The interaction of unfolding \( \alpha \)-lactalbumin and malate dehydrogenase with the molecular chaperone \( \alpha \)B-crystallin: a light and X-ray scattering investigation

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Abstract
Purpose: The molecular chaperone $\alpha B$-crystallin is found in high concentrations in the lens and is present in all major body tissues. Its structure and the mechanism by which it protects its target protein from aggregating and precipitating are not known. Methods: Dynamic light scattering and X-ray solution scattering techniques were used to investigate structural features of the $\alpha B$-crystallin oligomer when complexed with target proteins under mild stress conditions, i.e., reduction of $\alpha$-lactalbumin at 37 °C and malate dehydrogenase when heated at 42 °C. In this investigation, the size, shape and particle distribution of the complexes were determined in real-time following the induction of stress. Results: Overall, it is observed that the mass distribution, hydrodynamic radius, and spherical shape of the $\alpha B$-crystallin oligomer do not alter significantly when it complexes with its target protein. Conclusions: The data are consistent with the target protein being located in the outer protein shell of the $\alpha B$-crystallin oligomer where it is readily accessible for possible refolding via the action of other molecular chaperones. © 2010 Molecular Vision.

Keywords
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The interaction of unfolding α-lactalbumin and malate dehydrogenase with the molecular chaperone αB-crystallin: a light and X-ray scattering investigation

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Abstract

Purpose: The molecular chaperone αB-crystallin is found in high concentrations in the lens and is present in all major body tissues. Its structure and the mechanism by which it protects its target protein from aggregating and precipitating are not known.

Methods: Dynamic light scattering and X-ray solution scattering techniques were used to investigate structural features of the αB-crystallin oligomer when complexed with target proteins under mild stress conditions, i.e., reduction of α-lactalbumin at 37 °C and malate dehydrogenase when heated at 42 °C. In this investigation, the size, shape and particle distribution of the complexes were determined in real-time following the induction of stress.

Results: Overall, it is observed that the mass distribution, hydrodynamic radius, and spherical shape of the αB-crystallin oligomer do not alter significantly when it complexes with its target protein.

Conclusions: The data are consistent with the target protein being located in the outer protein shell of the αB-crystallin oligomer where it is readily accessible for possible refolding via the action of other molecular chaperones.

Introduction

Small heat shock proteins (sHsps) are a diverse family of intracellular molecular chaperones that are found in all organisms [1, 2]. In humans, they are present in many tissues at varying levels depending on the stage of development and the level of physiologic stress. The role of sHsps in cells is multi-faceted with their common theme of action being to interact with and stabilize partially folded states of other (target) proteins to prevent their aggregation and possible precipitation, for example under conditions of environmental stress such as elevated temperature, low pH and oxidation [2-5]. In vitro, sHsps prevent stress-induced aggregation of a variety of unrelated target proteins that undergo either disorderered (amorphous) or ordered (amyloid fibril) forms of aggregation [2, 6, 7].

The principal eye lens protein, α-crystallin, is a sHsp that comprises two closely related subunits, αA- and αB-crystallin, each of which is ~20 kDa in mass. In the human lens, the two isoforms are co-expressed in a ratio of 3:1 αA-:αB-crystallin [8]. The two isoforms form a heterogeneous oligomeric species of average mass of approximately 800 kDa and 150 Å in diameter [8]. The lens contains a very high concentration of protein (up to 450 mg/ml in the nucleus or center of the lens) encased in very long fiber cells. α-Crystallin is the primary protein component of the lens and can approach 50% of the total dry weight of the lens [9]. α-Crystallin has two important functions in the lens. First, in a structural role, it assists in the maintenance of short-range order in the lens cytoplasm, ensuring proper refraction of light and maintaining lens transparency [10]. Second, it acts as a molecular chaperone to maintain the solubility of the other classes of crystallin proteins, β- and γ-crystallin. It is also known to protect other non-crystallin lens proteins such as sorbitol dehydrogenase from both thermal aggregation and enzyme inactivation [11]. There is no protein turnover in the center of the lens, meaning that the crystallin proteins have to be very long lived; α-crystallin is involved in minimising lens protein precipitation over decades, and thereby the prevention of lens opacification and cataract formation [12].

Outside the lens, αB-crystallin is also expressed at significant levels [13] where it has a key role as a molecular chaperone. For example, in addition to its role in the prevention of cataract, αB-crystallin is of interest extralenticularly because its expression is associated with many other protein misfolding disorders. Thus, αB-crystallin is found in significant levels in the brains of patients with Alzheimer’s disease [14, 15], Parkinson disease [16, 17], in multiple sclerosis [18, 19], and in the ischemic heart [20]. In vitro, αB-crystallin protects target proteins against reduction-induced precipitation [21, 22], heat-induced aggregation [6, 23], enzyme inactivation [24-26], and amyloid fibril formation [7].

The polydispersity of α-crystallin means that its assemblages can vary considerably in their number of subunits. Cryo-electron microscopic (cryoEM) studies of the αB-crystallin oligomer [27] show that it contains a spherical protein shell of 80 to 180 Å in diameter surrounding a central cavity measuring 30 to 100 Å in diameter, as well as a region on the protein surface that is highly dynamic which arises from the flexible COOH-terminal extensions [28, 29]. The heterogeneous and dynamic nature of the αB-crystallin oligomer has precluded crystal formation and thereby precise atomic structural resolution of the protein and specific details about its mechanism of chaperone action.

Previously, we used low-angle X-ray scattering from solutions of extracted bovine α-crystallin on its own and in the presence of β-crystallin to study structural changes in α-crystallin during chaperone action as a function of temperature [30, 31]. The α-crystallin oligomer underwent extensive structural changes and became much larger at higher temperature, with a major transition at around 50 °C. We used the term ‘super aggregation’ to describe the enlargement of the α-crystallin oligomer with increasing temperature. Our results were consistent with earlier transmission electron microscopy, circular dichroism and non-denaturing gel electrophoresis studies of α-crystallin [32]. Interestingly, we also found that below 50 °C, a weak peak occurred between α-crystallin and β-crystallin implying that the β-crystallin subunits may be transiently localized in the exterior of the α-crystallin oligomer. At higher temperatures (i.e., under conditions of partial unfolding of β-crystallin), the β-crystallin subunits were more likely bound to the surface of the α-crystallin oligomer [31]. Our recent neutron scattering studies on the interaction of the target protein γE-crystallin with α-crystallin at 65 °C have provided further insight into the location of the target protein when interacting with α-crystallin under cataract conditions. Under these relatively harsh stress conditions, the data are consistent with γE-crystallin binding in the central cavity of the α-crystallin oligomer [33]. Our previous X-ray solution scattering experiments used extracted bovine α- and β-crystallins, both of which are comprised of several isoforms. While such experiments are relevant to the in vivo situation in the eye lens, they do not reflect the situation in other tissues where
only αB-crystallin is found. Furthermore, ascribing the observed experimental structural changes to particular species in such multi-component mixtures is difficult. Thus, the purpose of this study was to examine the interaction of recombinant αB-crystallin with the well characterized target proteins, α-lactalbumin (α-LA) and malate dehydrogenase (MDH). α-LA is a small monomeric milk protein (mass approximately 14 kDa) that contains four disulfide bonds in its native state. When these bonds are reduced, α-LA forms an intermediately folded (molten globule) state that aggregates and precipitates out of solution [34-36]. The precipitation of α-LA can be prevented by a sufficient quantity of αB-crystallin [34-36]. α-LA is an attractive target protein to study as its folding pathway, and its various intermediate states, have been well characterized. There have also been a variety of biophysical studies undertaken on the interaction of reduced α-LA with αB-crystallin including our detailed real-time spectroscopic and biophysical investigations in which we showed that αB-crystallin interacts with reduced, partially folded, monomeric α-LA to prevent its aggregation and precipitation [34-36] During this interaction, αB-crystallin acts on the destabilized molten globule form of α-LA, which consequently retains some secondary structure within the complex formed with the chaperone protein [34-36]. Malate dehydrogenase (MDH) is an enzyme involved in the citric acid cycle that catalyzes the conversion of malate to oxaloacetate and exists as a dimer or tetrameric enzyme comprised of identical subunits each of mass between 30 and 35 kDa [37]. Under mild thermal stress, MDH partially unfolds, aggregates and precipitates and therefore has been used as a target protein to investigate the chaperone activity of both α-crystallin and αB-crystallin [26,38].

Here we describe, for the first time, the use of X-ray solution scattering to investigate the complex formed between αB-crystallin and its target proteins α-LA and MDH. In addition, we have used dynamic light scattering (DLS) to characterize the size of the complex formed between αB-crystallin and α-LA. The X-ray scattering and DLS experiments enabled real-time measurements to be made of the structural alterations that occur when αB-crystallin interacts with reduced α-LA to prevent its aggregation and precipitation. Overall, the α-LA data presented herein are consistent with our previous studies [36] and the cryoEM studies of Stewart and coworkers [3,27,38]. We find that the size of the αB-crystallin oligomer is very similar to that determined from cryoEM measurements and that, in the complex formed between αB-crystallin and α-LA, the data are consistent with α-LA being located in the outer protein shell of the αB-crystallin oligomer. Similarly, from the light and X-ray scattering experiments of the interaction of thermally stressed MDH with αB-crystallin, we were able to monitor structural changes in real time of the proteins during chaperone interaction and conclude that a similar mode of interaction occurs as between reduced α-LA and αB-crystallin.

**Methods**

The vector pET24d(+) (Novagen, Madison, WI) containing the gene for expression of human αB-crystallin was a gift from Professor W. Boelens (University of Nijmegen, Netherlands). Human recombinant αB-crystallin was expressed and purified as described previously [39]. Calcium-depleted bovine α-LA (mitochondrial from porcine heart) was purchased from Sigma (Gillingham, UK). All other chemicals were of the highest grade.

**Light scattering and dynamic light scattering monitoring of the chaperone action of αB-crystallin against reduced α-LA**

Aggregation assays of αB-crystallin and α-LA were undertaken using methods outlined previously [22,35,36]. Briefly, α-LA (2 mg/ml) was dissolved in 50 mM phosphate buffer, 100 mM NaCl, 2.5 mM EDTA at pH 7.2, and incubated at 37 °C. Dithiothreitol (DTT, 20 mM) was added to the solution to induce reduction, unfolding and aggregation of the protein. Light scattering at 340 nm was monitored over time using a Cary 5000 UV/Vis/NIR spectrophotometer (Varian, Melbourne, Australia). The DLS measurements were recorded using a Zetavizer Nano ZS (Malvern Instruments, Worcestershire, UK). Accumulation times for each sample were determined automatically, and the temperature was controlled at 37.0 °C±0.1. The inbuilt software used the correlation function to calculate the z-average (intensity mean) hydrodynamic diameter ($D_H$), and the translational diffusional coefficient ($D_v$). The distributions of hydrodynamic diameters were calculated according to the Stokes-Einstein equation:

$$\frac{D_H}{kT} = \frac{3\pi\eta D_v}{4}$$  \hspace{1cm} \text{Equation 1}

where $k$ is the Boltzmann constant, $T$ is the absolute temperature, and $\eta$ is the solvent viscosity. The diameter measurements were converted to measurements of the hydrodynamic radius ($R_H$) for ease of comparison with the radius of gyration ($R_g$) measurements determined from the X-ray scattering data.

**Light scattering monitoring of the chaperone action of αB-crystallin against heat-stressed MDH**

Solutions containing 50 mM phosphate buffer, 100 mM NaCl, 2.5 mM EDTA at pH 7.5 were used. Two solutions were studied; the first contained 0.25 mg/ml MDH only, and the second contained 0.25 mg/ml MDH and 0.1 mg/ml αB-crystallin, i.e., a 2.5:1.0 w:w ratio of MDH: αB-crystallin. These concentrations are consistent with previous MDH and αB-crystallin chaperone studies [26,38]. The solutions were placed in 1 ml quartz cuvettes pre-heated to 42 °C in a Digilab Hitachi U-2800 spectrophotometer for 112 min and light scattering was measured at a wavelength of 360 nm The solutions were left for 5 min to equilibrate at 42 °C, as measured by a thermocouple (Cormark electronics Ltd., Littlehampton, UK).

**X-ray solution scattering measurements**

Low-angle X-ray solution scattering experiments were conducted at two synchrotrons. Initially, beamline X33 at the European Molecular Biology Laboratory (EMBL) of the Deutsches Elektronen Synchrotron (DESY), Hamburg, Germany, was used. The wavelength was $\lambda=1.5$ Å, with a sample to detector distance of 2.4 m, covering a scattering range of $0.09 \text{nm}^{-1} < q < 4.98 \text{nm}^{-1}$ ($q=4\pi \sin(\theta)/\lambda$). The detector was calibrated using the in-house EMBL software. Experiments were also undertaken at Station 2.1 of the Daresbury Synchrotron Radiation Source, Warrington, UK. The camera length was 5.25 m, with a corresponding scattering range of 0.00056 nm$^{-1}$ < $q$ < 0.72 nm$^{-1}$. The wavelength was $\lambda=1.5$ Å, and the detector was calibrated using hydrated rat tail tendon.

As with the light scattering assays, all experiments were performed in 50 mM phosphate buffer, 100 mM NaCl, 2.5 mM EDTA at pH 7.2. After an initial X-ray exposure, DTT was added (to a final concentration of 20 mM) to the solutions containing α-LA alone (2 mg/ml) and α-LA combined with αB-crystallin, and the samples were monitored with time. Samples containing mixtures of αB-crystallin and α-LA were prepared at 1:0, 1:1, and 1:10 w:w ratios, giving protein concentrations of 2 mg/ml αB-crystallin:0 mg/ml α-LA, 2 mg/ml αB-crystallin:2 mg/ml α-LA, and 2 mg/ml αB-crystallin:0.2 mg/ml α-LA, respectively. From our previous X-ray solution scattering studies [30,31], we found that the optimum signal to noise ratio of αB-crystallin and target protein is achieved with concentrations between 2 and 4 mg/ml. An initial X-ray exposure was acquired before the addition of DTT, then every 8 min for a total time of 136 min. The temperature was maintained at 37 °C.

For the MDH studies, X-ray exposures were taken every 8 min for a total of 112 min for solutions containing 2 mg/ml of MDH in the absence and presence of 2 mg/ml αB-crystallin in the same buffer as per the light scattering experiments. The temperature was maintained at 42 °C.

For both sets of experiments, the exposure times were 60 s each, and the temperature was kept constant with a thermostated circulating bath pumping fluid through the sample holder. The X-ray data were recorded and analyzed using the PRIMUS software package [40]. In all cases, sector integrations were performed with the origin at the position of the direct beam masked by the backstop. This method improves the signal-to-noise ratio at higher scattering angles, which is of particular significance for weakly scattering samples such as proteins in solution. All intensity profiles were corrected for background scattering. In all experiments, Guinier analysis was used to determine the average radius of gyration ($R_g$) of the protein aggregates as a
function of temperature following our earlier work [30, 31]. The $R_g$ value is derived from the Guinier region of the solution scattering X-ray intensity profiles. Such regions were used to fit the Guinier approximation Equation [41].

$$I = I_0 \exp (-4p^2 S^2 R_g^2 / 3) \quad \text{Equation 2}$$

Where, $I$ is the scattered intensity and $I_0$ the forward scattered intensity and $S$ is inverse square. The $R_g$ value is the root mean square distance of the electrons of the molecules in solution from the centers of their electronic masses and, therefore, is a measure of the overall size of the molecules. All $R_g$ values were found using PRIMUS software and satisfied the Guinier condition of $R_g \approx 1.3$. The ratio of $R_g$ and $R_H$, the hydrodynamic radius, leads to the dimensionless parameter $\rho$, i.e.

$$\rho = \frac{R_g}{R_H} \quad \text{Equation 3}$$

which is strongly dependent on the shape of the molecule [42].

**Results**

**αB-crystallin and α-LA**

**Light scattering experiments**— When reduced with DTT, α-LA (2 mg/ml) at 37 °C underwent amorphous aggregation and precipitation as monitored by light scattering at 340 nm (Figure 1). There was a lag phase of ~30 min followed by an exponential increase in light scattering over the remaining 200 min. The addition of α-crystallin at a 1:1 w:w ratio of α-LA:α-crystallin completely suppressed this increase in light scattering indicating that the chaperone prevents the precipitation of α-LA. Under these conditions, very similar behavior was observed in previous studies of the interaction between these two protein [22, 43]. α-B-Crystallin interacts in a chaperone manner with partially unfolded, reduced and monomeric α-lactalbumin forming a complex with it and thereby preventing its large-scale aggregation and precipitation. Our previous work has shown that complexation between the two proteins is established very quickly following the addition of DTT [24-28].

As monitored by DLS, the Z-average hydrodynamic radius of α-LA particles in solution, when incubated in the absence of α-crystallin increased immediately following the addition of DTT (i.e., there was no discernable lag phase; Figure 1). The lack of a lag phase in the DLS measurements compared to monitoring α-LA aggregation by light scattering at 340 nm (Figure 1) is reflective of the ability of DLS to detect very small changes in particle size that do not result in detectable levels of light scattering. As shown in Figure 2, the DLS measurements also allow for the determination of the $R_H$ value of individual peaks that contribute to the Z-average hydrodynamic radius, as well as the heterogeneity of each peak (as determined from the width of each peak at its base). The Z-average hydrodynamic radius of α-LA particles present in solution increased over the time-course of the assay, from ~40 Å (after 3 min, the first reading after DTT was added) to 860 Å by the end of the 200 min assay (Figure 1). When the individual components that contribute to this Z-average radius were examined, this increase was observed to be due to the formation of increasingly large and polydisperse mixtures of α-LA aggregates with $R_H$ values > 1000 Å (Figure 2B).

The Z-average hydrodynamic radius of α-crystallin alone in solution was 85±18 Å. The α-crystallin oligomers were found to range in size from a $R_H$ value of ~50 Å to 190 Å (based on the width of the peak at its base, see Figure 2A). When α-crystallin was added to DTT-treated α-LA at a 1:1 w:w ratio, i.e., conditions under which large scale aggregation of α-LA is prevented (Figure 1), there was a small increase in the Z-average radius of the solution from 82 Å to 100 Å over the first 60 min which then remained constant for the remaining 90 min of the assay (Figure 1). Figure 2C, which shows the size of the particles in solution, indicates that at 50 min there were two predominant species present in the 1:1 mixture of α-LA:α-crystallin, i.e., a smaller sized aggregate ($R_H$ ~100 Å with a peak width from 40 Å to 300 Å) and a minor, larger-sized aggregate ($R_H$>750 Å). In their studies of the interaction of α-LA and α-crystallin under reduction conditions, Bettleheim et al. [44], also observed these two different populations and demonstrated, using size exclusion HPLC and SDS-PAGE, that the peak at ~100 Å represents the complex formed between α-crystallin and α-LA and that the larger peak ($R_H$>750 Å) is attributable to aggregated α-LA alone. Interestingly, the investigations here show that when α-crystallin was present, the larger-sized aggregate of α-LA remained of similar size for the remainder of the assay. Moreover, the amount of this α-LA aggregate, as a proportion of the total number of particles in solution, remained small (i.e., ~14%; Figure 2C) unlike when the chaperone was not present (Figure 2B), where the large aggregates ($R_H$>1000 Å) represented most (i.e., >95%) of the particles in solution.

**X-ray scattering experiments**— X-ray scattering experiments of β-crystallin and α-LA solutions on their own and initially without the addition of DTT, each at 2 mg/ml, gave $R_g$ values of 58.2 (±0.1) Å and 22.4 (±0.2) Å respectively. The large difference in $R_g$ values between the two proteins is consistent with the much greater size of the β-crystallin oligomer (~650 kDa in mass) compared to the α-LA monomer (~14 kDa). Figure 3 shows the X-ray intensity scattering profiles of reduced α-LA, 32 and 72 min after the addition of DTT. To highlight the scattering features at higher angles, the intensity is plotted on a logarithmic scale. The increase in the X-ray intensity at low angles with time is an indication of aggregation [30, 31]. From Figure 3, it is apparent that in the region beyond $q=2$, the X-ray data become noisy and imprecise. In monodisperse systems, this is the region where information is obtained from the particle shape and Fourier transform and a shape reconstruction is feasible. From Figure 3, it is apparent that Fourier transforms from different-sized aggregating particles are overlaid and smear out the data meaning that shape reconstruction is not feasible, as is also the case in the polydisperse β-crystallin and α-LA mixtures under reducing conditions (data not shown).

The initial $R_g$ value of reduced α-LA (from data acquired as soon as possible after addition of DTT) is 26.9 (±0.2) Å which exhibits a steady increase with time (particularly after 50 min associated with the formation of large light scattering aggregates, Figure 1) to a value of 108.0 (±0.2) Å at 104 min and 185.0 (±0.6) Å after 136 min (Figure 4). In contrast, the solution containing β-crystallin and reduced α-LA at a 1:1 w:w ratio showed only a slight increase in the $R_g$ value from 55.1 (±0.1) Å initially to 60.4 (±0.1) Å 104 min after addition of DTT (Figure 4), through to a value of 78.2 (±1.1) Å at 136 min after addition of DTT, i.e., significantly lower than that measured for α-LA in the absence of β-crystallin. From our previous work [34, 36], it is well established that under these conditions and time frame, β-crystallin is an efficient chaperone for reduced α-LA (see Figure 1) whereby it interacts and complexes with α-LA to prevent it from aggregating. Thus, these $R_g$ values correspond to the complex formed between the two proteins and are not simply due to averaging of the $R_g$ values of the two components present. Evidence for this is that if the $R_g$ values simply reflected the aggregation of the two components, they would increase greatly over time due to the unfolding and aggregation of the reduced α-LA.

The initial $R_g$ value of the solution containing a 1:10 w:w ratio of the β-crystallin and α-LA mixture was 37.4 (±0.2) Å, which, because of the excess of α-LA, was much lower than the initial $R_g$ value (55.1 (±0.1) Å) of the 1:1 w:w ratio (Figure 4). The $R_g$ values for the 1:10 mixture then increased significantly with time following addition of DTT, reaching a value of 133.0 (±1.4) Å after 136 min, which is consistent with the inability of β-crystallin to function efficiently as a chaperone for reduced α-LA at this sub-stoichiometric ratio [36]. As a result, significant aggregation and precipitation of reduced α-LA occurs, as we have previously demonstrated [34, 36].

Figure 5 shows Kratky plots at the start and end of the X-ray solution scattering experiments for β-crystallin alone (Figure 5A), α-LA plus DTT (Figure 5B,C), α-crystallin plus α-LA at the 1:1 w:w ratio (Figure 5D,E) and α-crystallin plus α-LA at the 1:10 w:w ratio (Figure 5F,G). For these Kratky plots, the X-ray intensity scattering profiles (Figure 3) are plotted as $I(q)q^2$ against $q$, where $I$ is the scattered intensity and $q=4\pi\sin(\theta)/\lambda$. As can be seen in Figure 5A, β-crystallin alone shows a peak centered around $q=0.3$. The presence of such a peak indicates that the majority of particles in solution are globular [45] and therefore approximately spherical in shape. From Figure 5B,C, it is apparent that α-LA loses its globular structure upon unfolding, aggregation and precipitation associated with the reduction of its disulfide bonds and the adoption of a molten globule conformation [34, 36]. As a result, the peak centered at $q=0.7$ at 0 min in its Kratky plot is lost by 136 min after
addition of DTT. By contrast, the presence of a peak at q = 0.3 in the plots at 136 min for the α-B-crystallin plus α-LA mixtures at both the 1:1 and 1:10 w:w ratios (Figure 5E, G) clearly indicates that, under both conditions, the α-crystallin/α-LA complex has a spherical shape.

**α-B-crystallin and MDH**

**Light scattering experiments**— The chaperone activity of α-B-crystallin in solution under mild heating conditions at 42 °C was investigated with MDH as the target protein. Figure 6 shows that the light scattering of MDH at time with 42 °C is exponential following a lag period of 15 min. By contrast, the light scattering of a 2.5:1.0 w:w solution of MDH:α-B-crystallin shows only a very minimal increase over the same period, demonstrating that α-B-crystallin acts as a molecular chaperone to prevent the temperature-induced, partial unfolding and aggregation of MDH.

**X-ray scattering experiments**— Figure 6 also shows the half width at half maximum of MDH alone and for a 1:1 w:w mixture with α-B-crystallin during incubation at 42 °C. The initial mean value of MDH is 47.7 (+/−0.3) Å which exhibits a steady increase with time to a value of 59.9 (+/−0.4) Å at 48 min. The half width value then increases rapidly to 71.1 (+/−0.5) Å at 56 min. After this, the half width value increases further to 83.9 (+/−0.8) Å at 104 min. The initial mean value of a 1:1 w:w mixture of MDH and α-B-crystallin is 53.8 (+/−0.1) Å. As with the mixture of α-LA and α-B-crystallin, this value is lower than the half width of MDH alone (58.2 (+/−0.1) Å), which suggests that the α-B-crystallin MDH molecules are not as large in the average half width value for both proteins in solution. At the initial time point, it is unlikely that the two proteins strongly interact as the MDH molecules will not lower than that for α-LA. Thus, due to the heterogeneity of the two systems, the parameter (i.e., half width) of the MDH complex was found to have a range of 65 to 100 Å (P. Stewart, Vanderbilt University, Nashville, TN personal communication) compared to that previously reported for the α-crystallin oligomer (comprising both the α- and α-crystallin subunits) as measured by DLS [44]. Thus, these experimental techniques all gave very similar results with respect to the size of α-B-crystallin oligomer.

The DLS and X-ray solution scattering data for reduced α-LA and its 1:10 w:w mixture with α-B-crystallin imply that α-B-crystallin interacts with destabilized, partially unfolded α-LA molecules early on in the latter’s aggregation (off-folding) pathway, i.e., well before large scale aggregation occurs for reduced α-LA. These findings support our previous conclusions determined from NMR studies of the interaction of these two proteins [34–36]. Our present data confirm that the interaction between the two proteins is established very quickly following addition of DTT since there is no significant change in the half width and radius values with time from those obtained immediately after the addition of the reducing agent. For example, the DLS measurements indicate that reduced α-LA and α-B-crystallin form a stable complex (of RH ~100 Å) within the dead time of the experiment (~5 min), and this prevents the increase in α-LA aggregate size that leads to precipitation when the chaperone is absent. Bettleheim et al. [44] reported similar results for the interaction of α-crystallin with reduced α-LA. In addition, in the presence of α-B-crystallin, the larger α-LA aggregate (RH =750 Å) does not continue to grow in size, as occurred when the chaperone was absent (Figure 2B and Figure 4C), and it constitutes only a small percentage of the total number of particles in solution. These large particles are most likely too few in number to be detected by light scattering at 340 nm (due to its decreased sensitivity compared to DLS) and therefore a change in light scattering is not observed when α-LA is incubated in the presence of α-B-crystallin. Higher concentrations of α-crystallin completely suppress the formation of this aggregated form of α-LA [44,52].

Other spectroscopic and biophysical studies on this system have also come to the conclusion that α-B-crystallin acts early on to prevent the aggregation of destabilized α-LA molecules [34–36]. The X-ray solution scattering data for a 1:10 w:w ratio mixture of α-B-crystallin:α-LA show that time-dependent aggregation of this mixture (as monitored by the change in half width values) is only slightly inhibited compared to the situation with reduced α-LA and α-B-crystallin. The α-LA dependent aggregation almost doubles during incubation at 37 °C from a 1:10 w:w ratio of the two proteins (Figure 1, Figure 2, and Figure 4). The rationale for these observations is that at a sub-stoichiometric 1:10 w:w ratio, α-B-crystallin is not capable of completely suppressing reduced α-LA aggregation whereas it does so very effectively at a 1:1 w:w ratio [36].

Interestingly, our DLS studies showed that the size of the α-crystallin-α-LA complex (i.e., a RH value of 100±22.5 Å) was similar to that of the α-crystallin oligomer alone (a RH value of 85±18 Å) although the target protein-chaperone complex was more heterogeneous (compare Figure 2A and Figure 2C). In support of our DLS measurements, cryoEM images of the α-crystallin-α-LA complex also show that it is comparable in size to the α-crystallin oligomer (see Figure 7 in Hales et al. [53] and Figure 2 in Horwitz [31], i.e., the radius of the α-crystallin-α-LA complex was found to be within a range of 65 to 100 Å (P. Stewart, Vanderbilt University, Nashville, TN personal communication) compared to that of the α-crystallin oligomer itself of 40 to 90 Å [27,53]. Horwitz et al. [39] also found no significant difference in the size of the α-crystallin-α-LA complex (compared to the α-crystallin oligomer alone) by gel filtration chromatography. By contrast, at the end of the X-ray scattering experiment we found that the 1:1 w:w mixture of reduced α-LA and α-B-crystallin had a RG value of 78.2 (+/−1.1) Å (compared to α-crystallin alone (58.2 ±0.1) Å). In agreement with the latter value, Skouri-Panet et al. [54] used SAXS to examine the temperature and pressure-dependent changes in the structure of α-sHsps and found that a 3.7 mg/ml solution of α-crystallin had a RG value of 61.1 ± 23 °C, in a solution containing a phosphate buffer at pH 6.8. On the other hand Spinozzi et al. [55] found the RG value to be 52 Å in a solution containing a TRIS buffer. The slight variation in RG values between this study and ours most likely arises from the different buffer conditions used since the size of the α-crystallin oligomer is dependent on solvent conditions [56].

Intuitively, one would expect that an association between the two proteins during α-sHsp chaperone action would lead to the formation of a complex of increased size, yet we have measured size that, while being more polydisperse, there is no significant increase in the size of the α-crystallin-α-LA complex compared to the α-crystallin oligomer alone, but there is an increase in the RG value of the α-LA complex. To rationalise the DLS, cryoEM and X-ray solution scattering data, we must consider the parameter RH, which relates the radius of gyration (RG) with the hydrodynamic radius (RH; see Equation 3) and describes the distribution of mass and shape of the molecule. The calculated standard deviation of p is 22% due to the polydispersity of the α-B-crystallin oligomer and the α-B-crystallin-α-LA complex and therefore the range in RG values of these particles in solution. Based on the DLS and X-ray solution scattering data at 150 min, the p value for α-crystallin alone is 0.68±0.14 (i.e., RG=58.2±0.1 Å and RH=85±18 Å) and for the α-crystallin-α-LA complex, p is 0.78±0.18 (i.e., RG=78.2±1.1 Å and RH=100±22.5 Å). Thus, due to the heterogeneity of the two systems, the parameter p is the same and corresponds to a value approximating that of a solid sphere (i.e., p, solid sphere = 0.76, p, hollow sphere = 1.0) [42,57].

In other words, the distribution, size and shape of the α-crystallin oligomer are very similar whether it is bound target protein or not, since the hydrodynamic radius (RH; Figure 1 and Figure 2) and spherical shape (p value and Kratky plots, Figure 5) are not significantly altered upon complex formation between α-crystallin-α-LA complex.

Our findings that the globular (spherical) shape of α-B-crystallin is not capable of completely suppressing reduced α-LA aggregation whereas it does so very effectively at a 1:1 w:w ratio [36].

Discussion

α-B-crystallin is a member of the Hsp family that shares a conserved approximately 90 amino acid ‘α-crystallin’ domain and acts as a molecular chaperone protein by preventing the stress-induced aggregation and precipitation of target proteins. Crystallographic structures of two non-metazoan α-sHsps are available [1,46] along with the structure of a Hsp from a flatworm, Tsp36 [47]. Recently, the crystal structure of the mammalian ‘α-crystallin’ domain has been solved [48]. Several structural similarities exist between these α-sHsps, most notably for the two non-metazoan α-sHsps, which are both approximately spherical aggregates with a large central cavity. The α-crystallin oligomer also has a cavity at its center, as determined by EM studies [27,49,50], and as was predicted by simple consideration of mass distribution in the aggregate [51]. The conclusions from the cryoEM studies agree very well with those derived from the solution-based DLS and X-ray scattering data presented herein for the α-B-crystallin oligomer. The mass of the α-crystallin monomer is ~20 kDa and the cryoEM data were acquired on samples which contained α-crystallin oligomers of mass around 650 kDa, i.e., comprising approximately 32 subunits [49]. The cryoEM data indicate that the diameter of the α-crystallin oligomer is 147 (±28) Å, i.e., a radius of 74 (±14) Å [49]. In agreement with these values, from our DLS studies (Figure 2), we found that α-B-crystallin has a 2-average hydrodynamic radius (RH) of 85±18 Å, which is also very similar to that previously reported for the α-crystallin oligomer (comprising both the α- and α-crystallin subunits) as measured by DLS [44]. Thus, these experimental techniques all gave very similar results with respect to the size of α-B-crystallin oligomer.
following formation of a complex with α-LA (see Figure 5) is consistent with previous cryo-EM studies [53]. During chaperone action, this most likely arises from positioning the reduced α-LA molecules on the surface of the αB-crystallin aggregate [3,39,53] i.e., within the oligomer’s protein shell. The partially folded α-LA molecules could be located in the fenestrations that are on the surface of the αB-crystallin oligomer [31] and the non-mammalian shsp oligomers [1,46,57]. As a result, the target protein would be readily accessible for refolding via the action of other molecular chaperones (e.g., Hsp70), in a process that requires ATP hydrolysis, when cellular conditions allow [58]. The dynamic, flexible and maleable nature of the αB-crystallin oligomer, particularly on its surface where the flexible COOH-terminal extensions are located [28,49,51] would facilitate the incorporation of the target protein (in this case α-LA) within its outer protein shell [5]. Indeed, our NMR studies show that the flexibility of the COOH-terminal extension is altered significantly upon binding of α-LA [56] implying that the extensions and bound α-LA molecules are localized nearby to each other. Stengel et al. [61] have recently used sophisticated mass spectrometry methods to investigate the oligomeric states of a non-mammalian shsp, pea Hsp18.1, when interacting with a target protein, luciferase, under mild temperature stress. Unlike αB-crystallin, Hsp18.1 adopts a well defined oligomer (a 12-mer) in the absence of target protein at room temperature. Upon chaperone interaction with luciferase, however, Hsp18.1 forms a highly heterogeneous range of complexes containing different stoichiometries of luciferase and Hsp18.1. It is proposed that this temperature-induced heterogeneity of Hsp18.1 facilitates its interaction with a range of target proteins and also enables the target proteins to be readily accessible for refolding upon complexation. For αB-crystallin, the polydispersity is already present at physiologic temperatures which enables it to readily interact with target proteins in the absence of temperature stress. As a result, αB-crystallin is ‘primed’ for interaction with a diversity of target proteins. Indeed, X-ray crystallography has recently revealed that the polydispersity of αB-crystallin is facilitated by the presence of a nine amino acid palindromic sequence centered around P160 of the COOH-terminal region that participates in inter-subunit interactions via alignment in both directions of its sequence in both directions [62].

Similar results were observed in comparing the X-ray scattering results (Rg values versus time) for the two target proteins (α-LA and MDH) in the presence of αB-crystallin (Figure 4 and Figure 6, respectively). Thus, the interaction between the target proteins and αB-crystallin under different mild stress conditions (reduction at 37 °C and heating at 42 °C) leads to a stable complex that has a Rg value that varies little with time and is not significantly different to that of the αB-crystallin oligomer on its own. It is concluded that the arguments presented above for the interaction of α-LA with αB-crystallin also apply for the interaction of MDH with αB-crystallin. Using small-angle neutron scattering (SANS) experiments in conjunction with isotopic substitution and contrast matching techniques, we recently investigated the interaction of a lens target protein, γc-crystallin, with both the α-crystallin oligomer, under harsh thermal stress, i.e., 65 °C [33]. Under these conditions, we concluded that at γc-crystallin is located within the central cavity of the α-crystallin oligomer. As discussed above, 65 °C is well above the temperature at which α-crystallin undergoes a major rearrangement of its secondary, tertiary and quaternary structures which, coupled with the protein’s inherent dynamism and its porous nature due to the ‘fenestrations’ on its surface [27,30,49,50], would facilitate ready access of target proteins to the central cavity of the α-crystallin oligomer. Thus, α-crystallin may have different protective chaperone mechanisms depending on the stress conditions i.e., target proteins depending bind within the central cavity or to the surface of the outer shell depending on external environmental factors, including the size of the target protein and whether the target protein is aggregating amyorphously or to form amyloid fibrils, the pH, the type of stress (e.g., reduction or elevated temperature), the rate of target protein aggregation and the temperature. Indeed, our studies have shown the importance of these factors in determining the efficiency of chaperone action of αB-crystallin against target proteins [28,50,63,64].

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