sequence to morph into differently shaped binding sites. These results indicate that the observations made previously on the single examples of many-to-one and one-to-many signaling generalize to many hundreds of other well characterized protein-protein interactions.

1337-Pos
Energy Landscape Analysis Reveals Residual Order in Histone Tail Dynamics
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Histone tails are highly flexible N terminal protrusions of histone proteins which help to fold DNA into dense superstructures known as chromatin. On a molecular scale histone tails are polyelectrolites with high degree of conformational disorder, allowing them to function as bio-molecular ''switches", regulating various genetic regulatory processes. Because of being intrinsically disordered, the structural and dynamical aspects of histone tails are still poorly understood. In this work we have carried out 3 microsecond all atom replica exchange molecular dynamics (REMD) simulations of four histone tails, H4, H3, H2B and H2A, to probe for their intrinsic conformational preferences. Our subsequent energy landscape analysis demonstrated that some tails are not fully disordered, but contain residual secondary structure elements. In particular, H4 formed beta hairpins, H3 and H2B adopted alpha helical elements while H2A was fully disordered. We also carried out polymer physics based analysis of the histone tails' conformational ensembles. We found an intriguing re-entrant contraction-expansion of the tails upon heating, which is caused by the way mobile counterions associate with the protein chains at various temperatures.

1338-Pos
Connecting Unfolded Protein Dynamics and Aggregation
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Solving the ''protein folding problem'' will require a better understanding of the allowed conformations and dynamics of the unfolded state. Fundamentally, unfolded protein dynamics should be determined by the hydrophobic pattern of each sequence. As proof of this hypothesis, we have constructed a random wormlike chain model that is re-weighted to favor random conformations that have residues of similar hydrophobicity in close proximity. This model gives remarkable quantitative agreement with measurements of intramolecular contact formation for various sequences. Furthermore we have found that the intramolecular diffusion coefficient of various sequences in folding conditions vary by ~ 3 orders of magnitude. We find such dynamics qualitatively agree with those predicted by molecular dynamics in implicit solvent. The fastest sequences are intrinsically disordered proteins and peptides and the slowest sequences are well-behaved proteins with folding times of at least 1 ms. In between these two regimes are sequences prone to aggregation in which intramolecular diffusion is just fast enough to expose hydrophobes to solvent long enough to form bimolecular interactions.

1339-Pos
Investigation of the Intrinsically Disordered Protein IA3 by Multiple SDSL-EPR Techniques
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Intrinsically disordered proteins (IDPs) are proteins that contain little to no secondary or tertiary structure and are often functional proteins that are essential in biological systems. Many IDPs undergo a conformational change where the lack of intrinsic structure is relieved upon binding to its target protein. Due to the very nature of unstructured proteins, characterization of the conformational propensities and function of these proteins present a major challenge. Site-directed spin labeling (SDSL) coupled with electron paramagnetic resonance (EPR) spectroscopy is a valuable tool in characterizing the mobility and conformational changes of proteins. This combined technique, however, is not often used to investigate intrinsically disordered proteins (IDPs). IA3 is a 68 residue IDP that has been extensively characterized by various biophysical techniques and was used in this study as a model system to show SDSL-EPR may be employed to characterize conformational changes in IDPs. The TFE-induced disordered-to-α-helical transition of IA3 was monitored by various SDSL-EPR techniques. CW-EPR experiments were performed at X-, Q-, and W-band resonant frequencies and reveal conformational changes can be observed at all three frequencies, with the W-band spectra revealing the most striking changes in dynamics upon inducing α -helical secondary structure by TFE. Low temperature CW-EPR and DEER distance measurements were also performed to evaluate the ability of these methods in characterizing the induced α-helical conformation.

1340-Pos
Coupled Folding and Binding of pKID with KIX Domain Investigated by Multicanonical Molecular Dynamics Simulation in Explicit Solvent
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Intrinsically disordered proteins (IDPs) do not form any rigid tertiary structures alone. Most of them bind corresponding proteins and fold into an ordered structure to play important roles in biological functions such as signal transduction cascades. The phosphorylated kinase induced domain (pKID) is one of IDPs. The pKID adapts an ordered helical structure while binds to the KIX domain. This structural property of pKID is called ''coupled folding and binding''. From nuclear magnetic resonance (NMR) studies a kinetics model of the coupled folding and binding has been proposed: there are four states of pKID (disordered, encounter complex, intermediate, and ordered helical structure). However, the details of these states at atomic level were still unclear.
In order to obtain the free energy landscapes and the stable complex structures at various temperatures for the system, where pKID, the KIX domain, water and ions are included and they can interact with each other, multicanonical molecular dynamics (McMD) simulation was performed with the all-atom model in explicit solvent. McMD simulation is one of generalized ensemble methods, which can search conformational space much wider than a conventional molecular dynamics simulation, as well as the replica-exchange molecular dynamics (REMD). In this study, starting from completely disordered and unbound states of pKID, the helices of pKID were reproduced as approaching to the KIX domain at adequate low temperature. Additionally, we found a different binding mode from the NMR model. Interestingly, this binding mode is similar to that of the activation domain of the mixed lineage leukemia (MLL) transcription factor upon binding to the KIX domain.

1341-Pos
On the Origins of Intrinsically Disordered Proteins
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A large number of proteins are sufficiently unstable that their full three dimensional structure cannot be resolved. The origins of this intrinsic disorder are not well understood, but its ubiquitous presence undercuts the principle that a protein's structure determines its function. Here, we present a quantitative theory that makes novel predictions regarding the role of intrinsic disorder in protein structure and function. In particular, we discuss the implications of analytical solutions of a series of fundamental thermodynamic models of protein interactions in which disordered proteins are characterized by positive folding free energies. We validate our predictions by assigning protein function using the Gene Ontology classification, in which Protein Binding, Catalytic Activity and Transcription Regulator Activity are the three largest functional categories, and performing genome-wide surveys of both the amount of disorder in these functional classes and binding affinities for both prokaryotic and eukaryotic genomes. Specifically, without assuming any a priori structure-function relationship, the theory predicts that both Catalytic and low-affinity Binding ($K_d > 10^{-7}$ M) proteins prefer ordered structures, while only high-affinity Binding proteins (found mostly in eukaryotes) can tolerate disorder. Relevant to both Transcription and signal transduction, the theory also explains how increasing disorder can tune the binding affinity to maximize the specificity of promiscuous interactions. Collectively, these studies provide insight into how natural selection acts on folding stability to optimize protein function.

1342-Pos
Concerted Involvement of Long-Range Electrostatic Interactions and Fly-Catsing in Recognition of Intrinsically Disordered Proteins
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Intrinsically disordered proteins (IDPs) are the functional proteins that exist as dynamical ensemble under physiological conditions. IDPs play crucial roles in cellular signaling and regulation, and often fold upon binding to specific targets. In particular, the ability to interact with multiple targets appears to be a hallmark of IDPs. Two ideal mechanisms, namely fly-casting/induced folding and conformational selection, are possible for coupled folding and binding to the specific targets. Evidence recently accumulated to suggest that induced folding might be prevalent in IDP recognition. Our recent analysis reveals that both IDPs and the vicinity of their binding sites on the substrate surface are enriched with charges. Our hypothesis is the long-range electrostatic interactions between these charged residues triggers the promotion of the unfolding of residual structure in unbound IDPs and enhance the binding efficiency via the fly-casting effects. Go-like coarse-grained protein model has been used to investigate the interaction mechanisms of IDP complex pKID:KIX. Initial results appear to support the concerted involvement of long-range electrostatic interactions and fly-casting.