

REVIEW

View Article Online
View Journal | View Issue

Cite this: *Biomater. Sci.*, 2024, **12**, 581

Hydroxyl-rich branched polycations for nucleic acid delivery

Mengrui Su,^{†a} Zichen Hu,^{†a} Yujie Sun,^a Yu Qi,^{*b,c} Bingran Yu ^{*a} and Fu-Jian Xu ^{*a}

Recently, nucleic acid delivery has become an amazing route for the treatment of various malignant diseases, and polycationic vectors are attracting more and more attention among gene vectors. However, conventional polycationic vectors still face many obstacles in nucleic acid delivery, such as significant cytotoxicity, high protein absorption behavior, and unsatisfactory blood compatibility caused by a high positive charge density. To solve these problems, the fabrication of hydroxyl-rich branched polycationic vectors has been proposed. For the synthesis of hydroxyl-rich branched polycations, a one-pot method is considered as the preferred method due to its simple preparation process. In this review, typical one-pot methods for fabricating hydroxyl-rich polycations are presented. In particular, amine-epoxide ring-opening polymerization as a novel approach is mainly introduced. In addition, various therapeutic scenarios of hydroxyl-rich branched polycations *via* one-pot fabrication are also generalized. We believe that this review will motivate the optimized design of hydroxyl-rich branched polycations for potential nucleic acid delivery and their bio-applications.

Received 26th August 2023,
Accepted 14th November 2023

DOI: 10.1039/d3bm01394d

rsc.li/biomaterials-science

1. Introduction

Nucleic acid delivery is an approach that utilizes therapeutic nucleic acids, including DNA, RNA, and oligonucleotides, to treat

conditions such as cancers, immunodeficiency, and many human malignancies.^{1–7} To achieve optimal therapeutic effect, the nucleic acids require efficient delivery to the targeted cells and efficient vectors are necessary. As we know, vectors can be mainly divided into two categories: viral and non-viral vectors. Viral vectors, such as adenoviruses and retroviruses, are restricted due to their uncontrolled bio-risks.^{8–10} Non-viral vectors mainly include liposomes,^{11,12} peptides,^{13–17} polycations,^{18–22} and organic-inorganic hybrid materials.^{7,23,24} Among non-viral vectors, polycations have attracted more attention due to their low manufacturing cost and robust modifiability. However, significant cytotoxicity and unsatisfactory transfection performance still

^aKey Laboratory of Biomedical Materials of Natural Macromolecules (Beijing University of Chemical Technology, Ministry of Education), Beijing Laboratory of Biomedical Materials, Beijing University of Chemical Technology, Beijing 100029, PR China. E-mail: yubr@mail.buct.edu.cn, xuff@mail.buct.edu.cn

^bChina Meat Food Research Center, Beijing Academy of Food Sciences, Beijing 100068, PR China. E-mail: qiyu_0501@163.com

^cBeijing Forestry University, Beijing, 100083, PR China

[†]These authors contributed equally to this work.



Mengrui Su

Mengrui Su received her Master's degree in Materials Science and Engineering from Beijing University of Chemical Technology in 2023. She is currently a doctoral student supervised by Prof. Bingran Yu at Beijing University of Chemical Technology. Her current research interests are focused on the design and synthesis of cationic gene delivery systems and their application in biomedicine.



Zichen Hu

Zichen Hu received his BSc degree in Polymer Materials and Engineering from Jilin University in 2022. He is currently a Master's student supervised by Prof. Bingran Yu at Beijing University of Chemical Technology. His current research interests are focused on the design and synthesis of antibacterial polymers and their application in biomedicine.

hinder their applications.^{25–29} In addition, positively charged polycations can easily absorb negatively charged proteins in the extracellular fluid, which reduces the efficiency *via* which the vectors can complex with nucleic acids.^{30–32} Therefore, how to develop polycationic vectors for safe delivery and efficiently compressing nucleic acids has become a focus of research.

Polycationic vectors mainly refer to polymers with positively charged groups, which rely on electrostatic interaction to complex nucleic acids.^{33–35} Conventional polycationic vectors such as poly(2-*N,N*-dimethylaminoethylmethacrylate) (PDMAEMA)^{36–38} and polyethyleneimine (PEI)^{31,39–41} have already been proven to have outstanding complexation capability due to their high positive charge density, while unignorable cytotoxicity, protein absorption behavior, and high hemolysis rate caused by the positive charges compromise their transfection efficiency and bio-compatibility. As a strategy to overcome these shortcomings, the fabrication of hydroxyl-rich polycationic vectors is suggested.^{42,43} Primarily, researchers mainly modify hydroxyl-rich structures, such as polyethylene glycol, on polycationic vectors.^{44,45} The obtained polycations were found to exhibit negligible cytotoxicity, low protein absorption capability, and good blood compatibility benefited from abundant hydroxyl groups. However, such a post-modification method usually requires two steps to complete, preparation of the polycations and post modification. The whole process is complicated and the yield of the final product is unacceptable. With the deepening of research, the “one-pot” method is gradually becoming a better choice. Instead of post-modification, the one-pot method focuses on the design and preparation of the building blocks, and hydroxyl structures can be obtained from efficient, green, and mild reactions of the building blocks.^{46–48} One example is the amine-epoxide ring-opening reaction, which is based on the reaction of primary amino groups and epoxy groups.^{49,50} With the opening of the epoxy groups, secondary amino groups and hydroxyl groups will occur simultaneously. Positively-charged secondary amino groups can complex with the negatively-charged nucleic acids, and hydroxyl groups provide hydrophilicity and biocompatibility.

In this brief review, we mainly introduce the recent progress in the fabrication of hydroxyl-rich polycations *via* a one-pot

method. The typical one-pot methods for fabricating hydroxyl-rich polycations are presented. In particular, the amine-epoxide ring-opening reaction as a novel and efficient approach has been mainly highlighted. The therapeutic scenarios, including cancer, diabetes mellitus, liver fibrosis, anti-bacterial and wound healing treatments, of hydroxyl-rich branched polycations *via* the one-pot method are subsequently generalized. Hopefully, this review will inspire continuous endeavors in exploring more simple and efficient ways to develop various hydroxyl-rich branched polycations and enriching their biological applications for nucleic acid delivery.

2. One-pot method for fabricating hydroxyl-rich polycations

As an efficient and convenient synthetic strategy, the one-pot method can achieve a multi-step reaction process continuously without intermediate separation, which not only economizes cost, but also guarantees the yield of final product. In this section, many one-pot methods for the preparation of hydroxyl-rich polycations *via* ring-opening reactions of poly(glycidyl methacrylate) (PGMA), Michael addition reactions, and amine-epoxide ring-opening polymerization are generalized. Notably, amine-epoxide ring-opening polymerization is mainly introduced.

2.1 Amino-epoxy group reaction of PGMA

PGMA is usually obtained by atom transfer radical polymerization (ATRP) or reversible addition-fragmentation chain transfer polymerization (RAFT) of glycidyl methacrylate monomers. Our previous work synthesized PGEA or PGED by using ring opening reactions of ethanolamine (EA) or ethylenediamine (ED) with epoxy groups in PGMA, which achieved poly-hydroxylation of PGMA successfully.^{51–55} Inspired by the synthetic methods of PGEA and PGED, many aminated building blocks were introduced into the preparation of hydroxyl-rich polycation carriers, together with EA or ED through ring-opening



Yujie Sun

Yujie Sun received her BSc degree from Yanshan University in 2018. She is currently a doctor student under the supervision of Prof. Fu-Jian Xu and Prof. Bingran Yu at Beijing University of Chemical Technology. Her research interests focus on photo-dynamic antibacterial polymers and cationic polymers for bio-macromolecule delivery.



Qi Yu

Qi Yu received his PhD degree in Materials Science and Engineering from Beijing University of Chemical Technology in 2019, under the supervision of Prof. Fujian Xu. His current research focuses on the design and construction of functionalized cations and their biological applications in gene delivery.



Fig. 1 Strategies for the preparation of hydroxyl-rich polycations for co-ring-opening reactions of PGMA.

reactions to fabricate functionalized PGEA or PGED later. In this part, we summarize the one-pot methods for the fabrication of hydroxyl-rich polycationic vectors (Fig. 1).

For example, Li *et al.* used 3-aminophenyl boronic acid (APBA) and EA reacting with PGMA through the one-pot method to obtain functionalized PGEA, which lays the foundation for further works.⁵⁶ Moreover, synthetic aminated building blocks, tetra-aminophthalocyanine (TAPc-Zn) and aminated tetraphenylethylene (TPE-NH₂), were designed and constructed by Sun *et al.* and Qi *et al.*^{57,58} Then, PGPGEA and TPE-PGEA/TPE were obtained *via* a one-pot method by ring-opening reactions of PGMA, EA with TAPc-Zn and TPE-NH₂, which conferred photodynamic therapy and fluorescent imaging capabilities on the PGPGEA and TPE-PGEA/TPE while delivering the nucleic acids.

Although such a one-pot method based on ring-opening reactions of PGMA could be utilized for the fabrication of

hydroxyl-rich polycationic vectors, the degradability of the polycationic vectors has not been achieved owing to the backbone of PGMA. Therefore, considering the advantages of degradable biomaterials, novel strategies of one-pot methods for preparing hydroxyl-rich polycationic vectors are necessary.

2.2 Michael addition reactions

In order to achieve the degradability of polycations, researchers have made many attempts, including the design of polyamides and polyesters.^{59–61} However, the preparation of these polycations requires stringent reaction conditions, such as anhydrous and oxygen-free environments. Therefore, an easy-to-implement one-step method for the preparation of degradable polycations is urgently needed. Meanwhile, Michael addition reactions have attracted attention and proved to be a promising construction strategy.^{62–65} Specifically, a nucleophilic system, such as amino groups, could react with an electro-



Bingran Yu

Bingran Yu is a professor at Beijing University of Chemical Technology. He obtained his PhD in inorganic chemistry from Lanzhou University in 2013 and then joined Beijing University of Chemical Technology. Dr Yu has contributed significantly to the field of design, construction, and application of drug/gene delivery vectors and novel antibacterial polymers. He has published more than 30 articles in international journals as the corresponding author.



Fu-Jian Xu

Fu-Jian Xu obtained his PhD degree in biomolecular engineering in 2006 from National University of Singapore. He joined Beijing University of Chemical Technology, China as a Professor in 2009. He was a recipient of the National Science Foundation for Distinguished Young Scholars (NSFC, 2013) and Cheung Kong Distinguished Professor (Ministry of Education of China, 2014). His research interests focus on functional biomacromolecules.

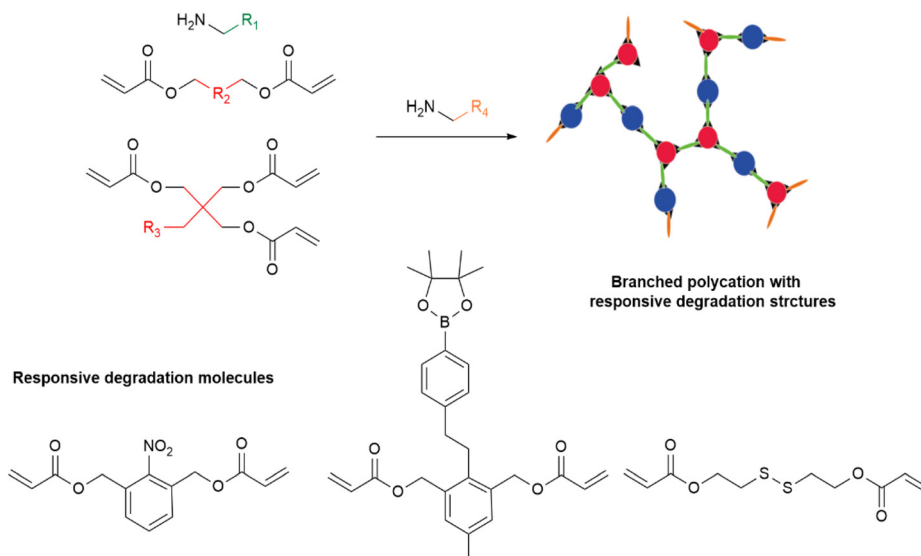


Fig. 2 One-pot Michael addition reaction for the construction of degradable branched hydroxyl-rich polycations.

philic system, like vinyl groups, *via* a one-pot conjugate addition reaction under relaxed conditions.^{66,67} Polycations with flexible structures, including linear, branched, and three-dimensional structures, could be fabricated by Michael addition reactions. In addition, degradable groups are easy to obtain by Michael addition reactions to realize the degradability of the polycations (Fig. 2).

Lu *et al.* synthesized a branched poly(β -amino ester) (PBAE) with backbone-embedded phenylboronic acid (PBA) *via* an amino-vinyl Michael addition reaction, which made the polycation degradable when driven by an oxidizing environment.⁶⁸ Moreover, through an A2 + B3 + C3-type Michael addition reaction, Duan *et al.* synthesized a hydroxyl-rich polycation BPAE-NB by the use of 4-amino-1-butanol, trimethylolpropane triacrylate, and 2-nitro-1,3-phenylene bis(methylene) diacrylate. Owing to the introduction of 2-nitrobenzene moieties, the BPAE-NB backbone demonstrated good degradability when exposed to UV light, which promoted intracellular gene release and diminished the toxicity of the materials in the post-transfection state.⁶⁹ Rui *et al.* used the bisulfide bond-containing monomer 2,2-disulfanediylbis(ethane-2,1-diyl) diacrylate (BR6) to create a series of reductively degradable quaternary polycations (rBEAQs) using a facile one-pot Michael addition reaction, and impressive nucleic acid binding affinity, cellular uptake efficiency, release performance, and functional nucleic acid delivery were proven.⁷⁰

However, the Michael addition process does not yield hydroxyl groups, and hence requires the utilization of building blocks that contain hydroxyl groups, which severely limits the variety of building blocks.

In addition, many chemically active groups, such as hydroxyl, amino, and vinyl groups, are often involved in Michael addition, which means that side reactions and the production of by-products can easily occur, making the whole

process inefficient. Therefore, how to expand the types of building blocks and improve the reaction efficiency has become a new problem.

2.3 Amine-epoxide ring-opening polymerization

Inspired by the Michael addition reaction, a one-pot method of amine-epoxide ring-opening polymerization was proposed.⁷¹ While retaining the amine-epoxy ring-opening reaction, this strategy abandons the polymerizations based on ATRP or RAFT, and designs building blocks containing a variety of responsive groups to achieve the degradability of the hydroxyl-rich polycation for nucleic acid delivery.⁷² Moreover, the terminal groups of the polycations are enriched, which is beneficial to the decoration of the vector and further improves the performance of the vectors.⁷³

2.3.1 Polymerization method. The amine-epoxide ring-opening reaction is characterized as a straightforward and gentle chemical transformation. Theoretically, two poly-functional building blocks, poly-amine and poly-epoxide molecules, are used to prepare polycations through step growth, where the epoxy rings undergo nucleophilic assault by the amino groups, resulting in the cleavage of the epoxy groups and the formation of secondary amino groups and hydroxyl groups simultaneously in one step (Fig. 3).^{74–76} Positively-charged secondary amino groups can complex negatively-charged nucleic acids, and the hydroxyl groups provide hydrophilicity and biocompatibility. In comparison with ring-opening reactions of PGMA and Michael addition reactions, amine-epoxide ring-opening polymerization can easily achieve the degradation of polycations through the introduction of responsive moieties in the building blocks, the choice of building blocks is wider, and the reaction is more efficient. In addition, this polymerization does not rely on harsh reaction conditions, and the reaction process is mild and green.

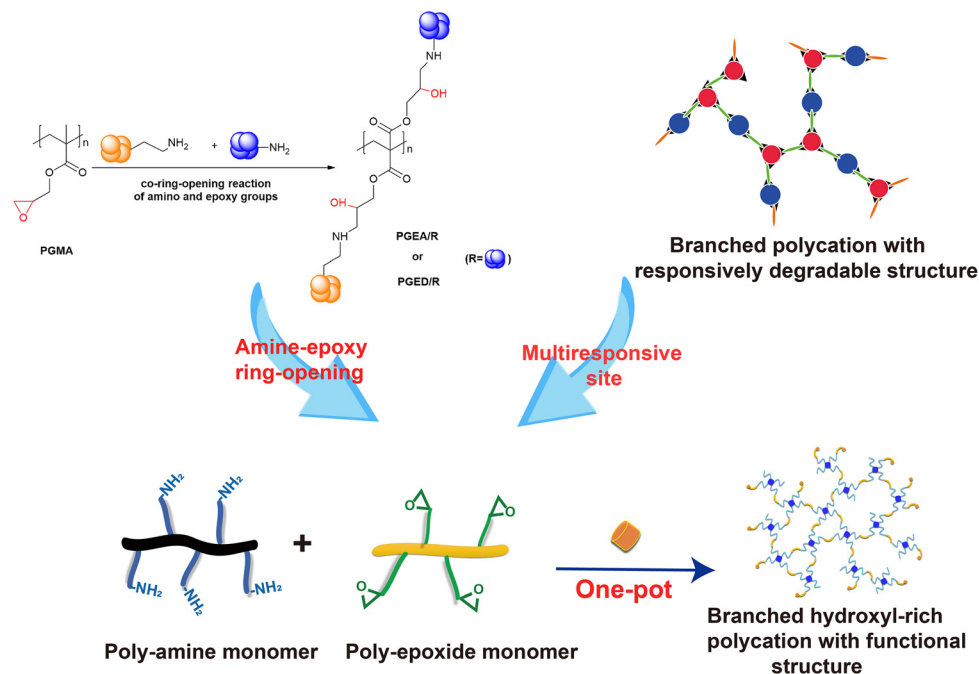


Fig. 3 One-pot poly-amine and poly-epoxy ring-opening reaction for the construction of functionalized branched hydroxyl-rich polycations.

The preparation of polycations through amine-epoxide ring-opening polymerization has been reported. Zeng *et al.* used the conventional aminoglycoside antibiotic gentamicin and poly (ethylene glycol) diglycidyl containing double epoxy groups *via* an amino-epoxy ring-opening polymerization to synthesize a branched polycation GPEG in one pot. ED as a capping agent was added when the polymerization was finished to consume the remaining epoxy groups.⁷⁷ Moreover, the branched polycation TE was synthesized by the amine-epoxide ring-opening polymerization of ED and triglycidyl isocyanurate (TGIC) including three epoxy groups, which had good nucleic acid condensation ability and low cytotoxicity. Compared with the branched PEI, the optimal transfection efficiency of TE in the presence of serum was significantly higher.⁷¹

2.3.2 Introduction of a responsive moiety. In consideration of the degradability of vectors, polycations need structures that are responsive to stimuli factors such as chemical factors (redox, pH, gases, *etc.*),^{30,78,79} biological factors (enzymes, receptors, *etc.*)^{16,80,81} or physical factors (temperature, magnetic, *etc.*),^{82–84} which will bring degradable capabilities to the polycations. Therefore, responsive moieties should be introduced into poly-amine and poly-epoxide building blocks for amine-epoxide ring-opening polymerization. We have developed several novel poly-amine or poly-epoxide building blocks involving responsive moieties, such as disulfide groups or *ortho* ester groups, for the preparation of degradable hydroxyl-rich branched polycations sensitive to reductive or acidic stimuli (Fig. 4). The introduction of responsive moieties for constructing degradable polycations based on molecular design makes polycations easier to apply in various physiological conditions.

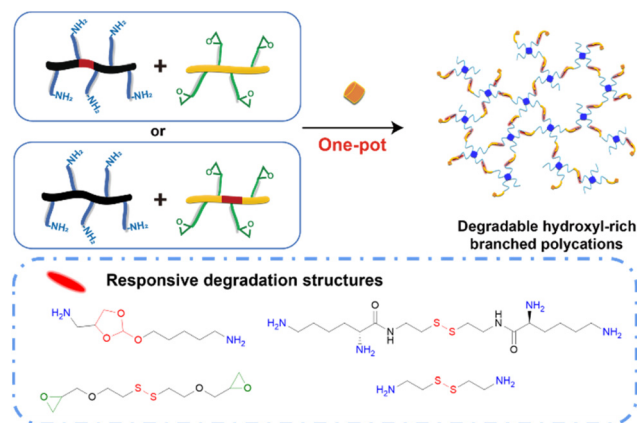


Fig. 4 One-pot poly-amine and poly-epoxy ring-opening reaction for the construction of degradable branched hydroxyl-rich polycations.

The progress of the typical responsive (redox-responsive and acid-responsive) degradable polycations for nucleic acid delivery will be introduced in detail. Huang *et al.* synthesized hydroxyl-rich branched polycation SS-HPs through amine-epoxide ring-opening polymerization involving disulfide-bonded HDDE and three polyaminoglycosides, which exhibited reductive degradability when used for gene therapy in glioma treatment.⁸⁵ Chen *et al.* used lysine-decorated cystamine (CA) and TGIC to fabricate a hydroxyl-rich branched polycation *via* amine-epoxide ring-opening polymerization, which demonstrated reductive degradability when used for osteosarcoma therapy.⁸⁶ Qi *et al.* used a synthetic poly-amine molecule involving an *ortho* ester linkage and TGIC to fabricate an acid-

moieties can be flexibly introduced into the building blocks through molecular design to achieve the degradability of the polycations, and the abundant active groups at the ends of the polycation provide a guarantee for further functionalization of the vector. Next, we summarize the different application scenarios of hydroxyl-rich polycations prepared by amine-epoxide ring-opening polymerization as nucleic acid delivery vectors.

3. Various scenarios for hydroxy-rich polycation application

With the continuous development of research, the application of nucleic acid delivery for disease treatment has gained widespread attention and shown good therapeutic effects. Based on the hydroxyl-rich polycations constructed by a one-pot method, researchers have innovatively designed several nucleic acid delivery systems with multiple functions against different application scenarios, such as malignant tumors, diabetes, and epidermal wounds. In this chapter, recent progress regarding the application of hydroxyl-rich polycations fabricated by a one-pot method is described.

As cancer is a fatal disease, researchers have been exploring different treatment methods for malignant tumors and gene therapy has been proven to be an effective strategy.^{90–94} Degradable hydroxy-rich polycations are widely used in nucleic acid delivery systems for tumor therapy due to their multiple responsiveness to the tumor microenvironment (TME).^{95–97} The TME is a special physiological environment required for tumor cell growth, and its salient features are higher glutathione (GSH) expression levels^{98,99} and a more acidic intracellular environment (pH: 5.0–5.5)^{100–102} compared with normal cell locations. As previously introduced, hydroxyl-rich polycations can be introduced with various responsive moieties *via* design of the building blocks, so responsive degradability can be successfully achieved influenced by the physiological microenvironment of tumor cells.^{103–105} The degradability of polycations is conducive to realizing the release of nucleic acids at the site of action, and greatly reduces the toxicity of their side effects on the organism.

Reduction and acid-responsive degradable nucleic acid delivery systems have been mainly constructed. Huang *et al.* prepared several hydroxyl-rich branched polycations (SS-HPs) by amine-epoxide ring-opening reaction of disulfide-containing bis-epoxy compounds with aminoglycosides containing multiple amino groups (Fig. 6a).⁸⁵ As verified by agarose gel electrophoresis assays, the SS-HPs showed an excellent degradation profile, which is attributed to the introduction of disulfide bonds (Fig. 6b). As shown in Fig. 6c, the SS-HPs demonstrated higher gene transfection efficiency in different cell lines than the un-disulfide HPs and branched PEI, which was mainly attributed to the reduction-responsive disulfide bonds in the SS-HP. Because of their reduction responsive

Post modification structures

The reaction scheme shows the synthesis of post-modified hydroxyl-rich branched polycations. The starting materials are a linear polycation with four terminal amino groups (NH₂) and a diol with two terminal hydroxyl groups (OH). The reaction proceeds in two steps: i) reaction with a pink triangle (representing a cross-linker) and ii) reaction with an orange hexagon (representing a cross-linker). The final product is a branched polycation with multiple terminal amino groups and hydroxyl groups, labeled "Post modified hydroxyl-rich branched polycations".

This journal is © The Royal Society of Chemistry 2024

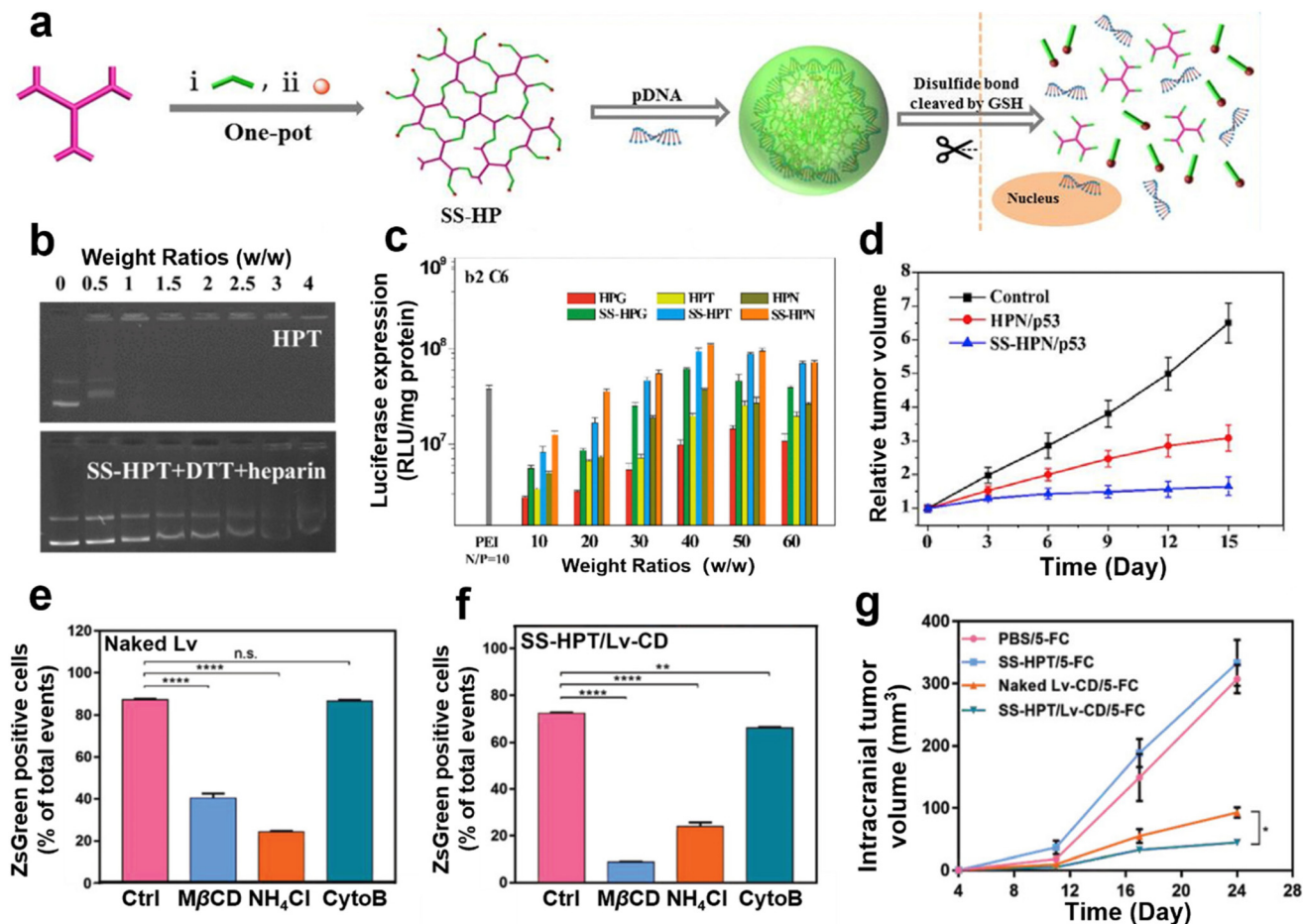


Fig. 6 (a) Schematic of the preparation of multifunctional aminoglycoside degradable branched polycations (HPs). (b) Electrophoretic mobility of HPs/pDNA in a reduced environment with and without DTT simulation, exemplified by SS-HPN. (c) Transfection performance assay of HPs/pDNA in C6 cells. (d) Tumor volume curves in tumor-bearing mice after different groups of treatment. Reprinted with permission from ref. 85. Copyright 2016, Elsevier. (e) Quantitative FACS analysis of naked Lv and (f) SS-HPT/Lv-20 mediated reporter gene ZsGreen expression in C6 cells after treatment with different endocytosis inhibitors. (g) Tumor volume curves of tumor-bearing rats during different treatment groups. Reprinted with permission from ref. 106. Copyright 2019, WILEY-VCH Verlag GmbH & Co. KGaA, Weinheim.

degradability, SS-HPs can easily realize the release of nucleic acids. Subsequently, the classical anticancer gene p53 was further selected as the therapeutic nucleic acid to evaluate the *in vivo* antitumor performance of the SS-HPs (Fig. 6d). The results showed that the SS-HPs have the ability to efficiently deliver the p53 gene, and due to the reduction-sensitive property of the disulfide bonds, the p53 gene was effectively released into the cytoplasm. SS-HPs/p53 complexes significantly inhibited the growth of C6 glioma and did not produce significant systemic toxicity during treatment.

Moreover, an SS-HP/Lv-CD system was constructed by coating CD suicide gene-engineered lentivirus (Lv-CD) with SS-HP for use against glioma by Shao *et al.*,¹⁰⁶ which showed good biocompatibility. Cellular uptake assays by using C6 glioma cells showed that SS-HP/Lv could enter into the cells efficiently and release Lv particles in a wider range of pathways than exposed Lv (Fig. 6e and f). In addition, SS-HP/Lv demonstrated significantly higher transfection efficiency,

which led to the expression of more CD proteins, further enabling SS-HPT/Lv-CD to significantly increase the sensitivity of glioma cells to 5-fluorocytosine (5-FC) (Fig. 6g). Furthermore, by evaluating the tumor volume and survival time of mice after treatment for glioma *in situ*, the results demonstrated that SS-HP/Lv-CD with 5-FC most effectively inhibited tumor growth and prolonged the survival time of the mice.

Encouraged by the above work, more versatile therapeutic nucleic acid delivery systems were constructed to achieve efficient treatment of cancer. Chen *et al.*⁸⁶ modified biomolecule lysine with cystamine to synthesize SS-Lys, a polyamine compound with a disulfide bond. Following amine-epoxide ring-opening reaction of SS-Lys with TGIC, a reduction-responsive degradable hydroxyl-rich branched polycation named SS-LHP was fabricated. A branched polylysine gene vector (CC-HP) without disulfide bonds was used as a control. *pKillerRed* gene (*pKR*) was delivered by the polycation

to estimate antitumor capability of the nucleic acid delivery system. *KillerRed* protein^{42,107} is a red fluorescent protein that can generate singlet oxygen ($^1\text{O}_2$) under visible light irradiation, which further enables the effects of PDT (Fig. 7a). Gel electrophoresis and atomic force microscopy were used to assess the degradation of the SS-LHP/pDNA complexes in a reducing environment. Compared to the non-degradable polycation CC-HP, SS-LHP was unable to encapsulate pDNA and SS-HP/pDNA did not exhibit a homogeneous and regular nanostructure in a reductive environment because of the existence of disulfide bonds. Owing to the degradability of polycations, the accelerated release of pDNA and good transfection efficiency were proved. Finally, SS-LHP carrying *pKR* plasmid

was used for *in vitro* photodynamic anticancer evaluation. SS-HP/pKR transfected C6 glioma cells successfully expressed *KillerRed* protein and promoted apoptosis of tumor cells after light exposure (Fig. 7b).

Osteosarcoma (OS) is the most common primary malignant bone tumor and is highly metastatic. Expression of miR-22 is significantly downregulated during OS development, making miR-22 a promising therapeutic target for OS. A novel class of reduction-sensitive hydroxyl-rich polycationic vector (TC) was constructed by a one-pot method through anime-epoxide ring-opening reaction of the epoxy group of cystamine with TGIC to deliver miR-22 and miR-30d, respectively, for the treatment of OS (Fig. 7c).¹⁰⁸ Due to its abundant secondary amino and

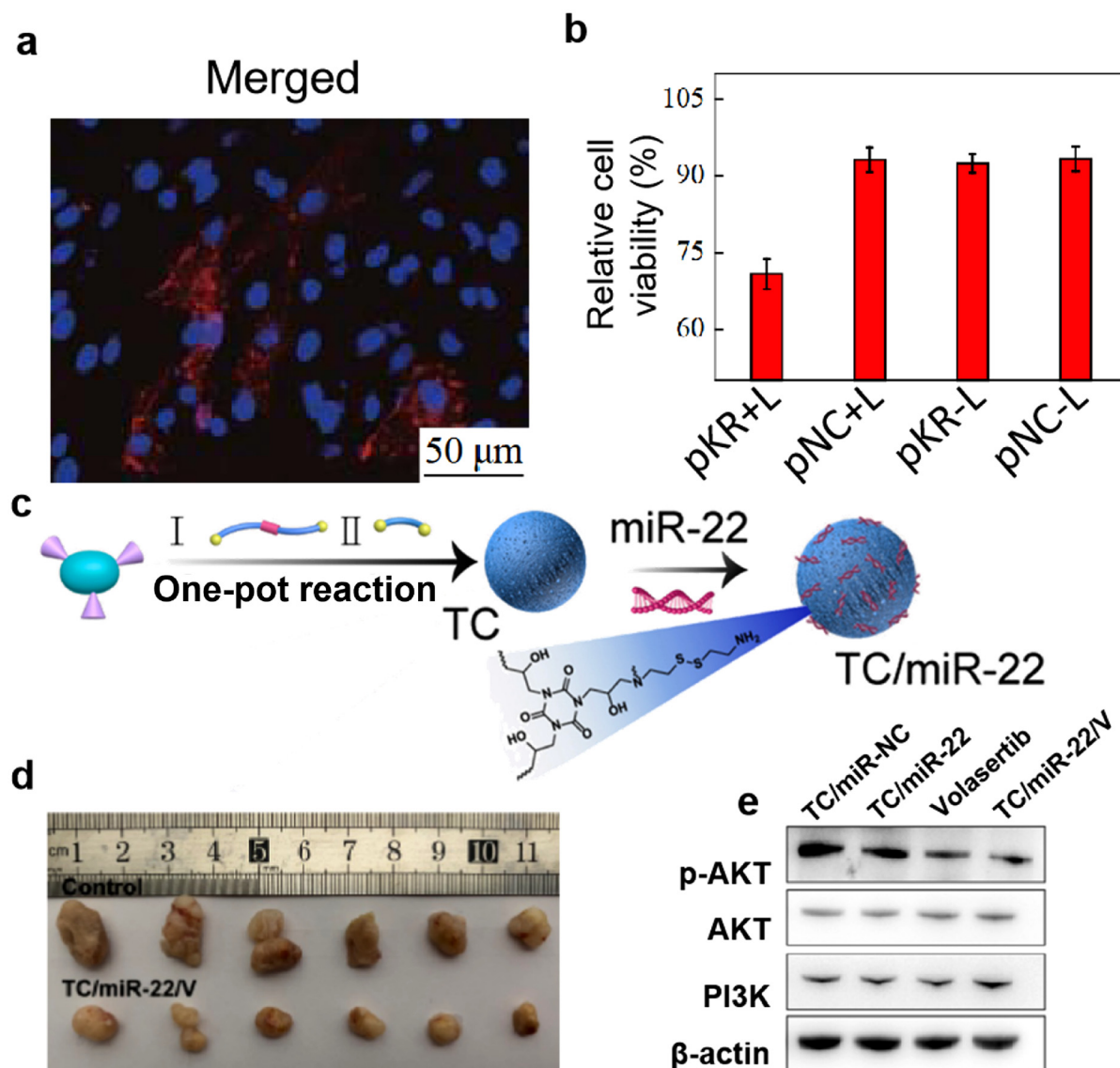


Fig. 7 (a) Representative images of SS-HP-mediated *KillerRed* expression in C6 cells. (b) Relative cellular activity of SS-HP/pKR and SS-HP/pNC in C6 cells in the presence or absence of light. Reprinted with permission from ref. 86. Copyright 2019, Science Press. (c) Preparation process of reductively reactive polycationic TC with TC/miR-22. (d) Tumor images of drug-resistant PDX model mice after treatment with different treatment groups. (e) Relative expression of p-AKT protein in Saos-2 cells with different treatment groups. Reprinted with permission from ref. 108. Copyright 2023, KeAi Publishing.

hydroxyl groups, TC has good nucleic acid condensation ability and biocompatibility. The hydrated particle size of TC particles increased when treated with DTT, indicating that TC had a good reduction-responsive degradability. Moreover, TC showed good transfection performance in the OS cell line (Saos-2) and biocompatibility in the osteoblast cell line (MC3T3-E1). By inhibiting the PI3K/AKT pathway, the use of TC-mediated miR-22 transfection of Saos-2 cells achieved good anti-tumor effects *in vitro*, not only inhibiting the proliferation and migration of Saos-2 cells, but also inducing the promotion of apoptosis of Saos-2 cells. Combined with the anticancer drug Volasertib treatment, TC/miR-22 complexes were able to enhance the efficacy of treatment. *In vivo* results confirmed that TC/miR-22 achieved excellent anti-tumor effects and effectively reduced lung metastasis of tumor cells in both *in situ* OS and patient-derived xenograft (PDX) models. To investigate the improvement of the efficacy of chemotherapy-resistant OS, a drug-resistant PDX model was established. In the treatment of TC/miR-22/Volasertib, increasing the dosage of miRNA was not only able to inhibit the growth of drug-resistant tumors (Fig. 7d), but also did not produce significant cytotoxicity. At the end of treatment, in order to investigate the mechanism of action of TC/miR-22, western blot assay was used to analyze the regulation of AKT protein phosphorylation by TC/miR-22 in the PI3K/AKT pathway of Saos-2 cells, and it was found that both TC/miR-22 and Volasertib could effectively inhibit the expression of AKT protein, thus affecting the PI3K/AKT pathway (Fig. 7e).

3.2 Targeted nucleic acid delivery systems for disease therapy

3.2.1 Cancer therapy. Considering the microenvironment of the tumor site, some chemical structures sensitive to the stimulated environment were introduced into the design of building blocks for the preparation of degradable hydroxyl-rich branched polycationic nucleic acid delivery systems and combined with chemotherapeutic drugs to further enhance the therapeutic effect on tumors. Qi *et al.* used acid-sensitive *ortho* ester diamine and TGIC preparing ARPs *via* amine-epoxide ring-opening reaction, then fluorinated alkyl chains were linked at the end of the ARP to obtain ARP-F with one-pot decoration (Fig. 8a).⁸⁷ GPC analysis and ¹H NMR spectra proved the impressive acid-responsive degradability of ARP-F (Fig. 8b). Next, pCas9-surv, a plasmid designed to knock out the apoptosis suppressor *survivin* gene, was introduced. pCas9-surv, mediated by ARP-F, showed excellent tumor suppressive properties *in vitro* and *in vivo*. In addition, ARP-F/pCas9-surv in combination with temozolomide (TMZ), a widely used cancer-related drug, was able to increase the sensitivity of cancer cells to anticancer drugs, further enhancing the tumor suppressor activity (Fig. 8c).

In the process of gene therapy, how to make nucleic acids reach target cells without interfering with other normal cell physiological behaviors^{109–111} is a key scientific issue. Therefore, in addition to being degradable to ensure its biological safety, the nucleic acid delivery vector also needs to have the ability to target the cells of the lesion. As for the

method of introducing targeting molecules into vectors, the aforementioned one-pot decoration method is widely used due to its simplicity of implementation and high efficiency of reaction.

According to previous research, the boronic acid group has affinity to the nucleus, showing a good tumor cell uptake efficiency and nucleus targeting ability. Shao *et al.* decorated a PBA moiety on a hydroxyl-rich branched polycation (SS-HPT) to get a PBA-functionalized branched polyaminoglycoside (SS-HPT-P).¹¹² In the fluorescent imaging of tumor model animals, it was observed that the SS-HPT-P/pCas9-surv complexes were rapidly enriched into tumors due to the good targeting ability of PBA. The typical pCas9-surv delivery mediated by SS-HPT-P shows good gene editing performance. Due to the targeting ability of PBA, SS-HPT-P/pCas9-surv is significantly effective in inhibiting the proliferation and migration of lung tumor cells *in vitro* and *in vivo* when combined with TMZ.

In addition, lactose, as a common natural disaccharide, composed of one glucose molecule and one galactose molecule, is able to specifically bind to the asialoglycoprotein receptor (ASGPr) on the surface of hepatic carcinoma cells (HCCs). Qi *et al.*⁸⁹ prepared a reduced-responsive hydroxyl-rich branched polycationic vector named LBP *via* amine-epoxide ring-opening polymerization of aminated lactose with TGIC for the treatment of *in situ* hepatic carcinoma (Fig. 8d). LBP demonstrated significant enrichment at the liver site *via in vivo* imaging characterization (Fig. 8e). Subsequently, *in vitro* and *in vivo* anti-tumor experiments demonstrated that the LBP-mediated CRISPR/Cas9 system (LBP/pCas9-*survivin*) not only effectively inhibits the proliferation of HCCs, but also accelerates cell apoptosis through effective gene editing. LBP/pCas9-*survivin* gene editing also enhances the anti-tumor effect and sensitizes the anticancer drugs (Fig. 8f).

3.2.2 Others. Except for cancer therapy, we also designed several targeting nucleic acid delivery systems against other diseases such as diabetic mellitus and liver fibrosis. Diabetes mellitus is a disease of the metabolism, featuring persistent hyperglycemia due to insufficient insulin secretion or insulin resistance. Conventional therapeutic strategies,^{113,114} like insulin injection, are prone to causing hypoglycemia and lipohypertrophy due to inflexible dose control and improper injection technique. Therefore, new treatment methods for diabetes are urgently required.¹¹⁵ Based on the cell replacement of non-beta cells to beta-like cells, reprogrammed hepatic parenchymal cells by ectopic expression of specific transcription factors such as Ngn3, Pdx1, Mafa, and Neurod1 have been proposed as a novel strategy to treat diabetes mellitus. In addition, phenylboronic acid (PBA) and its derivatives have been proven to demonstrate strong affinity with sialic acid (SA), which is highly expressed in hepatic parenchymal cells. Therefore, Pan and Shao *et al.*¹¹⁶ introduced a PBA moiety into a hydroxyl-rich branched polycation (SS-HPT-P) *via* one-pot decoration (Fig. 9a). The therapeutic plasmid pNPMN-PBase encoding Ngn3, Pdx1, Mafa, and Neurod1 expressed factors were constructed at the same time. In *in vitro* and *in vivo* experiments, SS-HPT-P/pNPMN-PBase complexes showed good hepatocyte

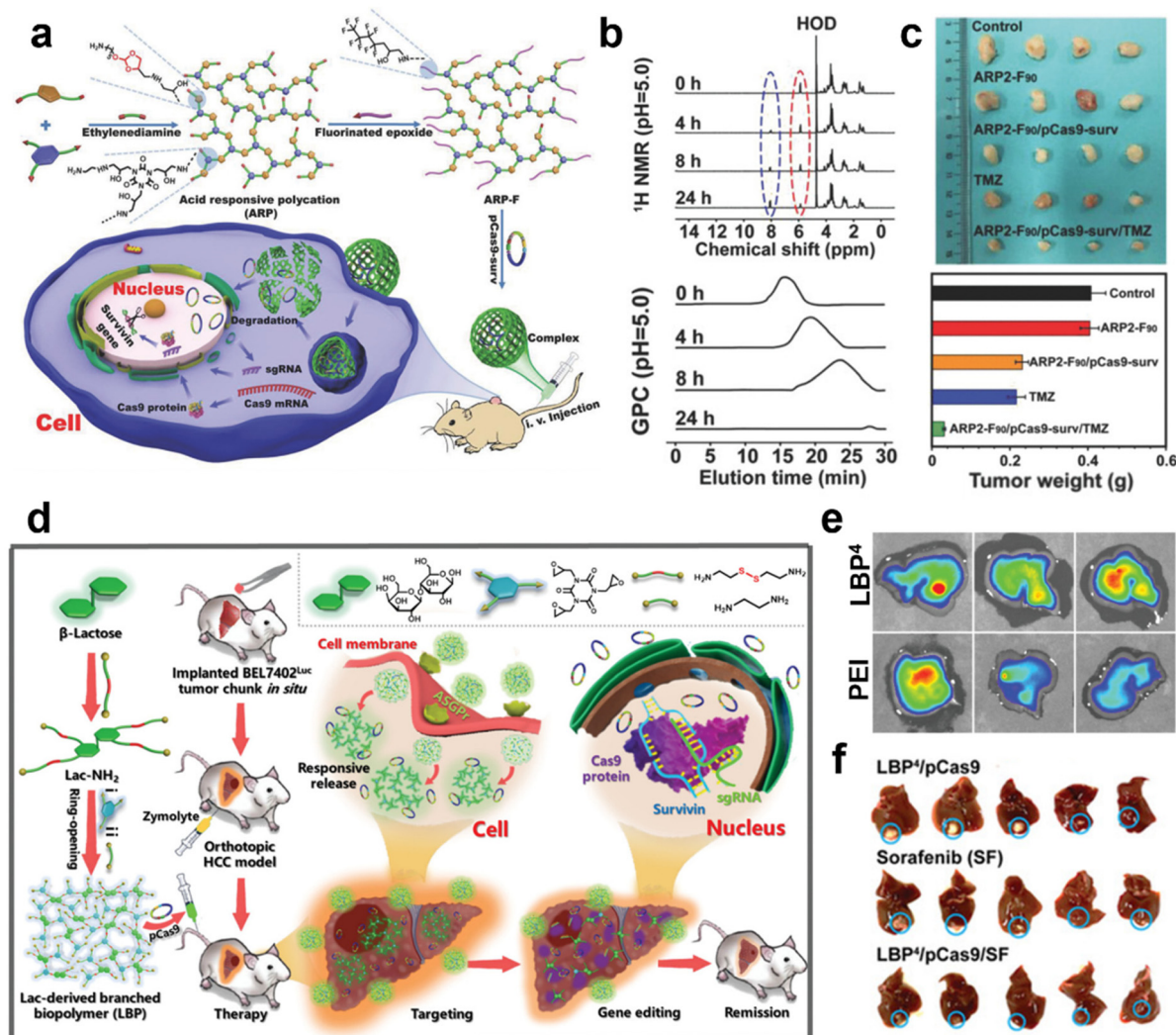


Fig. 8 (a) Preparation of branched polycationic ARP-F with its *in vivo* delivery of nucleic acids with delivery of pCas9-surv for gene editing. (b) Degradation properties of ARP-F *in vitro* under simulated acidic conditions. (c) *In vivo* assay of ARP-F/pCas9-surv against mouse tumors. Reprinted with permission from ref. 87. Copyright 2018, WILEY-VCH Verlag GmbH & Co. KGaA, Weinheim. (d) Preparation of a lactose-derived branched biopolymer (LBP) and schematic representation of the delivery and gene editing process of pCas9-survivin for the treatment of HCC *in situ*. (e) Representative fluorescence images of the liver in the LBP-treated group. (f) Representative liver images for each treatment group. Reprinted with permission from ref. 89. Copyright 2020, WILEY-VCH Verlag GmbH & Co. KGaA, Weinheim.

targeting properties (Fig. 9b), which induced insulin-negative hepatocyte cells to reprogram into insulin-positive beta-like cells by tandem expression of four specific transcription factors related to pancreatic development or insulin expression (Fig. 9c and d). This nucleic acid delivery system provided an effective strategy for cell replacement therapy for diabetes.

Liver fibrosis is a repair response to liver injury and is a key aspect in several liver diseases. Recent research studies revealed that the pro-fibrotic cytokine transforming growth factor (TGFβ1) is involved in almost all critical dimensions of liver fibrosis development. Therefore, down-regulating TGFβ1 expression has the potential to treat liver fibrosis. Wu *et al.*⁷³ constructed a targeting hydroxyl-rich branched polycationic vector (SS-HPVA) based on the one-pot decoration of branched polyaminoglycoside (SS-HPT). As a HSC targeting molecule, VA

was introduced to the SS-HPT. Multispectral photoacoustic tomography and *in vivo* fluorescent imaging verified that the SS-HPVA/pshRNA-TGFβ1 complexes demonstrated superior liver targeting ability, especially in the hepatic fibrosis model (Fig. 9e). On this basis, it was demonstrated that this system could exhibit good therapeutic effects on liver fibrosis by down-regulating TGFβ1 expression through non-invasive assays and histopathological analysis (Fig. 9f). Therefore, this gene silencing targeting delivery system is expected to provide a new strategy for the clinical treatment of liver fibrosis.

3.3 Nucleic acid delivery systems for antibacterials and wound healing

Growth factor delivery has attracted high attention for the treatment of severe skin wounds and the promotion of wound

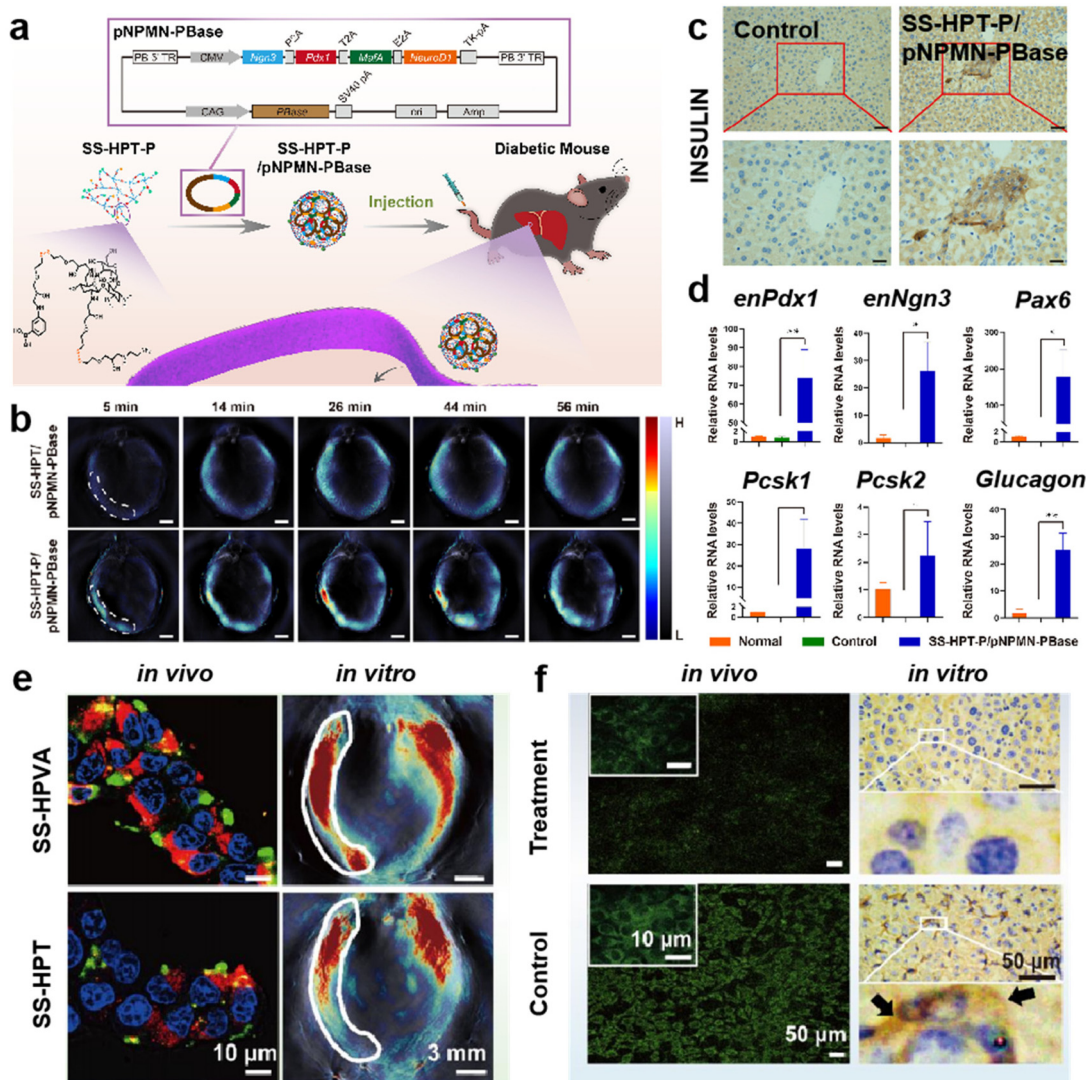


Fig. 9 (a) Schematic illustration of the preparation process of hydroxyl-rich branched polycation SS-HPT-P, and SS-HPT-P/pNPMN-PBase targeting the liver of diabetic mice for gene reprogramming therapy. (b) Representative MSOT cross-sectional images of mice at different times. (c) Representative images of immunohistochemical staining of liver insulin. The images are observed upstream at 1000 times magnification and downstream at 2000 times magnification. (d) Quantitative real-time detection of mRNA expression of pancreatic transcription factors and insulin expression/secretion-related genes after treatment in different groups. Reprinted with permission from ref. 116. Copyright 2022, Elsevier. (e) Endocytosis fluorescence imaging of SS-HPVA/pDNA with the SS-HPT/pDNA complex in HSCs (left) and representative cross-sectional photoacoustic images of liver fibrosis model mice after tail vein injection of SS-HPT/pshRNA-TGFβ1 or SS-HPVA/pshRNA-TGFβ1. (f) Immunofluorescence imaging of TGFβ1 cells after different treatments (left) and immunohistochemical sections (right). Reprinted with permission from ref. 73. Copyright 2023, Elsevier.

healing.^{117,118} Thus, Chen *et al.* developed a hydroxyl-rich branched polycation (SKP) *via* amine-epoxide polymerization of disulfide bond-modified lysine (SK) and TGIC.¹¹⁹ A transfection assay in the human umbilical vein endothelial cell line (HUVEC) indicated that SKP demonstrated good transfection efficiency compared with PEI. When vascular endothelial growth factor (VEGF) plasmid was delivered by SKP, it was further found that the SKP/pVEGF complexes successfully increase cell proliferation, and a western blot assay detected higher VEGF expression compared with the control group. Moreover, the SKP/pVEGF complexes could be coated on a pre-

treated clinical stent to produce a pVEGF-eluting stent (S-SKP/pVEGF). The *in vivo* re-endothelialization and anti-restenosis performance of S-SKP/pVEGF were evaluated *via* implanting stents into rabbit abdominal aortas.

Hydroxyl-rich branched polycations have also achieved antimicrobial activities and wound healing promotion. Aminoglycosides and their derivatives are active antimicrobial agents. A series of hydroxyl-rich reduction-responsive polycations (SS-HPG, SS-HPT and SS-HPN) synthesized based on aminoglycosides, including gentamicin, tobramycin, and neomycin, have been reported.⁸⁵ Moreover, an Eosin Y (EY)-based

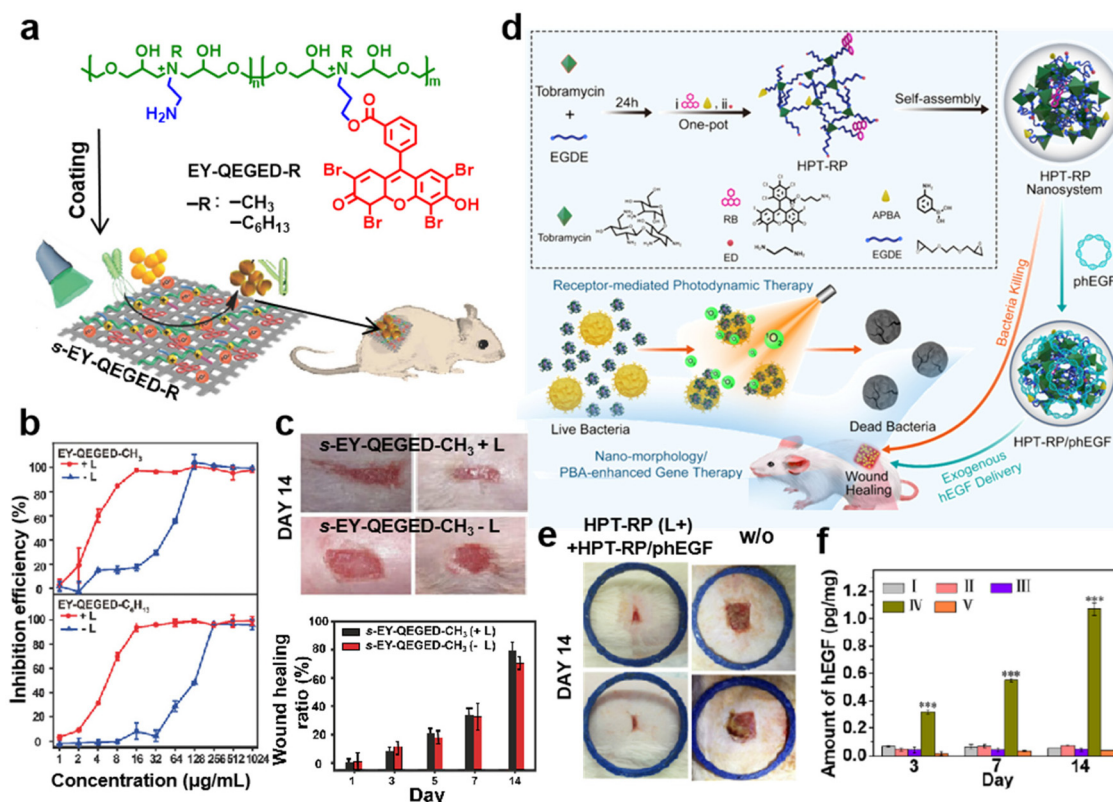


Fig. 10 (a) Construction of EY-based antimicrobial polycations (EY-QEGED-R, R = CH₃ or C₆H₁₃) by amino-epoxy ring-opening reaction and their antimicrobial properties after light exposure for wound dressings. (b) The inhibition effect of the antimicrobial polycations on *S. aureus* in the presence or absence of light. (c) Representative images and ratio of wound healing of a rat wound model on day 1 and day 14 after different groups of treatments. Reprinted with permission from ref. 120. Copyright 2018, WILEY-VCH Verlag GmbH & Co. KGaA, Weinheim. (d) Schematic of the preparation of HPT-RP/phEGF. (e) Bacteriostatic effect of HPT, HPT-R and HPT-RP in the presence or absence of light in the case of *S. aureus*. (f) Representative pictures of the rat infected-skin-defect model on day 14 after different groups of treatments. (g) Quantification of the expression levels of the excised tissues around the wounds at day 3, 7 and 14 after treatment in different groups. Group I–V are: Control, HPT-RP(L-), HPT-RP(L+), HPT-RP(L+) + HPT-RP/phEGF and w/o. Reprinted with permission from ref. 88. Copyright 2021, Elsevier.

antibacterial polycation (EY-QEGED-R, R = CH₃ or C₆H₁₃) with versatile types of functional components including quaternary ammonium, photosensitizer, primary amino, and hydroxyl species was readily synthesized by Zhu *et al.* based on amine-epoxide ring-opening reactions (Fig. 10a).¹²⁰ A remarkable synergistic antibacterial activity of the polycation owing to the combined photodynamic and quaternary ammonium antibacterial effects was proven (Fig. 10b). EY-QEGED-R also could be readily coated on nonwoven fabrics *via* an adhesive layer of polydopamine demonstrating antifouling capability against dead bacteria. Notably, EY-QEGED-R more significantly inhibited wound inflammation and promoted wound healing on an infected rat model under light conditions (Fig. 10c).

Therefore, it is necessary to develop new delivery systems based on hydroxyl-rich branched polycations with a photosensitizer that have superiority for antibacterial, wound healing promotion and delivery performances. One bifunctional hydroxyl-rich branched polycation (HPT-RP) was constructed through the one-pot dual conjugation of a photosensitizer and bacterium/cell-affinity phenylboronic acid onto hyperbranched polyaminoglycosides, achieving antimicrobial properties while

realizing its application to encoding epidermal growth factor (EGF) plasmid delivery (Fig. 10d).⁸⁸ HPT-RP realized high antibacterial ability owing to the receptor-mediated photodynamic therapy, and impressive gene transfection efficiency because of phenylboronic acid conjugation. In a rat infected-skin-defect model, HPT-RP exhibited the remarkable performance of anti-infective/EGF-delivery and accelerated healing (Fig. 10e). More importantly, a new insight into rat-EGF production revealed the underlying mechanism of extraneous EGF delivery and endogenous EGF production (Fig. 10f).

4. Conclusions

Owing to their convenient fabrication process and flexible designability, one-pot methods have shown promising potential in the construction of hydroxyl-rich polycationic vectors for nucleic acid delivery. In this brief review, we mainly summarized three one-pot methods for the synthesis of hydroxyl-rich polycations for nucleic acid delivery: co-ring-opening reactions of PGMA, Michael addition reactions, and amine-epoxide ring-

opening polymerization. Notably, amine-epoxide ring-opening polymerization as a novel approach was mainly introduced. In addition, various therapeutic scenarios used by multifunctional hydroxyl-rich polycation applications, including cancer, diabetes mellitus, liver fibrosis, antibacterial and wound healing treatments, were also generalized. We believe that this review will shed light on the design of hydroxyl-rich polycations and expand the biological applications of cationic gene carriers in the field of nucleic acid delivery.

Author contributions

Mengrui Su and Zichen Hu performed the literature survey and edited the original draft; Yu Qi edited the original manuscript and made original draft revision; Bingran Yu and Fujian Xu conceived the theme of this review, made the original draft revision, and supervised the work.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by the National Key R&D Program of China (grant number 2021YFB3800900), National Natural Science Foundation of China (grant numbers 52073013, 22122501, 52221006, and 22005019), Beijing Natural Science Foundation (grant number 7222011), Beijing Outstanding Young Scientist Program (BJJWZYJH01201910010024), and Beijing Postdoctoral Research Foundation (2022-6ZZ-113).

References

- 1 X. M. Cai, R. Dou, C. Guo, J. R. Tang, X. J. Li, J. Chen and J. Y. Zhang, *Pharmaceutics*, 2023, **15**, 1502.
- 2 S. A. Dilliard and D. J. Siegwart, *Nat. Rev. Mater.*, 2023, **8**, 282–300.
- 3 A. Gupta, J. L. Andresen, R. S. Manan and R. Langer, *Adv. Drug Delivery Rev.*, 2021, **178**, 113834.
- 4 Z. G. Lu, J. Shen, J. Yang, J. W. Wang, R. C. Zhao, T. L. Zhang, J. Guo and X. Zhang, *Signal Transduction Targeted Ther.*, 2023, **8**, 39.
- 5 A. S. Piotrowski-Daspi, A. C. Kauffman, L. G. Bracaglia and W. M. Saltzman, *Adv. Drug Delivery Rev.*, 2020, **156**, 119–132.
- 6 Y. Tian, M. V. Tirrell and J. L. LaBelle, *Adv. Healthcare Mater.*, 2022, **11**, 2102600.
- 7 Y. Y. Wang, P. K. Shahi, R. S. Xie, H. L. Zhang, A. A. Abdeen, N. Yodsanit, Z. Q. Ma, K. Saha, B. R. Pattnaik and S. Q. Gong, *J. Controlled Release*, 2020, **324**, 194–203.
- 8 K. Isgrig, D. S. McDougald, J. L. Zhu, H. J. Wang, J. Bennett and W. W. Chien, *Nat. Commun.*, 2019, **10**, 427.
- 9 Y. Kim, D. Y. Lee, J. U. Choi, J. S. Park, S. M. L. ee, C. H. Kang and C. H. Park, *Sci. Rep.*, 2023, **13**, 12365.
- 10 H. E. Maunder, J. Wright, B. R. Kolli, C. R. Vieira, T. T. Mkandawire, S. Tatoris, V. Kennedy, S. Iqbal, G. Devarajan, S. Ellis, Y. Lad, N. G. Clarkson, K. A. Mitrophanous and D. C. Farley, *Nat. Commun.*, 2017, **8**, 14838.
- 11 Y. A. Aksoy, B. Y. Yang, W. J. Chen, T. Hung, R. P. Kuchel, N. W. Zammit, S. T. Grey, E. M. Goldys and W. Deng, *ACS Appl. Mater. Interfaces*, 2020, **12**, 52433–52444.
- 12 J. W. Du, Y. Shao, Y. Hu, Y. W. Chen, J. H. Cang, X. Chen, W. Q. Pei, F. Q. Miao, Y. Q. Shen, M. Muddassir, Y. Zhang, J. Q. Zhang and G. J. Teng, *ACS Appl. Mater. Interfaces*, 2021, **13**, 23396–23409.
- 13 Y. Cheng, R. C. Yumul and S. H. Pun, *Angew. Chem., Int. Ed.*, 2016, **55**, 12013–12017.
- 14 M. M. Khan, N. Filipczak and V. P. Torchilin, *J. Controlled Release*, 2021, **330**, 1220–1228.
- 15 E. M. McErlean, M. Ziminska, C. M. McCrudden, J. W. McBride, S. P. Loughran, G. Cole, E. J. Mulholland, V. Kett, N. E. Buckley, T. Robson, N. J. Dunne and H. O. McCarthy, *J. Controlled Release*, 2021, **330**, 1288–1299.
- 16 Y. Wang, Y. Qi, X. D. Guo, K. Zhang, B. R. Yu and F.-J. Xu, *J. Mater. Chem. B*, 2023, **11**, 4121–4130.
- 17 D. X. Yin, M. J. Zhang, J. X. Chen, Y. Y. Huang and D. H. Liang, *Chin. Chem. Lett.*, 2021, **32**, 1731–1736.
- 18 Q. Q. Fan, Z. Yang, Y. H. Li, Y. Y. Cheng and Y. W. Li, *Adv. Funct. Mater.*, 2021, **31**, 2101646.
- 19 J. H. Li, J. Qian, Y. Y. Xu, S. Yan, J. Shen and M. Z. Yin, *ACS Sustainable Chem. Eng.*, 2019, **7**, 6316.
- 20 N. Zhang, Y. Wang, R. Wu, C. Xu, J.-J. Nie, N. N. Zhao, B. R. Yu, Z. J. Liu and F.-J. Xu, *Small*, 2019, **15**, 1904017.
- 21 Q. Zhang, C. N. Shen, N. Zhao and F.-J. Xu, *Adv. Funct. Mater.*, 2017, **27**, 1606229.
- 22 J. Zhao, F. Han, P. Zhao, X. J. Wen and C. Lin, *J. Mater. Chem. B*, 2017, **5**, 6119–6127.
- 23 N. N. Zhao, W. L. Fan, X. Y. Zhao, Y. J. Liu, Y. Hu, F. Duan and F.-J. Xu, *ACS Appl. Mater. Interfaces*, 2020, **12**, 11341–11352.
- 24 X. Y. Zhao, K. L. Guo, K. Zhang, S. Duan, M. W. Chen, N. N. Zhao and F.-J. Xu, *Adv. Mater.*, 2022, **34**, 2108263.
- 25 J. Chen, K. Wang, J. Y. Wu, H. Y. Tian and X. S. Chen, *Bioconjugate Chem.*, 2019, **30**, 338–349.
- 26 H. Fang, Z. Guo, L. Lin, J. Chen, P. Sun, J. Wu, C. Xu, H. Tian and X. Chen, *J. Am. Chem. Soc.*, 2018, **140**, 11992–12000.
- 27 H. M. Liu, H. Wang, W. J. Yang and Y. Y. Cheng, *J. Am. Chem. Soc.*, 2012, **134**, 17680–17687.
- 28 Y. Y. Pu, H. H. Yin, C. H. Dong, H. J. Xiang, W. C. Wu, B. G. Zhou, D. Du, Y. Chen and H. X. Xu, *Adv. Mater.*, 2021, **33**, 2104641.
- 29 C. N. Wu, J. H. Li, W. D. Wang and P. T. Hammond, *ACS Nano*, 2018, **12**, 6504–6514.

- 30 Y. L. Cheng, D. L. Sellers, J. K. Y. Tan, D. J. Peeler, P. J. Horner and S. H. Pun, *Biomaterials*, 2017, **127**, 89–96.
- 31 Q. Yang, S. Liu, X. Liu, Z. H. Liu, W. Xue and Y. Zhang, *Acta Biomater.*, 2019, **96**, 436–455.
- 32 C. Zhao, L. Shao, J. Lu, X. Deng and Y. Wu, *ACS Appl. Mater. Interfaces*, 2016, **8**, 6400–6410.
- 33 Y. J. Feng, Z. P. Guo, J. Chen, S. J. Zhang, J. Y. Wu, H. Y. Tian and X. S. Chen, *Biomater. Sci.*, 2022, **10**, 6230–6243.
- 34 W. C. Zhang, Y. X. Zhang, Y. L. Luo, S. B. Chen, Q. Y. Huang, Z. T. Cao, M. Liang and X. Z. Yang, *Nano Today*, 2022, **43**, 101418.
- 35 N. Zheng, X. Q. Luo, Z. Y. Zhang, A. G. Wang and W. Z. Song, *ACS Appl. Mater. Interfaces*, 2021, **13**, 27513–27521.
- 36 Y. L. Cheng, H. Wei, J. K. Y. Tan, D. J. Peeler, D. O. Maris, D. L. Sellers, P. J. Horner and S. H. Pun, *Small*, 2016, **12**, 2750–2758.
- 37 S. T. G. Street, H. C. Parkin, L. Shopperly, J. Chrenek, K. Letwin, S. M. Willerth and I. Mannes, *Biomater. Sci.*, 2023, **11**, 3512–3523.
- 38 E. C. Tan, J. Lv, J. J. Hu, W. W. Shen, H. Wang and Y. Y. Cheng, *J. Mater. Chem. B*, 2018, **6**, 7230–7238.
- 39 B. C. Jiang, H. He, L. Yao, T. Zhang, J. P. Huo, W. Sun and L. C. Yin, *Biomacromolecules*, 2017, **18**, 877–885.
- 40 Y. Jin, W. Yu, W. K. Zhang, C. Wang, Y. Liu, W. E. Yuan and Y. Feng, *J. Colloid Interface Sci.*, 2023, **648**, 287–298.
- 41 Z. W. Zhou, M. H. Zhang, Y. D. Liu, C. Z. Li, Q. Y. Zhang, D. Oupicky and M. J. Sun, *Biomacromolecules*, 2018, **19**, 3776–3787.
- 42 C. Xu, W. T. Hu, N. Zhang, Y. Qi, J.-J. Nie, N. N. Zhao, B. R. Yu and F.-J. Xu, *Biomaterials*, 2020, **248**, 120031.
- 43 J. Y. Zhu, X. Zeng, S. Y. Qin, S. S. Wan, H. Z. Jia, R. X. Zhuo, J. Feng and X. Z. Zhang, *Biomaterials*, 2016, **83**, 79–92.
- 44 J. Kim, Y. C. Kang, S. Y. Tzeng and J. J. Green, *Acta Biomater.*, 2016, **41**, 293–301.
- 45 A. M. Martinez, R. H. F. V. de Souza, M. S. Petronio, G. O. Martins, J. C. Fernandes, M. Benderdour, V. A. O. de Tiera and M. J. Tiera, *J. Nanostruct. Chem.*, 2022, 1–20.
- 46 G. Ahn, S. Kweon, C. Yang, J. E. Hwang, K. Kim and B. S. Kim, *J. Polym. Sci., Part A: Polym. Chem.*, 2017, **55**, 4013–4019.
- 47 X. F. Pan, Y. Y. Liu, Z. J. Li, S. D. Cui, H. Gebru, J. X. Xu, S. Q. Xu, J. Q. Liu and K. Guo, *Macromol. Chem. Phys.*, 2017, **218**, 1600483.
- 48 P. F. Song, Y. L. Chen, Y. L. Li, J. P. Ma, L. Y. Wang and R. M. Wang, *Macromol. Rapid Commun.*, 2020, **41**, 2000436.
- 49 Y. Zhao, B. R. Yu, H. Hu, Y. Hu, N. N. Zhao and F.-J. Xu, *ACS Appl. Mater. Interfaces*, 2014, **6**, 17911–17919.
- 50 Y. Zhi, C. Xu, D. D. Sui, J. Du, F.-J. Xu and Y. L. Li, *Adv. Sci.*, 2019, **6**, 1900023.
- 51 Y. J. Huang, H. Hu, R. Q. Li, B. R. Yu and F.-J. Xu, *ACS Appl. Mater. Interfaces*, 2016, **8**, 3919–3927.
- 52 J.-J. Nie, B. K. Qiao, S. Duan, C. Xu, B. Y. Chen, W. J. Hao, B. R. Yu, Y. L. Li, J. Du and F.-J. Xu, *Adv. Mater.*, 2018, **30**, 1801570.
- 53 Y. L. Ren, R. Q. Li, Y. R. Cai, T. Xia, M. Yang and F.-J. Xu, *Adv. Funct. Mater.*, 2016, **26**, 7314–7325.
- 54 C. Xu, B. R. Yu, H. Hu, M. N. Nizam, W. Yuan, J. Ma and F.-J. Xu, *Biomater. Sci.*, 2016, **4**, 1233–1243.
- 55 C. Xu, Y. Z. Z. Zhang, K. Xu, J.-J. Nie, B. R. Yu, S. J. Li, G. Cheng, Y. L. Li, J. Du and F.-J. Xu, *Nat. Commun.*, 2019, **10**, 3184.
- 56 R. Q. Li, H. Q. Song and F.-J. Xu, *Polym. Chem.*, 2015, **6**, 6208–6218.
- 57 Y. Qi, C. Xu, M. N. Nizam, Y. Li, B. R. Yu and F.-J. Xu, *Polym. Chem.*, 2016, **7**, 5630–5640.
- 58 Y. Sun, H. Hu, N. N. Zhao, T. Xia, B. R. Yu, C. A. Shen and F.-J. Xu, *Biomaterials*, 2017, **117**, 77–91.
- 59 L. Rotolo, D. Vanover, N. C. Bruno, H. E. Peck, C. Zurla, J. Murray, R. K. Noel, L. O'Farrell, M. Arainga, N. Orr-Burks, J. Y. Joo, L. C. S. Chaves, Y. H. Jung, J. Beyersdorf, S. Gumber, R. Guerrero-Ferreira, S. Cornejo, M. Thoresen, A. K. Olivier, K. T. M. Kuo, J. C. Gumbart, A. R. Woolums, F. Villinger, E. R. Lafontaine, R. J. Hogan, M. G. Finn and P. J. Santangelo, *Nat. Mater.*, 2023, **22**, 369.
- 60 L. Ye, H. M. Liu, X. Fei, D. Ma, X. Z. He, Q. Y. Tang, X. Zhao, H. B. Zou, X. J. Chen, X. M. Kong and P. F. Liu, *Nano Res.*, 2022, **15**, 1135–1144.
- 61 C. N. Zhang, R. Jin, P. Zhao and C. Lin, *Acta Biomater.*, 2015, **22**, 120–130.
- 62 D. M. Lynn and R. Langer, *J. Am. Chem. Soc.*, 2000, **122**, 10761–10768.
- 63 Y. Liu, Y. F. Li, D. Keskin and L. Q. Shi, *Adv. Healthcare Mater.*, 2019, **8**, 1801359.
- 64 Y. Rui, D. R. Wilson, J. Choi, M. Varanasi, K. Sanders, J. Karlsson, M. Lim and J. J. Green, *Sci. Adv.*, 2019, **5**, eaay3255.
- 65 Q. Y. Zhu, H. M. Shen, Z. J. Yang and H. B. Ji, *Chin. J. Catal.*, 2016, **37**, 1227–1234.
- 66 W. R. Cheng, D. C. Wu and Y. Liu, *Biomacromolecules*, 2016, **17**, 3115–3126.
- 67 B. D. Mather, K. Viswanathan, K. M. Miller and T. E. Long, *Prog. Polym. Sci.*, 2006, **31**, 487–531.
- 68 R. J. Lu, Y. J. Zheng, M. R. Wang, J. H. Lin, Z. Y. Zhao, L. Chen, J. H. Zhang, X. Liu, L. C. Yin and Y. B. Chen, *Acta Biomater.*, 2022, **152**, 355–366.
- 69 S. Z. Duan, D. S. Cao, X. D. Li, H. F. Zhu, M. Lan, Z. Z. Tan, Z. Y. Song, R. Y. Zhu, L. C. Yin and Y. B. Chen, *Biomater. Sci.*, 2020, **8**, 290–301.
- 70 Y. Rui, D. R. Wilson, K. Sanders and J. J. Green, *ACS Appl. Mater. Interfaces*, 2019, **11**, 10472–10480.
- 71 S. Duan, B. R. Yu, C. X. Gao, W. Yuan, J. Ma and F.-J. Xu, *ACS Appl. Mater. Interfaces*, 2016, **8**, 29334–29342.
- 72 Q. F. Zhang, Q. Y. Yu, Y. Y. Geng, J. Zhang, W. X. Wu, G. Wang, Z. W. Gu and X. Q. Yu, *ACS Appl. Mater. Interfaces*, 2014, **6**, 15733–15742.
- 73 T. Wu, Y. Qi, C. Xu, D. D. Sui and F.-J. Xu, *Nano Today*, 2023, **50**, 101887.

- 74 Y. Qi, C. Xu and F.-J. Xu, *WIREs Nanomed. Nanobiotechnol.*, 2020, **12**, e1631.
- 75 F.-J. Xu, *Prog. Polym. Sci.*, 2018, **78**, 56–91.
- 76 Y. C. Zheng, S. P. Li, Z. L. Weng and C. Gao, *Chem. Soc. Rev.*, 2015, **44**, 4091–4130.
- 77 Q. Zeng, Y. W. Zhu, B. R. Yu, Y. J. Sun, X. K. Ding, C. Xu, Y. W. Wu, Z. H. Tang and F.-J. Xu, *Biomacromolecules*, 2018, **19**, 2805–2811.
- 78 Z. Guo, S. Li, M. Lv, Z. H. Liu and W. Xue, *ACS Biomater. Sci. Eng.*, 2017, **3**, 2420–2430.
- 79 C. R. Wang, X. X. Wang, L. L. Du, Y. L. Dong, B. Hu, J. H. Zhou, Y. L. Shi, S. P. Bai, Y. Y. Huang, H. Q. Cao, Z. C. Liang and A. J. Dong, *ACS Appl. Mater. Interfaces*, 2021, **13**, 2218–2229.
- 80 B. Mondal, T. Dutta and S. S. Gupta, *Chem. Commun.*, 2021, **57**, 109–112.
- 81 M. R. Schwartz, A. C. Debski and R. J. Price, *J. Controlled Release*, 2021, **339**, 531–546.
- 82 C. Sardo, E. F. Craparo, B. Porsio, G. Giammona and G. Cavallaro, *Polymers*, 2019, **11**, 889.
- 83 G. B. Yang, J. J. Liu, Y. F. Wu, L. Z. Feng and Z. Liu, *Coord. Chem. Rev.*, 2016, **320**, 100–117.
- 84 R. L. Zhu, A. P. Qian, J. T. Yan, W. Li, K. Liu, T. Masuda and A. F. Zhang, *ACS Appl. Mater. Interfaces*, 2018, **10**, 13258–13263.
- 85 Y. J. Huang, X. K. Ding, Y. Qi, B. R. Yu and F.-J. Xu, *Biomaterials*, 2016, **106**, 134–143.
- 86 Y. M. Chen, Y. Qi, W. T. Hu, Z. Di, B. R. Yu and F.-J. Xu, *Acta Polym. Sin.*, 2019, **50**, 602–612.
- 87 Y. Qi, H. Q. Song, H. H. Xiao, G. Cheng, B. R. Yu and F.-J. Xu, *Small*, 2018, **14**, 1803061.
- 88 M. Y. Shao, Y. Q. Fan, K. Zhang, Y. Hu and F.-J. Xu, *Nano Today*, 2021, **39**, 101224.
- 89 Y. Qi, Y. L. Liu, B. R. Yu, Y. Hu, N. S. Zhang, Y. Zheng, M. Yang and F.-J. Xu, *Adv. Sci.*, 2020, **7**, 2001424.
- 90 J. L. Gao, T. Luo and J. K. Wang, *Nat. Commun.*, 2021, **12**, 5311.
- 91 R. W. Herzog and M. Suzuki, *Cell*, 2020, **186**, 893–893.
- 92 F. Li, N. C. Song, Y. H. Dong, S. Li, L. H. Li, Y. J. Liu, Z. M. Li and D. Y. Yang, *Angew. Chem., Int. Ed.*, 2022, **61**, e202116569.
- 93 S. Norberg, E. M. Von Euw, G. Parry, S. Highfill, Z. Franco, J. L. Gulley and C. S. Hinrichs, *J. Clin. Oncol.*, 2022, **40**, 16.
- 94 T. Zheng, W. T. Wang, M. Mohammadniaei, J. Ashley, M. Zhang, N. L. Zhou, J. Shen and Y. Sun, *Adv. Sci.*, 2022, **9**, 2103812.
- 95 M. H. Andersen, *Semin. Immunopathol.*, 2023, **45**, 253–264.
- 96 T. Suwa, M. Kobayashi, J. M. Nam and H. Harada, *Exp. Mol. Med.*, 2021, **53**, 1029–1035.
- 97 Q. F. Zhang, G. Z. Kuang, W. Z. Li, J. L. Wang, H. Z. Ren and Y. J. Zhao, *Nano-Micro Lett.*, 2023, **15**, 44.
- 98 E. K. Bang, G. Gasparini, G. Molinard, A. Roux, N. Sakai and S. Matile, *J. Am. Chem. Soc.*, 2013, **135**, 2088–2091.
- 99 K. Luo, C. X. Li, L. Li, W. C. She, G. Wang and Z. W. Gu, *Biomaterials*, 2012, **33**, 4917–4927.
- 100 E. Boedtkjer and S. F. Pedersen, *Annu. Rev. Physiol.*, 2020, **82**, 103–126.
- 101 S. Taylor, E. P. Spugnini, Y. G. Assaraf, T. Azzarito, C. Rauch and S. Fais, *Drug Resistance Updates*, 2015, **23**, 69–78.
- 102 C. C. Wang, P. R. Zhao, G. L. Yang, X. Y. Chen, Y. Q. Jiang, X. W. Jiang, Y. L. Wu, Y. Y. Liu, W. A. Zhang and W. B. Bu, *Mater. Horiz.*, 2020, **7**, 1180–1185.
- 103 D. P. Feldmann, Y. L. Cheng, R. Kandil, Y. R. Xie, M. Mohammadi, H. Harz, A. Sharma, D. J. Peeler, A. Moszczynska, H. Leonhardt, S. H. Pun and O. M. Merkel, *J. Controlled Release*, 2018, **276**, 50–58.
- 104 Y. H. Li, K. Tang, X. Zhang, W. Pan, N. Li and B. Tang, *Chem. Commun.*, 2022, **58**, 8754–8765.
- 105 G. Yang, Y. Liu, J. Chen, J. Ding and X. Chen, *Acc. Chem. Res.*, 2022, **3**, 1232–1247.
- 106 W. H. Fan, M. Y. Shao, J. W. Zhang, G. S. Jin, F.-J. Xu and F. S. Liu, *Adv. Funct. Mater.*, 2019, **29**, 1807104.
- 107 Z. X. Liao, Y. C. Li, H. M. Lu and H. W. Sung, *Biomaterials*, 2014, **35**, 500–508.
- 108 D. F. Chen, C. Y. Lei, W. F. Liu, M. Y. Shao, M. Z. Sun, J. X. Guo, J. J. Cao, P. Luo, Y. W. Luo, B. R. Yu, R. X. Wang, S. Duan and F.-J. Xu, *Bioact. Mater.*, 2023, **28**, 376–385.
- 109 S. Padmakumar, A. D'Souza, N. N. Parayath, B. S. Bleier and M. M. Amiji, *J. Controlled Release*, 2022, **352**, 121–145.
- 110 E. Wagner, R. Kircheis and G. F. Walker, *Biomed. Pharmacother.*, 2004, **58**, 152–161.
- 111 S. T. Xie, W. D. Sun, T. Fu, X. S. Liu, P. Chen, L. P. Qiu, F. L. Qu and W. H. Tan, *J. Am. Chem. Soc.*, 2023, **145**, 7677–7691.
- 112 M. Y. Shao, Y. Qi, D. D. Sui and F.-J. Xu, *Biomater. Sci.*, 2021, **9**, 7104–7114.
- 113 K. Chevreul, K. B. Brigham and C. Bouche, *Global Health*, 2014, **10**, 6.
- 114 M. R. Khanna, J. Kovalevich, V. M. Y. Lee, J. Q. Trojanowski and K. R. Brunden, *Alzheimers Dement.*, 2016, **12**, 1051–1065.
- 115 L. S. Satin, S. A. Soleimanpour, E. M. Walker and L. Isom, *Pharmacol. Rev.*, 2021, **73**, 1001–1015.
- 116 Y. Q. Pan, M. Y. Shao, P. Li, C. Xu, J.-J. Nie, K. Zhang, S. Wu, D. D. Sui and F.-J. Xu, *J. Controlled Release*, 2022, **343**, 420–433.
- 117 H. Q. Song, M. Y. Shao, Y. Li, X. J. Ding and F.-J. Xu, *Adv. Healthcare Mater.*, 2019, **8**, 1800889.
- 118 Y. W. Zhu, M. Y. Lin, W. T. Hu, J. K. Wang, Z. G. Zhang, K. Zhang, B. R. Yu and F.-J. Xu, *Angew. Chem., Int. Ed.*, 2022, **61**, e202200535.
- 119 W. J. Ye, Y. M. Chen, W. X. Tang, N. Zhang, Z. H. Li, Z. J. Liu, B. R. Yu and F.-J. Xu, *ACS Appl. Mater. Interfaces*, 2019, **11**, 28307–28316.
- 120 Y. W. Zhu, C. Xu, N. Zhang, X. K. Ding, B. R. Yu and F.-J. Xu, *Adv. Funct. Mater.*, 2018, **28**, 1708709.