Comparison of DNA binding across protein superfamilies

**Running title:** Analysis of DNA-binding superfamilies

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Abbreviations: TF = Transcription Factor, ZF = C2H2/C2HC zinc fingers, HE = Homing endonucleases, RE = Restriction endonucleases, LR = lambda repressor-like, H = Homeodomain-like, P53 = p53-like, WH = Winged helix, GR = Glucocorticoid receptor-like, RHH = Ribbon-helix-helix, RMSD = root mean square deviation, IAS = interface alignment score, DBD = DNA binding domain
ABSTRACT

Specific protein-DNA interactions are central to a wide group of processes in the cell and have been studied both experimentally and computationally over the years. Despite the increasing collection of protein-DNA complexes, so far only a few studies have aimed at dissecting the structural characteristics of DNA binding among evolutionarily related proteins. Some questions that remain to be answered are: a) what is the contribution of the different readout mechanisms in members of a given structural superfamily, b) what is the degree of interface similarity among superfamily members and how this affects binding specificity, c) how DNA-binding protein superfamilies distribute across taxa, and d) is there a general or family-specific code for the recognition of DNA. We have recently developed a straightforward method to dissect the interface of protein-DNA complexes at the atomic level and here we apply it to study 175 proteins belonging to 9 representative superfamilies. Our results indicate that evolutionarily unrelated DNA-binding domains broadly conserve specificity statistics, such as the ratio of indirect/direct readout and the frequency of atomic interactions, therefore supporting the existence of a set of recognition rules. It is also found that interface conservation follows trends that are superfamily-specific. Finally, this paper identifies tendencies in the phylogenetic distribution of transcription factors, which might be related to the evolution of regulatory networks, and postulates that the modular nature of zinc finger proteins can explain its role in large genomes, as it allows for larger binding interfaces in a single protein molecule.
INTRODUCTION

The specific interactions between short DNA sequences and proteins are a central feature of a wide group of processes in cell biology and organism development. Therefore, the study of the mechanisms of specific DNA binding by dedicated proteins has raised most attention. In addition to genetic, biochemical and molecular biology approaches, it seems clear that a systematic study of the characteristics of the complexes formed between proteins and DNA at the atomic scale will provide a better understanding of the recognition process. To date several reports have shed some light into the structural and functional characteristics of DNA-binding protein families \(^1\)-\(^5\) and the sequences recognized by DNA binding domains (DBD) \(^6\)-\(^10\). These studies have resulted in important contributions describing the interplay between DNA and protein during the recognition process and the structural determinants both at the protein and DNA contact surfaces responsible for specific recognition.

The high scientific relevance of the problem of protein-DNA recognition has contributed to a great increase in the number of high-quality structures of DNA-binding proteins reported in the Protein Data Bank \(^11\). The structures, especially those of their complexes with DNA, have provided valuable insight into the stereochemical principles of binding, including how particular base sequences are recognized and how the DNA structure is often modified on binding \(^3\). The availability and steady growth of structural data of protein-DNA complexes has constituted the ground for a group of computational studies describing the characteristics of the amino acid-base interactions that determine binding specificity \(^12\),\(^13\), the different types of readout mechanisms involved in DNA recognition \(^14\),\(^15\) and the evolutionary conservation of the residues located at contact interfaces \(^16\)-\(^18\).

Protein-DNA interaction can be seen as a reaction in which one or more protein domains dock to the major and/or minor grooves of a DNA double helix. It is known that specificity is
determined by the contribution of direct readout \(^1\) — i.e., associated to direct atomic contacts formed between atoms from amino acid side chains and nitrogen bases —, indirect readout \(^14,15\) — i.e., mediated by the conformational changes undergone by DNA and the contribution of residues that are not in direct contact — and desolvation of the contact interface upon binding. The diverse studies carried out so far have suggested that the relative contributions of each one of the mechanisms related to specificity are different for each DNA binding protein. It has also been claimed that the combination of the intermolecular and intramolecular readout energies leads to an enhanced specificity of recognition. The existence of a “universal” or “generic” protein-DNA recognition code at the atomic level has been proposed based on the strength of contact preferences \(^12\). Nevertheless, many amino acids form favorable contacts with different bases, making it necessary to generalize a deterministic recognition code to a probabilistic binding profile maximizing the likelihood of observed protein-DNA contacts \(^19\). However, several other reports question the existence of such kind of generic code for protein-DNA interaction \(^20\), while others argue that a family-specific code might exist \(^3\).

In a wide-ranging study of all the available structures of DBDs \(^21\), Siggers \textit{et al.} were able to cluster these domains according to the geometric conservation of the contact interface with DNA. They found that, with few exceptions, proteins within a structural family form definite clusters. Another remarkable conclusion from this work is that, although proteins with similar folds tend to dock in similar ways, important differences are observed that seem to correlate with the level of sequence conservation at the docking interface \(^22\). Siggers \textit{et al.} also proved that homologous interfaces tend to maintain certain contacts, even if this requires a distortion of the DNA. However, this study was mainly focused on the geometric properties at the protein side of contact interfaces, and thus was unable to address some interesting questions such as: a) what is the relative contribution of specific amino acids and bases at the interface to specific
binding, and b) what nitrogen bases contribute the most to indirect readout recognition and how this contribution affects the specificity of recognition. We have recently developed a simple methodology to generate atomistic representations of protein-DNA interfaces \(^{23}\), which has previously been used with fairly good results to generate structure-based models of transcription factor binding sites. This computational protocol, named DNAPROT, permits a detailed structural dissection of the interfaces accounting for direct and indirect readout of DNA, including the contribution of the interactions mediated by water molecules and allowing sampling and optimization of amino acid side-chains rotamers \(^{24}\) and has successfully been applied to a representative set of crystallographic structures and homology models.

This paper addresses some of the aforementioned issues and systematically explores the conservation of structural features of binding interfaces, centering the study both at the protein and DNA sides of docked complexes. In particular, the DNAPROT methodology is applied to 9 superfamilies from the Structural Classification of Proteins (SCOP) \(^{25}\) and the results demonstrate that evolutionarily unrelated DNA-binding domains conserve important specificity statistics, such as the ratio of indirect/direct readout and the frequency of atomic interactions, but also unveil patterns that are superfamily-specific. Although it had already been described that prokaryotic genomes have a dominant proportion of Winged helix transcription factors, known to be functional as dimers, and that larger metazoan genomes are enriched in zinc finger proteins, here we propose that this evolutionary trend is related to the modular nature of zinc fingers, which can be concatenated to ensure enough binding specificity, by means of larger interfaces, in a single protein molecule.
MATERIALS AND METHODS

Set of protein-DNA complexes and SCOP annotation

The subset of protein-DNA complexes in the Protein Data Bank \(^{11}\) (release May 15\(^{th}\) 2009) was downloaded and the accompanying list of clusters of protein chains with 95\% of sequence identity (under the derived_data directory) was parsed to define a non-redundant set of 175 monomeric complexes. For each cluster, the chain with best resolution was taken, considering NMR structures only when no crystallographic structures were available. For homodimeric complexes, only one chain was taken, the one appearing first in the corresponding non-redundant list. All non-redundant chains found as part of heterodimers or higher order complexes were considered. The protein sequence of each non-redundant complex was searched against SUPERFAMILY (version 1.69) using the superfamily.pl script with default parameters \(^{26}\) in order to be assigned to SCOP superfamilies, which are expected to share evolutionary history \(^{25}\), and to precisely define domain boundaries. To minimize sampling problems, only those superfamilies containing more than 5 complexes were further considered in this work. The full list of 175 complexes is available as Supplementary Material and their mean sequence identity percentages are shown in Table I. Complexes 1au7_B, 1e3o_C, 1ic8_A, 2d5v_B, 2h8r_A, 1fok_A and 2o61_A were not considered as they contain two DNA-binding domains from different SCOP superfamilies.

Atomic dissection of interfaces

The DNAPROT algorithm was applied to each monomeric complex in order to analyze the binding interface in terms of direct — i.e., atomic interactions: hydrogen bonds, water-mediated hydrogen bonds and hydrophobic interactions — and indirect — i.e., sequence-specific DNA geometry deformations — readout. It is important to note that DNAPROT considers only those atomic interactions that are sequence-specific, those that involve amino acid side-chains and...
purine/pyrimidine rings. Among hydrophobic interactions, only thymine C7 interactions are considered. Electrostatic interactions also play a major role in protein-DNA binding, but only a small fraction are expected to contribute to specific recognition, and these correspond to the interface hydrogen bonds mentioned earlier. Using this methodology we obtained a structure-based position weight matrix for each interface, in which the direct/indirect relative contribution of each nitrogen base is assessed. Briefly, the saturating mutation strategy implemented in DNAPROT iteratively evaluates the interaction potential of a given protein-DNA complex while each base at the crystallographic site is mutated by the other three bases. Each single mutant is processed to obtain the contacts and deformation contribution of the given base and the analysis of all possible mutants renders a matrix in which the direct and indirect readout contributions are linearly combined by means of a deformation weight. Indirectly recognized base pairs are defined as those columns in the abovementioned indirect readout position specific matrix in which at least one nucleotide has a frequency greater than 40 in a 0 to 100 scale — e.g. the prior frequency for all four nucleotides is by default 25. The DNA motif bound by any protein in this dataset is defined as the shortest oligonucleotide that encompasses all directly read bases plus all indirectly read bases that are less than 3 nucleotides away. This was necessary to adequately handle multimeric PDB structures that include several protein chains bound to the same DNA duplex. For the calculation of the indirectly readout fraction, motifs shorter than 4 nucleotides were not considered. This cut-off was chosen empirically based on the knowledge of the characteristics of TF-binding sites, which usually correspond to regions of 3–6 nucleotides. Motifs of 3 nucleotides were excluded to avoid overestimating indirect readout fractions in the case of interfaces with few dissected atomic interactions.

**Multiple alignment of DNA-binding superfamilies**

The first step was to split all domains contained in the set of complexes of each superfamily, as
many proteins contained more than one domain — e.g. the set of 13 C2H2/C2HC zinc finger (ZF) proteins yielded 35 domains. The domain boundaries reported by SUPERFAMILY, as explained above, were generally followed, but were manually corrected for some concatenated ZF proteins. These domains were then structurally aligned by MAMMOTHmult \(^{30}\), with the aim of putting all binding interfaces in the same frame of reference, and were further analyzed by using the Interface Alignment software \(^{21}\), which reports interface alignment scores for pairs of complexes. Further multiple alignments were calculated using SUPERFAMILY hidden Markov models. DANGLE 0.63 (http://kinemage.biochem.duke.edu/software) was employed to calculate side chain torsion angles. In order to assign single interaction roles to interface residues, which can be easily displayed in a multiple alignment, the following rules were applied in this order of priority:

1) if the residue forms 1 or more hydrogen bonds it is called a hydrogen bond residue
2) if the residue forms 1 or more hydrophobic interactions it is called a hydrophobic residue
3) if the residue forms 1 or more water-mediated hydrogen bonds it is called a water-mediated residue

Rotamers were clustered using the CPAN Algorithm::Cluster module (http://search.cpan.org/dist/Algorithm-Cluster/) by requiring two side chains to be in the same cluster if both their \(\chi_1\) and \(\chi_2\) angles were less than 40° away \(^{31}\). Multiple alignments and superposition PDB files are available at http://www.eead.csic.es/compbio/suppl/dna_families/mammoth.zip and http://www.eead.csic.es/compbio/suppl/dna_families/alignments.zip, respectively.

**Distribution of DNA-binding proteins at genomic scale**

We downloaded the genomic annotations for predicted transcriptions factors from release 2.0 of
DBD (http://www.transcriptionfactor.org) \(^{32}\), completing this information with the genomic assignments for restriction and homing endonucleases, which were kindly provided to us by curator Derek Wilson. This set was filtered in the following steps:

1) Genomes in which the subset of DNA-binding proteins (DBPs) that belong to our defined set of 9 superfamilies accounts for < 50% of the annotated DBPs were discarded. This was done to avoid species in which the repertoire of DBPs is not adequately represented by these 9 superfamilies.

2) Genomes with less than 100 annotated ORFs were discarded.

3) Genomes with less that 10 annotated TFs were also rejected.

4) Phyla with less than 4 genomes were ignored, with the exception of Nematoda, for which only 3 genomes are available.

The remaining 490 annotated proteomes were used to generate Figure 4.
RESULTS AND DISCUSSION

This section presents the results of several analyses carried out with a non-redundant set of 175 protein-DNA complexes obtained as explained in Materials and Methods. This set contains 9 protein superfamilies, which are now listed together with their abbreviated names and the number of complexes included in each superfamily: C2H2/C2HC zinc fingers (ZF, 13), Homing endonucleases (HE, 11), Restriction endonuclease-like (RE, 16), lambda repressor-like (LR, 12), Homeodomain-like (H, 38), p53-like (P53, 17), Winged helix (WH, 38), Glucocorticoid receptor-like (GR, 22) and Ribbon-helix-helix (RHH, 8).

Contribution of indirect readout to DNA recognition across superfamilies

The first question that this paper aims to address is: how important indirect readout mechanisms are in different DNA-binding superfamilies? For this purpose we applied the DNAPROT algorithm to all members of the set described above and estimated how many base pairs of the bound DNA motif are, on average, recognized by means of sequence-specific deformations of the DNA duplex. These indirect readout estimations correspond to energetic potentials of deformation associated to DNA base steps.

The results are displayed in Figure 1, with two observations to be made: a) the fraction of base pairs that are indirectly read within DNA motifs — i.e., the median values of the distributions represented by the box plots — is typically small, with an average of 20% and b) restriction endonucleases have a substantially larger proportion of indirectly readout bases (depicted as filled hexagons in the interface of Figure 1B). Overall, the contribution of indirect readout mechanisms, as dissected by our methodology, is rather small. However, it is important to note that these are superfamily generalizations — i.e., individual complexes may depart from the superfamily-shared behavior as implied by the range of variation in each superfamily. For instance, among members of the C2H2/C2HC zinc finger superfamily, the Wilms tumor
suppressor was found to substantially distort DNA upon binding, and hence more than half of its DNA motif is subject to indirect readout mechanisms. Two of the superfamilies included in the present study correspond to proteins with enzymatic activity, the restriction and homing endonucleases. As can be seen in Figure 1, for these two cases we obtained the highest and lowest indirect readout contributions to binding specificity, which is in agreement with the experimental data obtained for some members of these superfamilies. A recent study of DNA recognition by restriction endonucleases underscored the relevance of this indirect type of reading of DNA, as they proved that a mutant enzyme deficient of direct contacts showed no loss of sequence specificity. There are also other examples, such as the reports of the molecular structure of BgIII and NaeI, which found large DNA rearrangements that occur upon specific binding. Therefore, the structural evidence at hand indicates that restriction enzymes extensively utilize indirect readout to bind DNA sequences very specifically. Enzyme MunI (PDB: 1d02) turns out to be a special case, as it induces a kink on its target oligonucleotide, but this deformation is not translated into a significant increase in binding specificity as estimated by DNAPROT (see Supplementary Figure 2).

For the homing endonucleases, those that yield the lowest fraction of indirect readout, our findings are in accordance with the biological function of these enzymes, engaged in processes of specific chromosome cleavage. Proteins within this family contain a high number of direct contacts, as can be seen in Figure 2, with a relatively low contribution of indirect reading of DNA. Figure 1 also shows that the median indirect readout contributions of the others superfamilies, most of which are transcription factors, are somewhere in-between the values obtained for the enzymes. Although the contribution of indirect reading to specificity is not fully understood yet, the analysis of our results and previous reports suggests that the generic function of the superfamily — e.g. whether the members recognize very specific sequences, a large
number of different sequences, a restricted number of similar but not identical sequences or being completely unspecific — might be related to the ratio of indirect readout. While some reports propose that for some specific transcription factors 40, and restriction endonucleases 34, the indirect reading of DNA is case-dependent, the evidence presented here implies that the structural constraints of superfamilies impose limits on the indirect readout fractions of individual DBDs. In the case of restriction endonucleases, proteins with a common core fold 41 but with highly different DNA-binding regions 3, that must bind well defined sequences with high specificity, indirect readout can be used as a “pre-screening” mechanism during target site location by reducing the search space according to the deformation propensity of the sites 34. For homing endonucleases, which include a group of proteins with marked structural differences among them 39, indirect readout may not be such determinant because direct contacts alone may account for an efficient recognition. In the case of transcription factors indirect readout contributions correspond to intermediate values possibly related to multiple recognition scenarios — i.e., large regulons for global regulators or a compact group of similar sequences for local regulators 42 —, and are also likely influenced by the limitations that the DBD imposes on the number of sequences than can be recognized, as described for zinc fingers 20.

**Contribution of direct readout (atomic interactions) to DNA recognition across superfamilies**

This section analyzes direct readout, probably a more tractable mechanism than indirect readout, because its energetic contribution is made of pairs of interfacial atoms of both the protein and DNA molecules that interact. The results depicted in Figure 2 correspond to the frequency of sequence-specific atomic interactions within protein-DNA interfaces. Perhaps the most important observation is that there is an almost constant contribution of hydrogen bonds, water-mediated hydrogen bonds and hydrophobic interactions to the binding interface across
superfamilies. However, it seems clear that hydrogen bonds are the main source of specific interactions (on average they account for 72% of interactions), while the relevance of hydrophobic interactions is minor. A similar scenario is discovered when atomic pair potentials are used, providing further evidence in the same direction (see Supplementary Materials). It should be noted that only thymine C7 hydrophobic interactions were considered, as only these contacts were found to confer specificity in previous work \(^\text{43}\), and this explains this reduced contribution.

These results suggest that the molecular basis of direct readout follows general principles, that are shared by unrelated DNA-binding proteins, and therefore support the existence of a set of recognition rules, a code, at the atomic level, in agreement with the observations of Luscombe \textit{et al.} \(^\text{12}\). This set of rules can be associated to a general set of atomic interactions among chemical groups at the interface, involving the same amino acids and bases in somehow similar chemical contexts from one DBD to another. However, interface residues often contact several nitrogen base groups simultaneously, and the interface architecture imposes geometric restrictions that favor some particular contacts over others. For these reasons, it is not generally possible to translate these atomic preferences into a one-to-one residue-base code, which would necessarily be affected by superfamily-specific conditions, as previously claimed \(^\text{44,45}\). For instance, interactions between asparagine and adenine account for more than 14% in Homeodomains but only for 4% among Winged helix transcription factors. The same applies for the interactions between lysine and guanine at the interfaces of GR, H and WH proteins, where the summed relative contribution of water-mediated and direct hydrogen bonds accounts for 26%, 7% and just 1% respectively.

As in the previous section, it is worth noting that there are individual complexes that show an array of interactions clearly different from that of their superfamily. Among lambda repressors,
the interface of repressor protein P22 c2 (PDB: 2r1j) shows only one direct hydrogen bond per monomer, and recognition seems to be substantially driven by a combination of indirect readout, a key hydrophobic contact, and several water-mediated bonds ⁴⁶.

In addition to this, Figure 2 also shows the average number of atomic interactions in each superfamily (see the right vertical axis). In this respect, it would appear that there are two types of superfamilies: a) those in which the average number of contacts per monomer is between 3 and 7 and b) C2H2/C2HC zinc fingers and Homing endonucleases, with more than 14 interactions per monomer. The first type includes superfamilies, such as Homeodomains and Winged helices, which are known in many cases to be only functional as dimers. Thus, it can be assumed that these proteins will often be binding to DNA targets with an average number of interactions around 12. C2H2/C2HC zinc fingers can achieve a similar number of contacts without necessarily requiring the formation of dimers, but instead are usually made of several domains in tandem in the same polypeptide. For instance, the Wilms tumor suppressor mentioned earlier contains four canonical zinc fingers in a row ³³. Homing endonucleases, usually embedded in introns or inteins, appear to have the largest number of interactions at the interface, and this is consistent with the fact that these double-stranded DNases bind to extraordinarily large recognition sites, from 12 to 40 nucleotides long ⁴⁷. If data from Figure 1 and Figure 2 are combined, it could be argued that restriction enzymes (RE) belong to a third type of DNA-binding proteins, with an intermediate number of contacts at the interface and a very important indirect readout component.

**Structural comparison of DNA-binding interfaces**

The next step in our analysis is to compare structural determinants within each superfamily, with the aim of updating and extending the pioneering work of Pabo and Nekludova ⁴⁵. As a prerequisite each superfamily has to be put in a common frame of reference, and a natural way
of achieving this is by means of structure fitting. The degree of similarity of these proteins can be calculated at the domain scale, by means of average root mean square deviations (RMSD), or by focusing on the subset of interface residues, those that mediate direct readout, in which case the interface alignment score (IAS) of Siggers *et al.* 21 can be employed. In addition, we ask if aligned interface residues play a similar interface role across members of the superfamily or whether they have a similar spatial arrangement, which can be measured in terms of side chain torsion angles. Figure 3 shows the superposition and corresponding multiple alignment computed by MAMMOTHmult 30 for 35 C2H2/C2HC zinc finger domains, extracted from the 13 members of that superfamily, and serves as a guide to Table I, which summarizes the structural analyses performed on the 9 superfamilies subject of this study. There are 7 interface columns in this superfamily, of which 6 (86%) are shared among several domains. Among these core interface positions, there is one dedicated exclusively to hydrophobic interactions in two domains (column 18), while the rest have mixed uses, dominated by residues that make specific hydrogen bond interactions with nitrogen bases. Figure 3C summarizes the rotamer clusters found for interface residues aligned in column 20. While there are four clusters, the first cluster is the largest and includes different amino acids extracted from different complexes.

A quantitative evaluation of the structural statistics of binding interfaces in all 9 superfamilies can be found in Table I. Overall, structural analysis finds that ZFs are the smallest DNA-binding domains, which are very similar between them under both RMSD and IAS metrics, and have a very compact set of core interface residues, as described in the literature 5. In addition, zinc finger proteins contain on average 2.7 domains per protein chain, in contrast with most other superfamilies, which contain on average only one DNA-binding domain per chain. There are however other observations to be made to this table. For instance, in terms of RMSD, p53-like and Restriction endonuclease (RE) domains are found to be the most divergent, as expected for
their low percentages of sequence identity \cite{22}, and indeed these superfamilies include domains which cannot be superposed in frame. Multiple alignments based on SUPERFAMILY hidden Markov models confirm the observed structural divergence (see Table I and Supplementary Material). Winged helix domains are also divergent, but their superposition is in frame. In terms of IAS, proteins from the Glucocorticoid receptor superfamily are found to have very similar interfaces, in contrast with WHs, that appear to have a very flexible way of binding to the major groove of DNA. With respect to interface size, Ribbon-helix-helix and ZF proteins stand out for being compact DNA binders, which tend to use a small subset of residues to drive specific recognition. On the other end of the scale are REs, which have a large number of interface positions. Despite their different interface sizes, proteins from these superfamilies have an average of 55% of interacting residues in the core. These are positions observed in several members of the superfamily, which suggest that a common binding architecture might exist. However, REs display a different trend, as they seem to have almost unique arrays of interface residues when compared to each other. We now analyze the interaction roles of interface residues, which can display different recognition uses. Homeodomain and RHH domains tend to have conserved roles for their interface residues, whilst the remaining superfamilies are able to use the same interface position to have, say, a hydrogen bond in one complex and a hydrophobic interaction in another complex.

Finally, Table I also indicates that most (83 to 100%) interface positions across superfamilies can be expected to have at least two side chain rotamer conformations in the same superfamily. The data in Table I can also be used as a guide for comparative modelling exercises. For instance, the selection of protein templates, which should then be aligned to a query sequence in order to build a three-dimensional molecule \cite{48}, will be affected by the degree of conservation of binding interfaces, as suggested by previous reports \cite{49,50}. In addition, the variable size of the
core interface across superfamilies is expected to affect the outcome of methods that predict interface residues \(^5\). Finally, the finding that interface side-chains cluster in a few groups can be exploited in molecular modelling exercises to drive conformational searches.

The structural analysis of evolutionarily related DNA binding proteins yields the following conclusions. As with indirect readout, it is observed that the DBD of proteins from a given superfamily share structural features, but there is variability within superfamilies. It seems that DBDs with a common ancestor have evolved to develop flexible binding interfaces that allow equivalent positions to bind to different nitrogen bases and accommodating variants of rotamer conformations, while retaining some key core positions, such as glutamine 50 in Homeodomains \(^{45}\). Despite the overall conservation of geometry within superfamilies, the number of interface changes required to distinguishing one DBD from the other can be very small, as illustrated by CRP and FNR, two homologous TFs in *Escherichia coli* that can be interconverted with just a couple of point mutations \(^{52}\).

**Distribution of DNA-binding proteins across genomes**

In this section we wish to focus on the distribution of DNA-binding proteins (DBPs) across completely sequenced genomes, with the aim of searching for evolutionary trends that might complement our structure-based analysis. Figure 4 summarizes the proportion of annotated DBPs that belong to any of the considered superfamilies among 490 proteomes grouped in 15 phyla, which have been sorted by mean proteome size. A first inspection indicates that the relative proportion of endonucleases is rather low in most genomes, with only a little presence of these domains in higher organisms. Our data show that homing endonucleases are more abundant in Tenericutes and extremophiles of the phylum Euryarchaeota than in other bacteria, presumably because their opportunities for homing are limited \(^{38}\). HEs are also scarce among metazoans, possibly because their segregated germ lines impedes horizontal transmission \(^{38}\).
With respect to restriction endonucleases, these domains are more frequent in Tenericutes and Crenarcheota, although are frequently found in all bacterial genomes, as shown in Figure 4.

The bar plot also shows that Homeodomains are the only superfamily of TFs which have an important (10% or more) presence across phyla, from prokaryotes to Archaea and eukaryotes. Winged helix TFs are also present in all phyla, but while they are the dominant proteins in prokaryotes and Archeaea (with proteome sizes below 5000 bases), they represent just a minor (5%) fraction of TFs in metazoan genomes. Lambda receptor proteins represent 7-23% of TFs in organisms with small proteomes, but are less than 1% of metazoan TFs. The opposite case are Glucocorticoid receptor TFs, which are almost absent in small genomes but account for 13% and 37% of TFs in Arthropoda and Nematoda, respectively, with a minor fraction in Chordata. Finally, while C2H2/C2HC zinc fingers are almost absent in small genomes, they seem to be the preferred TFs of metazoan organisms, as they account for 27% to 76% of annotated TFs.

This data, together with the observations in Figure 2, suggests that transcriptional regulation has followed two distinct evolutionary paths in small and large genomes. In prokaryotic genomes, such as *Escherichia coli*, regulation is dominated by Winged helix TFs, that only accumulate a sufficient number of interface contacts after dimerization, a reaction which is usually dependent on some effector signal. Instead, metazoan organisms, with genomes that can be several orders of magnitude larger, have chosen modular C2H2/C2HC zinc finger domains, that can be easily concatenated in evolution and ensure enough binding specificity — i.e. enough atomic interactions spanning a longer oligonucleotide, in a single protein molecule. Furthermore, it is well documented that ZFs can still interact with other proteins and form multimeric complexes. 
CONCLUSION

After analyzing a comprehensive collection of protein-DNA complexes, we estimate that the average contribution of indirect readout to specific binding is approximately of one every five DNA bases, with the notable exception of restriction enzymes, which double its contribution. Furthermore, proteins from the same superfamily often display uneven indirect readout behaviors. With respect to direct readout, hydrogen bonds dominate DNA recognition, with a minor fraction of hydrophobic interactions. The constant contribution of atomic interactions across superfamilies supports the existence of a general set of recognition rules at the atomic level. It also appears that most superfamilies have a number of atomic interactions per monomer near 6, except Homing endonucleases and C2H2/C2HC zinc finger, which have more than 14 contacts on average.

Comparison of proteins from the same superfamily by means of structural fits indicates that some superfamilies show larger interface variability than others, particularly restriction enzymes and p53-like proteins. Aligned interface residues of Homeodomain and RHH domains are less likely to switch their interaction type than the rest, and, in general, their side chains cluster in rotamer groups.

In summary, while direct readout employs a generic code of recognition at the atomic level, the architecture of individual DNA-binding proteins show subtle superfamily deviations that determine a more case-specific way of DNA recognition. This corresponds to an scenario in which the same set of rules — i.e., the interactions between specific groups from amino acids and nitrogen bases — can be tuned at the structural level, involving deformation of DNA, the geometry of the binding domain and rotamer variants of amino acid side-chains, to ensure a specific binding and versatility of DNA recognition.

Finally, a survey of the frequency of transcription factor superfamilies across 15 phyla finds
clear patterns of distribution, confirming that prokaryotic genomes are preferentially regulated
by Winged helices, whilst metazoans are rich in zinc finger TFs. Here we propose that the
modular nature of zinc finger domains, which can be concatenated to ensure enough binding
specificity in a single protein molecule, can explain this evolutionary trend.
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FIGURE LEGENDS

Figure 1. Relevance of indirect readout across representative SCOP superfamilies. A) Box plots of the distribution of frequencies of indirectly recognized motif positions in 9 different DNA-binding SCOP superfamilies, calculated from the corresponding number of monomeric complexes in parenthesis, shown next to its abbreviation. The boxes correspond to the interquartile range, the median is represented with a bold line and the minimum and maximum values are marked with whiskers connected with a dashed line. Outliers of the distribution are represented with open dots. B) Atomic interface graph of restriction enzyme BgIII 35 (PDB: 1dfm_B ), with 7 base pairs in the bound DNA motif involved in indirect readout mechanisms, shown as filled bases. The arrows represent atomic interactions at the interface — i.e., solid arrows = hydrogen bonds, dashed arrows = water-mediated bonds, dotted arrows = hydrophobic interactions — and are displayed to illustrate the fact that motif positions are often recognized by means of both indirect and direct readout.

Figure 2. Direct readout explained in terms of atomic interactions at the protein-DNA interface. The bar plot depicts the mean contribution of hydrogen bonds, water-mediated bonds and hydrophobic interactions to direct readout across the set of 9 SCOP superfamilies (see abbreviations in Figure 1). A line plot is overlaid showing the average number of interface atomic interactions for each superfamily, according to the secondary Y-axis at the right end of the diagram.

Figure 3. Alignment and superposition of 35 C2H2/C2HC zinc finger domains. A) Structure-based multiple sequence alignment, with ruler on top and interface summaries at the bottom. Column 17, an example of multiuse interface position, contains 23 residues that form hydrogen bonds, while 2 residues take part in water-mediated bonds and another 2 form hydrophobic interactions. These 27 equivalent interface residues can be grouped in 5 side chain
rotamer clusters. Column 18 is an example of interface residue dedicated exclusively to hydrophobic contacts, observed in only two fingers (alanine in 2i13_A_3 and histidine in 1g2d_C_2) that belong to the same rotamer cluster. In this panel, ‘*’ means >20 residues and ‘+’ means >10 residues. B) Backbone cartoon of the corresponding structural superposition. C) Summary of 4 side chain rotamer clusters found for interface residues aligned in column 20 of the multiple alignment.

**Figure 4. Genomic distribution of DNA-binding superfamilies.** Bar plot of the distribution of 9 DBD superfamilies across the annotated genomes of 15 phyla. The data for each phyla was derived from the number of genome sequences shown in parenthesis. The fraction annotated as homeodomains is shown in number, as this superfamily corresponds to at least 10% of the annotated TFs in all phyla. A line plot is overlaid that shows the mean proteome size of each phyla, as the secondary Y-axis describes. The Arthropoda, Nematoda and Chordata phyla are summarized in the text as metazoans. This figure was prepared with data from the DBD database (http://www.transcriptionfactor.org)\textsuperscript{32}. 
**Figure 1**

A

- Ribbon-helix-helix (RHH, 6)
- Glucocorticoid receptor (GR, 21)
- Winged helix (WH, 36)
- p53-like (P53, 17)
- Homeodomain-like (H, 40)
- lambda repressor-like (LR, 15)
- Restriction endonuclease (RE, 15)
- Homing endonucleases (HE, 11)
- C2H2/C2HC zinc fingers (ZF, 13)

Below is a graph showing the frequency (indirect readout base pairs in site) of various protein domains.

B

- DNA sequences with indicated bases and sites labeled with A, G, T, and C.
Figure 2

Frequency of interface atomic interactions:
- Water-mediated hydrogen bonds
- Hydrophobic interactions
- Contacts/complex

Average interactions per monomer:
- ZF: 18
- HE: 16
- RE: 14
- LR: 12
- H: 10
- P53: 8
- WH: 6
- GR: 4
- RHH: 2
### Table I: Structural descriptors of 9 DNA-binding superfamilies. SCOP domains with boundaries defined by SUPERFAMILY\(^ {26} \) were superposed with MAMMOTHmult\(^ {30} \). The mean number of domains extracted per complex is shown in the second column, while the length of the resulting multiple alignments is given in the third column. Restriction enzymes and p53-like domains are marked with asterisks to indicate that their multiple rigid superpositions include domains that cannot be fit in frame, which affects the calculated statistics. The mean RMSD column shows the mean core size (in residues) and the resulting root mean square deviation of all pairwise domain structural alignments generated in the course of the progressive multiple alignment. The next column gives the mean sequence identity of these pairwise alignments. The mean IAS columns presents the mean protein-DNA interface similarity score of Siggers et al.\(^ {21} \). The remaining columns give more detailed information about the interfaces, as shown in Figure 3, and are all calculated in the frame of reference defined by the structural superpositions of

<table>
<thead>
<tr>
<th>superfamily (SCOP domains)</th>
<th>domains/protein</th>
<th>multiple alignment length (columns)</th>
<th>mean RMSD (Å, residues)</th>
<th>mean % sequence identity</th>
<th>mean IAS</th>
<th>interface (columns)</th>
<th>%core columns</th>
<th>%multiuse columns</th>
<th>%different rotamers columns</th>
</tr>
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<tr>
<td>HE (14)</td>
<td>1.4</td>
<td>216</td>
<td>3.23 (83)</td>
<td>19</td>
<td>3.65</td>
<td>33</td>
<td>67</td>
<td>68</td>
<td>89</td>
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<tr>
<td>ZF (35)</td>
<td>2.7</td>
<td>35</td>
<td>1.66 (26)</td>
<td>38</td>
<td>3.46</td>
<td>7</td>
<td>86</td>
<td>83</td>
<td>100</td>
</tr>
<tr>
<td>H (47)</td>
<td>1.1</td>
<td>191</td>
<td>2.87 (44)</td>
<td>16</td>
<td>2.53</td>
<td>37</td>
<td>54</td>
<td>35</td>
<td>100</td>
</tr>
<tr>
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<td>4.21</td>
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<td>44</td>
<td>43</td>
<td>100</td>
</tr>
<tr>
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<td>94</td>
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<tr>
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<td>3.34</td>
<td>17</td>
<td>41</td>
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<td>2.44</td>
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<td>75</td>
<td>0</td>
<td>100</td>
</tr>
<tr>
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<td>3.42 (69)</td>
<td>13</td>
<td>3.05</td>
<td>26 [34]</td>
<td>50 [21]</td>
<td>23 [29]</td>
<td>83 [100]</td>
</tr>
</tbody>
</table>
domains. Bracketed statistics for restriction enzymes and p53-like domains correspond to hidden Markov multiple alignments computed with SUPERFAMILY, calculated to demonstrate that different alignments still unveil large differences among domains. First, the *interface* column states how many columns of the original multiple alignment include interface residues, those that establish atomic interactions with nitrogen bases. Then, the *core* column shows the fraction of interface columns shared by at least two protein-DNA complexes in the same superfamily. Next, the *multiuse* column shows the fraction of core columns that includes residues that form atomic interactions of different types in different complexes. Finally, the *different rotamer* column states the fraction of core columns that include interface residues whose side chains belong to different rotameric states.