

Redox-responsive hyaluronic acid-functionalized graphene oxide nanosheets for targeted delivery of water-insoluble cancer drugs

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Background: Gefitinib (Gef), an important epidermal growth factor receptor (EGFR), is used to treat lung cancer, but low water solubility and poor bioavailability severely limit its application in cancer therapy.

Methods: In this study, nano-graphene oxide (NGO) was decorated with hyaluronic acid (HA) by a linker cystamine dihydrochloride containing disulfide bonds (-SS-), followed by the incorporation of gefitinib, thus, constructing a HA-functionalized GO-based gefitinib delivery system (NGO-SS-HA-Gef). Subsequently, studies of biological experiments in vitro and in vivo were performed to investigate the therapeutic effect of the system in lung cancer.

Results: The HA-grafted GO nanosheets possessed enhanced physiological stability, admirable biocompatibility, and no obvious side effects in mice and could act as a nanocarrier for the delivery of gefitinib to tumor. Cellular uptake and intracellular cargo release assays showed that the uptake of NGO-SS-HA by A549 cells was facilitated via CD44 receptor-mediated endocytosis, and that more drug was released from NGO-SS-HA in the presence of GSH than in the absence of GSH. The target-specific binding of NGO-SS-HA to cancer cells with redox-responsive cargo release significantly enhanced the abilities of gefitinib-loaded GO nanosheets to induce cell apoptosis, suppress cell proliferation, and inhibit tumor growth in lung cancer cell-bearing mice.

Conclusion: The results demonstrated the potential utility of NGO-SS-HA-Gef for therapeutic applications in the treatment of lung cancer.

Keywords: nano-graphene oxide, gefitinib, hyaluronic acid, CD44, redox-responsive

Introduction

Lung cancer, one of the most common cancers in the world, has become the leading cause of cancer-related death globally, with non-small-cell lung cancer accounting for ~80%.¹⁻³ Hence, increasing attention has been paid to the study of drugs against lung cancer. In the past few decades, researchers have developed many anti-lung cancer drugs to extend patient survival time, such as paclitaxel,^{4,5} docetaxel,^{6,7} and cisplatin.^{8,9} In addition, a few drugs specifically targeted against lung cancer have been used in clinical treatment, such as gefitinib (Gef).¹⁰⁻¹² Gef can restrict the activity of epidermal growth factor receptor tyrosine kinase (EGFR-TK), enhance the apoptosis of tumor cells, and then inhibit tumor growth. Although Gef has shown strong potential in the treatment of lung cancer, it is limited in this application by some of its clear defects,¹³⁻¹⁵ such as low water solubility, poor bioavailability, and side effects. Therefore, developing a Gef delivery system is necessary to overcome these shortcomings and improve its bioavailability.

Graphene oxide (GO), a prominent type of two-dimensional material, has attracted tremendous attention in many fields, including biomedicine, in recent years because

of its unique properties.^{16–20} However, GO aggregates rapidly in solutions rich in salts or proteins, which hinders the application of GO in biomedicine. Sun et al first reported that nano-graphene oxide (NGO) decorated with polyethylene glycol (PEG) could be dispersed evenly in both physiological solution and cell medium and act as a nanocarrier for doxorubicin loading in intracellular imaging and drug delivery.²¹ Subsequently, Song et al presented hyaluronic acid (HA)-decorated GO as a nanocarrier for targeted and pH-responsive anticancer drug delivery.²² HA, a water-soluble mucopolysaccharide, has excellent biodegradability, biocompatibility, and non-immunogenicity. Moreover, it can also specifically bind the cluster determinant 44 (CD44) receptor, which is overexpressed on the surface of various tumor cells.^{23–25} HA function not only improves the stability of GO in PBS and cell medium but also enhances the uptake of GO by CD44 receptor-overexpressing cells via the specific interaction between the CD44 receptor and HA.

However, similar to PEG,²⁶ HA forms a shell on the surface of nanosheets after combination with NGO, and this shell could restrict the release of a loaded drug from the nanocarrier, which seriously attenuates the therapeutic efficacy of the nanosheets. Therefore, it is necessary to utilize special means to remove the HA barrier for accelerating drug release, such as a redox-responsive mechanism. The redox environment in a tumor cell is an important parameter that may determine the response of a tumor to certain chemotherapeutic agents and radiation. Various intracellular molecules may contribute to the overall redox status in tissues, including glutathione (GSH).^{26–29} GSH, a tripeptide consisting of glutamic acid, cysteine, and glycine, can be found in almost all cells in the body, and the intracellular concentration of GSH is significantly higher than its extracellular concentration.^{30,31} A triggering mechanism based on the evident difference in GSH concentration could therefore be utilized to remove the diffusion barrier caused by HA.

In the current work, a new NGO-based drug delivery system was developed and decorated with HA via disulfide bonds (Figure 1A). The resulting NGO-SS-HA-Gef can not only improve the above shortcomings of Gef in the treatment of lung cancer but also promote the accumulation of nanosheets in tumor sites and facilitate drug release from the nanosheets in response to tumor-relevant GSH. Briefly, HA was conjugated onto the surface of NGO with a linker, cystamine dihydrochloride. The successful preparation of NGO-SS-HA was confirmed by infrared (IR) spectroscopy, UV–vis spectroscopy, and atomic force microscopy (AFM). Then, Gef was physically loaded onto NGO-SS-HA via π – π stacking and hydrophobic interactions to afford a nanosheet complex, NGO-SS-HA-Gef. The surface-engineered structure

of NGO-SS-HA-Gef was found to accelerate the release of Gef from the nanosheets in the presence of GSH. Next, rhodamine B (RB), a fluorescent dye, was loaded onto the NGO-SS-HA to generate a nanohybrid, NGO-SS-HA-RB, which was used to investigate the cellular uptake of NGO-SS-HA and intracellular cargo release from the nanosheets. The *in vitro* cytotoxicity of NGO-SS-HA-Gef against cells was evaluated in a dose- and time-dependent manner. Finally, a series of *in vivo* studies were conducted to investigate the tissue biodistribution of NGO-SS-HA-Gef and evaluate its efficacy in tumor therapy.

Experimental section

Materials

The native graphite flake was purchased from Alfa Aesar ($\geq 99.8\%$, Ward Hill, MA, USA). NaCl ($\geq 99.0\%$), H_3PO_4 ($\geq 85.0\%$), H_2SO_4 (98.0%), H_2O_2 ($\geq 30.0\%$), and HCl ($\geq 36.0\%$) were acquired from Sinopharm Chemical Reagent Co. Ltd (Shanghai, China). Cystamine dihydrochloride and hexamethylenediamine (HMD) were purchased from Bide Pharmatech Ltd (Shanghai, China). Hyaluronic acid (MW = 35 kDa) was obtained from Fureda Biological Technology Co., Ltd (Jinan, China). Gefitinib was purchased from Meilun Biological Technology Co., Ltd (Dalian, China). Cell Counting Kit-8 (CCK-8) and Ham's F-12K (Kaighn's) Medium were purchased from Dingguo Changsheng Biotechnology Co., Ltd (Beijing, China). Glutathione reduced ethyl ester (GSH-OEt, $\geq 90\%$) and glutathione (98%) were purchased from Sigma-Aldrich Co. (St Louis, MO, USA). All other reagents were purchased from J&K Scientific Ltd (Shanghai, China) and used without further purification.

Preparation of NGO

GO was prepared from native graphite flake following an improved Hummer's method.³² The resulting GO was mixed with NaOH aqueous solution (4 mg/mL), and the mixed solution was sonicated at 1,000 W for 3 hours to convert OH groups to COOH groups. The resulting GO-COOH solution was neutralized with dilute HCl, dialyzed in water for 1 week to remove impurities, and filtered through a 0.45 μm filter to fully remove any large-scale GO-COOH for NGO-COOH.

Synthesis of NGO-SS-HA

HA was dispersed in 2-(N-Morpholino) ethanesulfonic acid buffer (pH 6.2) followed by the addition of hydrochloride (EDC) (50 mg) and 1-hydroxybenzotriazole (70 mg). The mixed solution was stirred vigorously for 1 hour. The activated HA was added dropwise into 20 mL of cystamine dihydrochloride aqueous solution (56 mg/mL) over 10 minutes, and then the solution was stirred overnight.

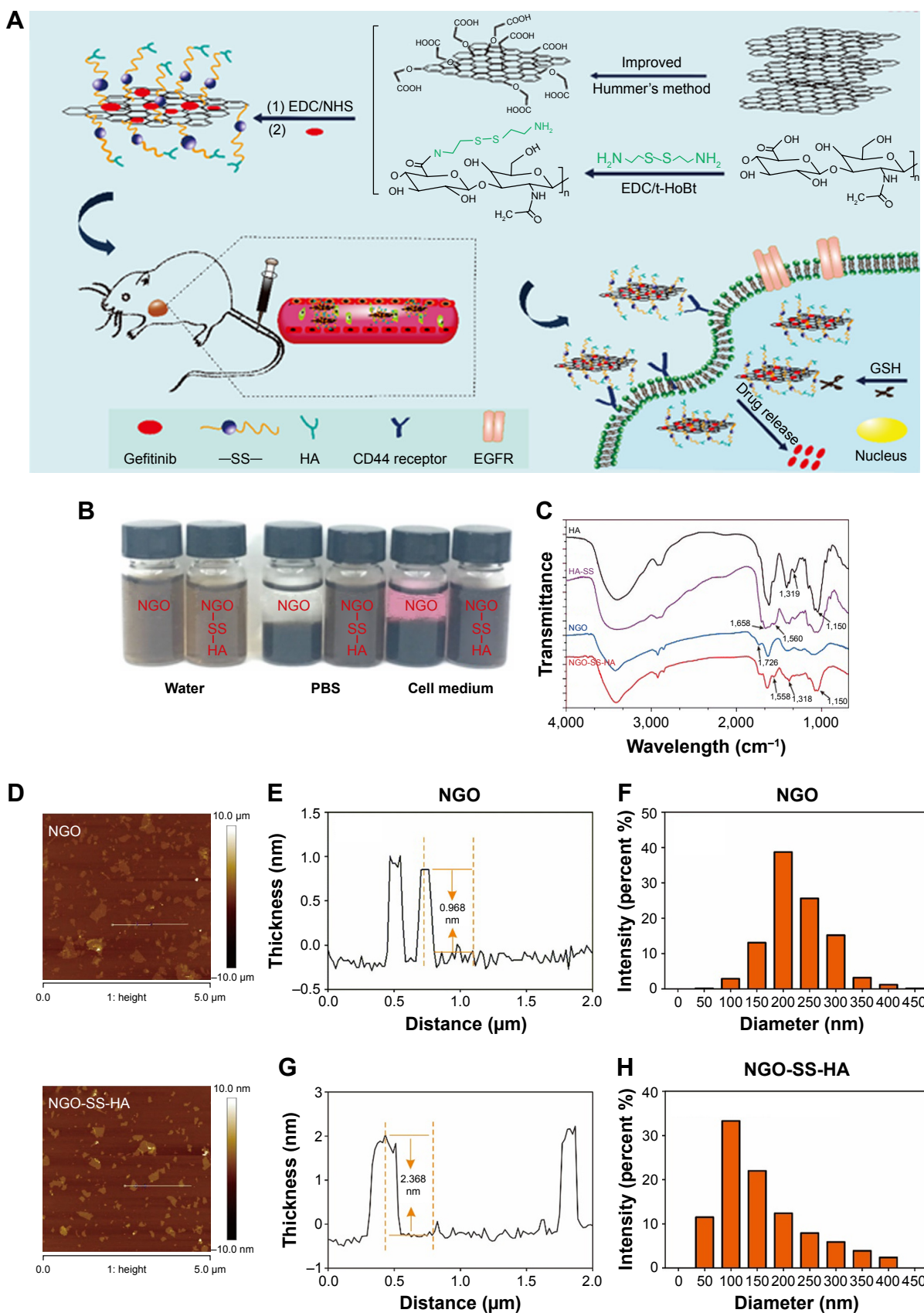


Figure 1 (A) Scheme showing the fabrication process of NGO-SS-HA-Gef, its capability to target cancer cells, and intracellular drug release triggered by redox responsiveness to GSH. (B) Images of NGO and NGO-SS-HA in water, PBS, and cell medium for 1 week. FTIR spectra (C) of HA, HA-SS, NGO, and NGO-SS-HA; AFM images (D) of NGO and NGO-SS-HA. (E and G) The thickness of NGO and NGO-SS-HA measured by AFM. (F and H) Histograms of NGO and NGO-SS-HA diameters, respectively. **Abbreviations:** NGO, nano-graphene oxide; HA, hyaluronic acid; Gef, gefitinib; GSH, glutathione; FTIR, Fourier transform infrared; AFM, atomic force microscopy.

HA-SS-NH₂ was acquired by freeze-drying after dialyzing against water for 2 days.

For the synthesis of NGO-SS-HA, 10 mL of NGO solution (1 mg/mL) was incubated with EDC (15 mg) and *n*-hydroxysuccinimide (NHS, 18 mg) for 1 hour. The resulting *N*-hydroxysuccinimide-activated NGO was washed with water several times by ultrafiltration to remove residual EDC and NHS, and then HA-SS-NH₂ aqueous solution (2 mg/mL) was added. After sonication for 30 minutes and stirring for 24 hours, non-conjugated HA-SS-NH₂ was removed by ultrafiltration. The resulting NGO-SS-HA was stored at 4°C until use.

Loading of Gef on NGO-SS-HA

NGO-SS-HA aqueous solution (0.4 mg/mL) was mixed with various concentrations of Gef in dimethyl sulfoxide (DMSO) solution and stirred for 24 hours. Excess Gef precipitated as solid was removed by centrifugation at 5,000 rpm for 20 minutes. The supernatant was filtered through a 0.45 µm filter to fully remove any solids and then washed five times by ultrafiltration to remove the small amount of solubilized Gef. The obtained NGO-SS-HA-Gef was re-suspended in PBS and stored until use. The UV-vis spectra of NGO-SS-HA before and after drug loading were recorded to determine the drug loading ratio.

GSH-induced Gef release in vitro

To evaluate the pH and redox-responsive release property, NGO-SS-HA-Gef in PBS at different pH values (5.5 and 7.4) and GSH concentrations (0 µM, 2 µM and 10 mM) was loaded into dialysis tubes. At designated time points, 0.2 mL of each dialysis solution was removed, and the same volume of fresh corresponding buffer was added. The amount of Gef released was measured by the UV-vis spectra at a wavelength of 343 nm.

Cell culture

A549 cells and HELF cells were purchased from the Cell Resource Center of Shanghai Institute for Biological Sciences (Chinese Academy of Sciences). A549 cells were cultured in Ham's F12K medium (Sigma-Aldrich Co., Ltd.), and HELF cells were cultured in DMEM. The medium was supplemented with 10% (v/v) fetal bovine serum (Invitrogen) and 1% penicillin/streptomycin. The cells were all maintained at 37°C in a humidified atmosphere with 5% CO₂.

Cytotoxicity of NGO-SS-HA against cells in vitro

The cytotoxicity of NGO-SS-HA against cells was measured using the CCK-8 assay. A549 and HELF cells were seeded into 96-well cell culture plates at 1×10⁴/well for 24 hours and incubated with NGO-SS-HA at various concentrations for another

24, 48, and 72 hours. Next, the absorbance of each well was measured at 450 nm with a microplate reader (Tecan, Mänedorf, Switzerland). The cell viability was computed by the following equation: Cell viability = (OD_{treated}/OD_{control})×100%.

Hemolysis assays in vitro

Blood samples (obtained from a healthy mouse) were stabilized with ethylenediaminetetraacetic acid. Red blood cells were separated from serum by centrifugation at 1,000 rpm for 10 minutes, washed several times with PBS, and then diluted with PBS. Then, 0.3 mL of diluted suspension was mixed with 1) 0.9 mL of PBS as a negative control, 2) 0.9 mL of deionized water as a positive control, or 3) 0.9 mL of NGO-SS-HA dispersions at various concentrations. Next, all the mixtures were shaken at 100 rpm for 2 hours at 37°C and centrifuged at 14,000 rpm. The absorbance of the supernatants at 541 nm was measured by UV-vis spectroscopy. Hemolysis percentage (%) = (A_{treated} - A_{negative})/(A_{positive} - A_{negative})×100%.

Cellular uptake of NGO-SS-HA

To study the mechanism of the nanosheets entering into cells, RB was used as a fluorescent probe to determine the ability of the HA-modified GO nanosheets to specifically bind CD44 receptor-overexpressing cancer cells. The loading of RB onto NGO-SS-HA was similar to that of Gef.

A549 (CD44 receptor-positive) and HELF cells (CD44 receptor-negative) were seeded separately into confocal dishes at densities of 1×10⁵ and 2×10⁵ cell/dish and allowed to grow for 24 hours. Next, the cells were treated with 0 or 5 mg/mL HA for 2 hours and then incubated with NGO-SS-HA-RB ([RB] = 5 µg/mL) for another 2 hours. Then, each dish was washed with PBS three times, and paraformaldehyde (4%) was added to fix the cell morphology. After staining with DAPI, confocal fluorescence images of the cells were acquired by confocal laser scanning microscope (CLSM) in the two channels relevant to DAPI (405 nm) and RB (550 nm).

The cellular uptake of the nanosheets by cells were also measured by fluorescence activated cell sorting (FACS). A549 and HELF cells pre-seeded in six-well plates were incubated with 0 or 5 mg/mL HA for 2 hours. The cells were washed with PBS three times and then incubated with NGO-SS-HA-RB ([RB] = 5 µg/mL) for another 2 hours. After washing with PBS three times, the cells were harvested with trypsin and re-suspended in PBS for FACS analyses.

Redox-dependent intracellular cargo release

To clarify the mechanism of intracellular drug release from the nanosheets, RB was again used as the fluorescent probe.

Briefly, A549 cells were seeded into a confocal dish at a density of 1×10^5 for 24 hours. Next, the cells were treated with fresh cell medium containing GSH-OEt at various concentrations for 2 hours and incubated with NGO-SS-HA-RB ([RB] = 5 $\mu\text{g}/\text{mL}$) for another 2 hours. After washing with PBS twice, fluorescence images of the cells were acquired by CLSM.

For the FACS analyses, A549 cells pre-seeded in six-well plates were treated with GSH-OEt for 2 hours, followed by incubation with NGO-SS-HA-RB ([RB] = 5 $\mu\text{g}/\text{mL}$) for another 2 hours. These cells were analyzed by flow cytometry after trypsinization and washing with PBS three times.

Evaluation of intracellular Gef levels

A549 cells were divided into four groups: group 1 with cells only, group 2 with cells incubated with HA+ NGO-SS-HA-Gef, group 3 with cells treated with NGO-SS-HA-Gef only, and group 4 with cells incubated with GSH-OEt+ NGO-SS-HA-Gef. After the corresponding treatments, the cells were washed with PBS three times, harvested after trypsinization, and centrifuged at 1,500 rpm for 10 minutes. Then, the cells were re-suspended in 1 mL of methanol, sonicated, and centrifuged at 15,000 rpm. Finally, the supernatants were evaluated by an absorbance spectrometer to determine the concentration of Gef.

Therapy in vitro

In the current work, the CCK-8 assay was used to investigate the cytotoxic effects of the NGO-SS-HA-Gef nanosheets on A549 cells. First, A549 cells pre-seeded into 96-well plates were incubated with 0 or 10 mM GSH-OEt for 2 hours and washed with PBS three times. NGO-SS-HA-Gef with various Gef concentrations was added to these 96-well plates and incubated with the cells for 24, 48, and 72 hours. The absorbance of each well was measured at 450 nm on a microplate reader to determine the cell viability.

Detection of the cell cycle and cell apoptosis

To further study the therapeutic efficacy of the nanosheets, the cell cycle and cell apoptosis after various treatments were analyzed by FACS. For the cell cycle assay, A549 cells pre-seeded in 6-well plates were treated with 0 or 10 mM GSH-OEt for 2 hours and washed with PBS three times. The cells were incubated with Gef or NGO-SS-HA-Gef at the same Gef concentrations for 24, 48, and 72 hours. After trypsinization and washing twice with ice-cold PBS, the cells were fixed with cold 70% ethanol for 24 hours at 4°C. Subsequently, the fixed cells were washed with PBS twice by centrifugation at

1,500 rpm and stained with a solution containing 1% (v/v) Triton X-100, 0.01% RNase, and 0.05% propidium iodide (PI) for 30 minutes in the dark for flow cytometry analyses.

For the detection of cell apoptosis, A549 cells pre-seeded in six-well plates were treated with or without 10 mM GSH-OEt for 2 hours. After washing with PBS, fresh cell medium containing Gef or NGO-SS-HA-Gef ([Gef] = 10 $\mu\text{g}/\text{mL}$) was added and incubated with the cells for 24, 48, and 72 hours. Then, the cells were trypsinized, washed with PBS, and stained with fluorescein isothiocyanate-labeled Annexin V and PI. The percentage of apoptotic cells was quantified using flow cytometry.

Tissue biodistribution in vivo

All studies involving animals were carried out in accordance with the NSFC regulation concerning the care and use of experimental animals approved by the Institutional Animal Care and Use Committee of Fuzhou University.

The tissue biodistribution of NGO-SS-HA-Gef was evaluated in mice bearing lung tumors and compared with that of free Gef. To develop the tumor model, A549 cells (5×10^6) suspended in 50 μL of PBS were subcutaneously injected into the forelimbs of female BALB/c nude mice. When the mean tumor volume reached $\sim 200 \text{ mm}^3$, the nude mice were randomly divided into six groups ($n=3/\text{group}$) and treated with NGO-SS-HA-Gef or free Gef (dissolved in PBS containing 0.5% DMSO) at a dose of 20 $\mu\text{g}/\text{mL}$ Gef. At designated time points (1, 2, and 12 hours), all mice were sacrificed, and tissue samples (heart, liver, spleen, lung, kidney, and tumor) were collected and weighed. The concentration of Gef in tissue samples was determined by HPLC. The abovementioned samples (100 mg) were ground with 1 mL of methanol, and 100 μL of the homogenate was mixed with 900 μL of internal standard solution (0.1 $\mu\text{g}/\text{mL}$ erlotinib in methanol) for HPLC analysis with a C18 column (4.6 \times 150 mm, 5 mm). The mobile phase was methanol/ ammonium acetate aqueous solution = 70:30 (v/v), and the detection wavelength was 343 nm.

Therapy in vivo

To study the therapeutic efficacy of NGO-SS-HA-Gef in depth, HA-modified GO nanosheets with HA that cannot be separated in high-concentration GSH solution were developed as a control. HA was conjugated onto the surface of NGO with HMD via EDC/NHS method. Gef was loaded on to NGO-HMD-HA using a similar method as the loading of Gef onto NGO-SS-HA.

The tumor model was developed as described earlier. After the mean tumor volume reached $\sim 100 \text{ mm}^3$, the nude

mice were randomly divided into five groups ($n=5/\text{group}$). Each group of nude mice was intravenously injected with 200 μL of PBS containing 0.5% DMSO, Gef, NGO-HMD-HA-Gef, or NGO-SS-HA-Gef ($[\text{Gef}] = 20 \mu\text{g}/\text{mL}$) every 3 days for 24 days. The tumor sizes were measured by a Vernier caliper and calculated as follows: V (volume, mm^3) = length (mm) \times width² (mm^2)/2. After 24 days, all the mice were sacrificed, and the major organs were collected for hematoxylin and eosin (H&E) and immunohistochemical (IHC) staining.

Results and discussion

Synthesis of NGO-SS-HA

GO was synthesized from graphite flakes using the improved Hummer's method,³² and NGO was obtained by filtering through a 0.45 μm filter. NGO was stable in pure water for a month but aggregated in the presence of salts, especially in PBS (Figure 1B). In this study, HA, a type of hydrophilic mucopolysaccharide, was conjugated with NGO to enhance the stability of NGO in PBS and cell medium. However, we found that HA could generate a shell barrier on the surface of NGO after conjugation, which limited drug release. To address this problem, we conjugated HA with NGO via a disulfide linker. The resulting NGO-SS-HA not only was evenly dispersed in PBS and cell medium but also accelerated the drug release in response to tumor-relevant GSH.

Characterization of GO was achieved by X-ray diffraction (Figure S1) and Raman spectroscopy (Figure S2). After carboxylation of GO, the oxygen content was found to increase from 51.36% to 58.87% (Figure S3), and the zeta potential of GO also changed from -37.5 to -53.0 mV (Figure S4), owing to the conversion of the OH groups into COOH groups. When HA was conjugated with NGO by a disulfide bond, the Fourier transform infrared (FTIR) spectrum confirmed the presence of HA on the NGO nanosheets (Figure 1C). Briefly, the FTIR spectrum of HA-SS showed peaks at $1,658 \text{ cm}^{-1}$ and $1,560 \text{ cm}^{-1}$ associated with amide bonds, confirming that cystamine dihydrochloride was coupled with HA through amide bonds. Next, HA-SS was conjugated with NGO via the EDC/NHS method. The spectrum showed disappearance of the peak at $1,736 \text{ cm}^{-1}$ for NGO and the presence of the characteristic peaks of HA-SS at $1,318 \text{ cm}^{-1}$ and $1,150 \text{ cm}^{-1}$ (skeletal acetal valence band), suggesting successful coupling of HA on the surface of NGO. UV-vis spectroscopy is also a good technique for the characterization of NGO-SS-HA (Figure S5).

We then used AFM to characterize NGO nanosheets before and after HA functionalization (Figure 1D and E).

As-prepared NGO nanosheets were found to have an average thickness of ~ 1 nm, suggesting the complete exfoliation of the NGO nanosheets down to individual ones.³³ After HA functionalization, the average thickness of the NGO nanosheets increased to ~ 2.4 nm (Figure 1G), whereas their average diameter decreased from ~ 250 to ~ 125 nm (Figure 1F and H), mainly because of sonication.

Drug loading and redox-responsive drug release in vitro

GO, a 2D material, has the advantage of an ultrahigh surface area,^{34–36} which facilitates the loading of Gef onto nanosheets. To investigate the potential Gef loading ability of NGO-SS-HA nanosheets, NGO-SS-HA solution (0.4 mg/mL) was mixed with various concentrations of Gef and stirred for 24 hours at pH 7.0. UV-vis spectroscopy was employed to determine the Gef loading efficiency. The absorbance of NGO-SS-HA-Gef in 343 nm (Figure 2A) increased with increasing weight ratio (between Gef and NGO-SS-HA). When the ratio rose from 2 to 2.5, the absorbance was found to not increase further (Figure 2B), indicating that maximum drug loading was reached and was $\sim 13.8\%$.

Many experimental factors are involved in drug release from nanocarriers,³⁷ such as nanoparticle size, pH, temperature, redox response, and drug-carrier interactions. To study the behavior of Gef release from the nanosheets, we investigated the drug release change as the function of pH and GSH concentration through a dialysis method. At a pH of 5.5 without GSH, 30.8% of Gef was released from the nanosheets over 72 hours (Figure 2C). Upon exposure to 10 mM GSH, 60.1% of Gef was released from the nanosheets under the same conditions, and the rate of drug release was 1.95 times faster than that in the absence of GSH, suggesting that the GSH level in tumor cells can trigger Gef release from NGO-SS-HA-Gef. When 1 or 0.1 mM GSH was present in the release buffer, NGO-SS-HA-Gef also exhibited redox-responsive release with a cumulative percent release of $>44.1\%$ or 39.2% over 72 hours, respectively. Interestingly, we also found that the cumulative drug release with 10 μM GSH was similar to that without GSH, indicating that the linkages between HA and NGO in NGO-SS-HA-Gef would be relatively stable in vivo in the circulation.

In the control experiments performed at pH 7.4 (Figure 2D), drug release acceleration was also observed in the presence of various concentrations of GSH, but the cumulative release curve of Gef over 72 hours was similar to that at pH 7.4 under the same conditions, suggesting that pH had almost no effect on Gef release from the nanosheets.

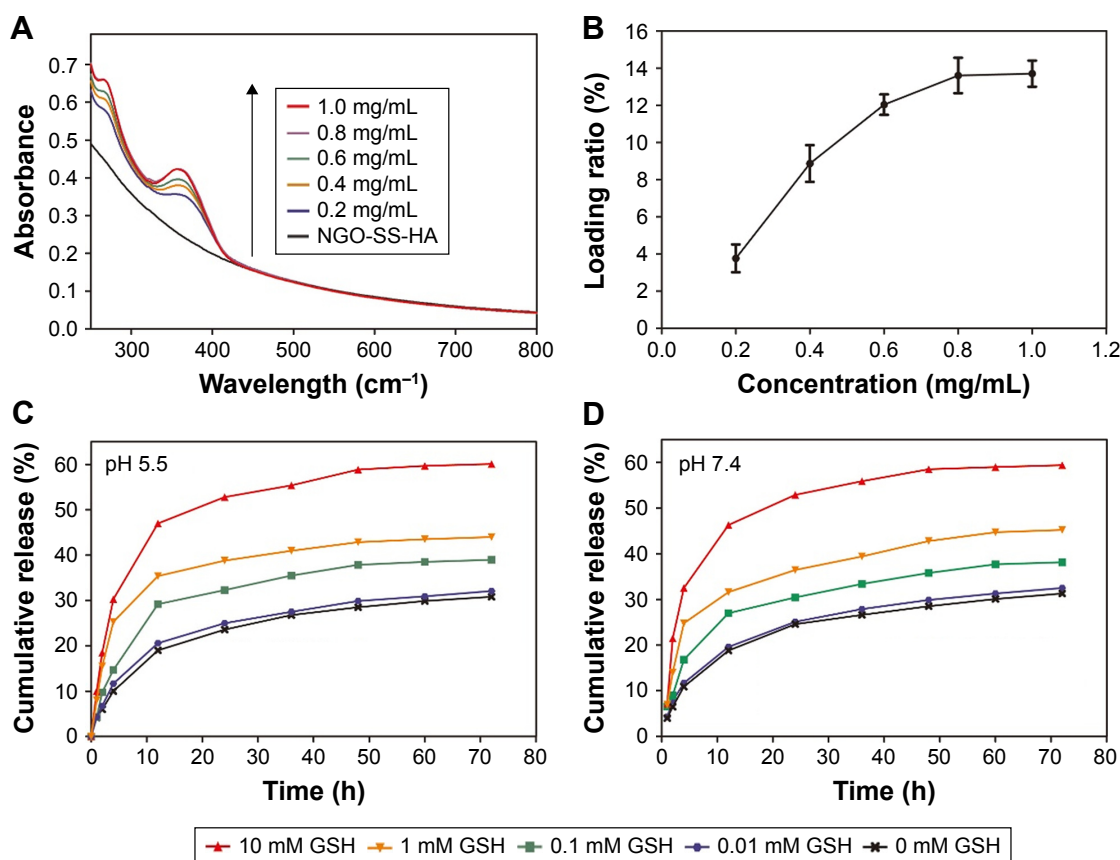


Figure 2 (A) UV-vis spectra of NGO-SS-HA-Gef obtained at different drug loading concentrations after the removal of free Gef. (B) Plots of the ratio of Gef loaded onto NGO-SS-HA at various concentrations of Gef. (C, D) Gef release from NGO-SS-HA-Gef at pH 5.5 and 7.4 in the presence of GSH as a function of time. **Abbreviations:** NGO, nano-graphene oxide; HA, hyaluronic acid; Gef, gefitinib; GSH, glutathione.

Therefore, the abovementioned results illustrated that NGO-SS-HA-Gef can be used to limit Gef release during blood circulation and accelerate intracellular drug release in tumor cells based on its redox responsiveness to GSH.

Biocompatibility in vitro

Biocompatibility is an essential concern in the development of nanomaterials for biomedical applications.³⁸ Therefore, evaluating the cytotoxicity of NGO-SS-HA was necessary. After incubation with high concentrations of NGO-SS-HA for 72 hours, >88.0% of the A549 cells remained alive, even when the nanosheet concentration reached 400 $\mu\text{g}/\text{mL}$ (Figure 3A). Similar results were observed for HELF cells (Figure 3B), further confirming that NGO-SS-HA had low cytotoxicity against cells.

In addition, a hemolysis assay was employed to evaluate the blood compatibility of NGO-SS-HA. After various treatments, we found that the hemolysis percentage of NGO-SS-HA was below 4% at nanosheet concentrations ranging from 25 to 400 $\mu\text{g}/\text{mL}$ (Figure 3C), indicating that NGO-SS-HA had admirable blood compatibility with cells.

As a result, the low cytotoxicity and hemolytic activity of the NGO-SS-HA nanosheets contribute to its application in biological fields.

Cellular uptake and intracellular drug release

The cellular association of the nanosheets was determined using RB as a fluorescent probe.³⁹ RB was loaded onto the surface of NGO-SS-HA through π - π stacking and hydrophobic interactions, and a new peak at 575 nm appeared in the UV-vis spectra (Figure S6). Figure S7 shows that the fluorescence of aqueous NGO-SS-HA-RB solution was weaker than that of free RB at the same RB concentration, suggesting partial fluorescence quenching caused by fluorescence resonance energy transfer.⁴⁰

To verify the targeting specificity, both A549 (CD44 receptor-positive) and HELF cells (CD44 receptor-negative) were incubated with NGO-SS-HA-RB for 2 hours, and confocal images of these cells were obtained by CLSM at an excitation wavelength of 550 nm. A strong red fluorescence was evident in A549 cells after incubation with

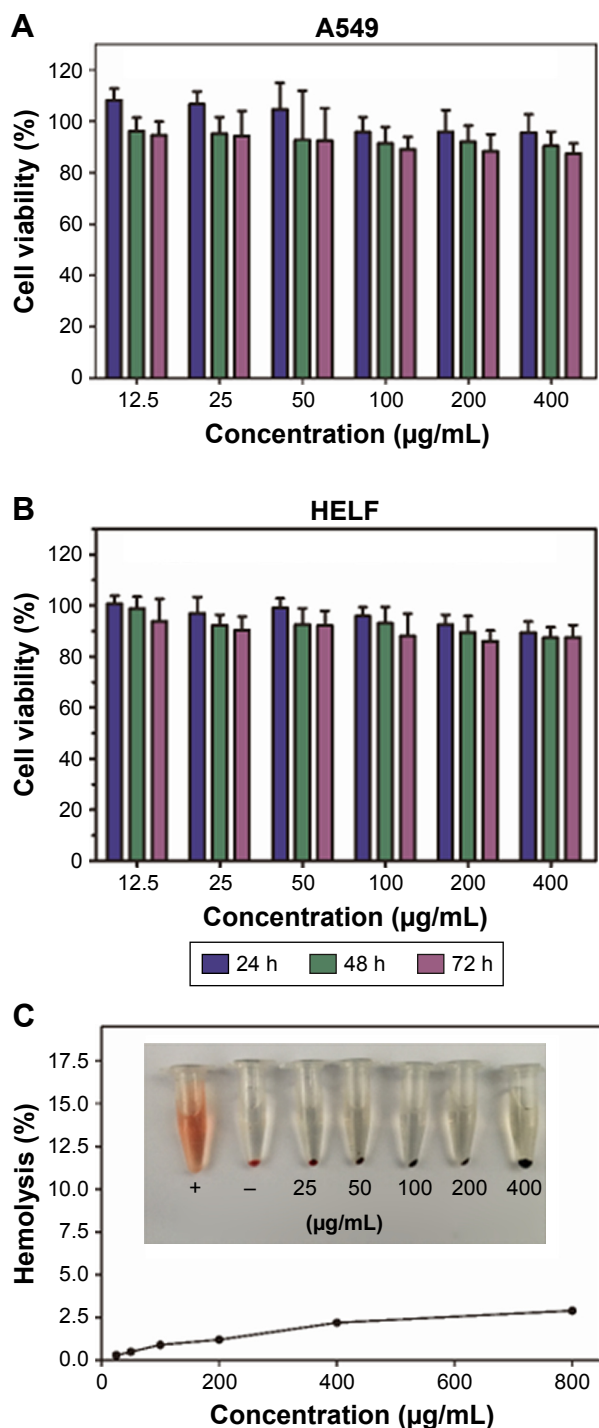


Figure 3 Cell viability of (A) A549 cells and (B) HELF cells incubated with various concentrations of NGO-SS-HA from 24 to 72 hours. (C) Hemolysis percentage of RBCs incubated with NGO-SS-HA at various concentrations for 2 hours. Inset: image of direct observation of hemolysis by NGO-SS-HA.

Abbreviations: NGO, nano-graphene oxide; HA, hyaluronic acid; Gef, gefitinib; h, hours.

NGO-SS-HA-RB, but a very weak fluorescence signal was detected in HELF cells (Figure 4A). Upon pre-treatment with HA, the fluorescence of A549 cells was greatly reduced, suggesting that the cellular uptake of NGO-SS-HA-RB was

associated with HA CD44-receptor-mediated endocytosis and could be blocked by excess free HA. The fluorescence intensity of these cells was also quantitatively detected by FACS. As expected, the results of flow cytometry (Figure 4B and C), consistent with the CLSM data, further confirmed that the uptake of NGO-SS-HA by A549 cells was facilitated by the specific interaction between CD44 receptor and HA.

To study the redox-responsive property of the nanosheets with disulfide linkages, GSH-OEt was included in the drug release experiment.²⁶ GSH-OEt, a derivative of GSH, had no cytotoxicity to A549 cells (Figure S8) and could act as an external enhancer of the cellular GSH level. A549 cells were pre-treated with various concentrations of GSH-OEt (0, 0.01, 0.1, 1, and 10 mM) for 2 hours, incubated with NGO-SS-HA-RB ([RB] = 5 µg/mL) for another 2 hours, and then imaged by CLSM (Figure 4D). We found that the intracellular red fluorescence became increasingly strong with increasing GSH-OEt concentration, indicating that more RB was released from the nanosheets in the presence of GSH, mainly because of the removal of the HA shell via the cleavage of disulfide bonds caused by GSH in a dose-dependent manner. Furthermore, we also studied GSH-dependent cellular RB release by FACS. Similar to the CLSM results, FACS (Figure 4E) revealed that stronger fluorescence intensity could be observed in the cells pre-treated with 10 mM GSH-OEt than the cells pre-treated with lower concentrations of GSH-OEt. These results suggested that the drug release from NGO-SS-HA could be accelerated by GSH-mediated degradation of the disulfide linkages between HA and NGO.

Based on the abovementioned results, we directly investigated the concentration of intracellular Gef by UV-vis spectroscopy. As shown in Figure 4F, cells incubated with NGO-SS-HA-Gef contained substantially more Gef than cells pre-incubated with HA. More importantly, upon pre-incubation of cells with 10 mM GSH-OEt, the intracellular concentration of Gef increased and was 2.05 times higher than that in the absence of GSH-OEt under the same conditions, consistent with the drug release results in vitro. Therefore, the results indicated that NGO-SS-HA-Gef could not only specially bind with CD44 receptor-overexpressing cancer cells but also accelerate Gef release from intracellular nanosheets based on its redox responsiveness to GSH.

Anti-proliferative assay in vitro

To study the cytotoxicity of NGO-SS-HA-Gef to cancer cells in vitro, we incubated A549 cells with NGO-SS-HA-Gef at various Gef concentrations for 24, 48, and 72 hours and then investigated cell viability by the CCK-8 assay. As shown in

Figure 5A, both free Gef and NGO-SS-HA-Gef inhibited the proliferation of A549 cells in a dose-dependent manner. NGO-SS-HA-Gef exhibited a greater cytotoxic effect on A549 cells than free Gef at the same Gef concentration, possibly due to HA-targeting-enhanced cellular uptake of Gef provided by the nanosheets. After the addition of GSH-OEt, NGO-SS-HA-Gef showed the strongest cytotoxicity, revealing that redox responsiveness may be an effective strategy to accelerate drug release in GSH-rich environments, thus resulting in enhanced antitumor activity. In addition, the effect of incubation time on the cytotoxicity of NGO-SS-HA-Gef was also investigated. The cytotoxicity of both Gef and NGO-SS-HA-Gef against A549 cells was found to become stronger with increasing incubation time (Figure 5B and C). Moreover, NGO-SS-HA-Gef in the presence of GSH exhibited the strongest anti-proliferative effect among the test groups. Therefore, we determined that NGO-SS-HA could function as an excellent nanocarrier of Gef by distinctly enhancing its cytotoxicity through CD44 receptor-mediated

endocytosis and by accelerating drug release based on the redox responsiveness of the carrier to GSH.

Cell cycle and cell apoptosis

The effects of NGO-SS-HA-Gef on cell proliferation may be explained in part by its effects on the cell cycle.⁴¹ Therefore, we treated A549 cells with NGO-SS-HA-Gef in a time-dependent manner and then analyzed the cell cycle. The percentage of A549 cells treated with Gef for 24 hours in the G0/G1 phase was ~60.53%, which was lower than that of the controls, indicating that Gef can cause G0/G1 arrest of A549 cells (Figure 6A). Following treatment of the A549 cells with NGO-SS-HA-Gef at the same Gef concentration, the percentage of cells in the G0/G1 phase increased to ~71.21%, indicating that NGO-SS-HA-Gef can enhance Gef-induced G0/G1 arrest. In other words, NGO-SS-HA-Gef could better prevent the cells from entering the S phase, during which some proteins involved in DNA replication are copied and thus inhibit cell proliferation. When the cells were

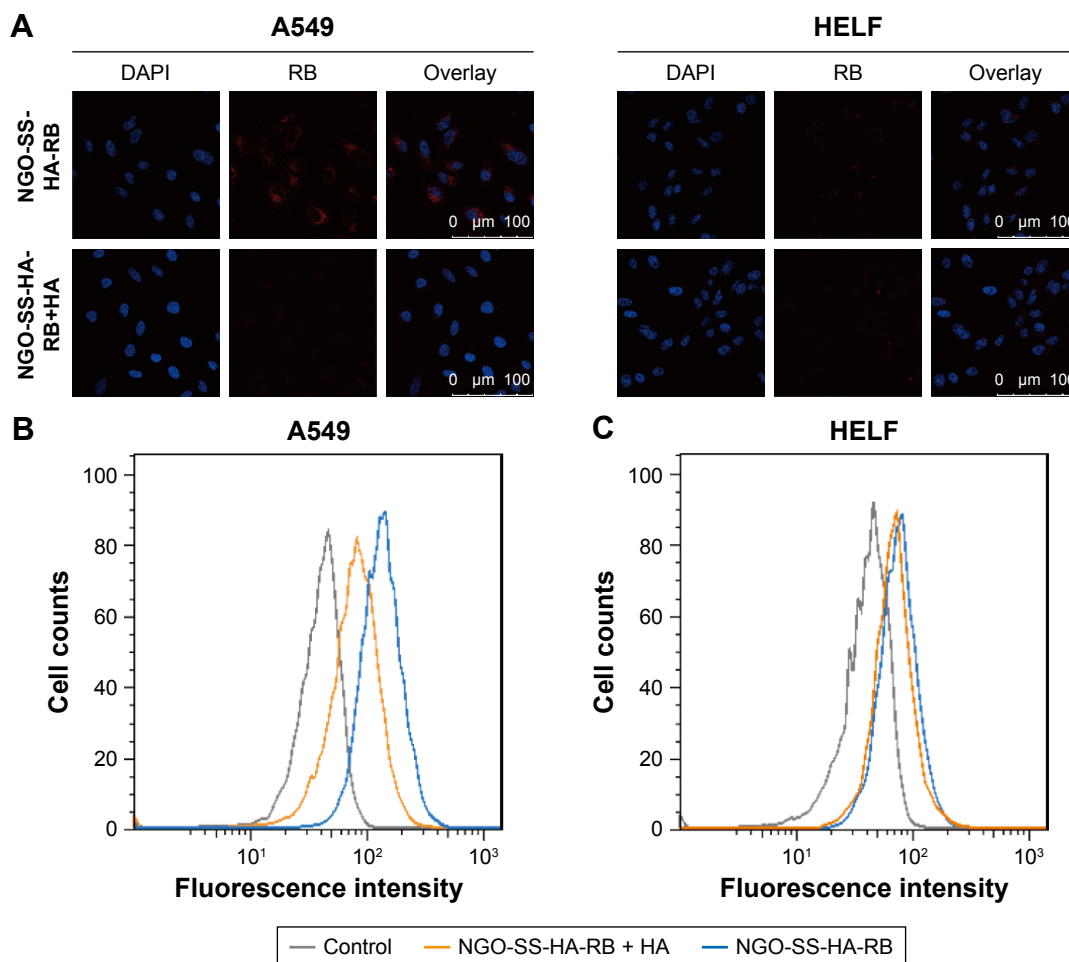


Figure 4 (Continued)

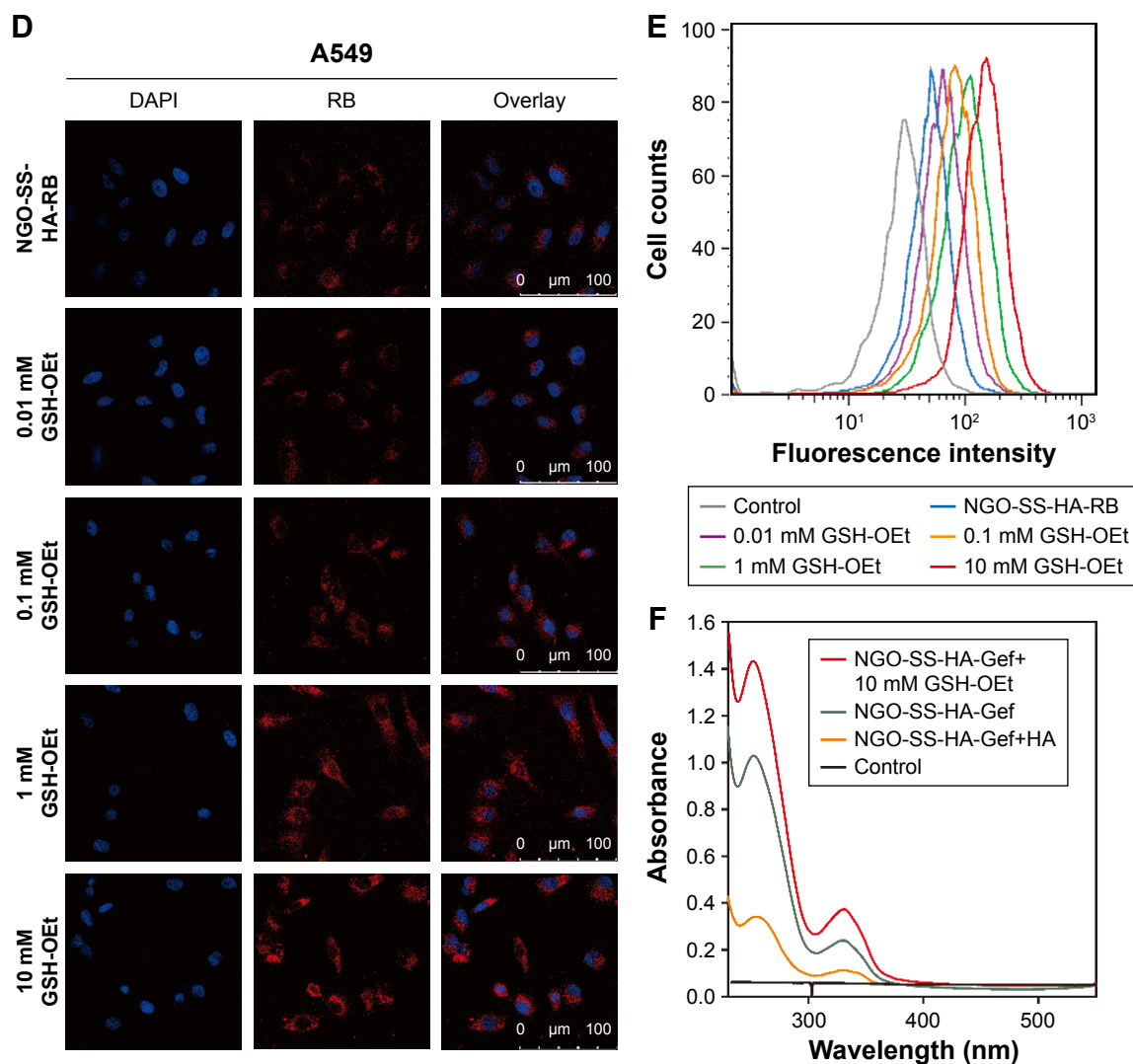


Figure 4 Confocal images (A) and flow cytometric analysis (B and C) of A549 and HELF cells with or without pre-treatment with excess HA incubated with NGO-SS-HA-RB for 2 hours. Confocal images (D) and flow cytometric measurement (E) of RB release from in A549 cells pre-treated with various concentrations of GSH-OEt; UV-vis spectra (F) of intracellular Gef concentration after A549 cells were treated with various treatments.

Abbreviations: NGO, nano-graphene oxide; HA, hyaluronic acid; Gef, gefitinib; RB, rhodamine B; GSH-OEt, glutathione reduced ethyl ester.

pre-incubated with 10 mM GSH-OEt, the percentage of cells in the G0/G1 phase continued to increase from ~71.21% to ~79.83%, suggesting that NGO-SS-HA-Gef more strongly arrested cells in the G0/G1 phase due to accelerated drug release based on its redox responsiveness to GSH. On the other hand, after increasing the incubation time, we found (Figure 6B and C) that the percentage of cells in the G0/G1 phase in each group increased in a time-dependent manner. Together with the abovementioned results, NGO-SS-HA-Gef was found to enhance G0/G1 arrest of A549 cells in a time-dependent manner, especially in the presence of GSH, thus preventing cells from progressing to the S phase and inhibiting their proliferation.

To explore whether the cell growth inhibition *in vitro* observed was due to enhanced apoptosis, the proportion of apoptotic cells was determined by FACS using Annexin V/PI staining.⁴² After incubation with NGO-SS-HA-Gef for 24 hours, the NGO-SS-HA-Gef-treated A549 cells exhibited an apoptotic rate of ~15.02%, which was significantly higher than that in cells treated with free Gef (Figure 6D). After pre-treatment with 10 mM GSH-OEt, the apoptotic rate of cells increased to ~25.42%. The significant cell-killing effect was attributed to the HA-targeting-enhanced cellular uptake of Gef-loaded nanosheets and GSH-triggered drug release. When the incubation time was increased to 48 or 72 hours, the apoptotic rate of each group also increased (Figure 6E

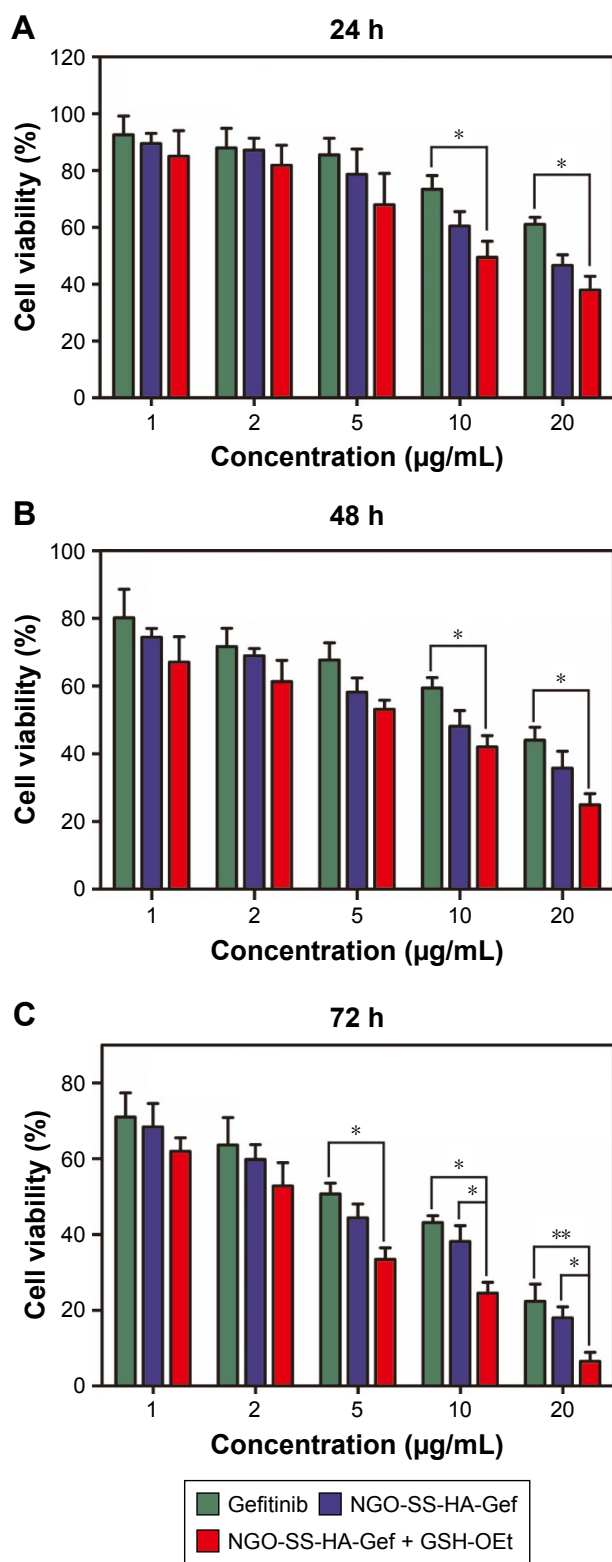


Figure 5 Cell viability of A549 cells after incubation with free Gef, NGO-SS-HA-Gef, or NGO-SS-HA-Gef + 10 mM GSH-OEt for 24 (A), 48 (B), and 72 hours (C). * $P < 0.05$, ** $P < 0.01$ (NGO-SS-HA-Gef + GSH-OEt vs free Gef and NGO-SS-HA-Gef only). The differences between groups were considered statistically significant when the P -value was < 0.05 .

Abbreviations: NGO, nano-graphene oxide; HA, hyaluronic acid; Gef, gefitinib; RB, rhodamine B; GSH-OEt, glutathione reduced ethyl ester.

and F), suggesting that the effects on the apoptotic rate of A549 cells are time-dependent, which is consistent with the in vitro results of the anti-proliferative assay.

Tissue biodistribution in vivo

Tissue biodistribution of Gef was assessed in A549 tumor-bearing mice. Gef was found to disperse rapidly, that is, within 1 hour, into the major organs of mice (heart, liver, spleen, lung, and kidney) after intravenous injection of free Gef or NGO-SS-HA-Gef (Figure 7A). The amount of Gef in the major organs followed the order: liver $>$ lung $>$ spleen $>$ kidney $>$ heart. The Gef contents in major organs reached maximum levels after 2 hours (Figure 7B) and decreased at 12 hours (Figure 7C), mainly owing to the gradual degradation and metabolism of the abovementioned two formulations in vivo.

At all the time points, the Gef content in the tumors of mice after intravenous injection with NGO-SS-HA-Gef was higher than that in the tumors of mice injected with free Gef. In particular, at 2 hours, the Gef level in the tumors of the NGO-SS-HA-Gef-treated mice was 2.12-fold ($P < 0.05$) greater than that in the tumors of the mice treated with free Gef. This result was attributed to the efficient accumulation of NGO-SS-HA-Gef in the tumors through a combination of passive (enhanced permeability and retention effect) and active (HA as a targeting moiety for the CD44 receptor) targeting.

Antitumor efficacy in vivo

Encouraged by the fascinating therapeutic effect of NGO-SS-HA-Gef in vitro, we next conducted comparative efficacy studies in vivo using the nanosheets. After the tumor volume reached $\sim 100 \text{ mm}^3$, these nude mice were randomly divided into five groups with five mice per group. The mice were intravenously injected with PBS, NGO-SS-HA, free Gef, NGO-HMD-HA-Gef, or NGO-SS-HA-Gef every 3 days. During 24 days of treatment, the body weight and the tumor volume of each group were recorded every time. The average body weight of each group showed no significant variation (Figure 8A), suggesting that HA-grafted NGO exerted no significant side effects. The mice treated only with NGO-SS-HA experienced rapid tumor volume increases as a function of time, similar to the observations in PBS-treated mice, indicating that NGO-SS-HA induced a negligible effect on tumor growth (Figure 8B). Moreover, compared with free Gef, NGO-HMD-HA-Gef was observed to significantly improve anticancer efficacy, as verified by a lower tumor growth rate. Furthermore, owing to HA CD44-receptor-mediated tumor targeting and GSH-triggered drug

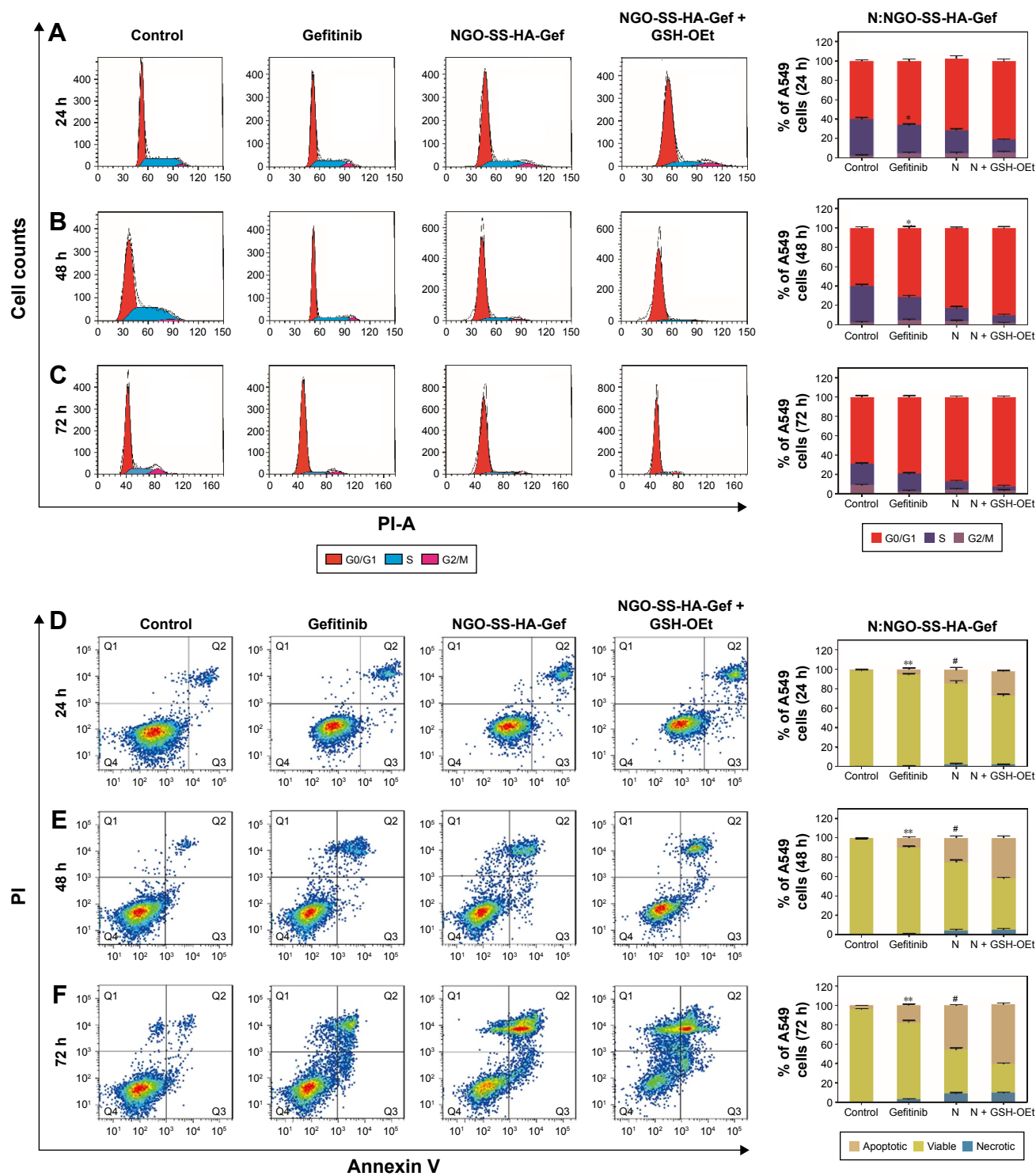


Figure 6 (A–C) Cell cycle distribution (G0/G1, S, and G2/M) was determined after various treatments by flow cytometry. A549 cells were treated with free Gef or NGO-SS-HA-Gef ([Gef] = 10 µg/mL) in the presence or absence of GSH-OEt. Controls remained untreated. **(D–F)** Apoptotic cells were detected by staining with FITC-Annexin V/PI after various treatments. Controls remained untreated. * $P < 0.05$, ** $P < 0.01$ compared with the Gef group. # $P < 0.05$ compared with the NGO-SS-HA-Gef only group. **Abbreviations:** NGO, nano-graphene oxide; HA, hyaluronic acid; Gef, gefitinib; RB, rhodamine B; GSH-OEt, glutathione reduced ethyl ester; FITC, fluorescein isothiocyanate; PI, propidium iodide.

release, mice treated with NGO-SS-HA-Gef experienced the lowest tumor volume growth among these groups, indicating that NGO-SS-HA-Gef can significantly inhibit the tumor growth, which is consistent with the results of the CCK-8 assay. After 24 days of various treatments, all the mice were

sacrificed, and their tumors (Figure 8C) and major organs were collected and sectioned for histological analysis. H&E staining of the major organs (Figure 8D) showed that there was no appreciable organ damage. The therapeutic efficacy of NGO-SS-HA-Gef in A549 tumor-bearing mice could

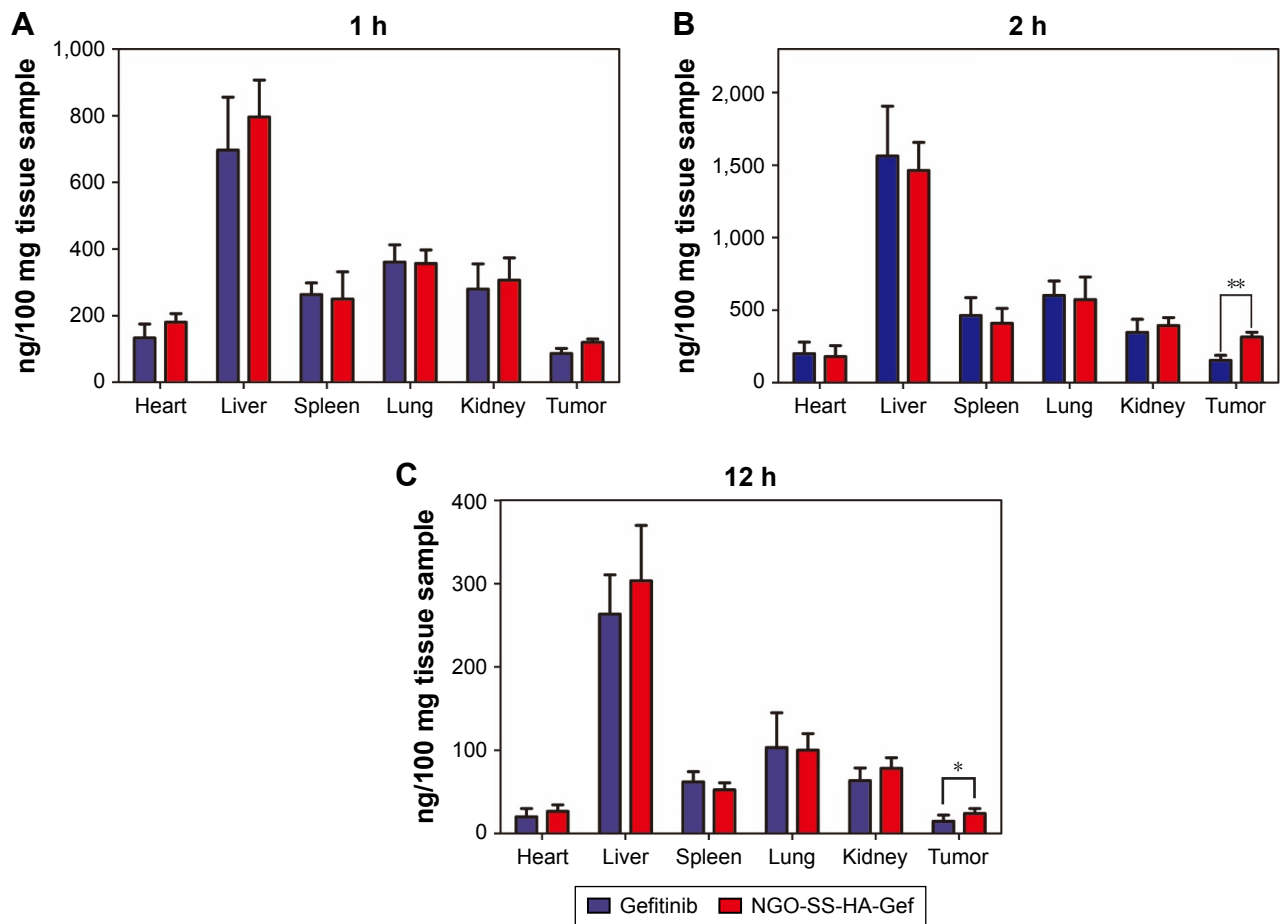


Figure 7 Tissue biodistribution of Gef after intravenous administration of NGO-SS-HA-Gef to A549 tumor-bearing mice: (A) 1, (B) 2 and (C) 12 hours following administration, with free Gef used as a control. Error bars were based on three parallel mice. Significant differences between groups are labeled with * $P < 0.05$ and ** $P < 0.01$.

Abbreviations: NGO, nano-graphene oxide; HA, hyaluronic acid; Gef, gefitinib.

also be detected by H&E staining of tumors. H&E staining (Figure 8E) revealed that the PBS group showed vigorous tumor growth, whereas heteromorphosis along with cell necrosis and lysis occurred in tumor cells in the Gef, NGO-

HMD-HA-Gef, and NGO-SS-HA-Gef groups. A large number of dead cells in the tumor were observed in the mice treated with NGO-SS-HA-Gef, indicating excellent antitumor efficacy. Gef, an important EGFR-TK inhibitor, can

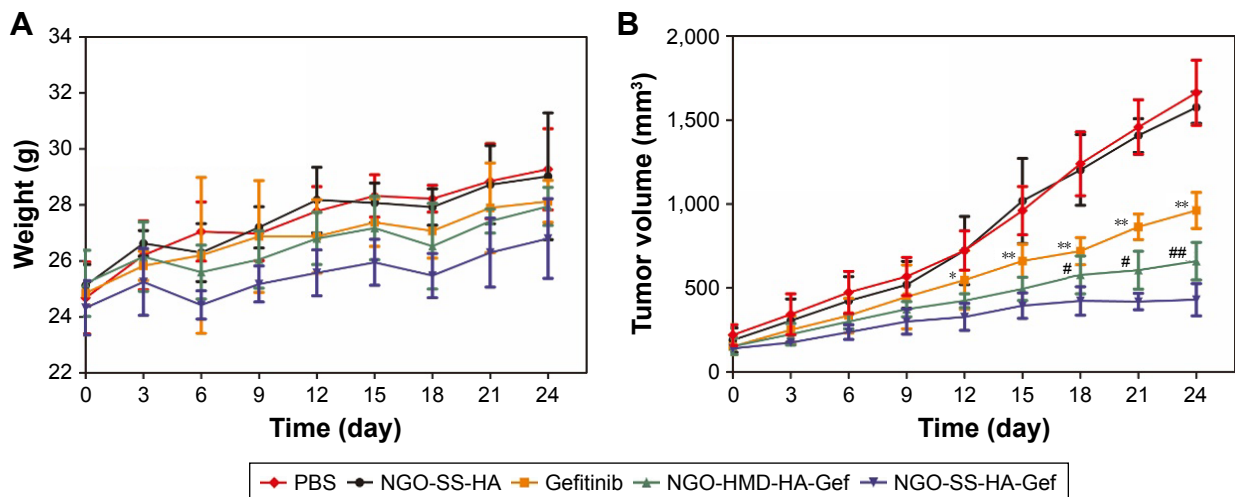


Figure 8 (Continued)

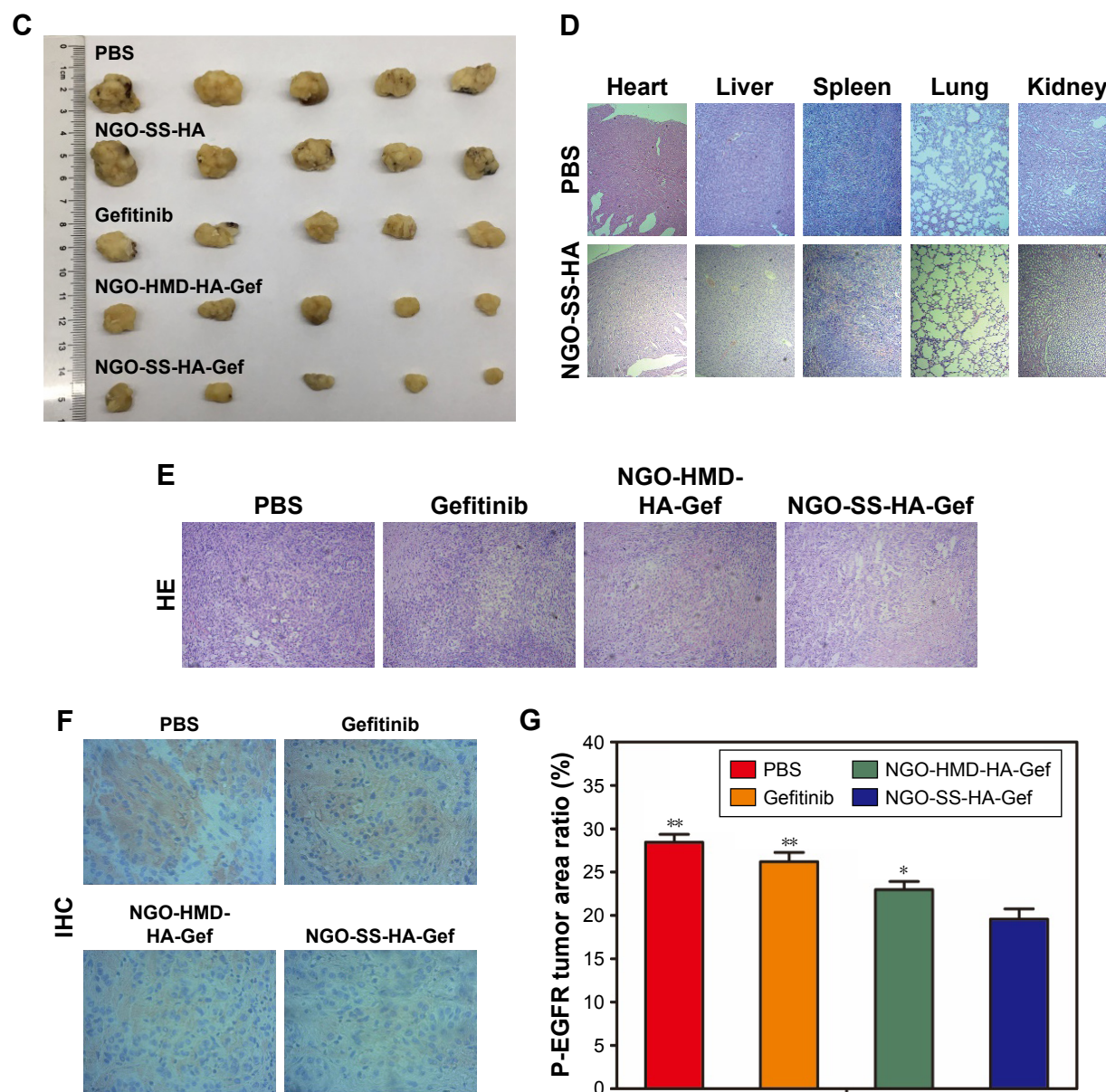


Figure 8 Comparative investigation of tumor inhibitory effects in vivo.

Notes: (A) Relative mouse body weight of each group as a function of time. (B) Tumor growth curves of different groups of mice after various treatments. Significant differences appeared between the NGO-SS-HA-Gef and the free Gef groups and are marked as * $P < 0.05$ and ** $P < 0.01$. Significant differences appeared between the NGO-SS-HA-Gef and NGO-HMD-HA-Gef groups and are labeled with # $P < 0.05$ and ### $P < 0.01$. (C) Photographs of tumors from the five groups. (D) H&E images of major organs from control mice and mice 24 days post-NGO-SS-HA injection. (E) H&E staining of tumors from different groups. (F) IHC analysis of the expression of P-EGFR in mice; P-EGFR was stained brown. (G) Quantification of the P-EGFR expression in mice after various treatments. * $P < 0.05$ and ** $P < 0.01$ compared with the NGO-SS-HA-Gef group.

Abbreviations: NGO, nano-graphene oxide; HA, hyaluronic acid; Gef, gefitinib; HMD, hexamethylenediamine; RB, rhodamine B; GSH-OEt, glutathione reduced ethyl ester; P-EGFR, phosphorylation of epidermal growth factor receptor tyrosine; H&E, hematoxylin and eosin; IHC, immunohistochemical.

significantly inhibit the phosphorylation of EGFR tyrosine (P-EGFR),⁴³ which is involved in cell growth, proliferation, differentiation, and so on. The expression of P-EGFR was examined by IHC staining. The results (Figure 8F and G) revealed that among the treatment groups, the NGO-SS-HA-Gef groups had the lowest expression of P-EGFR. These IHC results were correlated well with the antitumor capacity observed in the abovementioned studies.

Conclusion

We developed a redox-stimuli responsive targeted delivery system based on GO nanosheets, in which HA was conjugated with the nanosheets by disulfide bonds. The resulting NGO-SS-HA showed physiological stability, admirable biocompatibility, and low cytotoxicity in vitro and in vivo. Gef could be loaded on the surface of NGO-SS-HA through π - π stacking and hydrophobic interactions. However, the HA fixed

on the surface of the nanosheets could function as a shell barrier, limiting the Gef release from NGO-SS-HA-Gef. A redox-responsive mechanism, that is, cleavage of disulfide bonds in the presence of GSH, was used to accelerate the release of Gef from NGO-SS-HA-Gef, as evidenced by a drug release assay in vitro. Both flow cytometric analysis and confocal microscopy showed that the uptake of NGO-SS-HA by A549 cells was facilitated through CD44 receptor-mediated endocytosis and that intracellular drug release from the nanosheets could be induced in the presence of GSH. After incubation with A549 cells, NGO-SS-HA-Gef was found to significantly enhance the inhibitory activity of Gef against cell proliferation and induce cell apoptosis. Tumor-bearing mice treated with NGO-SS-HA-Gef experienced the lowest tumor growth rates among the test groups, indicating that injection of NGO-SS-HA-Gef induced the strongest tumor inhibition.

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Disclosure

The authors report no conflicts of interest in this work.

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