**Reads Meet Rotamers:**

**Structural Biology in the Age of Next Generation Sequencing**

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**Abstract: [[Maybe re-work the abstract after all of the changes/updates are introduced]]**

Structure has historically been interrelated with sequence, usually in the framework of comparing sequences across species sharing a common structural fold. However, the nature of information within the sequence and structure databases is evolving, changing the type of comparisons possible. In particular, we now have a vast amount of personal genome sequences from human populations and a larger fraction of new structures contain interacting proteins within large complexes. Consequently, we have to recast our conception of sequence conservation (considering more selection within the human population) and its relation to structure (now focusing more on interacting surfaces rather than folds). We cover this changing mindset here.

**Highlights:**

* Increasing amounts of sequencing data adds new dimension to study structural biology.
* Next generation sequencing facilitates analysis of genetic variations.
* Understanding evolutionary constraints acting on proteins remain elusive.
* Essential to integrate sequences, structures, and interaction networks information to rationalize the phenotypic impact of these variations.

**Introduction:**

The amount of genomic information is growing at an astonishing pace due to rapid improvements in next-generation sequencing (NGS) technology (Figure 1A) [1]. The nature of biological information stored within biological databases is undergoing a transformation (Figure 1). Before the completion of the human genome project in 2003, we had a large amount of genomic sequence information from different species as well as structural data. Since the technological advances in next-generation sequencing, the amount of human sequence information has grown at a rapid pace. Meanwhile, while structural biologists continue to deposit new structures in the PDB database, the pace of identifying new folds has slowed down indicating that few new folds remain undiscovered. However, the complexity of the structure in the PDB database continues to grow indicating that there is an increasing emphasis among structural biologists to treat biomolecules not as individual folds but rather as complex molecular machines that interact and regulate each another as they function within the cellular environment (Figure 1). Together, these trends suggest that the stage is set to integrate sequence and structural information to rationalize the effect of variants on protein function.

The identification and characterization of pathological disease-associated variants is an essential goal of genomic sequencing efforts [2,3]. A large number of medically-relevant mutations occur within proteins, some of which are available through databases such as the Online Database of Mendelian Inheritance in Man (OMIM) [4], the Human Gene Mutation Database (HGMD) [5], Humsavar [6], and ClinVar [7]. It is essential to utilize structural information for rationalizing the evolutionary pressure acting on these proteins as well as for developing drugs to combat the effects of disease-causing variants. However, it remains challenging to annotate the physical effects of these mutations on proteins due to the heterogeneous nature of functional constraints acting on a protein family. A protein-coding variant may cause local perturbations, or global changes in structure, or it may have a substantial impact on the protein-protein interaction (PPI) network, and each type of change adds a different layer of functional constraints on the protein. Such analyses are further complicated by the fact that we currently have incomplete knowledge of these constraints, and also by the fact that specific combinations of individually benign variants may cause disease.

Moreover, we expect the growing genomic information to be a valuable resource for structural biologists. In particular, as the amount of genomic data continues to grow, we envision a future in which biologists will utilize genetic variation within human population(s) to help interpret their structural data [8]. Population genetic analysis within human proteins has already been used to identify novel species-specific functional constraints within a protein family [9]. In addition, a number of fundamental insights about biological pathways can be garnered by analyzing newly discovered loci associated with a disease [10].

In this review article, we initially explain how genomic information is used to identify pathological disease associated variants as well as variants that are harmful to protein function even within healthy individuals. We later describe how structural information is utilized to understand the harmful effects of different variants. Finally, we discuss how it is necessary to integrate sequence and structural data into a holistic system or network perspective before we can predict whether any particular variant will be benign or whether it will have a phenotypic effect.

**Classical Sequence Comparison:**

Typically, structural biologists identify functionally constrained regions within a protein family by comparing homologous sequences from different species (Figure 2a) [11,12]. They focus on changes that take place over longer evolutionary timescales by comparing the dominant sequence within each species rather than focusing on intra-species changes. Nucleotides that do not change across different species are conserved over millions of years and are hence considered to be functionally important. Due to redundancy within the genetic code, some of the changes in the coding regions are silent as they occur without a corresponding change in the protein sequence (synonymous changes). While exceptions do exist, all synonymous changes and a majority of the nonsynonymous changes are expected to be neutral. A small fraction of the nonsynonymous changes can, however, either be harmful (deleterious) or beneficial to the fitness of the species. The ratio of nonsynonymous to synonymous variants (dN/dS) is commonly utilized to characterize the selection pressure on the coding regions of the genome (Figure 2) [13]. If the dN/dS ratio for a coding region is less than 1, it indicates that a few of these mutations are harmful or deleterious and that these changes are under negative selection. On the other hand, a dN/dS ratio exceeding unity indicates that evolution is promoting a change in the protein sequence and that this protein is under positive selection [9]. Proteins undergoing positive selection may improve the fitness of an organism in different environments.

**Introduction to Population Sequencing [[rephrase?]]:**

The vast amounts of genomic and exome sequences available are providing unique opportunities to characterize genetic variation within the human population. The exome comprises the coding sequences of all protein-coding genes and constitutes approximately 1% of the total genomic sequence [14]. Due to the reduced cost of exome sequencing and better-characterized clinical relevance of variation within the coding regions of the genome, it is more widely used for genetic diagnosis. Variants within an individual’s genome are either acquired at birth (germline mutations) or during the person’s lifetime (somatic mutations) as a consequence of errors during cell division. While germline mutations are typically present in every cell of the person, somatic mutations only affect certain cells and are typically not passed on to the next generation. There are approximately 74 *de novo* (new)variants that occur during each generation [15]. As only germline mutations are passed on to the next generation, somatic mutations are not under conventional evolutionary selection.

The human genome exhibits extensive variation [16-19]. On average, any individual genome contains 20,000-25,000 coding variants (Table 1), of which 9,000-11,000 are nonsynonymous. The frequency with which a particular variant or allele occurs within a population is used to characterize the evolutionary pressure acting on it as common variants (minor allele frequency greater > 5%) are expected to be benign. However, rare variants (minor allele frequency < 0.5%) are rare either because they are harmful (deleterious) to a protein’s function or because the variant has been introduced recently into the population. The ratio of common to rare variants is often used as a proxy to characterize the evolutionary pressure acting on a locus. ~~As deleterious variants are under negative selection, the frequency with which a particular variant or allele occurs in a particular population can be used as a proxy to characterize the evolutionary pressure on it.~~ Although most of the variants within any particular individual are common, most coding variants manifest as distinct single nucleotide variants (SNVs), each of which occurs very rarely within the human population. About 25-50% of the rare non-synonymous variants within healthy individuals are estimated to be deleterious, suggesting that the human proteome is highly robust to a large number of non-specific perturbations and because most rare deleterious variants are heterozygous implying that the cell also contains a functional copy of the gene [18,19].

Despite the fact that new genomic data is still being produced, about 200,000-500,000 previously unobserved SNVs are still discovered after each personal genome is sequenced, suggesting that there is no saturation observed on human polymorphism data yet [18,19]. Indeed, the number of rare variants continues to grow even after the 1000 Genomes Consortium and Exome Aggregation Consortium data (60,706 individuals) [20] data has become available. As deleterious mutations tend to occur at very low frequencies, we need to continue sequencing a large number of individuals to characterize and catalog these variants and their frequencies within the human population.

As such, we can turn to intra-human comparisons to uncover more human- or domain-specific features (Figure 2). [[Maybe elaborate on Fig 2 a little more here. Or maybe ref fig 2 right after the end of the prev para, since the ideas are so closely connected]] There is, however, an important distinction between interpreting inter- and intra-species conservation due to the huge disparities in the associated evolutionary timescales (Figure 2a-c). While performing such an analysis, one can also align homologous coding regions not only between individuals (Figure 2b), but also within a single human genome (i.e., paralogs), such as proteins originating from the same structural domain family (Figure 2c). In particular, this can be used to elucidate domain-specific features. [[ANS2JC – need a couple of sentences explaining Fig 2d and 2e here – especially 2e]]

Similar to the dN/dS ratio in cross-species comparisons, selective pressure on coding regions can be quantified using fraction of synonymous to nonsynonymous polymorphisms (pN/pS) at any site (Figure 2). In addition, evolutionary pressure can also be quantified during intra-species comparison using the ratio of rare to common variants at each site as rare variants are under negative selection. A statistically significant depletion of common variants as compared to rare variants implies that the site is under higher selective pressure.Furthermore, genomic variants that are increasing in frequency within a human population (positive selection) may help identify a novel gain-of-function event (such as a new protein-protein interaction). Some of these domain-specific events may be beneficial to the species. Comparative genetics/genomics studies have already uncovered a growing list of genes that might have experienced positive selection during the evolution of human and/or primates [9]. These genes offer valuable inroads into understanding the biological processes specific to humans, as well as the evolutionary forces that gave rise to them. It is also important to note that some variants occur in a correlated fashion within the population and these variants are said to be under linkage disequilibrium (LD). Note also that LD is statistically easier to observe for common variants than for rare ones.

~~[[Remove this text?]] There is one additional confounding factor to consider while identifying disease-associated variants. Genes associated with a disease are identified by detecting deleterious variants that are affecting genes within diseased individuals more often than in healthy populations. This might be misleading, however, because the variants associated with this gene might be correlated with other unanalyzed variants in the genome. Hence, all variants (including the variants within a gene) statistically associated with a disease might not be causative and additional analysis may be required to identify the real disease-causing mutations. We need to annotate the effect of individual variants, however, before we can predict the outcome of a large number of variants.~~

**Deleterious Effects of Variations on Protein Function:**

The patterns of conservation displayed by proteins are the product of a vast array of constraints active throughout its evolutionary history. In this regard, to understand the physical effects that cause a variant to be harmful, we need to consider the multitude of underlying constraints acting on the protein family. Such constraints are often intrinsic to the structure itself: they may include the need to maintain the integrity of functional hinge regions or interior packing geometry or the ability to regulate a protein through post-translational modifications at specific sites. They may also entail that residues at an interaction interface remain topologically compatible with those in the corresponding interface of an interaction partner. We can utilize the structural information in the PDB database to assess the effect of mutations on a protein’s stability as nonsynonymous changes that occur within the core of the protein or variants that disrupt the secondary structure of the protein could reduce its stability. Several computational tools based on sequence conservation (inter-species or intra-species) and/or several structural features (the physicochemical characteristics of the amino acid change, solvent accessibility, secondary structure, active site annotations, and protein-protein interfaces) were developed to predict the deleterious effect of sequence variations on a protein’s function [22-25]. Disease-associated mutations are highly enriched for residues in the interior of proteins (22% of all mutations in HGMD and OMIM), and active sites of proteins [16-19].

In terms of applying such a catalogue of rules as a means of understanding human disease-associated variants, the fibroblast growth factor receptor provides a case-in-point, several variants in which have been linked to craniofacial defects (Figure 3). The evolutionary constraints listed here provide sensible rationales for how many of these disease-associated variants may impart deleterious effects. Importantly, these constrains may act in synergistic ways rather than through isolated mechanisms \cite{23364837, 22153503}. However, the mechanisms for several other disease-associated variants fail to map to this catalogue, thereby underscoring the need to more comprehensively document sources of constraint. This more comprehensive documentation must transcend the native structure itself to include folding pathways, allosteric regulation, and the functional roles of disordered regions or conformational transitions. Such mutations that affect the thermodynamic stability of different allosteric states of a protein [26] are typically ignored while predicting the deleteriousness of a putative variant.

~~Mutations may not only affect the native state of the protein but could also affect the stability of intermediates within the folding pathway. Such considerations typically ignored while assessing the effect of mutations on a protein’s structure. In addition, mechanistic insight into the mutation- induced structural changes requires knowledge of the folding kinetics, which still remains elusive in these models. Finally, while mutations that occur on the active site of the protein reduce efficiency or ablate function entirely, mutations that are distant from an active site may also affect protein efficiency [26].~~

**Networks as a Framework for Understanding Deleterious Variants:**

While structural and sequence information are invaluable in providing a rationale for the deleterious effects of certain disease-causing and rare variations, it is often difficult to interpret the phenotypic effects of an individual variant without considering the broader cellular context. As proteins are extensively involved in protein-DNA interactions (gene regulatory networks), protein-RNA interactions (post-transcriptional regulation), and protein-protein interactions (PPI) within the cellular milieu, variants that disrupt these interactions could potentially affect the viability of the cell. We refer the reader to comprehensive essays on the phenotypic effect of noncoding variation [27,28], and focus instead on deleterious effects of variants on the protein-protein interaction (PPI) network here.

Various experimental and computational approaches have been applied to characterize the PPI network in several model organisms and human beings [29,30] and these networks have been invaluable in interpreting the role of evolutionary constraints on a protein family \cite{}. In the PPI network, a node represents a protein, while an edge represents an interaction between the two proteins connected by the edge. Proteins that are highly interconnected in PPI networks (hubs) are under strong negative selection constraints while proteins at the periphery of the network are under positive selection in humans [31]. Proteins that are more central in an integrated “multinet” formed by integrating biological networks from different context (PPI, metabolic, post-translational modification, gene regulatory network, etc.) are under negative selection within human populations [32]. In agreement with this, perturbations to hub proteins are more likely to be associated with diseases than non-hub proteins [33]. The PPI networks are organized in a modular fashion as proteins associated with the same function are more likely to interact with one another [34] and proteins associated with similar diseases tend to occur within the same module [33]. The system properties of the network have also been useful in interpreting how the human proteome is robust even in the presence of a large number of deleterious variants within healthy individuals. [[Is this really true?! I thought that the opposite was actually the case. What’s written here is very counter-intuitive…]]Most deleterious variants observed in healthy individuals occur in peripheral regions of the interactome. Such limited[[?]] effects may result as a consequence of compensatory mutations or functional redundancy [35]. On the other hand, cancer-associated somatic deleterious variations occur in the internal regions of the interactome and tend to have larger structural consequences on the PPI network.

[[Minimize the disc in the next few sentences – a lot of this seems very obvious (or maybe just to us)?]]The interactome provides a convenient platform to measure the impact of a deleterious variant on the cell. As shown in Figure 4 [[ANS2SK: Need a couple of examples as suggested by Reviewer 1]], a deleterious variant can either remove a protein (such a node effect would naturally also result in the removal of all the associated edges) from the PPI network by making a protein nonfunctional or it could lead to the loss of just one or more of its interactions (edgetic effects). Mutations at a PPI interface can have drastic effects on the biomolecular binding constant and several sequence and structure-based methods have been proposed to identify these interaction hotspots [36,37]. Even though we have incomplete information on the number of domain-domain interactions (Figure 1), it has been predicted that about 12% of all the HGMD and OMIM mutations occur at a PPI interaction [38] while approximately 28% of experimentally-tested HGMD missense mutations affect one or more interactions, thus underscoring the importance of these interactions for annotating rare variants and disease-associated mutations [39].

In an effort to bridge the information gained from individual structures with network properties of the interactome, Kim, et al., [40] combined the experimentally determined interactome with structural information from the iPfam database to form the structural interaction network (SIN) and were able to obtain a higher-resolution understanding of the selection constraints on the hubs. Using structural information, the hubs were classified into different groups based on the number of distinct interfaces utilized for biomolecular complex formation and they showed that the number of distinct interfaces is a better proxy for evolutionary pressure acting on the hub rather than the number of edges in the PPI network. [[This is a good place to more fully link to the previous discussion on metrics for conservation (ie, the first section)]]Consistent with this interpretation, hub proteins in the PPI network contain a higher fraction of disease-causing mutations on their solvent exposed surface, as compared to non-hub proteins suggesting that a larger fraction of a hub’s disease-associated mutations could affect its interactions [40].

Hub proteins interact with a large number of partners and tend to be more flexible and conformationally heterogenous than non-hub proteins [41]. Furthermore, the number of distinct interfaces in hub proteins is correlated with degrees of conformational heterogeneity [41]. To the extent that variants may enable or disable certain conformational states from being visited, such mutations could potentially affect protein complex formation and signaling pathways, and this has not yet been examined very closely. As deleterious mutations that affect hubs in networks tend to have a larger effect on the structures, they would also cause large changes in the PPI network. Proteins can utilize different interfaces for different (sets of) interactions, so multiple mutations on the same protein can be associated with drastically different diseases depending on the afflicted interface. Such mutations would have different edgetic effects on the protein’s interaction network - by breaking or weakening one of its interactions while the rest of its interactions remain intact - and a large proportion of HGMD and OMIM mutations are predicted to have edgetic effects on the PPI network [39,42].

It should also be noted that the hubs in PPI networks also tend to contain higher degrees of disordered regions, and these regions typically become well-ordered upon ligand or protein binding [49,50]. Disease-associated mutations are enriched within disordered regions of the protein as they could affect post-translational modifications and/or protein-protein interaction sites \cite{24830552, 22080206}. The assessment of a mutation on the activity of an intrinsically disordered protein is even more challenging because it would be dependent upon the effects of these mutations upon the unfolded ensemble or the structure gained in the presence of its interaction partner. Due to their inherent flexibility, the unfolded ensembles of disordered proteins are especially difficult to characterize using either experimental or computational techniques [51,52], making variant annotation in the context of disordered proteins an uphill task. However, the phenotypic effect of mutations on the functional viability of a disordered protein is important because a number of proteins also change their interaction partners in a tissue-specific manner based upon the dominant isoform of the protein in that tissue \cite{22749400}. Recent evidence suggests that many mutations occurring on these alternatively-spliced disordered motifs may drive cancer [53]. Therefore it is important to understand the phenotypic effects of sequence variations in the disordered regions.

Ultimately, we want to develop an integrative framework to understand the effects of deleterious variants on the phenotype of the cell. However, a mutation typically displays tissue-specific phenotypic effects, hence an understanding of functional constraints on a protein should also incorporate tissue-specific information. While the gene regulatory network is being mapped out in a developmental time point and cell type-dependent fashion by several international consortia [43,44] the PPI network is largely treated in a static fashion. Recent work has tried to integrate proteome and gene expression profiles with PPI networks to create tissue-specific networks \cite{24550720, 23399932}[45]. However, these studies typically neglect the protein isoform even though the protein’s interactions are dependent on its isoform [46,47]. A structural study on the effect of sequence variations on isoform-dependent PPI complexes has not been performed and would improve the prediction of phenotypic effects due to missense mutations. However, it is likely that the high costs in resources associated with studying isoform-specific assays in various cell types have impeded these types of studies. We anticipate that isoform-specific protein-protein interaction network annotation will become easier and more accessible in the near future, which will present new opportunities to better annotate such networks.

**Conclusions:**

[[The main theme of bridging the sequence constraints (from next-gen sequencing) to structural properties and nets is very thin here. It is ok to keep as is, but I feel like we should not wait until the conclusion section to integrate these 2 themes – we should see if we can more explicitly integrate these 2 themes within the main body itself, and then summarize/conclude that integration here]]The exponential growth in genomic data has demonstrated that a surprisingly large amount of genomic variation is present within the human population, and this data has also helped identify a vast number of rare variants and disease-associated variants. Though the motivation of developing methods to annotate the effects of variants that cause human disease are clear, it remains challenging to do so as it requires bridging disparate sources of information together to understand the functional constraints on a protein family. It is essential to utilize structural information to rationalize the effect of variants. The network properties of the protein in addition to sequence and structural information regarding the nonsynonymous amino acid changes need to be considered within a single framework before predicting the phenotypic impact of an amino acid change.

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**Figure Captions:**

**Figure 1:** The pace of novel fold discovery has begun to saturate, and while the volume of X-Ray crystal structures and structurally-resolved protein-protein interactions has continued to grow. However, the pace with which personal genomic sequencing databases are growing is considerably greater than the pace at which structure databases are growing.

Figure 2: Evolutionary conservation in different contexts. (a) Examining sequence conservation of homologous sequences across multiple species looks at a longer evolutionary timescale. (b) Examining the enrichment of rare variants (or depletion of common variants) in the same genomic element across multiple individuals within a single species or population looks at a shorter evolutionary timescale. (c) Examining sequence conservation of similar (d) domain sequences within a single genome, (e) reveals species- and domain-specific conservation that might be important to the structure or function of the domain family. Genomic variant profile across the domain can be further analysed, e.g. by comparing the number of non-synonymous (ns) relative to synonymous (s) variants and comparing the number of rare (r) relative to common (c) variants. Here, we use ankyrin domains found in the human genome as an example. The sequence profile of an ankyrin repeat motif is painted green to yellow, corresponding to the structure of each of six ankyrin repeats found in the human Notch 1 ankyrin domain (PDB ID: 1YYH).

**Figure 3:** (A)The fibroblast growth factor receptor is shown in complex with FGF2 (PDB 1IIL), along with the loci of HGMD variants (orange spheres). (B)Various structural annotations (i.e., a “catalogue of constraints”) are shown in sequence space. Hinge residues are taken from HingeMaster [54], buried residues are identified using NACCESS [55], protein-protein interaction residues are defined to be those within 4.5 Angstroms of the co-crystallized growth factor, and post-translational modification sites are taken from UniProt. HGMD loci shown as holo circles coincide with the catalogue of constraints, and may thus likely be rationalized in light of such constraints. However, a large number of HGMD loci (shown in filled orange circles) fail to overlap with these annotations, highlighting the need to consider alternative sources of constraint.

**Figure 4:** Various mechanisms of SNP-induced disruption in protein-protein interaction networks. A SNP that destabilizes a hub protein can ablate all associated interactions (A). SNPs disrupting different interfaces of the hub may interfere with interactions active in different tissues (B, C). Blue (hub protein), Yellow (nodes expressed in tissue1), Green (nodes expressed in tissue2), Turquoise (node expressed in tissue3).

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