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Advancing cell surface modification in mammalian cells with synthetic molecules

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Biological cells, being the fundamental entities of life, are widely acknowledged as intricate living machines. The manipulation of cell surfaces has emerged as a progressively significant domain of investigation and advancement in recent times. Particularly, the alteration of cell surfaces using meticulously crafted and thoroughly characterized synthesized molecules has proven to be an efficacious means of introducing innovative functionalities or manipulating cells. Within this realm, a diverse array of elegant and robust strategies have been recently devised, including the bioorthogonal strategy, which enables selective modification. This review offers a comprehensive survey of recent advancements in the modification of mammalian cell surfaces through the use of synthetic molecules. It explores a range of strategies, encompassing chemical covalent modifications, physical alterations, and bioorthogonal approaches. The review concludes by addressing the present challenges and potential future opportunities in this rapidly expanding field.

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1. Introduction

Mammalian cells (hereinafter referred to as “cells”), as natural constituents of organisms, have been propelled into the spotlight in the biomedical field, primarily due to their unique characteristics, such as biosynthesis ability, communication and interaction ability, and migration ability, among others.^{1–4} Over the past few decades, the manipulation of cells has provided a powerful tool to enhance our understanding of the

underlying mechanisms governing various biological behaviors in basic research and has also promoted the development in biomedical applications, such as medical diagnosis and cell-based therapy.⁵ For instance, mesenchymal stem cells (MSCs), red blood cells (RBCs), and macrophages exhibit promising capabilities as delivery vehicles to transport diagnostic molecules or therapeutic agents.^{6–8} Immune cells, including T cells and natural killer (NK) cells, have emerged as highly prominent candidates for tumor immunotherapy owing to their specific cytotoxicity against tumor cells while sparing normal cells.^{9,10}

Despite these exciting achievements, the functions of natural cells themselves are limited. The cell surface, also known as the cell membrane, is a highly heterogeneous and dynamic milieu comprising lipids, proteins, carbohydrates, and their complexes, which governs numerous intracellular and

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extracellular processes.¹¹ Simultaneously, the complex cell surface provides plenty of opportunities for further modification aimed at achieving particular functionalities, a process referred to as cell surface modification. This process serves as a powerful means to facilitate the biomedical application of natural cells.¹² One notable example is the universal blood, which involves the modification of red blood cell surfaces to impede the recognition of antigenic sites by antibodies, thereby preventing immune responses caused by blood type incompatibility.^{13–15} Chemical manipulation of cell behavior and function through modification of cell surfaces using precisely synthesized and well-characterized synthetic molecules, such as polymers, is a captivating area of research.¹⁶ Various strategies have been devised for this purpose. While several reviews have provided summaries of certain strategies, there is a noticeable dearth of comprehensive discussions specifically centered on the strategy of cell surface modification using synthetic molecules.^{16–18}

Hence, this paper provides a comprehensive review of the latest advancements in strategies related to the modification of cell surfaces accompanied by a discussion on their biomedical

applications, with a particular emphasis on developments from 2017 onwards. These strategies encompass a range of approaches, including chemical covalent approaches, physical techniques, and bioorthogonal methods for synthetic molecules (Fig. 1). The synthetic molecules discussed include specific functional groups, synthetic functional small molecules, synthetic polymers, synthetic nanoparticles, synthetic cell coatings, and synthetic DNA, among others. Lastly, the paper discusses the existing challenges and potential future prospects in this rapidly expanding field.

2. Chemical covalent modification

Chemical covalent modification is a strategic approach that entails the utilization of the chemically reactive functionalities found on the surface of the cell membrane. The cell membrane, a multifaceted chemical structure composed of lipids, proteins, carbohydrates and other components, offers a diverse array of functional groups that can be employed for chemical covalent binding.¹⁹ Previous researches have predominantly focused on employing amine, thiol, and vicinal diol groups present on amino acid residues within proteins or sugar residues as the most frequently utilized groups (Fig. 2).²⁰ The stable attachment of synthetic molecules and the absence of cell pretreatment are the primary benefits of this approach, making it a simple yet effective method for modifying cell membranes. However, it is widely recognized that directly modifying cell membranes with reactive functional groups through covalent bonds can potentially impair the functionality of membrane proteins and subsequently impair cellular functions. Consequently, when employing this strategy, careful attention must be paid to both cell viability and effector functions.

2.1 Amine-mediated covalent modification strategy

Amine groups ($-NH_2$) are mainly present at the lysine residues of proteins and the N-terminus of polypeptide chains. They are



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Fig. 1 Mammalian cell surface modification strategies. (A) Chemical covalent modification strategies. (B) Physical modification strategies. (C) Bioorthogonal modification strategies.

widely used for chemical modification of cell membrane surfaces due to their ease of chemical covalent modification and mild reaction conditions. The amine-mediated covalent binding strategy can be achieved through two primary pathways: acylation or alkylation. Generally, these reactions exhibit rapidity and selectivity, resulting in the formation of stable bonds (such as amide or secondary amine bonds) and high yields.

Among various kinds of reagents, *N*-hydroxysuccinimide (NHS) ester is the most frequently used to covalently bind to $-NH_2$ on cell membranes. Recently, Cai *et al.* modified human umbilical vein endothelial cells (HUVECs) and human skin fibroblasts (HSFs) with succinimide ester-methoxy polyethylene glycol (NHS-mPEG), resulting in a significant enhancement of cell migration ability and motility through reduction of the focal adhesion area.²¹ In a separate study shown in Fig. 3A, Wang *et al.* employed acrylic acid NHS ester (NHS-AA) to immobilize vinyl onto cell membranes, followed by free radical polymerization to covalently attach polymers to the membrane.²² After subsequent ion exchange and electroless deposition (ELD), the polymer-functionalized cells could be converted into metallic biocomposites, which can be applied in the fields of biosensors, electronics, and energy. Sulfo-NHS ester is a more suitable reagent for covalent reactions with $-NH_2$ groups on the cell

membrane due to its enhanced water solubility and negative charge, which reduces the transmembrane permeability of the sulfo-NHS ester. For instance, Jasiewicz *et al.* employed sulfo-succinimidyl 4-(*N*-maleimidomethyl) cyclohexane-1-carboxylate (sulfo SMCC), a heterobifunctional crosslinker, to modify MSCs by covalently binding to the amines on the cell membrane and subsequently decorating them with heterodimerizing leucine zippers.²³ In addition, sulfo-NHS-biotin was also commonly utilized to modify the cell membrane where it serves as a “bridge” and facilitates the decoration of the cell membrane with cargoes through streptavidin–biotin interaction.^{24–29} In addition to NHS ester derivatives, other types of reagents have been developed for covalently binding to $-NH_2$ including cyanuric chloride and benzotriazole carbonate.^{30,31}

2.2 Thiol-mediated covalent modification strategy

Thiol groups ($-SH$), mainly located on the cysteine residues of amino acids in proteins, are one of the most potent nucleophiles, stronger than amino groups. Thiol groups are frequently employed for the covalent modification of cell membranes. Maleimide derivatives, which form stable thioether bonds with thiol groups through an energetically favorable Michael addition reaction, are the most widely used reaction reagents because they exhibit high stability and chemoselectivity with





Fig. 2 Illustration of chemical covalent modification through the reactions between functional groups on cells and synthetic molecules.

thiol groups. The significant advantage of this strategy lies in the extensive availability of commercially accessible reagents and linkers.

Early research was conducted by Irvine's team which focused on surface modification of various cell types containing thiols. They investigated the application of liposomes and liposome-like nanoparticles containing maleimide terminal groups for T cells, hematopoietic stem cells, and other cell surface modifications.^{32–36} Recently, Wang *et al.* developed PEGylated solid lipid nanoparticles functionalized with maleimide end groups (SLN-PEG-Mal).³⁶ As is shown in Fig. 3B, by exploiting the reaction between maleimide and sulfhydryl groups on the surface of RBCs, the researchers successfully enhanced the adsorption of modified nanoparticles onto RBCs, leading to significant alterations in the properties and morphology of RBCs. Moreover, these nanoparticle-loaded RBCs exhibited a remarkable ability to be engulfed by macrophages, thereby demonstrating promising potential for targeted drug delivery to macrophages. Wang *et al.* utilized 2-iminothiolane (Traut's agent), a thiolation reagent, to introduce extra free thiol groups by capping primary amines with thiol groups. This allows them to modify platelets with PD-L1 antibody, thereby reducing post-surgical tumor recurrence and metastasis.³⁵

Research has been conducted to combine maleimide with other functional components in order to develop multifunctional nanoparticles.³⁸ For instance, Luo *et al.* synthesized double-bound magnetic nanoparticles (DBMN) containing PEG-

Mal, hyaluronic acid (HA), and Fe_3O_4 .³⁸ Following a simple incubation, DBMN was able to anchor onto the cell membrane through a Michael addition reaction between the Mal component and sulfhydryl groups on the T cell surface, resulting in magnetized T cells (DBMN-T). Under external magnetic field guidance, DBMN-T exhibited excellent targeting ability. Additionally, HA could bind to highly expressed CD44 on tumor cells, promoting recognition and killing of tumor cells.

In addition to the reaction between maleimides and thiols, cell surface modification could also be achieved through the exchange between disulfide bonds and thiols.^{39–43} Wayteck *et al.* incorporated thiol-reactive phospholipids into liposome bilayers with a pyridyldithiopropionate (PDP) head group, which was capable of forming reducible disulfide bonds with thiol groups exposed on the cell surface, thereby enabling the covalent coupling of liposomes.³⁹

2.3 Vicinal diol-mediated covalent modification strategy

Vicinal diol groups are abundant on the cell membrane and primarily originate from sialic acid (SA), mannose, and galactose residues within glycoproteins and the extracellular matrix. Phenylboronic acid (PBA) derivatives form unique dynamic covalent bonds with vicinal diol groups of the cell membrane and are affected by pH, making them of substantial interest. However, only sialic acids can be efficiently coupled to PBA under physiological conditions, while other diol groups require alkaline reaction conditions with a pH



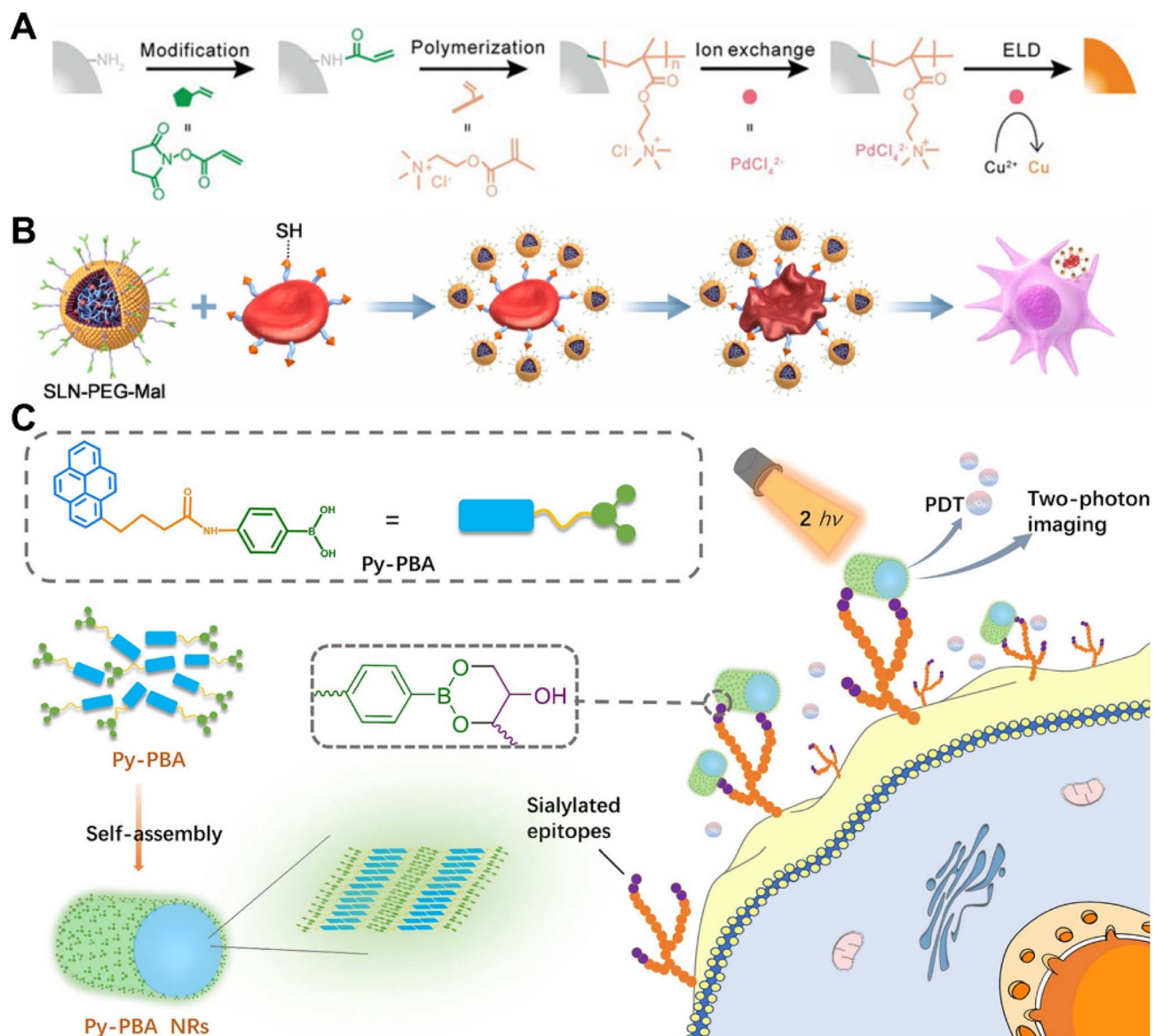


Fig. 3 Methods based on chemical covalent modification. (A) Schematic illustration of realizing polymer-assisted cell metallization by the use of reactions between amino groups on cell membranes and NHS-AA.²² Copyright 2021, John Wiley & Sons, Inc. (B) Schematic illustration of modifying RBCs with SLN-PEG-Mal through reactions between maleimide and thiol groups, leading to their engulfment by macrophages.³⁶ Copyright 2023, Elsevier. (C) Schematic illustration of combining Py-phenylboronic acid (PBA) NRs to the cell membrane by vicinal diol groups for two-photon imaging of cell surface sialic acids and PDT.³⁷ Copyright 2021, American Chemical Society.

value higher than the pK_a of PBA. Therefore, most studies have focused on the reaction between PBA derivatives and SA on the cell membrane surface. Tao and colleagues reported a novel fluorescent polymer containing PBA through the combination of multicomponent reactions (MCRs) with reversible addition-fragmentation chain transfer (RAFT) polymerization.^{44,45} Specifically, the Hantzsch reaction, a classical four-component reaction, was carried out simultaneously with RAFT polymerization to create innately fluorescent 1,4-dihydropyridine (1,4-DHP). The formed fluorescent polymer is suitable for cell membrane conjugation and imaging through the interaction between phenylboronic acid and sialic acid on the cell membrane. In our

laboratory, we utilized a copolymer containing PBA groups to modify silicon nanowire arrays which exhibited a high capture capacity for cells overexpressing SA on the membrane. This modification also allowed for high efficiency of intracellular delivery of diverse biomacromolecules.⁴⁶ Additionally, overexpression of SA has been demonstrated in various tumors, including lung, melanoma, colon, and breast cancers.⁴⁷ Consequently, the SA-PBA reaction was commonly employed for tumor cell imaging, targeting, and capturing.^{37,48-54} Li *et al.* developed self-assembled nanorods of PBA-functionalized pyrene (Py-PBA NRs), which possess a highly efficient and specific imaging feature of SA on the cell membrane. Three cell lines with different expression levels of



SA were utilized to demonstrate this imaging ability. Additionally, the nanorods exhibited efficient generation of $^1\text{O}_2$ under two-photon irradiation, providing potential possibilities for tumor therapy (Fig. 3C).³⁷ Furthermore, PBA derivatives with low pK_a values have been developed to enhance the applicability of this strategy to cell species characterized by low expression levels of sialic acids. A series of PBAs with different substituents were synthesized, and it was demonstrated that the introduction of electron-withdrawing groups, such as fluoro and nitro, effectively decreased the pK_a even to 4.2.⁵⁵ However, it should be noted that the optimal binding pH may not always exceed the pK_a of PBAs, particularly in complex multicomponent systems.⁵⁶

In addition to PBA derivatives, benzoxaborole (BA), a cyclic hemi-ester of boronic acid, can also be utilized for cell surface modification *via* the covalent reaction with vicinal diol.⁵⁷ Morgese *et al.* have successfully modified supramolecular polymers containing BAs onto the surface of human RBCs *via* the covalent reaction between BAs and SA.⁵⁸ The specific interactions between functional copolymers and the cell surface were further visualized in real time using total internal reflection fluorescence microscopy.

2.4 Other functional groups-mediated covalent modification strategy

Carboxyl groups are abundantly present on the cell membrane, mainly distributed at the residues of aspartic acid (Asp) and glutamic acid (Glu) within membrane proteins, as well as at the C-terminus of polypeptide chains. However, the modification of membranes using carboxyl groups necessitates pre-activation of these groups, typically employing an activator known as 3-(ethyliminomethylideneamino)-*N,N*-dimethylpropan-1-amine (EDC), which causes significant harm to mammalian cell viability. Consequently, the utilization of carboxyl groups for cell modification is often limited. Recently, Ma *et al.* developed a novel probe (3-phenyl-2*H*-azirine) that effectively labeled carboxyl groups on the surface of living cells. This presented new possibilities for chemical modification utilizing carboxyl groups present on the cell membrane.⁵⁹

In addition to utilizing existing groups for direct modification of the cell membrane, strategies have been devised to convert commonly present but difficult-to-modify functional groups into easily modifiable ones through mild oxidation or reduction reactions on the cell surface under gentle conditions. For example, researchers have found that a mild oxidation reaction using NaIO_4 can convert diols on cell surfaces into aldehydes, which can be treated as active sites for subsequent cell modification.^{60–65} Recently, Liu *et al.* proposed a novel cell surface engineering platform using classical thiazolidine chemistry to combine small molecules containing aminothiols moieties with cells pretreated with aldehyde groups on their surface by NaIO_4 .⁶⁴

Efforts have also been devoted to converting disulfide bonds (S–S) on cell membranes to thiol groups and tris(2-carboxyethyl) phosphine (TCEP) is a widely used mild reducing agent in this strategy. Researchers have utilized the strategy to modify

synthetic materials including chondroitin sulfate (CS), PEG, mesoporous silica nanoparticles, and silver nanoclusters on the cell membrane for a diverse range of applications.^{66–70}

3. Physical modification

In addition to the chemical covalent binding strategy, physical approaches such as hydrophobic insertion, membrane fusion, electrostatic interaction, and layer-by-layer self-assembly offer versatile and easy ways to introduce synthetic molecules to the cell membrane while maintaining cellular physiology.

3.1 Hydrophobic insertion

The cell membrane skeleton is a phospholipid bilayer structure allowing the spontaneous insertion of synthetic materials with hydrophobic anchors or “tails” driven by the hydrophobic effect.^{71–73} The commonly used hydrophobic anchors include lipids (phospholipids^{21,26,73–80} and cholesterol^{81–92}), alkane chains^{93–98} and oleyl chains.^{99–105} They can be categorized into single and multiple anchors based on the number of hydrophobic anchors. Shi *et al.* constructed a polyvalent antibody mimic (PAM) for engineering NK cells with highly efficient targeting, adhesion and killing effects for tumor cells.⁷¹ The DNA initiator (DI) with a single anchor was displayed on the NK cell membrane by the hydrophobic insertion approach. Subsequently, a DNA scaffold was synthesized and hybridized with multiple aptamers *in situ* forming PAM-engineered NK cells. Sun *et al.* reported a DNA-assisted bottom-up self-assembly approach for achieving precise control over the lateral and vertical distributions of T cell activation ligands on RBCs and constructing RBCs-based artificial antigen presenting cells (aAPCs) which could effectively activate and expand T cells.¹⁰⁷ DNA strands with a cholesterol end group were inserted into the membranes by hydrophobic interaction and then bound with T cell activation ligands through specific DNA hybridization as well as biotin–avidin interaction. The vertical distributions of T cell activation ligands can be easily manipulated by adjusting the length of DNA strands, while the lateral distributions were achieved through biotin–avidin interaction. The subsequent study shown in Fig. 4 employed the approach to construct lymphocyte-based aAPCs exhibiting homologous targeting functionality for personalized cancer immunotherapy.¹⁰⁶ Zhao *et al.* developed a surface-anchored framework for sheltering the epitopes on Rhesus D (RhD)-positive RBCs.¹⁰² RBCs were modified with horseradish peroxidase containing a single oleyl chain *via* hydrophobic insertion, thereby catalyzing the reaction of H_2O_2 to construct a polysialic acid (PSA)-tyramine framework on the RBC membrane. The crosslinking framework successfully achieved transfusion of the modified RBCs to RhD-negative recipients without eliciting immunogenicity, by effectively balancing the modified fluidity of RBC membranes and shielding of RhD antigens.

In addition to the single anchor, hydrophobic insertion moieties with more anchors were developed. Niu *et al.* reported the first effort for cytocompatible controlled radical polymerization (CRP) techniques.⁷² Chain-transfer agents



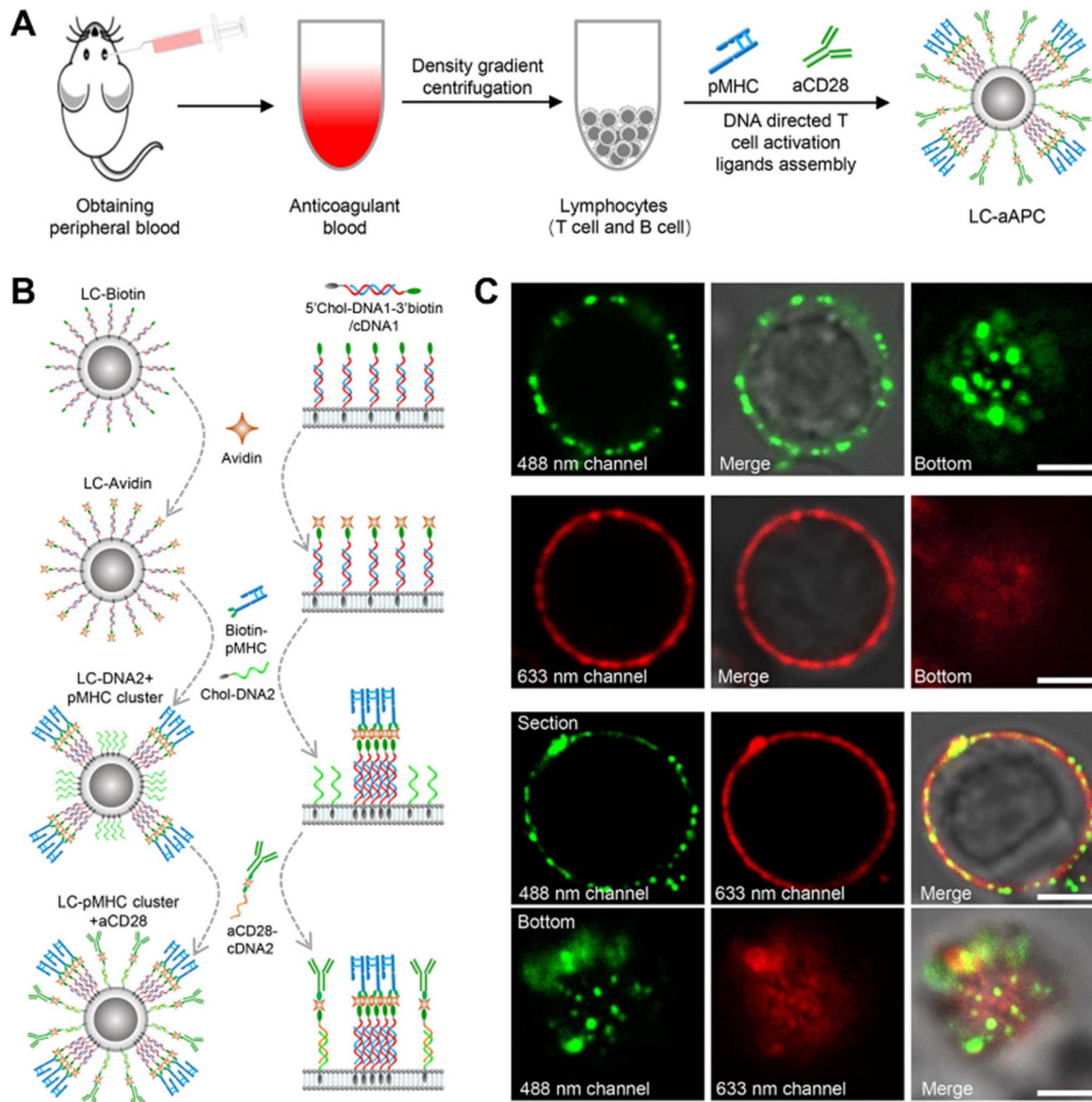


Fig. 4 Construction of LC-aAPCs with lymphocytes from peripheral blood.¹⁰⁶ (A) Schematic diagram of constructing lymphocyte-based aAPCs from peripheral blood for personalized tumor immunotherapy. (B) Schematic illustration of DNA-mediated bottom-up assembly of pMHC-I and aCD28 on lymphocytes, including hydrophobic insertion and specific DNA hybridization as well as biotin–avidin interaction. (C) Confocal microscope images showing distribution of pMHC-I and aCD28 on the surface of lymphocytes. Copyright 2022, John Wiley & Sons, Inc.

with a two-tailed hydrophobic anchor, 1,2-distearoyl-*sn*-glycero-3-phosphoethanolamine (DSEP), were successfully modified on the cell membranes with the hydrophobic insertion strategy, thereby realizing polymerization to be initiated directly in the live cell surface while maintaining high cell viability. The strategy effectively enhanced the efficiency of grafting polymers compared to the traditional grafting-to methods and offered novel possibilities for modulating cellular interactions.

Additionally, the method was also utilized for T cell modification with liposomal nanoparticles, as described in Hao *et al.*'s investigation.⁷³ The tetrazine (Tre) groups with two-tailed lipids (DSPE) were inserted into T cells and subsequently drug liposomes with bicyclo nonyne (BCN) were modified on the cell membranes of T cells *via* click reaction while preserving the intact functionality of T cells. A platform (Fig. 5A) for cell membrane engineering with modular polymers was developed by our group.⁹¹ The study employed cholesteryl-methacrylate as one of the monomers for





Fig. 5 Representative examples based on hydrophobic insertion and membrane fusion strategies. (A) Schematic illustration of the construction of a platform for the manipulation of cell behaviors, by using hydrophobic insertion to bind modular polymers to the cell surface.⁹¹ Copyright 2019, American Chemical Society. (B) Schematic showing the biomimetic LIFT approach to engineer the plasma membrane by membrane fusion.¹⁰⁸ Copyright 2022, John Wiley & Sons, Inc.



constructing modular polymers with multiple anchors through one-pot RAFT copolymerization, along with deoxy-2-(methacrylamido)glucopyranose (MAG) as a hydrophilic monomer and adamantane carbonyl methacrylate (Ada) as a guest monomer. In addition to the introduction of functional molecules through host-guest units, the residence time of the modular polymers could also be regulated on the cell membrane by adjusting the content of cholesterol modules.

The hydrophobic insertion strategy is considered a simple, powerful, and less invasive approach for cell surface modification. However, functional synthetic materials introduced to the cell membrane surface through hydrophobic insertion are prone to loss during membrane flow and endocytosis, thereby limiting their long-term presence on the cell membrane. The hydrophobic insertion moieties with multiple anchors may offer a promising strategy for achieving relatively stable and long-time modification.

3.2 Membrane fusion

Unlike the strategy of hydrophobic insertion into the cell membrane through hydrophobic anchors, cell surface modification is achieved through liposomes loaded with synthetic materials or functional groups diffusing and mixing with the cell membrane in the membrane fusion strategy.

Sarkar *et al.* developed a versatile platform technology for the modification of cell membranes.¹⁰⁹ Biotinylated lipid vesicles were utilized for the incubation with MSCs, leading to the attachment of biotin on the cell surface *via* vesicle fusion. The biotin moieties serve as binding sites for subsequent ligands. Yousaf and colleagues conducted a series of studies that utilized the membrane fusion strategy to introduce chemical functional groups onto the cell membrane.^{110–118} The prepared lipid, containing either ketone or oxamine molecules, underwent spontaneous insertion and fusion into the cell membrane, resulting in the modification of cells with either ketone or oxamine molecules for subsequent bio-orthogonal ligation reactions.¹¹⁰ Additionally, the bioorthogonal molecules, ketone or oxamine, could also be modified onto different populations of cells using the same method to regulate the cell-cell interaction and generate 3D tissue-like structures.¹¹¹ Following this, the researchers employed a membrane fusion approach to create and modify cell membrane surfaces with bioorthogonal chemical molecules possessing diverse characteristics, including photoresponsive and redox-responsive cleavage. The primary emphasis of their investigation was on the utilization of these modified cells in the field of three-dimensional tissue engineering. Membrane fusion strategies have recently been extensively used as a potent tool for modifying cell membranes in various investigations.

Zheng *et al.* designed core-shell membrane-fusing liposome (MFL) containing NK cell-activating glycans, Lewis X trisaccharide (LeX), and loaded it into a thermosensitive hydrogel which could be released responsively through the tumor microenvironment. Subsequently, the released MFL was fused with tumor cell membranes, realizing the

modification of tumor membranes with Lex which could enhance the anti-tumor effects.¹¹⁹ Shi *et al.* designed T-cell-targeting fusogenic liposomes by conjugating ROS-scavenging groups, 2,2,6,6-tetramethylpiperidine (TEMP) and T-cell-targeting anti-CD3 F(ab')₂ fragments to the surface of liposomes, in which TEMP groups were designed for neutralizing ROS and protecting T cells from an oxidation-induced loss of activity. In the meantime, the procedure would result in paramagnetic transition of TEMP to TEMPO molecules, allowing for the measurement of the *in situ* activity of T cells, enabling a better understanding of engineering T cells for cancer treatment.¹²⁰ Lin *et al.* reported a liposomal fusion-based transport (LiFT) strategy to anchor functional DNA strands on the inner face of the cell membrane, addressing the previous lack of suitable synthetic tools to engineer the intracellular interior (Fig. 5B).¹⁰⁸ In subsequent studies, the group combined membrane-anchored catalysts with the previously reported LiFT strategy, through which they were able to prepare the corresponding fusion liposome catalyst through a simple strategy, achieving precise position control of the catalyst on the cell membrane.¹²¹ The drug molecules generated by this method may have higher drug delivery efficiency than traditional methods using drug delivery vehicles. Furthermore, by integrating targeting motifs into the outer surface of liposomes, cell-specific membrane engineering can be achieved for potential targeted drug delivery.

3.3 Electrostatic interaction

Cell surface modification through electrostatic interactions is an appealing strategy that capitalizes on the negative charge conferred mainly by sialic acid residues in the carbohydrate layer, along with the phosphatidylserine on the plasma membrane. Cationic polymers such as polyethyleneimine (PEI), poly-L-lysine (PLL) and chitosan (CS) are often utilized in the strategy.^{122–126} For instance, Choi and colleagues achieved the development of silica coating on mammalian cells by modifying PEI on the cell membrane through electrostatic interactions, serving as a catalytic template for silicification.¹²⁶ In a subsequent study, TiO₂ shells were developed for the cytoprotective encapsulation of Jurkat T cells.¹²² This method could effectively protect the T cells in the shell while simultaneously preserving their functionality, including cell division, juxtacrine interactions and cytokine secretion. Upon administration into the organism, the lymphocytes' therapeutic capabilities are effectively reinstated through the rupture of the protective shell. The TiO₂-inducing peptide, (RKK)₄D₈ (R: arginine, K: lysine, D: aspartic acid), was deposited on the surface of Jurkat cells *via* electrostatic interactions to facilitate the formation of bio-inspired TiO₂ using titanium bis(ammonium lactato)dihydroxide (TiBALDH) as a precursor. However, interactions with most cationic polymers readily lead to the destruction of the cell membrane, resulting in pronounced cytotoxicity and cellular damage. To address this issue, cationic polymers can be modified with biocompatible molecules, such as grafting PEG or alginate, to mitigate the detrimental effects on cell viability.¹²⁷





Fig. 6 Methods based on layer-by-layer self-assembly. (A) Schematic illustration of MSCs nanofilms prepared using positively charged PLL, negatively charged HA and RGD, and the functions exhibited by the modified cells.¹²⁸ Copyright 2017, American Chemical Society. (B) Schematic illustration of mammalian cell nanoencapsulation conducted by LbL self-assembly between GA and GB, and subsequent thiol-maleimide reaction. GSH could be added for on-demand release.¹²⁹ Copyright 2017, Elsevier.

Despite the overall negative charge of the cell membrane surface, a few cationic sites on the plasmalemma still exist which can be modified with negatively charged materials.^{130,131}

For instance, Thomsen *et al.* modified T cells with negatively charged degradable poly(lactic acid) (PLA) nanoparticles with electrostatic adsorption.¹²⁵ Furthermore, the modification of



negatively charged materials can be achieved through a synergistic combination of electrostatic interactions, hydrogen bonding and hydrophobic interactions.^{132–134} It is important to note that the presence of a negatively charged cell membrane hinders the uptake of negatively charged nanoparticles by cells.

3.4 Layer-by-layer (LBL) self-assembly

LBL self-assembly strategies have been developed on the basis of electrostatic interaction and widely employed for the construction of cell coatings, in which oppositely charged materials are sequentially deposited onto the cell membrane through electrostatic interaction along with hydrogen bonding, van der Waals forces, *etc.*^{21,127,128,135–147} As shown in Fig. 6A, Hong and colleagues developed LBL self-assembled nanofilms for cell surface modification of viable MSCs. Positively charged PLL was layer-by-layer assembled with negatively charged hyaluronic acid (HA) and arginine-glycine-aspartic acid (RGD) to fabricate nanofilms, which not only provided biochemical signals but also offered mechanical support for MSCs without interfering with the stemness of MSCs.¹²⁸ Subsequent studies have demonstrated the successful construction of nanofilms on the surface of human induced pluripotent stem cells (iPSCs) and immune cells such as AML-12 cells and peripheral blood mononuclear cells (PBMCs) *via* the LBL self-assembly strategy.^{140,148} Gels can be formed *via* LBL to coat or encapsulate cells. Chen and colleagues proposed a gentle approach (Fig. 6B) to achieve the nanoencapsulation of individual mammalian cells.¹²⁹ The gelatin coatings, which mimic the extracellular matrix (ECM), are formed through LBL self-assembly between positively charged gelatin type A (GA) and negatively charged gelatin type B (GB) on the cell membrane surface. Additionally, the outer layer of PEG was further constructed using thiol-maleimide click chemistry which could be degraded on-demand by the addition of the reducing agent glutathione (GSH). Subsequent studies involved the development of an enzyme-responsive nano-coating for encapsulating individual living cells, which was prepared through layer-by-layer self-assembly of oppositely charged gelatin-poly(ethylene glycol)maleimide and the incorporation of cysteine-terminated peptide sequences (CGGPLGLAGGC) *via* click reaction.¹³⁵ Moreover, the peptide chain could undergo enzymolysis upon exposure to high concentrations of matrix metalloproteinase-7 (MMP-7), which is frequently overexpressed in tumors, leading to the release of encapsulated cells.

4. Bioorthogonal modification

Despite the abundance of functional groups on the surface of the cell membrane that are amenable to chemical covalent modification, the utilization of non-specific covalent modification strategies may have detrimental effects on the viability and functionality of normal cells. Additionally, physical strategies are limited by the short residence time of synthetic molecules. In contrast, bioorthogonal chemistry offers a highly efficient and selective approach that takes place within a mild physiological environment, without disrupting intrinsic

biochemical processes. This strategy represents a substantial advancement in terms of both cell viability and the stability of modifications. Here, we provide an overview of recent developments in the integration of metabolism with copper-free click chemistry, Halo-Tag proteins, and enzyme-mediated approaches.

4.1 Metabolic glycan labeling strategy

Since the pioneering work of Bertozzi and colleagues, who introduced exogenous glycans into the cell membrane glyco-calyx, there has been a gradual development of strategies for metabolic glycan labeling to modify the cell membrane.^{149,150} In this strategy, unnatural sugars containing functional groups such as azide, alkyne, thiol and alkene are internalized by cells and the chemically reactive functional groups are “installed” on the glycan residues on the cell membrane by metabolic pathways.^{150–156} In recent years, glycans containing azide groups represented by *N*-azidoacetylmannosamine-tetraacetate (Ac₄-ManNAz) have gained the most widespread adoption with the development of copper-free “click” azide-alkyne reactions due to their high selectivity, synthetic simplicity and commercial availability. Tomas and colleagues have conducted a series of investigations on the cell surface modification with polymers *via* the metabolic glycan labeling strategy.^{157–159} In a recent study, Tomas *et al.* who proposed the “engineering cells to capture polymers” strategy incubated tumor cells with Ac₄-ManNAz for 96 hours to obtain azido-modified cancer cells which could capture chemotherapeutic polymers covalently and this strategy significantly augmented the concentration specifically targeted towards the tumor cell membrane whilst optimizing therapeutic efficacy by reducing systemic toxicity and enhancing selectivity.¹⁵⁸ In addition to polymers, the metabolic glycan labeling method effectively facilitates the modification of nanoparticles on the live cell membrane which is an attractive strategy for drug delivery.^{160–162} Zhou *et al.* successfully modified an oligomeric proanthocyanidin loaded liposome on the membrane of MSCs (MSC-Lipo-OPC) *via* metabolic labeling combined with the click chemistry strategy (Fig. 7A).¹⁶⁰ The MSC-Lipo-OPC could control the progression of inflammation due to the excellent abilities to scavenge free radicals and effectively prevent the formation of radiation-induced pulmonary fibrosis. Chen *et al.* developed polyvalent spherical aptamer (PSA) engineered macrophages which could effectively recognize tumor cells and inhibit tumor growth.¹⁶¹ PSA which has superior affinity and specificity to tumor cells was constructed through covalent reaction of gold nanoparticles (AuNPs) with AS1411 aptamer and DBCO groups, and was subsequently modified on macrophage membranes *via* metabolic labeling. Moreover, Lamoot *et al.* developed a 2-step click strategy for achieving highly specific cell surface conjugation of nanoparticles. In the study, cells were incubated with *N*-azidoacetylmannosamine-tetraacetylated (Ac₄ManN₃) to present azido groups on cell membrane (Fig. 7B).¹⁶² Subsequently, sulfo-6-methyl-tetrazine-dibenzyl cyclooctyne (Tz-DBCO) was exploited as a “bridge” between azide-modified cells and *trans*-cyclooctene (TCO) functionalized nanoparticles





Fig. 7 Methods based on the metabolic glycan labeling strategy. (A) Schematic illustration of binding Lipo-OPC to MSCs which were pre-incubated with Ac₄ManNAz and presented azido groups on the cell membrane.¹⁶⁰ Copyright 2023, Elsevier. (B) Overview of the 2-step click strategy for achieving highly specific cell surface conjugation of nanoparticles.¹⁶² Copyright 2020, John Wiley & Sons, Inc.

to realize nanoparticle-engineered cells exhibiting extremely low non-specific background binding.

In addition to *in vitro* applications for modifying cell membranes, researchers have also conducted studies to achieve this process *in vivo*.^{163,164} Wang *et al.* conducted an interesting study by labeling and modulating DCs and regulating DC-T cell

interactions *in vivo*.¹⁶³ They synthesized Ac₄ManAz nanoparticles overcame the limitations of Ac₄ManAz utilized *in vivo* such as poor encapsulation and water solubility. The Ac₄ManAz nanoparticles and granulocyte-macrophage colony-stimulating factor (GM-CSF) were loaded into an injectable alginate gel for the purpose of realizing *in situ* recruitment and azide labeling of



dendritic cells. Dibenzocyclooctyne (DBCO)-labelled immunomodulatory agents, such as tumor antigens, adjuvants, and cytokines could be modified on the DC membrane *via* click chemistry *in vivo* thereby effectively enhancing the subsequent T cell activation and tumor killing process. Additionally, Tu *et al.* employed Ac₄ManAz nanoparticles for *in situ* labeling of tumor cell membranes with azido groups, followed by the binding of chlorin e6 (Ce6), a commonly used photosensitizer, *via* click chemistry.¹⁶⁴ This approach effectively enhanced the therapeutic efficiency of photodynamic therapy. Recently, Chen and colleagues reported a cell-type-specific labeling approach *in vivo*.¹⁶⁵ In this study, the cardiomyocyte was specifically labeled without any interference from other cardiac cell types, which provided a powerful tool for cell-type selective modification. Gong *et al.* accomplished *in situ* PEGylation of CAR-T cells through the utilization of the metabolic glycan labeling strategy.¹⁶⁶ When the molecular weight of PEG reached 600 000, it effectively hindered the intercellular interactions among CAR-T cells, tumor cells, and monocytes, thereby attenuating the secretion of cytotoxic cytokines and ameliorating the symptoms associated with cytokine release syndrome (CRS).

Compared to the direct covalent binding with functional groups on the cell membrane, the metabolic glycan labeling

strategy effectively enhances the density of reactive sites on cells, but it is a time-consuming process that can take several days. The significant advantage of metabolic glycan labeling combined with bioorthogonal reactions is that it transits cell surface modification from nonspecific to specific, enabling cell surface modification *in situ* and *in vivo*. This is still an emerging field, with immense potential for further development and expansion.

4.2 Halo-Tag protein

Halo-Tag protein (HTP) is an engineered protein derived from the bacterial haloalkane dehalogenase, which selectively reacts with alkanes containing a terminal chloride group (chloroalkanes) forming a covalent bond.^{167,168} Similar protein recognition tags, such as SNAP tags¹⁶⁹ and ACP tags,¹⁷⁰ could also be utilized in cell surface modification, but they will not be extensively discussed in this section. A two-step approach was utilized in the HTP strategy: the expression of HTP on the cellular membrane is achieved *via* genetic engineering methods and further combined with cargoes containing the chloroalkane. HTP is commonly used in protein isolation and purification, molecular imaging, molecular interactions *etc.* in most reported studies and was first utilized for cell surface



Fig. 8 Methods based on other bioorthogonal strategies. (A) Schematic illustration of displaying synthetic glycopolymers on HeLa cell membranes using HTP anchors.¹⁷² Copyright 2019, American Chemical Society. (B) Schematic illustration of Kell C-terminal sortase labeling with GGG-carrying antigen peptides.¹⁷³ Copyright 2017, National Academy of Sciences. (C) Schematic illustration of transferring biomacromolecules to glycocalyx on the surface of living cells with fucosyltransferase.¹⁷⁴ Copyright 2018, American Chemical Society.



modification by Pulsipher *et al.*¹⁷¹ They proposed a long-lived cell membrane engineering strategy utilizing HTP as an anchor for modifying embryonic stem cells (ESCs) with heparan sulfate (HS), which was covalently modified on the ESC membrane and stayed for more than one week.

Subsequently, our group developed a series of studies *via* the HTP strategy.^{172,175,176} Tumor cells were modified with specific glycopolymers *via* the HTP fusion technique combined with RAFT polymerization (Fig. 8A). The glycopolymers that were modified on tumor cells could bind to lectins on dendritic cells or macrophages which effectively enhanced the tumor immune response.¹⁷² An interesting discovery was that the migration of the tumor cells modified with glycopolymers could be affected. Specifically, compared with the unmodified tumor cells, the migration direction was altered and diffusion slowed down which offered novel insights pertaining to the management of cancer metastasis.¹⁷⁵ A following study was carried out and we constructed glycopolymers modified DCs *via* the HTP strategy. Enhanced interactions were discovered between glycopolymer modified DCs and T cells which effectively promoted the T cell activation and proliferation, providing a novel approach to designing more efficient DC vaccines.¹⁷⁶

The HTP strategy for cell surface modification is still in its infancy. It is noteworthy due to the strong stability of binding between HTP expressed on the cell membrane and its corresponding ligand, thereby enabling sustained modifications that persist for over a week, offering a suitable method for long-time and stable cell surface modification. However, the implementation of HTP expression necessitates the manipulation of gene transfection, a process that is intricate and time-consuming, thereby inapplicable to certain challenging-to-transfect cell types such as primary cells.

4.3 Enzyme-mediated strategy

Enzyme-mediated modification of cell membranes represents a novel approach for *in situ* modification of candidate materials, reacting with pre-existing structures on the cell surface under the specific catalysis of enzymes. Specifically, certain enzymes such as oxidoreductases (galactose oxidases¹⁷⁷), glycosyltransferases (sialyltransferases,^{178,179} galactosyltransferases, *N*-acetyl-glucosaminyl transferases and fucosyltransferases^{174,180}), transpeptidases (butelases and sortases^{173,181,182}), transglutaminases (TGases¹⁸³) *etc.* have been utilized for the modification of cell membranes, representing an appealing approach due to their remarkable specificity and high yield. For example, galactose oxidases can specifically convert the endogenous terminal galactoses or *N*-acetylgalactosamine residues on the cell surface into aldehyde groups, facilitating subsequent reactions between aldehyde groups and aminoxy-functional molecules.¹⁷⁷ Glycosyltransferases are primarily utilized for the modification of pre-existing sugars on cell membranes, thereby facilitating the introduction of non-natural sugars. Moreover, it's worth noting that in comparison to metabolic engineering approaches, glycosyltransferases, particularly sialyltransferases and fucosyltransferases, offer a novel method for introducing greater kinds and intricacy of sugars on the cell membrane

surface. In the presence of transpeptidases, molecules bearing recognition motifs can be directly conjugated to either the N or C termini of membrane proteins. For example, as shown in Fig. 8B, Pishesha *et al.* reported a strategy for inducing antigen-specific tolerance by utilizing the transpeptidase sortase to covalently conjugate disease-associated autoantigens onto red blood cells (RBCs), thereby attenuating the contribution of major subsets of immune effector cells to immunity in an antigen-specific manner.¹⁷³ Li *et al.* focused on fucosyltransferase and transferred bio-macromolecules to the glyco-calyx on the surface of living cells, which represented faster speed, better biocompatibility, and less interference to cells (Fig. 8C).¹⁷⁴ Through this method, they constructed two antibody-cell conjugates, which exhibited significant improvements in the process of targeting and killing anti-cancer immune responses.

5. Conclusion and outlook

Modifying cell surfaces with tailor-made and well-characterized synthesized molecules can effectively introduce novel functionalities or manipulate cells. This offers a powerful tool to overcome challenges encountered in cell-based biomedical applications. In this review, we present a comprehensive overview of the latest advances in cell surface modification using synthetic molecules. We summarize the typical strategies, including chemical covalent modifications, physical alterations, and bioorthogonal approaches (Table 1), along with the advantages, disadvantages, and applicable conditions of each strategy. The chemical covalent strategy offers a straightforward and versatile approach for achieving stable and long-lasting surface modification.¹⁸⁴ However, the strategy has the potential to adversely impact cell activity and functionality. The physical modification strategy provides a non-invasive and cytocompatible approach. However, modifications achieved through physical interactions, such as electricity and hydrophobicity, are relatively short-term and unstable. It is important to note that the two methods mentioned above are non-specific, lacking precision in cell surface modification and potentially increasing the risk of adverse effects during practical applications. Therefore, bioorthogonal chemistry provides a valuable strategy for the selective and highly biocompatible incorporation of synthetic molecules onto cell surfaces, even enabling cell surface modification *in vivo* – a remarkable development. However, the approaches used to introduce bioorthogonal groups, whether *via* genetic engineering or metabolic engineering, are time-consuming.

Despite notable advancements in the utilization of synthetic compounds for cell surface modification, there remain unresolved challenges and prospects for further investigation. One such challenge pertains to the inherent detrimental impact of exogenous synthetic compounds bound to the cell surface on cellular functionality, albeit with varying degrees of severity. Hence, it is of utmost importance to meticulously choose a suitable strategy for modifying cells, taking into consideration the particular cell type and application scenarios. Subsequently, it becomes imperative to assess and describe the condition of



Table 1 A summary of the advancements in strategies for the cell surface modification with synthetic molecules in this review

	Strategy	Cell type ^a	Synthetic molecules modified on the cell surface
Chemical covalent modification strategies	Amine groups (-NH ₂)	HUVECs	PEG, ²¹ biotin-streptavidin ²²
		C2C12 cells	Poly(methacryloyloxy)ethyltrimethylammonium chloride (PMETAC) ²²
		MSCs	Heterodimerizing leucine, ²³ chitosan nanoparticles, ²⁴ sialyl Lewis X ²⁹
		RBCs	Biotin-streptavidin, ²⁶ PEG ^{30,31}
	Thiol groups (-SH)	T cells	Nanoparticles ³²
		Jurkat and NK cells	Biotin-streptavidin ²⁷
		RBCs	Nanoparticles ³⁶
		DCs	Bovine serum albumin (BSA) protein shells ⁴¹
	Vicinal diol groups	B16 cells	BSA protein shells ^{41,43}
		T cells	Nanoparticles ^{33,34,38,39}
		L929 cells	Fluorescent polymer ⁴⁴
		MCF-7, HepG2 and HeLa cells	Pyrene derivative ³⁷
	Carboxyl groups Convert diols into aldehydes	B16 cells	Gadolinium DO3A amide ⁴⁸
		HDFs	DEGMA or NVP polymers ⁴⁹
HepG2 cells		BSA nanoparticles ⁵⁰	
B16, AB22 and ZL34 cells		Contrast agents (Gd- and ⁶⁸ Ga-DOTA-EN) ⁵²	
PC-3, DU145 and Jurkat cells		Sialic acid-imprinted fluorescent core-shell SiO ₂ particles ⁵³	
MCF-7, HeLa and PC-3 cells		Polymer nanoparticles ⁵⁴	
RBCs		Supramolecular polymers ⁵⁸	
MCF-7 cells		Reactive probe ⁵⁹	
HDFs		Biotin-streptavidin ⁶⁰	
HT29 and MDA-MB-231 cells		Maltol hydrazide ⁶¹	
Convert disulfide bonds (S-S) to thiol groups	HepG2 cells	Dendrimer hydrazides ⁶²	
	MCF-7 cells	PEI conjugated with multiple hydrazide groups ⁶³	
	HeLa cells	Peptide and protein ⁶⁴	
	Human iPSC-derived-MSCs	<i>p</i> -Benzoquinone/ethylenediamine polymer ⁶⁵	
	HepG2 cells	ECM coating ⁶⁶	
	HeLa cells	DNA bridge complex-templated silver nanoclusters (DNA bridge-AgNCs) ⁶⁷	
	NEs	Fluorescent dye, polymer, and nanoparticles ⁶⁹	
	HUVECs and HSFs	Liposomal stimulator of interferon genes (STING) agonists ⁷⁰	
	T cells	PEG ²¹	
	DCs and PC-3 cells	Tetrazine ⁷³	
Physical modification strategies	Hydrophobic insertion	CCRf-CEM cells, splenocytes, melanoma, human MSCs and beta cells	Cucurbit[7]uril-based supramolecular polymer ⁷⁴
		RAW264.7 cells and L-O2 cells	Synthetic peptide ⁷⁶
		Human T cells and B cells, erythrocytes, hepatocytes, L929 and HEK293T cells	β-Cyclodextrin (β-CD) and adamantane (ADA) ⁷⁷
		MCF7, A549, LUDLU-1 cells	Cell-penetrating peptide ⁷⁹
	Hydrophobic insertion	HeLa cells	Aptamer ⁸⁰
		MCF-7 cells	DNAzyme, ⁸² DNA, ⁸⁹ multicomponent polymer, ⁹¹ aptamer, ⁸⁰ core-shell upconversion nanoparticles ⁹²
		SubT1 cells	Fluorescein isothiocyanate (FITC), ⁸³ core-shell upconversion nanoparticles ⁹²
		MDA-MB-231 cells	Biotin-PEG ⁸⁴
		NK cells, ⁸¹ Jurkat, NK, and Ramos cells, ⁸⁶ CCRf-CEM cells, ⁸⁷ U937 cells, ⁸⁸ RBCs, ¹⁰⁷ lymphocytes ¹⁰⁶	Nitrilotriacetic acid (NTA) ⁸⁵
		MSCs	DNA
Hydrophobic insertion	Th9 cells, 4T1 cells	Hyperbranched polyglycerol (HPG) covalently modified with vasculature binding peptides (VBPs) ⁹³	
	Jurkat cells	Peptides ^{94,95}	
	3T3 cells	ssDNA ⁹⁷	
			Poly(oxanorbornene) block copolymers, ⁹⁸ PEG ¹⁰⁴



Table 1 (Contd.)

	Strategy	Cell type ^a	Synthetic molecules modified on the cell surface
Bioorthogonal strategies	Electrostatic interaction	HepG2 cells	Peptides, ⁹⁹ PEG ¹⁰⁰
		RIN cells	PEG ¹⁰¹
		Fibroblast cells	Ketone and oxyamine, ^{110,116,118} dialdehyde, ¹¹² hydroquinone ¹¹⁷
		NK cells	Lewis X trisaccharide ¹¹⁹
		T cells	2,2,6,6-Tetramethylpiperidine groups ¹²⁰
		HeLa cells	DNA ¹⁰⁸
		HUVEC	Poly-L-lysine (PLL) ²¹
		Jurkat cells	TiO ₂ coating ¹²²
		hASCs	Chitosan derivative ¹²⁴
		HeLa, 3T3, and Jurkat cells	Poly(ethyleneimine) (PEI) ¹²⁶
	Layer-by-layer (LBL) self-assembly	T cells	PLA nanoparticles ¹²⁵
		HeLa cells	Cationic gelatin and anionic gelatin ^{129,135}
		MSCs	PLL, hyaluronic acid (HA), and arginine-glycine-aspartic acid (RGD), ¹²⁸ gelatin, ¹³⁵ silk fibroin polyelectrolyte ¹³⁹
		RBCs	Chitosan- <i>graft</i> -phosphorylcholine, HA and PLL-PEG ¹³⁶
		DPCs	Gelatin and alginate ¹³⁷
		MIN6 cells	Gelatin and fibronectin (FN) ¹³⁸
		L929 cells	Silk fibroin polyelectrolyte ¹³⁹
		AML-12 cells and PBMCs	PLL and HA ¹⁴⁰
		T cells	Chitosan and alginate ¹⁴⁷
		iPSCs	FN, heparin (Hep), and chondroitin sulfate (CS) ¹⁴⁸
Metabolic glycan labeling strategy	Jurkat cells	Probes, ¹⁵¹ lipid nanoparticles, ¹⁶² polymer-antibody conjugates ¹⁵⁶	
	HEK 293T cells	Probes ¹⁵⁴	
	MCF-7 cells	β-CD and aptamer ¹⁵⁵	
	A549 cells	Poly(hydroxyethyl acrylamide), ¹⁵⁷ polycation, ¹⁵⁸ poly(hydroxyethyl acrylamide) (pHEA) ¹⁵⁹	
	MCF-7 cells	Polycation ¹⁵⁸	
	MSCs	Antioxidant liposome ¹⁶⁰	
	Macrophages	Polyvalent spherical aptamer ¹⁶¹	
	DCs	Antigens, adjuvants and cytokines ¹⁶³	
	4T1 cells	Chlorin 6 (ref. 164)	
	HeLa cells	Probes ¹⁸³	
Halo-Tag protein	B16 cells, ¹⁷⁵ DCs, ¹⁷⁶ HeLa cells ¹⁷²	Glycopolymers	
	Enzyme-mediated strategy	B cells	Biotin ¹⁷⁷
		NK cells	Bio-macromolecules, ¹⁷⁴ Sialyl Lewis X and CD22-specific ligands ¹⁸⁵
	T cells	Bio-macromolecules ¹⁷⁴	
	Mouse lung endothelial cells	Synthetic ligands and biotin ¹⁷⁸	
	CHO cells	Probes, ¹⁸¹ bio-macromolecules ¹⁷⁴	
	RBCs	Probes, ¹⁸² peptide ¹⁷³	
	HEK 293T cells	Probes ¹⁸¹	
	HeLa cells	Probes ¹⁸³	

^a Abbreviation for the full cell name or cell species: HUVECs: human umbilical vein endothelial cells; C2C12 cells: murine myoblasts; MSCs: mesenchymal stem cells; RBCs: red blood cells; Jurkat cells: acute T-cell leukemia cells; NK cells: natural killer cells; DCs: dendritic cells; B16 cells: murine melanoma cells; L929 cells: mouse fibroblast cells; MCF-7 cells: human breast adenocarcinoma cells; HepG2 cells: human liver cancer cells; HeLa cells: human cervical cancer cells; HDFs: human dermal fibroblasts; AB22 cells: mouse mesothelioma cells; ZL34 cells: human mesothelioma cells; PC-3 and DU145 cells: human prostate cancer cells; iPSCs: induced pluripotent stem cells; HT-29: human colon cancer cells; MDA-MB-231 cells: human breast cancer cells; NEs: neutrophils; HSFs: human skin fibroblasts; CCRF-CEM cells: human acute lymphoblastic leukemia cells; RAW 264.7 cells: mouse leukemia cells of monocyte macrophages; L-O2 cells: human normal liver cells; HEK293T cells: human embryonic kidney cells; A549 cell: human lung cancer cells; LUDLU-1 cells: human Caucasian lung squamous carcinoma cells; SubT1 cells: human CD4 expressing T-lymphoblastoid cells; U937: histiocytic lymphoma cells; Th9 cells: helper T cell 9; 4T1 cells: mouse breast cancer cells; 3T3 cells: mouse embryonic fibroblasts; RIN cells: mouse insulinoma cells; hASCs: human adipose-derived mesenchymal stem cells; DPCs: dermal papilla cells; MIN6 cells: pancreatic β-cells; AML-12 cells: human lung cancer cells; PBMCs: peripheral blood mononuclear cells; CHO cells: Chinese hamster ovary cells.

the modified cells, encompassing cell viability, phenotype, and associated functionalities. It is noteworthy that the cell surface constitutes a dynamic membrane structure, wherein synthetic

molecules may undergo endocytosis or excretion by the cell. Consequently, it is crucial to monitor the destiny of synthesized molecules during and post cell surface modification. In



practical scenarios, the presence of synthetic molecules on cellular surfaces carries the potential for immune activation and subsequent clearance by the immune system, thereby considerably restricting their capacity to modify cell surfaces *in vivo*. Consequently, it becomes crucial to implement suitable adjustments to synthetic molecules to ensure their compatibility with *in vivo* applications. Additionally, a significant hurdle lies in selecting and designing molecules that possess both biocompatibility and augmented functionality for specific applications. Determining the optimal chemical group, structure, and sequence becomes essential in this regard. Therefore, the availability of databases serving as a toolbox for researchers to facilitate informed molecule selection is highly desirable.

There are several possible avenues for future research. One potential area of exploration is the development of more precise and targeted methods for modifying cell surfaces. There is a need to develop more proficient and potent methodologies for the selective and highly biocompatible integration of synthetic molecules onto cell surfaces. Furthermore, additional research is necessary to gain a better understanding of the influence of synthetic molecules on cellular functionality and to optimize modification strategies tailored to specific cell types and applications. An additional area of research that holds promise for the future is the advancement of synthetic molecules that possess improved biocompatibility and biofunctionality, enabling their application in the modification of cell surfaces. Notable examples of these molecules encompass functional nucleic acids, targeting aptamers, and polymers characterized by well-defined structures and chain sequences. The utilization of artificial intelligence (AI) can be facilitated by the establishment of databases containing comprehensive information regarding ligand–receptor interactions specific to cells, as well as the attributes associated with each modification technique. This integration of AI can aid in the design of optimal, customized molecules and the selection of appropriate methods for modification.

Author contributions

Conceptualization: H. Chen, G. Chen and H. Yang; writing the original draft: H. Yang, L. Yao and Y. Wang; validation: H. Yang, L. Yao and Y. Wang; review and editing: H. Chen and G. Chen.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 L. Tang, Y. Zheng, M. B. Melo, L. Mabardi, A. P. Castaño, Y.-Q. Xie, N. Li, S. B. Kudchodkar, H. C. Wong, E. K. Jeng, M. V. Maus and D. J. Irvine, *Nat. Biotechnol.*, 2018, **36**, 707–716.
- 2 C. W. Shields, M. A. Evans, L. L.-W. Wang, N. Baugh, S. Iyer, D. Wu, Z. Zhao, A. Pusuluri, A. Ukidve, D. C. Pan and S. Mitragotri, *Sci. Adv.*, 2020, **6**, eaaz6579.
- 3 G. Q. Daley, *Cell Stem Cell*, 2012, **10**, 740–749.
- 4 M. A. Fischbach, J. A. Bluestone and W. A. Lim, *Sci. Transl. Med.*, 2013, **5**, 179ps177.
- 5 H.-R. Jia, Y.-X. Zhu, Q.-Y. Duan and F.-G. Wu, *Chem. Soc. Rev.*, 2021, **50**, 6240–6277.
- 6 L. Zheng, X. Hu, H. Wu, L. Mo, S. Xie, J. Li, C. Peng, S. Xu, L. Qiu and W. Tan, *J. Am. Chem. Soc.*, 2020, **142**, 382–391.
- 7 Z. Wu, B. Esteban-Fernández de Ávila, A. Martín, C. Christianson, W. Gao, S. K. Thamphiwatana, A. Escarpa, Q. He, L. Zhang and J. Wang, *Nanoscale*, 2015, **7**, 13680–13686.
- 8 D. Dapkute, M. Pleckaitis, D. Bulotiene, D. Daunoravicius, R. Rotomskis and V. Karabanovas, *ACS Appl. Mater. Interfaces*, 2021, **13**, 43937–43951.
- 9 M. H. Kershaw, J. A. Westwood and P. K. Darcy, *Nat. Rev. Cancer*, 2013, **13**, 525–541.
- 10 D. Zhang, Y. Zheng, Z. Lin, X. Liu, J. Li, H. Yang and W. Tan, *Angew. Chem., Int. Ed. Engl.*, 2020, **59**, 12022–12028.
- 11 H. Watson, *Essays Biochem.*, 2015, **59**, 43–69.
- 12 K. Adebowale, R. Liao, V. C. Suja, N. Kapate, A. Lu, Y. Gao and S. Mitragotri, *Adv. Mater.*, 2023, 2210059.
- 13 S. Yang, H. Choi, D. T. Nguyen, N. Kim, S. Y. Rhee, S. Y. Han, H. Lee and I. S. Choi, *Adv. Ther.*, 2023, **6**, 2300037.
- 14 S. Mansouri, Y. Merhi, F. M. Winnik and M. Tabrizian, *Biomacromolecules*, 2011, **12**, 585–592.
- 15 N. A. A. Rossi, I. Constantinescu, R. K. Kainthan, D. E. Brooks, M. D. Scott and J. N. Kizhakkedathu, *Biomaterials*, 2010, **31**, 4167–4178.
- 16 M. C. Arno, *Macromol. Rapid Commun.*, 2020, **41**, e2000302.
- 17 D. Wu, K. Yang, Z. Zhang, Y. Feng, L. Rao, X. Chen and G. Yu, *Chem. Soc. Rev.*, 2022, **51**, 1336–1376.
- 18 S. Roy, J. N. Cha and A. P. Goodwin, *Bioconjugate Chem.*, 2020, **31**, 2465–2475.
- 19 B. Kellam, P. A. De Bank and K. M. Shakesheff, *Chem. Soc. Rev.*, 2003, **32**, 327–337.
- 20 J. Park, B. Andrade, Y. Seo, M. J. Kim, S. C. Zimmerman and H. Kong, *Chem. Rev.*, 2018, **118**, 1664–1690.
- 21 F. Cai, Y. Ren, J. Dai, J. Yang and X. Shi, *Macromol. Biosci.*, 2023, **23**, e2200379.
- 22 W. Wang, Q. Gan, Y. Zhang, X. Lu, H. Wang, Y. Zhang, H. Hu, L. Chen, L. Shi, S. Wang and Z. Zheng, *Adv. Mater.*, 2021, **33**, e2102348.
- 23 N. E. Jasiewicz, K. C. Mei, H. M. Oh, P. Chansoria, D. A. Hendy, E. E. Bonacquisti, E. M. Bachelder, K. M. Ainslie, H. Yin, L. Qian, B. C. Jensen and J. Nguyen, *Adv. Healthcare Mater.*, 2023, **12**, e2201094.



- 24 M. Xu, S. Asghar, S. Dai, Y. Wang, S. Feng, L. Jin, F. Shao and Y. Xiao, *Int. J. Biol. Macromol.*, 2019, **134**, 1002–1012.
- 25 C. M. Csizmar, J. R. Petersburg and C. R. Wagner, *Cell Chem. Biol.*, 2018, **25**, 931–940.
- 26 X. Sun, X. Han, L. Xu, M. Gao, J. Xu, R. Yang and Z. Liu, *Small*, 2017, **13**, 1701864.
- 27 B. Wang, J. Song, H. Yuan, C. Nie, F. Lv, L. Liu and S. Wang, *Adv. Mater.*, 2014, **26**, 2371–2375.
- 28 P. Gong, W. Zheng, Z. Huang, W. Zhang, D. Xiao and X. Jiang, *Adv. Funct. Mater.*, 2013, **23**, 42–46.
- 29 D. Sarkar, P. K. Vemula, G. S. L. Teo, D. Spelke, R. Karnik, L. Y. Wee and J. M. Karp, *Bioconjugate Chem.*, 2008, **19**, 2105–2109.
- 30 A. J. Bradley, K. L. Murad, K. L. Regan and M. D. Scott, *Biochim. Biophys. Acta*, 2002, **1561**, 147–158.
- 31 M. D. Scott, K. L. Murad, F. Koumpouras, M. Talbot and J. W. Eaton, *Proc. Natl. Acad. Sci. U. S. A.*, 1997, **94**, 7566–7571.
- 32 B. Huang, W. D. Abraham, Y. Zheng, S. C. Bustamante López, S. S. Luo and D. J. Irvine, *Sci. Transl. Med.*, 2015, **7**, 291ra294.
- 33 M. T. Stephan, S. B. Stephan, P. Bak, J. Chen and D. J. Irvine, *Biomaterials*, 2012, **33**, 5776–5787.
- 34 M. T. Stephan, J. J. Moon, S. H. Um, A. Bershteyn and D. J. Irvine, *Nat. Med.*, 2010, **16**, 1035–1041.
- 35 C. Wang, W. Sun, Y. Ye, Q. Hu, H. N. Bomba and Z. Gu, *Nat. Biomed. Eng.*, 2017, **1**, 0011.
- 36 X. Wang, X. Meng, K. Mao, H. Chen, X. Cong, F. Liu, J. Wang, S. Liu, Y. Xin, G. Zhu, H. Tan, Y. G. Yang and T. Sun, *Biomaterials*, 2023, **300**, 122187.
- 37 T. Li and Y. Liu, *Anal. Chem.*, 2021, **93**, 7029–7036.
- 38 Z. Luo, L. Luo, Y. Lu, C. Zhu, B. Qin, M. Jiang, X. Li, Y. Shi, J. Zhang, Y. Liu, X. Shan, H. Yin, G. Guan, Y. Du, N. Cheng and J. You, *J. Nanobiotechnol.*, 2022, **20**, 261.
- 39 L. Wayteck, H. Dewitte, L. De Backer, K. Breckpot, J. Demeester, S. C. De Smedt and K. Raemdonck, *Biomaterials*, 2016, **77**, 243–254.
- 40 T. Li, W. Gao, J. Liang, M. Zha, Y. Chen, Y. Zhao and C. Wu, *Anal. Chem.*, 2017, **89**, 8501–8508.
- 41 L. He, H. Wang, Y. Han, K. Wang, H. Dong, Y. Li, D. Shi and Y. Li, *ACS Appl. Mater. Interfaces*, 2019, **11**, 47750–47761.
- 42 R. Bej, A. Ghosh, J. Sarkar, B. B. Das and S. Ghosh, *Chembiochem*, 2020, **21**, 2921–2926.
- 43 Y. Wen, Y. Liu, F. Guo, Y. Han, Q. Qiu, Y. Li, H. Dong, T. Ren and Y. Li, *Biomater. Sci.*, 2021, **9**, 973–984.
- 44 Q. Sun, G. Liu, H. Wu, H. Xue, Y. Zhao, Z. Wang, Y. Wei, Z. Wang and L. Tao, *ACS Macro Lett.*, 2017, **6**, 550–555.
- 45 Q. Zhang, Y. Zhang, Y. Zhao, B. Yang, C. Fu, Y. Wei and L. Tao, *ACS Macro Lett.*, 2015, **4**, 128–132.
- 46 Y. Qu, Y. Zheng, L. Yu, Y. Zhou, Y. Wang, Y. Zhang, Q. Yu and H. Chen, *Adv. Funct. Mater.*, 2020, **30**, 1906362.
- 47 M. N. Christiansen, J. Chik, L. Lee, M. Anugraham, J. L. Abrahams and N. H. Packer, *Proteomics*, 2014, **14**, 525–546.
- 48 C. Rajan, J. Seema, Y. W. Chen, T. C. Chen, M. H. Lin, C. H. Lin and D. W. Hwang, *Biomedicines*, 2021, **9**, 1459.
- 49 A. J. Amaral and G. Pasparakis, *Chem. Commun.*, 2015, **51**, 17556–17559.
- 50 J. Wang, W. Wu, Y. Zhang, X. Wang, H. Qian, B. Liu and X. Jiang, *Biomaterials*, 2014, **35**, 866–878.
- 51 H. Liu, Y. Li, K. Sun, J. Fan, P. Zhang, J. Meng, S. Wang and L. Jiang, *J. Am. Chem. Soc.*, 2013, **135**, 7603–7609.
- 52 J. Martinelli, L. Tei, S. Geninatti Crich, D. Alberti and K. Djanashvili, *Molecules*, 2021, **26**, 1730.
- 53 S. Shinde, Z. El-Schich, A. Malakpour, W. Wan, N. Dizeyi, R. Mohammadi, K. Rurack, A. Gyorloff Wingren and B. Sellergren, *J. Am. Chem. Soc.*, 2015, **137**, 13908–13912.
- 54 J. Kim, Y. M. Lee, H. Kim, D. Park, J. Kim and W. J. Kim, *Biomaterials*, 2016, **75**, 102–111.
- 55 J. Yan, G. Springsteen, S. Deeter and B. Wang, *Tetrahedron*, 2004, **60**, 11205–11209.
- 56 W. Chen, X. Zhen, W. Wu and X. Jiang, *Sci. China: Chem.*, 2020, **63**, 648–664.
- 57 B. Dhawan, G. Akhter, H. Hamid, P. Kesharwani and M. S. Alam, *J. Mol. Struct.*, 2022, **1252**, 132057.
- 58 G. Morgese, B. F. M. de Waal, S. Varela-Aramburu, A. R. A. Palmans, L. Albertazzi and E. W. Meijer, *Angew. Chem., Int. Ed. Engl.*, 2020, **59**, 17229–17233.
- 59 N. Ma, J. Hu, Z. M. Zhang, W. Liu, M. Huang, Y. Fan, X. Yin, J. Wang, K. Ding, W. Ye and Z. Li, *J. Am. Chem. Soc.*, 2020, **142**, 6051–6059.
- 60 J. Sinclair and A. K. Salem, *Biomaterials*, 2006, **27**, 2090–2094.
- 61 A. Ciupa, P. A. De Bank and L. Caggiano, *Chem. Commun.*, 2013, **49**, 10148–10150.
- 62 D. Zhao, S. M. Ong, Z. Yue, Z. Jiang, Y. C. Toh, M. Khan, J. Shi, C. H. Tan, J. P. Chen and H. Yu, *Biomaterials*, 2008, **29**, 3693–3702.
- 63 S. M. Ong, L. He, N. T. Thuy Linh, Y. H. Tee, T. Arooz, G. Tang, C. H. Tan and H. Yu, *Biomaterials*, 2007, **28**, 3656–3667.
- 64 X. Liu, Y. Wang, B. Ye and X. Bi, *Chem. Sci.*, 2023, **14**, 7334–7345.
- 65 B. B. Chen, X. Y. Wang and R. C. Qian, *Chem. Commun.*, 2019, **55**, 9681–9684.
- 66 I. Koh, I. Yong, B. Kim, D. Choi, J. Hong, Y. M. Han and P. Kim, *ACS Biomater. Sci. Eng.*, 2020, **6**, 813–821.
- 67 L. Li, B. Han, Y. Wang, J. Zhao and Y. Cao, *Biosens. Bioelectron.*, 2019, **145**, 111714.
- 68 J. Cha, H. Kim, N. S. Hwang and P. Kim, *ACS Appl. Mater. Interfaces*, 2018, **10**, 35676–35680.
- 69 H. Kim, K. Shin, O. K. Park, D. Choi, H. D. Kim, S. Baik, S. H. Lee, S. H. Kwon, K. J. Yarema, J. Hong, T. Hyeon and N. S. Hwang, *J. Am. Chem. Soc.*, 2018, **140**, 1199–1202.
- 70 M. Hao, L. Zhu, S. Hou, S. Chen, X. Li, K. Li, N. Zhu, S. Chen, L. Xue, C. Ju and C. Zhang, *ACS Nano*, 2023, **17**, 1663–1680.
- 71 P. Shi, X. Wang, B. Davis, J. Coyne, C. Dong, J. Reynolds and Y. Wang, *Angew. Chem., Int. Ed. Engl.*, 2020, **59**, 11892–11897.
- 72 J. Niu, D. J. Lunn, A. Pusuluri, J. I. Yoo, M. A. O'Malley, S. Mitragotri, H. T. Soh and C. J. Hawker, *Nat. Chem.*, 2017, **9**, 537–545.



- 73 M. Hao, S. Hou, W. Li, K. Li, L. Xue, Q. Hu, L. Zhu, Y. Chen, H. Sun, C. Ju and C. Zhang, *Sci. Transl. Med.*, 2020, **12**, eaaz6667.
- 74 F. Huang, J. Liu and Y. Liu, *Chem. Sci.*, 2022, **13**, 8885–8894.
- 75 C. Gao, Q. Cheng, J. Li, J. Chen, Q. Wang, J. Wei, Q. Huang, S. M. Y. Lee, D. Gu and R. Wang, *Adv. Funct. Mater.*, 2021, **31**, 2102440.
- 76 A. Yoshihara, S. Watanabe, I. Goel, K. Ishihara, K. N. Ekdahl, B. Nilsson and Y. Teramura, *Biomaterials*, 2020, **253**, 120113.
- 77 C. Gao, Q. Cheng, J. Wei, C. Sun, S. Lu, C. H. T. Kwong, S. M. Y. Lee, Z. Zhong and R. Wang, *Mater. Today*, 2020, **40**, 9–17.
- 78 E. Kim, J. C. Kim, K. Min, M. Goh and G. Tae, *ACS Appl. Mater. Interfaces*, 2018, **10**, 24431–24439.
- 79 Y. Teramura, S. Asif, K. N. Ekdahl, E. Gustafson and B. Nilsson, *ACS Appl. Mater. Interfaces*, 2017, **9**, 244–254.
- 80 M. O. Altman, Y. M. Chang, X. Xiong and W. Tan, *Sci. Rep.*, 2013, **3**, 3343.
- 81 X. Wang, C. Wen, B. Davis, P. Shi, L. Abune, K. Lee, C. Dong and Y. Wang, *ACS Appl. Mater. Interfaces*, 2022, **14**, 3900–3909.
- 82 R. C. Qian, Z. R. Zhou, W. Guo, Y. Wu, Z. Yang and Y. Lu, *J. Am. Chem. Soc.*, 2021, **143**, 5737–5744.
- 83 X. Chen, X. Zhang, H. Y. Wang, Z. Chen and F. G. Wu, *Langmuir*, 2016, **32**, 10126–10135.
- 84 W. Meier, *Langmuir*, 2000, **16**, 1457–1459.
- 85 Y. Zheng, T. Wegner, D. Di Iorio, M. Pierau, F. Glorius and S. V. Wegner, *ACS Chem. Biol.*, 2023, **18**, 1435–1443.
- 86 C. Lv, Y. Li, M. Zhang, Y. Cheng, D. Han and W. Tan, *Nano Lett.*, 2023, **23**, 1167–1174.
- 87 B. Davis, K. Lee, X. Wang and Y. Wang, *Biomacromolecules*, 2023, **24**, 3193–3202.
- 88 Y. Fang, Y. Yan, S. Bi, Y. Wang, Y. Chen, P. Xu, H. Ju and Y. Liu, *Anal. Chem.*, 2022, **94**, 13205–13214.
- 89 S. F. Jones, H. Joshi, S. J. Terry, J. R. Burns, A. Aksimentiev, U. S. Eggert and S. Howorka, *J. Am. Chem. Soc.*, 2021, **143**, 8305–8313.
- 90 L. Zhang, F. Wang, Q. Li, L. Wang, C. Fan, J. Li and Y. Zhu, *Nano Lett.*, 2020, **20**, 3521–3527.
- 91 Y. Gu, B. Liu, Q. Liu, Y. Hang, L. Wang, J. L. Brash, G. Chen and H. Chen, *ACS Appl. Mater. Interfaces*, 2019, **11**, 47720–47729.
- 92 Y. Kong, F. Yuan, F. Yang, C. Zhang and Y. Xian, *Anal. Chem.*, 2022, **94**, 12024–12032.
- 93 J. H. Jeong, J. J. Schmidt, R. E. Kohman, A. T. Zill, R. J. DeVolder, C. E. Smith, M. H. Lai, A. Shkumatov, T. W. Jensen, L. G. Schook, S. C. Zimmerman and H. Kong, *J. Am. Chem. Soc.*, 2013, **135**, 8770–8773.
- 94 T. Chen, Y. Xue, S. Wang, J. Lu, H. Zhou, W. Zhang, Z. Zhou, B. Li, Y. Li, Z. Wang, C. Li, Y. Eloy, H. Sun, Y. Shen, M. D. Diarra, C. Ge, X. Chai, H. Mou, P. Lin, X. Yu and Z. Ye, *Bioact. Mater.*, 2023, **23**, 508–523.
- 95 L.-H. Liu, W.-X. Qiu, Y.-H. Zhang, B. Li, C. Zhang, F. Gao, L. Zhang and X.-Z. Zhang, *Adv. Funct. Mater.*, 2017, **27**, 1700220.
- 96 C. Y. Lo, B. R. Weil, B. A. Palka, A. Momeni, J. M. Canty Jr and S. Neelamegham, *Biomaterials*, 2016, **74**, 19–30.
- 97 R. J. Weber, S. I. Liang, N. S. Selden, T. A. Desai and Z. J. Gartner, *Biomacromolecules*, 2014, **15**, 4621–4626.
- 98 D. C. Church and J. K. Pokorski, *Angew. Chem., Int. Ed. Engl.*, 2020, **59**, 11379–11383.
- 99 Z. Rao, M. Sasaki and T. Taguchi, *Colloids Surf., B*, 2013, **101**, 223–227.
- 100 T. Taguchi, Z. Rao, M. Ito and M. Matsuda, *J. Mater. Sci.: Mater. Med.*, 2011, **22**, 2357–2363.
- 101 M. Ito and T. Taguchi, *Acta Biomater.*, 2009, **5**, 2945–2952.
- 102 Y. Zhao, M. Fan, Y. Chen, Z. Liu, C. Shao, B. Jin, X. Wang, L. Hui, S. Wang, Z. Liao, D. Ling, R. Tang and B. Wang, *Sci. Adv.*, 2020, **6**, eaaw9679.
- 103 S. Sakai and M. Taya, *ACS Macro Lett.*, 2014, **3**, 972–975.
- 104 K. Kato, C. Itoh, T. Yasukouchi and T. Nagamune, *Biotechnol. Prog.*, 2004, **20**, 897–904.
- 105 L. Zhang, G. Liu, K. Lv, J. Xin, Y. Wang, J. Zhao, W. Hu, C. Xiao, K. Zhu, L. Zhu, J. Nan, Y. Feng, H. Zhu, W. Chen, W. Zhu, J. Zhang, J. Wang, B. Wang and X. Hu, *Adv. Sci.*, 2021, **8**, 2003348.
- 106 L. Sun, F. Shen, Z. Xiong, H. Yang, Z. Dong, J. Xiang, Q. Gu, Q. Ji, C. Fan and Z. Liu, *J. Am. Chem. Soc.*, 2022, **144**, 7634–7645.
- 107 L. Sun, F. Shen, J. Xu, X. Han, C. Fan and Z. Liu, *Angew. Chem., Int. Ed. Engl.*, 2020, **59**, 14842–14853.
- 108 M. Lin, Y. Chen, S. Zhao, R. Tang, Z. Nie and H. Xing, *Angew. Chem., Int. Ed. Engl.*, 2022, **61**, e202111647.
- 109 D. Sarkar, P. K. Vemula, W. Zhao, A. Gupta, R. Karnik and J. M. Karp, *Biomaterials*, 2010, **31**, 5266–5274.
- 110 D. Dutta, A. Pulsipher, W. Luo, H. Mak and M. N. Yousaf, *Bioconjugate Chem.*, 2011, **22**, 2423–2433.
- 111 D. Dutta, A. Pulsipher, W. Luo and M. N. Yousaf, *J. Am. Chem. Soc.*, 2011, **133**, 8704–8713.
- 112 S. Elahipanah, P. J. O'Brien, D. Rogozhnikov and M. N. Yousaf, *Bioconjugate Chem.*, 2017, **28**, 1422–1433.
- 113 W. Luo, A. Pulsipher, D. Dutta, B. M. Lamb and M. N. Yousaf, *Sci. Rep.*, 2014, **4**, 6313.
- 114 W. Luo, N. Westcott, D. Dutta, A. Pulsipher, D. Rogozhnikov, J. Chen and M. N. Yousaf, *ACS Chem. Biol.*, 2015, **10**, 2219–2226.
- 115 P. J. O'Brien, S. Elahipanah, D. Rogozhnikov and M. N. Yousaf, *ACS Cent. Sci.*, 2017, **3**, 489–500.
- 116 P. J. O'Brien, W. Luo, D. Rogozhnikov, J. Chen and M. N. Yousaf, *Bioconjugate Chem.*, 2015, **26**, 1939–1949.
- 117 A. Pulsipher, D. Dutta, W. Luo and M. N. Yousaf, *Angew. Chem., Int. Ed. Engl.*, 2014, **53**, 9487–9492.
- 118 D. Rogozhnikov, P. J. O'Brien, S. Elahipanah and M. N. Yousaf, *Sci. Rep.*, 2016, **6**, 39806.
- 119 C. Zheng, Q. Zhong, W. Song, K. Yi, H. Kong, H. Wang, Y. Tao, M. Li and X. Chen, *Adv. Mater.*, 2023, **35**, 2206989.
- 120 C. Shi, Q. Zhang, Y. Yao, F. Zeng, C. Du, S. Nijiati, X. Wen, X. Zhang, H. Yang, H. Chen, Z. Guo, X. Zhang, J. Gao, W. Guo, X. Chen and Z. Zhou, *Nat. Nanotechnol.*, 2023, **18**, 86–97.
- 121 Y. Chen, T. Wu, S. Xie, Y. Bai and H. Xing, *Sci. Adv.*, 2023, **9**, eadg2583.



- 122 W. Youn, E. H. Ko, M. H. Kim, M. Park, D. Hong, G. A. Seisenbaeva, V. G. Kessler and I. S. Choi, *Angew. Chem., Int. Ed. Engl.*, 2017, **56**, 10702–10706.
- 123 T. Thomsen and H. A. Klok, *ACS Appl. Bio Mater.*, 2021, **4**, 2293–2306.
- 124 M. M. Maciel, T. R. Correia, V. M. Gaspar, J. M. M. Rodrigues, I. S. Choi and J. F. Mano, *Adv. Funct. Mater.*, 2021, **31**, 2009619.
- 125 T. Thomsen, R. Reissmann, E. Kaba, B. Engelhardt and H. A. Klok, *Biomacromolecules*, 2021, **22**, 3416–3430.
- 126 J. Lee, J. Choi, J. H. Park, M. H. Kim, D. Hong, H. Cho, S. H. Yang and I. S. Choi, *Angew. Chem., Int. Ed. Engl.*, 2014, **53**, 8056–8059.
- 127 J. T. Wilson, W. Cui, V. Kozlovskaya, E. Kharlampieva, D. Pan, Z. Qu, V. R. Krishnamurthy, J. Mets, V. Kumar, J. Wen, Y. Song, V. V. Tsukruk and E. L. Chaikof, *J. Am. Chem. Soc.*, 2011, **133**, 7054–7064.
- 128 D. Choi, H. Lee, H.-B. Kim, M. Yang, J. Heo, Y. Won, S. S. Jang, J. K. Park, Y. Son, T. I. Oh, E. Lee and J. Hong, *Chem. Mater.*, 2017, **29**, 2055–2065.
- 129 J. Yang, J. Li, X. Li, X. Wang, Y. Yang, N. Kawazoe and G. Chen, *Biomaterials*, 2017, **133**, 253–262.
- 130 P. P. Wibroe, A. C. Anselmo, P. H. Nilsson, A. Sarode, V. Gupta, R. Urbanics, J. Szebeni, A. C. Hunter, S. Mitragotri, T. E. Mollnes and S. M. Moghimi, *Nat. Nanotechnol.*, 2017, **12**, 589–594.
- 131 A. Barbul, K. Singh, L. Horev–Azaria, S. Dasgupta, T. Auth, R. Korenstein and G. Gompper, *ACS Appl. Nano Mater.*, 2018, **1**, 3785–3799.
- 132 A. C. Anselmo, V. Gupta, B. J. Zern, D. Pan, M. Zakrewsky, V. Muzykantov and S. Mitragotri, *ACS Nano*, 2013, **7**, 11129–11137.
- 133 T. Thomsen, A. B. Ayoub, D. Psaltis and H. A. Klok, *Biomacromolecules*, 2021, **22**, 190–200.
- 134 S. Y. Han, H. Lee, D. T. Nguyen, G. Yun, S. Kim, J. H. Park and I. S. Choi, *ACS Appl. Bio Mater.*, 2021, **1**, 2000037.
- 135 J. Yang, Y. Yang, N. Kawazoe and G. Chen, *Biomaterials*, 2019, **197**, 317–326.
- 136 S. Mansouri, Y. Merhi, F. M. Winnik and M. Tabrizian, *Biomacromolecules*, 2011, **12**, 585–592.
- 137 J. Wang, Y. Miao, Y. Huang, B. Lin, X. Liu, S. Xiao, L. Du, Z. Hu and M. Xing, *Adv. Healthcare Mater.*, 2018, **7**, 1700447.
- 138 Y. Fukuda, T. Akagi, T. Asaoka, H. Eguchi, K. Sasaki, Y. Iwagami, D. Yamada, T. Noda, K. Kawamoto, K. Gotoh, S. Kobayashi, M. Mori, Y. Doki and M. Akashi, *Biomaterials*, 2018, **160**, 82–91.
- 139 O. Hasturk, J. K. Sahoo and D. L. Kaplan, *Biomacromolecules*, 2020, **21**, 2829–2843.
- 140 J. Hwang, D. Choi, M. Choi, Y. Seo, J. Son, J. Hong and J. Choi, *ACS Appl. Mater. Interfaces*, 2018, **10**, 17685–17692.
- 141 L. P. B. Guerzoni, Y. Tsukamoto, D. B. Gehlen, D. Rommel, T. Haraszti, M. Akashi and L. De Laporte, *Biomacromolecules*, 2019, **20**, 3746–3754.
- 142 A. Matsuzawa, M. Matsusaki and M. Akashi, *Langmuir*, 2013, **29**, 7362–7368.
- 143 J. Sun, Y. Ren, W. Wang, H. Hao, M. Tang, Z. Zhang, J. Yang, Y. Zheng and X. Shi, *ACS Biomater. Sci. Eng.*, 2020, **6**, 2336–2345.
- 144 K. Sasaki, T. Akagi, T. Asaoka, H. Eguchi, Y. Fukuda, Y. Iwagami, D. Yamada, T. Noda, H. Wada, K. Gotoh, K. Kawamoto, Y. Doki, M. Mori and M. Akashi, *Biomaterials*, 2017, **133**, 263–274.
- 145 W. Li, T. Guan, X. Zhang, Z. Wang, M. Wang, W. Zhong, H. Feng, M. Xing and J. Kong, *ACS Appl. Mater. Interfaces*, 2015, **7**, 3018–3029.
- 146 H. Lee, N. Kim, H. B. Rheem, B. J. Kim, J. H. Park and I. S. Choi, *Adv. Healthcare Mater.*, 2021, **10**, 2100347.
- 147 S. Zhao, L. Zhang, J. Han, J. Chu, H. Wang, X. Chen, Y. Wang, N. Tun, L. Lu, X. F. Bai, M. Yearsley, S. Devine, X. He and J. Yu, *ACS Nano*, 2016, **10**, 6189–6200.
- 148 U. Han, Y. J. Kim, W. Kim, J. H. Park and J. Hong, *Nanoscale*, 2019, **11**, 13541–13551.
- 149 L. K. Mahal, K. J. Yarema and C. R. Bertozzi, *Science*, 1997, **276**, 1125–1128.
- 150 R. M. F. Tomas and M. I. Gibson, *ACS Macro Lett.*, 2020, **9**, 991–1003.
- 151 T. L. Hsu, S. R. Hanson, K. Kishikawa, S. Wang, M. Sawa and C. H. Wong, *Proc. Natl. Acad. Sci. U. S. A.*, 2007, **104**, 2614–2619.
- 152 S. G. Sampathkumar, A. V. Li, M. B. Jones, Z. Sun and K. J. Yarema, *Nat. Chem. Biol.*, 2006, **2**, 149–152.
- 153 E. Saxon and C. R. Bertozzi, *Science*, 2000, **287**, 2007–2010.
- 154 A. K. Spate, J. E. Dold, E. Batroff, V. F. Schart, D. E. Wieland, O. R. Baudendistel and V. Wittmann, *ChemBiochem*, 2016, **17**, 1374–1383.
- 155 P. Shi, E. Ju, Z. Yan, N. Gao, J. Wang, J. Hou, Y. Zhang, J. Ren and X. Qu, *Nat. Commun.*, 2016, **7**, 13088.
- 156 A. Uvyn, R. De Coen, O. De Wever, K. Deswarte, B. N. Lambrecht and B. G. De Geest, *Chem. Commun.*, 2019, **55**, 10952–10955.
- 157 R. M. F. Tomas and M. I. Gibson, *Biomacromolecules*, 2019, **20**, 2726–2736.
- 158 R. M. F. Tomas and M. I. Gibson, *Chem. Sci.*, 2021, **12**, 4557–4569.
- 159 R. M. F. Tomas, B. Martyn, T. L. Bailey and M. I. Gibson, *ACS Macro Lett.*, 2018, **7**, 1289–1294.
- 160 H. Zhou, Y. Zhang, P. Pei, W. Shen, X. Yi and K. Yang, *Biomaterials*, 2023, **300**, 122202.
- 161 Y. Chen, P. Gao, W. Pan, M. Shi, S. Liu, N. Li and B. Tang, *Chem. Sci.*, 2021, **12**, 13817–13824.
- 162 A. Lamoot, A. Uvyn, S. Kasmi and B. G. De Geest, *Angew. Chem., Int. Ed. Engl.*, 2021, **60**, 6320–6325.
- 163 H. Wang, M. C. Sobral, D. K. Y. Zhang, A. N. Cartwright, A. W. Li, M. O. Dellacherie, C. M. Tringides, S. T. Koshy, K. W. Wucherpennig and D. J. Mooney, *Nat. Mater.*, 2020, **19**, 1244–1252.
- 164 Y. Tu, Y. Dong, K. Wang, S. Shen, Y. Yuan and J. Wang, *Biomaterials*, 2020, **259**, 120298.
- 165 X. Fan, Q. Song, D.-e. Sun, Y. Hao, J. Wang, C. Wang and X. Chen, *Nat. Chem. Biol.*, 2022, **18**, 625–633.



- 166 N. Gong, X. Han, L. Xue, R. El-Mayta, A. E. Metzloff, M. M. Billingsley, A. G. Hamilton and M. J. Mitchell, *Nat. Mater.*, 2023, DOI: [10.1038/s41563-023-01646-6](https://doi.org/10.1038/s41563-023-01646-6).
- 167 F. Chai, D. Cheng, Y. Nasu, T. Terai and R. E. Campbell, *Biochem. Soc. Trans.*, 2023, **51**, 1585–1595.
- 168 C. G. England, H. Luo and W. Cai, *Bioconjugate Chem.*, 2015, **26**, 975–986.
- 169 A. Keppler, S. Gendreizig, T. Gronemeyer, H. Pick, H. Vogel and K. Johnsson, *Nat. Biotechnol.*, 2003, **21**, 86–89.
- 170 N. George, H. Pick, H. Vogel, N. Johnsson and K. Johnsson, *J. Am. Chem. Soc.*, 2004, **126**, 8896–8897.
- 171 A. Pulsipher, M. E. Griffin, S. E. Stone and L. C. Hsieh-Wilson, *Angew. Chem., Int. Ed. Engl.*, 2015, **54**, 1466–1470.
- 172 Q. Liu, S. Jiang, B. Liu, Y. Yu, Z. A. Zhao, C. Wang, Z. Liu, G. Chen and H. Chen, *ACS Macro Lett.*, 2019, **8**, 337–344.
- 173 N. Pishesha, A. M. Bilate, M. C. Wibowo, N.-J. Huang, Z. Li, R. Deshycka, D. Bousbaine, H. Li, H. C. Patterson, S. K. Dougan, T. Maruyama, H. F. Lodish and H. L. Ploegh, *Proc. Natl. Acad. Sci. U. S. A.*, 2017, **114**, 3157–3162.
- 174 J. Li, M. Chen, Z. Liu, L. Zhang, B. H. Felding, K. W. Moremen, G. Lauvau, M. Abadier, K. Ley and P. Wu, *ACS Cent. Sci.*, 2018, **4**, 1633–1641.
- 175 L. Zhu, R. Feng, G. Chen, C. Wang, Z. Liu, Z. Zhang and H. Chen, *ACS Appl. Mater. Interfaces*, 2022, **14**, 4921–4930.
- 176 L. Yu, R. Feng, L. Zhu, Q. Hao, J. Chu, Y. Gu, Y. Luo, Z. Zhang, G. Chen and H. Chen, *Sci. Adv.*, 2020, **6**, eabb6595.
- 177 T. N. C. Ramya, E. Weerapana, B. F. Cravatt and J. C. Paulson, *Glycobiology*, 2013, **23**, 211–221.
- 178 C. J. Capicciotti, C. Zong, M. O. Sheikh, T. Sun, L. Wells and G. J. Boons, *J. Am. Chem. Soc.*, 2017, **139**, 13342–13348.
- 179 S. Hong, C. Yu, P. Wang, Y. Shi, W. Cao, B. Cheng, D. G. Chapla, Y. Ma, J. Li, E. Rodrigues, Y. Narimatsu, J. R. Yates III, X. Chen, H. Clausen, K. W. Moremen, M. S. Macauley, J. C. Paulson and P. Wu, *Angew. Chem., Int. Ed. Engl.*, 2021, **60**, 3603–3610.
- 180 T. Zheng, H. Jiang, M. Gros, D. Soriano del Amo, S. Sundaram, G. Lauvau, F. Marlow, Y. Liu, P. Stanley and P. Wu, *Angew. Chem., Int. Ed. Engl.*, 2011, **50**, 4113–4118.
- 181 T. Tanaka, T. Yamamoto, S. Tsukiji and T. Nagamune, *Chembiochem*, 2008, **9**, 802–807.
- 182 J. Shi, L. Kundrat, N. Pishesha, A. Bilate, C. Theile, T. Maruyama, S. K. Dougan, H. L. Ploegh and H. F. Lodish, *Proc. Natl. Acad. Sci. U. S. A.*, 2014, **111**, 10131–10136.
- 183 C. Lin and A. Y. Ting, *J. Am. Chem. Soc.*, 2006, **128**, 4542–4543.
- 184 U. S. Patil, H. Qu, D. Caruntu, C. J. O'Connor, A. Sharma, Y. Cai and M. A. Tarr, *Bioconjugate Chem.*, 2013, **24**, 1562–1569.
- 185 S. Hong, C. Yu, P. Wang, Y. Shi, W. Cao, B. Cheng, D. G. Chapla, Y. Ma, J. Li, E. Rodrigues, Y. Narimatsu, J. R. Yates III, X. Chen, H. Clausen, K. W. Moremen, M. S. Macauley, J. C. Paulson and P. Wu, *Angew. Chem., Int. Ed. Engl.*, 2021, **60**, 3603–3610.

