5, ESI<sup>†</sup>).

As shown in Fig. 1c, PGNPs in water have strong absorption in the NIR region, with a bathochromic shift relative to PDPP3T in THF, which is ascribed to the aggregation of PDPP3T molecules. The absorption coefficient of PGNPs in water at 808 nm is as high as  $60.0 \text{ L g}^{-1} \text{ cm}^{-1}$ , which is favorable for their photothermal application. To evaluate the photothermal performance of PGNPs in water, the temperature of the solution was recorded and compared with the variation of the concentrations and laser power. When the concentrations were



**Fig. 2** Cytotoxicity against HepG2 cells. Cell viabilities of HepG2 cells treated with PNPs or PGNPs (a) without or (b) with 808 nm laser irradiation ( $1.0 \text{ W cm}^{-2}$ , 10 min). Statistical significance:  $** P \leq 0.01$ . (c) Fluorescence images of live (green) and dead (red) HepG2 cells co-stained with Calcein-AM and PI after being incubated with DMEM, PNPs ( $5 \mu\text{g mL}^{-1}$  of PDPP3T), GNPs ( $0.5 \mu\text{g mL}^{-1}$  of GA) or PGNPs ( $5 \mu\text{g mL}^{-1}$  of PDPP3T and  $0.5 \mu\text{g mL}^{-1}$  of GA) respectively, and then treated with or without laser irradiation ( $1.0 \text{ W cm}^{-2}$ ) for 10 min. Scale bars, 100  $\mu\text{m}$ . (d) FCM analysis of early and late apoptosis of HepG2 cells after the same treatments as in (c).

increased from 0 (pure water) to  $5 \mu\text{g mL}^{-1}$  and the power density of the 808 nm laser was fixed at  $1.0 \text{ W cm}^{-2}$ , the increase in temperature revealed an obvious concentration-dependent tendency, as shown in Fig. 1d. The temperature elevation of the PGNPs could reach  $28 \text{ }^\circ\text{C}$  at a low concentration of  $5 \mu\text{g mL}^{-1}$  after laser irradiation for 10 min. A laser power-dependent temperature increase was also detected with a constant concentration of  $5 \mu\text{g mL}^{-1}$  (Fig. 1e). The photothermal conversion efficiency of the PGNPs calculated referring to previous methods<sup>69,70</sup> is 36% (Fig. S6, ESI<sup>†</sup>), further demonstrating their excellent photothermal performance. To validate the photothermal stability of PGNPs, 6 cycles of heating-cooling were conducted, and the elevation in temperature (Fig. 1f) and the sizes of the PGNPs (Fig. S7, ESI<sup>†</sup>) remained consistent. The high absorption coefficient, the excellent photothermal conversion behavior and the great photothermal stability of the PGNPs are beneficial to the potential application of the PGNPs in PTT.

### Cellular uptake

To demonstrate the efficient cellular uptake of PGNPs by HepG2 cells, NR was selected as the fluorescent probe<sup>78</sup> and loaded in the PGNPs (NR@PGNPs). As indicated in Fig. S8 (ESI<sup>†</sup>), the photoluminescence (PL) spectra of the PGNPs and NR@PGNPs as well as the picture of the NR@PGNPs in water substantiated the successful loading of NR. Then, CLSM was employed to observe the internalization of HepG2 cells incubated with NR@PGNPs for 0.5 and 2 h. The cell nuclei were stained with Hoechst 33258. As shown in Fig. S9a (ESI<sup>†</sup>), the fluorescence intensity of NR@PGNPs in the cells increases from 0.5 to 2 h, indicating efficient cellular uptake of PGNPs by HepG2 cells. FCM was also carried out to quantify the cellular uptake, and a time-dependent internalization of NR@PGNPs was detected (Fig. S9b, ESI<sup>†</sup>). The efficient internalization of PGNPs by cells is the firm foundation to exert their therapeutic effect.

### In vitro PTT efficacy

To evaluate the influence of GA on cytotoxicity, the viabilities of cells treated with PNPs or PGNPs with or without 808 nm laser

irradiation were determined by MTT assays. For cells treated with PNPs or PGNPs ( $0\text{--}5 \mu\text{g mL}^{-1}$ ) for 24 h in the absence of laser irradiation, there was no significant cytotoxicity detected (Fig. 2a). After laser irradiation ( $1.0 \text{ W cm}^{-2}$ , 10 min), a dramatic decline in viability was observed with the increase in concentration ( $3\text{--}5 \mu\text{g mL}^{-1}$ ) (Fig. 2b). More importantly, the existence of GA in the PGNPs significantly decreased the viabilities of HepG2 cells relative to PNPs, with half maximal inhibitory concentration ( $\text{IC}_{50}$ ) values of 4.3 and  $3.4 \mu\text{g mL}^{-1}$  for PNPs and PGNPs, respectively, demonstrating that the inhibition of HSP90 could enhance the PTT effect observably. However,

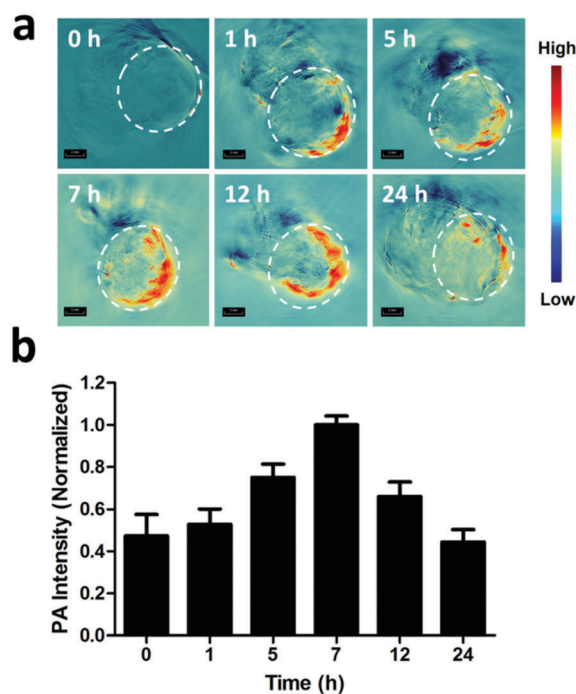


Fig. 4 (a) PA images of the tumor (in the white dashed circles) taken at specified time points after systemic administration of PGNPs ( $1.5 \text{ mg kg}^{-1}$  of PDPP3T). Scale bar, 3 mm. (b) Quantified signal intensities of the tumor.

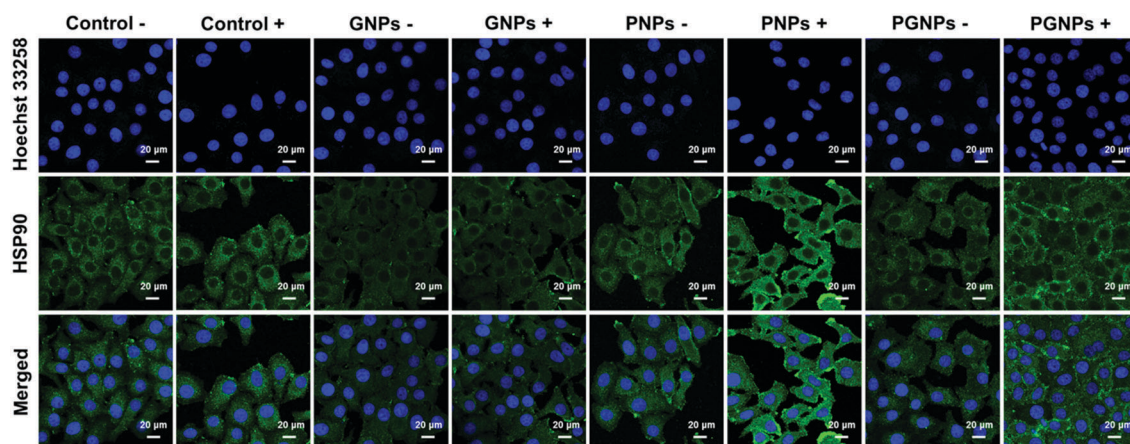


Fig. 3 The expression of HSP90 in HepG2 cells 1 h post various treatments determined by immunofluorescence staining. The HepG2 cells were treated with DMEM, GNPs, PNPs or PGNPs under 808 nm laser irradiation ( $1.0 \text{ W cm}^{-2}$ , 10 min) or not.



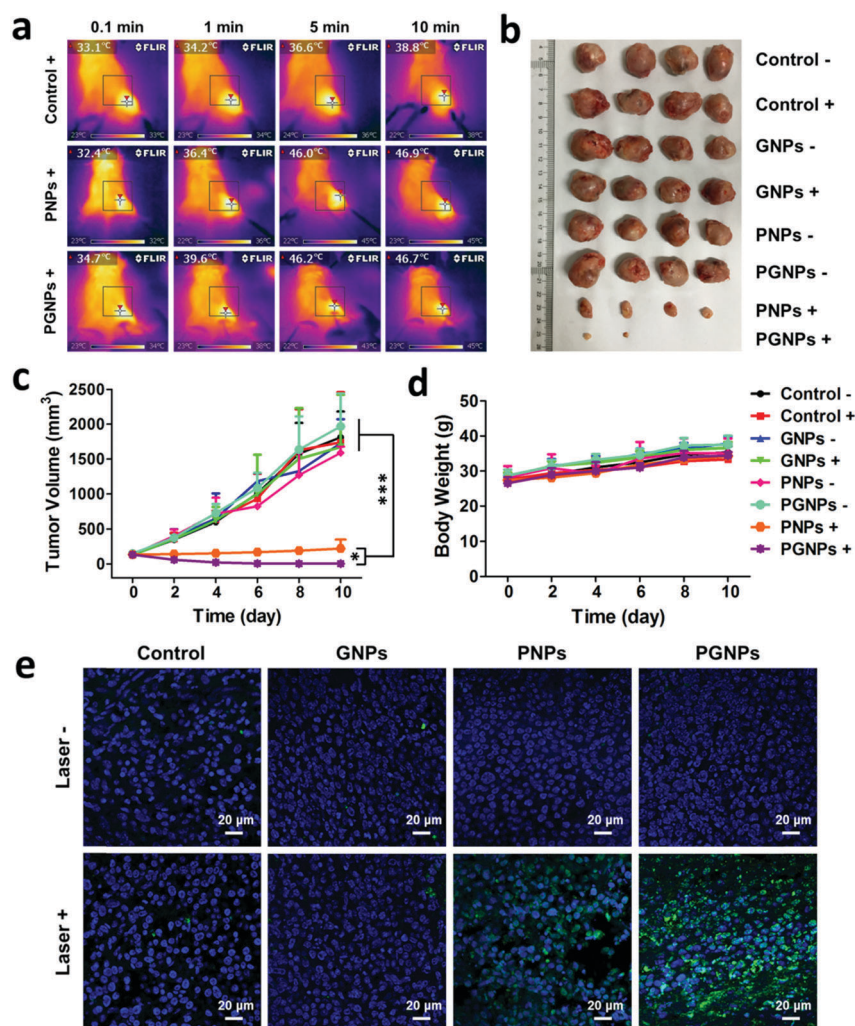
laser irradiation makes no difference to the viability of cells treated with GNPs (Fig. S10, ESI†).

In order to visualize the *in vitro* PTT effect, cells were stained with calcein-AM and PI to differentiate live cells (green) and dead ones (red). Cells were treated with PBS (Control group), PNPs ( $5 \mu\text{g mL}^{-1}$  of PDPP3T), GNPs ( $0.5 \mu\text{g mL}^{-1}$  of GA) or PGNPs ( $5 \mu\text{g mL}^{-1}$  of PDPP3T and  $0.5 \mu\text{g mL}^{-1}$  of GA), followed by laser irradiation (Laser +) or not (Laser -). Fig. 2c showed that most cells in the Control (Laser -/+), PNPs (Laser -), GNPs (Laser -/+) and PGNPs (Laser -) groups were alive with green fluorescence. Nearly all the cells in the PGNPs (Laser +) group were dead and exhibited red fluorescence, and the cell death was much more than that for the PNPs (Laser +) group (Fig. 2c), which was in line with the MTT results (Fig. 2b). Then, cell apoptosis induced by different treatments was examined quantitatively with an Annexin-V FITC/PI staining assay *via* FCM analysis. It could be seen from Fig. 2d that there was no obvious difference between cells treated with PBS without or with

laser irradiation, indicating that the laser irradiation only has a negligible effect on cell apoptosis. The cells incubated with PNPs, GNPs or PGNPs without laser irradiation also exhibited insignificant apoptosis. For the PTT effect of PDPP3T alone, cells treated with PNPs and laser irradiation showed early and late apoptosis of 9.24% and 8.40%, respectively. In contrast, PGNPs caused the early and late apoptosis of 58.4% and 23.4%, respectively upon laser irradiation, probably due to the synergistic effect of PDPP3T and GA. The results in Fig. 2 intuitively confirmed the enhanced PTT efficacy of PGNPs with 808 nm laser irradiation by inhibiting HSP90.

### The expression of HSP90

To determine the expression level of HSP90 in HepG2 cells, cells from various treatments were applied for immunofluorescence staining analysis. For conciseness, “Laser +/-” is abbreviated as “+/-”. As shown in Fig. 3 and Fig. S11 (ESI†), the expression of HSP90 was significantly upregulated in the cells



**Fig. 5** *In vivo* therapy of tumors. (a) IR images of mice irradiated by 808 nm laser irradiation ( $1.0 \text{ W cm}^{-2}$ ) after intravenous injection of PBS, PNPs or PGNPs ( $1.5 \text{ mg kg}^{-1}$  of PDPP3T). (b) A photograph of the excised tumors from each group. (c) Growth profiles of tumors and (d) changes of body weight after different treatments. (e) TUNEL staining of tumor slices from mice subjected to various treatments 2 d later. Statistical significance: \* $P \leq 0.05$ ; \*\*\* $P \leq 0.001$ .

incubated with PNP after PTT treatment (PNPs +), demonstrating that PTT-induced heating stress could indeed upregulate the expression of HSP90 by cancer cells. Much lower HSP90 level was observed for cells treated with PGNPs under laser irradiation (PGNPs +) relative to that for cells in the PNP + group, although it was higher than that for other groups, especially the GNPs -/+ groups. These results should be ascribed to the GA-mediated inhibition of the temporarily stressfully overexpressed HSP90 during PTT of cancer.

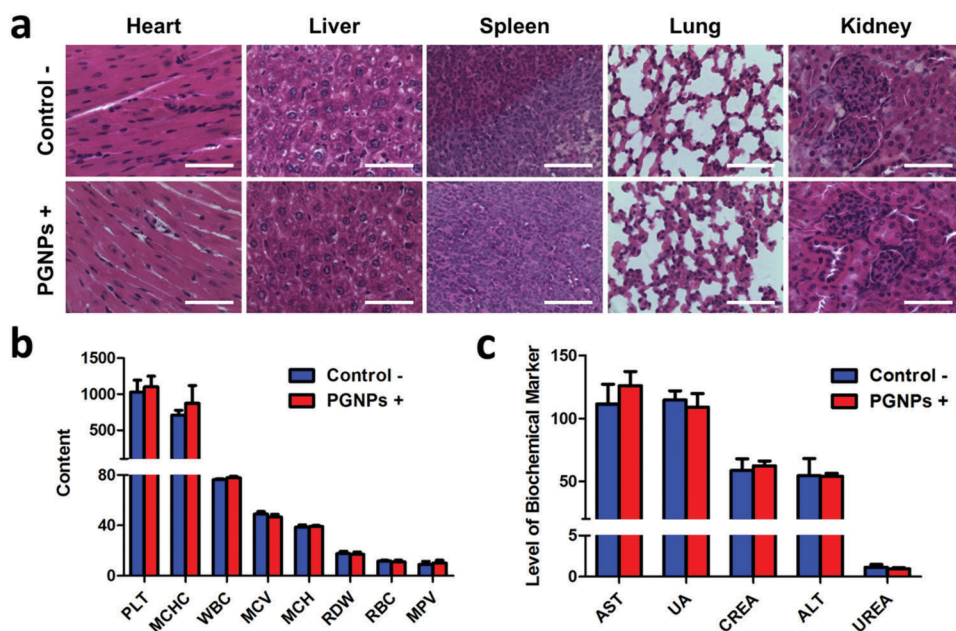
### PA imaging performance of PGNPs

As reported, the process of photothermal transformation is accompanied by the generation of PA waves.<sup>42</sup> Therefore, the PA imaging performance of the PGNPs was further employed to investigate the biodistribution of the nanoparticles. Firstly, the PA signals of PGNPs of varying concentrations were determined, exhibiting a linearly strengthened tendency with increasing concentration (Fig. S12, ESI<sup>†</sup>). Next, *in vivo* PA imaging was implemented on HepG2 tumor-bearing nude mice after intravenous injection with PGNPs (1.5 mg kg<sup>-1</sup> of PDPP3T) for different time periods (0, 1, 5, 7, 12 and 24 h). As shown in Fig. 4, the PA signals of PGNPs in the tumor region displayed a time-dependent variation with the maximum accumulation at 7 h post injection, which was very important for determining the optimal time for laser irradiation to guarantee the best therapeutic effect. After injection for 24 h, the mice were sacrificed, and the major organs (heart, liver, spleen, lungs and kidneys) of the mice were harvested. The *ex vivo* PA data in Fig. S13 (ESI<sup>†</sup>) illustrate that PGNPs are mainly accumulated in the liver and spleen. These results well certified the PA imaging

ability of the PGNPs, suggesting the probability of employing the nanoparticles for PA imaging-guided cancer therapy.

### *In vivo* antitumor activity and biosafety

Encouraged by the enhanced cytotoxicity of PGNPs in PTT *in vitro* and their tumor accumulation, we then studied the *in vivo* PTT on H22 tumor bearing KM mice. When the tumor volumes of mice reached 100–150 mm<sup>3</sup>, they were randomly divided into eight groups the same as in the *in vitro* experiments: Control -/+; GNPs -/+; PNP -/+ and PGNPs -/+. The mice were intravenously injected with PBS, GNPs (0.15 mg kg<sup>-1</sup> of GA), PNP (1.5 mg kg<sup>-1</sup> of PDPP3T) or PGNPs (1.5 mg kg<sup>-1</sup> of PDPP3T and 0.15 mg kg<sup>-1</sup> of GA) respectively, and then they were treated with or without 808 nm laser irradiation (1.0 W cm<sup>-2</sup>, 10 min). Upon laser irradiation, an infrared (IR) thermal camera was employed to monitor the temperature changes of mice in the Control +, PNP + and PGNPs + groups (Fig. 5a). The tumors in the Control + group showed a slight increase in temperature upon laser irradiation. The temperature of the tumors in the PNP + and PGNPs + groups manifested a rapid rise in the first 5 min, then kept nearly unchanged between 46 and 47 °C, which validated the photothermal activity of the PNP and PGNPs *in vivo*. The tumor volumes and body weight of the mice were measured every 2 d during the subsequent 10 d. The mice of the eight groups were sacrificed, and the tumors were collected at day 10. The tumor sizes in the PNP + and PGNPs + groups were much smaller than those in other groups, and the tumors in the PGNPs + group were the smallest or even disappeared completely (Fig. 5b and Fig. S14, ESI<sup>†</sup>). As shown in Fig. 5c, the tumor volumes of the mice in the



**Fig. 6** (a) H&E staining of major organs (heart, liver, spleen, lungs and kidneys) of mice in the Control - and PGNPs + groups. Scale bars: 200 μm. (b) Hematology data and (c) serum biochemical analysis of kidney and liver function parameters of the mice ( $n = 4$ ) in the Control - and PGNPs + groups after treatment for 10 d. PLT, platelet; MCHC, mean corpuscular hemoglobin concentration; WBC, white blood cell count; MCV, mean corpuscular volume; MCH, mean corpuscular hemoglobin; RDW, red cell distribution width; RBC, red blood cell; MPV, mean platelet volume; AST, aspartate aminotransferase; UA, uric acid; CREA, creatinine; ALT, alanine transaminase; and UREA, urea nitrogen.

Control  $-/+$ , GNPs  $-/+$ , PNP $s$   $-$  and PGNPs  $-$  groups increased rapidly during the entire period. Under irradiation, PNP $s$  could inhibit the growth of tumors upon irradiation to a large extent, but a mild increase in tumor volume was also observed. While PGNPs could decrease the volumes of all the tumors under irradiation, and some of them were even completely ablated. Next, the H&E staining of H22 tumor slices collected 2 d post treatment further verified the most severe cell damage in the PGNPs + group with that of the PNP $s$  + group in second place, while no remarkable damage was detected in the other groups (Fig. S15, ESI $^{\dagger}$ ). To determine the cell apoptosis levels of tumor cells, the terminal deoxynucleotidyl transferase-mediated dUTP-biotin nick end labeling (TUNEL) staining assay was carried out on tumor slices (Fig. 5e). As expected, the slice from the PGNPs + group showed the brightest green fluorescence, standing for the highest level of cell apoptosis, followed by that from the PNP $s$  + group. However, the tumor slices from the other groups possessed a negligible level of TUNEL positive cells. It can be concluded that PGNPs are more effective than PNP $s$  in inducing apoptosis of cancer cells under 808 nm laser irradiation in the hyperthermia range. The body weight of all the mice was in a slow upward trend (Fig. 5d), suggesting that these treatments had no significant acute toxicity to the animals. Moreover, as revealed by the H&E staining of normal tissues (heart, liver, spleen, lungs and kidneys) (Fig. 6a), the complete blood panels (Fig. 6b) and serum biochemistry (Fig. 6c), there is no obvious differences between the Control  $-$  and PGNPs + groups, indicating the great systemic safety of treatment with PGNPs and laser irradiation.

## Conclusions

In summary, we have constructed PGNPs integrating the photo-thermal properties of PDPP3T with inhibition towards HSP90 for highly potent cancer therapy. PGNPs possess a high photo-thermal conversion efficiency of 36% under 808 nm laser irradiation, and have excellent physiological stability as well as photostability. Furthermore, the PA imaging ability of the PGNPs makes imaging-guided cancer therapy possible. More importantly, the PGNPs could induce more severe cell apoptosis *in vitro* and effective inhibition of tumor growth *in vivo* than PNP $s$  without the inhibitor of HSP90. This work demonstrates a new design to improve the PTT efficacy of PTAs based on semiconducting polymer nanoparticles, thus realizing more effective cancer therapy.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

This project was supported by the National Natural Science Foundation of China (Project No. 51522307).

## Notes and references

- H. Luo, Q. Wang, Y. Deng, T. Yang, H. Ke, H. Yang, H. He, Z. Guo, D. Yu, H. Wu and H. Chen, *Adv. Funct. Mater.*, 2017, **27**, 1702834.
- B.-K. Wang, X.-F. Yu, J.-H. Wang, Z.-B. Li, P.-H. Li, H. Wang, L. Song, P. K. Chu and C. Li, *Biomaterials*, 2016, **78**, 27–39.
- L. Cheng, C. Wang, L. Feng, K. Yang and Z. Liu, *Chem. Rev.*, 2014, **114**, 10869–10939.
- X. Cai, J. Liu, W. H. Liew, Y. Duan, J. Geng, N. Thakor, K. Yao, L.-D. Liao and B. Liu, *Mater. Chem. Front.*, 2017, **1**, 1556–1562.
- M. P. Melancon, M. Zhou and C. Li, *Acc. Chem. Res.*, 2011, **44**, 947–956.
- X. Zhen, C. Xie and K. Pu, *Angew. Chem., Int. Ed.*, 2018, **57**, 3938–3942.
- Y. Yang, W. Zhu, Z. Dong, Y. Chao, L. Xu, M. Chen and Z. Liu, *Adv. Mater.*, 2017, **29**, 1703588.
- C. Wang, L. Xu, C. Liang, J. Xiang, R. Peng and Z. Liu, *Adv. Mater.*, 2014, **26**, 8154–8162.
- P. Huang, J. Lin, W. Li, P. Rong, Z. Wang, S. Wang, X. Wang, X. Sun, M. Aronova, G. Niu, D. Leapman Richard, Z. Nie and X. Chen, *Angew. Chem., Int. Ed.*, 2013, **52**, 13958–13964.
- J. Li, C. Xie, J. Huang, Y. Jiang, Q. Miao and K. Pu, *Angew. Chem., Int. Ed.*, 2018, **57**, 3995–3998.
- Q. Dong, X. Wang, X. Hu, L. Xiao, L. Zhang, L. Song, M. Xu, Y. Zou, L. Chen, Z. Chen and W. Tan, *Angew. Chem., Int. Ed.*, 2017, **57**, 177–181.
- Z. Yang, W. He, H. Zheng, J. Wei, P. Liu, W. Zhu, L. Lin, L. Zhang, C. Yi, Z. Xu and J. Ren, *Biomaterials*, 2018, **161**, 1–10.
- Q. Ban, T. Bai, X. Duan and J. Kong, *Biomater. Sci.*, 2017, **5**, 190–210.
- Y. Liu, X. Ji, J. Liu, W. L. Tong Winnie, D. Askhatova and J. Shi, *Adv. Funct. Mater.*, 2017, **27**, 1703261.
- W. Huang, Y. Huang, Y. You, T. Nie and T. Chen, *Adv. Funct. Mater.*, 2017, **27**, 1701388.
- X. Deng, K. Li, X. Cai, B. Liu, Y. Wei, K. Deng, Z. Xie, Z. Wu, P. A. Ma, Z. Hou, Z. Cheng and J. Lin, *Adv. Mater.*, 2017, **29**, 1701266.
- V. J. Pansare, S. Hejazi, W. J. Faenza and R. K. Prud'homme, *Chem. Mater.*, 2012, **24**, 812–827.
- W. Cheng, J. Nie, N. Gao, G. Liu, W. Tao, X. Xiao, L. Jiang, Z. Liu, X. Zeng and L. Mei, *Adv. Funct. Mater.*, 2017, **27**, 1704135.
- K. Yang, L. Feng, X. Shi and Z. Liu, *Chem. Soc. Rev.*, 2013, **42**, 530–547.
- S. Thota, Y. Wang and J. Zhao, *Mater. Chem. Front.*, 2018, **2**, 1074–1089.
- W. M. Girma, S.-H. Tzing, P.-J. Tseng, C.-C. Huang, Y.-C. Ling and J.-Y. Chang, *ACS Appl. Mater. Interfaces*, 2018, **10**, 4590–4602.
- S. Zhang, Q. Huang, L. Zhang, H. Zhang, Y. Han, Q. Sun, Z. Cheng, H. Qin, S. Dou and Z. Li, *Nanoscale*, 2018, **10**, 3130–3143.



- 23 N. Yan, X. Wang, L. Lin, T. Song, P. Sun, H. Tian, H. Liang and X. Chen, *Adv. Funct. Mater.*, 2018, **28**, 1800490.
- 24 Y. Zhan, Y. Liu, H. Zu, Y. Guo, S. Wu, H. Yang, Z. Liu, B. Lei, J. Zhuang, X. Zhang, D. Huang and C. Hu, *Nanoscale*, 2018, **10**, 5997–6004.
- 25 H. S. Jung, P. Verwilt, A. Sharma, J. Shin, J. L. Sessler and J. S. Kim, *Chem. Soc. Rev.*, 2018, **47**, 2280–2297.
- 26 X. Cheng, R. Sun, L. Yin, Z. Chai, H. Shi and M. Gao, *Adv. Mater.*, 2016, **29**, 1604894.
- 27 Y. Shi, M. Liu, F. Deng, G. Zeng, Q. Wan, X. Zhang and Y. Wei, *J. Mater. Chem. B*, 2017, **5**, 194–206.
- 28 X. Liu, H. Su, W. Shi, Y. Liu, Y. Sun and D. Ge, *Biomaterials*, 2018, **167**, 177–190.
- 29 K. Chang, Y. Liu, D. Hu, Q. Qi, D. Gao, Y. Wang, D. Li, X. Zhang, H. Zheng, Z. Sheng and Z. Yuan, *ACS Appl. Mater. Interfaces*, 2018, **10**, 7012–7021.
- 30 H. Kim, S. Beack, S. Han, M. Shin, T. Lee, Y. Park, S. Kim Ki, K. Yetisen Ali, H. Yun Seok, W. Kwon and K. Hahn Sei, *Adv. Mater.*, 2018, **30**, 1701460.
- 31 J. Meng, X. Chen, Y. Tian, Z. Li and Q. Zheng, *Chem. – Eur. J.*, 2017, **23**, 17521–17530.
- 32 R. Vankayala and C. Hwang Kuo, *Adv. Mater.*, 2018, **30**, 1706320.
- 33 C. Leng, X. Zhang, F. Xu, Y. Yuan, H. Pei, Z. Sun, L. Li and Z. Bao, *Small*, 2018, **14**, 1703077.
- 34 Z. Liu, Q. Xu, Y. Li and W. Chen, *Mater. Chem. Front.*, 2017, **1**, 538–541.
- 35 Q. Jin, J. Liu, W. Zhu, Z. Dong, Z. Liu and L. Cheng, *ACS Appl. Mater. Interfaces*, 2018, **10**, 332–340.
- 36 Y. Jiang, D. Cui, Y. Fang, X. Zhen, P. K. Upputuri, M. Pramanik, D. Ding and K. Pu, *Biomaterials*, 2017, **145**, 168–177.
- 37 D.-D. Li, J.-X. Wang, Y. Ma, H.-S. Qian, D. Wang, L. Wang, G. Zhang, L. Qiu, Y.-C. Wang and X.-Z. Yang, *ACS Appl. Mater. Interfaces*, 2016, **8**, 19312–19320.
- 38 S. Li, X. Wang, R. Hu, H. Chen, M. Li, J. Wang, Y. Wang, L. Liu, F. Lv, X.-J. Liang and S. Wang, *Chem. Mater.*, 2016, **28**, 8669–8675.
- 39 J. Zhang, C. Yang, R. Zhang, R. Chen, Z. Zhang, W. Zhang, S.-H. Peng, X. Chen, G. Liu, C.-S. Hsu and C.-S. Lee, *Adv. Funct. Mater.*, 2017, **27**, 1605094.
- 40 T. Sun, J.-H. Dou, S. Liu, X. Wang, X. Zheng, Y. Wang, J. Pei and Z. Xie, *ACS Appl. Mater. Interfaces*, 2018, **10**, 7919–7926.
- 41 T. Dong, K. Wen, J. Chen, J. Xie, W. Fan, H. Ma, L. Yang, X. Wu, F. Xu, A. Peng and H. Huang, *Adv. Funct. Mater.*, 2018, **28**, 1800135.
- 42 Z. Cao, L. Feng, G. Zhang, J. Wang, S. Shen, D. Li and X. Yang, *Biomaterials*, 2018, **155**, 103–111.
- 43 B. Guo, G. Feng, N. Manghnani Purnima, X. Cai, J. Liu, W. Wu, S. Xu, X. Cheng, C. Teh and B. Liu, *Small*, 2016, **12**, 6243–6254.
- 44 C. Xie, X. Zhen, Q. Lei, R. Ni and K. Pu, *Adv. Funct. Mater.*, 2017, **27**, 1605397.
- 45 Q. Miao, C. Xie, X. Zhen, Y. Lyu, H. Duan, X. Liu, J. V. Jokerst and K. Pu, *Nat. Biotechnol.*, 2017, **35**, 1102.
- 46 J. Li, X. Zhen, Y. Lyu, Y. Jiang, J. Huang and K. Pu, *ACS Nano*, 2018, **12**, 8520–8530.
- 47 Y. Lyu, D. Cui, H. Sun, Y. Miao, H. Duan and K. Pu, *Angew. Chem., Int. Ed.*, 2017, **56**, 9155–9159.
- 48 Q. Miao and K. Pu, *Adv. Mater.*, 2018, **30**, 1801778.
- 49 Y. Jiang and K. Pu, *Acc. Chem. Res.*, 2018, **51**, 1840–1849.
- 50 X. Zhen, C. Xie, Y. Jiang, X. Ai, B. Xing and K. Pu, *Nano Lett.*, 2018, **18**, 1498–1505.
- 51 S. K. Calderwood, *Tumor Ablation*, Springer, Netherlands, Dordrecht, 2013, pp. 29–37.
- 52 M. Koishi, S. I. Yokota, T. Mae, Y. Nishimura, S. Kanamori, N. Horii, K. Shibuya, K. Sasai and M. Hiraoka, *Clin. Cancer Res.*, 2001, **7**, 215–219.
- 53 S.-i. Yokota, M. Kitahara and K. Nagata, *Cancer Res.*, 2000, **60**, 2942–2948.
- 54 D. Yoo, H. Jeong, S.-H. Noh, J.-H. Lee and J. Cheon, *Angew. Chem., Int. Ed.*, 2013, **52**, 13047–13051.
- 55 J. Zhou, M. Li, Y. Hou, Z. Luo, Q. Chen, H. Cao, R. Huo, C. Xue, L. Sutrisno, L. Hao, Y. Cao, H. Ran, L. Lu, K. Li and K. Cai, *ACS Nano*, 2018, **12**, 2858–2872.
- 56 J. W. Fisher, S. Sarkar, C. F. Buchanan, C. S. Szot, J. Whitney, H. C. Hatcher, S. V. Torti, C. G. Rylander and M. N. Rylander, *Cancer Res.*, 2010, **70**, 9855–9864.
- 57 D. R. Ciocca, N. Cayado-Gutierrez, M. Maccioni and F. D. Cuello-Carrion, *Curr. Mol. Med.*, 2012, **12**, 1183–1197.
- 58 M. Rybiński, Z. Szymańska, S. Lasota and A. Gambin, *J. R. Soc., Interface*, 2013, **10**, 20130527.
- 59 S. W. Lee, J. W. Lee, J. H. Chung and J. K. Jo, *World J. Mens Health*, 2013, **31**, 247–253.
- 60 W. Xu and L. Neckers, *Clin. Cancer Res.*, 2007, **13**, 1625–1629.
- 61 T.-y. Lin, W. Guo, Q. Long, A. Ma, Q. Liu, H. Zhang, Y. Huang, S. Chandrasekaran, C. Pan, K. S. Lam and Y. Li, *Theranostics*, 2016, **6**, 1324–1335.
- 62 W. Guo, P. Reigan, D. Siegel, J. Zirrolli, D. Gustafson and D. Ross, *Cancer Res.*, 2005, **65**, 10006–10015.
- 63 A. Kamal, L. Thao, J. Sensintaffar, L. Zhang, M. F. Boehm, L. C. Fritz and F. J. Burrows, *Nature*, 2003, **425**, 407.
- 64 J. Davenport, J. R. Manjarrez, L. Peterson, B. Krumm, B. S. J. Blagg and R. L. Matts, *J. Nat. Prod.*, 2011, **74**, 1085–1092.
- 65 J. C. Bijleveld, A. P. Zoombelt, S. G. J. Mathijssen, M. M. Wienk, M. Turbiez, D. M. de Leeuw and R. A. J. Janssen, *J. Am. Chem. Soc.*, 2009, **131**, 16616–16617.
- 66 X. Chen, Z. Zhang, Z. Ding, J. Liu and L. Wang, *Angew. Chem., Int. Ed.*, 2016, **55**, 10376–10380.
- 67 C. Dou, J. Liu and L. Wang, *Sci. China: Chem.*, 2017, **60**, 450–459.
- 68 R. Zhao, C. Dou, J. Liu and L. Wang, *Chin. J. Polym. Sci.*, 2017, **35**, 198–206.
- 69 T. Sun, J. Qi, M. Zheng, Z. Xie, Z. Wang and X. Jing, *Colloids Surf., B*, 2015, **136**, 201–206.
- 70 W. Wang, L. Wang, Y. Li, S. Liu, Z. Xie and X. Jing, *Adv. Mater.*, 2016, **28**, 9320–9325.
- 71 Y. Li, C. Jiang, D. Zhang, Y. Wang, X. Ren, K. Ai, X. Chen and L. Lu, *Acta Biomater.*, 2017, **47**, 124–134.

- 72 Y. Zhang, Z. Yang, X. Tan, X. Tang and Z. Yang, *AAPS PharmSciTech*, 2017, **18**, 1987–1997.
- 73 Y. Yang, L. Yang, Q.-D. You, F.-F. Nie, H.-Y. Gu, L. Zhao, X.-T. Wang and Q.-L. Guo, *Cancer Lett.*, 2007, **256**, 259–266.
- 74 X. Zhu, H. Zhang, Y. Lin, P. Chen, J. Min, Z. Wang, W. Xiao and B. A. Chen, *J. Chemother.*, 2009, **21**, 666–672.
- 75 M. K. Pandey, B. Sung, K. S. Ahn, A. B. Kunnumakkara, M. M. Chaturvedi and B. B. Aggarwal, *Blood*, 2007, **110**, 3517–3525.
- 76 J. Chen, H.-Y. Gu, N. Lu, Y. Yang, W. Liu, Q. Qi, J.-J. Rong, X.-T. Wang, Q.-D. You and Q.-L. Guo, *Life Sci.*, 2008, **83**, 103–109.
- 77 L. Zhao, Q.-L. Guo, Q.-D. You, Z.-Q. Wu and H.-Y. Gu, *Biol. Pharm. Bull.*, 2004, **27**, 998–1003.
- 78 J. Wu, L. Zhao, X. Xu, N. Bertrand, W. Choi II, B. Yameen, J. Shi, V. Shah, M. Mulvale, L. MacLean James and C. Farokhzad Omid, *Angew. Chem., Int. Ed.*, 2015, **54**, 9218–9223.