

## Personalized Biochemistry and Biophysics

Brett M. Kroncke,<sup>†,‡</sup> Carlos G. Vanoye,<sup>§</sup> Jens Meiler,<sup>‡,||</sup> Alfred L. George, Jr.,<sup>§</sup> and Charles R. Sanders<sup>\*,†,‡</sup>

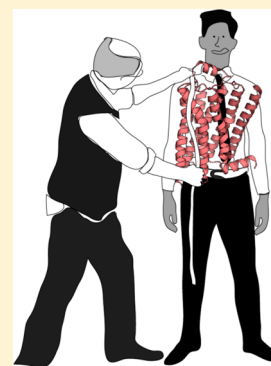
<sup>†</sup>Department of Biochemistry, Vanderbilt University School of Medicine, Nashville, Tennessee 37232, United States

<sup>‡</sup>Center for Structural Biology, Vanderbilt University, Nashville, Tennessee 37232, United States

<sup>§</sup>Department of Pharmacology, Northwestern University Feinberg School of Medicine, Chicago, Illinois 60611, United States

<sup>||</sup>Departments of Chemistry, Pharmacology, and Bioinformatics, Vanderbilt University, Nashville, Tennessee 37232, United States

**ABSTRACT:** Whole human genome sequencing of individuals is becoming rapid and inexpensive, enabling new strategies for using personal genome information to help diagnose, treat, and even prevent human disorders for which genetic variations are causative or are known to be risk factors. Many of the exploding number of newly discovered genetic variations alter the structure, function, dynamics, stability, and/or interactions of specific proteins and RNA molecules. Accordingly, there are a host of opportunities for biochemists and biophysicists to participate in (1) developing tools to allow accurate and sometimes medically actionable assessment of the potential pathogenicity of individual variations and (2) establishing the mechanistic linkage between pathogenic variations and their physiological consequences, providing a rational basis for treatment or preventive care. In this review, we provide an overview of these opportunities and their associated challenges in light of the current status of genomic science and personalized medicine, the latter often termed precision medicine.



More than 80 million validated rare and common single-nucleotide variations (SNVs) in human genomes have already been catalogued (Figure 1).<sup>1,2</sup> Of these, more than 150000 have been documented to predispose individuals to “simple” diseases that exhibit Mendelian (monogenic) inheritance,<sup>3</sup> such as cystic fibrosis and familial early onset Alzheimer’s disease. Others impact the responsiveness of patients to certain drug treatments (pharmacogenomics).<sup>4–6</sup> For example, rare variations in the gene encoding the vitamin K epoxide reductase complex subunit 1 (VKORC1) influence the dosing requirements of the commonly used anticoagulant warfarin, which targets this protein.<sup>7</sup> Still other SNVs are risk factors for complex or sporadic disorders. For example, the  $\epsilon 4$  variant (encoding Cys112Arg) of apolipoprotein E, which is present in roughly 30% of all human genomes, is a risk factor for the common (sporadic) form of Alzheimer’s disease.<sup>8</sup> These advances in medical genetics and human genomics are having transformative effects on medicine and biology.

In clinical medicine, genetic testing for selected gene variations has become the standard of care in many circumstances, particularly when the information provided is actionable as the basis for prophylactic treatment or improved patient care (cf. refs 9 and 10). Recently, a new paradigm of human genetic analysis involving complete sequencing of whole exomes or genomes has emerged as being capable of providing comprehensive lists of rare and common variants present in patient genomes that may be disease-predisposing and/or have pharmacogenomic relevance. However, many of these variants will have uncertain clinical significance.

The ever-increasing speed and decreasing cost of exome and genome sequencing are already providing information about patient genetics that was, until very recently, inaccessible to

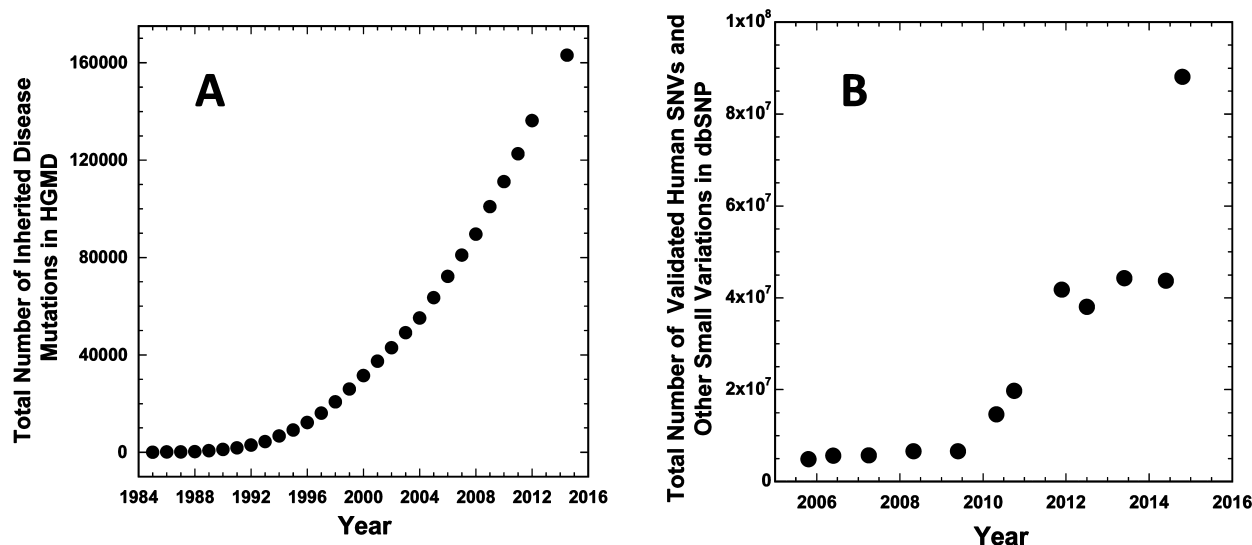
more traditional medical genetic tools.<sup>11</sup> As of early 2015, the utilization of such information in the practice of medicine is at a tipping point. A case study documented by Richard Lifton and colleagues at Yale University<sup>12</sup> provides a glimpse of both the potential and complexities associated with efforts to bring personal genomic information relating to rare mutations to clinical medicine.

In 2013, a 1-week-old baby was admitted to the Yale-New Haven hospital, presenting with fever and diarrhea. Tests for infections were negative. The child’s condition progressively worsened, during which time whole exome sequencing of the patient and his parents was initiated. Because there was no obvious family history of inherited disease and the parents seemed to be healthy, it was thought that the child’s disorder might be caused by a *de novo* mutation, a noninherited germline mutation. Unfortunately, 23 days after the onset of symptoms and 1 day before exome sequencing was completed, the infant died of alveolar hemorrhage. Additional tests revealed extensive physiological damage consistent with an aggressively activated immune system. Comparison of the infant’s exome with those of the parents revealed no informative *de novo* mutations. However, two days after the funeral the father developed a high fever, respiratory distress syndrome, and other symptoms of a highly activated immune system, but without an evident infection. It was only then revealed that the father had suffered from periodic incidents of high fever throughout his life. Similarities in the symptoms and test results between father and son prompted reanalysis of their exomes to search for shared

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**Figure 1.** (A) Growth with time in the total number of identified human mutations that result in inherited (Mendelian) monogenic disorders. Data from the Human Gene Mutation Database.<sup>3</sup> (B) Growth with time in the total number of discovered human genome variations (SNVs and other small scale variations) as logged in the dbSNP Database.<sup>2</sup> The small decreases in the number of variations seen in this plot for some time points reflect the consequences of changes in the reference genome and its annotation with time.

genetic variations. Among 34 previously undocumented shared protein-altering variations was a missense mutation in the *NLRC4* gene encoding a valine to alanine substitution in the nucleotide binding domain (NBD) of the NLRC4 protein, an ATPase that promotes inflammasome assembly. On the basis of the crystal structure of NLRC4,<sup>13</sup> it was deemed probable that the wild-type valine side chain stabilizes the inactive ADP-bound state of NLRC4, an interaction that is disrupted by mutation to alanine, promoting ADP-for-ATP exchange and constitutive activation of inflammasome assembly. NBD variations that trigger gain of function in a related ATPase, NLPR3, had previously been documented<sup>14</sup> to result in dysregulated inflammasome assembly and autoinflammatory syndromes with symptoms similar to those of the father and son of this case study. Subsequent experiments confirmed the pro-inflammatory “gain-of-function” nature of the Val341Ala mutation in transfected macrophages. The condition shared by the father and son in this case study has been named “syndrome of enterocolitis and autoinflammation associated with mutation in *NLRC4*” (SCAN4). A contemporaneous published study reported an unrelated *NLRC4* mutation that resulted in a very similar autoinflammatory syndrome.<sup>15</sup> Given that similar syndromes have previously been shown to be relieved by drugs that target interleukin-1 $\beta$ , SCAN4 is probably a treatable disorder, although the father in this case declined this therapy.

One can imagine an imminent future in which the time line for connecting the dots between genetics and diagnosis will be accelerated to the point where there will routinely be little lag time between disease presentation and the availability of actionable genomic insight, even when very rare variations are involved. Indeed, case studies akin to that summarized above but with happier outcomes can already be cited.<sup>16,17</sup> The sequencing of genomes in children and their parents may soon become routine, and analyses performed in the perinatal period may allow the compiling of genetic variations in a newborn along with identities of the variant proteins and their predicted function. Moreover, the similarity of the rare variations in genes such as *NLRC4* to mutations in homologous proteins

previously demonstrated to be disease-associated could be flagged by bioinformatic algorithms. Much effort is underway to develop the infrastructure and tools to address the practical challenges to effective personal genome-informed medicine, sometimes termed “personalized medicine” or “precision medicine”.

In this work, we highlight some of the opportunities and challenges associated with this burgeoning area of genomic medicine that invite the participation of biochemists, biophysicists, and structural biologists. In terms of scope, we generally focus on mutations that occur in germline DNA rather than on somatic mutations that may cause cancer<sup>18</sup> or contribute to other disorders.<sup>19</sup> However, many of the principles described here also apply to somatic mutations. We also focus on proteins but recognize that many disease-promoting gene variations impact RNA molecules (see Table 1 and ref 20).

## ■ VARIATION IN PERSONAL GENOMES AND HUMAN DISEASE

Table 1 provides a compilation of genome statistics based on 2015 estimates. In that table, we focus on the fraction of the ~4 million variations in a typical human genome that impact the exome, which encodes the proteome (Table 1). Most exomic mutations are SNVs, and each individual has 20000–25000 exomic SNVs, approximately half of which are silent at the protein level. Although most silent variants are probably not disease-linked, some are. This seems to be due either to aberrant RNA splicing<sup>21</sup> or to alteration of the identity of the codon translated during protein synthesis. The latter can induce protein misfolding stemming from an aberrant change in the translation rate and kinetically linked cotranslational folding due to the switch from a common codon to a synonymous rare codon (or vice versa).<sup>22–24</sup>

The most common disease-promoting mutations are found among the 10000–13000 nonsynonymous SNVs (nsSNVs) that encode an amino acid change or, more rarely, introduce an aberrant stop codon.<sup>25</sup> Of these, it is estimated that for each individual, 100–500 nsSNVs will cause loss of protein function

**Table 1. Variations in an Individual Human Genome (estimates as of 2015; per person unless otherwise noted)<sup>a</sup>**

no. of nucleotide base pairs	3.2 × 10 <sup>9</sup>
no. of protein coding genes	~23500
no. of probable essential proteins	2500–5000
no. of exons	180000
size of exome (bp)	30000000
no. of DNA sequence variant (DSVs)	4000000
no. of <i>de novo</i> variants (noninherited germline)	30–100
no. of single-nucleotide variants (SNVs)	3500000
no. of copy number variants (CNVs) and other genome structural variants	10 <sup>4</sup> –10 <sup>5</sup>
no. of nonsynonymous SNPs (nsSNVs, encode changes in amino acid)	10000–13000
no. of synonymous SNPs (sSNVs, alter codon, but not amino acid)	10000–12000
no. of in-frame insertions/deletions (indels) in exons	190–210
no. of aberrant stop codons in exons	25–100
no. of frameshifts in exons	220–250
no. of splice disruption variants	40–50
no. of probable protein loss-of-function (LOF)-inducing nsSNVs	250–500
no. of heritable protein LOF-inducing nsSNVs already logged in the human gene mutation database (HGMD)	40–100
no. of homozygous LOF-inducing nsSNVs	40–85
no. of homozygous LOF-inducing nsSNVs already in the HGMD	3–24
no. of probable disease-causing homozygous LOF-inducing nsSNVs	1–5
percent of healthy individuals carrying a medically actionable homozygous disease mutation	1–4
no. of genetic variations logged as being linked to a Mendelian (monogenic) disorder in the HGMD (total to date)	163000
no. of missense/nonsense variations in HGMD linked to Mendelian disorders (total to date)	91000
no. of RNA splicing–impacting variations in HGMD linked to Mendelian disorders (total to date)	15000
no. of small insertions and deletions recorded in HGMD linked to Mendelian disorders (total to date)	38000

<sup>a</sup>Parts of this table were adapted from Table 1 of ref 25. Most other estimates were gleaned from refs 26–28 and 123 or <http://www.hgmd.cf.ac.uk/ac/index.php>.

(~15% involving premature stop codons, and the rest single-amino acid changes), with 20–85 of these nsSNVs being homozygous.<sup>26–28</sup> A great many disease-promoting nsSNVs have already been documented for Mendelian disorders<sup>3,29</sup> and as risk factors for complex disorders.<sup>30</sup> However, for any personal genome, there will be numerous nsSNVs that must initially be classified as “variants of unknown significance” (VUS) because of uncertainty regarding how the variant will impact protein function and whether there will be physiological or pharmacogenomic consequences for the patient. Some VUS will encode previously undocumented mutations in proteins that are known to be subject to disease-associated mutations, making these VUS prime suspects as disease-promoting variations. Of course, numerous VUS—even those in known disease-linked genes—will be harmless neutral variations.

One can imagine the numerous dilemmas in clinical medicine that will arise from the discovery of VUS in patients. Consider the case in which a child is found to have a newly discovered missense mutation in the gene that encodes KCNQ1, a voltage-gated potassium channel that is an essential component of the cardiac action potential. Many of the several hundred known genetic variants that alter the amino acid

sequence of KCNQ1 predispose individuals to an arrhythmia known as long-QT syndrome.<sup>31</sup> Arrhythmic episodes may go undetected for years but may result in sudden death, often triggered by swimming or exercise.<sup>32</sup> Should the child in our imaginary case study be pre-emptively treated with antiarrhythmic drugs<sup>33</sup> or a medical device [e.g., internal cardioverter defibrillator (ICD)], which have their own risks? Should the child be permitted to swim? Dilemmas such as these may become commonplace in the very near future. There is clearly an imperative for the continued development of methods that can rapidly determine or predict whether new VUS are harmless or predispose a patient to disease and, if so, how.

## ■ DEFECTIVE PROTEINS AND DISEASE

When considering how protein-altering nsSNVs result in disease, it should be kept in mind from gene knockout studies of model organisms that only 10–20% of all proteins appear to be essential for birth and survival.<sup>34–36</sup> Also, recall that humans are diploid organisms such that there are normally two copies (alleles) of each gene, with the exception of genes on the X and Y chromosomes. Homozygous mutations impact both alleles, whereas heterozygous variations impact only one allele.

Many disease-linked nsSNVs resulting in protein LOF are homozygous, such that loss of that protein’s function is complete. The classic example is provided by inherited mutations in the gene encoding the cystic fibrosis transmembrane regulator (CFTR), a chloride ion channel.<sup>37</sup> In contrast, individuals heterozygous for CF mutations do not suffer from the disease because the amount of functional protein originating from the wild-type allele is sufficient to fulfill an essential physiological function of this protein in maintaining proper salt balance in lung alveoli. However, for some proteins, even heterozygous mutations can result in LOF-based disease because of the reduced total functional output of that protein in critical tissues (cf., ref 38) or because the mutant protein can form oligomers with the wild-type protein, rendering it inactive. This latter possibility applies, for example, to heterozygous mutations in peripheral myelin protein 22 (PMP22) that result in the inherited peripheral neuropathy, Charcot-Marie-Tooth disease. Mutant forms of PMP22 misfold and are targeted for degradation. Because they can form oligomers with the wild-type gene product, some of the wild-type protein is targeted along with the misfolded mutant protein for degradation.<sup>39</sup>

LOF due to nsSNVs can result from perturbations of global protein structure, active site structure, posttranslational modifications, trafficking, dynamics, and/or protein–protein or protein–ligand interactions.<sup>40</sup> However, there is considerable evidence that the most common cause of protein LOF is reduced thermodynamic stability, leading to protein for which the folding equilibrium is shifted toward the nonfunctional unfolded state, possibly coupled to irreversible aggregation (thermal instability) and/or degradation by cellular quality control.<sup>41–44</sup>

Less common than disease-linked LOF variants are nsSNVs that result in protein that still functions, but in a dysregulated manner, so-called gain-of-function (GOF) mutants. These include the variation in the NLRC4 ATPase described above in the Yale case study and oncogenic mutations in proteins such as the Ras GTPase and the epidermal growth factor receptor family of tyrosine kinases, which often drive cancer tumor cell development and proliferation.<sup>45,46</sup> Other examples include a number of G protein-coupled receptors that are subject to disease-causing mutations that induce constitutive activation,



resulting in unregulated signaling of normally hormone-regulated pathways.<sup>47</sup> These mutations are often heterozygous, reflecting the fact that unregulated protein function triggered by a GOF mutation in a single allele is sometimes sufficient to trigger pathophysiology. Consider the vasopressin V2 receptor (V2R) located in kidney collecting duct epithelia, which controls uptake of water into the bloodstream and formation of urine. GOF mutations in V2R result in constitutive signaling that causes too much water to be absorbed into the blood from the kidney, resulting in a rare disorder characterized by dilute blood and concentrated urine.<sup>48</sup> The opposite disorder (diabetes insipidus, which causes hyperosmolar blood and dilute urine) is caused by much more common LOF mutations in V2R that result in misfolding.<sup>49</sup>

A third general mechanism by which nsSNVs promote disease is promotion of toxic protein aggregation.<sup>50–52</sup> The “toxic gain of function” associated with these aggregates is typically unrelated to the native function of the affected protein. This seems to be the case for the amyloid- $\beta$  polypeptides of Alzheimer’s disease, which form both soluble oligomers that are toxic to neurons and insoluble amyloid aggregates that induce chronic inflammation<sup>53,54</sup> in brain tissue. The immediate precursor of the amyloid- $\beta$  polypeptides is C99, the 99-residue C-terminal domain of the amyloid precursor protein. Some familial Alzheimer’s disease mutations in C99 alter its site specificity for cleavage by  $\gamma$ -secretase, increasing the production ratio between the longer and more toxic forms of the amyloid- $\beta$  polypeptide relative to shorter and less toxic forms.<sup>55</sup> Other inherited mutations in C99, located in its amyloid- $\beta$  domain, make the released amyloid- $\beta$  more prone to aggregate.<sup>55</sup> Mutations can also promote toxic misfolding through indirect mechanisms. For example, it is generally thought that familial AD mutations in  $\gamma$ -secretase alter its scissile site specificity in C99 so that the production ratio of long to short forms of amyloid- $\beta$  is increased.<sup>56</sup>

Finally, it should not be overlooked that other genetic variations besides nsSNVs can result in protein LOF, GOF, or toxic aggregates. Some genes may be completely deleted or inactivated by variations that alter their transcription, while for other genes, protein overdosage may occur when there are more than the usual two copies of the gene or when the gene is subject to aberrantly upregulated transcription.<sup>57</sup> While single-site mutations in PMP22 are one cause of Charcot-Marie-Tooth disease, this disorder more commonly results from trisomy (a third copy) of the wild-type gene, resulting in overexpression of wild-type PMP22.<sup>58</sup>

Mutations that alter either the chaperone or degradative components of cellular protein folding quality control systems can have adverse impacts on the folding of specific client proteins or, more generally, on cellular proteostasis.<sup>59,60</sup> The introduction of stop codons into protein open reading frames leads to the production of truncated proteins that may fail to function and possibly misfold, but alternatively may be functional but dysregulated (GOF).<sup>61,62</sup> Frameshift mutations lead to proteins with partial wild-type sequence, followed by incorrect amino acids. Chromosomal translocation can lead to the generation of fusion proteins composed of two normally independent proteins (or fragments thereof), sometimes having a highly toxic or aberrantly proliferative function.<sup>63</sup>

## ■ DETERMINING OR PREDICTING THE IMPACT OF GENETIC VARIATIONS ON PROTEINS

Much effort is already being dedicated to developing the infrastructure of databases, methods, protocols, training, counseling approaches, actuarial resources, bioethics, and laws that are needed for personalized genomic information to be widely employed in patient treatment (cf. refs 64–66). Molecular bioscientists such as biochemists, biophysicists, and structural biologists are already making important contributions to this process<sup>67</sup> (e.g., see lists of already available online resources in refs 43 and 68–70), which may well span decades. It is to be hoped that such investigators will make major contributions to the very important task of assessing *exactly how* genetic variations impact biomolecular folding, structure, function, and interactions. This information will often be essential to the practice of personalized medicine. For example, there is compelling need to distinguish between true disease-promoting genetic variations and neutral variations to avoid treatment of healthy patients who should therefore not be subjected to expensive, unhelpful, and potentially harmful and expensive pre-emptive treatment. Consider also a disease such as cystic fibrosis for which there are more than 1000 different gene variations known to result in LOF of the CFTR protein.<sup>71</sup> It is clear that different mutations can induce LOF through different mechanisms, some (such as the common  $\Delta F508$  mutant) by inducing misfolding and others by altering channel properties within correctly folded protein. Given that CF therapeutic drugs are in various states of development that address different classes of potential defects in the CFTR protein,<sup>72,73</sup> it may eventually be possible to mechanistically classify each CFTR disease mutation to ensure correct therapeutic matching for each specific CFTR mutation. Given that most drugs have unwanted off-target side effects, there is an added imperative to avoid treatment of unresponsive patients.

The ability to predict with great accuracy and mechanistic detail the possible outcome of nsSNVs in terms of protein LOF, GOF, altered protein–protein interactions, or toxic aggregation would be a great resource. Indeed, there are already numerous algorithms available, often deployed in the form of publicly accessible web servers that make such predictions.<sup>69,70,74–83</sup> Given that destabilization of proteins leading to LOF may be the most common mechanism by which genetic variations promote disease, it is not surprising that a number of these algorithms are devoted to prediction of mutation-induced protein stability changes. As of late 2014, the authors easily identified several dozen online programs/servers that predict protein stability either as a primary function or as a trait that is factored into overall predictions of whether a particular genetic variation is pathogenic. The very best of the stability predictive algorithms, which can be of formidable sophistication, are able to correctly and quantitatively predict mutation-induced changes in protein stability roughly 85% of the time for *relatively small proteins of known three-dimensional structure*.<sup>74,84–89</sup> However, this success rate drops by ~15% when the protein’s three-dimensional structure is not known. In this case, algorithms must rely instead on homology models, sequence conservation patterns, and/or other variables.<sup>78</sup> Moreover, the benchmarking used to evaluate the success of these predictions usually lacks data for multidomain proteins, which are especially common in higher organisms. The same deficit holds for membrane proteins, which comprise roughly

20% of all proteins and upward of 50% of the targets of currently used drugs. This problem is fundamental as the rules that govern protein stability may well be altered for interfacial residues in multidomain proteins<sup>90–92</sup> and transmembrane segments in membrane proteins.<sup>93–96</sup> Therefore, aforementioned methods that are optimized for small soluble proteins would be expected to display decreased accuracy. There remains a compelling need for additional experimental studies of protein stability, especially for membrane proteins and multidomain proteins, for which there may not currently be sufficient data available even to validate and benchmark stability prediction methods.

While most previous effort has been dedicated to targeted studies of particularly common or representative disease variants of a given protein (such as the  $\Delta F508$  form of CFTR), experimental studies of modest panels of disease mutant forms of a given protein have been described (cf. refs 97–104). One might flinch at the thought of conducting experimental studies on hundreds of mutant forms a single protein, such as the more than 500 known retinitis pigmentosa-linked mutant forms of the ABCA4 flippase<sup>105</sup> or the more than 15000 cancer-related variants of the p53 tumor suppressor.<sup>106</sup> However, the potential power and efficiency of high-throughput experimental methods should not be overlooked when considering how best to test the impact of potential disease-linked sequence variations on protein function, structure, stability, intermolecular interactions, and tendency to aggregate. The high value attached to the availability of a high-quality experimental structure for the protein of interest or a close homologue should also be emphasized, given that our ability to predict mutation-induced perturbations of normal protein traits is often dramatically enhanced by the availability of such a structure.<sup>68,107</sup>

It has recently been estimated that 35% of all residues in the human proteome have been structurally characterized, through experimental structural determination or being inferred from modeling to homologous protein templates of known experimental structure.<sup>108,109</sup> This highlights the tremendous frontier of structural space in the human proteome that remains to be characterized. At the same time, the numerous available experimental structures and homology models for human proteins represent an immediate resource for use in assessing VUS.

As of mid-2015, there are still numerous human proteins for which biochemical and biological function and regulation remain poorly understood, a fact that will confound the degree to which genome variations affecting these proteins can be analyzed for personal medical purposes. The continued need for basic science studies of such proteins reflects the unwavering importance of basic biological research in providing the foundations of molecular medicine.

An additional category of insight needed for personalized medicine to which biochemists and biophysicists can contribute is in assessing how partial or complete GOF or LOF for a variant protein alters the biochemical, cellular, and physiological pathways associated with that protein and how the dysfunction or dysregulation of those pathways impacts cellular and, ultimately, organismal fitness.<sup>110,111</sup> Because a great many of these pathways are not isolated but are integrated within complex webs, this realm of analysis will overlap with that of systems biology.

## ■ CONFOUNDING FACTORS IN TRANSLATING RESULTS OF PROTEIN STUDIES INTO PERSONALIZED MEDICAL DECISION MAKING

When considering results for variant proteins, it can be very difficult to predict whether correct assessment of protein function informs on whether a variation actually predisposes the human subject to a related disease.<sup>112,113</sup> Indeed, it is now appreciated that many “disease mutations” compiled in databases such as HGMD are probably neutral variations or of low penetrance, meaning they result in disease in only a fraction of the mutation carriers.<sup>114</sup> There are a variety of confounding factors, even for mutations associated with “simple” monogenic Mendelian diseases. Among these are data, noted earlier, that show that many proteins seem to be dispensable, at least under some conditions. It may be that life has to some degree evolved so as to be robust by generating molecules with redundant functions and compensatory mechanisms to avoid placing too much importance on single gene products.<sup>115</sup> Moreover, in any individual, each genetic variation occurs within the context of the thousands of other genetic variations, some of which may exert a compensatory effect that nullifies the possible physiological consequences of GOF or LOF in another protein. Perhaps this is the case for the dementia-free centenarian who was recently found to be homozygous for the Alzheimer’s-promoting apolipoprotein E  $\epsilon 4$  variant.<sup>116</sup> The complex genetics of some disorders, where disease occurs only when multiple disease-predisposing variations collude, makes such maladies difficult to predict or mechanistically dissect.<sup>117</sup> In complex diseases such as type II diabetes, environmental factors such as diet, exercise, gender, smoking, lifetime exposure to toxins or infectious agents, age, and prior medical history can work, variously, either to reinforce or to offset disease-promoting genetic factors.<sup>118</sup> Epigenetics can also complicate the linkage of genome variations with personal medical traits.<sup>119</sup> Finally, because protein expression levels can be highly variable from cell to cell, tissue to tissue, and patient to patient, the actual degree of net physiological GOF or LOF associated with an otherwise well-characterized variant protein may be very difficult to predict in an individual.<sup>120</sup>

## ■ THE DEVELOPING INTERFACE AMONG BIOMOLECULAR SCIENCE, BIOINFORMATICS, AND PERSONALIZED MEDICINE

In light of the contributions that can be made by biochemists and biophysicists to the knowledge infrastructure that will support personalized medicine, we would like to offer a couple suggestions regarding how this interface can be enhanced.

Excellent communication between bioinformaticists and biomolecular scientists will increasingly be required. It will be critical to develop methodologies that combine bioinformatics capabilities in large scale data analysis, machine learning, and biostatistics with data gleaned from biochemical and biophysical experiments and simulations. Bioinformaticists are often well-equipped for tapping into the databases, theory, and information provided by the studies of both geneticists and biomolecular scientists. This is admirably reflected, for example, by the sophistication of some of the available protein stability and pathogenicity prediction methods. However, basic scientists sometimes do not have the required training in statistics and elementary computer programming needed to access and fully understand/assess many bioinformatic analyses,

even when the topic (such as protein stability assessment) is familiar. This deficit might well be addressed by injection of a modest dose of formal training in statistics, computer programming, and bioinformatics into graduate programs in the basic biological/biomedical sciences. Along these same lines, a modicum of up-to-date didactic training in the areas of genetics and genomics should perhaps also be regarded as an essential curricular objective.

There also is the continued need to expand the interface between the basic biomedical sciences and the engineering-tinged discipline of systems biology. This is essential to the continued development of methods to assess whether and how a particular variant protein will actually result in disease. Moreover, systems approaches will be required to determine the degree to which multiple genetic variations synergistically promote health problems or alter pharmacogenomic profiles in ways that cannot be predicted on the basis of summing the impacts of the same variants in isolation. The relationship between a single protein and the fitness of the host organism may sometimes be exceedingly complex. At the same time, all diseases arise from problems that are, ultimately, molecular in origin.

## CONCLUSIONS

Personalized medicine is not a new concept but is being transformed by the advent of widespread exome/genome sequencing. Some of the paradigms for how genome information will be employed are already established, such as the fact that genetics dictates that the optimal treatment for some patients is sometimes different than for others. It is also clear that analysis of genome information will allow rational preemptive treatment of disease prior to the symptoms. Genetic information may, in favorable cases, be used to relieve the anxiety of individuals who fear a particular disorder because of family history or other factors. An interesting emerging area of genetic analysis involves the search for genetic factors that are *protective*, lowering the risk for various diseases and/or increasing life expectancy.<sup>121</sup>

Beyond patient care, information from whole genome analysis may well come to be routinely factored into important and highly personal decisions such as those associated with human reproduction. This highlights currently unresolved issues such as the question of how much patients should be told about their genomes and those of family members, who should have access to personal genomic information, and how this information can lawfully and ethically be used in decision making. One wonders whether the impact of the genome on everyday life is currently where personal computers and the computer networks were circa 1980. At that time, it was apparent that the electronic innovations were here to stay, but few could foresee the extraordinary degree to which information technology has come to impact society only 35 years later. It seems probable that widespread genome sequencing will ultimately shape civilization in ways that extend well beyond what we can foresee. There are and will continue to be numerous opportunities for biochemists, biophysicists, structural biologists, and other basic scientists to generate knowledge and approaches that will be critical to the successful ongoing deployment of the genomics revolution. In closing, we suggest that while this revolution will have immediate impact, it is likely to unfold over decades, maybe even centuries. Societal expectations in terms of imminent widespread improvements in healthcare should be tempered. In this regard, it may be helpful

to ponder a 2001 observation by Sir David Weatherall, a pioneer in classical genetic studies of hemoglobin disorders such as anemias and thalassemias:<sup>122</sup>

*"The one regret is that the remarkable advances in human genetics, and in the evolution of molecular medicine in general...have done little to help patients with these distressing diseases... The lack of a more definitive approach to the cure of the haemoglobin disorders, despite so much knowledge about their molecular pathology, should not surprise us. The story of the advancement of medical practice over the last century tells us that there is always a long lag period before developments in the basic sciences reach the clinic."*

## AUTHOR INFORMATION

### Corresponding Author

\*E-mail: chuck.sanders@vanderbilt.edu. Telephone: (615) 936-3756. Fax: (615) 936-2211.

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