Disordered nucleome: Abundance of intrinsic disorder in the DNA- and RNA-binding proteins in 1121 species from Eukaryota, Bacteria and Archaea

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Intrinsically disordered proteins (IDPs) are abundant in various proteomes, where they play numerous important roles and complement biological activities of ordered proteins. Among functions assigned to IDPs are interactions with nucleic acids. However, often, such assignments are made based on the guilty-by-association principle. The validity of the extension of these correlations to all nucleic acid binding proteins has never been analyzed on a large scale across all domains of life. To fill this gap, we perform a comprehensive computational analysis of the abundance of intrinsic disorder and intrinsically disordered domains in nucleomes (~548,000 nucleic acid binding proteins) of 1121 species from Archaea, Bacteria and Eukaryota. Nucleome is a whole complement of proteins involved in interactions with nucleic acids. We show that relative to other proteins in the corresponding proteomes, the DNA-binding proteins have significantly increased disorder content and are significantly enriched in disordered domains in Eukaryotes but not in Archaea and Bacteria. The RNA-binding proteins are significantly enriched in the disordered domains in Bacteria, Archaea and Eukaryota, while the overall abundance of disorder in these proteins is significantly increased in Bacteria, Archaea, animals and fungi. The high abundance of disorder in nucleomes supports the notion that the nucleic acid binding proteins often require intrinsic disorder for their functions and regulation.

Keywords:
DNA-binding proteins / Intrinsically disorder / Intrinsically disordered proteins / Nucleome / RNA-binding proteins / Systems biology

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1 Introduction

A significant portion of any given proteome is ascribed to biologically active proteins that do not have unique 3D structures as a whole or in part [1–6]. The dynamic conformational ensembles of such intrinsically disordered proteins...
Significance of the study

Intrinsically disordered proteins and protein regions are abundant across all domains of life and their functions complement the functional repertoire of structured (ordered) proteins. In spite of the functional importance of protein-nucleic acids interactions and the fact that intrinsically disordered proteins were shown to be very common in some classes of RNA- and DNA-binding proteins, the question on how abundant the disorder is in the nucleiome (whole complement of proteins involved in interaction with nucleic acids) remains open. We report a first-of-its-kind comprehensive analysis of the abundance of intrinsic disorder and intrinsically disordered domains in nucleiomes of over 1100 species from Archaea, Bacteria and Eukaryota. We investigate intrinsic disorder from multiple perspectives that include enrichment in the overall amount of disorder and the amount of disordered domains, relation to the nucleic acid binding and enrichment in a comprehensive set of functional classes of RNA- and DNA-binding proteins. Our results provide a strong support to the notion that the nucleic acid binding proteins often require intrinsic disorder for their functions and regulation.

2 Materials and methods

2.1 Datasets and annotation of DNA- and RNA-binding proteins

We collect 1674 complete proteomes and the corresponding 20 619 215 proteins from the release 2015_08 of UniProt [31]. We remove viruses and categorized the remaining proteomes into the Archaea, Bacteria and Eukaryota domains of life. To reduce bias towards certain overpopulated in the UniProt species and to reduce redundancy, we aggregate all the taxonomic identifiers to the second last level of the taxonomy, which most commonly is the genus. We select one largest proteome from each group of these identifiers, consequently sampling species at the genus level. We annotate DNA- and RNA-binding proteins in these proteomes based on the annotations from the Gene Ontology (GO) database [32] that are linked in the UniProt. We include proteins that have molecular function specified as “DNA binding” or “RNA binding”, and the direct child terms of the “DNA binding” and “RNA binding” terms in the gene ontology network. To assure that the number of DNA- and RNA-binding proteins is sufficiently large to perform statistical analysis, we remove the proteomes...
where the total number of DNA-binding proteins and RNA-binding proteins is below 20. The resulting dataset includes 7,912,445 proteins from 1121 complete proteomes (species). The number of species and proteins in each domain of life and in the eukaryotic kingdoms that are included in our dataset, and the number and fraction of the DNA- and RNA-binding proteins aggregated into the domains and kingdoms of life are shown in Supporting Information Table S1. To summarize, the nucleome analyzed in our study includes 548,091 DNA- and RNA-binding proteins. The complete list of considered species is given in Supporting Information Table S2.

The annotations of the DNA- and RNA-binding proteins that we use include both manually annotated proteins and proteins that are annotated automatically in UniProt using sequence similarity to the manually annotated entries. The coverage drops dramatically if we only use the manually annotated entries. Out of the considered 1121 proteomes only 1% of eukaryotic species has over 50% of its proteins manually annotated while such level of coverage is not even present in Archaea and Bacteria (Supporting Information Fig. S1A). Moreover, only 6, 3 and 2% of the species in Archaea, Eukaryota and Bacteria, respectively, have over 20% of their proteins reviewed. Similar lack of reviewed entries is true for the proteins annotated as DNA- and RNA-binding. About 30, 15 and 5% of species in Archaea, Bacteria and Eukaryota, respectively, have at least 20% of their all DNA- and RNA-binding proteins annotated manually and these numbers drop to 6, 1 and 1% when we require at least 50% coverage (Supporting Information Fig. S1B). More importantly, our analysis shown in the Results section reveals that results based on a limited set of the reviewed proteomes and proteins are very similar to the results on the complete set of all proteomes and proteins.

2.2 Annotation and computational characterization of intrinsic disorder

Given the large scale of our analysis, we perform the annotation of disordered regions with high-throughput predictors. We utilize a consensus of five such predictors including two versions of IUPred [33] designed to find long and short IDR s and three versions of Espritz [34] that predict intrinsic disorder annotated based on structures solved via X-ray crystallography, the NMR-derived structures and the Disprot database [35]. These methods were shown to provide good predictive performance in a recent large-scale assessment with AUC values around 0.77 [36]. We use majority vote consensus where a given residue is predicted as disordered if majority of the methods (three or more out of the five methods) predict it as disordered; otherwise the residue is predicted as structured. The motivation to use of the consensus comes from empirical observations which reveal that this leads to further increase in the predictive performance when compared to the use of individual predictors [36–38]. The same consensus was recently used in related works [6, 27, 39]. The predictions were filtered by removing disordered segments with less than four consecutive residues, which is in agreement with other studies [6, 27, 40]. We note that such annotations of disordered regions can be collected from two databases: MobiDB [41,42] and DIANA [43].

We use the putative disorder to compute disorder content (DC) per protein, which is defined as the fraction of disordered residues in the given protein chain. The DC for a proteome is the average of the DC values of its proteins. We calculate relative disorder content (RDC), which quantifies the DC in the DNA- and RNA-binding proteins relative to the overall DC in their corresponding proteome. RDC is calculated per protein as:

$$RDC_{protein} = \frac{\text{sign} \left( DC_{bind} - DC_{proteome} \right)}{\min\left( DC_{bind}, DC_{proteome} \right) - 1} \times 100\%$$

where $DC_{bind}$ is the DC of a given DNA- or RNA-binding protein; $DC_{proteome}$ is the DC of the corresponding proteome; and sign is the sign of the difference between $DC_{bind}$ and $DC_{proteome}$ where the value is $+(-)$ when $DC_{bind} > DC_{proteome}$ ($DC_{bind} < DC_{proteome}$) and $RDC_{protein} = 0$ if $DC_{bind} = DC_{proteome}$.

We also compute RDC per proteome as:

$$RDC_{proteome} = \frac{\text{sign} \left( \text{ave}DC_{bind} - DC_{proteome} \right)}{\min\left( \text{ave}DC_{bind}, DC_{proteome} \right) - 1} \times 100\%$$

where $\text{ave}DC_{bind}$ is the average DC of all DNA- or all RNA-binding proteins in a given proteome.

The $RDC_{protein}$ ($RDC_{proteome}$) values are defined as a ratio between DC of the binding chain (or a set of binding chains) and its (their) proteome and it quantifies the amount of enrichment (if value is positive) or depletion (if negative) of the disorder in the binding protein(s) relative to the content of the proteome. The values of RDC vary between negative and positive infinity and have intuitive interpretation, e.g., given $DC_{bind} = 0.5$ and $DC_{proteome} = 0.25$, $RDC_{protein} = 100\%$ which shows that the DC of the protein is $100\%$ higher than the DC of its proteome; given $DC_{bind} = 0.25$ and $DC_{proteome} = 0.5$, $RDC_{protein} = -100\%$ which means that the DC of the protein is $100\%$ lower than that of its proteome.

We measure correlation between the average DC in proteome and the average DC in its DNA- or RNA-binding proteins over a given domain or kingdom of life to investigate whether enrichment of depletion of disorder in the binding proteins is consistent. The correlation is quantified with the Pearson correlation coefficient (PCC):

$$PCC = \frac{\text{covariance}(\text{ave}DC_{bind}, DC_{proteome})}{\sigma_{\text{ave}DC_{bind}} \cdot \sigma_{DC_{proteome}}}$$

where $\text{ave}DC_{bind}, DC_{proteome}$ are computed over proteomes in a given domain or kingdom and $\sigma$ is the standard deviation.
We annotate long disordered regions (LDRs) which are defined as having 30 or more consecutive disordered residues [3, 44-46]. These regions are recognized as a distinct class of biologically functional IDP domains [3, 47, 48], which means that proteins with LDRs are likely to carry functions through disorder. We use these annotations to compute fraction of proteins with LDRs in DNA- or RNA-binding proteins in a given proteome, in entire proteomes and in domains and kingdoms of life. The latter results are summarized in Supporting Information Table S1.

We compute mean net charge (MNC) of proteins and LDRs, which can be approximated as [4, 49]:

\[
\text{MNC} = \frac{[(N_R + N_K) - (N_E + N_D)]}{\text{length}}
\]

where \(N_R\), \(N_K\), \(N_E\) and \(N_D\) are the counts of arginine, lysine, glutamic acid and aspartic acid residues and length is the length of a given protein sequence of a protein or LDR.

### 2.3 Evaluation of the annotations of disorder

We compare the consensus of the five methods that is utilized in this study with two other consensuses to demonstrate that our results are not affected by a potential noise coming from the use of different types of annotations of disorder by various prediction methods. The first alternative consensus is based on the ten methods that are included in MobiDB database [41, 42]: two versions of IUPred [33], three versions of Esprit [34], two flavors of DisEMBL [50], GlobPlot [51], VSL2b [52] and jRONN [53]). The second consensus includes these ten methods and another recently published method: DynaMine [54, 55], totalling in 11 predictors. We compare the predictions of the three consensuses on two popular model genera: *Escherichia* and *Drosophila* (Supporting Information Fig. S2). The results show that the three consensuses generate similar distributions of DC values for these genera, which suggests that our putative annotations of disorder are not affected by the selection of methods in the consensus.

Moreover, we also test predictive quality of these consensuses on the same genera by comparing their predictions with the results generated by the ANCHOR method [56, 57], which predicts protein binding residues located in the disordered regions. The predictions of the disordered residues should significantly overlap with the binding residues that are annotated with ANCHOR. To test that we compute the fraction of binding residues that are predicted by ANCHOR and that are also predicted as disordered by each of the three consensuses. We compute a ratio between this fraction and the fraction of randomly generated binding residues that are also predicted as disordered; we generate the same number of binding residues and regions as generated by ANCHOR and place them randomly in protein sequences excluding the locations of the ANCHOR’s predictions. The ratio equal 1 would indicate that disorder is equally abundant in the randomly picked residues and in the disordered protein binding regions coming from ANCHOR, while ratios > 1 would indicate that disorder is enriched among the ANCHOR-generated binding residues. On the *Escherichia* we obtain ratios of 6.02, 4.13 and 3.20 for the consensuses with 5, 10 and 11 methods, respectively, while for *Drosophila* the ratios are 3.47, 2.64 and 2.39, respectively. These high values demonstrate the expected substantial enrichment of intrinsic disorder in the disordered protein binding regions. This suggests that the considered consensuses generate accurate predictions, which in in agreement with the prior empirical evaluations of these methods [36, 58].

### 2.4 Analysis of statistical significance

We assess statistical significance of the relationship between enrichment or depletion of intrinsic disorder in the DNA- or RNA-binding proteins relative to the disorder in the corresponding proteomes for each domain or kingdom of life. We quantify this using paired tests that evaluate significance of differences between \(\text{aveDC}_{\text{bind}}\) and \(\text{DC}_{\text{proteome}}\) (between the DC the DNA- or RNA-binding proteins and the whole proteome) paired for the same proteomes and over all proteomes in a given domain or eukaryotic kingdom. We also evaluate the significance of difference in the fraction of DNA- or RNA-binding proteins with LDRs between the DNA- or RNA-binding proteins and the whole proteome, again paired for the same proteomes and over all proteomes in a given domain or eukaryotic kingdom. First we test normality of the values of \(\text{aveDC}_{\text{bind}}, \text{DC}_{\text{proteome}}\) and the fractions using Anderson–Darling test at \(p\)-value of 0.05. We use the Student’s paired \(t\)-test for the normal data; otherwise we apply the Wilcoxon signed-rank test.

### 2.5 Functional annotations of the DNA- and RNA-binding proteins

We functionally annotate the considered DNA- and RNA-binding proteins using the GO annotations collected from the release 2015_08 of UniProt. We only consider reviewed UniProt entries and aggregate the annotations by the corresponding domains of life. We generate a list of all molecular functions from GO and Enzyme Commission (EC) numbers associated with the RNA- and DNA-binding proteins in the UniProt for each of the three domains of life. To allow for statistically sound estimates of the abundance of intrinsic disorder we include a given functional annotation in our analysis if the count of proteins that it covers > 20. In case of the enzyme types, we focus the analysis of eukaryotic species since the coverage of the EC numbers in the Archaea and Bacteria is relatively low. Moreover, we also collected annotations of four specific functional classes of RNA- and DNA-binding proteins for which the count of proteins > 20: transcription factors, histones, ribosomal proteins and splicing factors. These were identified by the corresponding keywords in the
Figure 1. Relationship between average disorder content in an entire proteome (species) and average disorder content in the corresponding DNA-binding (panel A) and RNA-binding proteins (panel B) for the considered 1121 proteomes. Each marker represents a proteome (species). The black diagonal line represents the positions where the average disorder content in the DNA-binding or RNA-binding proteins equals to the average disorder content in the entire proteome. Markers above (below) the diagonal line correspond to species that have enriched (depleted) disorder content in the binding proteins when compared to all proteins.

protein name. In total, we consider 36 molecular functions in Archaea (17 for DNA-binding and 19 for RNA-binding proteins), 85 molecular functions in Bacteria (34 for DNA-binding and 51 for RNA-binding proteins), 194 molecular functions in Eukaryota (117 for DNA-binding and 77 for RNA-binding proteins), 23 enzyme types (10 for DNA-binding and 13 for RNA-binding proteins) and four functional classes in Eukaryota. Some of these functional annotations are common for both DNA- and RNA-binding proteins and across the three domains of life.

3 Results

3.1 Enrichment in intrinsic disorder in the DNA- and RNA-binding proteins

We compare the DC computed per proteome (DC\textsubscript{proteome}) with the content computed for the DNA- or RNA-binding proteins (aveDC\textsubscript{bind}) in the same proteome. The results for the considered 1121 proteomes are grouped by their taxonomic annotations into Archaea, Bacteria, Eukaryota and four eukaryotic kingdoms including animals, plants, fungi and Protista (Fig. 1). Points located above the black diagonal line indicate species for which abundance of intrinsic disorder in the nucleic acid binding proteins is enriched compared to their overall abundance of disorder. Our analysis suggests that DNA-binding proteins are enriched in disorder in proteomes from all eukaryotic kingdoms (Fig. 1A). The RNA-binding proteins are enriched in proteomes from Bacteria, Archaea and in most of the Eukaryota, in particular, in the majority of animal and fungi proteomes (Fig. 1B). The amount of enrichment is substantial and can be as high as 100%, which we observe for some animal and plant proteomes for the DNA-binding proteins and for some bacterial proteomes for the RNA-binding proteins.

We aggregate these results for each domain and eukaryotic kingdom of life, compute the change in the DC in the RNA- and DNA-binding proteins relative to the content in their proteomes, assess statistical significance of these changes and compute correlation between the DC in proteomes and in the corresponding nucleic acid binding proteins (Table 1). The results reveal that the median increase of the DC in the DNA-binding proteins equals 46% in Eukaryota, with animal and plant proteomes having the largest median increases at 63 and 49%, respectively. The median DC in these proteins in proteomes from Eukaryota is 0.3 with 0.32 in animals, 0.31 in fungi and 0.28 in plants. This means that about a third of residues in these DNA-binding proteins are disordered. The enrichment in the disorder in the DNA-binding proteins in Eukaryota and all considered here eukaryotic kingdoms is statistically significant (p-value < 0.01). Interestingly, the correlations between DC in the DNA-binding proteins and
Table 1. The content and enrichment of the disorder in the DNA- and RNA-binding proteins in the considered 1121 proteomes. All measurements are computed at the proteome (species) level and grouped into the corresponding domains of life and eukaryotic kingdoms. We report the 20\textsuperscript{th} centile (c20), 50\textsuperscript{th} centile (median) and 80\textsuperscript{th} centile (c80) over all species in a given domain/kingdom. We include values of disorder content in the RNA- and DNA-binding proteins, relative disorder content (content in the binding proteins relative to the content in a given species), fraction of proteins with LDRs, relative fraction of proteins with LDRs and Pearson Correlation Coefficient (PCC) values and statistical significance of differences between the overall disorder content and disorder content in the binding proteins, and between the overall fraction of proteins with LDRs and fraction of the binding proteins with LDRs. The Eukaryotic kingdoms are sorted by the median value of the relative disorder content.

<table>
<thead>
<tr>
<th>Binding type</th>
<th>Domain or kingdom of life</th>
<th>Disorder content in binding proteins</th>
<th>Relative disorder content</th>
<th>PCC</th>
<th>Significance of differences in content (p-value)</th>
<th>Fraction of binding proteins with LDRs</th>
<th>Relative fraction of proteins with LDRs</th>
<th>PCC</th>
<th>Significance of differences in fraction (p-value)</th>
</tr>
</thead>
<tbody>
<tr>
<td>DNA</td>
<td>Archaea</td>
<td>0.06 [0.05, 0.14]</td>
<td>–4% [–17%, 12%]</td>
<td>0.95</td>
<td>0.0720</td>
<td>0.05 [0.02, 0.19]</td>
<td>17% [–19%, 87%]</td>
<td>0.96</td>
<td>8.63E–06</td>
</tr>
<tr>
<td></td>
<td>Bacteria</td>
<td>0.07 [0.05, 0.09]</td>
<td>–9% [–24%, 5%]</td>
<td>0.88</td>
<td>7.84E–42</td>
<td>0.06 [0.03, 0.11]</td>
<td>–16% [–60%, 15%]</td>
<td>0.89</td>
<td>6.77E–13</td>
</tr>
<tr>
<td></td>
<td>Eukaryota</td>
<td>0.30 [0.24, 0.34]</td>
<td>46% [29%, 65%]</td>
<td>0.75</td>
<td>1.30E–107</td>
<td>0.70 [0.57, 0.80]</td>
<td>72% [52%, 88%]</td>
<td>0.84</td>
<td>4.05E–151</td>
</tr>
<tr>
<td></td>
<td>Animals</td>
<td>0.32 [0.25, 0.35]</td>
<td>63% [46%, 76%]</td>
<td>0.89</td>
<td>1.15E–59</td>
<td>0.69 [0.56, 0.79]</td>
<td>79% [70%, 90%]</td>
<td>0.88</td>
<td>3.34E–72</td>
</tr>
<tr>
<td></td>
<td>Plants</td>
<td>0.28 [0.26, 0.31]</td>
<td>49% [36%, 66%]</td>
<td>0.76</td>
<td>6.21E–52</td>
<td>0.62 [0.55, 0.70]</td>
<td>88% [58%, 102%]</td>
<td>0.91</td>
<td>3.19E–65</td>
</tr>
<tr>
<td></td>
<td>Fungi</td>
<td>0.31 [0.25, 0.35]</td>
<td>37% [29%, 48%]</td>
<td>0.60</td>
<td>2.08E–16</td>
<td>0.76 [0.66, 0.84]</td>
<td>68% [53%, 83%]</td>
<td>0.70</td>
<td>4.75E–20</td>
</tr>
<tr>
<td></td>
<td>Protista</td>
<td>0.22 [0.15, 0.32]</td>
<td>18% [4%, 55%]</td>
<td>0.77</td>
<td>1.39E–05</td>
<td>0.54 [0.39, 0.71]</td>
<td>44% [27%, 77%]</td>
<td>0.82</td>
<td>1.04E–12</td>
</tr>
<tr>
<td>RNA</td>
<td>Archaea</td>
<td>0.09 [0.07, 0.19]</td>
<td>46% [28%, 69%]</td>
<td>0.97</td>
<td>2.45E–28</td>
<td>0.06 [0.03, 0.25]</td>
<td>48% [8%, 154%]</td>
<td>0.97</td>
<td>5.04E–13</td>
</tr>
<tr>
<td></td>
<td>Bacteria</td>
<td>0.12 [0.10, 0.15]</td>
<td>56% [32%, 88%]</td>
<td>0.83</td>
<td>2.25E–298</td>
<td>0.14 [0.09, 0.21]</td>
<td>102% [43%, 184%]</td>
<td>0.81</td>
<td>3.38E–225</td>
</tr>
<tr>
<td></td>
<td>Eukaryota</td>
<td>0.21 [0.16, 0.26]</td>
<td>4% [–10%, 18%]</td>
<td>0.76</td>
<td>0.0015</td>
<td>0.54 [0.43, 0.64]</td>
<td>34% [15%, 50%]</td>
<td>0.81</td>
<td>3.66E–84</td>
</tr>
<tr>
<td></td>
<td>Animals</td>
<td>0.21 [0.16, 0.25]</td>
<td>13% [–10%, 25%]</td>
<td>0.84</td>
<td>0.0052</td>
<td>0.52 [0.41, 0.61]</td>
<td>37% [25%, 49%]</td>
<td>0.85</td>
<td>7.17E–43</td>
</tr>
<tr>
<td></td>
<td>Fungi</td>
<td>0.20 [0.18, 0.21]</td>
<td>6% [–7%, 19%]</td>
<td>0.63</td>
<td>0.0001</td>
<td>0.49 [0.42, 0.55]</td>
<td>54% [16%, 71%]</td>
<td>0.85</td>
<td>2.23E–41</td>
</tr>
<tr>
<td></td>
<td>Plants</td>
<td>0.24 [0.18, 0.27]</td>
<td>4% [–8%, 14%]</td>
<td>0.32</td>
<td>0.2659</td>
<td>0.60 [0.49, 0.67]</td>
<td>31% [15%, 47%]</td>
<td>0.47</td>
<td>3.44E–10</td>
</tr>
<tr>
<td></td>
<td>Protista</td>
<td>0.16 [0.12, 0.20]</td>
<td>–5% [–22%, 8%]</td>
<td>0.81</td>
<td>0.0201</td>
<td>0.44 [0.27, 0.59]</td>
<td>18% [0%, 47%]</td>
<td>0.79</td>
<td>0.0004</td>
</tr>
</tbody>
</table>
in the whole proteomes are very high, 0.6 or higher, in all domains of life; this can be observed in Fig. 1A. This means that proteomes that have lower or higher overall amount of disorder will also have similarly decreased or increased DC in the DNA-binding proteins. Results for the RNA-binding proteins show that organisms from Archaea and Bacteria are characterized by substantially larger amounts of disorder in these proteins compared to their overall amount of disorder. The median enrichment values of disorder in the RNA-binding proteins among these species are 46 and 56%, respectively, and these increases are statistically significant ($p$-value $< 0.01$). The enrichment in Eukaryota is relatively small at 4% ($p$-value $= 0.002$), with the exception of animal species that have the median enrichment at 13% ($p$-value $= 0.005$). Similarly to the results for the DNA-binding proteins, the DC in the proteomes is highly correlated with the disorder in the RNA-binding proteins; see also Fig. 1B. The overall distribution of the RDC values for the RNA- and DNA-binding proteins in the considered domains and kingdoms of life is shown in Fig. 2A. It confirms that DNA-binding proteins are enriched in disorder in virtually all Eukaryotes while having no bias in the Archaea and Bacteria (black bars). Also, the RNA-binding proteins are universally and strongly enriched in disorder in Bacteria and Archaea and slightly enriched in the animals (gray bars).

We analyze further details of the enrichment in the intrinsic disorder in DNA- and RNA-binding proteins using histograms of the RDC for the nucleic acid binding proteins (Supporting Information Fig. S3). The $x$-axis shows RDC where positive values denote enrichment and negative depletion in DC in the nucleic acid binding proteins relative to the content of their proteomes. The first and last intervals on the $x$-axis denote DNA- and RNA-binding proteins that are structured and highly disordered, respectively. The abundance values for each interval, which are in the form of error bars, give the distribution of the fraction of proteins from specific proteomes in a given domain of life including the 5th, 20th, 50th (median), 80th and 95th centiles. Our analysis demonstrates that the differences in the DC range from a strong depletion to a strong enrichment. We note a bias towards enrichment in the DC for the RNA-binding proteins and for the DNA-binding proteins in Eukaryota; this is visible as a shift towards larger fractions of binding proteins (higher values on the $y$-axis) for the RDC values above 0 in Supporting Information Fig. S3. Majority of these DNA- and RNA-binding proteins have their relative enrichment in the DC below 250% (the median fraction of the binding proteins drops to low values for the $x$-axis values over 250%; see Supporting Information Fig. S3), although some binding proteins are enriched in disorder by as much as over 500%. This is true for the RNA-binding proteins in Archaea and Bacteria and stems from the overall lower amounts of disorder in these two domains of life compared to Eukaryota. However, a relatively substantial fraction of RNA-binding proteins (between a few and about 15%) and DNA-binding proteins (several to about 15% in Archaea and Bacteria and up to about 10% in Eukaryota) are structured. Thus, although overall we observe enrichment in the disorder in the nucleic acid binding proteins...
proteins, some of them are fully structured. Interestingly, both RNA- and DNA-binding proteins from animals (gray lines in Supporting Information Fig. S3E and F) are characterized by histograms that indicate increase in the enrichment (depletion) that is higher (lower) than the overall values for Eukaryota over the entire range of the RDC values.

3.2 Enrichment in intrinsically disordered domains in the DNA- and RNA-binding proteins

Overall, about 39% of proteins in Eukaryota, 10% in Archaea and 9% in Bacteria have LDRs; i.e., IDR having 30 or more consecutive disordered residues, which constitute disordered domains (Supporting Information Table S1). We analyze differences in the abundance of proteins with LDRs between the DNA- and RNA-binding binding proteins and the overall proteomes. Table 1 shows that similarly to the DC, the DNA-binding proteins are significantly enriched in LDRs in Eukaryota \( p \)-value < 0.01). The overall median enrichment equals 72%, and the median enrichment for animals and plants is 79 and 88%, respectively. About 68% of DNA-binding proteins in the eukaryotic species have LDRs compared to only about 12% in Archaea and 8% in Bacteria (Supporting Information Table S1). Interestingly, the RNA-binding proteins across all domains of life are significantly enriched in LDRs, see Table 1 \( p \)-value < 0.01). The median enrichment values range between 18% in Protista and 102% in Bacteria, and they are at 37% and 54% for animals and fungi, respectively. Our analysis reveals that about 50% of RNA-binding proteins in Eukaryota have disordered domains, 15% in Bacteria and 11% in Archaea (Supporting Information Table S1). In spite of having relatively small magnitude, the latter two numbers are significantly higher than those for the corresponding complete proteomes, see Table 1 \( p \)-value < 0.01). Figure 2B shows the distribution of the fraction of proteins that have LDRs and further confirms the notion that nearly all species in the three domains of life are enriched in the disordered domains in the RNA-binding proteins (gray bars), while this enrichment for the DNA-binding proteins holds only for the Eukaryotes (black bars). Taken together, we show that the amount of disordered domains is significantly enriched in the RNA-binding proteins across all three domains of life while the DC in these proteins is also significantly enriched in Bacteria, Archaea and animals. Furthermore, DNA-binding proteins in Eukaryota are significantly enriched in the disorder and in the disordered domains.

3.3 Comparison of results based on reviewed and a combined set of reviewed and automatically annotated DNA- and RNA-binding proteins

Since our analysis is based on both manually and automatically (based on sequence alignment) annotated nucleic acid binding proteins, we compare these results with a small subsets of manually reviewed annotations. The coverage rates of reviewed proteins are very low in most proteomes (Supporting Information Fig. S1) and thus to secure sufficient amount of data we select the proteomes with the coverage rates of at least 20%. Consequently, our analysis includes 31 such proteomes with five in Archaea, 18 in Bacteria and eight in Eukaryota. We compare the distribution of the per proteome values of DC and fraction of proteins with LDRs in the DNA- and RNA-binding proteins in each domain of life when considering the reviewed and all (reviewed and automatically annotated) proteins (Supporting Information Fig. S4); these data can be compared with Table 1 that is based on the complete set of 1121 proteomes. The values are very similar between the analysis using reviewed and all entries. For instance, Supporting Information Fig. S4A shows that the median DC in the DNA-binding proteins in species in Archaea for both reviewed and all proteins is 5.6% (6% in Table 1), in Bacteria for both is also equal 5.6% (7% in Table 1) and in Eukaryota it equals 30.1 and 33.8% for all and reviewed entries, respectively (30% in Table 1). The content in the RNA-binding proteins in species in Archaea for the reviewed and all proteins is 9.0 and 10.5%, respectively (9% in Table 1), in Bacteria is 11.0 and 9.7%, respectively (12% in Table 1), and in Eukaryota is 25.0 and 25.1%, respectively (21% in Table 1). We observe similar results for the fractions of proteins with LDRs. For instance, in Eukaryota 69.4 and 71.9% of the reviewed and all DNA-binding proteins, respectively, have LDRs (70% in Table 1) and 58.0 and 58.6% of the reviewed and all DNA-binding proteins, respectively, have LDRs (54% in Table 1). Overall, these numbers for the reviewed and all binding proteins from the selected 31 proteomes and from the all binding proteins in the full set of 1121 proteomes are similar and preserve the relative (between the domains of life) similarities and differences. This suggests that the inclusion of the automatically assigned annotations does not affect the overall conclusions.

3.4 Peculiarities of charge distribution

Figure 3 summarizes the MNC differences of proteins from the whole genomes and RNA- and DNA-binding proteins in Archaea, Bacteria and Eukaryota (black bars). As expected, the nucleic acid binders are characterized by higher net charge. This is because the protein-nucleic acid binding events involve interactions between the positively charged amino acids of the nucleosome members with the negatively charged phosphate backbone of DNA or RNA. This also agrees with the fact that net charge was found predictive for the DNA- and RNA-binding proteins and sites [59–61]. Furthermore, we investigate the values of the MNC in the LDRs localized in the DNA- and RNA-binding proteins compared to the LDRs in the whole genomes (gray bars in Fig. 3). Similar to the analysis for the whole proteins, there is a visible increase in the net charge for the disordered regions in the RNA-binding
proteins in Archaea and Bacteria and to a lesser extend in Eukaryota. This means that the disordered domains are not only enriched in the RNA-binding proteins in the three domains of life (Fig. 2B) but they are possibly directly involved in the binding. However, in spite of the substantial enrichment of the LDRs in the DNA-binding proteins in Eukaryota (Fig. 2B), these regions have similar net charge when compared with the overall population of LDRs (Fig. 3).

3.5 Analysis of functional annotations in the nucleome

We analyze the enrichment in the disordered content for a set of nucleic acid binding proteins with well-annotated (manually reviewed) in the UniProt functional classes. We compute the median RDC per protein in each functional annotation (molecular functions from GO, EC numbers and the four specific functional classes) and each type of binding and select the annotations that are substantially depleted or enriched in disorder, i.e., the RDC > 50% or < −50%, respectively, when compared to the content in the same domain of life. In Eukaryota, we raise the threshold to 200% given the large number of enriched and depleted in disorder annotations in this domain of life. Consequently, out of 36 considered molecular functions in Archaea, 19 are either enriched or depleted in disorder with eight for the DNA-binding proteins (Fig. 4A) and 11 for the RNA-binding proteins (Fig. 4B). Similarly, in Bacteria we selected 30 out of 85 considered molecular functions, 11 for the DNA-binding proteins (Fig. 4C) and 19 for of the RNA-binding proteins (Fig. 4D). Finally, using the higher threshold, in Eukaryota we found 46 enriched or depleted molecular functions, 34 in the DNA-binding proteins (Fig. 4E) and 12 in the RNA-binding proteins (Fig. 4F), compared to the 194 considered annotations. We also found nine out of 23 considered types of enzymes in Eukaryota that are depleted or enriched in disorder, four in the DNA-binding proteins and five in RNA-binding proteins (Fig. 4G).

In Archaea (Fig. 4A and B) we found 3′-5′ exonucleases, polymerases, ribosomal proteins and ribonucleases to be enriched in disorder, while the 5′-3′ exonucleases, endonucleases, proteins associated with magnesium and ATP binding, and some of the ligases are depleted in disorder. In Bacteria (Fig. 4C and D), single stranded DNA-binding and double stranded RNA-binding proteins, ribosomal proteins, phosphatases and some mRNA-binding proteins are enriched in disorder. The list of molecular functions in Bacteria that are depleted in disorder includes magnesium binding proteins, 5′-3′ exonucleases, some of the hydrolases, polymerases, ligases, nucleotidytransferases, sulfotransferases, adenylyltransferases and methyltransferases. In Eukaryota (Fig. 4E and F) majority of the molecular functions are enriched in disorder and they include transcription factor and chromatic binding proteins, beta-catenin, ubiquitin, p53 and heparin binders, some of the kinases and methyltransferases and snRNA binders. The annotations that are depleted in disorder in Eukaryota include some of the ligases and ribonucleoside binding proteins. Analysis of the various types of enzymes (Fig. 4G) is consistent with the analysis based on the GO’s molecular functions. We found that some of the methyltransferases and lyases are enriched in disorder, while a class of enzymes that act on ester bonds and some of the polymerases and ligases are depleted in disorder. Our analysis of the four specific functional classes that include histones, transcription factors, splicing factors and ribosomal proteins reveals that they are all substantially enriched in disorder in Eukaryota, with the median RDC of 317, 195, 355 and 47%, respectively.
Figure 4. Median relative disorder content per protein for the enriched or depleted in disorder functional annotations of the DNA-binding and RNA-binding proteins. Panels A and B, C and D, E and F show results for the GO's molecular functions for the DNA- and RNA-binding proteins in Archaea, Bacteria and Eukaryota, respectively. Panel G shows results for the enzyme types for the DNA-binding and RNA-binding proteins in Eukaryota. Black and gray bars corresponding to the DNA-binding and RNA-binding proteins, respectively. Solid and hollow bars corresponding to functions where the disorder is enriched and depleted, respectively. The x-axis is in the logarithmic scale.

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4 Discussion

Summarizing, for the first time, we performed a comprehensive computational analysis of the abundance of intrinsic disorder and intrinsically disordered domains in ~548,000 nucleic acid binding proteins found in 1121 complete proteomes of species from the three domains of life. These whole complements of proteins involved in interactions with nucleic acids (DNA and RNA) comprise nucleomes of different species. On average, the nucleome accounts for ~8.7% of the corresponding proteome. Although the 1121 analyzed proteomes contain, on average, ~16% of proteins with LDRs, the corresponding fractions are significantly larger among the DNA- and RNA-binding proteins: 23.6 and 24.7% of them, respectively, contain LDRs. Overall, our study reveals that relative to other proteins in the corresponding proteomes, the DNA-binding proteins are characterized by the significantly increased DC and are significantly enriched in disordered domains in Eukaryotes (p-value < 0.01) but not in Archaea and Bacteria. The RNA-binding proteins are significantly enriched in disordered domains in Bacteria, Archaea and Eukaryota (p-value < 0.01), while the overall abundance of disorder in these proteins is significantly increased in Bacteria, Archaea and animals. We also show that proteins in nucleomes are noticeably more charged than other proteins in corresponding proteomes and that the nucleosomal LDRs typically retain this bias toward higher content of charged residues. A very noticeable exception is the LDRs of the DNA-binding proteins in Eukaryota. In fact, although these proteins are characterized by the substantial enrichment in LDRs (Fig. 2B), their putative LDRs are characterized by the net charge similar that of other eukaryotic LDRs (Fig. 3). This finding points out to the intriguing possibility that the disordered domains of the eukaryotic DNA-binding proteins can have moonlighting functions other than DNA-binding and that these LDRs can also be used for regulation. This may also be true in reverse, meaning that the disordered domains that are not annotated as DNA-binding could have this as a “secondary” function.

Importantly, our analyses on disorder-function association show that among the highly disorder-enriched nucleic acid binding proteins are histones (over 300% enrichment), transcription factors (~200% enrichment), ribosomal proteins (~50% enrichment) and splicing factors (over 300% enrichment). These findings are in excellent agreement with the results of previous studies. In fact, earlier bioinformatics analyses revealed that many transcription factors are highly enriched in intrinsic disorder [23–25], with amino acid sequences of 401 human transcription factors being ~50% disordered [25]. Histone tails (N-terminal domains of core histones and C-terminal domains of linker histones) were long known to be important members of the IDP realm [62]. They contain multiple sites of various posttranslational modifications that modulate the structure of chromatin [63, 64] and constitute the basis of the histone code [64–68]. Interestingly, a recent large-scale analysis of the members of histone family, where 2007 non-redundant chains from 746 species were studied, revealed that majority of the histone family members were predicted to be mostly disordered, with intrinsic disorder extending far beyond the limits of the aforementioned tails [26]. Another recent bioinformatics analysis of the 3411 ribosomal proteins from 32 species revealed that >35% of these proteins are completely disordered and almost all remaining ribosomal proteins contain disordered domains [27]. Finally, computational analyses showed that ~50% of the 109 yeast spliceosomal proteins were predicted to be mostly disordered, 44 and 48% were expected to be moderately (10% ≤ DC < 30%) and highly disordered (disorder ≥ 30%), respectively, and that only ~8% of the yeast spliceosomal proteins were expected to be highly ordered proteins containing less than 10% of disordered residues [29].

Concluding, our study provides compelling evidence for the high abundance of disorder in nucleomes of species from three domains of life. Therefore, this work provides a strong support to the notion that the nucleic acid binding proteins often require intrinsic disorder for their functions and regulation.

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5 References


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