

Cite this: *Nanoscale Adv.*, 2021, 3, 1789

Folding and self-assembly of short intrinsically disordered peptides and protein regions

Pablo G. Argudo *^a and Juan J. Giner-Casares ^b

Proteins and peptide fragments are highly relevant building blocks in self-assembly for nanostructures with plenty of applications. Intrinsically disordered proteins (IDPs) and protein regions (IDRs) are defined by the absence of a well-defined secondary structure, yet IDPs/IDRs show a significant biological activity. Experimental techniques and computational modelling procedures for the characterization of IDPs/IDRs are discussed. Directed self-assembly of IDPs/IDRs allows reaching a large variety of nanostructures. Hybrid materials based on the derivatives of IDPs/IDRs show a promising performance as alternative biocides and nanodrugs. Cell mimicking, *in vivo* compartmentalization, and bone regeneration are demonstrated for IDPs/IDRs in biotechnological applications. The exciting possibilities of IDPs/IDRs in nanotechnology with relevant biological applications are shown.

Received 10th November 2020
Accepted 17th January 2021

DOI: 10.1039/d0na00941e

rsc.li/nanoscale-advances

Introduction

Proteins and their fragments as short peptides are being revealed as relevant players in nanoscience and self-assembly.¹ Peptide derivatives can be designed to reach any nanostructure, from nanotubes² to vesicles.³ Given the excellent biocompatibility and the biological origin of peptides, the forefront field of application for self-assembled nanostructures based on peptides is biomedicine and controlled delivery.^{4,5} Other fields such as optical materials also benefit from protein-based nanostructures.⁶ The directed design of peptides can greatly enhance intrinsic properties and biological activity against bacteria.^{7,8} Hybrid derivatives including peptides for self-assembly allow the solubilization of hydrophobic molecules relevant in the field of biological applications as antimicrobial peptides.⁹ Purposefully designed peptides can show self-assembly upon being triggered by certain stimuli such as fluid forces.¹⁰ Stimuli can be used to promote the self-assembly of peptide-based nanostructures when required, *e.g.*, under a pH variation in tumor cells.¹¹ Peptide derivatives are also relevant when considering biomineralization at interfaces.^{12,13} Peptides can form hybrids with relevant nanostructures based on carbon for subsequent self-assembly into intriguing materials.^{14,15}

The structural features of proteins and peptide fragments are undoubtedly the focus of a large body of research. The traditional view assumes that proteins were flexible and reach a final shape depending on the targeting molecule.¹⁶ A more recent view accepts that proteins and peptides appear in an ensemble of

conformations with certain regions showing a higher degree of disorder. Thus, intrinsically disordered proteins (IDPs) can be described as the maximum degree of disorganization presented in a protein. IDPs are isolated polypeptide chains with no stable tertiary structure while still being functional. Note that not the whole protein necessarily lacks a fixed structure. The unstructured segments are the so-called intrinsically disordered regions (IDRs).^{17–19} IDRs were present in at least 2.0% of archaeal, 4.2% of eubacterial and 33.0% of eukaryotic proteins.²⁰ IDPs/IDRs are generally found in nature. The analysis of *ca.* 3500 protein species from three different domains of life (archaea, bacteria, and eukaryotes) and viruses allowed an estimation of the prevalence of IDPs/IDRs. Viruses had the widest spread of the proteome disorder content (7.3 to 77.3%). Eukaryotic cells present a higher ratio of IDPs/IDRs to the total content of proteins than prokaryotic cells. A higher eukaryotic proteome disorder might be used by nature to deal with the increased cell complexity due to the appearance of various cellular compartments.²¹

The state of the field and the current trends in the study of intrinsically disordered proteins and regions are summarized herein. This review is focused on short IDRs or completely disordered peptides. After a brief introduction of the structural basics of proteins and peptides, several examples of IDPs/IDRs are introduced. Relevant experimental and computational methods for characterization are discussed. An excellent book by Tanford and Reynolds is strongly recommended for a longer analysis of how protein structure characterization was approached at early stages.²²

Structures

Proteins are 'naturally occurring and synthetic polypeptides having molecular weights greater than about 10 000 Da'.²³ From

^aUniversité de Bordeaux, CNRS, Bordeaux INP, LCPO, 16 Avenue Pey-Berland, 33600 Pessac, France. E-mail: Pablo.Gomezargudo@enscbp.fr

^bDepartamento de Química Física y T. Aplicada, Instituto Universitario de Nanoquímica IUNAN, Facultad de Ciencias, Universidad de Córdoba (UCO), Campus de Rabanales, Ed. Marie Curie, E-14071 Córdoba, Spain



this IUPAC official definition, it can be clearly inferred that peptides are small chains of amino acids linked by peptide bonds.²⁴ Peptides can be folded into several structures due to the interaction between the atoms of the backbone, described as a secondary structure in proteins. The most common structures are α -helix²⁵ and β -sheet²⁶ structures, which also can appear along β -turns²⁷ and omega loops.²⁸ Other structures are also rarely found, e.g., 3_{10} -helix²⁹ or π -helix.³⁰ Proteins with a longer peptide chain allow subsequent folding leading to ternary and quaternary structures. No further folding but self-assembly processes appear in short peptides. Self-assembly is defined as the autonomous organization of components into ordered patterns or structures.³¹ The short peptide chains act as building blocks, adding value to the final structure. Several final structures can be obtained after a first folding and a second self-assembly organization. Examples of these highly ordered nanostructures can be nanoparticles, nanotubes, nanofibers, vesicles, gels or films.³² However, this behaviour cannot be directly extrapolated to IDPs/IDRs. Due to their completely disordered random coil structure, these peptides miss their folding process.³³ Thus, their final self-assembled conformation is going to be related not only to their chemical inner composition but also to the media that surround them and the interactions that take place.

In this section, we give a brief overview of the most extended folding and self-assembly structures that can be obtained for IDRs or peptides and the bindings or recognition processes responsible for the final conformation. Additional examples can be found in the following chapters of the review.

Helices

As the α -helix is the most extended structure, this peptide bonding organization is a mainly right-handed helical structure. The N–H group of each amino acid forms a hydrogen bond with the C=O group of the amino acid placed in the previous fourth position. This Pauling–Corey standard model can be expressed as $i + 4 \rightarrow i$. Specifically, the α -helix measures about 5.4 Å in width, has 3.6 amino acid residues per turn and each two continuous residues adopt a total dihedral angle (ϕ , ψ) of around -105° . Similar structures 3_{10} -helix ($i + 3 \rightarrow i$)²⁹ and π -helix ($i + 5 \rightarrow i$)³⁴ can be found in the -75° and -130° region, respectively. For a deep helix structure analysis, see Barlow *et al.*³⁵

Zsila *et al.* observed unstructured peptide LL-37 folding into an α -helix structure by non-covalent association of anti-inflammatory drugs, pigments, bile salts and food dyes. By hydrogen bonding and salt bridges, Lys and Arg amino acids were able to interact with a wide range of small molecules, resulting in multimeric complexes (Fig. 1).³⁶ Moreover, hemin and bile pigments were able to force the 26 amino acid IDP melittin, the major bee venom component, to fold into parallel β -sheet structures. In contrast, an α -helix promoting effect was observed with the also disordered but more cationic hybrid derivative 15 amino acid long CM15. The Trp and Phe residues induced π - π stacking interactions with the porphyrin dye.³⁷ León *et al.* studied the 11-mer repeat disordered unit P1LEA-22

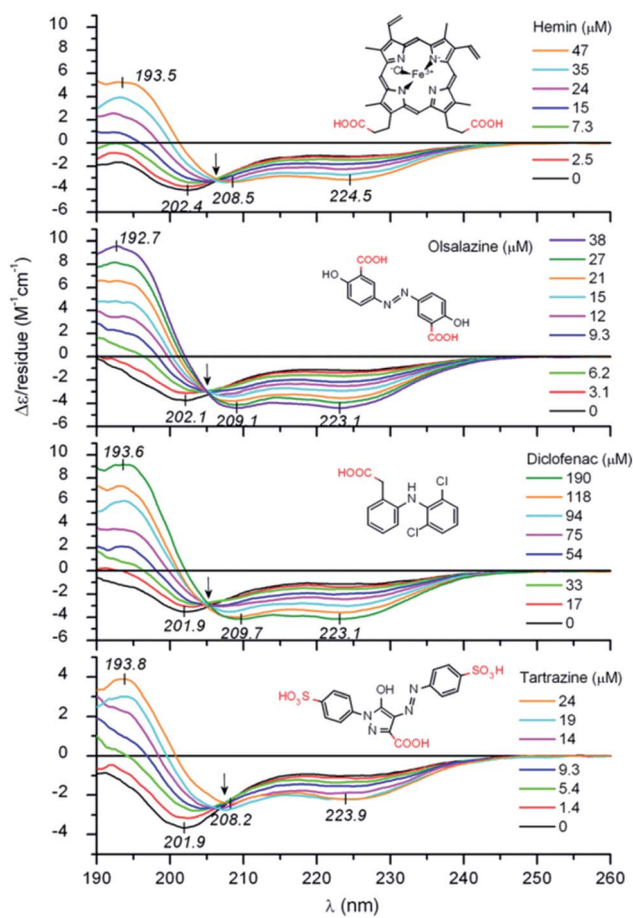


Fig. 1 UV-Vis IDP LL-37 disorder-to-helix transitions under the addition of various organic compounds. Reprinted with permission from Zsila *et al.*³⁶ Copyright 2019 Elsevier B.V.

behaviour at different temperatures and salt concentrations. Compared to similar 11-mer peptides, the presence of several Ala amino acids enabled the addition of FeCl_3 to enhance a PPII helical structure. The same behaviour was observed after choline chloride addition. A higher percentage of the PPII structure at a low temperature was found.³⁸ Fealey *et al.* investigated the structural dependence of synaptotagmin 1's IDR (Syn1) after dielectric constant and phosphorylation changes. A reduced dielectric constant promoted helix formation in neutrally charged core region residues. Lys–Asp acid salt bridges contributed to the stabilization of a transient secondary structure. However, phosphorylation in this region resulted in the formation of salt bridges, unsuitable for helix formation.³⁹ On a higher scale, Johnson *et al.* observed how IDP 4E-BP1 folded into an α -helix upon binding to its protein ligand, eIF4E. H-bond thiol–aromatic interaction between Phe₅₈ and Cys₆₂ at 4E-BP1 stabilized the helix.⁴⁰ Saglam *et al.* also observed the disorder-to-helix transition of the disordered peptide p53 in the presence of its protein receptor MDM2. A single α -helix was formed by induced fit if the unfolded state of the peptide was more stable than its folded state or at elevated MDM2 concentrations. The folding process was otherwise dominated by conformational selection.⁴¹ Jephthah *et al.* studied the N-



terminal MgtA IDR, referred to as KEIF. While disordered in aqueous solution, the helical content of this peptide increased if added to an organic solvent, similar to an aqueous solution containing anionic vesicles,⁴² leading to similar results to histatin 5. For this peptide, the presence of the His-Ser-His residue sequence was directly related to the α -helical structure formation.⁴³

Sheets

Peptides in a β -sheet conformation zig-zag in a more extended conformation with ϕ and ψ angles in the -140° and 130° range, compared to the -60° and -45° of α -helix ones. The β -sheet axial distance between adjacent residues is 3.5 Å, while in the α -helix it is 1.5 Å. In a β -sheet, two or more polypeptide chains run alongside each other and are linked in a regular manner by hydrogen bonds between the main chain C=O and N-H groups, while the side R chains point outward. The variations of the structure can be described depending on the strand orientation. All strands run towards the same direction in a parallel β -sheet conformation, while the strands are all alternated in an antiparallel β -sheet conformation. Mixed β -sheets show parallel and antiparallel strands. For extended structural characterization, we recommend the Salemmé review.⁴⁴

Coskuner *et al.* studied the relevance of the Tyr residue at the IDP A β 42. After Tyr₁₀-Ala mutations, the formation of β -sheet structures greatly diminished in the presence of adenosine triphosphate. They concluded that, after the Tyr₁₀-Ala mutation, a decreased in the reactivity of A β 42 toward various ligands and self-oligomerization in aqueous environments denoted the high structural control of the Tyr amino acid.⁴⁵ Takekiyo *et al.* studied the β -sheet folding of the A β fragment A β ₁₋₁₁ in the presence of the ionic liquid (IL) 1-butyl-3-methylimidazolium thiocyanate. IDR Asp and Glu residues interacted with the IL imidazolium region, which led to their oligomerization. Moreover, ILs with lower denaturing ability could not promote the aggregation.⁴⁶ Boopathi *et al.* showed how disordered A β 42 was affected by Zn²⁺ and Cu²⁺. Zn²⁺ had a higher hydrophobic behaviour compared to Cu²⁺, directly related to the fastest self-assembly of A β 42-Zn²⁺. Zn²⁺ increased the solvation free energy due to a higher tendency of forming the β -sheet structure at the Leu₁₇-Ala₄₂ residues.⁴⁷

Micelles and vesicles

Micelles and vesicles are colloidal dispersions formed by the supramolecular assembly of amphiphilic molecules in a liquid media. These colloids usually show a spherical shape, with different molecule organizations. In aqueous solution, micelles are surfactants that have their hydrophilic part in the outer region of the sphere-like structure, in direct contact with the solvent. The hydrophobic section is placed at the core, surrounded by the hydrophilic one. In the case of an organic solvent, the orientation of the molecules will shift, leading to a reverse micellar structure. Vesicles, on the other hand, are spherical capsules formed by one or more bilayers entrapping an aqueous medium while being surrounded by an aqueous solution. In comparison, the number of molecules required to

form a micelle and the size are mostly lower compared to a vesicle. Micelles are in the tens of nanometres diameter range, and vesicles are in the hundreds or even thousands. For an in depth understanding of the formation and design of both, we recommend the latest Lu *et al.*⁴⁸ and Has *et al.*⁴⁹ reviews, respectively.

Ivanović *et al.* analysed the micellar behaviour of *n*-dodecyl- β -D-maltoside (DDM). Due to this peptide intrinsically disorder behaviour, a moderate shape fluctuation was observed in its self-assembly, leading to final DDM ellipsoidal, oblate and prolate, conformations.⁵⁰ Accardo *et al.* synthesized two disordered peptide amphiphiles (PAs). PAs were characterized by two alkyl chains connected directly or by a linker to the R11 IDP, denoted as (C18)₂-R11 and (C18)₂-L1-R11, respectively. Presenting an ordered core and a 'disordered' surface, (C18)₂-L1-R11 self-assembled into micelles (~16 nm diameter) and small unilamellar vesicles (~200 nm diameter), while the (C18)₂-R11 PA was able to form only vesicles. With a mainly β -strand conformation of both PAs, the addition of the linker L1 gave a closer-to-liquid behaviour to the structure.⁵¹ Klass *et al.* imitated the process using diblock polymers that contained a hydrophobic and an intrinsically disordered hydrophilic domain (Fig. 2). These low polydispersity 27 nm diameter micelles were found to be formed across a broad range of pH (3.7–9.7), ionic strength (0–200 mM), and temperature conditions (25–70 °C). The authors concluded that at pH 7.9 or higher, significant heterogeneity and polydispersity in the micellar diameters were observed, due to the most collapsed state of the hydrophilic IDP portion. The micellar volume decreased reversibly with increasing temperature according to the interplay of intermolecular interactions of the hydrophobic tails and water with the hydrophilic headgroups. Finally, no obvious trends were observed after changing different salt concentrations.⁵² Acosta *et al.* proposed the use of antimicrobial peptides (AMPs) as self-assembling domains to drive hierarchical organization of intrinsically disordered protein polymers (IDPPs) based on an elastin-like recombinamer (ELR) (Val-Pro-Gly-Ser-Gly)₅₀-(Ile-Pro-Gly-Val-Gly)₆₀. At 5 °C, the ELR alone did not form any nanostructure. However, the AMP-ELR polymers formed nanofibers. Differences in the size and shape of the nanostructures as a function of the AMP sequence used were also observed. At a physiological temperature of 37 °C, both ELR and AMP-ELR self-assembled into micellar structures. In AMP-ELR samples, a small portion of nanofibers were present due to early AMP assemble processes. Moreover, after the incubation of the AMP-ELR samples, the presence of the AMP drove a second self-assembly in the form of aggregates with globular or amorphous shapes depending on the AMP structure.⁵³ Rao *et al.* used several low complexity IDPs (LC-IDPs) to form vesicles. LC-IDRs of SM50, LSM34, MSP130, and Priliskin-39, in the presence of Ca²⁺, self-associated by ionic interactions leading to 100–300 nm diameter vesicles. Moreover, THF was applied as an orthogonal solvent instead of the mineral precursor, forming the same structures.⁵⁴ Going one step further, Costa *et al.* designed an IDP based on hydrophilic Val-Pro-Gly-Val-Gly and hydrophobic Val-Pro-Gly-Ser-Gly motifs. Self-assembly into spherical micelles was triggered by



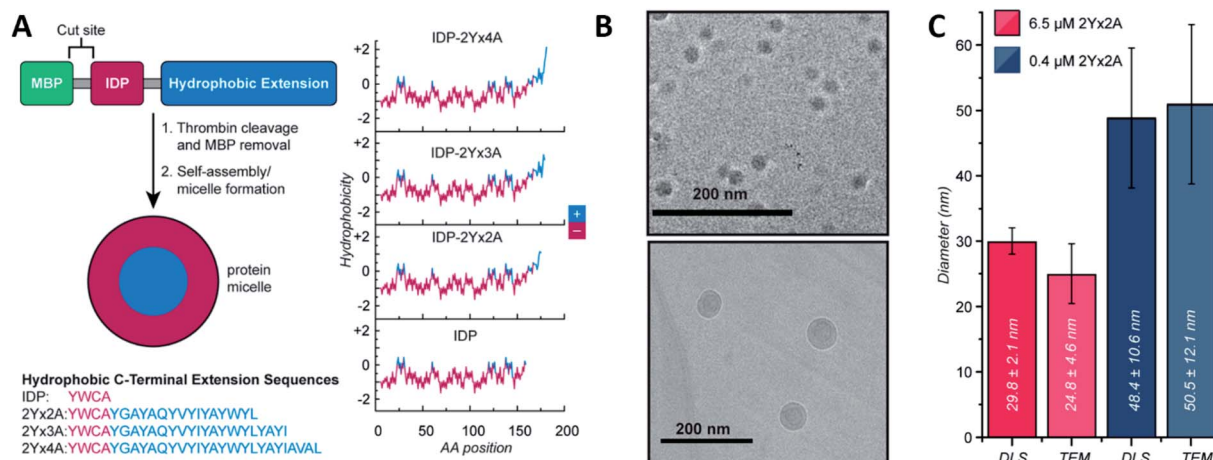


Fig. 2 IDP-based self-assembly behaviour. (A) Structure. IDP segment fused to different hydrophobic sequences and hydrophobicity plots of each final amphiphilic protein. (B) Cryo-TEM images of 6.5 μM (top) and 0.4 μM (bottom) IDP-2Yx2A micelles in PBS, pH 5.7. (C) comparison of DLS and cryo-TEM diameters obtained at different concentrations. Reprinted with permission from Klass *et al.*⁵² Copyright 2019 American Chemical Society.

increasing the temperature above its critical micelle temperature in bulk solution. This reversible process could be combined with a UV irradiation process, while peptides were in their micellar form for the formation of nanogels, after the addition of *para*-azidophenylalanine groups.⁵⁵

Fibrils

Fibrils are linear 10–100 nm diameter chains differentiated from filaments (their precursor) and fibers (their product). Fibrils are a well-known structure in the biological field, quite commonly found in the form of amyloid fibrils. The peptide is folded in a β -sheet parallel or antiparallel structure with an inter-strand distance of *ca.* 4.8 Å (N–H \cdots O=C hydrogen bonds between two consecutive peptide backbones). Different β -sheets stack in parallel. In function of the residues and packaging, its inter-sheet distance varies from 8.8 to 14.6 Å. This structure is elongated to protofilaments and subsequently twisted into multistrand helical mature amyloid fibrils. Depending on their torsion, fibrils can lead to crystal or nanotube structures. For a wider view of protein fibrils and amyloid fibrils specifically, we recommend the Fändrich review.⁵⁶

Humenik *et al.* observed the assembly of recombinant spider silk variants, denoted as eADF4(*Cn*), '*n*' being the number of C-modules. This C-module was a 35 amino acid segment rich in Gly and Pro residues and one poly-Ala stretch. While monomers showed an intrinsically disordered behaviour by themselves, they could self-assemble into cross β -sheet fibrils. For $C \geq 2$, the peptide folded towards antiparallel β -sheets followed by the formation of a nucleus *via* hydrophobic interactions of poly-Ala β -sheets. Finally, monomer addition occurred by the dock-lock mechanism forming the final fibril structure.⁵⁷ Hernik-Magon *et al.* studied the influence of the poly-L-glutamic acid (PLGA) length in the self-assembly process. Long chained (Glu)₂₀₀ molecules fibrillated more readily than short IDP (Glu)₅ fragments. While both started with an alpha structure, only β -(Glu)₂₀₀ amyloid tended to form well-defined twisted

superstructures. Moreover, their mixture accelerated the process. The intrinsically disordered pentapeptide, merged with structured (Glu)₂₀₀ chains, followed the PLGA's fibrillation pattern. At different mixture ratios, (Glu)₅ adopted a self-assembly β 2-fibril pattern normally accessible only to long-chained PLGA.⁵⁸ Similar results were reported by Zhang *et al.*, who theoretically explained how Glu/Asp-rich peptides aggregated in β -sheet structures and self-assembled into highly ordered amyloid fibrils.⁵⁹ Pan *et al.* showed the formation of amyloid-like fibrils from intrinsically disordered α -, β -, and k-caseins during heating (90 °C) at an acidic pH (2.0). The fibrillated caseins had increased contents of β -sheet organized structures with different nanomechanical properties and bulk viscosity.⁶⁰ Bakou *et al.* described the self-assembly of an intrinsically disordered polypeptide islet amyloid polypeptide (IAPP) fibril. Phe, Leu, and Ile were the residues directly related to the fibrillar structure formation.⁶¹ Larini *et al.* analysed a construct that included the PHF6* region of the neuronal-related IDP Tau. Specifically, Tau_{273–284} self-assembled into full-fledged fibrils.⁶² Adamcik *et al.* studied the third repeat fragment (R3) of this protein and obtained similar results. The 26-amino acid Tau-derived peptide could self-assemble into amyloid fibrils with a β -sheet-based structure without any external induction. Complete ordered 2D laminated flat ribbons with on average 18.7 protofilaments were observed with a lateral size of 149.7 nm and 3.8 nm thickness (Fig. 3). Moreover, ribbons of >350 nm lateral size and 45 protofilaments could be observed, the biggest reported to date.⁶³ The self-assembly of the complete microtubule-associated protein Tau into neurotoxic oligomers, fibrils, and paired helical filaments took place after the addition of polyanions, such as heparin, as described by Despres *et al.*⁶⁴ Dec *et al.* analysed the H-fragment, disordered in aqueous solutions, of predominantly ordered α -helical insulin. Thin and structurally homogenous fibrils with a typical parallel β -sheet conformation appeared upon lowering of the pH value. It was concluded that, due to its acidity, the Ala-rich



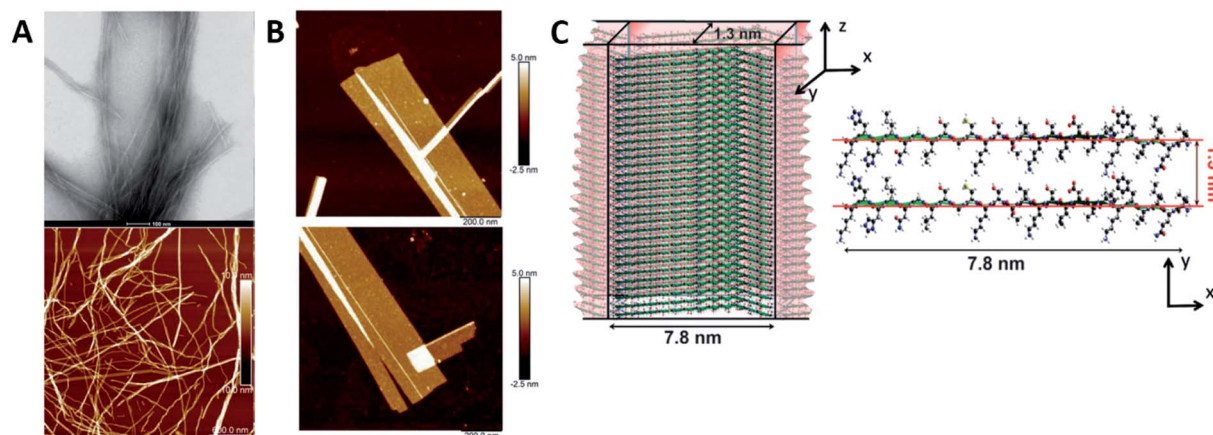


Fig. 3 R3 peptide (A) TEM (up) and AFM (low) images of self-assembled fibril structures in the presence of heparin. (B) High-resolution AFM images of flat multistranded ribbons in the absence of heparin. (C) Structural illustration of protofilaments. The distance between β -sheets is 1.3 nm with an off-set of ca. 0.4–0.6 nm corresponding to the peptide residues on both sides of the β -sheet. Reprinted with permission from Adamcik *et al.*⁶³ Copyright 2016 Wiley-VCH Verlag GmbH & Co.

chain portion played a crucial role in the aggregation of the whole H-fragment.⁶⁵ Kuhn *et al.* observed the p3 peptide behaviour and the IDP formed through alternative processing of Amyloid- β (A β).⁶⁶ The self-assembly of this peptide proved to form oligomers and fibrils at higher aggregation rates than A β . In addition, p3 fibrils exhibited cross- β -sheet amyloid structures. A final hydrophobic steric zipper was a convincing organization given that the two amyloidogenic hydrophobic patches of A β are also found in p3.⁶⁷ Focused on the A β IDP, Jana *et al.* showed how the addition of glycosylated Lys residues forced A β to self-assemble at early stages into protofibrillar conformations after folding of β -sheets. New and stronger inter-monomer salt bridging bindings took place in the glycosylated form with dispersion interactions playing no significant roles.⁶⁸

Other structures

Although most natural disordered peptides and proteins may have α -helix and β -sheet structures after folding and fibrils after self-assembling, some exceptions could be found. Mostly artificially synthesized, several IDPs/IDRs can be organized from more classical to complex structures, from rods to fractals, respectively.

Khatun *et al.* analysed 37-residue IDP amylin. While several typical structures were already reported for this peptide, such as fibrils⁶⁹ or micelles,⁷⁰ they discovered that also fractal self-assembly processes could occur (Fig. 4). Affected by the solvent and the media, results indicated the main role of the hydrophobic interactions in the fractal self-assembly and aggregation of amylin. Relevant interactions between the anisotropically distributed hydrophobic residues and polar/ionic residues on the solvent-accessible surface of the protein drove the process.⁷¹ Quiroz *et al.* synthesized two IDPPs that exhibited lower and/or upper critical solution temperature phase behaviour. The IDPPs were composed of the same corona-forming ELP block and different hydrophobic IDPP core-forming blocks with distinct hysteretic phase behaviours. While the ELP formed 20 to 30 nm diameter micelles, IDPPs

self-assembled into nanoparticles with a rod-like morphology, all thermodynamically controlled.⁷² Stehli *et al.* reported the formation of highly ordered spherulite structures after the self-

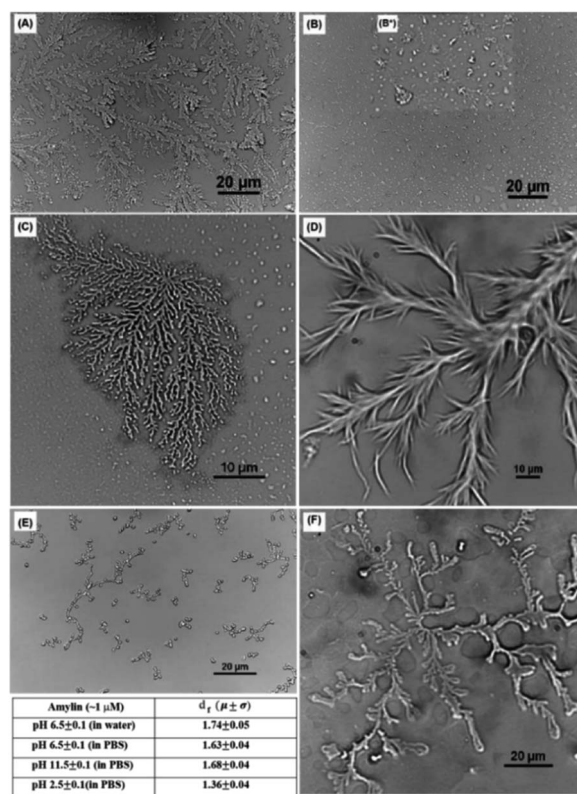


Fig. 4 Optical microscopic images of an amylin fractal observed in PBS buffer at pH 6.5 \pm 0.1 (A) at \sim 10 μM concentration, (B) at \sim 0.1 μM concentration, and (B*) inset showing the presence of different morphologies (C) at \sim 1 μM concentration; in PBS buffer at \sim 1 μM concentration (D) at pH 11.5 \pm 0.1 and (E) at pH 2.5 \pm 0.1; and (F) amylin fractal observed in DI water at \sim 1 μM concentration at pH 6.5 \pm 0.1. The table in the figure contains the d_f for the morphologies obtained with an optical microscope shown in (c)–(f). Reprinted with permission from Khatun *et al.*⁷¹ Copyright 2020 The Royal Society of Chemistry.



assembly of intrinsically disordered PLGA. PLGA monomers at early stages could exist in either a collapsed globular state or an extended random coil conformation. An α -helical conformation promoted the spherulite formation, while a random coil promoted the formation of an amyloid fibril.⁷³ Bishof *et al.* stated how the disordered low-complexity domain 1 (LC1) of the U1 small nuclear ribonucleoprotein (U1-70K) self-assembled into oligomers. The LC1 domain contains highly repetitive basic (Arg/Lys) and acidic (Asp/Glu) residues and behaves like a Gln/Asn-rich LC domain. These domains could form “glue” that drove the granule assembly processes.⁷⁴ Dooley *et al.* designed an IDP based on a repeats-in-toxin (RTX) domain that allowed it to gain a β -roll secondary structure in a calcium rich environment. Based on the adenylate cyclase region obtained from *Bordetella pertussis* bacteria, the repeating polypeptide consisted of two parallel β -sheet faces separated by flexible turn regions. Aspartic acid residues were included in the turn regions to promote the coordination of the calcium ions by the carboxylic groups. Moreover, after engineering, the addition of L-Leu and D-Leu residues to the structure made this IDP self-assemble into hydrogels with minimal impact on its calcium affinity.⁷⁵

Experimental techniques

While having biological activity, IDPs/IDRs lack a well-defined structure. A classical approach based on basic structural experimental techniques makes IDPs/IDRs impossible to characterize due to their highly heterogeneous conformational behaviour. Over the last few years, a large variety of physical techniques have been optimized and applied successfully to unveil the fundamental rules that make proteins fold into different structures and elucidate the conformational transition from their native disordered to a more structured state. Even more, the understanding of these transitions from a disordered to different secondary or tertiary structures can lead to remarkable changes in their implementation and future applicability. However, the mechanisms of these structure transitions are not fully understood, and a fundamental description of the kinetic and thermodynamic variables will undoubtedly help to recognize these transition processes and their relevance. The effect of the AA sequence, the length of the AA chain, and the biological media are the main parameters to be studied. Furthermore, the existence of domains showing this behaviour or IDRs is an intriguing question in the field.

In this section, we provide examples of key types of systems and insights that can be addressed using the most relevant and widely used methods, summarized in Table 2.

Nuclear magnetic resonance (NMR)

NMR spectroscopy is undoubtedly one of the most preeminent methods for determining the molecular behaviour of IDPs/IDRs at the atomic resolution. The use of NMR for protein structural characterization was developed in the 1970s and 80s by scientists such as Ernst,⁷⁶ Wüthrich,⁷⁷ Clore⁷⁸ or Gronenborn.⁷⁹ Nowadays, after its optimization, NMR is applied to the

biochemical field, specifically in the binding and self-assembly of peptides in aqueous media. In NMR spectroscopy, radio-frequency (RF) waves in the MHz range are applied to samples subjected to a strong and homogeneous magnetic field (B_0). Their active nucleus (such as ^1H , ^{13}C or ^{15}N) absorbs the electromagnetic radiation at a specific frequency characteristic of the isotope. To know more about the analysis of NMR spectra applied to peptides, how to assign the chemical shifts and how structural changes are reflected in the NMR signal, we recommend the Brutscher *et al.* chapter.⁸⁰ Briefly, the Hamiltonian $\langle H \rangle$ describes the energy of interaction of the nuclear spins with internal and external electric and magnetic fields. In isotropic systems, like IDPs, the fast-molecular reorientation will average out any orientation dependence of the interactions, and only the isotropic parts of this Hamiltonian solution will remain. Therefore, despite the complexity of these molecules, a well resolved spectra can be obtained. It can be easily explained by the absence of structure. Similar shifts will be obtained due to protons in the same chemical group having the same chemical environment.

Chaves-Arquero *et al.*, using $^1\text{H}_\alpha$ and $^{13}\text{C}_\alpha$ conformational shifts in solution, could observe the phosphorylation effect on the structural behaviour of two derived IDPs.⁸¹ While both had a random coil structure in water ($|\Delta\delta\text{H}_\alpha| \leq 0.05$ ppm and $|\Delta\delta\text{C}_\alpha| \leq 0.4$ ppm, being $\Delta\delta\text{H}_\alpha = \delta\text{H}_{\alpha,\text{observed}} - \delta\text{H}_{\alpha,\text{random coil}}$, ppm; and $\Delta\delta\text{C}_\alpha = \delta\text{C}_{\alpha,\text{observed}} - \delta\text{C}_{\alpha,\text{random coil}}$, ppm), the IDP T¹¹⁸-H1.0 and its phosphorylated derivative, pT¹¹⁸-H1.0, consisted of two helical regions in the presence of trifluoroethanol (TFE). Clear changes in $|\Delta\delta\text{H}_\alpha|$ as well as $|\Delta\delta\text{C}_\alpha|$ denoted it (T¹¹⁸-H1.0: $|\Delta\delta\text{H}_\alpha| = -0.29$ ppm and $|\Delta\delta\text{C}_\alpha| = 3.35$ ppm; pT¹¹⁸-H1.0: $|\Delta\delta\text{H}_\alpha| = 0.27$ ppm and $|\Delta\delta\text{C}_\alpha| = 2.92$ ppm, (Table 1)). Even more, the helix populations could be estimated from $\Delta\delta\text{H}_\alpha$ and $\Delta\delta\text{C}_\alpha$ averaged for the helical residues as well as the orientation between the helices. Both systems contained a >80% helical structure, with a perpendicular arrangement exclusively for the non-phosphorylated IDP. This structural difference is directly related to the different biological roles. Kosol *et al.* elucidated the structural basis of the Bamb_5917 protein. They could observe low heteronuclear Nuclear Overhauser Effects (NOEs) in the Bamb_5917 PCP domain, typical behaviour of an IDR with high picosecond-nanosecond flexibility ($\{^1\text{H}\}$ - ^{15}N NOE close to 0).⁸² Based on 2D CON^{Pro} experiments, Murrall *et al.* showed how to obtain the fingerprint of the Pro residue in IDPs,⁸³ a largely exploited AA used to prevent the formation of stable secondary structures. The lack of the peptide HN atom implied that this amino acid was not directly detectable in the commonly used 2D ^1H , ^{15}N NMR spectroscopy. 2D CON was the chosen experimental technique to determine the correlations between the backbone carbonyl carbon and nitrogen of neighbouring residues. Fast NMR assignments of IDPs can be done by combining 2D hNCA and 2D hNcoCA spectra as Sukumaran *et al.* presented for the human α -synuclein protein during its self-aggregation process.⁸⁴ By solid-state NMR measurements, Reichheld *et al.* were able to observe the conformational transition of elastin cross-linking domains during their self-assembly. The C_α and C_β chemical shift values of Ala and Lys AAs were used due to their particular sensitivity to backbone



Table 1 $[\theta]^{222\text{nm}}$, averaged $\Delta\delta$ values and α -helix populations estimated from $[\theta]^{222\text{nm}}$, and $\Delta\delta_{\text{H}\alpha}$ and $\Delta\delta_{\text{C}\alpha}$ for peptides T¹¹⁸-H1.0, pT¹¹⁸-H1.0, T¹⁴⁰-H1.0 and pT¹⁴⁰-H1.0 in aqueous solution and in 90% TFE at pH 5.5 and 25 °C. Reprinted with permission from Chaves-Arquero *et al.*⁸¹ Copyright 2020 Wiley-VCH Verlag GmbH & Co

Peptide	Conditions	$[\theta]^{222\text{nm}}$ (deg cm ⁻² dmol ⁻¹)	% helix ^a from $[\theta]^{222\text{nm}}$	Helix length ^a	$\Delta\delta_{\text{H}\alpha}$ [ppm]	% α helix from $\Delta\delta_{\text{H}\alpha}$	$\Delta\delta_{\text{C}\alpha}$ [ppm]	% α helix from $\Delta\delta_{\text{C}\alpha}$	Avgd% α helix ^{a,c}
T ¹¹⁸ -H1.0	H ₂ O	-69.9	8	105-115	-0.05 ^b	13 ^b	+0.12 ^b	4 ^b	9 ± 5 ^b
	90% TFE	-11 229.6	37		-0.29	75	+3.35	100	87 ± 13
pT ¹¹⁸ -H1.0	H ₂ O	86.4	7	105-115	-0.05 ^b	13 ^b	+0.15 ^b	5 ^b	9 ± 4 ^b
	90% TFE	-10 988.9	36		0.27	66	+2.92	95	81 ± 14
T ¹⁴⁰ -H1.0	H ₂ O	-915.8	10	141-147	-0.06 ^b	16 ^b	+0.21 ^b	7 ^b	12 ± 5 ^b
	90% TFE	-7137.4	26		-0.15	39	+1.58	51	45 ± 6
pT ¹⁴⁰ -H1.0	H ₂ O	1175.9	5	141-147	-0.06 ^b	16 ^b	+0.28 ^b	9 ^b	13 ± 4 ^b
	90% TFE	-8168.4	29		-0.16	40	+1.80	58	49 ± 9

^a Note that the CD-estimated helix percentages correspond to an average for all the peptide residues, whereas the NMR-estimated helix percentages relate to the residues within the helix. ^b Values measured at 5 °C. ^c Reported errors are standard deviations for the mean of the percentages obtained from the $\Delta\delta_{\text{H}\alpha}$ and $\Delta\delta_{\text{C}\alpha}$ values.

torsion angles. Lyophilized EP20-24 IDP Ala residues showed C_α-C_β cross-peaks with chemical shift values indicative of a larger α -helix population combined with a smaller random coiled one. On the other hand, the hydrated EP20-24 coacervate displayed a conspicuous C_α-C_β cross-peak with shifts indicative of a β -strand backbone conformation. If cross-linked with genipin, the polypeptide showed a very prominent cross-peak with chemical shift values characteristic exclusively of Ala α -helical conformations.⁸⁵ Using ¹H-¹⁵N HSQC and 3D HNCACB, Garry *et al.* studied the self-assembly of the Leucine-Rich Amelogenin Protein (LRAP) into a unique quaternary structure referred to as a 'nanosphere' in the presence of NaCl. They identified the specific residues involved in the early stages of the nanosphere assembly in this IDP by following its amide chemical shift perturbations as a function of salt concentration. The disappearance of amide cross peaks in the ¹H-¹⁵N HSQC spectrum at high NaCl concentrations likely reflected a restricted motion at the protein-protein interface.⁸⁶ With

these techniques, Beck Erlach *et al.* also studied the pressure and temperature effects on the self-assembly of intrinsically disordered human IAPP (hIAPP) and Alzheimer peptide A β ₁₋₄₀. The hIAPP N-terminal region displayed large differences in pressure sensitivity compared to A β , pinpointing to a different structural ensemble in this sequence element. A helical origin was related to hIAPP and an amyloid deposit to A β ₁₋₄₀.⁸⁷ Based on 2D [¹H ¹H] NOESY and TOCSY, Accardo *et al.* characterized the self-assembly and final organization of different IDP-amphiphilic molecules. In the case of (C18)₂-L1-R11, the spectra revealed NOEs between the CH₂ groups from the C18 alkyl chains and peptidic protons, which resonated at 3.55, 3.51, 2.55 and 2.42 ppm.⁵¹ Data indicated that the peptidic region did indeed interact with the C18 chains, endowing (C18)₂-L1-R11 with a certain degree of flexibility in solution. In the case of the (C18)₂-R11 peptide, NOEs between the CH₂ protons related to the C18 alkyl chains and either peptidic HN or aromatic protons appeared rather clear with a line broadening effect. This

Table 2 Most relevant IDP/IDR characterization methods

Technique	Structural observation	References
Nuclear magnetic resonance (NMR)	¹ H _α and ¹³ C _α signal shifts ($\Delta\delta_{\text{H}\alpha}$ and $\Delta\delta_{\text{C}\alpha}$)	53, 66, 72, 73, 83-90, 94, 99, 109, 115, 116 and 165
Circular dichroism (CD)	Maximums and minimums in the 190-250 nm CD region	38-42, 44, 45, 48, 53, 55, 59, 60, 62, 65, 67, 70, 71, 75, 77, 83, 87, 93, 94, 97, 99, 102, 105, 109, 112, 117, 119 and 120
Electron paramagnetic resonance (EPR)	Substituted Cys or coordinated Cu ²⁺ tracking	96-99
Fluorescence spectroscopy	Tryptophan (Trp, 300-450 nm), tyrosine (Tyr, 250-370 nm) and phenylalanine (Phe, 250-350 nm) shifts	39, 42, 53, 62, 64, 66, 69, 70, 71, 73, 75, 77, 87, 94, 101 and 102
Raman spectroscopy	Amide I (1630-1700 cm ⁻¹), amide III (1230-1310 cm ⁻¹) and backbone skeletal stretch (870-1150 cm ⁻¹) regions	104-107
Fourier transform infrared spectroscopy (FT-IR)	Amide I (1700-1600 cm ⁻¹) and amide II (1600-1500 cm ⁻¹) regions	48, 59, 60, 67, 75, 93, 106 and 109-112
Small-angle X-ray scattering (SAXS)	Form factor, Kratky plot and pair distance-distribution function (PDDF) shape	44, 45, 52, 65, 111, 115, 116, 117 and 120
Static and dynamic light scattering (SLS & DLS)	Gyration (R_g) and hydrodynamic radius (R_h)	44, 53, 54, 55, 56, 57, 73, 75, 88, 99, 119 and 120



confirmed the high tendency of this PA to self-associate in larger aggregates than (C18)₂-L1-R11 due to the lack of the connector L1. Hou *et al.* could characterize the self-assembly mechanism of A β IDPs in a global β structure and how the oxidation of some regions prevented this structure from being promoted to a random coil instead. The relevance of His residues in the self-assembly process was studied by oxidation processes and pH variations, key for a final β -structure.^{68,88}

Circular dichroism (CD)

IDPs, due to the lack of any significant organized secondary structure, show the characteristic CD spectrum of an unordered polypeptide, with a strong negative band near 200 nm and either a weak negative shoulder or a weak positive maximum near 220 nm. IDRs are also recognized by CD, even though they show ordered and unordered regions, making the IDR difficult to diagnose. For these regions, a limited proteolysis approach followed by CD is commonly used.⁸⁹ As IDPs/IDRs present dynamic conformations, measurements should be carried out not at a single and unique set-up but instead changing the chemical conditions and observing the peptide behaviour, as part of the determination of the intrinsically disordered nature. While we recommend Chemes *et al.* work for a complete understanding of the methodology to achieve the maximum the CD can contribute,⁹⁰ we will try to show the relevance of this powerful tool. This essential technique is often used as a preliminary to more complex methodologies such as NMR, for a deep secondary structure characterization. Without going further, some studies cited in the NMR section also carried out CD measurements as a starting point.

Chaves-Arquero *et al.* work is an example. Four IDPs derived from the C-terminal domain of Histone H1.0 showed a typical random coil strong minimum at *ca.* 195 nm and no other secondary structure features, confirming that they were predominantly disordered in water (Table 1 and Fig. 5). The addition of TFE caused an increase of structural organization on all the peptides with a progressive conversion of the 195 nm region into a maximum at *ca.* 197 nm, related to helix populations.⁸¹ Reichheld *et al.* used mainly CD to support the structural conformations observed in elastin. IDP EP20-24 and tropoelastin (its monomer), at 5 °C, had a similar behaviour with a strong minimum at 202 nm and another at 222 nm, characterized as the disordered and α -helical signals, respectively. At 37 °C or higher temperature values under physiological conditions, the absence of the 222 nm band denoted the significant loss of the α -helical structure. EP20-24 showed a totally different structure at temperatures above 40 °C after the mutation of the Lys monomers to Ala. The 202 nm signal was lost above 40 °C and a shift to a strong minimum at 217 nm was indicative of β -strand formation. Furthermore, a Lys-to-Tyr mutation made the IDP show at 5 °C a strong minimum at 217 nm with a very weak minimum at 202 nm, demonstrating that this mutant polypeptide was predominantly a β -strand even at low temperatures. Increasing the temperature to 37 °C resulted in the loss of the minimum at 202 nm, indicating that EP:Lys-to-Tyr had a greater propensity for β -strand formation

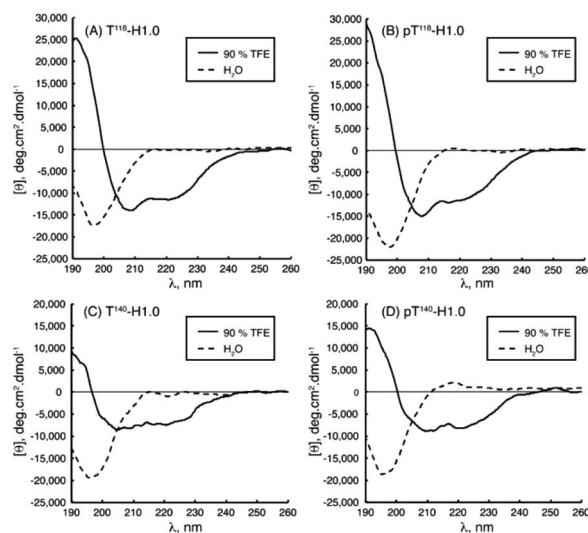


Fig. 5 CD spectra of peptides (A) T¹¹⁸-H1.0, (B) pT¹¹⁸-H1.0, (C) T¹⁴⁰-H1.0 and (D) pT¹⁴⁰-H1.0 in aqueous solution (dotted line) and in 90% TFE (black line) at pH 5.5 and 25 °C. Reprinted with permission from Chaves-Arquero *et al.*⁸¹ Copyright 2020 Wiley-VCH Verlag GmbH & Co.

than the EP:Lys-to-Ala mutant.⁸⁵ Sun *et al.*, while analysing the amyloid self-assembly fibrillation process of hIAPP_{8–20}, revealed a negative peak at 200 nm at 0 h and a negative peak at 218 nm at 48 h, denoting a structural transition from random coil to β -strands.⁹¹ Rivera-Najera *et al.* tested the disordered self-association behaviour predicted for PvLEA6 protein in its secondary structure. Temperature changes could also promote modifications in the protein conformation, as demonstrated by CD in far-UV light. A negative band at 197–200 nm related to a random coil unordered structure described this IDP at low temperatures, which was altered to a β -like one for higher temperatures, denoted by an increase in the CD negative signal between 215 and 220 nm.⁹² Dooley *et al.* used an intrinsically disordered peptide isolated from the repeats-in-toxin (RTX) domain to show how the addition of Ca²⁺ folded the system into a β -roll secondary structure based on two parallel β -sheet faces. In calcium-free environments, the spectra exhibited large negative peaks at 198 nm, indicative of a randomly coiled peptide. A random coil to β -sheet transition was evidenced by the emergence of a negative peak at 218 nm after Ca²⁺ titration, indicative of this disordered-to-folded transition.⁷⁵ Bakou *et al.* observed the key effects of Ala mutations on the conformation of IAPP. Mixtures of random coil and β -sheet/ β -turn structural elements were observed for freshly dissolved IAPP and mutant samples. CD spectral deconvolutions suggested IAPP 30–40% β -turn/ β -sheet, 50–60% random coil and 10% α -helix contents. In the case of most mutants, 30–50% β -turn/ β -sheet, 40–60% random coil and 5–15% α -helical distributions were determined. Together, CD studies provided evidence that (a) mutations are not related to changes in the structure and (b) β -strand-loop- β -strand conformers can be observed in major populations for IAPP compared to its mutants.⁶¹



Electron paramagnetic resonance spectroscopy (EPR)

EPR spectroscopy specifically detects unpaired electrons. The introduction of a site-directed spin labelling (SDSL) model at the protein of study is the common procedure when applied in the biological field. It is usually accomplished by Cys substitution mutagenesis followed by covalent modification of the unique sulphhydryl group with a selective nitroxide reagent. For an in-depth analysis of all EPR variety approaches depending on the system, see Weickert *et al.*⁹³

Pirman *et al.* characterized the disordered to α -helical transition of IA3 upon TFE addition. In this case, they modified the Cys AA side chain with methanethiosulfonate (MTSL), 4-maleimido-TEMPO (MSL) and 3-(2-iodoacetamido)-proxyl (IAP). They compared the peak-to-peak intensities of the low-field, $h_{(+1)}$, center-field, $h_{(0)}$, and high-field, $h_{(-1)}$, resonances. It was possible to conclude that (a) the line shapes obtained showed the following expected mobility trend: IAP > MTSL > MSL and (b) the addition of TFE produced a conformational change. The first conclusion was reached because the overall intensity of the signal is proportional to the mobility. The spin probe increases the motional average with the increase of intensity. The overall intensity of all the spectra decreased and was broader after the TFE addition. This decrease was directly related to the lower overall mobility of the spin label, arising from the conformational changes of the protein backbone.⁹⁴ Bund *et al.* used this technique to observe how copper induced a self-assembly process in the 18.5 kDa intrinsically disordered Myelin Basic Protein (MBP). Because Cu^{2+} is an EPR-active ion (with $S = 1/2$, $^{63/65}\text{Cu} : I = 3/2$), direct investigations of the interaction between MBP and Cu^{2+} are feasible. Comparing the continuous wave EPR spectra of Cu^{2+} /MBP with and without PBS, clear shifts of the peak positions as well as significant changes in the overall signal width were denoted. Multiple nitrogen coordinations of Cu^{2+} were indicated by the copper g- and hyperfine coupling values obtained in phosphate buffer. These results were in line with the values of Cu^{2+} coordinated to nitrogen atoms of imidazole rings of several His AAs and significantly differ from values for non-coordinated Cu^{2+} . To summarize, at a MBP : Cu^{2+} ratio > 1 : 2, a coordination process occurred with significant aggregation of MBP into larger particles of 100–

200 nm diameter in a PBS media.⁹⁵ Kaminker *et al.* directly probed the backbone changes between IDPs that allowed the control over their preferred conformation (Fig. 6). For oligomer structures, the use of peptide-linkers had a high tendency to form *trans* configurations around the amide bonds due to their more extended conformations (distance distribution of 8.3 to 20.5 Å). In contrast, the peptoid octamer adopted a more compact conformation, likely due to a relative increase in *cis* configurations around the amide bonds (11 to 23 Å). The DS-peptoid octamer showed an intermediate behaviour, as both *cis/trans* options were viable given its alternating sequence (9 to 22 Å).⁹⁶ Chinak *et al.* tracked the structural and aggregation features of the Human k-Casein fragment called lactaptin. The pH variation from 3.9 to 7.5 led to significant changes in the EPR spectrum of the main fraction and the appearance of a very wide line with a very short electron spin relaxation time. These results were in good agreement with the fact that most of the proteins form aggregates under physiological pH conditions, which greatly broadened the EPR spectra because (a) larger-size aggregates correspond to slow rotation and long correlation times and (b) modulation of the dipole-dipole interaction and the exchange interaction between spin labels in aggregates lead to short electron spin relaxation times.⁹⁷

Fluorescence spectroscopy

IDPs can be characterized by tracking the Trp residues using fluorescence spectroscopy. Trp is accessible to external fluorescence quenchers and has a redshifted fluorescence spectrum with a maximum at 340–353 nm. Moreover, interactions with other residues lead to a more rigid or more hydrophobic system, resulting in a displacement of the fluorescence maximum position to the blue region. This spectral effect can be used to evaluate not only self-assembly but also interaction processes. For a more comprehensive analysis, see Permyakov & Uversky.⁹⁸

Bakou *et al.* used fluorescence to analyse the binding between IDPs A β 40, IAPP and several alanine-altered IAPPs. Using N-terminal fluorescein-labelled IAPPs (Fluos-IAPP), the alanine-altered IAPP affinity with IAPP and A β 40 was quantified by fluorescence spectroscopic titrations. The results denoted that Fluos-IAPP, in the presence of A β 40 or IAPP, led to a significant enhancement of its fluorescence emission, while the alanine-altered IAPP showed no fluorescence changes after the addition of A β 40 or IAPP at the same low concentrations. The interactions were weaker when the number of Ala groups was increased, directly related to a decrease in the affinity. By mutating different AAs and looking at its effect on the fluorescence signal, they could conclude which residues were the ones taking part in the IAPP–IAPP self-assembly and IAPP–A β 40 binding.⁶¹ Rivera-Najera *et al.* also used fluorescence spectroscopy to track the local environment around the aromatic residues of the PvLEA6 IDP. They observed a decrease in the fluorescence intensity of the Tyr residue, indicative of a structure reorganization, going from a mostly rigid and hydrophobic environment to a more relaxed structure with possible tertiary interactions. This explained a transition from random coil and PII-like extended helices to β -like structures.⁹² Acharya *et al.*

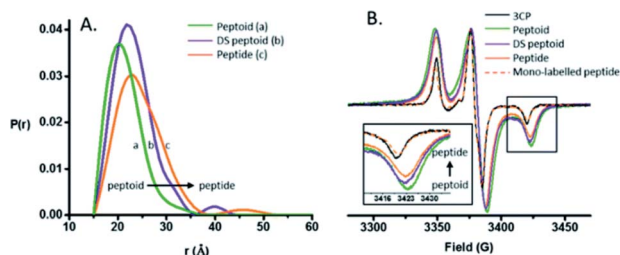


Fig. 6 (A) DEER distance distributions obtained for the peptide, peptoid, and DS-peptoid octamers. (B) Low temperature CW EPR of the trimer series overlaid with the mono-labeled peptide and 3CP free radical. Insets: zoom in of the high-field region. Reprinted with permission from Kaminker *et al.*⁹⁶ Copyright 2018 The Royal Society of Chemistry.



modified the α -synuclein IDP at positions 94 and 69 to promote Trp-Cys quenching. The binding was monitored using the fluorescence properties of the molecular tweezers CLR01. The incubation of α -synuclein alone over 6 h denoted a red shift in its fluorescence spectra, meaning that Trp was more solvent-exposed in oligomers than in the monomer form. Also, the increase in the sample turbidity denoted the formation of aggregates. When incubated with CLR01, increased values in the fluorescence of Trp₉₄ were observed, related to a high-affinity binding between them. No red shift was found, denoting CLR01 changed the oligomer structure such that the solvent exposure of Trp did not increase during oligomerization.⁹⁹ Zsila *et al.* studied Melittin binding with several bile pigments, in this case using the existence of a Trp residue in the molecule. At low pigment concentrations, the emission declined more sharply than at higher loadings, suggesting the formation of an initial complex serving as a scaffold for the binding of additional molecules and denoting the Trp stabilizing behaviour.¹⁰⁰

Raman spectroscopy

Conventional polarized visible Raman is generally considered a low-resolution technique, which can solely be used to discriminate between helices, β -sheets, turns, and random coils. Raman spectroscopy mostly exploits the structural sensitivity of the amide I mode to obtain dihedral angles adopted by folded and unfolded peptides. The amide I band region is located in the Raman shift range of *ca.* 1500 to 1750 cm^{-1} . Specifically, the $\sim 1650 \text{ cm}^{-1}$ region is typical of an α -helix, $\sim 1670 \text{ cm}^{-1}$ for β -sheet and $\sim 1680 \text{ cm}^{-1}$ for random coil structures. Recent advances in the field have substantially improved its usability. Chiral techniques like Raman optical activity (ROA) or UV resonance Raman spectroscopy (UVR) have been developed to probe the ordered and unordered structures of peptides and proteins. Insights in the study of IDPs by Raman spectroscopy can be found in the Zhu *et al.* review.¹⁰¹

Signorelli *et al.* used Raman spectroscopy to carry out the structural characterization of the p53 IDP with potential therapeutic applications. A careful analysis of the amide I Raman band revealed the presence of extended random coils and predominant β -sheet regions in its DNA binding domain (DBD). A wide curve at 1675 cm^{-1} for p53 and a maximum peak at 1669 cm^{-1} for the DBD were observed by peak deconvolution, explaining their predominant random coil and β -sheet organization. Even more, they observed how the addition of MeOH to the PBS aqueous media affected the amide I structure, with both p53 and DBD peptides showing a major contribution of the β -sheet structure. In contrast, if TFE was added, a final α -helix could be observed.¹⁰² McCaslin *et al.* used UVR to obtain new information concerning the structure and function of histatin 5 and its interactions with Zn^{2+} and Cu^{2+} (Fig. 7). Bands at 1315 cm^{-1} , 1334 cm^{-1} , 1371 cm^{-1} , and 1565 cm^{-1} were assigned to the His side chain and the bands at 1176 cm^{-1} , 1210 cm^{-1} , and 1612 cm^{-1} were related to Tyr due to the phenol/phenolate and imidazole side chain contributions. While adding Cu^{2+} did not change the histatin 5 self-assembled

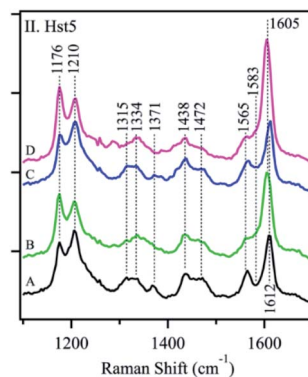


Fig. 7 UVR spectra of Hst-5 in the absence of added metals (A, black) and after the addition of Zn^{2+} (B, green), Cu^{2+} (C, blue), or a mixture of Zn^{2+} and Cu^{2+} (D, pink). Reprinted with permission from McCaslin *et al.*¹⁰³ Copyright 2019 Springer Nature.

structure, Zn^{2+} binding altered it. The 1315 cm^{-1} band underwent an upshift to 1334 cm^{-1} , and the 1371 cm^{-1} band was decreased in intensity. In addition, the 1565 cm^{-1} band was reduced in intensity, and a zinc-induced shoulder was present at 1583 cm^{-1} . Still, it was stated that more analysis were needed to finally elucidate the complete 3D structure of the IDP.¹⁰³ Rawat *et al.* observed possible different IAPP structures after its aggregation. The Raman spectra of oligomers suggested the presence of mostly an α -helix due to a clear peak at 1656 cm^{-1} and a weak peak at 1261 cm^{-1} , assignable to amide-I and amide-III, respectively. In the fibril state, there were strong peaks at 1668 cm^{-1} (in the amide-I region) and 1235 cm^{-1} (in the amide-III region), which confirmed their well-known β -sheet conformation. These results led to the understanding of why oligomers interact more with membranes, which prefer an α -helix structure.¹⁰⁴ Again, for IAPPs, La Rosa *et al.* used Surface Enhanced Raman Scattering (SERS) to validate their hIAPP self-aggregation simulations after the addition of silver nanoparticles to the system. The secondary structure of the amyloidogenic proteins was revealed. It showed how proteins self-aggregate from monomers to oligomers, and eventually into proto-fibrils and fibrils. Three signals were used for the identification of different protein backbone confirmations: amide I (stretching vibration of C=O ranging from 1600 to 1690 cm^{-1}), amide II (1480–1580 cm^{-1}) and amide III (1230–1300 cm^{-1}) both associated with the coupled C–N stretching and N–H bending vibrations of the peptide group.¹⁰⁵

Fourier transform infrared spectroscopy (FT-IR)

Infrared spectroscopy is a reliable tool to obtain information on the protein secondary structure and aggregation. Similarly to Raman, it mostly requires to identify and analyse the protein absorption components in the amide I (C=O stretching vibrations, 1700–1600 cm^{-1}) and amide II (C–N stretching vibrations in combination with N–H bending, 1600–1500 cm^{-1}) regions, leading to the final structure characterization. Interestingly, this spectroscopy allows the examination of proteins under different environmental conditions such as in solution or in the



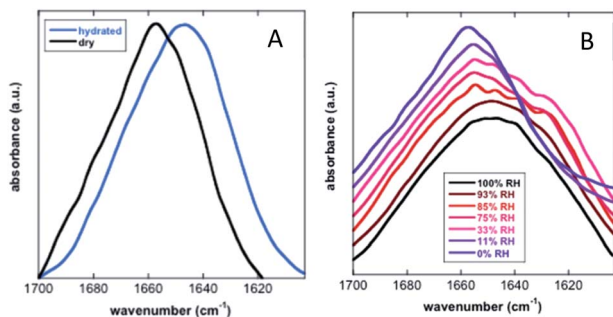


Fig. 8 FT-IR analysis of the secondary structure of TdLEA3. (A) Amide I region in the hydrated (D_2O , blue) and in the dry (black) state. (B) Amide I region at different relative humidities (RH). Reprinted with permission from Koubaa *et al.*¹⁰⁷ Copyright 2019 Springer Nature.

form of solid films. For a complete overview of this procedure and its spectra analysis, see Natalello *et al.*¹⁰⁶

Koubaa *et al.* investigated the structure of the 11-mer repeat motif TdLEA3, an IDR of the Late Embryogenesis Abundant (LEA) protein. The results showed that TdLEA3 was mostly disordered under aqueous conditions and acquired an α -helical structure in a dry medium (Fig. 8). Because H_2O overlaps in the amide I region, measurements in D_2O were carried out. Bands in the region between 1660 and 1650 cm^{-1} were assigned to the α -helix, between 1640 cm^{-1} and 1650 cm^{-1} to unordered regions and at around 1620 cm^{-1} to intermolecular β -sheet aggregates. The hydrated IDR was centred at 1648 cm^{-1} , indicating a mainly unstructured protein. Upon drying, this maximum shifted to 1657 cm^{-1} , indicating a more α -helix oriented structure.¹⁰⁷ Mohammad *et al.* utilized Attenuated Total Reflection Fourier Transform Infrared (ATR-FTIR) spectroscopy to study two intrinsically disordered protein α -synuclein variants, the IDP wildtype αS (αS -wt) and the naturally occurring splicing variant (αS - Δ exon3). A disordered state in the amide I spectra for both compounds as the initial state was observed. A slow aggregation process was observed over time, but with striking dissimilarities: αS -wt revealed two bands at 1665 cm^{-1} and 1618 cm^{-1} , whereas αS - Δ exon3 exhibited a broad band with a maximum at 1630 cm^{-1} . In the long term, both variants showed a conformational heterogeneity of secondary structures and aggregates but with some differences. The fibrillar aggregates dominated in αS -wt and the oligomers prevailed in αS - Δ exon3. αS -wt showed a very low frequency that indicated a well-ordered extended β -sheet and fibrils with strong hydrogen bonds formed between the backbone amide carbonyls along the fibril axis. For αS - Δ exon3, an absorption below 1630 cm^{-1} denoted the formation of typical β -structured aggregates.¹⁰⁸ Villarreal-Ramirez *et al.* used FT-IR to demonstrate how one IDR of dentin phosphoprotein (DPP), named P5, assumed different conformations when associated with Ca^{2+} or hydroxyapatite (HA). Furthermore, they showed that after P5 phosphorylation (P5P), DPP also adopted distinct conformations. In solution, P5 was disordered, while P5P displayed a more compact globular structure. P5 had a higher amide I intensity with a narrow band, whereas P5P had a broadened

amide I signal. Also, the P5 amide II band corresponding to the COO^- of the Asp side chain was less intense, whereas the absorbance for P5P increased due to the substitution of phosphoserine residues. In the presence of Ca^{2+} or HA, P5 adopted a random coil structure, whereas its phosphorylated counterpart had a more compact arrangement associated with conformations that showed β -sheet and α -helix motifs. P5, in the presence of Ca^{2+} or HA crystals, showed a slight decrease in the amide I region, whereas the amide II band intensity was increased. These changes were associated with a modest decrease in random coil and an increase in a beta turn structure. In P5P, after adding Ca^{2+} or HA, the amide I band was broadened due to the formation of β -sheet structures.¹⁰⁹ Vitali *et al.* analysed the structures of three IDPs (α -casein, Sic1 and α -synuclein) after interacting with silica NPs. α -Casein did not show conformational changes and continued being dominated by a peak at 1644 cm^{-1} , assigned to disordered structures. In the case of Sic1, the appearance of amide I shoulders at ~ 1638 cm^{-1} and 1689 cm^{-1} denoted a clear random coil to β -sheet transition. Finally, α -synuclein displayed a similar behaviour, the silica induced intermolecular interactions with a final β -sheet morphology (1627 cm^{-1} and 1696 cm^{-1}).¹¹⁰

Small-angle X-ray scattering (SAXS)

SAXS was initially used exclusively to qualitatively monitor folding/unfolding processes. Currently, unlike most other structural methods, this technique is applied to equilibrium and non-equilibrium mixtures to monitor kinetic processes extracting the quantitative information of IDPs. For the reminder of the main theoretical and experimental aspects of X-ray scattering applied to IDPs, see Bernadó and Svergun.¹¹¹

Jephthah *et al.* studied the N-terminal IDR of the magnesium-transporter-A protein (KEIF). By the analysis of the form factor, Kratky plot and Pair Distance-Distribution Function (PDDF), they could observe the natively unfolded behaviour of this region. The absence of a minimum in the first one and a maximum in the second graph were typical curve shapes of a fully flexible and extended peptide.⁴² Lenton *et al.* studied the phosphorylation effect on the recombinant human-like osteopontin (rOPN). A plateau at high q values in the Kratky plot and the asymmetrical shape of the PDDF confirmed its highly unfolded and flexible behaviour. Also, phosphorylation appeared to have minimal effect on the solution scattering of rOPN, reflected in an overall unchanged conformation at the SAXS resolution.¹¹² Didry *et al.* observed how a few amino acids in the IDR β -thymosin control the actin peptide self-assembly process. In a 1 : 1 stoichiometric ratio with actin, the β -thymosin inhibits its assembly by sequestering its monomers like thymosin- $\beta 4$. In other words, an exchange in the β -thymosin linker -Phe-Asn-Gln-Asp-Lys- with a -Phe-Asp-Lys-Ser-Lys- one decreased the β -thymosin:actin binding affinity, showing the last linker mentioned a different structure in the SAXS spectra.¹¹³ Cragnell *et al.* proposed a molecular mechanism of oligomerization directed by divalent cations. After adding Zn^{2+} to histatin 5, a clear relationship between the cation concentration and the IDP was observed (Fig. 9). The



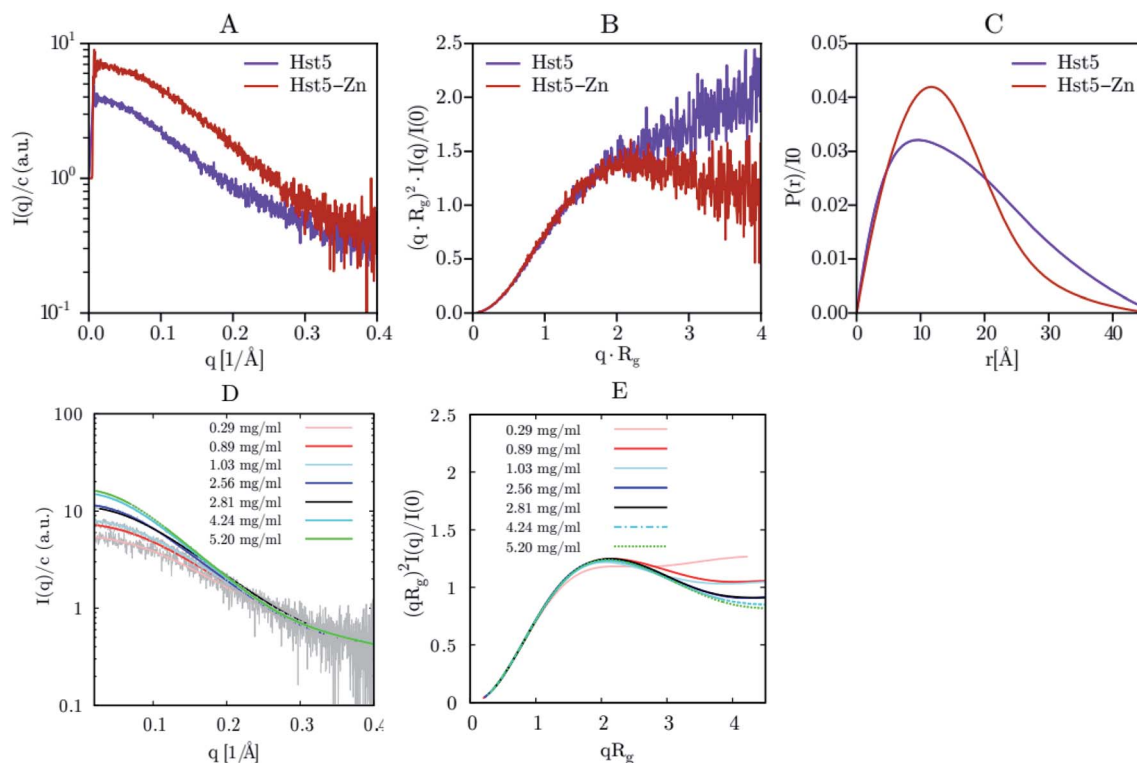


Fig. 9 SAXS analysis of Hst5 in the absence and presence of ZnCl_2 . (A) Comparison of the intensity function normalised by concentration for 0.9 mg mL^{-1} Hst5, in 20 mM MES-buffer, pH 6.7, 150 mM NaCl and 4 mM ZnCl_2 . (B) SAXS data shown as a dimensionless Kratky plot. (C) Plot of the intra-peptide distance distribution determined by indirect Fourier transform, for Hst5, with either NaCl (purple curves) or ZnCl_2 (red curves). (D and E) Concentration dependent SAXS-measurements of Hst5 in the presence of ZnCl_2 , showing the intensity curve normalised with protein concentration and the corresponding Kratky plot. Reprinted with permission from Cragnell *et al.*¹¹⁴ Copyright 2019 MDPI.

addition of Zn^{2+} resulted in an increase of $I_{(0)}$ corresponded to an increase in the measured molecular mass. Furthermore, a less linear plateau in the Kratky plot also concluded that the cation led to a compaction of the overall protein. To conclude, this compaction was also supported by a redistribution of the PDDF towards shorter distances in the protein, moving to a more Gaussian-like structure in the presence of zinc.¹¹⁴ Hardouin *et al.* observed the behaviour of the RNaseY N-terminal IDR (BsRNaseY). The resulting SAXS curve averaged on the plateau gave values for R_g and the maximal extension, D_{max} , significantly higher than those expected for a 176-residues compact protein. Therefore, the D_{max} and R_g values indicated a highly elongated shape. Moreover, PDDF and Kratky curves were significantly different from that of a fully unstructured protein. Coupling this qualitative information with the strong propensity of BsRNaseY to form coiled-coil structures, they could finally fit the SAXS model factor to a representative central coiled-coil conformation appended with flexible ends.¹¹⁵

Static and dynamic light scattering (SLS & DLS)

SLS and DLS can identify several different physical macromolecule parameters. SLS can measure molar masses within the 10^3 – 10^8 g mol^{-1} range, directly related to the state of association of IDPs in solution. On the other hand, DLS is an appropriate technique to monitor the expansion or compaction of protein molecules and their Stokes radius, R_s . The radius of

gyration, R_g , that could be obtained by SLS is not easily elucidated due to the small size of this systems. For a long and detailed explanation of the DLS and SLS basics applied to IDPs and IDRs, see Gast and Fiedler.¹¹⁶

Chinak *et al.* used DLS to elucidate the structure–activity relationship of an analogue of lactaptin, RL2. They studied the structural and aggregation features of this fairly large intrinsically disordered fragment of human milk κ -casein. This IDR, due to its Pro and Gln-enriched AA sequences, self-assembled into micellar formation or amyloid fibrils, preventing casein precipitation in milk. Changes in pH from 5.5 to 8.0 and the addition of NaCl led to a dramatic increase in the diameter distributions, related to its oligomerization ratio. Under extracellular environmental conditions, RL2 led to large 700 nm diameter oligomers, while at pH 5.5 (corresponding to early endosomes), RL2 was predominantly in monomeric/dimeric forms, and its oligomers had a size of *ca.* 200 nm. Also, the presence of physiological ionic strength caused RL2 to oligomerize at lower pH (*ca.* 430 nm and *ca.* 290 nm diameter with and without NaCl, respectively).⁹⁷ Zsila *et al.* observed how the addition of a drug or dye induced a self-organization on the cationic IDP CM15. After the addition of these ligands to the CM15 system in a ratio 1 : 1 or 1 : 2, an increase in the hydrodynamic radius of *ca.* 1000 nm was obtained, after the formation of large aggregates. The mutual charge neutralization within the complexes composed of cationic CM15 and its



anionic partners could be reached. As a consequence, the resulting adducts became less hydrophilic and were prone to aqueous aggregation. Further increase of the ligand concentration (2 : 1 ratio) decreased the broadness of their size distribution.¹¹⁷ Khatun *et al.* used both techniques, DLS and SLS, to observe the possible structures of the IDP human amylin. The peptide self-assembled in aqueous media, as demonstrated by the presence of a small percentage of protofilaments with diameter values from 200 to 400 nm along with matured fibrils (>1000 nm). After a short sonication step, a reduction in the average size of the protofilaments (to 100–200 nm) as well as the fibril maturation (*ca.* 1000 nm) were observed. If extended to 30 min, even smaller protofilaments (50–100 nm) and slightly smaller fibrils (1000 nm) were detected.⁷¹ To conclude, Shou *et al.* also analysed the conformation selection, in this case, of the IDP COR15A. SLS was used to obtain a hydrodynamic radius of 2.5 nm, which corresponded to a typical IDP. At glycerol concentrations above 5.47 osM, an increase in R_S up to 3.4 nm was observed, which is far above the scaling behaviour of IDPs and denoted the COR15A oligomer formation. After obtaining the R_g by DLS, the R_g/R_S ratio concluded that the IDP had a slightly oblate shape in the absence of glycerol (ratios between 0.875 and 0.987) and adopted a more elongated conformation at values of osmotic concentrations between 1 and 3 osM ($R_g/R_S > 1$). The aggregate structural change also could revert back to an oblate ellipsoid at higher glycerol concentrations.¹¹⁸

Force fields and simulations

Experimental techniques for the characterization of IDPs/IDRs offer information on an average conformation upon binding

or recognition process. The conformational ensembles of IDPs/IDRs still far exceed the number of available experimental observables. Thus, theoretical models are a suitable alternative for extracting detailed structural information at the atomic level, which experimental techniques cannot provide. Molecular Dynamics (MD) and Monte Carlo (MC) simulations are powerful tools to fill this gap. They produce a time sequence of atomic-level configurations and offer a potentially powerful complement to elucidate the key conformational characteristics of IDPs/IDRs. Even more, the atomistic details obtained from force field-based simulations can be used to help interpret experimental results. MD, while being the most commonly applied to IDPs/IDRs, rely on the accuracy of the underlying potential energy functions or force fields, so performing accurate structural characterization is a challenging task.

Various force fields have been developed to describe biomolecular structures in aqueous environments. In this section, we will show some of the currently relevant force fields used in recent studies focused on the IDP/IDR field, which are also summarized in Table 3.

Assisted model building with energy refinement (AMBER)

AMBER is a suite of biomolecular simulation programs which started to be designed in the late 1970s by Peter Kollman.¹¹⁹ The energy function form of this force field used in protein, nucleic acid and organic molecule simulations is described as:

$$E_{\text{total}} = \sum_{\text{bonds}} K_r (r - r_{\text{eq}})^2 + \sum_{\text{angles}} K_\theta (\theta - \theta_{\text{eq}})^2 + \sum_{\text{dihedrals}} \frac{V_n}{2} [1 + \cos(n\phi - \gamma)] + \sum_{\text{non-bonding}} \left[\frac{A_{ij}}{R_{ij}^{12}} - \frac{B_{ij}}{R_{ij}^6} + \frac{q_i q_j}{\epsilon R_{ij}} \right] \quad (1)$$

Table 3 Relevant force fields applied in IDP/IDR studies

Force fields	Parameter sets	Changes	References	
AMBER	ff99	First AMBER parameter set	150 and 166	
	ff99SB	Improved backbone torsional term	74, 129, 143, 148, 150 and 166	
	ff99SB*	Corrected backbone energy term	129, 143 and 146	
	ff99SB-ILDN	Improved side-chain torsion term	44, 45, 49, 125, 129, 131, 132, 134, 135, 136, 150–152, 166 and 167	
	ff99SB*-ILDN	Improved side-chain torsion term	128, 132, 137, 139, 147, 148 and 151	
	ff99SB-DISP	Corrected protein and water vdW terms	127, 128 and 136	
	ff14SB	Improved backbone and side chain	63, 126, 127, 130, 134, 137, 138, 149, 150 and 167	
	ff14IDPSFF	Corrected backbone torsional term	126, 127 and 138	
	ff03	Second AMBER parameter set	135 and 166	
	ff03w	Corrected backbone torsional term	132, 146 and 147	
	ff03ws	Modified protein–water interaction term	127, 128 and 139	
	CHARMM	CHARMM22	CHARMM parameter set	47, 70, 129, 143, 158 and 166
		CHARMM22*	Corrected backbone energy term	128, 136, 139, 146, 147, 148, 150–152 and 167
		CHARMM36	Modified backbone and side-chain torsional term	41, 67, 129, 143, 146, 147, 149 and 150
CHARMM36m		Corrected backbone conformational term	127–129, 132, 135–137, 144, 149 and 150	
CHARMM36IDPSFF		Corrected backbone torsional term	145	
GROMOS	GROMOS96 43a1	GROMOS parameter set	111, 150 and 152	
	GROMOS96 53a6	Improved hydration thermodynamics reproduction	107, 131, 150, 152, 158 and 166	
	GROMOS96 54a7	Improved torsional term and hydration energy	107, 131, 139, 150, 152, 158 and 159	
OPLS	OPLS-AA	OPLS parameter set	64, 139, 150, 158, 166 and 167	
	OPLS-AA/L	Refitted Fourier torsional term	135, 147 and 161	
	OPLS-AA/M	Refitted Fourier torsional term	164	
	OPLSIDPSFF	Corrected backbone torsional term	165	



In brief, the model represents the bonds and angles by a simple diagonal harmonic expression, the dihedral energies by a simple set of parameters (often only specified by the two central atoms) and the non-bonding energies, electrostatic and van der Waals (vdW) interactions, are only calculated between atoms in different molecules or for atoms in the same molecule separated by at least a three bond distance. The first non-bonding energy value is modelled by a coulombic interaction of atom-centred point charges while vdW is represented by a 6–12 potential.^{120,121} To use the AMBER force field, it is necessary to have the preliminary parameter values of the force field (*e.g.* force constants, charges, equilibrium bond lengths and angles). Several authors during the last few decades have applied different base parameter sets, improving the original force field and finally leading to several parameter sets optimized for each analysed system. Here, we show the strength and weaknesses of the AMBER force field using some examples of its application to IDPs/IDRs. To know more about this package of computer programs, see Case *et al.*¹²²

Based on relevant recent modifications of the force field for IDPs/IDRs, Chen group, based on AMBER ff99SB-ILDN, developed the AMBER ff99IDPs. They refined the IDPs sampling by transplanting residue-specific grid-based energy correction maps (CMAPs) corrections of eight disordered promoting residues (Ala, Arg, Gln, Glu, Gly, Lys, Pro, and Ser), improving the φ/ψ dihedral terms.¹²³ This approach was followed by Song *et al.*, who extended these CMAPs to all 20 amino acids, and proposed AMBER ff14IDPSFF, that raised its quality in the reproducibility of secondary chemical shifts of multiple short disordered proteins. 14 unstructured short peptides showed similar results

between the simulated C_α chemical shifts obtained by NMR and the ff14IDPSFF force field. As an example, ff14IDPSFF produced diverse β -sheet conformers for the Tau protein, consistent with previous experimental observations.¹²⁴ Continuing with this tendency, Song *et al.* also applied the CMAP approach and presented an environmental specific precise force field (ESFF1) to improve the accuracy and efficiency of MD simulations for both, IDPs and folded proteins.¹²⁵ Meanwhile, Robustelli *et al.* proposed another force field, AMBER ff99SB-DISP, that could describe ordered, disordered, and transitional regions. They were able to achieve this goal by modifying the water model and iteratively testing small changes in backbone torsion corrections and the strength of the backbone O–H Lennard-Jones (LJ) pair.¹²⁶ Best *et al.* started from AMBER ff03 and proposed AMBER ff03ws, strengthening the LJ potential for protein–water interactions and applying a scaling factor for protein–water interactions.¹²⁷ Lately, Yu *et al.* introduced a residue-specific protein force field, ff99SBnmr2, derived from ff99SBnmr1. A different balance at the backbone dihedral angle potentials quantitatively better reproduced the dihedral angle distributions from a set of experimental coil systems.¹²⁸

Focusing on their applicability, Henriques *et al.* applied AMBER ff99SB-ILDN and AMBER ff03ws to the IDP histatin 5, achieving a reasonable balance between protein–protein and protein–water dispersion interactions using a TIP4P-D and TIPAP/5 water model, respectively.¹²⁹ Pietrek *et al.* observed the local and overall dimensions of the IDP α -synuclein. By using AMBER ff99SB-ILDN, they could match the modelling structure to the NMR and SAXS experimental data, complemented with AMBER ff03ws simulations for the possible force field issues of

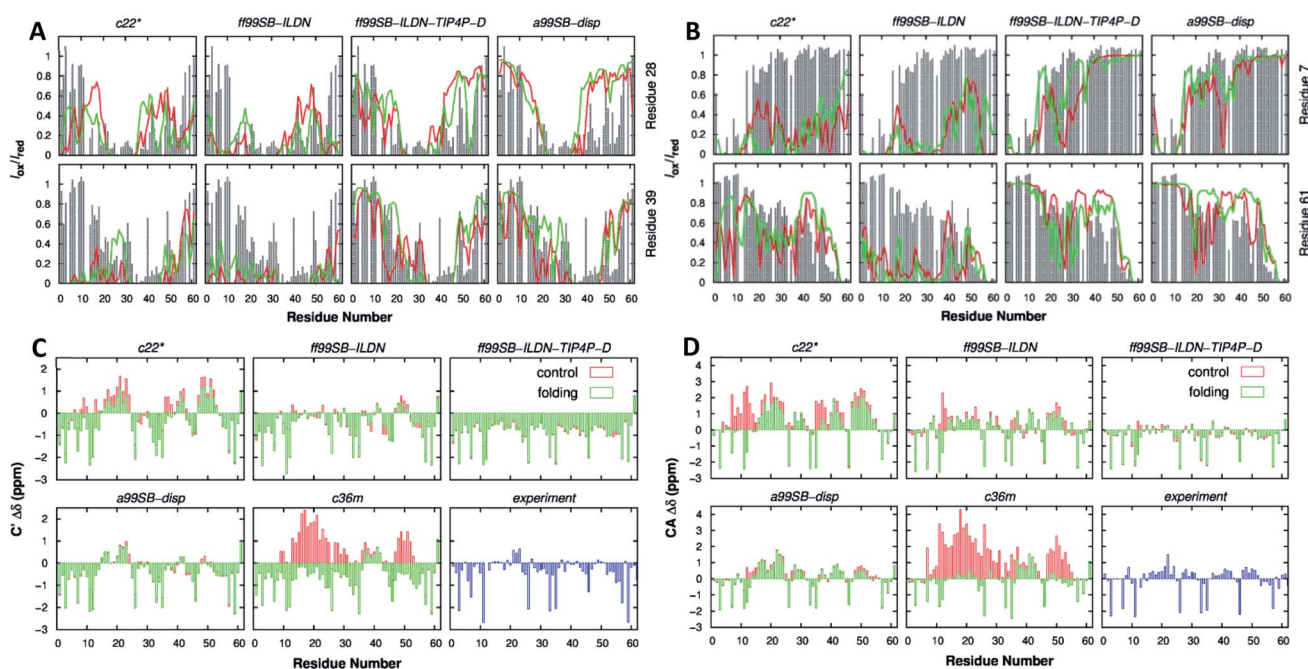


Fig. 10 p53 61-residue N-terminal TAD (A) calculated (lines) and experimental (gray bars) paramagnetic relaxation enhancement effects induced by paramagnetic spin labeling at residues 28 (top row) and 39 (bottom row) and (B) residues 7 (top row) and 61 (bottom row). (C) Secondary chemical shift analysis for C_α atoms and (D) C' atoms. Calculations were performed using independent control (red) and folding (green) simulations. Reprinted with permission from Lui *et al.*¹³⁴ Copyright 2019 American Chemical Society.



the full-length coiled structure.¹³⁰ Rieloff *et al.* also compared this force field with CHARMM36m but in the 15-residue-long N-terminal fragment of the IDP fragment (SN15n) before and after phosphorylation (SN15p). While both force fields agreed regarding the size and shape of SN15n, for SN15p the CHARMM36m force field denoted strong interactions in the form of hydrogen bonding between the phosphorylated amino acids and the Arg residues. AMBER ff99SB-ILDN showed a less compacted structure higher in helical content, closer to what the CD experimental data suggested.¹³¹ Joseph *et al.* compared AMBER ff99SB-ILDN with older force fields such as AMBER ff14ipq or AMBER ff14SB in the human CD4 receptor, an IDR. Overall, ff99SB-ILDN performed better than its predecessors in terms of reproducing the HN-NMR shifts and J coupling constants. In the N-terminal peptidic region, ff14SB predicted more helical structures and ff14ipq more disordered ones than those observed experimentally.¹³² Ouyang *et al.* analysed the conformational features of the p53 distinct activation domain 2 (TAD2) IDR with different force fields. They concluded that AMBER ff99SB-ILDN showed a structural dimension closer to that theoretically predicted, with a more heterogeneous conformation. This force field provided correct results for p53 TAD2, whereas other force fields led to a collapse of the system. Force fields like CHARMM27 tended to over-stabilize a helical structure, CHARMM36m produced a most expanded coil ensemble and OPLS-AA/L exhibited a strong preference on a β -sheet structure, far from experimental results.¹³³ In a later study, Lui *et al.* demonstrated that the AMBER ff99SB-DISP force field had the best agreement with the experimental data obtained for, in this case, the p53 61-residue N-terminal TAD (Fig. 10). The AMBER ff99SB-DISP force field seemed capable of faithfully recapitulating virtually all experimental characterization results, including the overall chain dimensions, residual secondary structures, and transient long-range ordering. CHARMM36m and CHARMM36mw (CHARMM36m with a new water force field) failed to generate converged ensembles despite using multiple microsecond simulation time scales. CHARMM22* generated overly compact structural ensembles and an overestimation of the residual helicity, like AMBER ff99SB-ILDN.¹³⁴ Kuzumanic *et al.* supported the use of AMBER ff99SB-DISP for the Von Willebrand Factor (VWF) study, being the force field that overall agreed best with the NMR data, followed by RSFF2+ and CHARMM36m ones. While all the force fields kept the β -sheets of the rigid VWF E' domain in place, the TIL' domain showed differences. By NMR, it could be inferred that this domain, as an IDR, lacked a secondary structure except for one 3_{10} -helical and three β -sheet regions. The AMBER ff99SB-DISP force field agreement with NMR came from the comparison of NOE distance restraints, chemical shifts, and backbone dihedral angles.¹³⁵ Duong *et al.* were successful in simulating short peptides with a Glu-Gly-Ala-Ala-X-Ala-Ala-Ser-Ser structure (X = Asp, Gln, Glu, His, Leu, Lys, Pro, Trp, Tyr). Two force fields were tested and, while AMBER ff14SB denoted an increased helical content, a coiled content was obtained for ff14IDPSFF, with the latter in higher agreement with NMR and CD experiments.¹³⁶ Henriques *et al.* also reported the improvement of AMBER ff03w compared to old AMBER and

GROMOS force fields for the histatin 5 model. Previous models exhibited considerable bias towards overly compact conformational ensembles (force field independent) and certain secondary structure motifs (force field dependent), over-stabilizing the structure.¹²⁹ Carballo-Pacheco *et al.* defended the use of AMBER ff03ws in the study of the aggregation and non-aggregation of the Alzheimer related $A\beta_{16-22}$ IDP. GROMOS 54a7 and OPLS-AA strongly over-stabilized protein-protein interactions. AMBER99SB*ILDN and CHARMM22* were also considered, even though they were not that accurate.¹³⁷

Chemistry at Harvard macromolecular mechanics (CHARMM)

CHARMM is a well-known and widely used set of force fields for molecular dynamics developed by Martin Karplus that can be used for DNA, RNA, lipids, drug-like molecules and especially proteins.¹³⁸ The general form of the potential energy function most commonly used in CHARMM for self-assembling amphiphilic peptide simulations is based on fixed point charges and described as:

$$\begin{aligned}
 E_{\text{total}} = & \sum_{\text{bonds}} K_r (r - r_{\text{eq}})^2 + \sum_{\text{angles}} K_\theta (\theta - \theta_{\text{eq}})^2 \\
 & + \sum_{\text{Urey-Bradley}} K_{\text{UB}} (S - S_{\text{eq}})^2 + \sum_{\text{dihedrals}} K_\phi [1 + \cos(n\phi - \gamma)] \\
 & + \sum_{\text{impropers}} K_\omega (\omega - \omega_{\text{eq}})^2 + \sum_{\text{non-bonding}} \left\{ \epsilon \left[\left(\frac{R_{ij}^{\text{min}}}{r_{ij}} \right)^{12} \right. \right. \\
 & \left. \left. - \left(\frac{R_{ij}^{\text{min}}}{r_{ij}} \right)^6 \right] + \frac{q_i q_j}{\epsilon_1 r_{ij}} \right\}
 \end{aligned} \quad (2)$$

While terms like bond stretches, angles, dihedral force, or non-bonded forces appear as in the basic AMBER force field, two more are added in CHARMM. These terms account for the out of plane bending and the Urey-Bradley component, a cross-term accounting for the angle bending using 1,3 non-bonded interactions in the harmonic potential.¹³⁹ A significant number of groups around the world are working on the development of the CHARMM package. Among them, the Charles L. Brooks III group deserves a special remark for their multiple improvements. Thus, we recommend their deep analysis of the program, from the basics to its implementation in different systems.¹⁴⁰

After the improvement of the CHARMM22 force field by MacKerell *et al.* in the form of CHARMM22*,¹³⁹ Best *et al.* proposed the CHARMM36 force field. They validated it by the comparison of: (i) simulations of eight proteins; (ii) backbone scalar couplings for each IDP/IDR; (iii) NMR residual dipolar couplings and scalar couplings for both the backbone and side-chains in folded proteins; (iv) folding equilibrium of peptides.¹⁴¹ Huang *et al.* presented the refinement of the CHARMM36 protein force field, the known CHARMM36m, improving the accuracy in generating polypeptide backbone conformational ensembles for intrinsically disordered peptides and proteins. The field was validated using a comprehensive set



of 15 peptides and 20 proteins. In general, the sampling of α -helical conformations in IDP ensembles generated with the CHARMM36m force field was significantly lower than in ensembles generated with CHARMM36, in agreement with experimental data. Examples of it were the arginine-serine peptide, the FG-nucleoporin peptide, a hen egg white lysozyme N-terminal fragment, and the N-terminal domain of HIV-1 integrase.¹⁴² On the other side, Liu *et al.* developed the CHARMM36IDPSFF, which showed an improvement over the CHARMM36 force field in 18 IDPs, even though some limitations were found in the radius of gyration of large disordered proteins and the stability of fast-folding ones.¹⁴³

Lazar *et al.* carried out 24-residue Ser/Arg-rich (SR22-45) MD simulations using CHARMM22* and CHARMM36. The histogram of R_g distributions, compared to experimental data, showed a higher than real compactness in the CHARMM36 model, making the CHARMM22* force field the way to go. Results that also supported the use of CHARMM22* for this IDR were also reported by Rauscher *et al.*^{144,145} Carballo-Pacheco *et al.* tested the ability of five force fields to model the IDP A β 42. Comparing their results to NMR experimental data, they observed how CHARMM22* was the best force field for reproducing C_α and HN chemical shifts associated with a β -hairpin structure. Particularly, CHARMM22* generated fewer compact conformations without the recalibration of protein-water interactions, as AMBER ff99SB*ILDN or AMBER ff03w. Older force fields like OPLS, GROMOS or CHARMM22 showed unreal

structures.¹⁴⁶ These results were updated by Krupa *et al.*, who applied the main force fields currently used. They concluded that CHARMM36m > CHARMM36 > CHARMM22* force field for the IDP A β 42. In CHARMM36m, the monomeric A β 42 structure was less stable and more hydrophilic compared to AMBER. That could be explained by water interactions, which played a much more important role in CHARMM compared to AMBER.¹⁴⁷ Man *et al.* compared 17 different force fields: 7 from the AMBER and GROMOS families, 3 from the CHARMM and one from the OPLS (Fig. 11). Applied to the seven-residue IDR fragment A β 16–22, just 5 force fields were able to denote the real amyloid peptide assembly by providing good balances in terms of structures and kinetics. Among them, all the CHARMM force fields included (CHARMM22*, CHARMM36 and CHARMM36m) reported great results. While the old AMBER force fields predicted α -helices, far from real, and the GROMOS-family formed β -sheets too rapidly, CHARMM force fields matched the CD and NMR experimental data.¹⁴⁸ Watts *et al.* compared the conformational space of the A β 1–40 dimers using several force fields. They concluded that CHARMM22* and CHARMM36 were the chosen ones for explaining the collapse of the central and C-terminal hydrophobic cores from residues 17–21 and 30–36 and reproduced a theoretically expected β -sheet-turn- β -sheet conformational motif.¹⁴⁹ These results were in agreement with Somavarapu *et al.*, who defended the use of CHARMM22* over every AMBER, GROMOS or OPLS force field.¹⁵⁰

Groningen molecular simulation (GROMOS)

The GROMOS force fields are united atom force fields, *i.e.* without explicit aliphatic (non-polar) hydrogens. Developed in 1978 for the dynamic modelling of biomolecules, it was a simulation computer program package released by the research group of Wilfred van Gunsteren, who also realized a substantial rewrite of it in 1996.^{151,152} Known for having two different versions, GROMOS software can be applied to aqueous or non-polar solutions of proteins, nucleotides, and sugars (GROMOS force field A-version) or to simulate gas phase isolated molecules (B-version). For an understanding of how GROMOS, in general, and GROMOS05, specifically, work, see Christen *et al.*¹⁵³

The force field was updated twice during the last decade, leading to GROMOS 53a6 and GROMOS 54a7. The first one was done by Oostenbrink *et al.*, introducing a new set of charges into the system to reproduce more accurately the hydration free enthalpies in water but with a drawback, an underestimation of the helical behaviour of peptides and proteins.¹⁵⁴ The second one was developed by Schmid *et al.* Several corrections were applied, being the most relevant one the adjustment of the torsional angle terms to correct the helical inaccuracy mentioned before.¹⁵⁵

While it is not the best force field for IDPs/IDRs compared to the AMBER and CHARMM families, authors such as Gerben *et al.* concluded that GROMOS96 54a7 and GROMOS 53a6, along with OPLS-AA, were the best force fields to explain the β -strand content in the intrinsically disordered amyloid β -peptide (A β). AMBER ff03 and CHARMM22 over-stabilized a helical

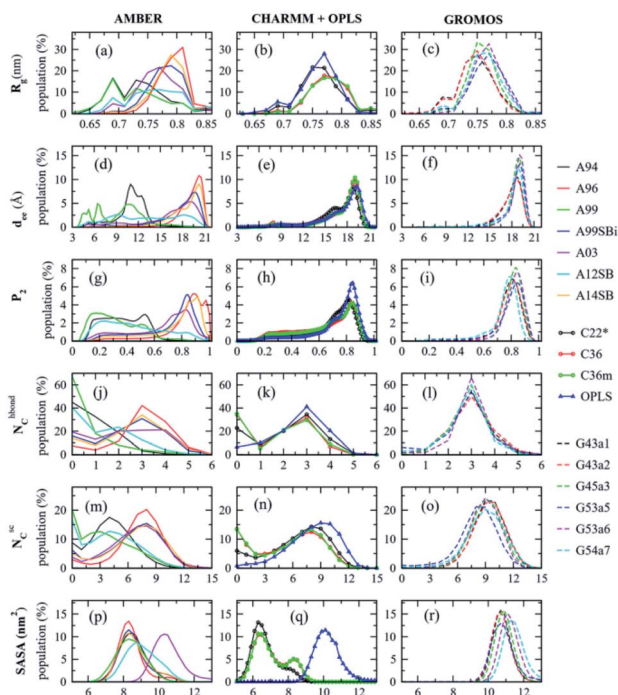


Fig. 11 A β 16–22 Dimer normalized distributions of the radius of gyration (R_g), the end-to-end distance (d_{ee}), the order parameter (P_2), the intermolecular backbone H-bonds (N_C^{bond}), the intermolecular side chain-side chain contacts (N_C^{sc}), and the solvent accessible surface area (SASA). Reprinted with permission from Man *et al.*¹⁴⁸ Copyright 2019 American Chemical Society.



structure and also could produce elongated A β structures, as for the last force field, far from what NMR shifts and R_g experimental data showed.¹⁵⁶ Also, GROMOS96 54a7 was used by Bandyopadhyay *et al.* to study two IDRs (the scaffolding protein GPB from *Escherichia* virus phix174, 1CD3, and the human coagulation factor Xa, 1F0R) as well as two IDPs (α -synuclein, α -syn, and amyloid beta, A β 42). 1CD3 showed three different structural conformations, α -syn had five and the last two peptides six. 1CD3 proposed structures were relatively more self-similar to each other by having the highest secondary structural as well as helical content, which made 1CD3 the closest peptide to the globular class out of all of them. α -Syn had more structural diversity yet a continuous transition behaviour. 1F0R and A β 42 both had appropriate diverse structural phases with a substantial self-similarity among the conformational phases.¹⁵⁷

Optimized potentials for liquid simulations (OPLS)

The OPLS force field was developed by William L. Jorgensen and started with a functional form very similar to the one used by AMBER. Currently used IDP-specific force fields have been mostly derived from the force fields of AMBER and CHARMM, while little attention has been paid to the widely used OPLS family. OPLS potential energy is expressed in a summary of 4 terms as:

$$E_{\text{total}} = E_{\text{bonds}} + E_{\text{angles}} + E_{\text{dihedrals}} + E_{\text{non-bonding}} \quad (3)$$

One of the most relevant differences that can be observed is at the non-bonded interactions; while the charges used in the OPLS force fields are empirical, in AMBER they are obtained on a case-by-case basis from fitting to electrostatic potential surfaces from *ab initio* 6-31G* calculations. For a complete understanding and a comparison between OPLS and other force fields, such as AMBER and CHARMM, see Jorgensen *et al.*¹⁵⁸

Focused just on the improvement of the force field and after the RSFF1 modifications of OPLS-AA/L by Jiang *et al.* and Xun *et al.*,^{159,160} Robertson *et al.* presented OPLS-AA/M. This force field demonstrated a significant improvement over previous OPLS-AA force fields. This model can be applied to normal peptides and IDPs out-performing previous OPLS-AA¹⁵⁸ and

OPLS AA/L¹⁶¹ dihedral parameters. Their ability to reproduce both gas phase conformer energies for longer peptides and aqueous phase experimental properties in molecular dynamics simulations was improved.¹⁶² The residue-specific force field OPLSIDPSFF, based on OPLS-AA/L, corrected the backbone dihedral term for all 20 residues by two-dimensional CMAPs (Fig. 12). IDPs and two short peptides were tested showing an agreement with NMR experimental results. The force field could obtain the β -sheet structures of GB1, while not stabilizing helix structures for the proteins AAQAA3 and GB1. In addition, the remaining disability of helical structures could be addressed by: (i) a novel CMAP refinement schedule, (ii) a more precise water model, or (iii) incorporating electronic polarization in a next step.¹⁶³

Smith *et al.* examined the dynamic behaviour of the IDR A β _{21–30} under seven force fields. Analysing the secondary structure, AMBER-family force fields, CHARMM27-CMAP, and GROMOS 53a6 were hindered in finding the local minima due to their enhancement of helical structures. OPLS-AA showed a substantially greater overall number of intrapeptide hydrogen bonds and suggested a metastable β -hairpin motif, associated with previous experimental results. Even more, OPLS was preferred as its sampling of Ramachandra space was more attuned to steric restrictions.¹⁶⁴ Man *et al.* also supported the application of the OPLS-AA force field, but in this case, for the A β _{1–42} IDR. While AMBER SB14 and CHARMM22* ensembles significantly overestimated the CD-derived helix content, OPLS-AA, followed by AMBER ff99SB-ILDN, denoted a more accurate β -hairpin secondary structure. In the 17–21 and 30–36 regions, 8% and 13% β -hairpin were observed by both force fields, respectively, while AMBER SB14 showed only 1.5% and CHARMM22* 5%.¹⁶⁵ Fluit *et al.* observed that OPLS-AA/L, along with AMBER ff99SB and AMBER ff99SB*, was the most suitable for studies of polyglutamine (polyQ) folding and aggregation when comparing 12 force fields. OPLS-AA/L denoted predominantly disordered and collapsed conformations in water. CHARMM22*, and CHARMM36 exhibited no obvious biases in secondary structures but do exhibited larger persistence lengths, leading to more extended, aspherical, and diffuse conformations in water. CHARMM27 predicted a large fraction of helical secondary structures. GROMOS96 54a7 appeared to under-stabilize α -helices and over-stabilize β -sheets while GROMOS96 53a6 also failed in predicting a large fraction of β -strand content.¹⁶⁶

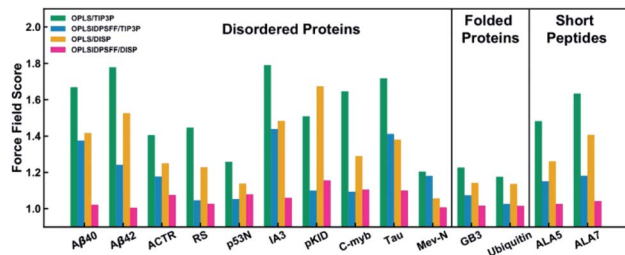


Fig. 12 Normalized force field scores (lower the better) for short peptides, folded proteins, and disordered proteins. OPLS and OPLS-SIDPSFF represent the original OPLS-AA/L and the new force field, respectively. DISP means the disp-TIP4PD solvent model. Reprinted with permission from Yang *et al.*¹⁶³ Copyright 2019 American Chemical Society.

Applications

Given all the possibilities that intrinsically disordered proteins or regions bring about, their applicability is also wide. Their multiple conformations, such as their core unique property position, make these biological structures highly relevant in the biomedical field. Already extended in several fields, IDP/IDR implementations are mostly in biology or biomedicine fields. Moreover, artificial disordered peptides are engineered to improve and tune up even more their performance and adapt their behaviour to specific functionalities. Due to the extension and all the possibilities that these short peptides structures can



support, and not being the focus of this review, only a representative small sample of bio-related recent studies are summarized below.

Focused on the biology field, cells present compartments called organelles to carry out their inner functions. However, membrane-less organelles formed *via* active liquid–liquid phase separation (LLPS) have garnered interest during the last few years. Proteins, peptides, and AAs can condense while being surrounded by a light phase, leading to a two-phase regime. Thermodynamically controlled, this process is based on intermolecular as well as water–water interactions, mostly by hydrogen bonding. However, this stimulus-responsive process is directed by external stimuli and environmental changes such as salt or molecule concentration, pH and temperature. Dzuricky *et al.* analysed a total of 63 IDPs that formed these membrane-less organelles in order to determine common structural features to exploit in future artificial IDPs. The octapeptide Gly–Arg–Gly–Asp–Ser–Pro–Tyr–Ser was the key to control LLPS processes by temperature and pH transitions. The formation and dynamics of their phase separation into coacervate droplets were controlled by two simple design parameters using *in vitro* and *in vivo* conditions: the molecular weight of the final octapeptide-based IDP and the aromatic : aliphatic ratio of residues in the octapeptide repeat.¹⁶⁷ Savastano *et al.* used the IDP Tau at the AT180 epitope to regulate the cell compartmentation and form liquid-like droplets. While Tau assembled into microtubules, AT180 underwent LLPS in solution and on the surface of the microtubules. From these results, phosphorylation processes were suggested as a mechanism to modulate the LLPS of IDPs in a condensate-mediated cytoskeletal assembly.¹⁶⁸ Metrick *et al.* reported the LLPS behaviour of the IDP UL11, from herpes simplex virus 1 (HSV-1). This tegument protein, while the process remains unclear, assembled as a biomolecular condensate in a complex network. Its disordered properties would form this membrane-less conformation, helping future biological processes such as membrane deformation during endocytosis.¹⁶⁹ Dogra *et al.* also formed membrane-less organelles. In this case, they were controlled by using a pH-responsive IDR comprising 10 imperfect repeats rich in hydrophobic, polar, and acidic residues. Based on Ala, Gly, Thr, Pro, Ser and Val residues, this Pmel17 protein disordered domain promoted the formation of liquid droplets at neutral cytosolic pH that formed solid aggregates. At a mildly acidic melanosomal pH, the monomers self-assembled into amyloid fibrils in a reversible way.¹⁷⁰ To study IDP LLPS relevance, Dignon *et al.* developed a model to predict temperature-dependent solvent-mediated interactions of each type of amino acid for further LLPS design. Sequences with an hourglass-shaped phase diagram or upper critical solution temperature behaviour generally were obtained for IDPs with more polar or charged residues than a typical IDP sequence.¹⁷¹ Using artificial simplified IDP models, Zhao *et al.* used elastin-like polypeptides (ELPs) in two compartmentalization strategies, namely bulk phase emulsion and cell-like compartment. ELP thermo-responsive phase transition properties allowed them to form membrane-less organelles *via* LLPS in the cellular milieu. This study is considered a significant step in the

building of cell-mimicking systems with a higher degree of hierarchical complexity.¹⁷² Faltova *et al.* conjugated soluble globular domains to low complexity domains (LCDs) of a few disordered amino acids. In this way, they developed molecular adhesives that enabled sensitive and controlled self-assembly processes into final supramolecular architectures. LCD regions, which contained a high fraction of charged and polar amino acids, led to liquid–liquid phase separation processes due to their colocalization behaviour while the globular domain maintained its functionality. These chimera proteins reversibly self-assembled into liquid droplets which evolved into irreversible protein aggregates and finally solid particles over time. Finally, they applied active porous solid particles as micro-reactors, releasing soluble proteins over time.¹⁷³ With a different application in mind, Urosev *et al.* used specific ELPs (hELPs) to restore the mechanical strength of fibrin networks, improve their clot development rate, reduce the plasmin degradation rate, and reduce the fibrin network pore size. IDPs mainly based on Val–Pro–Gly–X–Gly pentapeptides (with Ala, Glu and Val residues in guest X positions at a ratio of 2 : 8 : 1) coacervated at physiological temperature in β -spirals. The addition of a Gln residue to the N-terminal region, in the presence of the protein FXIIIa, covalently cross-linked the IDP by Lys–Gln interactions. After interacting with fibrinogen, thrombin and FXIII, hELP coacervates could be integrated into fibrin networks. These interactions took place through Gln- and Lys-residues on Fb γ -chains and α -chains, and AA cross-linked with hELP through its Gln- and Lys-blocks.¹⁷⁴ Hossain *et al.* used intrinsically disordered peptide-polymers (IDPPs) for post-translational modifications (PTMs) adding a lipid chain to encode non-equilibrium phase behaviour transitions, an emergent frontier in biomacromolecular engineering. The IDR was based on a tropoelastin (Gly–X–Gly–Val–Pro)₈₀ domain (containing a mixture of Ala : Val 2 : 8 in the X position), while the lipids tested were a canonical PTM (M-IDPP) and an azide (–N₃) non-canonical PTM (ADA-IDPP). Both IDPPs self-assembled into spherical micelles at room temperature. When heated above the lower critical solubility temperature (LCST) around 31 °C and then cooled again, the azide based-IDPP behaviour was totally different. Unlike myristic acid, the ADA chain could not efficiently pack inside the hydrophobic core due to the forced linear arrangement of the terminal azide group. With heat, an increase in the mobility could facilitate the rearrangement of ADA-IDPP, leading to the shifting of the spherical micelles into rod-like aggregates.¹⁷⁵ Wonderly *et al.*, based on a marine mussel IDP (Mfp), improved the adhesion and cohesion of peptidic structures by changing their backbone to a peptidic one.¹⁷⁶ Bulutoglu *et al.* designed a stimulus responsive peptide based on two domains. The first domain was an IDR that self-assembled into a β -roll conformation when binding to Ca²⁺ due to its Leu residues. The second one was also a repeats-in-toxin domain that could recognize the lysozyme protein in specific situations. Ca²⁺ ions were responsible of the β -roll formation and final gelation, while the protein binding helped to obtain even more robust hydrogel networks.¹⁷⁷

As stated before, IDPs/IDRs can participate in conditioning soft and hard extracellular matrices, among other structural



processes. One of the most relevant is based on biomineral-associated protein interactions with final biomedical applications. Rao *et al.* showed how IDRs appeared to not only regulate the finally formed biomineral structure, but also modulate the formation and stability of crystal precursors. Four unstructured peptides with a vesicular shape were able to control and inhibit crystallization processes *via*

a confinement-based mechanism. High Ca^{2+} concentrations forced organic–inorganic interactions and disorder-to-order transitions in these Gln, Thr or Ser rich peptides at high pH values. IDRs were able to interact with discrete mineral species and present lower free energy values, stabilizing and stopping the biomineralization process at intermediate structures between the Ca^{2+} ions and the final crystal conformation.⁵⁴ In

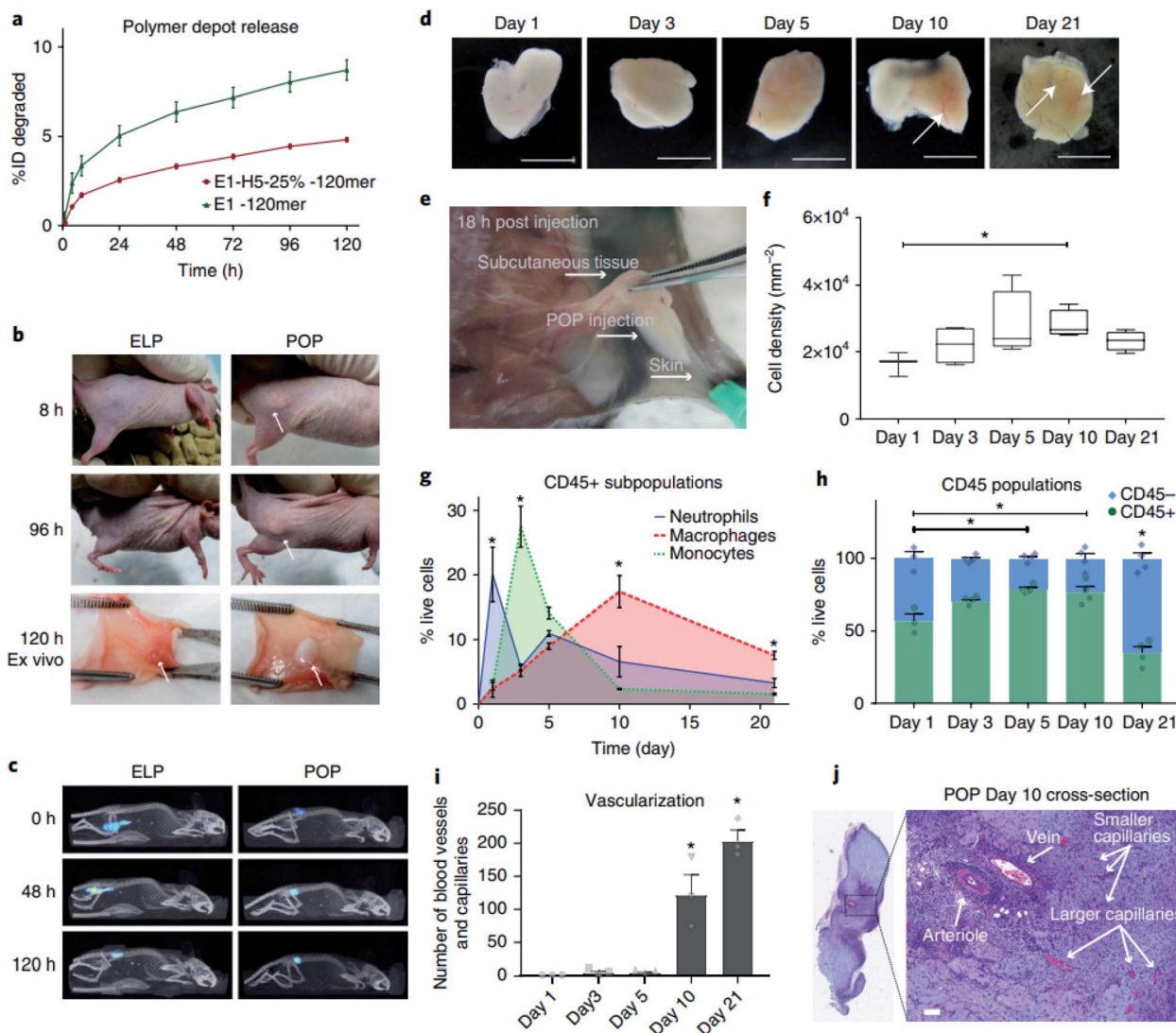


Fig. 13 *In vivo* stability and tissue incorporation of POPs: (a) ^{125}I radiolabelled E1-H5-25% POP subcutaneous injections were significantly more stable than their E1 counterparts, with just 5% of the injected dose (ID) degraded at 120 h; 200 μl 250 μM injections; $p < 0.05$ for all data points after 0 h, determined by two-tailed *t*-tests ($n = 6$ mice); data represent mean \pm s.e.m. (b) Whereas ELPs diffuse into the subcutaneous space, POP deposits were externally apparent, retaining the shape and volume of the initial injection up to dissection and *ex vivo* analysis. (c) Representative CT-SPECT images of the deposits confirm the increased diffusivity of ELPs and the increased stability of POPs. (d) POPs were injected into BL/6 mice and explanted for analysis over 21 days. Representative images are shown with arrows pointing at externally evident vascularization. Scale bars: 5 mm. (e) POPs rapidly integrated into the subcutaneous environment with sufficient strength to endure moderate extension less than 24 h after injection. (f) There is a high initial cell incorporation with some change over the observed time periods; for *, $p < 0.05$ determined by ANOVA with Tukey *post-hoc* (day 1 $n = 3$, days 3–21 $n = 4$); data presented as 10–90% box plots. (g) Flow cytometry for cells involved in innate immunity reveals subsequent spikes in neutrophils, inflammatory monocytes, and macrophages, with a loss in all haematopoietic cells (CD45+) by day 21; for *, $p < 0.05$ determined by ANOVA with Tukey *post-hoc* (day 1 $n = 3$, days 3–21 $n = 4$); data represent mean \pm s.e.m. (h) Population of haematopoietic-derived cells (CD45+) in time. (i) The loss in inflammation corresponds to an increase in vascularization, quantified by the number of visible capillaries in histological sections; for *, $p < 0.05$ as determined by ANOVA with Tukey *post-hoc* ($n = 3$); data represent mean \pm s.e.m. (j) An example tissue slice 10 days post injection shows an area of particularly high vascularization density (scale bar: 100 μm). Reprinted with permission from Roberts *et al.*¹⁷⁹ Copyright 2018 Springer Nature.



contrast, biomineralization processes can be enhanced and be directly applied to bone formation. Zhu *et al.* presented two biomolecules inspired by IDPs, denoted as P2 and P6, that helped the bone regeneration in 2D and 3D systems by increased biomineralization rates, cell attachment and proliferation. These rich-proline peptides were based on hydrophobic residues Leu, Met, Pro, and Val and polar Gln, His and Ser amino acids. The results showed how these amelogenin and ameloblastin hard tissue extracellular matrix protein imitations were more efficient than actual drugs such as Emdogain®.¹⁷⁸ Roberts *et al.* studied synthetic partially organized polymers (POPs) based on ELP IDRs (a Val-Pro-Gly-X-Gly pentapeptide) attached to helix polyalanine (Ala-Ala-Ala-Ala-Ala) regions for tissue recovery (Fig. 13). While ELPs alone formed micrometre-sized coalescing aggregates, leading to a colloidal suspension of liquid-like droplets, POPs underwent arrested phase separation into porous networks. Moreover, the lower size the disordered ELP region presented, the more fractal-like architecture they showed in PBS media. Depending on the helical percentage, the pore size could be tuned, going from *ca.* 30–50 μm pores (90% polyalanine) to *ca.* 3–5 μm pores (60%). *In vivo* mice studies showed how POPs rapidly and robustly were integrated into the sub-cutaneous space, creating mechanical connections with the surrounding tissues and finally promoting wound healing and tissue growth.¹⁷⁹ Recent studies by Chilkoti's group concluded how these POP structures self-assembled into fractal conformations. While using Val as a guest residue formed the already reported conformations, the use of Ala formed coacervate droplets with a physically crosslinked interconnected porous shell. The adjustment of the ELP/polyalanine ratio allowed the tuning of the porosity.¹⁸⁰

Conclusions

The self-assembly and functional features of IDPs/IDRs constitute a hot and exciting topic. The relationship between the protein structure and disorder with the biological function of IDPs/IDRs is also a rich research area. Eight amino acid residues have been identified as the main promoters for the disordered behaviour: Ala, Arg, Gln, Glu, Gly, Lys, Pro, and Ser. Advanced experimental characterization methods connecting the amino acid sequence with the resulting disordered structure are highly desirable. Computational models and force fields accounting for the unique properties of IDPs/IDRs are to be developed. Preliminary studies with IDPs/IDRs show their promising performance in different areas. Biological and biotechnological applications stand out as the forefront field. With detailed understanding of the nature of IDPs/IDRs, nanotechnology will be one step closer to replicate real complex biological media and apply self-assembled nanostructures in biology and biotechnology.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

P. G. A. acknowledges the Région Nouvelle Aquitaine, the Université de Bordeaux, Bordeaux INP and CNRS for financial support. Support from the Ministry of Science and Innovation of Spain through the MANA project CTQ2017-83961-R and JEANS project CTQ2017-92264-EXP is acknowledged. J. J. G.-C. acknowledges the Ministry of Science and Innovation of Spain for a "Ramon y Cajal" contract (RyC-2014-14956). The Andalusian Government (Consejería de Economía, Conocimiento, Empresas y Universidades, Junta de Andalucía) of Spain is acknowledged for the financial support through the UCO-1263193 Project.

Notes and references

- B. O. Okesola and A. Mata, *Chem. Soc. Rev.*, 2018, **47**, 3721–3736.
- A. Méndez-Ardoy, J. R. Granja and J. Montenegro, *Nanoscale Horiz.*, 2018, **3**, 391–396.
- Z. Gong, X. Liu, J. Dong, W. Zhang, Y. Jiang, J. Zhang, W. Feng, K. Chen and J. Bai, *Nanoscale*, 2019, **11**, 15479–15486.
- S. Li, Q. Zou, Y. Li, C. Yuan, R. Xing and X. Yan, *J. Am. Chem. Soc.*, 2018, **140**, 10794–10802.
- S. Li, R. Xing, R. Chang, Q. Zou and X. Yan, *Curr. Opin. Colloid Interface Sci.*, 2018, **35**, 17–25.
- D. Sanchez-deAlcazar, D. Romera, J. Castro-Smirnov, A. Sousaraei, S. Casado, A. Espasa, M. C. Morant-Miñana, J. J. Hernandez, I. Rodríguez, R. D. Costa, J. Cabanillas-Gonzalez, R. V. Martinez and A. L. Cortajarena, *Nanoscale Adv.*, 2019, **1**, 3980–3991.
- Z. Ye, X. Zhu, S. Acosta, D. Kumar, T. Sang and C. Aparicio, *Nanoscale*, 2019, **11**, 266–275.
- Z. Ye and C. Aparicio, *Nanoscale Adv.*, 2019, **1**, 4679–4682.
- M. Gontsarik, M. Mohammadtaheri, A. Yaghmur and S. Salentinig, *Biomater. Sci.*, 2018, **6**, 803–812.
- C. L. Hedegaard, E. C. Collin, C. Redondo-Gómez, L. T. H. Nguyen, K. W. Ng, A. A. Castrejón-Pita, J. R. Castrejón-Pita and A. Mata, *Adv. Funct. Mater.*, 2018, **28**, 1703716.
- Y. Cong, L. Ji, Y. Gao, F. Liu, D. Cheng, Z. Hu, Z. Qiao and H. Wang, *Angew. Chem., Int. Ed.*, 2019, **58**, 4632–4637.
- B. O. Okesola, S. Ni, B. Derkus, C. C. Galeano, A. Hasan, Y. Wu, J. Ramis, L. Buttery, J. I. Dawson, M. D'Este, R. O. C. Oreffo, D. Eglin, H. Sun and A. Mata, *Adv. Funct. Mater.*, 2020, **30**, 1906205.
- M. Cano and J. J. Giner-Casares, *Adv. Colloid Interface Sci.*, 2020, **286**, 102313.
- Y. Wu, B. O. Okesola, J. Xu, I. Korotkin, A. Berardo, I. Corridori, F. L. P. di Brocchetti, J. Kanczler, J. Feng, W. Li, Y. Shi, V. Farafonov, Y. Wang, R. F. Thompson, M.-M. Titirici, D. Nerukh, S. Karabasov, R. O. C. Oreffo, J. Carlos Rodriguez-Cabello, G. Vozzi, H. S. Azevedo, N. M. Pugno, W. Wang and A. Mata, *Nat. Commun.*, 2020, **11**, 1182.



- 15 M. Liutkus, A. López-Andarias, S. H. Mejías, J. López-Andarias, D. Gil-Carton, F. Feixas, S. Osuna, W. Matsuda, T. Sakurai, S. Seki, C. Atienza, N. Martín and A. L. Cortajarena, *Nanoscale*, 2020, **12**, 3614–3622.
- 16 B. A. Shoemaker, J. J. Portman and P. G. Wolynes, *Proc. Natl. Acad. Sci. U. S. A.*, 2000, **97**, 8868–8873.
- 17 C. González-Obeso, M. González-Pérez, J. F. Mano, M. Alonso and J. C. Rodríguez-Cabello, *Small*, 2020, 2005191.
- 18 H. Ruan, Q. Sun, W. Zhang, Y. Liu and L. Lai, *Drug Discovery Today*, 2019, **24**, 217–227.
- 19 I. Yruela and J. L. Neira, *Arch. Biochem. Biophys.*, 2020, **684**, 108328.
- 20 J. J. Ward, J. S. Sodhi, L. J. McGuffin, B. F. Buxton and D. T. Jones, *J. Mol. Biol.*, 2004, **337**, 635–645.
- 21 B. Xue, A. K. Dunker and V. N. Uversky, *J. Biomol. Struct. Dyn.*, 2012, **30**, 137–149.
- 22 C. Tanford and J. Reynolds, *Nature's Robots: A History of Proteins*, 2001.
- 23 International Union of Pure and Applied Chemistry, *Glossary of Class Names of Organic Compounds and Reactivity Intermediates Based on Structure*, IUPAC, 1995.
- 24 W. T. Miller, *Q. Rev. Biol.*, 1996, **71**, 563.
- 25 L. Pauling, R. B. Corey and H. R. Branson, *Proc. Natl. Acad. Sci. U. S. A.*, 1951, **37**, 205–211.
- 26 L. Pauling and R. B. Corey, *Proc. Natl. Acad. Sci. U. S. A.*, 1951, **37**, 729–740.
- 27 A. M. C. Marcelino and L. M. Gierasch, *Biopolymers*, 2008, **89**, 380–391.
- 28 J. S. Fetrow, *FASEB J.*, 1995, **9**, 708–717.
- 29 C. Tonlolo and E. Benedetti, *Trends Biochem. Sci.*, 1991, **16**, 350–353.
- 30 B. W. Low and H. J. Greenville-Wells, *Proc. Natl. Acad. Sci.*, 1953, **39**, 785–801.
- 31 G. M. Whitesides and B. Grzybowski, *Science*, 2002, **295**, 2418–2421.
- 32 J. J. Panda and V. S. Chauhan, *Polym. Chem.*, 2014, **5**, 4418–4436.
- 33 L. J. Smith, K. M. Fiebig, H. Schwalbe and C. M. Dobson, *Folding Des.*, 1996, **1**, 95–106.
- 34 P. Kumar and M. Bansal, *FEBS J.*, 2015, **282**, 4415–4432.
- 35 D. J. Barlow and J. M. Thornton, *J. Mol. Biol.*, 1988, **201**, 601–619.
- 36 F. Zsila, G. Kohut and T. Beke-Somfai, *Int. J. Biol. Macromol.*, 2019, **129**, 50–60.
- 37 F. Zsila, T. Juhász, S. Bősze, K. Horváti and T. Beke-Somfai, *Chirality*, 2018, **30**, 195–205.
- 38 D. Léon, M. P. Vermeuel, P. Gupta and M. R. Bunagan, *J. Pept. Sci.*, 2020, **26**, 1–7.
- 39 M. E. Fealey, B. P. Binder, V. N. Uversky, A. Hinderliter and D. D. Thomas, *Biophys. J.*, 2018, **114**, 550–561.
- 40 O. T. Johnson, T. Kaur and A. L. Garner, *ChemBioChem*, 2019, **20**, 40–45.
- 41 A. S. Saglam, D. W. Wang, M. C. Zwier and L. T. Chong, *J. Phys. Chem. B*, 2017, **121**, 10046–10054.
- 42 S. Jephthah, L. K. Månsson, D. Belić, J. P. Morth and M. Skepö, *Biomolecules*, 2020, **10**, 1–22.
- 43 S. Jephthah, J. Henriques, C. Cragnell, S. Puri, M. Edgerton and M. Skepö, *J. Chem. Inf. Model.*, 2017, **57**, 1330–1341.
- 44 F. R. Salemme, *Prog. Biophys. Mol. Biol.*, 1983, **42**, 95–133.
- 45 O. Coskuner and V. N. Uversky, *J. Chem. Inf. Model.*, 2017, **57**, 1342–1358.
- 46 T. Takekiyo, N. Yamada, T. Amo and Y. Yoshimura, *Pept. Sci.*, 2020, **112**, 1–8.
- 47 S. Boopathi, P. Dinh Quoc Huy, W. Gonzalez, P. E. Theodorakis and M. S. Li, *Proteins: Struct., Funct., Bioinf.*, 2020, **88**, 1285–1302.
- 48 Y. Lu, E. Zhang, J. Yang and Z. Cao, *Nano Res.*, 2018, **11**, 4985–4998.
- 49 C. Has and S. Pan, *J. Liposome Res.*, 2020, 1–22.
- 50 M. T. Ivanović, M. R. Hermann, M. Wójcik, J. Pérez and J. S. Hub, *J. Phys. Chem. Lett.*, 2020, **11**, 945–951.
- 51 A. Accardo, M. Leone, D. Tesauero, R. Aufiero, A. Bénarouche, J. F. Cavalier, S. Longhi, F. Carriere and F. Rossi, *Mol. Biosyst.*, 2013, **9**, 1401–1410.
- 52 S. H. Klass, M. J. Smith, T. A. Fiala, J. P. Lee, A. O. Omole, B. G. Han, K. H. Downing, S. Kumar and M. B. Francis, *J. Am. Chem. Soc.*, 2019, **141**, 4291–4299.
- 53 S. Acosta, Z. Ye, C. Aparicio, M. Alonso and J. C. Rodríguez-Cabello, *Biomacromolecules*, 2020, **21**, 4043–4052.
- 54 A. Rao, M. Drechsler, S. Schiller, M. Scheffner, D. Gebauer and H. Cölfen, *Adv. Funct. Mater.*, 2018, **28**, 1802063.
- 55 S. A. Costa, J. R. Simon, M. Amiram, L. Tang, S. Zauscher, E. M. Brustad, F. J. Isaacs and A. Chilkoti, *Adv. Mater.*, 2018, **30**, 1–9.
- 56 M. Fändrich, *Cell. Mol. Life Sci.*, 2007, **64**, 2066–2078.
- 57 M. Humenik, M. Magdeburg and T. Scheibel, *J. Struct. Biol.*, 2014, **186**, 431–437.
- 58 A. Hernik-Magoń, W. Puławski, B. Fedorczyk, D. Tymecka, A. Misicka, P. Szymczak and W. Dzwolak, *Biomacromolecules*, 2016, **17**, 1376–1382.
- 59 Y. Zhang, V. H. Man, C. Roland and C. Sagui, *ACS Chem. Neurosci.*, 2016, **7**, 576–587.
- 60 K. Pan and Q. Zhong, *Soft Matter*, 2015, **11**, 5898–5904.
- 61 M. Bakou, K. Hille, M. Kracklauer, A. Spanopoulou, C. V. Frost, E. Malideli, L. M. Yan, A. Caporale, M. Zacharias and A. Kapurniotu, *J. Biol. Chem.*, 2017, **292**, 14587–14602.
- 62 L. Larini, M. M. Gessel, N. E. Lapointe, T. D. Do, M. T. Bowers, S. C. Feinstein and J. E. Shea, *Phys. Chem. Chem. Phys.*, 2013, **15**, 8916–8928.
- 63 J. Adamcik, A. Sánchez-Ferrer, N. Ait-Bouziad, N. P. Reynolds, H. A. Lashuel and R. Mezzenga, *Angew. Chem., Int. Ed.*, 2016, **55**, 618–622.
- 64 C. Despres, J. Di, F. X. Cantrelle, Z. Li, I. Huvent, B. Chambraud, J. Zhao, J. Chen, S. Chen, G. Lippens, F. Zhang, R. Linhardt, C. Wang, F. G. Klärner, T. Schrader, I. Landrieu, G. Bitan and C. Smet-Nocca, *ACS Chem. Biol.*, 2019, **14**, 1363–1379.
- 65 R. Dec, M. Koliński, M. Kouza and W. Dzwolak, *Int. J. Biol. Macromol.*, 2020, **150**, 894–903.
- 66 A. J. Kuhn and J. Raskatov, *J. Alzheimer's Dis.*, 2020, **74**, 43–53.



- 67 A. J. Kuhn, B. S. Abrams, S. Knowlton and J. A. Raskatov, *ACS Chem. Neurosci.*, 2020, **11**, 1539–1544.
- 68 A. K. Jana, K. B. Batkulwar, M. J. Kulkarni and N. Sengupta, *Phys. Chem. Chem. Phys.*, 2016, **18**, 31446–31458.
- 69 B. Konarkowska, J. F. Aitken, J. Kistler, S. Zhang and G. J. S. Cooper, *FEBS J.*, 2006, **273**, 3614–3624.
- 70 R. P. R. Nanga, J. R. Brender, S. Vivekanandan and A. Ramamoorthy, *Biochim. Biophys. Acta, Biomembr.*, 2011, **1808**, 2337–2342.
- 71 S. Khatun, A. Singh, S. Maji, T. K. Maiti, N. Pawar and A. N. Gupta, *Soft Matter*, 2020, **16**, 3143–3153.
- 72 F. G. Quiroz, N. K. Li, S. Roberts, P. Weber, M. Dzuricky, I. Weitzhandler, Y. G. Yingling and A. Chilkoti, *Sci. Adv.*, 2019, **5**, 1–12.
- 73 D. Stehli, M. Mulaj, T. Miti, J. Traina, J. Foley and M. Muschol, *Intrinsically Disord. Proteins*, 2015, **3**, 1–12.
- 74 I. Bishof, E. B. Dammer, D. M. Duong, S. R. Kundinger, M. Gearing, J. J. Lah, A. I. Levey and N. T. Seyfried, *J. Biol. Chem.*, 2018, **293**, 11047–11066.
- 75 K. Dooley, B. Bulutoglu and S. Banta, *Biomacromolecules*, 2014, **15**, 3617–3624.
- 76 W. P. Aue, E. Bartholdi and R. R. Ernst, *J. Chem. Phys.*, 1976, **64**(5), 2229–2246.
- 77 K. Wüthrich, *Nat. Struct. Biol.*, 2001, **8**, 923–925.
- 78 G. M. Clore, *eMagRes*, 1996, 1–7.
- 79 A. M. Gronenborn and G. M. Clore, *Biochem. Pharmacol.*, 1990, **40**, 115–119.
- 80 B. Brutscher, I. C. Felli, S. Gil-Caballero, T. Hošek, K. Rainer, A. Piai, R. Pierattelli and Z. Sölyom, *Adv. Exp. Med. Biol.*, 2015, 49–122.
- 81 B. Chaves-Arquero, J. M. Pérez-Cañadillas and M. A. Jiménez, *Chem. - Eur. J.*, 2020, **26**, 5970–5981.
- 82 S. Kosol, A. Gallo, D. Griffiths, L. Valentic, T. R. Masschelein, J. Jenner, M. de los Santos, E. L. C. Manzi, P. K. Sydor, D. Rea, S. Zhou, V. Fülöp, N. J. Oldham, S.-C. Tsai, G. L. Challis and J. R. Lewandowski, *Nat. Chem.*, 2019, **11**, 913–923.
- 83 M. G. Murralli, A. Piai, W. Bermel, I. C. Felli and R. Pierattelli, *ChemBioChem*, 2018, **19**, 1625–1629.
- 84 S. Sukumaran, S. A. Malik, S. Sharma, K. Chandra and H. S. Atreya, *Chem. Commun.*, 2019, **55**, 7820–7823.
- 85 S. E. Reichheld, L. D. Muiznieks, R. Stahl, K. Simonetti, S. Sharpe and F. W. Keeley, *J. Biol. Chem.*, 2014, **289**, 10057–10068.
- 86 W. B. Garry, J. T. Barbara, J. Roberts, L. S. Malcolm and W. J. Shaw, *Biochim. Biophys. Acta*, 2010, **1804**, 1768–1774.
- 87 M. Beck Erlach, H. R. Kalbitzer, R. Winter and W. Kremer, *Biophys. Chem.*, 2019, **254**, 106239.
- 88 L. Hou, H. Shao, Y. Zhang, H. Li, N. K. Menon, E. B. Neuhaus, J. M. Brewer, I.-J. L. Byeon, D. G. Ray, M. P. Vitek, T. Iwashita, R. A. Makula, A. B. Przybyla and M. G. Zagorski, *J. Am. Chem. Soc.*, 2004, **126**, 1992–2005.
- 89 A. Fontana, P. P. De Laureto, B. Spolaore and E. Frare, *Identifying Disordered Regions in Proteins by Limited Proteolysis*, 2012, vol. 896.
- 90 L. B. Chemes, L. G. Alonso, M. G. Noval and G. De Prat-Gay, *Methods Mol. Biol.*, 2012, **895**, 387–404.
- 91 Y. Sun, A. Kakinen, Y. Xing, P. Faridi, A. Nandakumar, A. W. Purcell, T. P. Davis, P. C. Ke and F. Ding, *Small*, 2019, **15**, 1–10.
- 92 L. Y. Rivera-Najera, G. Saab-Rincón, M. Battaglia, C. Amero, N. O. Pulido, E. García-Hernández, R. M. Solórzano, J. L. Reyes and A. A. Covarrubias, *J. Biol. Chem.*, 2014, **289**, 31995–32009.
- 93 S. Weickert, J. Cattani and M. Drescher, *Electron Paramagn. Reson.*, 2019, **26**, 1–37.
- 94 N. L. Pirman, E. Milshteyn, L. Galiano, J. C. Hewlett and G. E. Fanucci, *Protein Sci.*, 2011, **20**, 150–159.
- 95 T. Bund, J. M. Boggs, G. Harauz, N. Hellmann and D. Hinderberger, *Biophys. J.*, 2010, **99**, 3020–3028.
- 96 R. Kaminker, I. Kaminker, W. R. Gutekunst, Y. Luo, S. Lee, J. Niu, S. Han and C. J. Hawker, *Chem. Commun.*, 2018, **54**, 5237–5240.
- 97 O. A. Chinak, A. V. Shernyukov, S. S. Ovcherenko, E. A. Sviridov, V. M. Golyshev, A. S. Fomin, I. A. Pyshnaya, E. V. Kuligina, V. A. Richter and E. G. Bagryanskaya, *Molecules*, 2019, **24**, 2919.
- 98 E. A. Permyakov and V. N. Uversky, in *Instrumental Analysis of Intrinsically Disordered Proteins: Assessing Structure and Conformation*, 2010, pp. 323–344.
- 99 S. Acharya, B. M. Safaie, P. Wongkongkathep, M. I. Ivanova, A. Attar, F. G. Klärner, T. Schrader, J. A. Loo, G. Bitan and L. J. Lapidus, *J. Biol. Chem.*, 2014, **289**, 10727–10737.
- 100 F. Zsila, T. Juhász, S. Bösze, K. Horváti and T. Beke-Somfai, *Chirality*, 2018, **30**, 195–205.
- 101 F. Zhu, N. W. Isaacs, L. Hecht and L. D. Barron, *Structure*, 2005, **13**, 1409–1419.
- 102 S. Signorelli, S. Cannistraro and A. R. Bizzarri, *Appl. Spectrosc.*, 2017, **71**, 823–832.
- 103 T. G. McCaslin, C. V. Pagba, J. Yohannan and B. A. Barry, *Sci. Rep.*, 2019, **9**, 1–14.
- 104 A. Rawat, B. K. Maity, B. Chandra and S. Maiti, *Biochim. Biophys. Acta, Biomembr.*, 2018, **1860**, 1734–1740.
- 105 C. La Rosa, M. Condorelli, G. Compagnini, F. Lolicato, D. Milardi, T. N. Do, M. Karttunen, M. Pannuzzo, A. Ramamoorthy, F. Fraternali, F. Collu, H. Rezaei, B. Strodel and A. Raudino, *Eur. Biophys. J.*, 2020, **49**, 175–191.
- 106 A. Natalello, D. Ami and S. M. Doglia, *Methods Mol. Biol.*, 2012, **895**, 229–244.
- 107 S. Koubaa, A. Bremer, D. K. Hinch and F. Brini, *Sci. Rep.*, 2019, **9**, 1–11.
- 108 M. A. Fallah, H. R. Gerding, C. Scheibe, M. Drescher, C. Karreman, S. Schildknecht, M. Leist and K. Hauser, *ChemBioChem*, 2017, **18**, 2312–2316.
- 109 E. Villarreal-Ramirez, D. Eliezer, R. Garduño-Juarez, A. Gericke, J. M. Perez-Aguilar and A. Boskey, *Bone*, 2017, **95**, 65–75.
- 110 M. Vitali, V. Rigamonti, A. Natalello, B. Colzani, S. Avvakumova, S. Brocca, C. Santambrogio, J. Narkiewicz, G. Legname, M. Colombo, D. Prospero and R. Grandori, *Biochim. Biophys. Acta, Gen. Subj.*, 2018, **1862**, 1556–1564.
- 111 P. Bernadó and D. I. Svergun, *Mol. BioSyst.*, 2012, **8**, 151–167.



- 112 S. Lenton, M. Grimaldo, F. Roosen-Runge, F. Schreiber, T. Nylander, R. Clegg, C. Holt, M. Härtle, V. García Sakai, T. Seydel and S. C. Marujo Teixeira, *Biophys. J.*, 2017, **112**, 1586–1596.
- 113 D. Didry, F. X. Cantrelle, C. Husson, P. Roblin, A. M. E. Moorthy, J. Perez, C. Le Clainche, M. Hertzog, E. Guittet, M. F. Carlier, C. Van Heijenoort and L. Renault, *EMBO J.*, 2012, **31**, 1000–1013.
- 114 C. Cragnell, L. Staby, S. Lenton, B. B. Kragelund and M. Skepö, *Biomolecules*, 2019, **9**, 168.
- 115 P. Hardouin, C. Velours, C. Bou-Nader, N. Assrir, S. Laalami, H. Putzer, D. Durand and B. Golinelli-Pimpaneau, *Biophys. J.*, 2018, **115**, 2102–2113.
- 116 K. Gast and C. Fiedler, *Methods Mol. Biol.*, 2012, **896**, 137–161.
- 117 F. Zsila, S. Bösze, K. Horváti, I. C. Szegvártó and T. Beke-Somfai, *RSC Adv.*, 2017, **7**, 41091–41097.
- 118 K. Shou, A. Bremer, T. Rindfleisch, P. Knox-Brown, M. Hirai, A. Rekas, C. J. Garvey, D. K. Hinch, A. M. Stadler and A. Thalhammer, *Phys. Chem. Chem. Phys.*, 2019, **21**, 18727–18740.
- 119 M. Levitt and V. Daggett, *Nat. Struct. Biol.*, 2001, **8**, 662.
- 120 P. K. Weiner and P. A. Kollman, *J. Comput. Chem.*, 1981, **2**, 287–303.
- 121 C. I. Bayly, K. M. Merz, D. M. Ferguson, W. D. Cornell, T. Fox, J. W. Caldwell, P. A. Kollman, P. Cieplak, I. R. Gould and D. C. Spellmeyer, *J. Am. Chem. Soc.*, 1995, **117**, 5179–5197.
- 122 D. A. Case, T. E. Cheatham, T. Darden, H. Gohlke, R. Luo, K. M. Merz, A. Onufriev, C. Simmerling, B. Wang and R. J. Woods, *J. Comput. Chem.*, 2005, **26**, 1668–1688.
- 123 W. Wang, W. Ye, C. Jiang, R. Luo and H. F. Chen, *Chem. Biol. Drug Des.*, 2014, **84**, 253–269.
- 124 D. Song, R. Luo and H. F. Chen, *J. Chem. Inf. Model.*, 2017, **57**, 1166–1178.
- 125 D. Song, H. Liu, R. Luo and H. F. Chen, *J. Chem. Inf. Model.*, 2020, **60**, 2257–2267.
- 126 P. Robustelli, S. Piana and D. E. Shaw, *Proc. Natl. Acad. Sci. U. S. A.*, 2018, **115**, E4758–E4766.
- 127 R. B. Best, W. Zheng and J. Mittal, *J. Chem. Theory Comput.*, 2014, **10**, 5113–5124.
- 128 L. Yu, D. W. Li and R. Brüschweiler, *J. Chem. Theory Comput.*, 2020, **16**, 1311–1318.
- 129 J. Henriques and M. Skepö, *J. Chem. Theory Comput.*, 2016, **12**, 3407–3415.
- 130 L. M. Pietrek, L. S. Stelzl and G. Hummer, *J. Chem. Theory Comput.*, 2020, **16**, 725–737.
- 131 E. Rieloff and M. Skepö, *J. Chem. Theory Comput.*, 2020, **16**, 1924–1935.
- 132 J. A. Joseph and D. J. Wales, *J. Phys. Chem. B*, 2018, **122**, 11906–11921.
- 133 Y. Ouyang, L. Zhao and Z. Zhang, *Phys. Chem. Chem. Phys.*, 2018, **20**, 8676–8684.
- 134 X. Liu and J. Chen, *J. Chem. Theory Comput.*, 2019, **15**, 4708–4720.
- 135 A. Kuzmanic, R. B. Pritchard, D. F. Hansen and F. L. Gervasio, *J. Phys. Chem. Lett.*, 2019, **10**, 1928–1934.
- 136 V. T. Duong, Z. Chen, M. T. Thapa and R. Luo, *J. Phys. Chem. B*, 2018, **122**, 10455–10469.
- 137 M. Carballo-Pacheco, A. E. Ismail and B. Strodel, *J. Chem. Theory Comput.*, 2018, **14**, 6063–6075.
- 138 M. Karplus, *Annu. Rev. Biophys. Biomol. Struct.*, 2006, **35**, 1–47.
- 139 A. D. MacKerell, D. Bashford, M. Bellott, R. L. Dunbrack, J. D. Evanseck, M. J. Field, S. Fischer, J. Gao, H. Guo, S. Ha, D. Joseph-McCarthy, L. Kuchnir, K. Kuczera, F. T. K. Lau, C. Mattos, S. Michnick, T. Ngo, D. T. Nguyen, B. Prodhom, W. E. Reiher, B. Roux, M. Schlenkrich, J. C. Smith, R. Stote, J. Straub, M. Watanabe, J. Wiórkiewicz-Kuczera, D. Yin and M. Karplus, *J. Phys. Chem. B*, 1998, **102**, 3586–3616.
- 140 B. R. Brooks, C. L. Brooks, A. D. Mackerell, L. Nilsson, R. J. Petrella, B. Roux, Y. Won, G. Archontis, C. Bartels, S. Boresch, A. Caflisch, L. Caves, Q. Cui, A. R. Dinner, M. Feig, S. Fischer, J. Gao, M. Hodoscek, W. Im, K. Kuczera, T. Lazaridis, J. Ma, V. Ovchinnikov, E. Paci, R. W. Pastor, C. B. Post, J. Z. Pu, M. Schaefer, B. Tidor, R. M. Venable, H. L. Woodcock, X. Wu, W. Yang, D. M. York and M. Karplus, *J. Comput. Chem.*, 2009, **30**, 1545–1614.
- 141 R. B. Best, X. Zhu, J. Shim, P. E. M. Lopes, J. Mittal, M. Feig and A. D. MacKerell, *J. Chem. Theory Comput.*, 2012, **8**, 3257–3273.
- 142 J. Huang, S. Rauscher, G. Nawrocki, T. Ran, M. Feig, B. L. De Groot, H. Grubmüller and A. D. MacKerell, *Nat. Methods*, 2016, **14**, 71–73.
- 143 H. Liu, D. Song, Y. Zhang, S. Yang, R. Luo and H. F. Chen, *Phys. Chem. Chem. Phys.*, 2019, **21**, 21918–21931.
- 144 T. Lazar, M. Guharoy, W. Vranken, S. Rauscher, S. J. Wodak and P. Tompa, *Biophys. J.*, 2020, **118**, 2952–2965.
- 145 S. Rauscher, V. Gapsys, M. J. Gajda, M. Zweckstetter, B. L. De Groot and H. Grubmüller, *J. Chem. Theory Comput.*, 2015, **11**, 5513–5524.
- 146 M. Carballo-Pacheco and B. Strodel, *Protein Sci.*, 2017, **26**, 174–185.
- 147 P. Krupa, P. D. Quoc Huy and M. S. Li, *J. Chem. Phys.*, 2019, **151**, 055101.
- 148 V. H. Man, X. He, P. Derreumaux, B. Ji, X. Q. Xie, P. H. Nguyen and J. Wang, *J. Chem. Theory Comput.*, 2019, **15**, 1440–1452.
- 149 C. R. Watts, A. Gregory, C. Frisbie and S. Lovas, *Proteins: Struct., Funct., Bioinf.*, 2018, **86**, 279–300.
- 150 A. K. Somavarapu and K. P. Kepp, *ChemPhysChem*, 2015, **16**, 3278–3289.
- 151 W. F. v. Gunsteren, *Biomolecular Simulation: GROMOS 96 Manual and User Guide*, 1996, vol. 44.
- 152 W. R. P. Scott, P. H. Hünenberger, I. G. Tironi, A. E. Mark, S. R. Billeter, J. Fennen, A. E. Torda, T. Huber, P. Krüger and W. F. Van Gunsteren, *J. Phys. Chem. A*, 1999, **103**, 3596–3607.
- 153 M. Christen, P. H. Hünenberger, D. Bakowies, R. Baron, R. Bürgi, D. P. Geerke, T. N. Heinz, M. A. Kastenholz, V. Kräutler, C. Oostenbrink, C. Peter, D. Trzesniak and



- W. F. Van Gunsteren, *J. Comput. Chem.*, 2005, **26**, 1719–1751.
- 154 C. Oostenbrink, A. Villa, A. E. Mark and W. F. Van Gunsteren, *J. Comput. Chem.*, 2004, **25**, 1656–1676.
- 155 N. Schmid, A. P. Eichenberger, A. Choutko, S. Riniker, M. Winger, A. E. Mark and W. F. Van Gunsteren, *Eur. Biophys. J.*, 2011, **40**, 843–856.
- 156 S. R. Gerben, J. A. Lemkul, A. M. Brown and D. R. Bevan, *J. Biomol. Struct. Dyn.*, 2014, **32**, 1817–1832.
- 157 A. Bandyopadhyay and S. Basu, *Biochim. Biophys. Acta, Proteins Proteomics*, 2020, **1868**, 140474.
- 158 W. L. Jorgensen, D. S. Maxwell and J. Tirado-Rives, *J. Am. Chem. Soc.*, 1996, **118**, 11225–11236.
- 159 F. Jiang, C. Y. Zhou and Y. D. Wu, *J. Phys. Chem. B*, 2014, **118**, 6983–6998.
- 160 S. Xun, F. Jiang and Y. D. Wu, *Bioorg. Med. Chem.*, 2016, **24**, 4970–4977.
- 161 G. A. Kaminski, R. A. Friesner, J. Tirado-Rives and W. L. Jorgensen, *J. Phys. Chem. B*, 2001, **105**, 6474–6487.
- 162 M. J. Robertson, J. Tirado-Rives and W. L. Jorgensen, *J. Chem. Theory Comput.*, 2015, **11**, 3499–3509.
- 163 S. Yang, H. Liu, Y. Zhang, H. Lu and H. Chen, *J. Chem. Inf. Model.*, 2019, **59**, 4793–4805.
- 164 M. D. Smith, J. S. Rao, E. Segelken and L. Cruz, *J. Chem. Inf. Model.*, 2015, **55**, 2587–2595.
- 165 V. H. Man, P. H. Nguyen and P. Derreumaux, *J. Phys. Chem. B*, 2017, **121**, 5977–5987.
- 166 S.-H. Chong, P. Chatterjee and S. Ham, *Annu. Rev. Phys. Chem.*, 2017, **68**, 117–134.
- 167 M. Dzuricky, B. A. Rogers, A. Shahid, P. S. Cremer and A. Chilkoti, *Nat. Chem.*, 2020, **12**, 814–825.
- 168 A. Savastano, D. Flores, H. Kadavath, J. Biernat, E. Mandelkow and M. Zweckstetter, *Angew. Chem., Int. Ed.*, 2021, **60**, 726–730.
- 169 C. M. Metrick, A. L. Koenigsberg and E. E. Heldwein, *mBio*, 2020, **11**, 1–22.
- 170 P. Dogra, A. Joshi, A. Majumdar and S. Mukhopadhyay, *J. Am. Chem. Soc.*, 2019, **141**, 20380–20389.
- 171 G. L. Dignon, W. Zheng, Y. C. Kim and J. Mittal, *ACS Cent. Sci.*, 2019, **5**, 821–830.
- 172 H. Zhao, V. Ibrahimova, E. Garanger and S. Lecommandoux, *Angew. Chem., Int. Ed.*, 2020, **59**, 11028–11036.
- 173 L. Faltova, A. M. Küffner, M. Hondele, K. Weis and P. Arosio, *ACS Nano*, 2018, **12**, 9991–9999.
- 174 I. Urosev, J. Lopez Morales and M. A. Nash, *Adv. Funct. Mater.*, 2020, **30**, 2005245.
- 175 M. S. Hossain, C. Maller, Y. Dai, S. Nangia and D. Mozhdzhi, *Chem. Commun.*, 2020, **56**, 10281–10284.
- 176 W. R. Wonderly, T. R. Cristiani, K. C. Cunha, G. D. Degen, J. E. Shea and J. H. Waite, *Macromolecules*, 2020, **53**, 6767–6779.
- 177 B. Bulutoglu, S. J. Yang and S. Banta, *Biomacromolecules*, 2017, **18**, 2139–2145.
- 178 H. Zhu, M. Gomez, J. Xiao, G. Perale, F. Betge, S. P. Lyngstadaas and H. J. Haugen, *ACS Appl. Bio Mater.*, 2020, **3**, 2263–2274.
- 179 S. Roberts, T. S. Harmon, J. L. Schaal, V. Miao, K. Li, A. Hunt, Y. Wen, T. G. Oas, J. H. Collier, R. V. Pappu and A. Chilkoti, *Nat. Mater.*, 2018, **17**, 1154–1163.
- 180 S. Roberts, V. Miao, S. Costa, J. Simon, G. Kelly, T. Shah, S. Zauscher and A. Chilkoti, *Nat. Commun.*, 2020, **11**, 1.

