Neat Protein Single-Chain Nanoparticles from Partially Denatured BSA

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ABSTRACT: The main challenge for the preparation of protein single-chain nanoparticles (SCNPs) is the natural complexity of these macromolecules. Herein, we report the suitable conditions to produce “neat” bovine serum albumin (BSA) single-chain nanoparticles (SCNPs) from partially denatured BSA, which involves denaturation in urea and intramolecular cross-linking below the overlap concentration. We use two disuccinimide ester linkers containing three and six methylene spacer groups: disuccinimidyl glutarate (DSG) and disuccinimidyl suberate (DSS), respectively. Remarkably, the degree of internal cross-linking can be followed simply and efficiently via $^1$H NMR spectroscopy. The associated structural changes as probed by small-angle neutron scattering (SANS) reveal that the denatured protein has a random-like coil conformation, which progressively shrinks with the addition of DSG or DSS, thus allowing for size control of the BSA-SCNPs with radii of gyration down to 5.4 nm. The longer cross-linker exhibits slightly more efficiency in chain compaction with a somewhat stronger size reduction but similar reactivity at a given cross-linker concentration. This reliable method is applicable to a wide range of compact proteins since most proteins have appropriate reactive amino acids and denature in urea. Critically, this work paves the way to the synthesis of “neat”, biodegradable protein SCNPs for a range of applications including nanomedicine.

1. INTRODUCTION

Single-chain nanoparticles (SCNPs) are discrete polymer chains folded or collapsed via intramolecular cross-linking under dilute conditions. Due to their small size, softness, and internal compartmentalization, they have potential applications in catalysis, sensing, and drug delivery as well as in all-polymer nanocomposites. Up to date, most efforts have been focused on SCNPs obtained from synthetic macromolecules. More recently, there has been great interest in expanding this technology to biodegradable and biocompatible polymers, including proteins.

Numerous synthesis strategies of SCNPs from synthetic (co)polymers have been reported, which typically involve two steps: functionalization of a linear precursor and subsequent internal cross-linking. Each cross-link between two reactive functional sites of the chain generates an internal loop, and globally, the precursor chain reduces its size. The resulting shrinking can be monitored by experimental techniques of different degrees of complexity, e.g., size exclusion chromatography (SEC) and dynamic light scattering (DLS). These are indirect methods that have shown that the length and flexibility of the cross-linker have a significant impact in the chain folding. However, changes in the polymer conformation, i.e., an increase in internal particle compactness that goes along with the size reduction, are best probed directly with small angle X-ray scattering (SAXS) or small angle neutron scattering (SANS) techniques, thanks to the scattering vector (Q) dependence of the measured intensities. Direct structural determination by SAXS/SANS with the aid of simulations and scaling analysis has shown that the usual

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morphology of SCNPs is sparse with internal loops that leads to the presence of local “pockets”. Furthermore, unlike globular proteins whose folding is given by defined interactions, the collapse of synthetic SCNPs is the result of a stochastic process leading not only to a less controlled compaction but also to a polydispersity of resulting topologies.

In addition to their applications as soft synthetic nanoparticles, synthetic SCNPs are considered as simple models to mimic the more compositionally complex intrinsically disordered proteins (IDPs). The main advantage of synthetic SCNPs as IDP models is their simpler chemical composition, making it possible to isolate excluded volume effects particularly relevant in crowded environments. However, real proteins are compositionally more complex due to the chemical diversity of the proteinogenic amino acids. Thus, the synthesis of polypeptide single-chain nanoparticles bridges the gap between synthetic SCNPs and proteins, thereby extending their potential applications. In particular, protein SCNPs could find interest in nanomedicine, where gene therapy is its largest emergent field.

To synthesize “neat” protein-based SCNPs, we chose the protein BSA (bovine serum albumin) because it has several advantages as a precursor. First, it consists of a well-characterized long polypeptide single chain with approximately 10% of reactive lysine residues homogeneously distributed along the chain that are available for cross-linking. Furthermore, globular BSA denatures in the presence of chemical denaturants such as urea and adopts a random-like coil conformation. Urea denatures proteins by breaking the H-bonds and by increasing the solvation of hydrophobic groups. Last but not least, serum albumins, due to their availability, low cost, and low immunogenicity, are great candidates for the synthesis of nanoparticles. In fact, protein nanoparticles with radii ranging from tens to hundreds of nanometers have been made from BSA for applications in targeted drug delivery and pH sensing. In this article, we report a procedure to push down the size to that of a neat polypeptide single chain. In a recent work, “hybrid” SCNPs of human serum albumin stabilized with PEG grafts were synthesized with a hydrodynamic radius of down to 9.5 nm. Here, we show that it is possible to obtain “neat” SCNPs from unPEGylated BSA with radius of gyration of down to 5.4 nm. With this aim, we have adopted the strategy to cross-link internally the lysine residues of denatured BSA using two cross-linkers of different lengths: disuccinimidyl suberate (DSS) and disuccinimidyl glutarate (DSG) and exploited our previous experience in the preparation and characterization of synthetic unimolecular SCNPs.

2. MATERIALS AND METHODS

2.1. Materials. Bovine serum albumin (BSA), disuccinimidyl glutarate (DSG), urea, sodium chloride (NaCl), phosphate-buffered saline (PBS, powder, pH 7.4, for preparing 1 L solutions), and dimethyl sulfoxide were obtained from Sigma-Aldrich. Disuccinimidyl suberate (DSS) was purchased from Acros Organics. Deionized water was obtained from a Barnstead TII Pure Water System. Deuterium oxide (D$_2$O, 99.9% D), deuterated urea (D$_4$, 98% D), and deuterated dimethyl sulfoxide (d$_6$-DMSO, 99.8% D) were obtained from Eurisotop and used without further purification. Deuterated PBS (d-PBS) powder was prepared from hydrogenated PBS powder by exchanging the hydrogen atoms for deuterium, dissolving the salts in D$_2$O, followed by freeze-drying. Similarly, the BSA for SANS was dissolved in D$_2$O to remove exchangeable protons, and after 1 day of incubation, the solution was lyophilized. This process was performed twice.

2.2. Sample Preparation and BSA-SCNP Formation. Denatured BSA solutions were prepared by mixing 10 vol % of a solution of BSA (10 mg/mL) in PBS and 90 vol % of a solution of urea (6.67 M) and NaCl (0.57 M) in PBS. The final composition thus was BSA (1 mg/mL), urea (6 M), and NaCl (0.6 M) at pH 7.4. BSA-SCNPs were formed by the addition of cross-linker in DMSO (with concentrations of 5–8 wt %) and stirred for at least 12 h. SANS samples were prepared with partially hydrogenated BSA, hydrogenated cross-linkers in deuterated DMSO, deuterated urea, d-PBS, and D$_2$O. The structural characterization with SANS and SAXS (see the SI) was performed directly in the reacting medium. Samples for $^1$H-NMR were dialyzed against decreasing concentrations of urea until the proteins were in water and subsequently dialyzed against water. Then, the protein SCNPs were lyophilized and redissolved in D$_2$O.

$^1$H nuclear magnetic resonance ($^1$H NMR) spectra were recorded at room temperature on a Bruker spectrometer operating at 400 MHz using D$_2$O as solvent. The solution density was measured with a DMA 4500 M apparatus from Anton Paar. The pH was measured with a Laquatwin pH meter from Horiba.
Table 1. Summary of the Main Properties of Native BSA, (Partially) Denatured BSA, and BSA-SCNPs as Determined by $^1$H NMR and SANS Measurements

| sample     | cross-linker spacer, R | cross-linker(mol %) | app. lysine conversion(%) | $I(0)^c$ (cm$^{-1}$±0.003) | $R_p$ (nm) | $\nu^d$ (±0.01)
|------------|------------------------|---------------------|---------------------------|-----------------------------|------------|---------------------
| native BSA |                        |                     |                           |                             |            | 0.073               |
| denatured BSA |                      |                     |                           |                             |            | 3.7 ± 0.1           |
| SCNP #1    | (CH$_2$)$_3$          | 4.6                 | 51                        | 0.109                       | 6.65 ± 0.5 | 0.45                |
| SCNP #2    |                        | 9.1                 | 63                        | 0.100                       | 6.50 ± 0.6 | 0.43                |
| SCNP #3    |                        | 18                  | 74                        | 0.107                       | 5.86 ± 0.3 | 0.42                |
| SCNP #4    |                        | 28                  | 83                        | 0.102                       | 5.46 ± 0.3 | 0.39                |
| SCNP #5    | (CH$_2$)$_6$          | 4.6                 | 42                        | 0.122                       | 6.07 ± 0.2 | 0.47                |
| SCNP #6    |                        | 9.1                 | 64                        | 0.146                       | 6.09 ± 0.4 | 0.46                |
| SCNP #7    |                        | 18                  | 80                        | 0.141                       | 5.51 ± 0.5 | 0.40                |
| SCNP #8    |                        | 28                  | 86                        | 0.142                       | 5.42 ± 0.3 | 0.43                |

$^a$Mole percent of cross-linker with respect to the total concentration of protein residues. $^b$Relative contribution of peak $a$’ from $^1$H-NMR (see the SI). $^c$Parameters obtained from fitting the SANS data with a generalized Gaussian random coil form factor (see eq 3 in the text). $^d$The BSA in its native conformation has the form of a prolate ellipsoid with a polar axis of $R_p$ = 7.3 nm and an equatorial axis of $R_e$ = 2.5 nm, with a corresponding radius of gyration of $R_g$ = 3.7 nm (see the SI).

2.3. Small-Angle Neutron Scattering. Small-angle neutron scattering (SANS) measurements were performed on the SANS-I instrument at the Spallation Neutron Source, SINQ, Paul Scherrer Institut, Switzerland. The mean wavelength of the incident neutron beam was $\lambda$= 5 Å with a wavelength resolution of approximately 10% at two-sample-to-detector distances of 4.5 and 18 m, respectively, to cover the data in the wave vector transfer $Q$ range of 0.03 to 0.23 nm$^{-1}$, with $Q = 4\pi\sin(\theta/2)/\lambda$, where $\theta$ is the scattering angle. Samples were measured in Quartz Hellma cells (110-QS) of 2 mm thickness. The measured SANS data were corrected and normalized to a cross-sectional unit using BERSANS-PC data processing software. Solvent backgrounds (d-PBS in D$_2$O and d-PBS, NaCl, and d-urea (6 M) in D$_2$O) were measured in the same cells and subtracted from the data. The background-corrected scattering curves were fitted to an ellipsoid in the case of BSA in its native conformation and, in the case of BSA and the SCNPs, in denaturing conditions with a model for generalized Gaussian coils.

3. RESULTS AND DISCUSSION

To synthesize neat BSA-SCNPs, we first denatured the protein in phosphate-buffered saline (PBS) with NaCl (0.6 M) and urea (6 M). We used a low protein concentration of 1 mg/mL well below the overlap concentration to avoid, as far as possible, intermolecular cross-linking events. We synthesized the BSA-SCNPs via addition of the cross-linkers DSS and DSG at 0–28 mol % with respect to the total protein amine group concentration. DSS and DSG react mainly with the lysine primary amine groups, leading to amide formation as shown in Figure 1A,B. Table 1 collects the library of prepared BSA-SCNPs and their main properties.

First, we examine the cross-linking reaction via $^1$H-NMR. Figure 1 shows the representative $^1$H-NMR spectra of the BSA-SCNPs with both cross-linkers compared to that of BSA. As expected in the case of high-molecular weight proteins, the $^1$H-NMR spectrum of globular BSA is complex since the most part of the protons give rise to broad unresolved resonances. Nevertheless, the main signatures due to cross-linking are visible. First, we observe changes in the chemical shift of the lysine a protons corresponding to the change from being linked to a primary amine to being bonded to an amide: disappearance of the peak a at 2.92 ppm and appearance of a new peak a’ at 3.09 ppm (see Figure 1C,D). This shift is much clearer in the cross-linking of poly-lysine where there is no interference of neighbor signals, confirming the peak assignment (see the Supporting Information (SI), Figure S1). The BSA in its native conformation has the form of a prolate ellipsoid with a polar axis of $R_p$ = 7.3 nm and an equatorial axis of $R_e$ = 2.5 nm, with a corresponding radius of gyration of $R_g$ = 3.7 nm (see the SI).
in d-urea (6 M)/NaCl (0.6 M)/D2O. Figure 2A shows that the scattering pattern of the native protein in buffer solution corresponds to rather small and compact objects, whereas denaturation leads to larger objects as seen from the shift of the curve toward lower Q values and the decrease of the intensity decay slope at higher Qs. The size, related to lower degrees of compactness within the object.

In general, the SANS intensity is expressed as

\[ I(Q) = \frac{1}{\nu U^{1/\nu}} \left( 1 - \frac{1}{\nu U} \right)^{\nu + 2} \left( \frac{1}{\nu U} \right)^{\nu + 1} \frac{Q^2}{R_g^2} / 6 \]

where \( U = (2\nu + 1)(2\nu + 2)Q^2R_g^2/6 \) and \( \nu(a, x) = \int x e^{-x} \exp(-t) dt \). This function allows taking into account different compaction degrees of the coil and excluded volume effects. In eq 3, \( R_g \) is the radius of gyration and \( \nu \) is the scaling exponent, which defines the relation between the polymer molar mass and its size \( R_g \approx M_{AV}^{1/\nu} \). \( \nu \) is 0.5 for an unordered polymer (which is the conformation of a linear polymer chain in theta-conditions), 0.6 for a swollen chain in a good solvent (the so-called self-avoiding-walk conformation), and 1/3 for a polymeric chain collapsed into a globule. Thus, the degree of compaction of the polypeptide chain is characterized through the scaling factor, \( \nu \), which determines the Q-dependence of the scattered intensity at intermediate and high Q-values.

The forward scattering, \( I(0) = \phi \Delta \rho^2 V_p P(Q) S(Q) \), contains information on the macromolecular volume \( V \approx M_{AV} / (N_{AV} d) \) (with \( M_{AV} \) as the molecular weight, \( N_{AV} \) as the Avogadro number, and \( d \) as the mass density).

Equation 3 fits well the data of the denatured BSA with a radius of gyration of \( R_g = 3.78 \) nm and a scaling exponent of \( \nu \approx 0.5 \). Thus, the chain has clearly expanded with respect to the native conformation. The scattering pattern of the BSA-SCNP synthesized with 28 mol % DSG depicted in Figure 2B shows a size reduction with respect to denatured BSA visible in a shift of the scattering pattern toward higher Q-values. The size reduction is confirmed by the fit of the experimental data using eq 3 with \( R_g = 5.46 \) nm and \( \nu = 0.39 \), indicating a higher degree of compaction.

We observe that there are no significant changes in the intensity at low Q after cross-linking, indicating that under the conditions employed, the cross-linking is mostly intrachain, and interchain bonding may be considered minor. A slight intensity increase due to the additional molecular weight provided by the cross-linker is expected, but it should be a maximum of a factor of 1.06 for DSG and 1.08 for DSS in case all lysine groups were cross-linked. Thus, the individual BSA chains collapse into unimolecular SCNPs, which demonstrates the success of our approach to synthesize polypeptide SCNPs.
The low-Q upturn in the intensity observed in both curves of BSA and SCNPs in denaturing media is attributed to the denaturing conditions rather than a significant contribution of interchain cross-link formation.

In addition to a size reduction upon cross-linking, the fit of the scattering data using eq 3 revealed a higher scaling exponent, from $\nu = 0.5$ for denatured BSA to $\nu = 0.39$ for the BSA-SCNP synthesized with 28 mol % DSG. To accentuate changes in the macromolecular shape, Figure 3 shows the SANS data in a Kratky plot. By rescaling with their respective radii of gyration, we eliminate the size effect and emphasize the differences in the fractal dimension at high Q. We note that differences at very low Q arise as a result of the rescaling with $R_g$ and we just focus on the high-Q behavior. Comparing the results of the denatured BSA with the SCNPs with 28 mol % DSG, we observe that the data at intermediate Q-values superimpose, whereas differences appear at high Q-values in the range of small length scales due to the internal cross-linking. In denaturing conditions, the scattering intensity of the BSA-SCNP decays steeper than that of the BSA, revealing a higher degree of compactness, but it does not reach the compaction degree of the BSA in its native state. In this representation, it is clear that the protein denatured with urea (6 M) has a conformation that is roughly a Gaussian random-like walk, as reported in other works,\textsuperscript{16,27} rather than a coil in good solvent. This may be attributed to the fact that BSA has intact disulfide cross-links in urea (6 M), which has an impact on the short-length scales of BSA even when denatured as observed by SANS.\textsuperscript{27} Moreover, previous SANS experiments have shown that higher concentrations of urea lead to even higher values of $R_g$ due to a higher degree of denaturation.\textsuperscript{16} Here, the $R_g$ value is slightly larger than the one reported for the same urea concentration but lower pH, closer to the isoelectric point of BSA.\textsuperscript{16} Thus, taking into account all these arguments, we conclude that the BSA protein is only partially denatured and it compacts more as a consequence of intramolecular cross-linking.

The SANS curves and the fits using the generalized Gaussian coil model for all the synthesized BSA-SCNPs are shown in Figure 4. The fit results are collected in Table 1. In all cases, the cross-linking reaction leads to a progressive size reduction and increased degree of compaction with increasing cross-linker concentration.

Figure 5 shows the radii of gyration obtained for all BSA-SCNPs as a function of the amount and type of cross-linker. The radii of gyration decrease from 7.38 to ca. 5.4 nm upon increasing the cross-linker concentration up to 28 mol % (relative to the total monomer concentration in BSA) according to the progressive collapse of the denatured BSA chain. We also observe from Figure 5 that the longer cross-linker is slightly more efficient in collapsing the polypeptide chain at low contents of cross-linker. The average cross-linker length in its most possible extended conformation can be estimated from the number of carbon bonds to be 0.89 and 0.51 nm for DSS and DSG, respectively. These values were calculated using the program ChemBio3D Ultra simulating the chemical structures in their all-trans conformation. Note that they should be taken cautiously because a priori, the actual conformation of the cross-linking segment in the SCNP is unknown.
The degree of cross-linking was estimated by means of H NMR, where we found a similar reaction extent for both DSS and DSG as a function of the added cross-linker concentration. $R_g$ is obtained from fitting a generalized Gaussian coil to the SANS data (eq 3 in the text). Dashed lines are fits to $R_g \propto x^{-7}$ with $y=0.11$ for DSG and 0.07 for DSS. The error bars represent the maximum width of uncertainty.

unknown. Nevertheless, we can compare them to the size of the denatured BSA—radius of gyration $R_g = 7.38$ nm and end-to-end distance $R_E = 18$ nm—and the effective monomer length taking into account the number of amino acids in BSA $N = 583$, $l = \sqrt{\frac{R_E^2}{N \nu}} = 0.74$ nm. The cross-linker lengths seem to be comparable to the effective monomer length. Our SANS results suggest that the longer cross-linker is able to form larger loops, leading to a stronger size reduction. This finding is in agreement with recent literature results, which report that shorter cross-linking molecules have been found to achieve less compaction due to a dominance of short-range loops. The overall size reduction of SCNPs compared to their linear precursors is expected to depend on solvent quality, molecular weight of the precursor, and cross-linking density. Theoretically, the size of elastic single-chain nanoparticles in solution is estimated to scale with the cross-linking density as $R_g \propto x^{-1/5}$ in good solvent and $R_g \propto x^{-1/8}$ in theta solvent. Here, in both cases, we obtain lower dependencies, namely, $R_g \propto x^{-0.11}$ and $\approx x^{-0.07}$ for DSG and DSS (dotted lines in Figure 5), respectively, probably due to lower reactivity of the cross-linker added that may yield less effective cross-links or due to the stiffness of the polypeptide chain.

4. CONCLUSIONS

This work provides a straightforward and versatile method to prepare protein-based SCNPs. We have synthesized polypeptide SCNPs based on BSA (bovine serum albumin) denatured with urea (6 M) and cross-linked below the overlap concentration with succinimide-type linkers with two spacer lengths. The degree of cross-linking was estimated by means of $^1$H-NMR spectra and peak decomposition; additional SANS analysis; complementary SAXS results (PDF)

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### Notes

The authors declare no competing financial interest.

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