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## Surface-based protein domains retrieval methods from a SHREC2021 challenge

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## ABSTRACT

Proteins are essential to nearly all cellular mechanism and the effectors of the cells activities. As such, they often interact through their surface with other proteins or other cellular ligands such as ions or organic molecules. The evolution generates plenty of different proteins, with unique abilities, but also proteins with related functions hence similar 3D surface properties (shape, physico-chemical properties, ...). The protein surfaces are therefore of primary importance for their activity. In the present work, we assess the ability of different methods to detect such similarities based on the geometry of the protein surfaces (described as 3D meshes), using either their shape only, or their shape and the electrostatic potential (a biologically relevant property of proteins surface). Five different groups participated in this contest using the shape-only dataset, and one group extended its pre-existing method to handle the electrostatic potential. Our comparative study reveals both the ability of the methods to detect related proteins and their difficulties to distinguish between highly related proteins. Our study allows also to analyze the putative influence of electrostatic information in addition to the one of protein shapes alone. Finally, the discussion permits to expose the results with respect to ones obtained in the previous contests for the extended method. The source codes of each presented method have been made available online.

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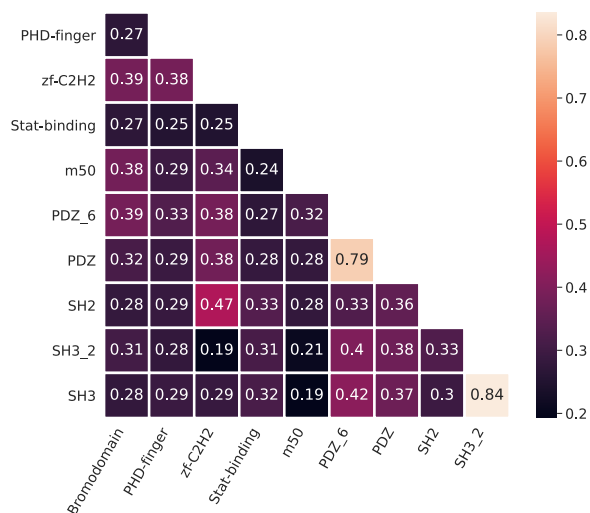
## 1. Introduction

Proteins are key molecular effectors at the cellular level. Proteins are linear assemblies of amino-acids that fold in specific, energy-driven 3D structures [1, 2] linked to their activity. Identifying similarities within protein structures is therefore of tremendous importance in various fields, from biochemistry to drug design. Numerous methods have been dedicated to structural similarity search of proteins in structural bioinformatics, that mainly rely on the analysis of the 3D point clouds defined by the 3D coordinates of their individual atoms [3, 4, 5, 6, 7]. These methods are mostly based on the conserved core structure of proteins, and therefore, may be inefficient to detect proteins sharing similar surface. The protein surface is a higher-level description of the protein structure that abstracts the underlying protein sequence, structure and fold into a continuous shape with geometric and chemical features that fingerprint its interactions with the other molecules of its environment (solvent, ligands, proteins, nucleic acids, ...) [8]. Methods able to detect protein surficial similarities are then of major importance.

Only a limited number of methods have been proposed so far:

- Sael *et al.* use 3D Zernike descriptors to detect either global or local similarities between protein's surfaces [9]. This method is able to use surficial physico-chemical properties like the electrostatic potential or the hydrophobicity [10].
- MaSIF (molecular surface interaction fingerprinting) is a geometric deep learning framework that allows to fingerprint biomolecular surfaces [11]. Both geometric and chemical features are extracted and embedded into numerical vectors which is subsequently processed in an application-dependent manner.
- FTIP (Farthest point sampling-enhanced Triangulation-based Iterative-closest-Point) is a global protein surface comparison method that uses the Farthest point sampling method to extract a subset of protein surfaces, and then uses a triangulation-based efficient Iterative-closest-Point algorithm to align these so-called feature-points [12].
- BioZernike [13] adopts a slightly different approach: instead of using the 3D point cloud formed by the atoms

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**Fig. 1. Structural similarity between the protein structure queries. The TM-score (in the (0, 1] range) measures the topological similarity between two protein structures: the higher the TM-score, the more similar the two structures. Scores below 0.17 correspond to unrelated proteins, while those above 0.5 usually indicate two structures having the same fold [15].**

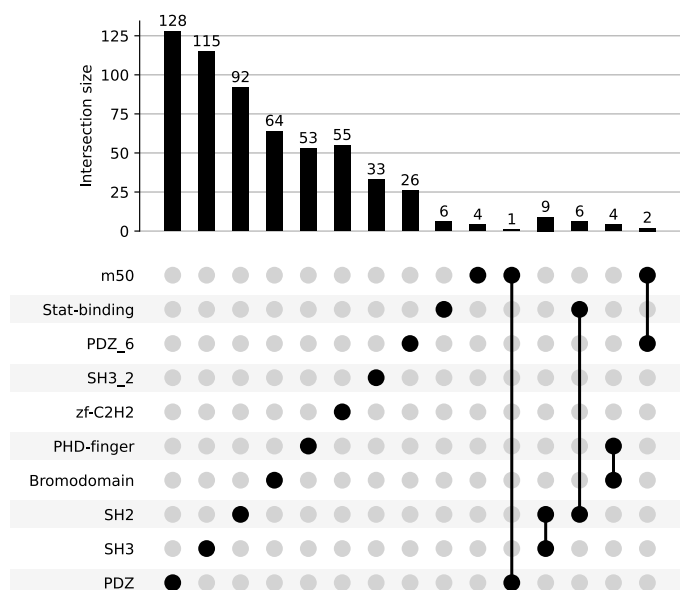
coordinates, it uses the density distribution. A 3D Zernike moment normalization procedure is applied to orient the density volumes to be compared, allowing for fast retrieval of proteins/protein assemblies.

The aim of this track is therefore to assess the performance of currently available methods and to stimulate the development of novel methods. To this end, the dataset encompasses (1) A variety of protein domains, with some of them closely related, to query the dataset. (2) A dataset of experimental structures that contain one or more domains. (3) A few protein shapes corresponding to protein that contains two of the query domains. (4) Two versions of the same dataset, one made of the protein shapes only, and the other with an additional physico-chemical property, the electrostatic potential, encoded along the shapes. We selected this surficial physico-chemical feature as it is the main driving force in many biological recognition processes, such protein-ligand and protein-protein interactions [14]. In the present work, we detail the dataset proposed by the challenge organizers to the participants and how it differs from the previously proposed datasets in Section 2. In Section 3, we describe the 5 methods submitted to the contest. The evaluation metrics are briefly introduced in Section 4, and the performance of the methods is presented in Section 5. Finally, we discuss the outcomes of the different submitted methods in Section 6.

## 2. The Dataset

### 2.1. Constitution of the SHREC'21 dataset

The SHREC'21 protein dataset is based on the Pfam 33.1 database [16]. Basically, this database classifies protein sequences into domains and families, that can be grouped into clans whenever they are evolutionarily related. Protein domains of structures from the Protein Data Bank (PDB [17, 18])



**Fig. 2. The Upset Plot of ten selected Pfam domains in SHREC2021 challenge datasets. The dataset is composed of 554 individual shapes, of which 22 bears two of the domains of the dataset.**

can therefore be attributed to a Pfam domain and, possibly, a clan. To build up the track dataset, we relied on this notion of domain, and manually selected 10 Pfam domains: the SH2 domain (PfamID PF00017), the SH3 domain (PfamID PF00018), the variant SH3 domain (SH3\_2, PfamID PF07653), the PDZ domain (PfamID PF00595), the PDZ\_6 domain (PfamID PF17820), the peptidase family M50 (m50, PF02163), the bromodomain family (PF00439), the DNA-binding domain of the STAT protein (STAT-binding, PF02864), the PHD-finger domain (PfamID PF000628), and the C2H2 Zinc-finger domain (zf-C2H2, PfamID PF00096).

For each selected domain, all corresponding structures from the PDB were listed, and the best resolution structures were retrieved to serve as a query for the track. When applicable, the NMR (Nuclear Magnetic Resonance) structures were assigned an arbitrary resolution of 2.25 Å [19], while structures with no resolution were discarded. The residues corresponding to the Pfam domains were then extracted from the selected structures when necessary, so that the selected domains were left alone. For example, only the DNA-binding domain of the STAT (Signal Transducer and Activator of Transcription) protein was kept as a query, its others domains being discarded.

The remaining structures were filtered according to their Uniprot [20] identifier, and duplicates were discarded to (1) Ensure a diversity of sequence structures among the dataset. (2) Limit the dataset size to a tractable size given the track timeline. Finally, only the best resolution structures for each Uniprot entry were kept. When NMR structures were selected, only the first model was considered. Unlike the query structures, we kept the other domains present in these structures that eventually constitute the dataset. Therefore, many dataset structures display several domains, at least one of which is one of the query domains.

For all structures (queries and dataset structures), we re-

moved all hetero-atoms, and unwanted chains. The resulting PDB structures were then protonated using pdb2pqr [21], using propka [22, 23] to compute the pKa values of the ionizable groups at pH=7.2. The solvent-excluded surface of all protonated structures were computed using the default parameters of EDTSurf [24, 25] except that inner cavities were discarded. We then computed the electrostatics using APBS suite [26], and used the *multivalue* to compute the electrostatic potential at the mesh vertices locations. Two datasets were then assembled, one with only the protein surface shapes (shape-only dataset) and one combining the protein surface shapes and electrostatics values (shape+electrostatic dataset). Similarly, two sets of query surfaces were produced (shape-only and shape+electrostatic). Each dataset (shape-only and shape+electrostatic) includes 554 molecular surfaces which were made available to the track participants, along with the 2 sets of 10 queries, on the track webpage (<http://shrec2021.drugdesign.fr>).

Regarding the dataset, it is important to note that SH3 and SH3\_2 domains were annotated as similar according to HHsearch (a tool commonly used to detect homologous proteins [27]), as well as the PDZ, PDZ\_6 and m50 domains. We present in Figure 1 the TM-scores matrix for all queries of the dataset. A TM-score above 0.5 indicates that the two structures are likely to share the same topology, while unrelated structures are usually associated to TM-scores below 0.17 [28]. SH3 and SH3\_2 query structures show a TM-score of 0.84, while, PDZ and PDZ\_6 query structures show a TM-score of 0.79. Surprisingly, the m50 query structure only has a TM-score of 0.28 and 0.32 with PDZ and PDZ\_6 structures, respectively. A visual inspection of these structures confirmed that the peptidase M50 is topologically different from both the PDZ and PDZ\_6 domains: while the peptidase M50 is mainly  $\alpha$ -helical, PDZ and PDZ\_6 are mixed  $\alpha$ - $\beta$  proteins. Overall, most query structures present an intermediate topological similarity with all other queries, as evidenced by the fact that all TM-scores range from 0.19 to 0.47, except for the aforementioned pairs of classes (SH3 / SH3\_2 and PDZ / PDZ\_6).

Finally, as the dataset encompasses multi-domain proteins, 22 dataset proteins display two of the query domains. Namely, 9 proteins of the dataset encompass both a SH2 and a SH3 domains, 6 proteins encompass both a SH2 and a Stat-binding domains, 4 proteins encompass both a bromodomain and a PHD-finger domain, 2 proteins encompass a PDZ\_6 and a peptidase family M50 domains, and one structure encompass a PDZ and a peptidase m50 domains. The final structure of the dataset and the size of each class is summarized in Figure 2.

## 2.2. Comparison to previous SHREC datasets on proteins

Compared to previous SHREC datasets dealing with protein structures or surfaces [29, 30, 31, 32, 33, 34], the SHREC'21 protein track dataset is characterized by three main aspects : (1) The presence of two datasets representing the same set of proteins, one shape-only and one shape+electrostatic dataset. (2) The close evolutionary relationship of some of the query domains, further characterized by a similar topology (Figure 1). (3) The intermediate similarity of the domains topologies (Figure 1). And (4) The use of individual domains to query a dataset of single- as well as multi-domain proteins shapes.

The main novelty of the SHREC'21 track is arguably the availability of protein surfaces with electrostatic values, which has been shown to improve the retrieval performance of protein surfaces [11, 35]. This additional feature might therefore allow to better distinguish structurally related proteins based on their surficial properties and improve the methods' performance.

### 2.3. Challenge proposed to the participants

SHREC, or 3D Shape Retrieval Challenges, are challenges primarily organized in order to evaluate the effectiveness of 3D-shape retrieval algorithms. A group organizes a challenge by building up a dataset, then proposes the challenge publicly to the community, and finally gathers, analyses and verifies the results. The theme of the challenge may vary from one to another, but all challenges take place in a limited time, which ranges from 1 to 1 ½ months.

In our contest, the participants were asked, given each of the 10 query surfaces, to retrieve the molecular surfaces of proteins from the dataset that encompass the same domain as the query. Each query-to-dataset-surface distance was expected to be expressed as a dissimilarity score. The results were therefore  $10 \times 554$  matrices of dissimilarity scores. Each participant was allowed to submit one dissimilarity matrix for each dataset: one matrix for the shape-only dataset, and one matrix for the shape+electrostatic dataset.

## 3. Participants and methods

Among the seven groups that initially registered to this track, only 5 were able to produce the results in time and returned a shape-only dissimilarity matrix. Only one method (3DZD, see 3.1) returned a dissimilarity matrix for the shape+electrostatic dataset. The other groups were not able to produce a satisfying training dataset or willing to develop their algorithm to handle the electrostatics values. In the following subsections, each group describes their new, respective methods.

### 3.1. Network trained with encoded 3DZD (3DZD) by Tunde Aderinwale, Charles Christoffer, Woong-Hee Shin, Genki Terashi, Xiao Wang & Daisuke Kihara

Our group submitted two (shape-only and shape+electrostatic) dissimilarity matrices of the target proteins to the 10 query proteins provided by the organizers. These methods are based on the 3D Zernike Descriptor (3DZD). 3DZD is the rotation-invariant shape descriptor derived from the coefficients of 3D Zernike-Canterakis polynomials [36].

#### 3.1.1. Summary of the 3DZD method

Similar to SHREC'20 [34], our group trained two types of neural network to output a score that measures the dissimilarity between a pair of protein shapes. Briefly, the first framework (the Extractor model) is structured into multiple layers: an encoder layer with 3 hidden units of size 250, 200 and 150, a feature comparator layer which computes the Euclidean distance, cosine distance, element-wise absolute difference and product, and a fully connected layer with 2 hidden units of size 100 and 50. There are multiple hidden units in each layer, and our group

uses the ReLu activation function in all except the output of the fully connected layer where the sigmoid activation function is used to output the probability that the two proteins belong to the same protein- or species-level in the SCOPE dataset classification [37, 38]. The second framework (the end-to-end model) is similar to the first except the feature comparator layer is removed and the output of the encoder is directly connected to the fully connected layer.

The network is trained on the latest SCOPE dataset of 259,385 protein structures. 2,500 protein structures were set aside for network validation. Proteins in Class I (Artifacts) were removed. Each of the two network frameworks is trained with two datasets. The first dataset is 3DZDs of the surface shape of proteins and the second one is feature vectors that concatenate 3DZD of shape and 3DZD of the electrostatic properties.

Our group examined the performance of the networks on the validation dataset to determine which models to use. For the shape-only dataset, we submitted predictions generated by the Extractor model. For the shape+electrostatic dataset, we submitted the average predictions between the Extractor model and the end-to-end model.

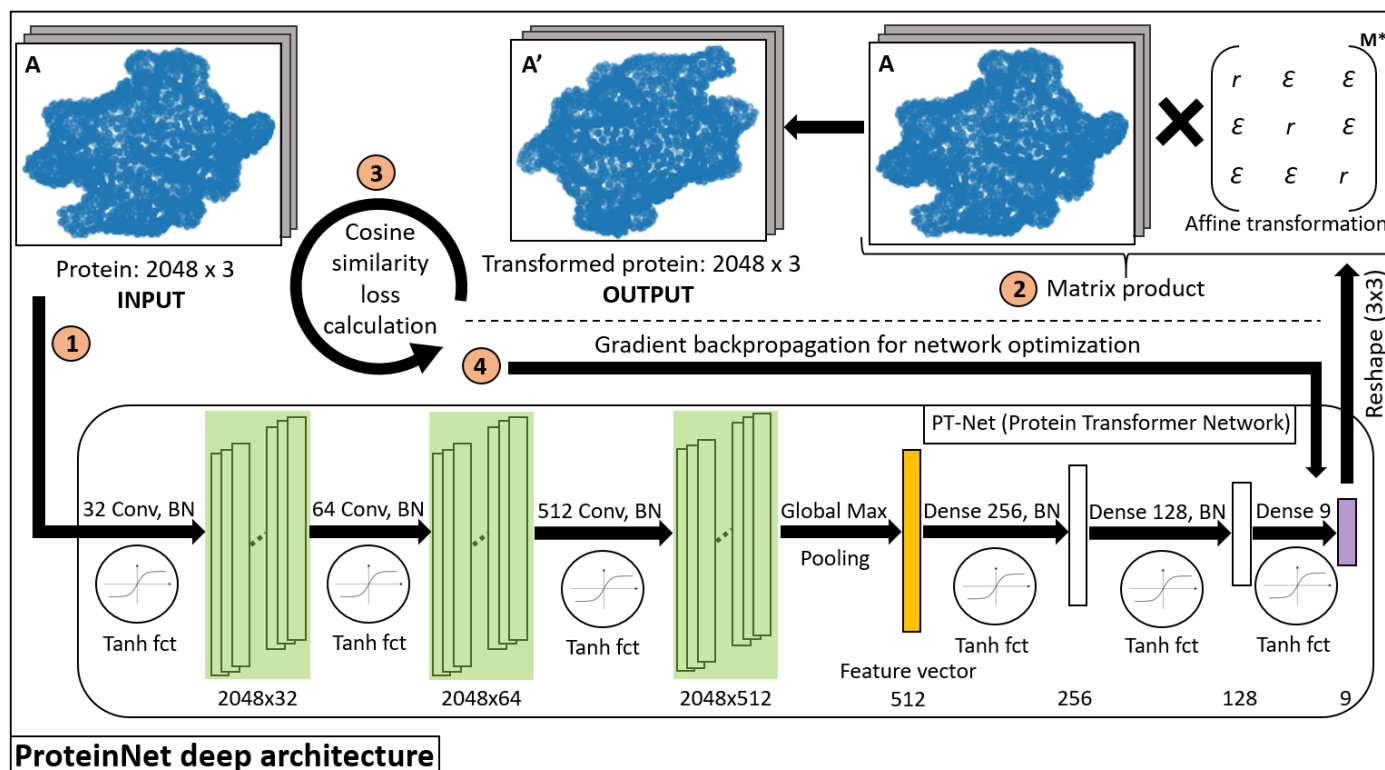
For each protein in the provided dataset, our group performs a pre-processing step as follows: (1) The PLY mesh data file is converted to a volumetric skin representation (Situs file) where points within 1.7 grid intervals are assigned with values that are interpolated from the mesh [9]. For the electrostatic features, the interpolated values are the potentials at the mesh vertices. For the shape feature, a constant of 1 is assigned to grids that overlap with the surface. (2) The resulting Situs file is fed into the EM-Surfer pipeline [39] to compute 3DZD.

#### 3.1.2. Runtimes and computational cost

It takes approximately 12–13 min to pre-process each file. Generating 3DZD took ~8.00 s on average for each protein on an Intel® Xeon® CPU E5-2630 0 @ 2.30GHz. The training of each models took 12 h. Dissimilarity prediction between two proteins using the trained model took ~0.22 s on average on a Nvidia® Titan X GPU. The averaging of the two matrices was almost instant and negligible. The code is available at the following url: [https://github.com/kiharalab/shrec\\_2021\\_shape\\_retrieval](https://github.com/kiharalab/shrec_2021_shape_retrieval).

### 3.2. ProteinNet: Deep learning based protein characterization from 3D point clouds (ProteinNet) by Halim Benhabiles, Karim Hammoudi, Adhane Cabani, Feryal Windal & Mahmoud Melkemi

Our group proposes a deep learning approach to calculate a protein descriptor from its 3D point cloud. To this end, we developed a variant of PointNet [40] which is a point cloud deep architecture dedicated for 3D classification and segmentation. We adapted this architecture in order to learn an affine transformation matrix that allows to align the coordinates of the input 3D protein point cloud into a canonical representation. The new representation maintains interesting properties demonstrated in [40], including invariance to rigid geometric transformations as well as point order permutations. The ProteinNet deep architecture is illustrated in Figure 3. More specifically, the architecture is based on a PT-Net module (Protein



**Fig. 3. ProteinNet deep architecture for protein point cloud transformation into canonical representation. Step (1): affine transformation matrix estimation. Step (2): protein point cloud transformation using the estimated affine matrix. Step (3): similarity calculation between the original protein point cloud (the input) and its transformed point cloud. Step (4): cosine similarity loss calculation between the original input protein point cloud and its transformation; and back-propagation over the network to optimize the estimation of the affine transformation matrix.**

Transformer Network) which is inspired from the T-Net (Transformer Network) module of the original PointNet architecture. The PT-Net module is trained to predict an affine transformation matrix  $M$  that is constrained to be close to an orthogonal matrix, namely  $|(M.M^t) - I| = 0$  (step 1 in Figure 3). The matrix  $M$  is used to transform the input protein into its canonical representation (step 2 in Figure 3). A cosine similarity loss between the original protein and the transformed one is then calculated (step 3 in Figure 3) in order to back-propagate the error over the network (step 4 in Figure 3) and optimize the matrix  $M$ .

### 3.2.1. PT-Net module

The module is composed of a sequence of 3 convolution blocks (32, 64 and 512 layers) followed by a global max pooling layer and 3 successive dense layers (256, 128 and 9). As shown in Figure 3, each convolution block as well as the dense layers (except the last one) undergo a batch normalization and a tangent hyperbolic activation function. The last dense layer of 9 units is reshaped to output the  $(3 \times 3)$   $M$  matrix (see details in [40]).

### 3.2.2. Data preparation and architecture training

All the proteins of the dataset of the track have been sampled to 2,048 points using a Poisson disk sampling technique [41] and then normalized into a zero-center unit sphere based on their respective minimum bounding spheres [42]. The architecture has then been trained using a batch size of 16 on 80% of

the dataset over 150 epochs and validated on the remaining 20% of the data. The training data were augmented on-the-fly (during the training process) by adding some geometric noise (e.g. random displacement of point coordinates in a limited interval).

### 3.2.3. Protein feature extractor

The trained ProteinNet model has then been exploited to calculate a protein feature descriptor, for each input protein, by extracting its intermediate Global Max Pooling hidden layer. This descriptor corresponds to a 1-dimension vector of 512 values.

### 3.2.4. Dissimilarity matrix computation

The dissimilarity matrix between the ten protein shape queries and the set of 554 protein shapes has been calculated using Euclidean distance between their respective 512 feature vectors.

### 3.2.5. Runtimes and computational cost

This framework has been developed in Python 3.7.6 using different libraries, namely Open3D 0.8.0.0, and Keras 2.2.4-tf on a TensorFlow-GPU 2.1.0 backend. The experiments have been conducted on an Intel Xeon® Gold® 5118 CPU@2.30 GHz with 128 GB of memory and NVIDIA® Tesla® T4 GPU with 16 GB of memory. The running times in sof each stage performed on CPU are reported in Table 1 for one protein. Table 2 shows the training times of the ProteinNet model trained on GPU. The code is available at the following url: <https://github.com/Benhables-JUNIA/ProteinNet>.



**Table 1. Running times in s using CPU for each stage of the ProteinNet framework obtained for one protein.**

Point cloud maximum and minimum sizes of two proteins	Point cloud sampling (2,048) and normalization	Feature descriptor (512) calculation of one protein	Distances from one protein to all protein dataset (554 proteins)
582,496 points	1.14	0.005	0.004
37,658 points	0.9		

**Table 2. Running times in s using CPU for each stage of the ProteinNet framework obtained for one protein.**

Deep learning model	ProteinNet
<b>Training data size</b>	499
<b>Epochs</b>	150
<b>Training time (s)</b>	155

3.3. Fisher Kernel agglomerated local Augmented Point Pair Feature Descriptors, trained with Gaussian Mixture Model (APPFD-FK-GMM) by Ekpo Otu, Reyer Zwiggelaar, David Hunter & Yonghuai Liu

Our group presents a novel APPFD-FK-GMM 3D shape retrieval method (see Figure 4) based on Fisher Kernel (FK) and Gaussian Mixture Model (GMM) agglomeration of the Augmented Point-pair Feature Descriptor (APPFD) [43]: a 3D key point shape descriptor that robustly captures the physical geometric characteristics of 3D surface regions. Previous APPFD binning technique involves bucketting each of the 6-dimensional features of the APPFD into a multi-dimensional histogram with at least 7 bins in each feature-dimension, resulting to approximately  $7^6 = 117649$ -dimensional final feature-vector (APPFD), which is very high-dimensional final descriptor.

In this work, we contribute a simpler approach, where each of the 6-dimensional feature is binned into a 1-dimensional histogram with 35 bins for each feature-dimension to produce a 210-dimensional local descriptor (APPFD) for every key point or local surface patch (LSP). Finally, the locally computed APPFDs are agglomerated into a compact code called the Fisher Vector (FV) with 4210 dimensions, which is  $L_2$  and power-normalized, and represents a single protein model, using the FK and GMM [44] framework.

The goal of the APPFD-FK-GMM method/contribution is to provide a straight-forward, efficient, robust, and compact representation, describing the geometry of 3D protein surfaces, with a knowledge-based (*i.e.* non-learning) approach. While a single protein surface in this challenge contains an average of 120,000 vertices and 200,000 triangular faces, our implementation address this very high data-structure by reducing 3D protein surface representation to 3,500 points sample.

### 3.3.1. Summary of The APPFD-FK-GMM Method

Our method involves two main stages: (1) Computing local APPFDs for selected key points on 3D protein surface. (2) Key points APPFDs aggregation with FK and GMM described below. Figure 4 shows the processing pipeline of the APPFD-

FK-GMM algorithm with complete implementation details provided in [44]. The reader is referred to [45], for further details regarding this method.

*Stage (1) - Computing Local APPFDs.* Following key points ( $p_{k_i}$ ) determination for each 3D protein surface, represented as point cloud ( $P$ ), the 4-dimensional feature,  $f_1 = (\alpha, \beta, \gamma, \delta)$  in [46] is augmented with a locally-extracted 2-dimensional feature:  $f_2(p_i, p_j) = (\phi, \theta)$  for every possible combination of point pair,  $p_i, p_j$  (*without* their estimated normals,  $n_i, n_j$ ) in the local surface patch (LSP),  $\{P_i, i = 1 : K\}$  around each key point  $\{p_{k_i}, i = 1 : K\}$  in  $P_s$ , where  $K$  is the number of key points. The extraction of  $f_2$  (see Figure 5) is because  $f_1$  is not robust enough to capture the entire geometric details of the underlying surface, whereas, the PPF approach opens up possibilities for additional feature space.

The angular projections:  $\theta$  and  $\phi$  in Figure 5 are derived by taking the scalar products of  $(\vec{S} \cdot \vec{V}_1)$  for  $\angle_1$ , and  $(\vec{S} \cdot \vec{V}_2)$  for  $\angle_2$  about a point  $p_i$  in a given LSP. Mathematically, scalar products defined in this manner are homogeneous (*i.e.* invariant) under scaling and rotation. Therefore,  $f_2$  is considered rotation and scale invariant for 3D shapes under rigid and non-rigid transformations [34].

Finally, a 6-dimensional  $f_3 = (f_2 + f_1)$  are locally obtained thus:  $f_3(p_i, p_j) = (f_2(p_i, p_j), f_1(p_i, p_j)) = (\phi, \theta, \alpha, \beta, \gamma, \delta)$ , and binned into a 1-dimensional histogram with bins = 35 in each feature-dimension, normalized and concatenated to give  $35 \times 6 = 210$ -dimensional single local APPFDs per LSP.

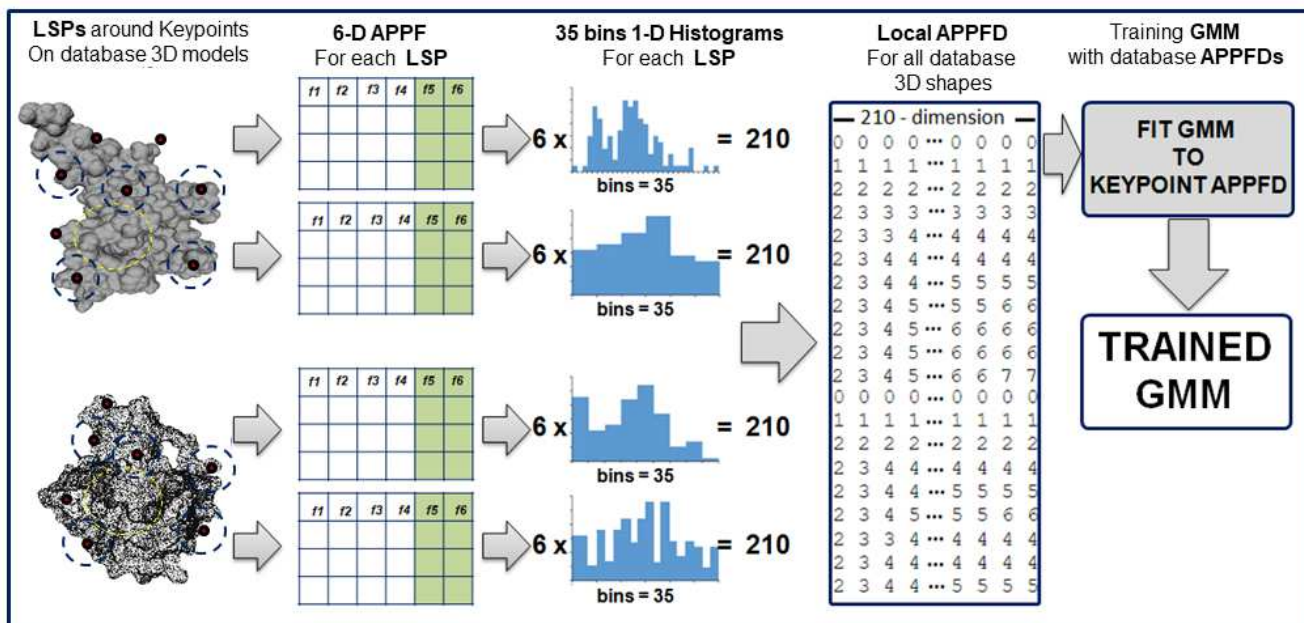
*Stage (2) - Key points APPFDs Aggregation with FK and GMM.* Here, the final descriptor (*i.e.* fisher-vector, FV) computation approach involves an initial step of training a GMM, given aggregated key points local APPFDs for all database 3D objects, then FK is applied on the trained model and a single protein's local APPFDs to derive a global signature (APPFD-FK-GMM) for the protein surface (see Figure 4).

### 3.3.2. Runtimes and computational cost

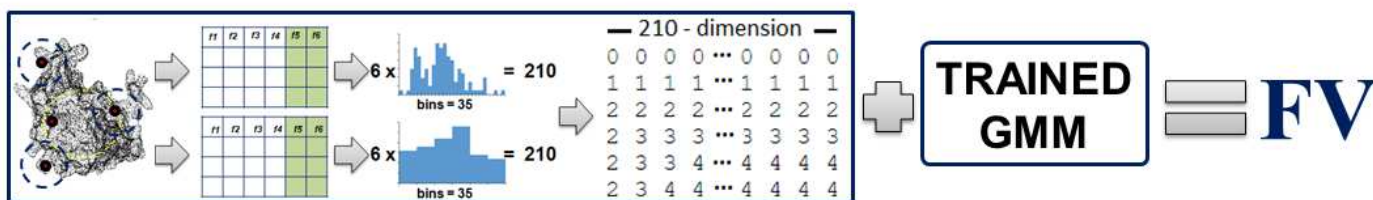
Our group submitted a dissimilarity matrix  $D = [10 \times 554]$ , where the entry  $D = [i, j]$  corresponds to the  $L_2$  distance from  $i^{th}$  FV in the *query* set to the  $j^{th}$  FV in the *collection* set.

While implementing the APPFD-FK-GMM for this task,  $K$  is specified by the parameter,  $vs = 0.20$ , which is the voxel size for point cloud down-sampling, while the radius parameter,  $r = 0.50$  specifies the size of  $P$ . Regarding point cloud size,  $P = 3,500$  points are sampled.

In conclusion, we present a pure Python 3.60 implementation code that computes the APPFD-FK-GMM method. All



Phase 1: Train GMM on database APPFDs



Phase 2: Compute Fisher Vector (FV) for a single 3D model

Fig. 4. APPFD-FK-GMM processing pipeline involving Phase 1 (fitting a GMM to all the keypoints or LSPs descriptor, *i.e.* local APPFDs from each 3D protein surface and for all database protein surfaces) and Phase 2 (computing a single compact descriptor called fisher-vector (FV) for each 3D protein by aggregating all its keypoints or local APPFDs using the fisher kernel (FK) framework and the trained GMM in Phase 1. Within each LSP around a keypoint, six different geometric features are first extracted, and each feature-dimension is binned into a 1D histogram with 35 bins, where all histograms are combined to form a 210-dimensional descriptor, *i.e.* local APPFD for each LSP. All such LSP descriptors from each 3D protein are compacted into a 4210-dimensional FV for that protein model, as in Phase 2.

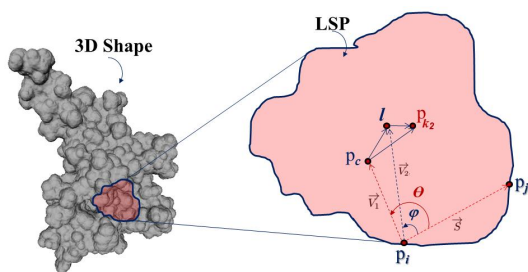


Fig. 5. Local Surface Patch (LSP),  $P_i$  with pairwise points  $(p_i, p_j)$  as part of a surflet-pair relation for  $(p_i, n_i)$  and  $(p_j, n_j)$ , with  $p_i$  being the origin.  $\theta$  and  $\phi$  are the angles of vectors projection about the origin,  $p_i$ .  $\theta$  is the projection angle from vector  $\langle p_i - p_j \rangle$  to vector  $\langle p_i - p_c \rangle$  while  $\phi$  is the projection angle from vector  $\langle p_i - p_j \rangle$  to vector  $\langle p_i - l \rangle$ . The LSP centre is given by  $p_c$ , keypoint is given as  $p_{k_i}$  where  $i = 2$ . Finally,  $l$  is the vector position of  $p_{k_i} - p_c$ . [34]

experiments were carried out under Windows® 7 desktop PC with Intel® Core® i7-4790 CPU @ 3.60GHz, 32GB RAM. It takes an average of 30 s to compute the APPFD-FK-GMM. The implementation code is available at the following url: <https://tinyurl.com/shrec21>.

### 3.4. Projected Wave Kernel Signature Maps (PWKSM) by Léa Sirugue & Matthieu Montès

This method is based on the 2D projection of the surface and the Wave Kernel Signature (WKS) descriptor. Wave Kernel Signature [47] is an isometric invariant descriptor that has been extensively improved and used in the field of computer vision [48, 49, 50, 51]. We have combined WKS with a 2D projection on a unit sphere [52]. Lowering one dimension of the space allows us to have a fast and dense comparison of the surface while having a smaller storage size for files.

*Descriptor calculation.* In a first step, the WKS descriptor is computed on the surface of the 3D object for each point of the



mesh. The surface is flattened on the unit sphere using a conformal transformation [52]. Then, the 2D spherical coordinates of the unit sphere are converted into 2D cartesian coordinates on the plane [53]. A maps of size  $(\theta_{max} - \theta_{min})/\delta, (\phi_{max} - \phi_{min})/\delta$  is created.  $\theta_{max}$  and  $\theta_{min}$  are the maximum and minimum values of  $\theta$  on the unit sphere and the same with  $\phi$ , each representing an angle coordinate.  $\delta$  is a coefficient to adapt for resolution. This type of projection is similar to topographic maps, that is why our group called this descriptor Projected Wave Kernel Signature Maps (PWKSM). An interpolation in the space of discrete integers is done to densify the maps. To reduce impact of deformation at the poles when converting to 2D cartesian coordinates, we computed 7 different maps with different pole orientations.

*Descriptor comparison.* A dense comparison is made using GPGPU sum reduction technique [54] [55] [56]. Each point's WKS of a PWKSM is compared to all points' WKS of another PWKSM. The Earth Mover's distance  $L$  is used to compare the WKS descriptor of each point. Then, the smallest distance between a point of a first PWKSM  $T$  and all points of a second PWKSM  $V$  is selected. The sum of all the smallest distances for each point of the first PWKSM are summed to create the score  $S_T$ . The same is done for computing  $S_V$ .

$$S_T(T, V) = \sum_{k_T=1}^{N_T} \min_{k_V} L(T(k_T), V(k_V)) \quad (1)$$

The final score is the average of  $S_T$  and  $S_V$  defined as follows:

$$S = \frac{S_T + S_V}{2} \quad (2)$$

### 3.4.1. Runtimes and computational cost

All the calculations were made on a computer based on a 64-bit OS with an Intel® Xeon® CPU @ 2.30GHz, a Nvidia® Quadro® k4200 GPU with 4GB and 32GB of RAM. Computing the WKS took on average 9 min and 31 s. It required on average 44 s to compute one PWKSM. The comparison of two surfaces (*i.e.* 7 versus 7 PWKSM) takes on average 23 s. The code is available at the following url: <https://gitlab.cnam.fr/gitlab/siruguej/PWKSM>.

## 3.5. Graph-based learning methods for Surface-based protein domains retrieval (DGCNN) by Huu-Nghia H. Nguyen, Tuan-Duy H. Nguyen, Vinh-Thuyen Nguyen-Truong, Danh Le, Hai-Dang Nguyen & Minh-Triet Tran

In this deep learning method, our group exploits the availability of protein class labels from [35] to optimize the representation of protein surfaces without any additional properties. Particularly, we designed a message-passing graph convolutional neural network (MPGCNN) with the Edge Convolution (EdgeConv) paradigm [57] for the protein classification objective. Then, the latent representation of protein surfaces from this neural network is used for the retrieval task in this track.

### 3.5.1. Data pre-processing

For the meshes in each 3D model of a protein surface, we first sample 512 points on the surfaces of the meshes based on the area of the meshes. Then, to re-assign the topological structures for sampled points, we connect each nodes with their  $k$ -Nearest Neighbors based on their original coordinates ( $k = 16$ ).

### 3.5.2. Edge Convolution

In this geometry-only setting, the initial node features is the coordinates of sampled points. Each protein surface is represented by a  $k$ -Nearest Neighbors graph generated in the pre-processing step with 512 vertices (nodes).

The module that performs the graph message-passing function is the EdgeConv layer [57]. In the EdgeConv layer, the information of a vertex  $i$  after layer  $l$  is calculated as follows:

$$x_i^{l+1} = \max_{j \in N} h(x_i^l, x_j^l) \quad (3)$$

where  $N$  is the neighboring vertices of vertex  $i$  with

$$h(x_i^l, x_j^l) = \text{ReLU}(\text{MLP}(x_i^l \oplus x_j^l)) \quad (4)$$

where ReLU is Rectified Linear Unit (in this implementation, we used LeakyReLU—a variant of ReLU), MLP is a standard multilayer perceptron (MLP), and  $\oplus$  is the concatenation operator.

In this implementation, our group uses a dynamic variant of EdgeConv instead of the standard EdgeConv described above. At each Dynamic EdgeConv layer, each vertex's  $k$ -Nearest Neighbors is re-calculated in the feature space produced by the previous layer, before applying the standard EdgeConv operation. After the graph has been recomputed, standard EdgeConv operation is performed.

After the pre-processing phase, the vertex features first go through 4 layers of Dynamic EdgeConv. The dimensions of output features for each vertex after these first-4 layers are 64, 64, 128, and 256, respectively. Then, the outputs of these 4 layers are concatenated to become a 512-dimensional vector for each vertex. This 512-dimension vector is then fed through another Dynamic EdgeConv layer, creating the output vector  $v$  with 512 dimensions. The feature vector  $v$  is pooled using the concatenation of the outputs of a *max-pooling* and a *mean pooling* layer to generate the first graph-level feature vector. This vector is passed through two MLP blocks with BatchNorm, Leaky-ReLU, and Dropout layers. Finally, the vector is passed through a Fully-Connected layer for classification.

The latent representation of the graph is extracted as vectors by removing the last Fully-Connected layer from the network. The retrieval task is then performed by exploiting the  $L2$ -distances between these vectors.

### 3.5.3. Runtimes and computational cost

This method is implemented in Python 3.8 [58], using Pytorch [59] and Pytorch Geometric [60] libraries. The experiments were carried out a machine with an Intel® Core® i7-8700K 6-core CPU Processor @3.70 GHz with 32 GB of RAM and an NVIDIA® TITAN V with 12 GB of VRAM. The training and test set's embedding extraction uses both the CPU and

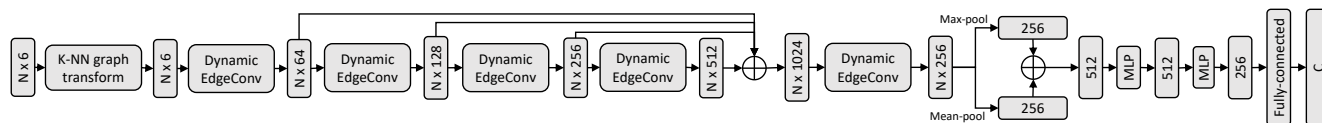


Fig. 6. Dynamic Edge Convolutional Neural Network

the GPU, while computation of distance matrix only uses the CPU. The detailed time report is represented in Table 3. The code is available at the following url: <https://github.com/huunghia160799/SHREC-protein-domains>.

Table 3. Time report of each step of the DGCNN method.

Training	Test Set Extraction	Matrix Computation	Total
≈ 1100 min	≈ 7 min	≈ 3 min	1173 min

#### 4. Evaluation metrics

We use common evaluation metrics to assess the performance of the proposed methods, most of which are used in other SHREC tracks [61] or similar works evaluating the performance in retrieval [62]. For each method, we compute the overall metrics (*i.e.* the metrics averaged over all queries) and the individual metrics (*i.e.* the metrics for each query) to provide a better understanding of the performance of each method for each query. Two composite classes are also presented: the SH3-like and the PDZ-like, which correspond to the grouping of the SH3 and SH3\_2 classes, and of the PDZ and PDZ\_6 classes, respectively. To set the dissimilarity values for these composite classes, for each entry of the dataset, we kept the minimal dissimilarity value from the SH3 / SH3\_2 queries and from the PDZ / PDZ\_6 queries.

##### 4.1. Nearest neighbor, First tier and Second Tier

These metrics measure the ratio of relevant objects among the  $k$  retrieved objects, and ranges in the interval  $[0, 1]$ . For the nearest neighbor (NN), only the first retrieved object is considered ( $k = 1$ ), while the top  $C$  objects are considered for the first tier (FT), and the top  $2 \times C$  objects for the second tier (ST). Here,  $C$  represents the cardinal of the class under investigation, *i.e.* the size of the class to which the query belongs. Higher values of nearest neighbor, first tier and second tier indicate better performance.

##### 4.2. Precision-Recall curves

The Precision (P) represents the fraction of relevant object retrieved compared to the top  $k$  retrieved objects:  $P = (\text{relevant} \cap \text{retrieved}) / \text{retrieved}$ . Therefore, precision can be evaluated at different intervals. The Recall (R) represents the fraction of relevant objects retrieved compared to the size  $C$  of

the class of the query:  $P = (\text{relevant} \cap \text{retrieved}) / \text{relevant}$ . Both metrics range from 0 to 1. Precision-Recall curve plots the precision values at given recall values, which produces, in an ideal case, an horizontal line at  $P = 1$  that spans the entire range of recall values.

##### 4.3. Confusion matrix

A confusion matrix (CM) is a square matrix, whose columns represents the different classes of the dataset and rows the class of the query. For each row, each element  $CM(i, j)$  gives the number of objects from class  $i$  retrieved using the query  $j$ , considering the top  $k = C$  retrieved objects,  $C$  being the size of the class corresponding to query  $j$ . The elements  $CM(i, i)$  in the diagonal of the confusion matrix indicates the objects classified correctly, while the off-diagonal indicates mis-classified elements. To ease the comparison between classes of different sizes, the numbers were normalized over the class sizes  $C$  of the queries. Consequently, the sum  $\sum_j CM(i, j)$  of all elements of each row equals 1.

##### 4.4. Reciprocal Rank and Mean Reciprocal Rank

The Reciprocal Rank (RR) measures the performance to find the first relevant item. For a given query, it equals to the inverse of the rank  $r$  of the first relevant item found:  $RR = 1/r$ . The Reciprocal Rank ranges from 0 (no relevant object found) to 1 (the first retrieved object is relevant). The Mean Reciprocal Rank (MRR) is the Reciprocal Rank averaged over all queries. This metric is useful as: (1) It is considered order-aware, contrary to the previous metrics. (2) Typical use cases only consider the few first retrieved items; therefore, the higher the reciprocal rank, the better the performance.

## 5. Results

Among the participants of the track, all teams returned a dissimilarity matrix for the shape-only dataset, and only one method (3DZD) was adapted to handle the shape+electrostatics dataset.

### 5.1. Shape-only challenge

The results for the shape-only dataset are presented in Table 4 and in Figures 8 and 7. Table 4 summarizes the performances of all submitted matrices for the shape-only dataset. For each metric (Nearest Neighbor, First Tier, Second Tier and Mean Reciprocal Rank), the highest value is indicated in bold. Given the dataset structure and the selected query domains, the best

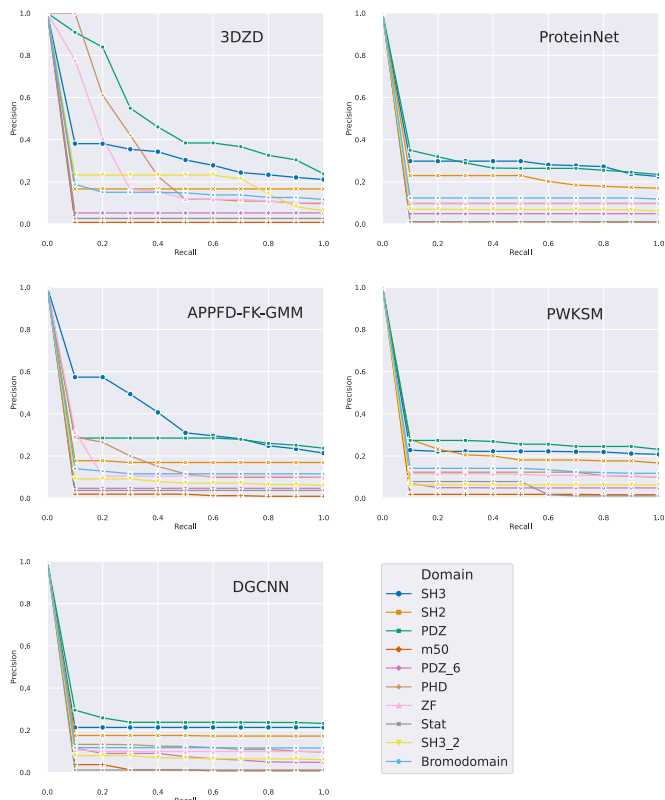


Fig. 7. Per-query precision-recall curves for the shape-only dataset, for each method. All plots are colored according to the legend on the bottom right of the figure.

Table 4. Summary of the average evaluation metrics for the shape-only dataset. The composite classes are excluded from the average; they are presented in Tables C.7 to C.10.

Method	Nearest Neighbor	First Tier	Second Tier	Mean Reciprocal Rank
3DZD	<b>0.5</b>	<b>0.160</b>	<b>0.292</b>	<b>0.523</b>
ProteinNet	0	0.088	0.195	0.126
APPFD	0.3	0.136	0.237	0.410
PWKSM	0.1	0.105	0.201	0.236
DGCNN	0	0.098	0.189	0.193

1 method achieves an overall level of 0.5 for the nearest neighbor  
 2 metric, 0.160 for the first tier, 0.292 for the second tier and  
 3 0.523 for the mean reciprocal rank. These results must be bal-  
 4 anced by the fact that a few classes have only a small number  
 5 of models (namely, the Stat-binding and m50 classes only have  
 6 6 and 4 members, respectively, see Figure 2), and thus impact  
 7 negatively the averaged results. For completeness, Tables C.7,  
 8 C.8, C.9 and C.10 in Appendix C contain the per-class evalua-  
 9 tion metrics for all methods.

10 The precision-recall curves for each individual classes (Fig-  
 11 ure 7) show that most methods display a similar behavior for  
 12 all classes, characterized by a quick drop of the precision at  
 13 low recall values. A few methods, however, show a differ-

ent pattern for a few classes (Figure 7): see the PDZ class for  
 14 the 3DZD method (green curve, top left plot) or the SH3 class  
 15 for the APPFD-FK-GMM method (dark blue curve, middle left  
 16 plot), for instance, whose corresponding curves display medium  
 17 precision values at medium recall. The confusion matrices for  
 18 all methods are shown in Figure 8. Combined with Figure 1,  
 19 they allow us to put the performance into perspective. For in-  
 20 stance, PDZ and PDZ\_6 domains are topologically very simi-  
 21 lar (TM-score: 0.79, Figure 1) and therefore were expected to  
 22 be confusing. When using the PDZ\_6 query, ProteinNet re-  
 23 trieved only 1 (4%) of the 26 PDZ\_6 shapes within the first  
 24 26 retrieved results, but also 12 (46%) shapes from the PDZ  
 25 class (Figure 8, middle top confusion matrix). More strikingly,  
 26 3DZD only found 1 (3%) of the 33 SH3\_2 shapes within the 33  
 27 first retrieved shapes using the SH3\_2 query, but the other 32  
 28 retrieved shapes belong to the SH3 class (Figure 8, second row  
 29 of the top left confusion matrix), which is closely related to the  
 30 SH3\_2 class (TM-score: 0.84, Figure 1).  
 31

## 5.2. Shape+electrostatics challenge

Table 5. Summary of the average evaluation metrics for the shape+electrostatics dataset. The composite classes are excluded from the average; they are presented in Tables D.11 to D.14.

Method	Nearest Neighbor	First Tier	Second Tier	Mean Reciprocal Rank
3DZD	0.5	0.160	0.321	0.454

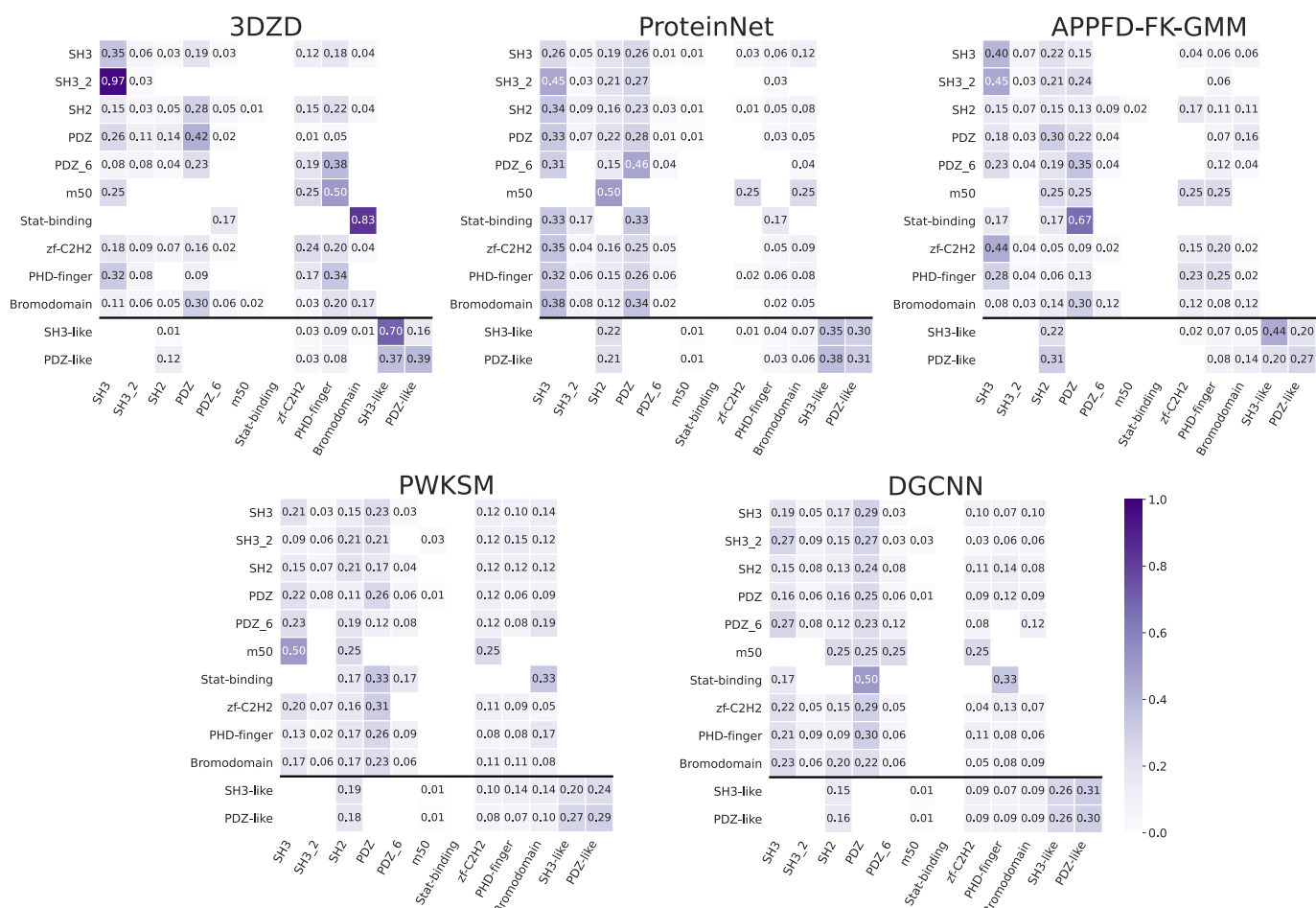
Similarly to the shape-only dataset, results for the  
 33 shape+electrostatics dataset are presented in Table 5 and Fig-  
 34 ures 9 and 10. Only one team returned a dissimilarity matrix  
 35 for the shape+electrostatics dataset. The evaluation metrics are  
 36 listed in Table 5. The results show similar trends compared to  
 37 the shape-only dataset, with a nearest neighbor of 0.5, a first tier  
 38 value of 0.16, a second tier value of 0.321 and a mean reciprocal  
 39 rank of 0.454. These metrics are similar to the results obtained  
 40 from the shape-only dataset for the 3DZD method (the second  
 41 tier value increased while the mean reciprocal rank decreased).  
 42 The per-class metrics are shown in Appendix D (Tables D.11,  
 43 D.12, D.13 and D.14).  
 44

45 The precision-recall curves (Figure 9) show a similar overall  
 46 behavior for the 3DZD method, whose performance improved  
 47 significantly for the SH3 domain but decreased significantly for  
 48 the PDZ domain (dark blue and green curves, respectively, left  
 49 plot of Figure 9). The confusion matrix (Figure 10) is in line  
 50 with the previous results, indicating that 3DZD performs simi-  
 51 larly in terms of overall performance but with a few differences  
 52 at the per-class results.

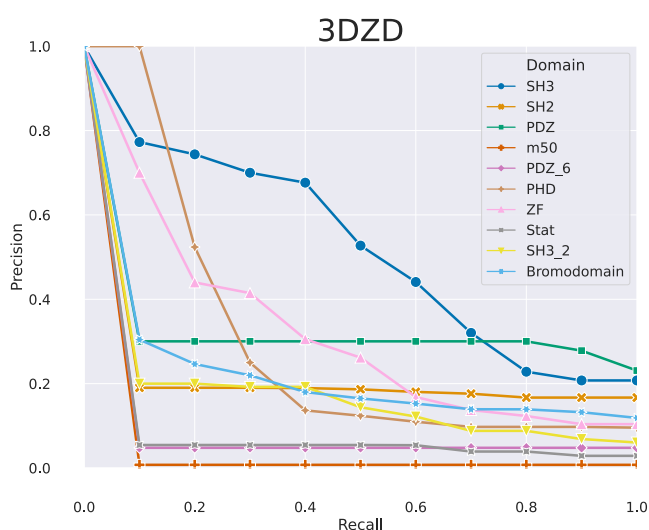
## 6. Discussion and concluding remarks

### 6.1. Shape-only dataset

Overall, the 3DZD method obtained the best results, in line  
 55 with the previous tracks on protein shapes, where this group  
 56



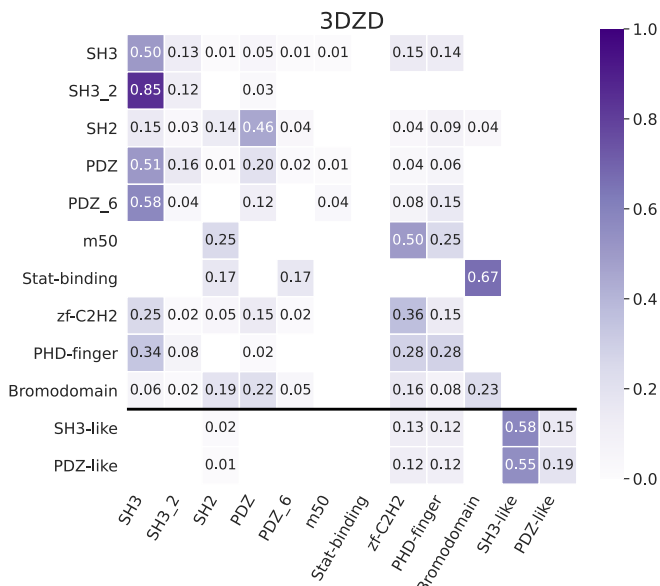
**Fig. 8. Confusion matrices of all methods for the shape-only dataset. The color-range is the same for all matrices. Confusion ranges from 0 (white background) to 1 (deep purple background). The original classes are separated from the composite classes (SH3-like and PDZ-like) by a black line.**



**Fig. 9. Per-query precision-recall curves for the shape+electrostatics dataset, for each method. All plots are colored according to the legend on the far right of the figure.**

similarly obtained overall good results [33, 34]. This method relies on the use of 3D Zernike polynomials, which has been successfully used to retrieve proteins based on their shapes [9] or their  $C\alpha$  coordinates [13]. It then uses a neural network trained on the SCOPe [37, 38] database, whose classification largely overlaps with the classification of the Pfam database [16]. As instance, in the SCOPe databases, the SH3 domain and the SH3\_2 domain are classified in two different SCOPe domains, similarly to the Pfam classification. The DGCNN used the data from another SHREC'21 track, the *Retrieval and classification of protein surfaces equipped with physical & chemical properties* track [35]. The organizers of this track, similarly to a previous track [33], proposed a set of shapes derived from NMR structures along their surficial physico-chemical properties to allow the participants to train their methods, and the resulting classification of the proteins was derived from the SCOPe database as well. The DGCNN and 3DZD methods were therefore trained on similar data, but produced different performance.

Another point to consider, is that the DGCNN method uses a sampling (down to  $\approx 8,000$  points) of the initial point clouds (see 3.5.1) that potentially resulted in a loss of information



**Fig. 10. Confusion matrix for the shape+electrostatics dataset. Confusion ranges from 0 (white background) to 1 (deep purple background). The original classes are separated from the composite classes (SH3-like and PDZ-like) by a black horizontal line.**

that might explain the difference of performance between these two groups (DGCNN and 3DZD). However, the ProteinNet and APPFD-FK-GMM methods use more severe down-sampling steps as well to reduce the number of points down to 2,048 for ProteinNet (see 3.2.2) and to 3,500 for APPFD-FK-GMM (see 3.3) with various outputs in terms of performance. These numbers should be compared to the initial meshes sizes, which range from 37,658 to 582,496 points. The APPFD-FK-GMM group, however, was able to better retrieve relevant results within the first hits, as evidenced by higher values of Nearest-neighbor and Mean Reciprocal Rank for the shape-only dataset (Table 4).

While some methods were able to maintain medium precision levels at medium recall values (see section 5.1), a few queries were difficult to handle for all methods. For the DNA-binding domain from the STAT protein family or the peptidase M50 domain, the low number of such surfaces (6 and 4, respectively) in the datasets explains the low performance observed for all methods. For the other queries, like the PDZ\_6 domain or the SH3\_2 domain, the explanation is the presence of closely related domains, the PDZ and SH3 domains, respectively. These confusing classes are significantly more populated (128 *versus* 26, for the PDZ / PDZ\_6 domains, and 115 *versus* 33, for the SH3 / SH3\_2 domains). This is supported by the confusion matrices, which showed that, for instance, the ProteinNet group retrieved a great amount of SH3 domains (and almost no SH3\_2 domains) within the top results using the SH3\_2 query, or the DGCNN group retrieved a significantly higher proportion of PDZ domains than PDZ\_6 domains within the first results using the PDZ\_6 query. In these cases, the high level of similarity between the domains coupled to the imbalanced size of the classes have negatively impacted the results. In the mean

time, these results highlight the limits of the currently available methods to distinguish between the most closely related proteins.

Also, we observed different results for order-aware (mean reciprocal rank) and order-unaware (nearest neighbor, first tier and second tier) metrics. While DGCNN, ProteinNet and PWKSM methods display similar values for order-unaware metrics, PWKSM displays a higher Mean Reciprocal Rank. When converted back to ranks, these results mean that, on average, PWKSM ranked better the first relevant match, but did find a similar amount of relevant items within the top results.

## 6.2. Shape+electrostatic dataset

While the shape of a protein is of tremendous importance, its surficial properties are important as well. Therefore, in this track, we generated the shape+electrostatic dataset which encompass both properties, in order to stimulate the development of such methods. However, only one groups returned a dissimilarity matrix for this dataset, namely the 3DZD group. Most groups that participated to this challenge come from the computer vision field. As such, most of the methods presented in this work are the result of methodological developments dedicated to the analysis of 3D point clouds. The development of new, specific methods to handle both shapes and electrostatics would require an amount of time far greater than the SHREC timeline. Nevertheless, we hope that this challenge along with the other challenge dedicated to protein shapes [35] would stimulate the development of such methods.

The overall results showed that the treatment of the electrostatics by the 3DZD marginally improved the results, compared to the shape-only results. Interestingly, the electrostatic potential impacted differently each class: it improved the ability of the 3DZD method to handle the SH3 or the Bromodomain classes, as evidenced by the precision-recall curves (Figures 7 and 9, Tables in Appendix C and Appendix D). The 3DZD method is derived from previous attempts from the same team to couple shape and electrostatics analysis to classify protein surfaces [10]. As noted in this exploratory work, electrostatics may be suitable to compare closely related proteins, while our datasets was mainly composed of loosely related proteins (Fig 1). The electrostatics potential feature is likely to be more beneficial for protein surface comparison of local features rather than global shapes. In such local cases (like comparison of catalytic or binding sites), local electrostatic hot spots would represent the major local feature rather than one of the many features of the global protein surface, as it is the case for the MaSIF method [11].

## 6.3. Current machine learning-based methods: pitfalls and challenges

Pitfalls of previously mentioned methods lies in the exploited protein datasets and their characteristics, notably the class imbalance as well as the high inter-class shape similarity. Indeed, the protein datasets are often highly imbalanced in terms of protein classes, which introduces a bias in the training process of these methods towards learning efficiently class representation. One common technique consists of using data augmentation to



1 overcome this lack of original data. Hence, it is important to  
2 bear in mind that several protein classes cannot be considered as  
3 representative groups of protein families. Moreover, our prob-  
4 lematic tackles a large quantity of classes composed of protein,  
5 which are *e.g.* visually highly similar. In such a case, a chal-  
6 lenge lies in the design of new methods with a high discrimi-  
7 nating power that allows to extract the most significant features  
8 for distinguishing between protein classes. In this sense, other  
9 aspects of the proteins (in addition to the shape) such as molec-  
10 ular properties and electrostatic properties could be considered.  
11 These parameters have to be carefully analyzed through exper-  
12 iments before envisaging method generalizations.”

#### 13 6.4. Concluding remarks

14 In conclusion, we have presented the results of the  
15 SHREC’21 challenge on *Surface-based protein domains re-*  
16 *trieval*. The number of participants remained stable compared  
17 to the last two years, indicating a constant interest of the shape  
18 retrieval community towards biologically relevant problems.  
19 Each group relied on different methods and theoretical back-  
20 ground with respect to recommended modeling/machine learn-  
21 ing practices [63, 64] in order to solve the problem proposed  
22 by the organizers, and represent a variety of approaches to the  
23 same problem. As a step towards open science, all partici-  
24 pants accepted to share their programs publicly with the com-  
25 munity. Overall, the results are decreased compared to simi-  
26 lar past tracks [34]. Indeed, two methods based on descriptors  
27 similar to 3DZD and APPFD-FK-GMM (3DZD and HAPPS,  
28 respectively) were presented in the SHREC’20 contest and per-  
29 formed very well (*e.g.* both methods exceeding 0.95 for the NN  
30 metric) on a problem similar to the shape-only problem (see Ta-  
31 bles 6 and 7 of [34]). However, the adapted versions (3DZD and  
32 APPFD-FK-GMM) did not reach the same level of performance  
33 by exploiting this new, particular dataset of proteins. This de-  
34 crease of performance (and low performances from the three  
35 other methods) reveals that this year dataset was particularly  
36 hard to analyze, and that there is still room for improvements.  
37 Among the proposed methods, we observed that 3 over 5 used a  
38 learning-based protocol at some point. This proportion is in line  
39 with last year track, and show that such approaches continue  
40 to be investigated as they usually improve the results. To this  
41 regard, the SHREC’21 track on *Retrieval and classification of*  
42 *protein surfaces equipped with physical & chemical properties*  
43 might highlight some interesting points on the best architecture  
44 to learn protein surficial properties [35]. Similarly, the organiz-  
45 ers of this track computed a set of additional chemical proper-  
46 ties (electrostatic potential, location of potential hydrogen bond  
47 donors and acceptors, hydrophobicity). In this track, the par-  
48 ticipants first used the surface geometry then the combination  
49 of the geometry and physico-chemical features of the protein  
50 surfaces. The results showed that all methods improved their  
51 results when using both the geometric and physico-chemical  
52 data compared the the geometry only. Particularly, the results  
53 generated by the machine learning based methods increased  
54 more compared to the other methods. As each of the physico-  
55 chemical feature was not considered individually, it remains  
56 hard to knowpredict whether one feature hasbeat a greater im-  
57 portance than the other. However, in their work, Gainza et al.

58 showed that the electrostatic potential have the greatest impact  
59 of the physico-chemical features they computed. In our work,  
60 the electrostatic potential was used by the 3DZD team as an  
61 additional feature to help the retrieval task. The results reveals  
62 only slightly improved results compared to the results from the  
63 shape-only dataset. As noted in [10], the electrostatic potential  
64 may be of better use to compare closely related proteins, rather  
65 than comparing loosely related proteins, as it is the case in our  
66 work. Alternatively, shapes and electrostatics may be used in  
67 hierarchical way, *i.e.* using first the shape then the electrostatics  
68 to achieve a better result.

69 This track reveals significantly lower results when compared  
70 to past tracks [31, 32, 33, 34]. However, satisfactory solutions  
71 exist to distinguish between loosely related proteins, or to iden-  
72 tify identical proteins with different conformations, based on  
73 their shapes only. Our work also reveals some limits of the  
74 methods used by the participants for the challenge. Very closely  
75 related proteins (such as SH3 and SH3\_2 protein domains),  
76 *i.e.* proteins displaying a high topological similarity and lim-  
77 ited variations of their amino-acid sequences, hence surfaces,  
78 are still difficult to separate in different classes, but some meth-  
79 ods distinguish them from the other classes. Also, when we  
80 consider the DNA-binding domain from the STAT proteins, no  
81 method was able to produce satisfactory results. While the  
82 DNA-binding domain used as a query has a globular shape,  
83 the STAT proteins are significantly bigger, with a non-globular  
84 shape, and have 3 additional domains (1 of which is a SH2 do-  
85 main, a domain included in the dataset), which means that only  
86 partial matches compared to the query may be achieved. This  
87 specific issue (the comparison of partially overlapping objects)  
88 may require further development.

89 In the future, this latter point could be the subject of a ded-  
90 icated SHREC track, and a good indicator of the overall pro-  
91 gresses made in the field of the retrieval of proteins based on  
92 their surfaces. Currently, most methods have difficulties to han-  
93 dle such cases, which are quite common. Solving this chal-  
94 lenge would be a step forward for the community. At the same  
95 time, explainable artificial intelligence (XAI) methods [65] may  
96 highlight the latent features responsible for good or bad pre-  
97 dictions, and help decipher the results of machine learning-  
98 based methods. XAI methods may help explain the perfor-  
99 mance difference observed for each class of protein, and pro-  
100 vide a human-interpretable representation of machine learning  
101 descriptors, and therefore help identifying the current limits of  
102 these algorithms. Finally, deciphering to which extend, if any,  
103 the standard physico-chemical features (electrostatics potential,  
104 charges distribution, hydrophobicity, *etc.*) improve the results  
105 may be the main focus of the next SHREC tracks devoted to  
106 protein surfaces.

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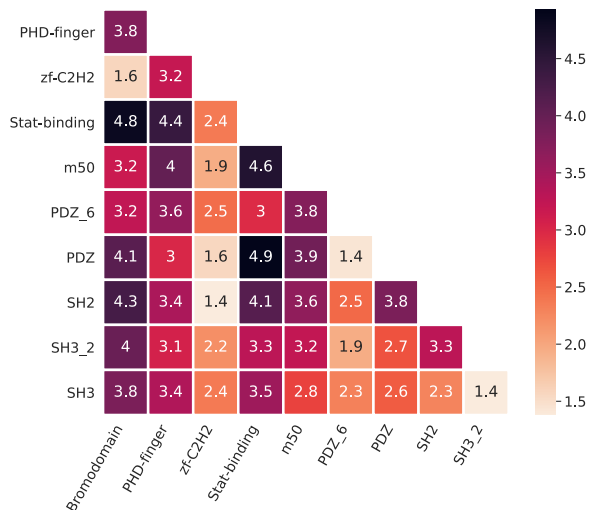
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**Fig. B.11. Ca-RMSD (Root Mean Square Deviations) between queries structures. The higher the RMSD, the more distant the structures.**

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## Appendix A. List of PDB structures used as queries for the dataset

## Appendix B. RMSD between queries structures.

## Appendix C. Evaluation metrics details for the shape-only dataset: reciprocal rank, per-class nearest-neighbor, first tier and second tier.

## Appendix D. Evaluation metrics details for the shape+electrostatics dataset: reciprocal rank, per-class nearest-neighbor, first tier and second tier.

**Table A.6. List of the Protein Data Bank [17, 18] structures used as queries for the track.**

Domain name	Pfam ID	PDB code	chain	residues	Reference
SH2 domain	PF00017	1P13	B	161-243	[66, 67]
SH3 domain	PF00018	1ABO	B	67-113	[68, 69]
Variant SH3 domain (SH3_2)	PF07653	5O99	B	474-527	[70, 71]
PDZ domain	PF00595	2HE2	B	421-499	[72, 73]
PDZ_6 domain	PF17820	3KHF	B	982-1034	[74]
Peptidase family M50 (m50)	PF02163	3B4R	B	111-186	[75]
Bromodomain	PF00439	6CW0	B	10-95	[76]
PHD-finger domain	PF00628	3KV5	D	39-88	[77, 78]
Zinc-finger domain, C2H2 type (zf-C2H2)	PF00096	4ISI	D	472-493	[79, 80]
STAT protein, DNA-binding domain (Stat-binding)	PF02864	5D39	D	277-413	[81, 82]

**Table C.7. Per-class nearest-neighbor for the shape-only dataset.**

Method	SH3	SH3_2	SH2	PDZ	PDZ_6	m50	STAT	zf-C2H2	PHD	Bromodomain	SH3-like	PDZ-like
Class size	115	33	92	128	26	4	6	55	53	64	148	154
3DZD	0	0	1	1	0	0	0	1	1	1	1	1
ProteinNet	0	0	0	0	0	0	0	0	0	0	1	0
APPFD	0	0	0	1	0	0	0	1	1	0	0	1
PWKSM	0	0	0	1	0	0	0	0	0	0	0	1
DGCNN	0	0	0	0	0	0	0	0	0	0	0	0

**Table C.8. Reciprocal rank for the shape-only dataset.**

Method	SH3	SH3_2	SH2	PDZ	PDZ_6	m50	STAT	zf-C2H2	PHD	Bromodomain	SH3-like	PDZ-like
Class size	115	33	92	128	26	4	6	55	53	64	148	154
3DZD	0.143	0.033	1	1	0.030	0.002	0.021	1	1	1	1	1
ProteinNet	0.071	0.036	0.25	0.5	0.25	0.005	0.002	0.010	0.071	0.067	1	0.5
APPFD	0.5	0.042	0.1	1	0.2	0.01	0.022	1	1	0.167	0.333	1
PWKSM	0.333	0.053	0.125	1	0.333	0.015	0.042	0.083	0.042	0.333	0.333	1
DGCNN	0.333	0.083	0.167	0.5	0.333	0.037	0.006	0.042	0.1	0.333	0.25	0.5

**Table C.9. Per-class first tier for the shape-only dataset.**

Method	SH3	SH3_2	SH2	PDZ	PDZ_6	m50	STAT	zf-C2H2	PHD	Bromodomain	SH3-like	PDZ-like
Class size	115	33	92	128	26	4	6	55	53	64	148	154
3DZD	0.35	0.03	0.05	0.42	0.00	0.00	0.00	0.24	0.34	0.17	0.70	0.39
ProteinNet	0.26	0.03	0.16	0.28	0.04	0.00	0.00	0.00	0.06	0.05	0.35	0.31
APPFD	0.40	0.03	0.15	0.22	0.04	0.00	0.00	0.15	0.25	0.13	0.44	0.27
PWKSM	0.18	0.06	0.21	0.26	0.08	0.00	0.00	0.11	0.08	0.08	0.20	0.29
DGCNN	0.19	0.09	0.13	0.25	0.12	0.00	0.00	0.04	0.08	0.09	0.26	0.30

**Table C.10. Per-class second tier for the shape-only dataset.**

Method	SH3	SH3_2	SH2	PDZ	PDZ_6	m50	STAT	zf-C2H2	PHD	Bromodomain	SH3-like	PDZ-like
Class size	115	33	92	128	26	4	6	55	53	64	148	154
3DZD	0.55	0.45	0.13	0.72	0.04	0.00	0.00	0.33	0.45	0.27	0.79	0.69
ProteinNet	0.55	0.09	0.43	0.51	0.08	0.00	0.00	0.02	0.15	0.13	0.68	0.60
APPFD	0.58	0.12	0.30	0.53	0.08	0.00	0.00	0.18	0.34	0.23	0.66	0.59
PWKSM	0.39	0.12	0.39	0.49	0.12	0.00	0.00	0.18	0.11	0.20	0.46	0.56
DGCNN	0.37	0.09	0.33	0.39	0.12	0.00	0.00	0.20	0.25	0.16	0.49	0.52



**Table D.11. Per-class nearest-neighbor for the shape+electrostatics dataset.**

Method	SH3	SH3_2	SH2	PDZ	PDZ_6	m50	STAT	zf-C2H2	PHD	Bromodomain	SH3-like	PDZ-like
Class size	115	33	92	128	26	4	6	55	53	64	148	154
3DZD	0	0	0	1	0	0	0	1	1	0	1	0

**Table D.12. Reciprocal Rank for the shape+electrostatics dataset.**

Method	SH3	SH3_2	SH2	PDZ	PDZ_6	m50	STAT	zf-C2H2	PHD	Bromodomain	SH3-like	PDZ-like
Class size	115	33	92	128	26	4	6	55	53	64	148	154
3DZD	0.5	0.167	0.333	1	0.010	0.003	0.030	1	1	0.5	1	0.5

**Table D.13. Per-class first-tier for the shape+electrostatics dataset.**

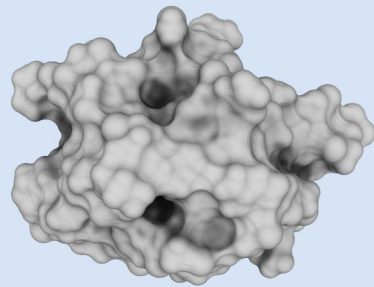
Method	SH3	SH3_2	SH2	PDZ	PDZ_6	m50	STAT	zf-C2H2	PHD	Bromodomain	SH3-like	PDZ-like
Class size	115	33	92	128	26	4	6	55	53	64	148	154
3DZD	0.50	0.12	0.14	0.20	0.00	0.00	0.00	0.36	0.28	0.23	0.58	0.19

**Table D.14. Per-class second tier for the shape+electrostatics dataset.**

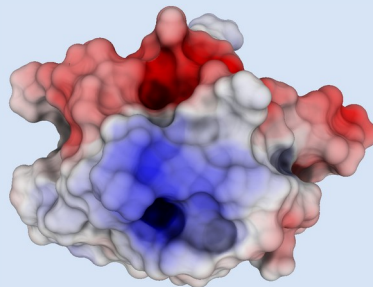
Method	SH3	SH3_2	SH2	PDZ	PDZ_6	m50	STAT	zf-C2H2	PHD	Bromodomain	SH3-like	PDZ-like
Class size	115	33	92	128	26	4	6	55	53	64	148	154
3DZD	0.68	0.36	0.38	0.55	0.00	0.00	0.00	0.51	0.36	0.36	0.74	0.74

Shape  
Retrieval  
Contest  
2021  
On  
Protein  
Surfaces

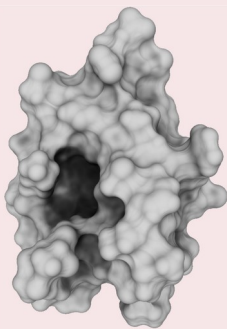
Queries: 10 proteins  
Datasets: 554 proteins



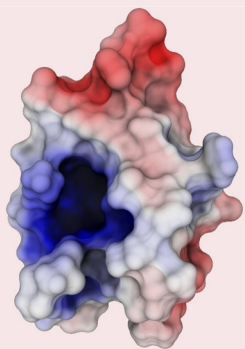
Shape only



Shape + electrostatics



Shape only



Shape + electrostatics

