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A comprehensive analysis in one run – in-depth conformation studies of protein–polymer chimeras by asymmetrical flow field-flow fractionation †

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Polymer-based protein engineering has enabled the synthesis of a variety of protein–polymer conjugates that are widely applicable in therapeutic, diagnostic and biotechnological industries. Accurate characterizations of physical–chemical properties, in particular, molar masses, sizes, composition and their dispersities are critical parameters that determine the functionality and conformation of protein–polymer conjugates and are important for creating reproducible manufacturing processes. Most of the current characterization techniques suffer from fundamental limitations and do not provide an accurate understanding of a sample's true nature. In this paper, we demonstrate the advantage of asymmetrical flow field-flow fractionation (AF4) coupled with multiple detectors for the characterization of a library of complex, zwitterionic and neutral protein–polymer conjugates. This method allows for determination of intrinsic physical properties of protein–polymer chimeras from a single, rapid measurement.

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1 Introduction

Protein–polymer conjugates are exceptional macromolecules that possess characteristics of both biotic and abiotic worlds. Synthetic polymers are conjugated to proteins to enhance the native properties of the protein, improve activity and increase stability in non-native environments. This has been a significant area of interest since the synthesis of the first protein–polymer chimera reported in 1977. This conjugate, and most since, was generated by covalently attaching poly(ethylene) glycol (PEG) to bovine serum albumin (BSA).¹ PEGylation is used to shield the antibody binding epitopes within a protein, while also providing protection from proteolytic enzymes.^{2,3} Preventing antibody binding is a crucial step in evading the immune system thereby increasing the plasma circulation time. The

ability of a polymer to shield the surface of a protein from immune complex formation, led to medicinal application of protein–polymer conjugates.⁴

In recent years, considerable attention has been paid to the creation of “smart conjugates” by incorporating stimuli responsive polymers to proteins.^{5–7} Examples include temperature^{5,7,8} and pH-responsive⁹ polymers that can change conformation in response to trigger. “Smart conjugates” have been employed for sensing and detection.^{10–12} Polymer-based protein engineering has also been used to increase protein stability at high temperatures, pH extremes, in organic solvents and in ionic liquids. These properties are highly attractive for industrial applications of protein–polymer chimeras.

Protein–polymer conjugates can be synthesized using two alternative strategies: “grafting to” and “grafting from”. In the “grafting to” approach, end-functionalized polymers are synthesized first to a desired molar mass and then grafted covalently to a protein surface.¹³ This approach, while widely used, can be a somewhat random process where grafting density and attachment sites are difficult to control. Additionally, purification of these conjugates can be laborious due to the difficulties associated with the removal of unreacted polymers.^{13–15} Polymer synthesis has also evolved over the past few decades, with the development of controlled radical polymerization (CRP).^{16,17} Using CRP considerably widens the synthetic horizon, allowing the incorporation of wide range of monomers, introduce plethora of functional groups and monomers with complex architecture. Atom-transfer radical polymerization (ATRP) and reversible addition–fragmentation chain transfer (RAFT) have been shown to be ideal methods to prepare

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protein–polymer chimeras. “Grafting from” approach consists of modifying proteins with small initiator groups⁹ or chain transfer agents,¹⁸ followed by polymer synthesis using the protein as macroinitiator. Comparatively, “grafting from” allows the conjugates to be prepared with higher grafting densities,¹⁹ higher yield⁹ and finer modification site control.^{20,21} Additionally, CRP methods provide precise control of polymer composition, functionality and topology.²² Therefore, not only linear homopolymers, but also random copolymers, gradient copolymers and block copolymers as well as branched structures can be grown from proteins.⁸ Due to the process of growing polymers from a protein surface, simpler purification processes, such as dialysis, are also possible.

While the synthetic strategies to make protein–polymer conjugates have evolved over the past decade, the development of precise and accurate characterization techniques has lagged considerably behind. Synthetic advances cannot achieve the desired impact without reliable characterization of the products.²³ Due to their hybrid nature, the characterization of intact protein–polymer conjugates is very difficult, and we have had to rely on controlled decomposition and characterization of the resulting synthetic and biologic fragments. Thus, the synthetic polymer chains are typically cleaved from the protein surface, followed by molar mass and dispersity (D) characterization using size exclusion chromatography (SEC).⁹ Once the single chain's molar mass is determined and assuming that the number of polymers per protein is known, the whole conjugates molar mass can be estimated. This method has several drawbacks which may affect the molar mass calculation: (i) SEC molar masses are mostly estimated in reference to calibration standards, which may have different solution properties, conformation, architecture, (ii) different chemical structures may lead to different interaction with separation column's material than the cleaved polymer chains.²⁴ Absolute molar masses can be obtained by the application of multi-angle static light scattering detector (MALS). However, limited size range, shear degradation and sample interaction of column-based SEC are still existent (see ESI, Fig. S14†). Various complementary assay techniques, as well as batch dynamic light scattering (DLS) studies, are also used to provide information about the quality of conjugation. With these techniques, however, only average values are obtained, and the co-existence of complex protein–polymer structures remains undetected. Thus, novel and versatile techniques are needed to gain a deeper insight into the true nature of protein–polymer chimeras.

An attractive alternative to column-based SEC is the asymmetrical flow field-flow fractionation (AF4) as a channel-based separation technique that separates molecules depending on differences in their Brownian motion. This gentle technique does not use a densely packed column for separation and is non-destructive to delicate samples such as bioconjugates due to significantly reduced shear forces.^{25–28} Channel-based separation can more accurately reflect physical properties of the sample in solution. When coupling AF4 to MALS, the absolute, intrinsic molar mass can be obtained based on the Rayleigh ratio.^{29,30} Since MALS does not require the use of calibration standards, protein–polymer conjugate molar mass, radius and

D can be determined as intact entities, without cleaving the polymers prior to analysis (Fig. 1). MALS can provide information on radius of gyration (R_g) and combining that with hydrodynamic radius (R_h) that is obtained from online DLS detector, can further provide an insight into conformation properties such as scaling behavior and ρ parameter (R_g/R_h), as well as information about structural complexity.³¹ In previous studies, the great potential of AF4 in combination with up to five detectors for characterization of complex macromolecular architectures and particles has been demonstrated.^{28,32,33} Light scattering detectors in combination with concentration and chemical structure sensitive detectors (UV-Vis) also enable the determination of the chemical composition distribution. Previous studies show the potential of AF4-MALS for the separation and characterization of conjugated structures and enabled a detailed study of the complexity of avidin–polymer conjugation. These studies began to unravel a variety of co-existing architectures that were driven by changes of parameters such as type of polymer, degree of biotinylation, molar ratios, etc.^{25,34–36}

Herein, we demonstrate the potential of using AF4 with four-fold detection (AF4-D4, Fig. 1) for the characterization of avidin–polymer intrinsic physical properties. The library of polymers was prepared by using surface-initiated ATRP and variation of the number of polymer chains (grafting density), polymer length (degree of polymerization, DP) and polymer type. The comprehensive study of molecular properties like molar mass, radius and conformations by AF4-D4 enabled a correlation between the synthetic strategies and the physical properties of the resulting polymer–protein chimeras.

2 Experimental

2.1 Materials

Avidin from egg white was purchased from Lee Biosolutions (Maryland Heights, MO). Bicinchoninic acid solution, copper(II) chloride, sodium ascorbate and poly(ethylene glycol)methyl ether methacrylate (OEGMA, average M_n 500) were purchased from Sigma Aldrich (St. Louis, MO), CBMA was purchased from TCI (Portland, OR). Protein surface active ATRP initiators were prepared as described previously.^{9,19} 4-(Bis(*N,N*-diethylaminoethyl)aminoethyl)morpholine (MMA) synthesized in a previous report was used as ligand.

2.2 Synthetic procedures

2.2.1 Attachment of ATRP initiators on the surface of avidin. Single-headed ATRP initiator was synthesized as previously described.⁹ Avidin–Br conjugates were synthesized by mixing avidin (160 mg, 0.01 mmol protein, 0.1 mmol primary amine groups) with single-headed bromine-functionalized ATRP initiator (0.7 mmol, 235 mg (7 eq.)) and were dissolved in 80 mL of 0.1 M sodium phosphate buffer, pH 8.0. The reaction was stirred at 4 °C for 2 h and then dialyzed against 25 mM sodium phosphate, pH 8 using dialysis tubing with a molecular weight cut off of 15 kDa, for 24 h at 4 °C.



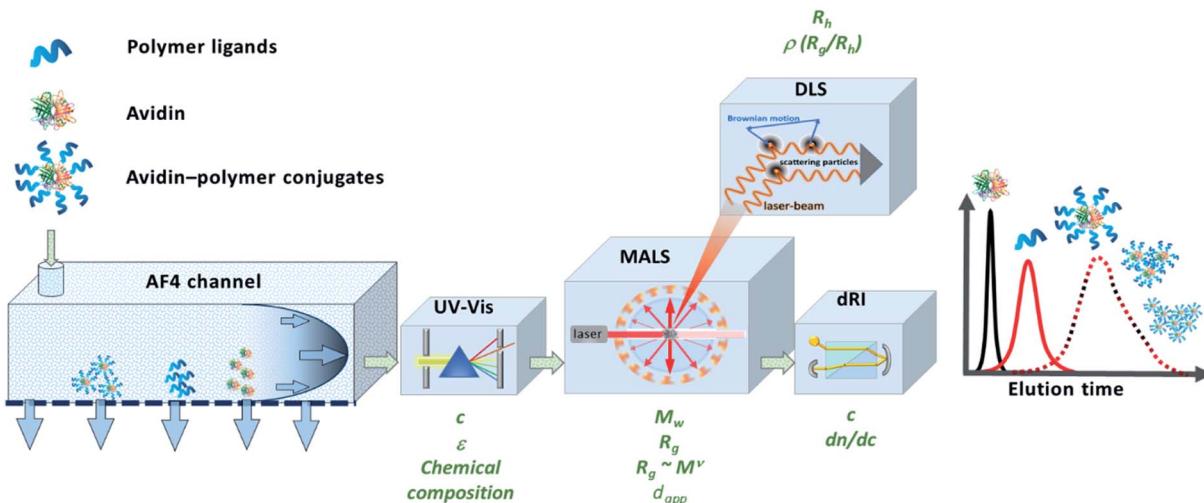


Fig. 1 Schematic representation of the instrumental setup of asymmetrical flow field-flow fractionation with fourfold detection system (AF4-D4) for the in-depth characterization of the molecular properties of avidin–polymer conjugates. Adapted with permission from J. Engelke, et al.³³ Copyright 2021 American Chemical Society.

Double-headed ATRP initiator was synthesized as previously described.¹⁹ Avidin–Br₂ conjugates were synthesized by mixing avidin (350 mg, 0.022 mmol protein, 0.22 mmol primary amine groups) with double-headed bromine-functionalized ATRP initiator (1.53 mmol, 915.7 mg (7 eq.) dissolved in 2 mL DMSO) in 170 mL of 0.1 M sodium phosphate buffer, pH 8.0. The reaction was stirred at 4 °C for 2 h and then dialyzed against 25 mM sodium phosphate, pH 8 using dialysis tubing with a molecular weight cut off of 15 kDa, for 24 h at 4 °C.

2.2.2 Preparation of Cu-MMA as deoxygenated catalyst solution. 100 mM CuCl₂ in deionized water (1.2 mL, 120 µmol) was bubbled with N₂ for 25 min and then 100 mM sodium ascorbate in deionized water (100 µL, 10 µmol) was added. MMA (49 µL, 144 µmol) was added to the copper suspension bubbled with N₂ for 3 min. The deoxygenated Cu-MMA solution was added to the synthesis vessel immediately.

2.2.3 ATRP from single-headed ATRP initiator modified avidin. A solution of monomer (36 mg for targeted DP 50 and 142 mg for DP 200 of CBMA, and 78 mg for DP 50 and 313 mg for DP 200 of OEGMA) and avidin-single-headed initiator conjugate (6.8 mg, 3.1 µmol of initiator) in 25 mM phosphate and 50 mM NaCl buffer (pH 8.0) was sealed and bubbled with N₂ for 30 min, then 350 µL of the deoxygenated Cu-MMA solution as mentioned above was added to the polymerization reactor under N₂ bubbling. The mixture was stirred at 4 °C for 4 h. The obtained conjugate was isolated by dialysis with 25 kDa molar mass cutoff dialysis tube in mixture of 25 mM phosphate buffer (pH 7.0) and deionized water at 4 °C, and then lyophilized.

2.2.4 ATRP from double-headed ATRP initiator modified avidin. A solution of monomer (59 mg for targeted DP 50 and 235 mg for DP 200 of CBMA, and 128 mg for DP 50 and 510 mg for DP 200 of OEGMA) and avidin-double-headed initiator conjugate (6.8 mg, 5.1 µmol of initiator) in 25 mM phosphate and 50 mM NaCl buffer (pH 8.0) was sealed and bubbled with

N₂ for 30 min, then 550 µL of the deoxygenated Cu-MMA solution as mentioned above was added to the polymerization reactor under N₂ bubbling. The mixture was stirred at 4 °C for 4 h. The obtained conjugate was isolated by dialysis with 25 kDa molar mass cutoff dialysis tube in mixture of 25 mM phosphate buffer (pH 7.0) and deionized water at 4 °C, and then lyophilized.

2.3 Analytical methods

2.3.1 MALDI-ToF. MALDI-ToF measurements were recorded using a PerSeptive Voyager STR MS with nitrogen laser (337 nm) and 20 kV accelerating voltage with a grid voltage of 90%. 500 laser shots covering the complete spot were accumulated for each spectrum. For determination of molecular weights of synthesized modified protein with initiator, sinapinic acid (10 mg mL⁻¹) in 50% acetonitrile with 0.4% trifluoroacetic acid was used as matrix. More details on MALDI-ToF studies can be found in ESI.[†]

2.3.2 Size exclusion chromatography (SEC). Number and weight average molar masses (M_n and M_w) and their dispersity D (M_w/M_n) of hydrolyzed polymer chains were estimated by SEC on a Water 2695 Series with a data processor, equipped with three columns (Waters Ultrahydrogel Linear Column, 500 and 250), using Dulbecco's Phosphate Buffered Saline with 0.02 wt% sodium azide as an eluent. Pullulan standards were used for molar mass determination. More details on hydrolysis and SEC can be found in ESI.[†]

2.3.3 Asymmetrical flow field-flow fractionation with four-fold detection (AF4-D4). AF4 measurements were carried out on an Eclipse DUALTEC system (Wyatt Technology Europe, Germany) with a 10 mM PBS buffer (pH 7.4) as mobile phase. Regenerated cellulose membrane (molecular weight cut-off 10 kDa) was employed as accumulation wall (Superon GmbH, Germany). The detection system consisted of a MALS detector (DAWN HELEOS II, Wyatt Technology, US) operating at



a wavelength of 659 nm with online DLS detector (QELS module, Wyatt Technologies, US) which is an add-on unit connected to the 99° angle of the MALLS, a variable wavelength detector (Agilent Technologies, US) set to 280 nm and an absolute refractive index (RI) detector (Optilab T-rEX, Wyatt Technology, US) operating at a wavelength of 658 nm. More details on AF4-MALS studies can be found in ESI.†

3 Results and discussion

3.1 Polymer-based protein engineering of avidin with zwitterionic and neutral polymers

Avidin–polymer conjugates were synthesized with varying polymer grafting density, polymer chain length and polymer type using grafting-from ATRP (Fig. 2). The avidin surface was first modified with either single or double-headed initiators as described previously.¹⁹ The number of modified amino groups was determined using MALDI-ToF mass spectrometry. Single-grafted avidin–polymer conjugates were synthesized from 35 single-headed initiator modified avidin (Fig. S2†) while double-grafted conjugates were produced from 29 double-headed initiator attached avidin (Fig. S3†). Two different chain lengths (short and long) of two different polymer types (zwitterionic poly(carboxybetaine methacrylate) and neutral poly(oligo(ethylene glycol) methacrylate)) were selected to be polymerized to avidin and analyzed by AF4-D4. To obtain a suitable comparison of the properties and solution behavior of polymers and protein–polymer conjugates, the non-conjugated polymers were also investigated using AF4-D4 (Table S1†). SEC analysis and bicinchoninic acid (BCA) protein assay of single-grafted polymers and avidin–polymer conjugates results were reported previously.³⁷ Avidin is a tetrameric protein with four identical subunits. For SEC analysis, when estimating overall avidin–polymer conjugate molar mass, the calculations used monomeric avidin with

polymers. Due to nondestructive nature of AF4, in a channel-based separation, avidin conjugates elute in their tetrameric form. Therefore, intrinsic conjugate molar masses determined from AF4-D4 are those of tetrameric avidin with polymers. For an easier comparison of AF4 results to conventional SEC and BCA assays, we recalculated the molar masses from SEC and BCA for tetrameric avidin–polymer conjugates (Table S2†). The polymer chain length was controlled by variation of the monomer to initiator ratio in ATRP solution (targeted DPs of 50 and 200). Conjugates were purified *via* dialysis and then lyophilized.

3.2 Avidin–polymer conjugates molar mass estimation using SEC and BCA protein assay

After purification, conjugates were characterized using BCA to determine protein content in the conjugates and then used to estimate conjugate molar mass and polymer DP. Targeted DPs were 50 and 200 and measured DPs from BCA assay were 32 and 62 for single-grafted avidin-pCBMA and 17 and 26 for double-grafted avidin-pCBMA, while for single-grafted avidin-pOEGMA DPs were 14 and 26 and for double-grafted avidin-pOEGMA conjugates they were 10 and 19 (Table S2†).

Acid induced polymer cleavage followed by SEC was performed to determine polymer's molar mass and *D*. For pCBMA DPs of 41 and 172 in single-grafted and 21 and 206 in double-grafted conjugates were found, while pOEGMA show DPs of 19 and 104 in single-grafted and 21 and 197 in double-grafted conjugates (Table S2†).

Our goal herein was to develop a characterization approach that could provide accurate and precise physical properties for tightly and weakly controlled ATRP reactions. The very large and disperse complexes that were generated in the more complex conjugates were particularly useful to test the limits of BCA, SEC, and later AF4.

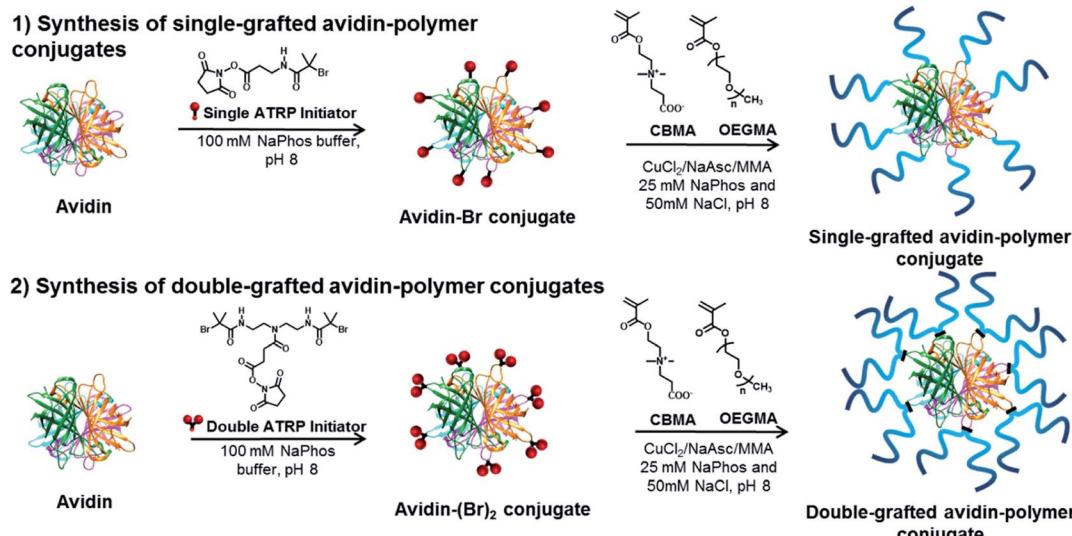


Fig. 2 Surface-initiated synthesis of single-grafted and double-grafted avidin conjugates using zwitterionic pCBMA and neutral pOEGMA polymers.



It was interesting that BCA assay and SEC showed significantly different DPs for longer polymer chains. The difference can be ascribed to different characterization principles and assumptions in result interpretation. Indeed, this is one of the central problems of current ways to characterize protein–polymer conjugates. In SEC, polymer–column enthalpic interactions (Fig. S14†) and calculation of molar mass in reference to a calibration standard can lead to a significant bias of the resulting data. It is critical to note that SEC separates according to size and not to molar mass. To accurately convert size into molar mass, calibration standards must be of identical conformation and density as the analyte.²⁴ Due to the lack of calibration standards for protein–polymer chimeras the polymers have to be cleaved from protein by acid treatment. Conjugates with high-grafting densities and long polymer chains can prevent the full cleavage and result in some polymers that are still linked to peptide fragments. This will increase heterogeneity of the analyte and thereby affect both D and the molar mass calculations.²³

In the BCA assay, characterization relies on accurate determination of protein concentration in the sample. Using protein concentration one can calculate the molar ratio of protein to polymer and conjugate's molar mass. Here the challenge is to accurately determine the protein content of conjugates that have high grafting polymer densities and long polymer chains, which then can result in misinterpretations of results.

3.3 AF4-D4 studies of avidin–protein conjugates

3.3.1 Separation and characterization of conjugates. The application of AF4 in combination with a multi-detection

system offers a variety of benefits in the characterization of bioconjugates.²⁷ A single run opens up a portfolio of parameters which can be simultaneously collected. The determination of absolute molar masses by AF4-MALS eliminates the need for relative molar mass calibration and increases the quality of the results. Compared to relative determination methods and related estimation of molar masses such as SEC of hydrolyzed polymers or BCA, AF4-MALS allows direct and absolute analysis of the unaffected conjugates. After the separation of the conjugate according to hydrodynamic diffusion properties concentration sensitive detection (UV and RI) and size-sensitive detection (MALS and DLS) is performed (Fig. 1).

The complete library of single-grafted polymers with varied physical and chemical properties and the corresponding conjugates have been characterized by AF4-D4, the results are summarized in Table 1. Additionally, more detailed interpretations and all fractograms can be found in the ESI.†

In Fig. 3, estimated conjugate molar masses of SEC and BCA assay are compared with the absolute molar masses determined by AF4-D4.

The molar masses obtained by SEC were significantly different from those determined by the BCA assay and the AF4-D4 (which were similar to each other). This strong discrepancy in SEC data may have been due to the incomplete hydrolytic cleavage of the polymer ligands and the resulting overestimation of the molar masses. Furthermore, molar mass determination is affected by the relative calibration with pullulan standards. The conformational properties of this polysaccharide are not comparable to those of zwitterionic pCBMA or neutral pOEGMA. The quantification of avidin by BCA and

Table 1 AF4-D4 results of avidin–polymer conjugates

	M_n^a (kg mol ⁻¹)	M_w^a (kg mol ⁻¹)	D (M_w/M_n)	Polymer chains per avidin ^b	R_g (nm)	R_h (nm)	ρ (R_g/R_h) ^c	Apparent density (kg m ⁻³)	Recovery ^e (%)	
Zwitterionic polymers										
C1	Single-grafted avi-pCBMA _{short}	383	517	1.35	7.9	16.3	15.0	0.97	22.0	93
C2	Single-grafted avi-pCBMA _{long}	741	1250	1.69	6.3	23.0	24.6	0.97	18.9	92
C3	Double-grafted avi-pCBMA _{short}	159	386	2.43	n.d.	17.2	16.5	0.94	14.0	95
C4	Double-grafted avi-pCBMA _{long}	395	867	2.19	n.d.	24.7	23.6	1.05	10.6	88
Neutral polymers										
C5	Single-grafted avi-pOEGMA _{short}	488	797	1.63	6.5	19.1	20.0	0.92	21.1	90
C6	Single-grafted avi-pOEGMA _{long}	641	1540	2.40	3.5	31.2	32.5	1.07	9.3	92
C7	Double-grafted avi-pOEGMA _{short}	298	904	3.03	2.3 (4.6 ^d)	28.9	28.0	0.91	6.6	91
C8	Double-grafted avi-pOEGMA _{long}	2320	11 800	5.09	7.8 (15.6 ^d)	109	66.6	1.38	1.7	97

^a Calculated with determined $dn/dc = 0.154$ mL g⁻¹ for pCBMA conjugates and 0.139 mL g⁻¹ for pOEGMA conjugates. ^b Average number of functionalities per avidin molecule (calculated by M_n of conjugate C, subtracted by M_n of macroinitiator (68 kg mol⁻¹) and divided by M_n of specific polymer F). ^c Determined at the peak maximum of DRI signal. ^d Single polymer segments per double-headed initiator. ^e Calculated by dn/dc and exact sample load.



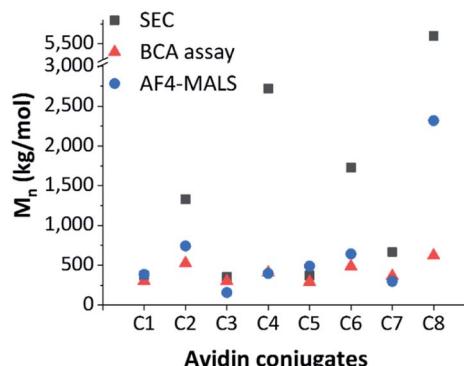


Fig. 3 Comparison of differently determined conjugate molar masses, estimated M_n by SEC (grey square), estimated M_n by BCA assay (red triangle) and absolute M_n by AF4-D4 (blue circle).

the estimation of related conjugated polymer amount was in better agreement with the AF4-D4 results. Only conjugate C8 showed deviation in the molar mass determined by these two methods. A deeper look into conformational properties helped clarify these inconsistencies, as discussed below.

It should be noted that the average number of effectively coupled polymer chains per avidin was less than the number of available sites for all conjugates, as determined by MALDI (accessibility of primary amines). ATRP of free single-grafted polymers and single-grafted avidin–polymer conjugates should yield molecules with similar polymer properties.

In addition, the synthesis and conjugation processes are statistical in nature, leading to different chain lengths of the polymers as indicated from the broad molar mass distribution determined *via* AF4-D4. Physical properties such as chain stiffness or flexibility, which are determined by the chemical nature, can have an influence on the dispersity and the resulting conjugate architecture. Therefore, the determination of the conformational properties is essential when seeking to understand the resulting conjugate dispersity. In former studies,^{25,35} it was demonstrated that differences in chemical composition and structural parameters caused a broad heterogeneity and dispersity instead of well-defined architectures.

3.3.2 Conformation at the nanoscale in one run. The process of data evaluation for conformational elucidation at the

nanoscale was similar for all architectures and is demonstrated using the conjugate of avidin and the long, zwitterionic pCBMA as an example (Fig. 2).

The fractograms in Fig. 4a show the separation of the individual components (native avidin, pCBMA_{long} (F2) and avidin-pCBMA_{long} conjugate (C2)) detected at different elution times. The later the elution, the lower the diffusion coefficient and, accordingly, the larger the hydrodynamic volume or molar mass. In addition to the absolute molar masses and radii (R_g and R_h) obtained from MALS and DLS detection, information about conformations at the molecular level was also obtained. For example, changes in the radii of gyration as a function of molar mass provided information about the scaling properties of the conjugate in the respective solvent. The slope ν (0.50) of the scaling plot for pCBMA in Fig. 4b was typical for statistical coils in theta-solvent (see ESI†).³¹ After the formation of conjugates, the scaling parameter of C2 ($\nu = 0.36$) revealed a more compact conformation that approximated a spherical shape. Due to the small size of native avidin and the narrow dispersity, no scaling plot could be generated.

Further data processing allowed the calculation of the apparent densities. These densities provided information about molecular compactness. Compared to the pure polymer chain, the conjugate exhibited a significantly higher d_{app} (Fig. 4c). For both, the polymer and the conjugate, it was noticeable that the d_{app} increased with increasing molar masses, passed through a maximum and then dropped again. Closer inspection of the ratio of R_g and R_h , known as the ρ parameter, revealed the conformation of the conjugates even more precisely and confirmed our previous observations (Fig. 4d). While $\rho(R_g/R_h)$ for the F2 was in the range for flexibly coiled chains, the conjugate possessed values of around 1. According to the literature, these $\rho(R_g/R_h)$ values are typical for spheres with rather heterogeneous surfaces. This behavior has also been observed for polymeric vesicles such as polymersomes.^{32,38,39}

3.3.3 Conformational changes depending on ligand variation. Comparative observation of the conformation of single-grafted conjugates with both zwitterionic pCBMA (C1 and C2) and neutral pOEGMA (C5 and C6) led to the conclusion that the chain length had a decisive influence on the structural properties of the conjugate. Thus, the longer the polymer chains, the larger the molar masses and radii and the broader the dispersity

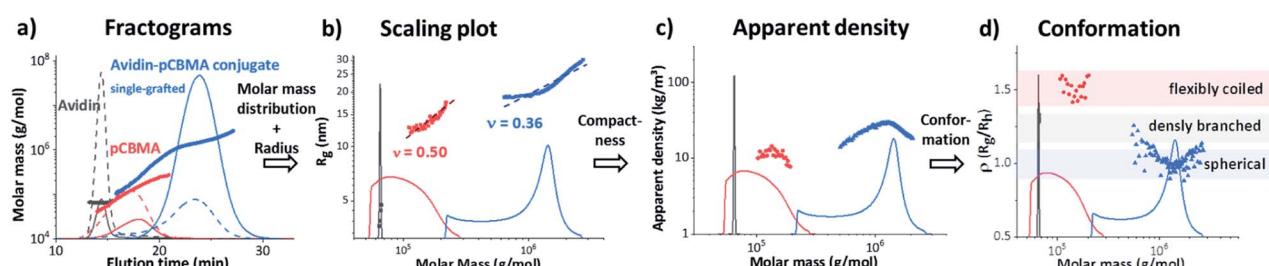


Fig. 4 Interpretation approach by AF4-D4 of native avidin (black), pCBMA_{long} (red) and single-grafted avidin-pCBMA_{long} conjugate C2 (blue), (a) fractograms with LS (solid line) and RI (dashed line) detector signals and molar masses vs. elution time, (b) scaling plots, radius of gyration (R_g) (symbols) and molar mass distributions (solid lines) vs. molar masses; (c) d_{app} , apparent densities (symbols) and molar mass distributions (solid lines) vs. molar masses; and (d) $\rho(R_g/R_h)$ parameter (symbols) and molar mass distributions (solid lines) vs. molar masses.



(Table 1). On the other hand, a decrease in density was observed (Fig. S10 and S12†). Since avidin forms a compact core around the mass centre of the conjugate, these data are to be expected. The single-grafted chains, in contrast, form a less dense shell around this core. With increasing chain length, the overall density of the conjugates decreased, while their size (R_g) increased. Regardless of the length of the polymer chain, the maximum density of all samples was at the maximum of the DRI signal. Thus, the major sample fraction possessed the highest apparent density.

The $\rho(R_g/R_h)$ parameters ($\rho = 0.97$) of the zwitterionic conjugates (C1 and C2) and the neutral C5 in this area were characteristic for globular, sphere-like conformations with a well-defined surface (Fig. 5). An increased heterogeneous surface was observed for the neutral avidin-pOEGMA conjugate with long polymer ligands (C6). Here, the ρ parameter was slightly higher ($\rho = 1.07$), which was characteristic for a compact, globular shape possessing a rough surface.

This behaviour was an indication that the conformational properties of the polymer chains had an influence on the conjugate shape. Although this may seem obvious, this is the first time that a single run technique could tease apart the intricacies of the protein–polymer conjugate formation. The scaling plots (Fig. S9†) of the individual polymers F2 and F4 confirmed this assumption. While the polymer chains of the zwitterionic pCBMA (F2) were clearly more flexible and more

coiled ($\nu = 0.5$), the neutral POEGMA (F4) showed a more elongated conformation ($\nu = 0.75$). In turn, the double grafted F6 of similar chemical origin as F4 shows again the typical reduction of the scaling parameter due to the compact, branching conformation. Finally, the more flexible polymer chains of C2 compensated for different degrees of polymerisation and generated a uniform conjugate surface. In contrast, the differences in chain length of the rigid polymer molecules after conjugation formed a rather rough surface for C6.

This becomes even clearer when considering the double-grafted conjugate structures, which contained higher numbers of polymer chains grown from double-headed initiators. The formation of double-grafted avidin-conjugates with short polymers (C3 and C7) led to comparable spherical conformations as observed for single-grafted C1 and C5 independent on the polymer chemistry. However, an increase of radii and a decrease in apparent density was observed for the double-grafted C3 and C7 (Fig. S11 and S13†).

A completely different behaviour was observed in the case of double-grafted, neutral avidin conjugate with long pOEGMA (C8). Here, molar masses, dispersity and radii were significantly higher compared to the other conjugates, but their apparent density was extremely low (Table 1). The reason for this can be found in their specific structure and conformation. The $\rho(R_g/R_h)$ parameter of C8 was significantly higher (1.38). This value is typical for irregular, clustered or branched chain conformations. In addition, an extremely low apparent density was observed. These two observations lead to the conclusion, that structures with a large hydrated shell around the avidin–polymer conjugate had been generated. A comparable behaviour was demonstrated for a similar polymer-conjugate system by molecular dynamic simulations.⁴⁰ A few highly hydrated, very long and stiff polymer chains, in addition to shorter chains resulted in high radii with low apparent densities. These structures can be classified as micelles, but with a heterogeneous polymer chain distribution (Fig. 5b). Additionally, the very broad distribution and high molar masses indicated the formation of a heterogeneous network, constructed of several avidin molecules. One reason for that could be aggregation. We systematically investigated the interaction of non-conjugated polymers with each other and with avidin by mixing variation of free polymer chains with native avidin. No shifts to higher molar masses or radii were observed and native avidin and free polymers eluted separately (Fig. S6d†). Thus, the interaction between single polymer chains and conjugates was excluded.

However, as mentioned earlier, the average number of available grafting sites for the polymer ligands per avidin molecule was 35 single-headed initiators and 29 double-headed initiators, respectively. Steric reasons could also have led to structures with longer polymer chains. At the same time, unreacted initiators were possibly still present.

Unfortunately, a statistical calculation of the polymer or avidin distribution within the conjugates was not possible due to absorption of both components at the applied wavelength ($\lambda = 280$ nm) of the UV detector. Here, a labelling of the protein could help. However, the introduction of a marker group could also change the solution properties and AF4 membrane

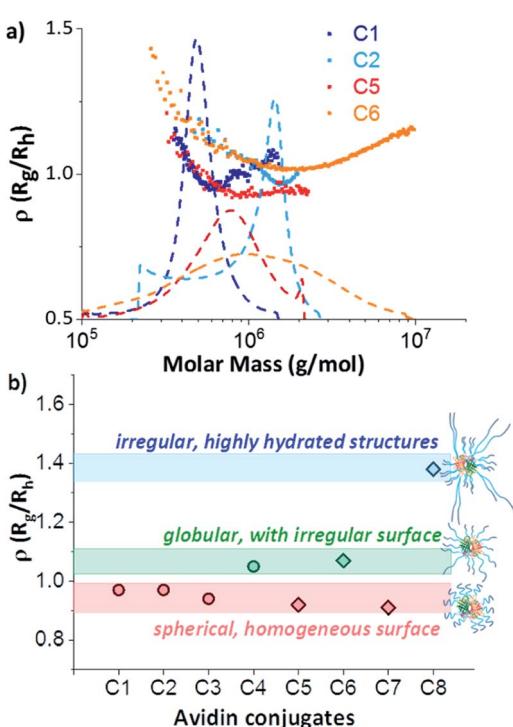


Fig. 5 (a) $\rho(R_g/R_h)$ parameters (symbols) and differential weight fractions (dashed lines) as a function of molar mass of single-grafted conjugates C1 (dark blue); C2 (bright blue); C5 (red) and C6 (orange); and (b) $\rho(R_g/R_h)$ parameters at concentration maximum of all zwitterionic (circles) and neutral (diamonds) avidin conjugates with proposed schematic structures.



interactions significantly. Furthermore, fluorescence effects in the wavelength range of the laser must be avoided when using light scattering detection.

4 Conclusions

AF4-D4 shows that the formation of polymer–protein conjugates depends on the physicochemical properties of the polymer chains. As an example, two polymer types (zwitterionic pCBMA as well as neutral pOEGMA) with different chain lengths, single-grafted and double-grafted conjugate architectures were generated and characterized. To correlate the conformation properties with the chemical nature, asymmetrical flow field flow fractionation in combination with a multi-detection system was used. The AF4-D4 hyphenation allows to generate the data needed for determination of a series of structural parameters, which can be correlated only if they are collected simultaneously. The data evaluation enabled a comprehensive elucidation of the structures with the help of the conformational properties and contributed to a remarkable understanding of the conjugation processes. It has been shown that polymer flexibility and hydration have a major influence on the properties in solution such as shape and apparent density.

Data availability

All relevant data is included in the paper and ESI.†

Author contributions

BK and HM prepared the polymers and conjugate samples and performed the SEC, BCA and MALDI-ToF experiments. SB performed the AF4 experiments and the conformation studies. BK and SB analysed the data and wrote the manuscript. KM and AR supervised the synthetic work, MALDI-ToF and BCA analyses. AL supervised the AF-D4 analyses and interpretation. All authors contributed to discussing the results and reviewing the manuscript.

Conflicts of interest

There are no conflicts to declare.

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