Influence of the chirality of short peptide supramolecular hydrogels in protein crystallogenesis†

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For the first time the influence of the chirality of the gel fibers in protein crystallogenesis has been studied. Enantiomeric hydrogels 1 and 2 were tested with model proteins lysozyme and glucose isomerase and a formamidase extracted from B. cereus. Crystallization behaviour and crystal quality of these proteins in both hydrogels are presented and compared.

Chirality is ubiquitous in nature and has major implications in cell interactions and biological processes. Within this topic, chiral discrimination on chiral surfaces has significant effects in protein adsorption and cellular adhesion and proliferation.1 It would be expected that similar chiral discrimination was feasible with other common macromolecular hydrogels, which either have no stereocenters (polyacrylamide) or are constituted diastereoisomerically with proteins, thus expecting different behaviors. It is worth noting that the structural simplicity of short peptides is essential for this purpose, allowing an easy access to their corresponding enantiomers. This property is not feasible with other common macromolecular hydrogels, which either have no stereocenters (polyacrylamide) or are constituted by the natural chiral product (agarose).

To test this hypothesis we selected two cysteine-based peptides, compound 1 (N,N'-di(benzoyl)-L-cysteine diamide) and its corresponding D enantiomer 2 (N,N'-di(benzoyl)-D-cysteine diamide) (Fig. 1) due to their capacity to self-assemble in neat water to give the corresponding hydrogels. They were tested with two model proteins, chicken HEWL lysozyme and glucose isomerase, and one target protein, a formamidase extracted from B. cereus being the first time that supramolecular peptide based hydrogels are used for this purpose. Additionally, the results were compared with crystal

† Electronic supplementary information (ESI) available: General experimental details, 1H NMR and 13C NMR of hydrogels 1 and 2, VT-NMR and DSC spectra of hydrogels 1 and 2 and crystallization and crystallography data. See DOI: 10.1039/c4cc09024a

Fig. 1 Characterization of hydrogels 1 and 2 by TEM microscopy, circular dichroism and rheology. TEM images of negatively stained dried hydrogels at 3 mM: (A) 1, (B) 2. Scale bar at 0.2 μm. (C) CD spectra and (D) rheology of hydrogels 1 and 2.

†† Open Access Article. Published on 29 January 2015. Downloaded on 13/01/2016 15:21:43.

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COMMUNICATION

ChemComm

Cite this: Chem. Commun., 2015, 51, 3862

Chem. Commun., 2015, 51, 3862--3865 This journal is © The Royal Society of Chemistry 2015

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growth in agarose gels to investigate their potential in protein crystallization.

Compounds 1 and 2 were selected from a family of 22 cysteine derivatives based on its capacity to form a stable and completely transparent hydrogel. TEM images of hydrogel 1 showed self-assembled nanofibers of diameters ranging from 10 to 20 nm and lengths that go up to μm in size (Fig. 1). Fibrous aggregates co-exist with regions where fibers are more dispersed. Hydrogel 2 presented fibers of the same aspect. To gain insight into the molecular arrangement of these hydrogels circular dichroism (CD) spectra were recorded (Fig. 1). Hydrogel 1 exhibited a negative band near 218 nm (n* transition), a positive band near 250 nm (n* disulfide band and/or π* of the aryl groups) and a negative band near 280 nm (π* of the aryl groups) in good agreement with the CD spectrum from other oligopeptide hydrogels identified as the β-sheet conformations. The CD spectrum of hydrogel 2 is the specular image of hydrogel 1 showing that the chirality of the peptides is conserved and transferred supramolecularly to the fibers. Hydrogels 1 and 2 were also analyzed using a regime of simple oscillatory shear stress through frequency (see ESI†) and amplitude sweep tests. Both hydrogels showed G' values (elastic modulus) higher than G" values (viscous modulus) indicating the predominant elastic behavior of the gels (Fig. 1). Based on the magnitude of the values of G', G" and σc (G' ≈ 30 Pa; G" ≈ 4 Pa; σc < 1 Pa) we can conclude that these hydrogels are extremely weak. The similar values of G' and G" for hydrogels 1 and 2 confirm that both enantiomeric hydrogels have the same rheological properties. VT-NMR studies on both hydrogels showed that the signals of the aromatic region were shifted 0.5 ppm downfield as the stress through frequency (see ESI†) and amplitude sweep tests. Both hydrogels showed G' values (elastic modulus) higher than G" values (viscous modulus) indicating the predominant elastic behavior of the gels (Fig. 1). Based on the magnitude of the values of G', G" and σc (G' ≈ 30 Pa; G" ≈ 4 Pa; σc < 1 Pa) we can conclude that these hydrogels are extremely weak. The similar values of G' and G" for hydrogels 1 and 2 confirm that both enantiomeric hydrogels have the same rheological properties. VT-NMR studies on both hydrogels showed that the signals of the aromatic region were shifted 0.5 ppm downfield as the temperature increased from 25 °C to 70 °C. The small values of Tg (1.84 s for the 7.74 ppm signal) and the negative values of nOe indicated that the 1H-NMR signals corresponded to aggregates of the di-peptide which are not higher enough to be NMR-silent. The stability of these aggregates indicated that Tg should be higher than 70 °C. In fact in the 75/25 H2O/DMSO mixture, the reported value of Tg for 1 was above 90 °C. 8 DSC scans in both directions of hydrogels 1 and 2 did not show any endothermic or exothermic peaks. These results indicated continuous gel to sol and sol to gel transitions (see ESI†).

The two enantiomeric hydrogels 1 and 2 were then tested as the supramolecular media for protein crystallization. Following the procedure described in ESI†, protein was allowed to diffuse in the gel prior to the addition of the precipitant. Fig. 2A shows a typical counter-diffusion pattern of lysozyme crystals grown in hydrogel 1. The effect of hydrogels 1 and 2 on the quality of lysozyme crystals was evaluated from a set of lysozyme crystals prepared at beam-lines Xaloc (ALBA) following the standard quality criterion. 11

Lysozyme crystals grown in hydrogel 2 were better than those obtained in hydrogel 1, reaching the resolution limit of the detector (0.95 Å) and having the best quality indicators in terms of I/σ(I), Rmerge and mosaicity (Fig. 2B). The excellent reproducibility of crystals grown in hydrogel 2 (Fig. S1 and Table S2, ESI†) is also worth mentioning. Moreover crystal quality indicators of crystals grown in agarose (able to produce protein crystals of high quality5αd) were similar to those obtained in hydrogel 1 but still inferior to those in hydrogel 2. The agarose gel has been proven to facilitate the soaking of different compounds5αd including cryoprotectants. In order to evaluate the possible cryo-protection capacity of these new hydrogels we compared the crystal quality of cryo-protected and “naked” (flash frozen without cryoprotectants) lysozyme crystals grown in hydrogel 1 and agarose from two data sets collected at beam-lines ID23 and BM30 (ESRF). It is important to notice that the polymeric nature of both agarose and hydrogel 1 allowed us to collect full data sets without the need to use any additive (Fig. S2, S3 and Table S3, ESI†).

A similar analysis was performed with glucose isomerase crystals grown in hydrogels 1, 2 and agarose. In this case crystals obtained in hydrogel 1 and agarose were more homogeneous and of better quality than those grown in hydrogel 2 (Fig. 2C, Table S4 and Fig. S4, ESI†). However, a second crystal form (P212121) was identified only in the experiments run with hydrogel 2 that co-exist with the other polymorph (P2222), the form normally found under similar crystallization conditions. Dauter and co-workers have described a new crystal form (P212121) of glucose isomerase and have solved its structure (PDB ID. 1OAD). Remarkably, the polymorph found in hydrogel 2 presents different unit cell parameters (Å): 86.00, 93.68, 99.22 versus 98.45, 129.59, 78.33 for a, b and c, respectively (Fig. S4, ESI†). To date there has been only one PDB entry (PDB ID. 1O1H) with a similar unit cell and space group but obtained under different crystallization conditions.

The quality of this primitive orthorhombic crystal was the highest of all glucose isomerase crystals tested, with the structure determined at 1.20 Å and refined to R/Rfree values of 10.73/13.22 (%) (Table S5, ESI†). When compared with the model used for molecular replacement (PDB. ID 1OAD) the deviation of the main chain...
is minimal (rmsd of 0.3). It is worth mentioning that this new polymorph has only been obtained in free solution using other precipitants, as in the case of PDB ID. 101H that crystallized using MPD, or in the presence of high agarose concentration.\textsuperscript{1a} In our case, under identical crystallization conditions, the $P_{2_1}2_12_1$ polymorph was obtained only with hydrogel 2 and not in agarose at 0.5% (w/v), our control, suggesting a novel interaction between the chiral gel fibers and the protein at the nucleation stage. It has been previously demonstrated that the nature of the gel fibers, such as agarose\textsuperscript{15} and silica,\textsuperscript{16} affect protein nucleation density (promoted and inhibited, respectively) with no effect on the final crystal packing i.e. formation of new polymorphs. To date, the correlation between the protein–fiber interaction and polymorphism has been unknown. Taking into account that both effects opens a question on the different roles that gel nature can have in the whole process of protein crystallogenesis.

It is also known that the incorporation of hydrogel fibers into the protein crystal lattice occurs with agarose, silica and PEG hydrogels.\textsuperscript{5,15,16a,c} To prove that the same effect also occurs with peptide hydrogels, dissolution experiments of lysozyme crystals grown in hydrogel 1 were carried out (Fig. 3). This result showed that effectively the peptide fibers are incorporated within the protein crystals. The uncorrelated behavior of incorporation and nucleation effects opens a question on the different roles that gel nature can have in the whole process of protein crystallogenesis.

Finally, formamidase extracted from \textit{B. cereus} was selected as a target protein to test its crystallizability using hydrogels 1, 2, and agarose as controls. Good diffracting crystals were already obtained by the capillary counter diffusion method diffracting to a maximum resolution of 1.78 Å (Table S6, ESI\textsuperscript{†}). From previous experiments we knew that agarose used in combination with capillary counterdiffusion did not improve or even fully impedied crystallization. Crystals grown under three different crystallization conditions (Table S7, ESI\textsuperscript{†}) were evaluated. Only cryo-protected crystals obtained in hydrogel 2 under the condition C18 of the GSK crystallization screening kit (Triana SBT, Granada, Spain) produced the best crystals to date (1.4 Å resolution limit, Table S6, ESI\textsuperscript{†}). On the other hand, hydrogel 1 inhibited nucleation while crystals grown in agarose were of low quality proving the different behavior of these hydrogels.

To sum up, we present the novel use of supramolecular hydrogels based on short-peptides in neat water for protein crystallization. These hydrogels, made of symmetrical L and D di-cysteine derivatives, were used to grow protein crystals of high quality. Differences in crystal quality and packing for the three proteins have been observed using enantiomeric hydrogels 1 and 2, suggesting a significant influence of the fibers’ chirality in the crystallization process. The formation of the orthorhombic polymorph of glucose isomerase and the growth of the highest quality crystals of formamidase exclusively in hydrogel 2 support this hypothesis. This fact highlights the relevance that chirality may have in protein crystallogenesis for X-ray structural determination. Moreover these supramolecular peptide based hydrogels are excellent mediators for protein crystallization producing lysozyme and glucose isomerase and formamidase crystals with excellent quality indicators. Peptide based hydrogels also acted as cryo-protectants avoiding the use of additives. Hence, these novel chiral gels would expand the field of protein crystallogenesis.

This research was funded by the MICINN (Spain) projects BIO2010-16800 (JAG), CTQ-2011.22455 (LAC & JMC), CTQ2012-34778 (JJDM & ALG), “Factoría Española de Cristalización” Consolider-Ingenio 2010 (JAG & MCM) and EDRF Funds (JAG, LAC & JMC), P12-FQM-2721 (LAC) Junta de Andalucía. JJDM thanks MICINN for a R&C Fellowship and MCM thanks CSIC for her JAE Fellowship. We would like to thank Dr S. Martinez-Rodriguez for providing the plasmids of formamidase and B. Morel and E. Sánchez-Cobos for their help with the CD studies. We are very grateful to the staff at beam-line BM30, ID29 (ESRF) and Xaloc (ALBA) for support during X-ray data collection.

Notes and references


![Fig. 3](image-url) Dissolution sequence of a lysozyme crystal grown in hydrogel 1. Crystal extracted from the gel was placed in a water drop and allowed to dissolve (t = 0 s). A red square is drawn around the remaining hydrogel. The last picture was taken after partially drying the remaining gel framework.


