



Data-driven supervised learning of a viral protease specificity landscape from deep sequencing and molecular simulations

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Biophysical interactions between proteins and peptides are key determinants of molecular recognition specificity landscapes. However, an understanding of how molecular structure and residue-level energetics at protein–peptide interfaces shape these landscapes remains elusive. We combine information from yeast-based library screening, next-generation sequencing, and structure-based modeling in a supervised machine learning approach to report the comprehensive sequence–energetics–function mapping of the specificity landscape of the hepatitis C virus (HCV) NS3/4A protease, whose function—site-specific cleavages of the viral polyprotein—is a key determinant of viral fitness. We screened a library of substrates in which five residue positions were randomized and measured cleavability of ~30,000 substrates (~1% of the library) using yeast display and fluorescence-activated cell sorting followed by deep sequencing. Structure-based models of a subset of experimentally derived sequences were used in a supervised learning procedure to train a support vector machine to predict the cleavability of 3.2 million substrate variants by the HCV protease. The resulting landscape allows identification of previously unidentified HCV protease substrates, and graph-theoretic analyses reveal extensive clustering of cleavable and uncleavable motifs in sequence space. Specificity landscapes of known drug-resistant variants are similarly clustered. The described approach should enable the elucidation and redesign of specificity landscapes of a wide variety of proteases, including human-origin enzymes. Our results also suggest a possible role for residue-level energetics in shaping plateau-like functional landscapes predicted from viral quasispecies theory.

protease | sequence–function mapping | substrate specificity | machine learning | molecular modeling

Predicting the impact of genetic diversity on molecular recognition specificity of enzymes is of fundamental importance in molecular biology and also has implications for the design of novel enzymes with controllable molecular recognition properties. The balance between mutational tolerance and functional plasticity is encapsulated in the notion of mutational landscapes (1), which are high-dimensional maps that relate the function of individual biomolecular variants to their functional and/or evolutionary fitness (2, 3). Recent empirically determined sequence–function mappings of proteins (4–12) have enabled the partial construction of mutational landscapes. Typically, sequence–function mapping of proteins and protein–protein interactions involves partial enumeration of the possible sequence diversity (for example, all single mutations and a subset of double mutations at a large number of protein residue positions) and high-throughput functional evaluation coupled with deep sequencing (13–17). The astronomical size of sequence space, however, limits comprehensive elucidation of sequence–function landscapes with any one experimental approach. Computational biophysical methods may, in principle, assist in creation and analysis of functional and fitness landscapes (18, 19). Indeed, mutational landscapes

of simple protein models, such as lattice models, have been extensively investigated using biophysical evolutionary theory and computational simulations (20–30), and connections with population genetics theories have been discovered (20, 31, 32). While pioneering and crucial insights have been obtained in these studies, chemically realistic atomic resolution structure-based elucidation of functional landscapes has remained elusive.

The genomes of several RNA viruses, e.g., human immunodeficiency virus (HIV) and hepatitis C virus (HCV), encode polyproteins, which are processed posttranslationally by viral proteases during maturation (33). The activity of HCV NS3 protease is key for viral maturation, as it cleaves exclusively at four specific sites in the viral polyprotein (Fig. 1A) to release individual nonstructural proteins (34) and also mediates inactivation of key human immunity proteins (35). The cleavage specificity of the protease is thus a key determinant of viral fitness, and its proper functioning includes negative specificity—the lack of cleavage of noncanonical sites on the viral protein and most host cell proteins—but how and whether these features are encoded in the protease–substrate interactions at a molecular level is currently unknown. Furthermore, while RNA viruses

Significance

Substrate specificity landscape of a protease enzyme is the set of all substrate sequences that are recognized/cut (and, as importantly, not recognized/cut) by the enzyme. Accurate and rapid elucidation of these landscapes for any given protease is key for the design of novel targeted proteases to prevent unwarranted off-target cleavage, and provides insight into the functional robustness of naturally occurring proteases. We developed a structure-guided approach for predicting protease substrate specificity landscapes, in which data from experiments in yeast and molecular simulations are combined using machine learning. Using this approach, we comprehensively map the sequence–energetics–function landscape of the hepatitis C virus NS3/4A protease and its drug-resistant variants.

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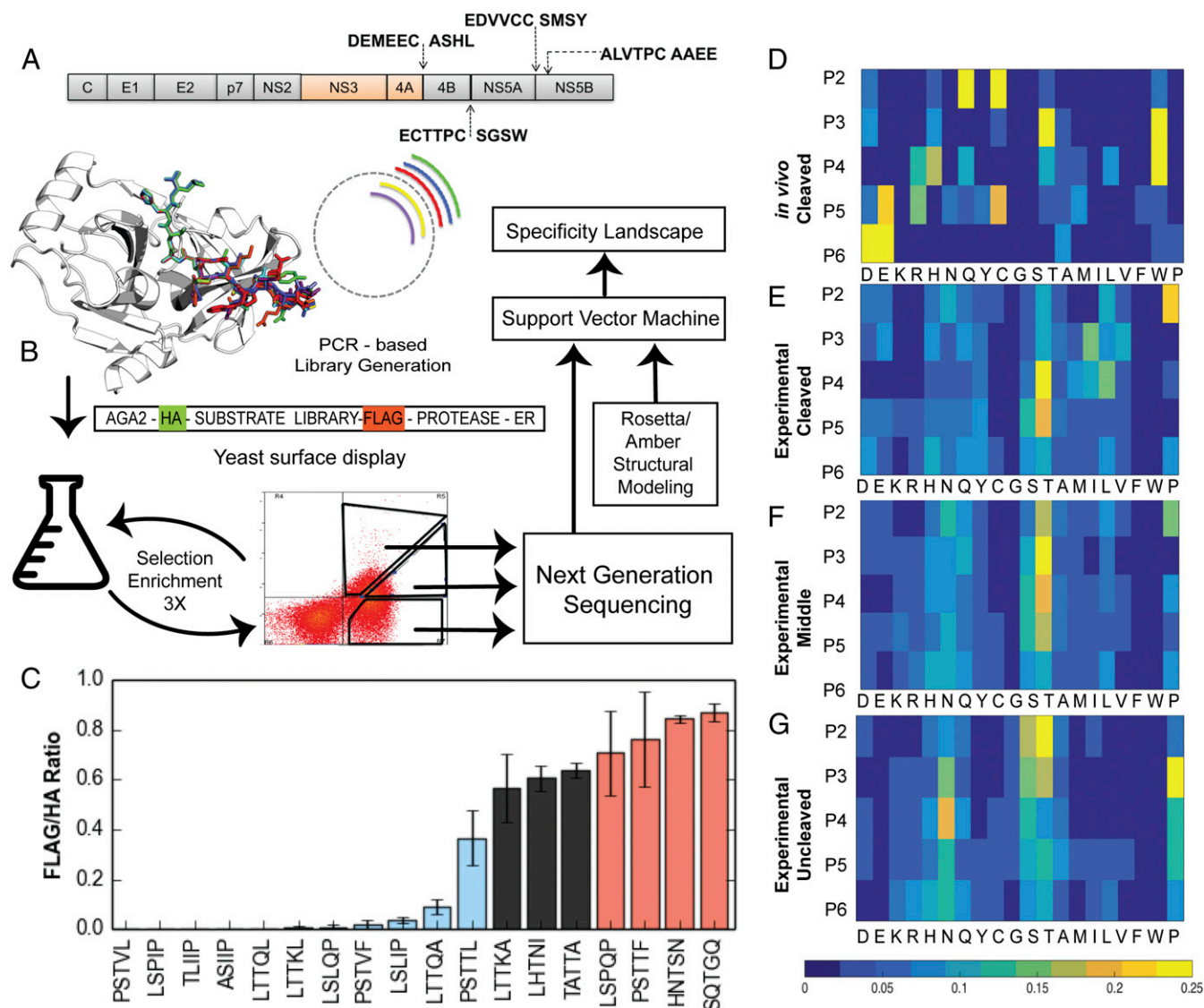


Fig. 1. Overview of workflow and results. (A) The HCV viral polyprotein depicting marked biological cleavage sites for the HCV NS3 protease. (B) Overview of the experimental and computational workflow. The construct shown (LY104) is the vector used for testing in the yeast-based assay. The substrate was cloned in the region between the FLAG and HA signaling tags. Extent of cleavage was measured as a ratio of FLAG/HA, with a ratio of 1 indicating that the substrate was uncleaved and a ratio of 0 indicating that the substrate was cleaved. (C) Validation of FACS gates for cleaved (blue), partially cleaved (black), and uncleaved (red) sequences using the yeast surface display assay. (D–G) Heatmaps showing per-position amino acid frequencies for (D) sequences taken from in vivo samples of HCV patients (8,726 sequences) compared with (E) sequences determined by our assay as cleaved (7,472 sequences), (F) sequences determined by our assay as partially cleaved (8,737 sequences), and (G) sequences determined by our assay as uncleaved (14,702 sequences).

such as HCV are believed to be under strong purifying selection against nonsynonymous mutations (36–38), due to the extremely high error rates of the associated RNA polymerases (39–41), these viruses can exist as a population of variants called quasispecies (42, 43) even within a single host individual (44). Indeed, spontaneous emergence of diverse mutations (including drug-resistant mutations) was demonstrated in continuous evolution studies of the protease (45) and in viral replicon assays coupled to ultradeep sequencing (46). Can molecular interaction fidelity be maintained in the face of a large mutational load, and what, if any, are the limits imposed on the allowed genetic diversity by the underlying molecular interactions? The degeneracy of the genetic code, the thermodynamic and kinetic stabilities of RNA and proteins, and the presence of molecular chaperones may all contribute to the mutational robustness of the structures of individual viral biomolecules under a high mutational load (41).

However, the mechanism by which key viral protein-based interactions—for example, protease–substrate interactions critical for viral propagation—may encode “fuzziness” (47) is not well understood.

Here, we use a combination of experimental (biochemical) and computational techniques to elucidate the specificity landscape of the interaction between HCV NS3/4A protease enzyme and its substrates. Using yeast surface display, next-generation sequencing, and a machine learning approach which combines features from experimental data and atomistic computational simulations (utilizing the Rosetta and Amber force fields) that we recently developed (48, 49), we construct the specificity landscapes (with cleavability assignments made for 3.2 million substrate pentapeptide sequences) of the HCV NS3/4A protease and three of its known drug-resistant variants (50).

Results

We mapped the protease–substrate interaction landscape for the HCV NS3/4A protease by considering all possible pentapeptide sequence combinations at positions P6 through P2 (Schechter and Berger nomenclature) (51) in the substrate. Positions P1 and P1', between which the scissile bond is present, were maintained as the canonical C and A, respectively. In the remainder of this paper, we refer to individual pentapeptide patterns (e.g., the canonical cleavage sites DEMEE, EDVVC, ECTTP, and ALVTP) and omit the identity of the P1, P1' residues. By mapping the substrate diversity for HCV NS3/4A wild-type protease as well as three drug-resistant protease mutants, we explore the interaction landscape for both protease and substrate diversity.

Exploration of the (P6-P2) Specificity Landscape of the HCV NS3/4A Protease Reveals a Diverse Specificity Profile. To mimic the viral intrachain arrangement of substrate libraries and the protease, we utilized a modified version of the assay described by Iverson and coworkers (52) (Fig. 1B and *SI Appendix, Fig. S1A*). A mutagenic library was created incorporating degenerate codons at P6 to P2 specificity defining substrate positions (53, 54). In this assay, substrates are transported to the surface of yeast cells in a cleavage-dependent manner: The degree of cleavage is estimated by measuring the relative levels of substrate-flanking FLAG and HA tags using fluorescent-labeled antibodies. We have previously used this assay to test known and novel substrates of the HCV protease (48). A first round of yeast surface display assay and Fluorescence Assisted Cell Sorting (FACS) was performed with an inactive protease variant (S139A) to select for high expression of library variants, remove sequences containing stop codons in the substrate region, and deplete substrate sequences that are cleaved by yeast endoplasmic reticulum (ER) proteases (55).

The resulting substrate variants from the preselection were subjected to rounds of yeast surface display assay and FACS with an active-protease-containing construct to select cleaved, partially cleaved, and uncleaved variants using three sorting gates (Fig. 1B), based on the relative levels of anti-HA and anti-FLAG fluorescence values. The FLAG/HA ratio ranges between 0 for completely cleaved substrates and 1 for completely uncleaved substrates. Sorting gates were defined based on the distribution of populations observed for known cleaved and uncleaved sequences (48). This procedure was coupled with rounds of growth and selection to improve the signal-to-noise ratio for variants in each pool. Specificity profiles of the unselected population and isolated functional variants were determined using next-generation sequencing technology (Illumina NextSeq).

We identified a total of ~379,000 unique sequences in the background pool corresponding to ~12% of the possible amino acid diversity (3.2 million). Analysis of technical replicates as well as the overlap between the sequence pools was used to determine a count threshold (normalized count of 11) to remove noise from the sequencing data (*SI Appendix, Supplementary Methods and Fig. S2*). Based on these criteria, we identified 7,472, 8,737, and 14,702 unique pentapeptide sequences in the cleaved, partially cleaved, and uncleaved pools, respectively. In parallel, we performed Rosetta simulations on all 3.2 million sequences in the P6 to P2 region to determine energetic features of the protease–substrate models. We then used a support vector machine (SVM) in a supervised machine learning approach to predict the complete protease–substrate interaction landscape that incorporated sequence information procured from the aforementioned library and Rosetta-generated energetic features (Fig. 1B).

Several novel substrates identified from the three variant populations were tested as clonal populations in the yeast surface display assay system (Fig. 1C and *SI Appendix, Fig. S3*) to validate that individual sequences fall into the gates used for selection from the library (*SI Appendix, Fig. S4 A–C*). Some

sequences were chosen because they were either different from any of the canonical cleavage sites at three or more positions (e.g., PSTVL) or implicated in epistatic networks (e.g., LSLIP; see *SI Appendix, Supplementary Discussion*), or were identified using a stricter enrichment threshold as described in *Materials and Methods*. A subset of these sequences was also tested in vitro to ensure that the cleavage properties observed in the yeast system were reproduced with purified protease and substrates (*SI Appendix, Fig. S4D*). The process of transforming quantitative cleavage efficiencies to discrete functional pools of cleaved, partially cleaved, and uncleaved substrates by definition involves a loss of information regarding relative binding efficiencies of substrates within each pool; however, clear differences in FLAG/HA ratio between the three categorizations are apparent (Fig. 1C).

We next analyzed the cleaved, partially cleaved, and uncleaved sequence sets obtained from deep sequencing. The cleaved specificity profile has greater diversity than the substrates identified from viral genomes sequenced from patient populations (*SI Appendix, Supplementary Methods and Fig. 1D*). For example, we observe that a more diverse subset of amino acids is tolerated at substrate positions P6 and P5 in our cleaved and partially cleaved pools (Fig. 1E and F) whereas the patient-derived genomes display a high enrichment of Asp and Glu specifically at these positions. The relative abundance (compared with all 20 amino acids, normalized to 1) of Asp at P6 is 0.08 in the cleaved population compared with 0.03 in partially cleaved and 0.02 in uncleaved populations; the relative abundance of Glu at position P5 in the cleaved population is 0.06 compared with 0.03 and 0.01 in partially cleaved and uncleaved populations. In contrast, these abundances are P6-Asp-0.47, and P5-Glu-0.35 in the patient-derived genome data. Overall, the viral genome-derived specificity profile has a lower information content (i.e., lower diversity) than that of the cleaved specificity profile obtained using our method (1.92 bits vs. 3.85 bits, out of a maximum 4.32 bits), pointing to the relative flatness of the extended substrate specificity profile (*SI Appendix, Table S10*) that can be recognized by the protease.

Strikingly, even though the actual sequences in each pool are chosen to be distinct (*SI Appendix, Fig. S9B*), the overall specificity profiles of the three sequence sets (cleaved, partially cleaved, and uncleaved) are similar (Fig. 1E–G): The cosine similarities (ranges between 0 and 1, dissimilar to similar) of cleaved to uncleaved, cleaved to partially cleaved, and partially cleaved to uncleaved are 0.61, 0.86, and 0.84, respectively. Thus, cleaved sequences are more similar overall to partially cleaved sequences than to uncleaved ones. Additionally, there are several differences in the enrichment of certain residues at each position in the three pools. For example, we found prolines enriched at position P2 in the cleaved (relative abundance of 0.2) and partially cleaved populations (relative abundance of 0.14) compared with the uncleaved set (relative abundance 0.04), which, in turn, prefers proline at P3 (0.23 relative abundance). These trends correspond well with the fact that two out of four canonical cleaved sequences have proline at P2 (ECTTP and ALVTP). While some of the above trends are also reflected in the sequences we tested during method validation (Fig. 1C), it is evident that overall sequence composition or individual positional enrichments cannot be directly used to predict the pool assignments of individual sequences. For example, His is somewhat enriched (relative abundance of 0.09 in cleaved population) at P6 in the cleaved sequence pool, but the sequence HNTSN is experimentally determined to be in the uncleaved pool (Fig. 1C and *SI Appendix, Fig. S3*). We conclude that interactions between amino acids at various substrate positions (mediated possibly through interaction networks in the protease) influence the cleavability, thereby motivating the need for an analysis of the specificity landscape using properties of whole pentapeptide sequences and models of their complexes with the protease.

opposed to cleaved < uncleaved << partially cleaved in the experimentally derived graph, further highlighting the relationship between mutational distance and function preservation.

Energetic Features Derived from Rosetta Modeling Enable Reconstruction of the Complete Protease–Pentapeptide Substrate Landscape. While the experimentally derived populations of the cleaved, partially cleaved, and uncleaved sequences display striking clustering patterns in sequence space, they include a small fraction of the entire sequence diversity in the P6 to P2 region (3.2 million sequences). To predict cleavability of all possible 3.2 million sequences in the interaction landscape, we used an SVM method that we developed previously (48). Briefly, each sequence was threaded onto a bound complex based on a modeled near-attack conformation, and the complex was then relaxed to accommodate the substrate while maintaining favorable catalytic geometry. Energy evaluation of each of the 3.2 million complexes was performed using Rosetta and Amber simulation packages.

A binary classification (cleaved/uncleaved) SVM was trained on two sets of sequences. The first set was a subset of experimentally identified sequences that were identified using more stringent criteria than the original set of experimentally categorized sequences (1,817 cleaved and 3,605 uncleaved sequences; see *Materials and Methods* for details). The second set included 196 cleaved sequences and 1,943 uncleaved sequences identified by Shiryayev et al. (59), and both sets together yielded 7,338 unique sequences. Training features consisted of structure-based features (energies of interaction) and sequence-based features (see *Materials and Methods* and *SI Appendix, Fig. S7A*). We initially cross-validated the SVM on the training set using an 80:20 split with 100 iterations, which yielded an average AUROC (area under receiver operating characteristic) of 0.96 (*SI Appendix, Fig. S7D*) indicating high recapitulation of training data (perfect performance would lead to an AUROC of 1). We then used the SVM to predict cleaved and uncleaved labels for the remaining 3,192,658 sequences. These predictions have a precision of 0.95 at a recall level of 0.89 for an overall accuracy of 0.95 (*SI Appendix, Fig. S7C*) for the experimentally derived assignments that were left out of the training set (5,906 cleaved sequences and 11,087 uncleaved sequences). For testing of SVM predictions, we selected six sequences that were not found within the experimentally defined set and had predicted distances greater than 2 from the SVM-calculated separation hyperplane. For all of the tested sequences, we find good agreement with the SVM-based predictions (Fig. 3A): The experimentally observed FLAG/HA ratios are ~ 0 and ~ 1 for the predicted cleaved and uncleaved test sequences,

respectively. To validate that the successful predictions are not localized to specific regions of sequence space, we visualized a subgraph of predicted cleaved sequences, present at a distance of >2 from the hyperplane constructed by the SVM learning procedure (*SI Appendix, Fig. S7B*). The experimentally identified cleaved sequences are distributed evenly across sequence clusters in the predicted cleaved population graph, further indicating that the machine learning procedure performs evenly across the sequence diversity of cleaved sequences.

Structural and Energetic Bases for Observed Specificity Patterns. Having obtained and validated predictions of cleavability by combining experimental and computational data using supervised learning, we turned to structural models of protease–substrate complexes to obtain insight into the underlying structural basis of observed specificity patterns. For example, a comparative analysis of the partially cleaved substrate “TATTA” and canonical substrate “EDVVC” reveals that the former, composed of small residues, does not completely occupy the substrate cavity volume, whereas EDVVC occupies the entire cavity (Fig. 3B and C). The lack of voids at the interface and several hydrogen bonds formed by the canonical substrate lead to better binding [binding interaction energy = -80.2 Rosetta energy units (Reu), as opposed to -77.5 Reu for TATTA], resulting in better cleavage for this substrate. Apart from sidechain-based interaction patterns, models also capture backbone conformational changes that affect the orientation of the substrate in the active site. For example, in the model corresponding to the sequence RPGPG (uncleaved), the proline present at P3 in RPGPG (Fig. 3D) bends the peptide chain away from the protease, resulting in breaking of the crucial backbone hydrogen bond patterns that are characteristic of protease–substrate interactions (60).

Connectivity Properties of the Experimentally Determined and Computationally Reconstructed Landscapes. Having computed the entire P6 to P2 specificity landscape, we next examined the connectivity patterns between cleaved and uncleaved sequences in this reconstructed landscape. As with the experimentally determined landscape, the reconstructed landscape also shows evidence of clustering between cleaved and uncleaved nodes (*SI Appendix, Fig. S5 H and I*): Most neighbors of (un)cleaved nodes are (un)cleaved. While the clustering properties are similar, the lack of a partially cleaved category in the computationally derived dataset and incompleteness of the experimental dataset also lead to some differences in the measured connectivity properties of these graphs (*SI Appendix, Fig. S5 F and G*).

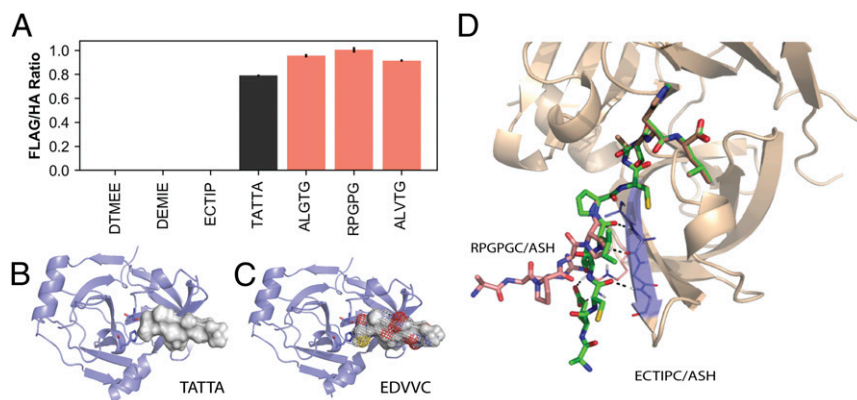


Fig. 3. Structural modeling provides insights into basis for cleavage. (A) Validation assay performed for three predicted cleaved, one partially cleaved, and three uncleaved sequences using a yeast surface display-based technique. (B and C) The volume occupied by (B) TATTA and (C) EDVVC. EDVVC occupies an optimal volume, making good contacts with the protease residue sidechains. TATTA fits in the available space but does not make optimal contacts, thus resulting in suboptimal interaction energetics making TATTA a suboptimal substrate. (D) Structure of two models, ECTIP (cleaved) and RPGPG (uncleaved).

ability to be reachable from/to canonical sequences that are at least three amino acid substitutions away in sequence space.

We note that the nature of the qualitative bins within which substrates are classified in our analyses precludes evaluating the evolutionary likelihood of individual mutational trajectories. In particular, paths that appear to be completely viable (i.e., contain no uncleaved nodes) may contain nodes that are of lower catalytic efficiencies and/or fitness than the starting point, and therefore are less likely to be traversed. Therefore, our analysis considers all paths between nodes and focuses on relative nonviable path fraction, e.g., nonviable path fraction between canonical and other canonical substrates vs. nonviable path fraction between canonical and novel substrates. An uncleaved node in a path is guaranteed to make it nonviable, but demonstrating that two nodes which have 100% viable paths between themselves form a neutral network would require more quantitative characterization of the catalytic efficiencies and molecular fitness of all of the nodes.

Specificity Landscapes of Drug-Resistant Protease Variants. As the NS3/4A protease plays a key role in the viral assembly and maturation process, it is a target for therapeutics that aim at neutralizing viral activity. However, due to prevalence of quasi-species that are lurking at low levels in the population (61),

several viral variants get exposed to the drug. Some of these develop resistance, and propagate to form drug-resistant mutants (DRMs). To investigate how drug-resistant variants of the protease affect the mutational robustness, we explored the specificity landscape for three DRMs: A156T, D168A, and R155K/A156T/D168A (Fig. 5). We find that the DRM proteases have similar specificity profiles to those of each other and to that of the wild-type protease, with cosine similarities of DRM specificity profiles to wild-type specificity profiles that range between 0.89 and 0.99 (Fig. 5 C–F). However, the sequences cleaved by the variant and wild-type proteases are distinct, with an 8 to 21% overlap in sequences between the each of the various DRMs and wild-type proteases (SI Appendix, Fig. S9 I and J). Upon comparing the graphical properties of the specificity landscapes of the various protease variants, we observe that substrates that are experimentally detected in the cleaved pools of a greater number of protease variants (more recognizable) are more reachable (higher pageranks) and more connected (higher degree) in each graph (Fig. 5 G and H). These data indicate that more recognizable substrates appear to be more robust to changes in the protease. Thus, functional clustering in sequence space appears to be a robust feature of the molecular recognition between the HCV NS3/4 protease variants and their substrates.

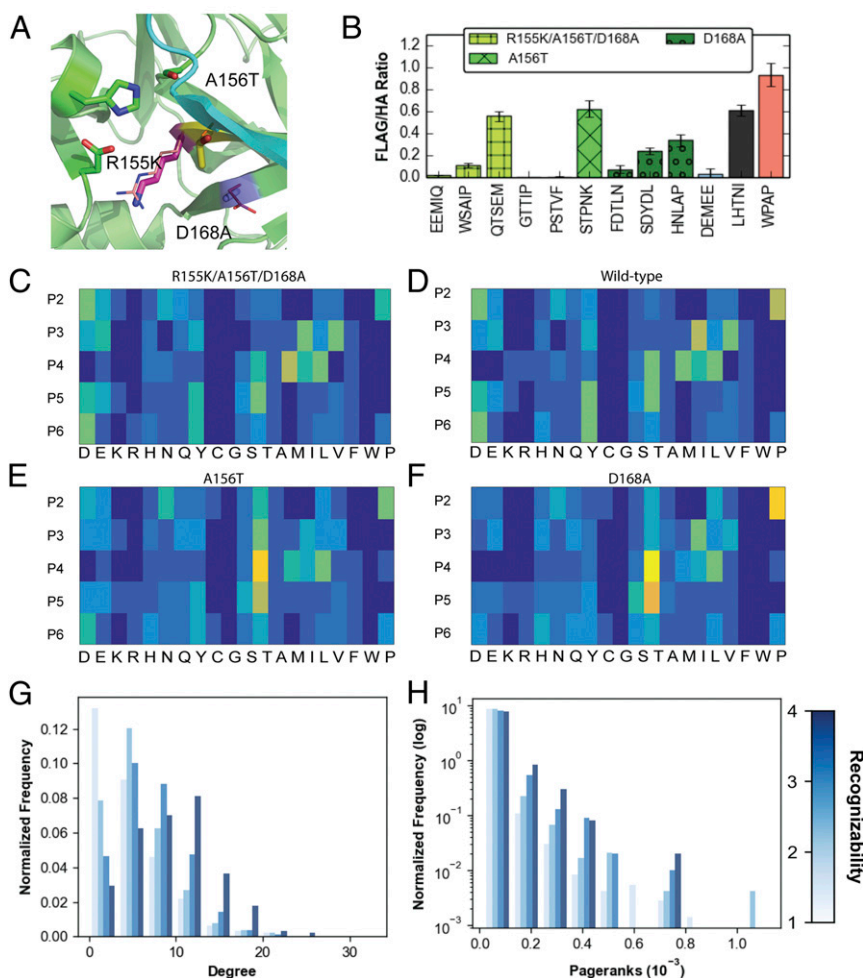


Fig. 5. DRMs have similar structures and similar specificity profiles. (A) Drug-resistant variant structures. Mutations are outlined in sticks, and WT residues are outlined in lines. Active site residues are represented as green sticks. (B) Validation assay performed using yeast surface display for each of the mutants. (C–F) Mutant cleaved sequence specificity profiles for the (C) triple mutant, (F) D168A, (E) A156T, and (D) wild type, showing that the mutants have very similar specificity profiles with slight variation compared with the WT. (G and H) Substrate sequences that are recognized by a greater number of variants have higher (G) degrees and (H) pageranks.

Discussion

We combined information gleaned from library screening in yeast, deep sequencing, and structure-based modeling, using a machine learning framework to delineate the protease–substrate interaction landscape of HCV NS3/4A protease. These results provide atomic-resolution insight into the bases for both positive and negative specificity. We used a yeast surface display-based assay that relies on the cleavage of the substrate region in the ER of yeast followed by cell sorting into gates and deep sequencing. We note that our assay is qualitative, and does not permit association of the detected signal from deep sequencing with quantitative cleavability of substrates. Indeed, while we have validated that assignments to the three different pools are accurate with at least ~20 individual sequences, the identified cleaved and partially cleaved substrates may represent a range of catalytic efficiencies within their pools. On the other hand, the assay construct with the protease and substrate on the same chain is a good representation of the situation in the virus, where the substrates of the protease are part of the same polyprotein (although both *cis* and *trans* cleavages occur), leading to high effective concentrations of substrates ($[S] \gg K_M$) in vivo. Under these saturating conditions in the virus and in our assay, we argue that selectivity and catalytic efficiency are both determined to a great extent by the goodness of fit of various substrates in the protease active site (i.e., by the relative binding between the different substrates). Similarly, our SVM-based machine learning approach to combine experimental and computational data also is not without errors, showing a false-positive rate of ~5 to 10% on the experimental data. While we have validated several predictions on individual sequences (Figs. 1, 3, and 5), it is possible that some individual sequences may be mispredicted. However, the overall trends regarding the connectivity patterns observed for the entire landscape should be robust to the misprediction noise. Further ongoing development of the computational and experimental methods that we utilized is expected to help make the approach outlined here more quantitatively accurate.

We note that our graph theoretic analyses (Fig. 4), which draw from similar previous analyses (10, 62), report qualitatively on the local topology of the recognition specificity landscape, which is related to, but distinct from, a molecular fitness landscape. This difference arises as functional bins, whose boundaries are derived from experiments and within which substrates are classified (cleaved, partially cleaved, uncleaved), serve as a qualitative proxy for the catalytic efficiency and associated molecular fitness; binning necessarily flattens functional differences within a pool, and there is loss of information. Overall clustering and connectivity properties thus qualitatively describe the specificity landscape. On the molecular fitness landscape, the likelihood/fitness of individual mutational trajectories may depend quantitatively on the relative catalytic efficiency of traversed nodes. Definitive demonstration of specific network topologies (e.g., a neutral network) would require a more quantitative approach for molecular fitness characterization.

Our results provide a biophysical baseline for understanding how HCV protease substrates may sample genetic diversity while

maintaining function. For example, our analysis suggests that viral evolution occurring at the substrate sites in the polyprotein could also contribute to drug resistance. Due to the high interconnectedness of partially cleaved and fully cleaved clusters, novel sequences that are better substrates of drug-resistant variants may easily arise. This mode of substrate coevolution-based drug resistance has been observed in HIV-1 (63). At the same time, our analysis of the dominant HCV sequences obtained from patients suggests that the protease–substrate interactions may also contribute to negative selection (*SI Appendix, Supplementary Discussion*) and help limit the acquisition of heterogeneity. The molecular interaction between the protease and substrates, while key for viral survival, does not directly determine evolutionary fitness and is one of the many evolutionary forces at play, especially in the “wild” (64). Other factors such as the impact of genetic diversity on intrahost population size, stability, size and structure of the viral RNA genome, and interactions between the host and viral machineries and other environment-dependent factors are also important to consider while considering the evolution of HCV (46). Thus, apart from the connectivity properties of the specificity landscape (Fig. 4), a confluence of various factors, including the lack of sampling of genetic diversity by the virus in the wild and a relatively small number of genomes sequenced, may contribute to the lack of genetic diversity observed in patient-derived sequences of HCV protease substrates.

In summary, our exploration of a viral molecular specificity landscape uncovers novel specificities for the HCV NS3/4A protease. The developed specificity landscape enumeration approach is general, and combining experimental deep sequencing and structural modeling at a matching high throughput, followed by supervised machine learning, may be useful for elucidating a significantly larger space of sequence–function relationships for a variety of other natural or designed protease enzyme systems.

Materials and Methods

See *SI Appendix* for detailed descriptions of experimental procedures and computational methods and for additional data: detailed overview (*SI Appendix, Fig. S1*), threshold determination (*SI Appendix, Fig. S2*), flow cytometry (*SI Appendix, Figs. S3 and S4*), graph metrics (*SI Appendix, Fig. S5*), experimentally derived graphs (*SI Appendix, Fig. S6*), SVM classification (*SI Appendix, Figs. S7, S16, and S17*), negative selection (*SI Appendix, Fig. S8*), population overlap (*SI Appendix, Fig. S9*), positive and negative epistasis (*SI Appendix, Fig. S10*), specificity profile comparisons (*SI Appendix, Fig. S11*), sorting replicates (*SI Appendix, Fig. S12*), sorting gates for all enriched populations (*SI Appendix, Fig. S13*), molecular cloning experimental overview (*SI Appendix, Fig. S14*), and sequence subsampling with more stringent criteria (*SI Appendix, Fig. S15*).

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