

FireProt^{ASR}: A Web Server for Fully Automated Ancestral Sequence Reconstruction

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Abstract

There is a great interest in increasing proteins' stability to widen their usability in numerous biomedical and biotechnological applications. However, native proteins cannot usually withstand the harsh industrial environment, since they are evolved to function under mild conditions. Ancestral sequence reconstruction is a well-established method for deducing the evolutionary history of genes. Besides its applicability to discover the most probable evolutionary ancestors of the modern proteins, ancestral sequence reconstruction has proven to be a useful approach for the design of highly stable proteins. Recently, several computational tools were developed, which make the ancestral reconstruction algorithms accessible to the community, while leaving the most crucial steps of the preparation of the input data on users' side. FireProt^{ASR} aims to overcome this obstacle by constructing a fully automated workflow, allowing even the unexperienced users to obtain ancestral sequences based on a sequence query as the only input. FireProt^{ASR} is complemented with an interactive, easy-to-use web interface and is freely available at <https://loschmidt.chemi.muni.cz/fireprotasr/>.

Key words: ancestral sequence reconstruction; ancestral enzymes; evolution; phylogeny-based analysis; protein stability

Introduction

Proteins are widely used in numerous biomedical and biotechnological applications. Native proteins have mainly evolved under mild intracellular conditions [1]. Therefore, their applicability is often limited in the harsh industrial environments characterized

by inhospitable temperature, extreme pH, high pressure or the presence of organic co-solvents. As a result, there is a continuous interest in increasing protein stability. New approaches in the field of protein engineering, such as fluorescence-activated cell sorting and microfluidics, have widened the throughput of

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directed evolution experiments. However, saturation mutagenesis of all positions and systematic re-combinations of many single-point mutations of the protein of interest is often out of reach.

In the past decades, various computational methods were designed to unburden costly and laborious experimental work by narrowing down the search space for potential stabilizing mutations. Most of those methods can be assigned to one of the three categories: (i) machine learning, (ii) force-field-based predictions and (iii) molecular evolution. Each category has its advantages and shortcomings [2]. Machine-learning methods are able to unearth hidden features and dependencies overreaching the current state of expert knowledge, while still struggling with the insufficient size, quality and diversity of the experimental data, essential for training and validation of statistically significant models. Force-field-based approaches are a robust solution for the prediction of protein stability; however, they rely on the high-resolution protein structures that are available for only a small fraction of the known proteins. Evolution-based approaches do not suffer from these limitations due to the rapid growth of the sequence databases. However, this continuous growth widens the search space and increases noise in the data, requiring laborious and time-demanding manual corrections from the side of the user with expert knowledge of the system of interest. Inexperienced user may not therefore utilize evolution-based methods effectively to obtain accurate and reliable results.

The two most widely used evolution-based methods for stability engineering are ancestral sequence reconstruction (ASR) and consensus design. Both methods start with the multiple-sequence alignment (MSA) of the set of relevant homolog sequences. Consensus design relies on the simple analysis of the conservation of the amino acids on the individual positions in the sequence alignment. As a result, it cannot account for the coevolution of the residues located in the sites responsible for the protein's activity [3] and is utilized mostly as a part of the hybrid workflows [4, 5]. In comparison, ASR goes much further by also considering evolutionary information depicted by the phylogenetic tree. This inclusion of the evolutionary distances inscribed into the phylogenetic tree is mostly negligent at the positions with low Shannon entropy; however, the discrepancies grow stronger with noisy MSA [6]. ASR is a probabilistic method that explores the deep evolutionary history of homolog sequences to reassemble protein's evolutionary trajectory [7]. ASR is able to unearth sequences of the long-extinct genes and organisms from which the current ones evolved and is, therefore, an invaluable tool in the field of evolutionary biology [8, 9]. ASR has also been shown to be a very effective strategy not only for thermostability engineering [10, 11], but also for improving other protein's characteristics such as specificity [12], activity, or expression [13]. Furthermore, ASR was previously proven to be an effective strategy for the stabilization of prokaryotic proteins [10, 11], as well as for the improvement of significantly more complex eukaryotic proteins such as cytochrome P450 [14, 15]. Two main algorithms, maximum-likelihood [16, 17] (ML) and Bayesian inference [18] (BI) were designed to infer ancestral sequence from MSA and phylogenetic tree. Many tools were built over the years to make those algorithms accessible to the community. However, the requirement of the MSA of carefully selected homologs and the rooted phylogenetic tree are still huge limiting factors for the general use of ASR method by the non-expert users.

FireProt^{ASR} addresses those limitations by introducing one-stop-shop solution for the ancestral sequence reconstruction. It covers all steps of ancestral inference including search for

homolog sequences, selection of the biologically relevant subset of the sequences, construction of the multiple-sequence alignment, construction and rooting of the phylogenetic tree and finally the ancestral inference with the use of ML. Our computational workflow is fully automated and removes the need for extensive expert knowledge of the system of interest as well as employed bioinformatics tools. Furthermore, a novel algorithm based on the localized weighted back-to-consensus analysis was utilized to resolve an issue of the ancestral gaps reconstruction. Assembled workflow and developed web server were thoroughly validated using: (i) *in-house* laboratory experiments, (ii) detailed comparison with three previously published studies and (iii) a large number of proteins representing structurally and functionally different families. FireProt^{ASR} does not require installation and settings of any software packages as the method is implemented in the interactive web interface freely available at: <https://loschmidt.chemi.muni.cz/fireprotastr/>.

Methods

Workflow description

The basic workflow of the FireProt^{ASR} method is outlined in Figure 1. To infer ancestral sequences representing all ancestral nodes of the evolutionary tree in a fully automated way, a set of biologically relevant homologous sequences must be collected from genomic databases and reduced to a suitable size (Phase 1). With the initial set of homologous sequences in hand, several state-of-the-art methods are utilized to construct a multiple-sequence alignment and a phylogenetic tree, which are then used to support the inference of ancestral nodes and reconstruction of ancestral gaps (Phase 2). The FireProt^{ASR} workflow requires no user intervention beyond providing a query sequence and (in the case of enzymes) selecting catalytic residues used to identify a biologically relevant set of homologous sequences. However, it is also possible to start a calculation with a user-defined initial set of homologous sequences, MSA, or even a phylogenetic tree instead of a single sequence, thus skipping the first phase of the calculation.

Phase 1: collection of the initial set of homologous sequences

The query sequence of the target protein in plain text or FASTA format is the only input required from the side of the user. Once the query sequence has been uploaded to the server and checked for validity, searches for the catalytic residues are performed automatically using SwissProt [19] and the Catalytic Site Atlas [20]. The user can also specify the catalytic residues by themselves if no/incorrect catalytic residues are found. Once the catalytic residues and query sequence have been specified, an *in-house* tool called EnzymeMiner [21] is used to collect an initial set of homologous sequences. EnzymeMiner first performs two rounds of PSI-BLAST [22] against the NCBI nr database [23] and then filters out all sequences lacking the designated catalytic residues, thereby ensuring the biological relevance of the remaining homologs. EnzymeMiner searches can yield up to tens of thousands of homologous sequences for large families. If no catalytic residues were selected or provided by the user, BLAST [24] will be used instead of EnzymeMiner, to obtain an initial set of homologous sequences with potentially lower quality.

Next, the FireProt^{ASR} reduces the set of homologous sequences to the required number, which is set to 150 sequences by default. Several filters are applied during this process. First, all homologs

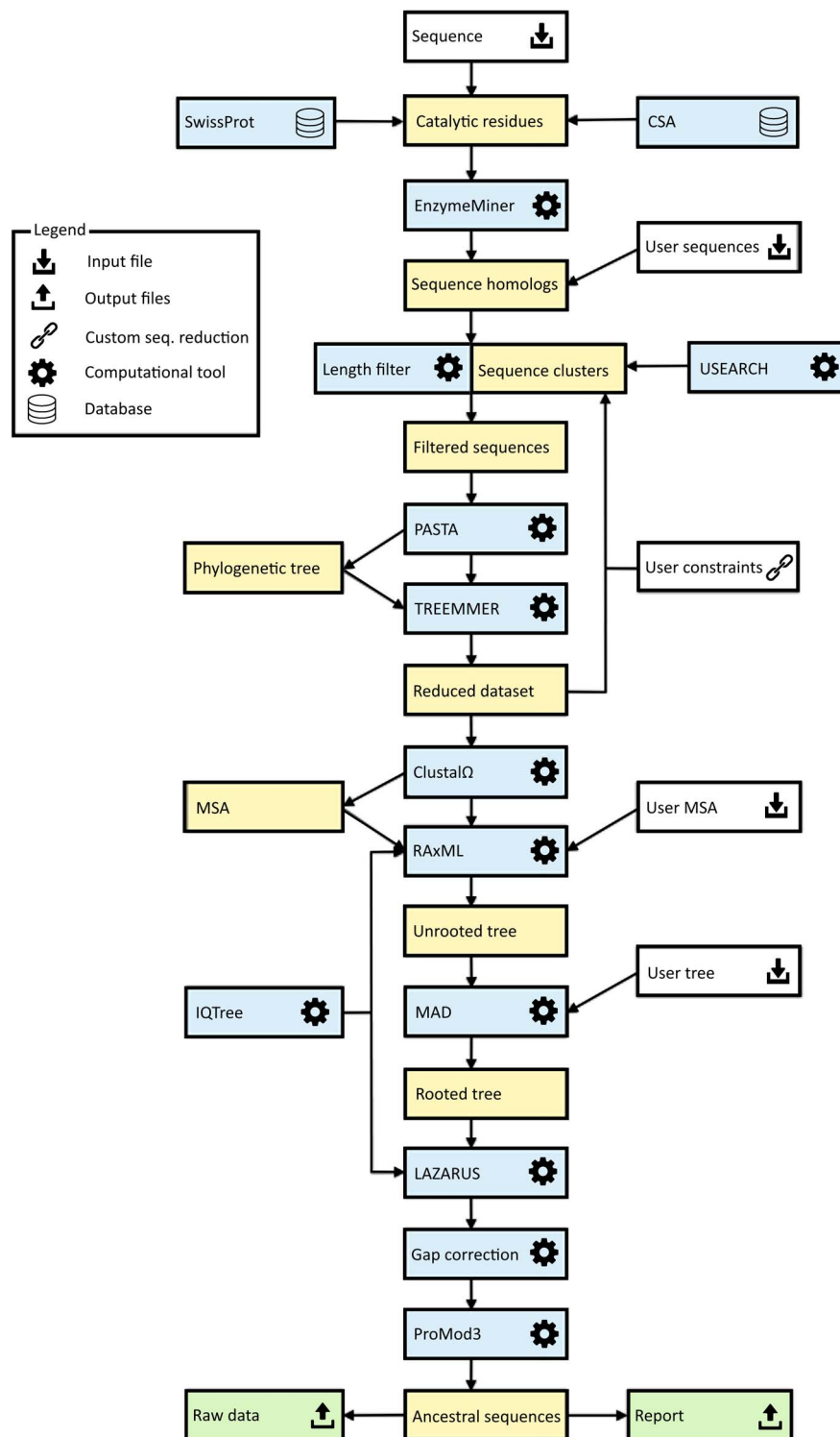


Figure 1. Workflow diagram for the FireProt^{ASR} method. The workflow has two phases: (1) collection of the initial set of homologous sequences and (2) ancestral sequence reconstruction. Colour coding: yellow denotes intermediate results and blue denotes computational tools. Grey and green denote inputs and outputs of the calculations, respectively.

with sequence lengths 20% higher or lower than that of the query sequence are excluded from the initial set. This sequence length normalization is done to remove potential outliers that could lead to a construction of a noisy MSA with many gaps. Second, all homologs whose sequence identity to the query

falls outside a certain range are removed from the initial set. By default, the upper and lower similarity limits are set to 90 and 30%, respectively. This step ensures that the phylogenetic tree is unbiased towards the query sequence while removing distant homologs that would degrade the quality of the sequence

alignment. Third, USEARCH [25] is used to cluster the remaining sequences with 90% sequence identity, and a single sequence is randomly selected from each cluster.

Applying these filters produces a diverse set containing hundreds to thousands of homologous sequences. An initial phylogenetic tree is quickly constructed with the PASTA software suite [26], using MAFFT [27] and the swift neighbour-joining algorithm implemented in FastTree 2.0 [28]. The resulting phylogenetic tree is then forwarded to Treemmer [29], which iteratively prunes leaves from the input tree until a specific number of leaves remains, while minimizing the loss of genetic diversity. The pruned tree is then displayed to the user via the interactive user interface, allowing the user to choose to exclude selected branches or even whole subtrees of the phylogenetic tree from further calculations.

Phase 2: ancestral sequence reconstruction

In the second phase, the ancestral sequences are inferred from the initial set of up to 150 homologs approved by the user. To begin with, a new MSA is constructed from the reduced set of homologous sequences. For this task, Clustal Ω [30] is utilized by default, but other methods will be available in upcoming versions of FireProt^{ASR}. For inference of the final phylogenetic tree, the best-fitting evolutionary matrix must be selected. This is done using one of the modules of the IQTREE package [31]. Alternatively, if the user prefers a specific evolutionary matrix for the biological system of interest, the appropriate model and all the relevant modifiers can be specified manually when setting up the calculation.

The evolutionary model and its parameter settings along with the MSA are then forwarded into RAxML [17], which is used to construct a robust phylogenetic tree. By default, fifty bootstraps are performed at the start of the maximum-likelihood search; since no outgroup is provided, the resulting phylogenetic tree is unrooted. Automated outgroup sequence selection is not straightforward, especially for prokaryotic proteins due to the high frequency of horizontal gene transfers. Rooting of the tree is therefore performed using a minimal ancestor deviation algorithm, which was shown to achieve comparable levels of accuracy to outgroup rooting in trees describing the evolution of eukaryotes, and to surpass both outgroup and midpoint rooting in the case of prokaryotes [32].

The MSA constructed with Clustal Ω , the selected evolutionary model, and the rooted phylogenetic tree from RAxML are used as inputs for the Lazarus method [33], which is implemented using the PAML software package [16]. The Lazarus method was re-implemented for FireProt^{ASR} to enable calculations to be performed without specifying outgroup. Consequently, ancestral sequences of all ancestral nodes are parsed from their posterior probabilities and provided to users in separate files in FASTA format. Additionally, BLASTp [24] is used to search for a template in the PDB database [34], and a model structure of the query sequence is constructed by homology modelling using the ProMod3 program [35]. This model is shown in the web interface to allow users to visualize the differences between the query sequence and the selected ancestor.

Finally, due to the large number of undesirable ancestral gaps inserted into ancestral sequences by Lazarus, a novel algorithm for ancestral gap reconstruction was designed for use in FireProt^{ASR}. This algorithm is based on the principle of localized weighted back-to-consensus because consensus analysis has proven to be an effective approach for increasing proteins' thermal stability [36–38]. To begin with, each terminal node of

the phylogenetic tree is assigned a binary vector of length equal to the length of the corresponding sequence in the MSA. Each position in this vector is assigned a value of -1 or 1 , indicating the presence of a gap or standard amino acid, respectively, at the corresponding position of the relevant sequence. On moving from the terminals towards the root of the tree, the probability of a gap in ancestral node A_n at position i is calculated as $A_{n_i} = \frac{A_{k_i} * t_1 + A_{l_i} * t_2}{t_1 + t_2}$, where A_{k_i} , A_{l_i} are the child nodes of A_n and t_1 , t_2 are the evolutionary distances between A_n and its child nodes. Taking t_3 to be the evolutionary distance between A_n and its parental node, its value can be updated based on the values of t_1 and t_2 as follows: $t_{3_new} = t_3 + \frac{t_1 + t_2}{2}$. This new value is computed before proceeding with the calculation for the parental node; its use increases the relative impact of well-branched subtrees and therefore limits the impact of lone sequences and small subtrees compared to that of well-represented ones. Finally, ancestral sequences are reconstructed based on the scores in the corresponding vector. Positions with values lower than 0 are assigned as gaps, and the remaining amino acids are selected based on their posterior probabilities as estimated by Lazarus. The nature of inconclusive positions with scores in the interval $\langle -0.1, 0.1 \rangle$ is determined based on the frequencies of gaps in the global alignment and the state of the parental node. To include the ancestral gap, frequencies of gaps in the global alignment should reach over 60%, or over 40% if the ancestral gap is present in the parental node sequence. The model case for a single position in the sequence alignment is shown in Figure 2.

Experimental validation

The workflow was experimentally validated using haloalkane dehalogenases as a model enzyme. This enzyme was selected as a typical representative of the α/β superfamily, counting over 100 000 proteins. The sequence of the haloalkane dehalogenase DhaA (UniProt ID P0A3G2) was used as the sole input for the calculation. Six different ancestral sequences were selected and experimentally characterized.

Chemicals and growth media

1-bromobutane and LB medium were purchased from Sigma-Aldrich Co. (St. Louis, MO, USA). IPTG was purchased from Duchefa Biochemie B.V. (Haarlem, The Netherlands). All chemicals used in this work were of analytical grade.

Expression in *Escherichia coli* BL21 (DE3)

Escherichia coli Dh5 α cells were obtained from Invitrogen and *Escherichia coli* BL21 (DE3) from New England Biolabs. The genes for the ancestral dehalogenases were synthesized and subcloned into the expression vector pET21b. The generated plasmids were transformed into chemo-competent *E. coli* BL21 (DE3) cells. Obtained colonies were used to prepare precultures by inoculation into 10 ml of LB medium (with 100 μ g/ml ampicillin) followed by overnight incubation at 37°C and 180 rpm. For expression of each variant, 1 l of LB medium supplemented with 100 μ g/ml ampicillin was inoculated with 5 mL of the appropriate pre-culture (1/200). The flasks were incubated at 37°C and 180 rpm until OD₆₀₀ 0.6–0.8 was reached, then incubated at 20°C for 30 min. β -D-1-thiogalactopyranoside (IPTG, 0.2 mM) was then added for induction, and the culture was incubated at 20°C and 180 rpm overnight. Finally, the culture was harvested by centrifugation at 4500 \times g, 4°C for 15 min, after which the cell pellets were frozen at -80°C until further use.

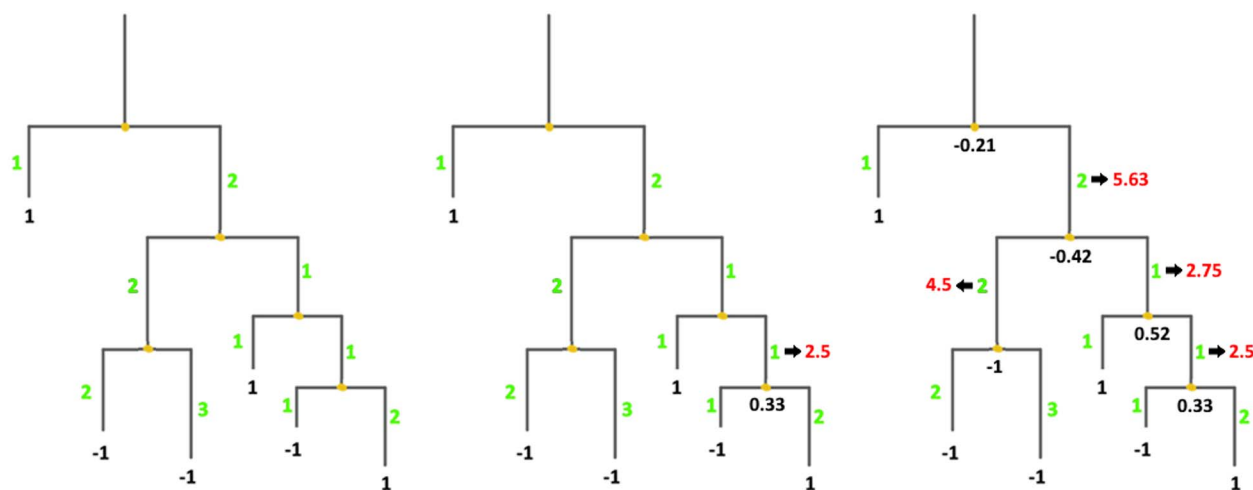


Figure 2. Ancestral gaps reconstruction algorithm. Green colour denotes the initial branch lengths of the phylogenetic tree. Black numbers indicate the values of the vectors of the terminal and the ancestral sequences at the given position in the multiple sequence alignment. Red values show the modified branch lengths that are updated after the calculation of the underlying ancestral node.

Protein purification

The cell pellets were suspended in 50 ml of equilibration buffer (20 mM phosphate buffer pH 7.5 containing 0.5 M NaCl and 10 mM imidazole) and disrupted by sonication with a Hielscher UP200S ultrasonic processor (Hielscher, Germany) four times for 4 min each. Disrupted cells were centrifuged at $13\,000 \times g$ and 4°C for 1 h (Laborzentrifugen, Germany). The crude extract was then collected, filtered and loaded onto a Ni-NTA Superflow Cartridge (Qiagen, Germany) in equilibration buffer. Unbound and weakly bound proteins were washed out using increasing imidazole concentrations. The target enzyme was eluted with purification buffer containing 300 mM of imidazole. The eluted protein was dialyzed three times overnight against 50 mM of phosphate buffer (pH 7.5), after which its purity was checked by SDS-polyacrylamide gel electrophoresis (SDS-PAGE). About, 15% polyacrylamide gels were stained with Instant Blue (Fluka, Switzerland). Protein concentrations were determined by NanoDrop (Sigma-Aldrich, USA). The enzymes were lyophilized using a vacuum pump system for long-term storage.

Circular dichroism (CD) spectroscopy

CD spectra were recorded at 20°C using a spectropolarimeter Chirascan (Applied Photophysics, United Kingdom). Data were collected from 190 to 260 nm, at 100 nm/min with a 1-s response time and 1-nm bandwidth using a 0.1-cm quartz cuvette. Each spectrum shown is the average of five individual scans and was corrected for absorbance caused by the buffer. Collected CD data were expressed in terms of the mean residue ellipticity (θ_{MRE}), which was calculated using the equation:

$$\theta_{\text{MRE}} = \frac{\theta_{\text{obs}} \cdot M_w \cdot 100}{n \cdot c \cdot l}$$

where θ_{obs} is the observed ellipticity in degrees, M_w is the protein molecular weight, n is number of residues, l is the cell path length, c is the protein concentration (0.2 mg/ml) and the factor 100 originates from the conversion of the molecular weight to mg/dmol.

Thermal denaturation

Thermal unfolding was followed by monitoring the ellipticity at 224 nm over the temperature range of $20\text{--}94^\circ\text{C}$, with a resolution of 0.1°C at a heating rate of $1^\circ\text{C}/\text{min}$. Recorded thermal denaturation curves were roughly normalized to represent signal changes between approximately 1 and 0 and fitted to sigmoidal curves using Origin 6.1 (OriginLab Corporation, USA). The melting temperature (T_m) was evaluated as the midpoint of the normalized thermal transition.

Enzymatic haloalkane dehalogenase activity

Dehalogenation activity was assayed using the colorimetric method of Iwasaki et al. [49]. The release of halide ions was analyzed spectrophotometrically at 460 nm using an Eon microplate reader (BioTek, USA) after reaction with mercuric thiocyanate and ferric ammonium sulfate. The reactions were performed at 37°C in 25-ml Reacti Flasks closed with Mininert Valves. The reaction mixtures consisted of 10 ml 100 mM glycine buffer (pH 8.6) and 10 μl of the substrate 1-bromobutane. Reactions were initiated by adding the enzyme to a final concentration of 0.01 (DhaA 172Loc), 0.0065 (DhaA 172Glob), 0.0052 (DhaA 230Glob), 0.028 (DhaA 238Loc) or 0.014 mg/ml (DhaA 238Glob). Reactions were monitored by withdrawing 1 ml of samples from the reaction mixture after 0, 5, 10, 15, 20 and 30 min. The samples were immediately mixed with 0.1 ml of 35% nitric acid to stop the reaction. Dehalogenation activities were quantified as rates of product formation over time. Each activity was measured in three independent replicates.

Enzymatic luciferase activity

Luminescence activity measurements were performed with a FLUOstar OPTIMA Microplate reader (BMG Labtech, Germany) using coelenterazine as the substrate at 37°C . A 25 μl of sample of purified enzyme at a concentration of about 1 mg/ml was placed into a microtiter plate well. After baseline collection for 10 s, the luminescence reaction was initiated by adding 225 μl of 8.8 μM coelenterazine in reaction buffer (100 mM potassium phosphate buffer, pH 7.5). Luminescence was recorded for 72.5 s,

and each sample was measured in at least three independent experiments. The areas of the resulting luminescence intensity peaks in relative luminescence units (RLU) were converted into values in units of RLU/mg/s.

Results

Web server input

The only required input to the web server is a query sequence of the target protein in plain text or FASTA format. Alternatively, one can upload a FASTA file containing an initial set of sequence homologs or a multiple sequence alignment (MSA). Rooted and unrooted phylogenetic trees in the standard Newick format can also be provided. When performing calculations in basic mode, only the table containing the essential residues is available to the user. Essential residues are identified automatically by searching in SwissProt [19] and mCSA [20]. However, the initial selection can be changed by the user. The default values and settings of individual computational tools are optimized to provide reliable results for most systems. Operating in advanced mode expands the list of modifiable parameters to include those related to: (i) the thresholds of the homolog identity filters and sequence clustering, (ii) selection of the evolutionary model and (iii) construction of the phylogenetic tree. Advanced mode allows experts to fine-tune the calculation's parameters based on the studied biological system, which may be useful when dealing with particularly small or large protein families.

Selection and reduction

Upon submission, a unique identifier is assigned to each job to track the calculation. The 'calculation browser' informs the user about the status of the individual steps in the ancestral sequence reconstruction workflow. Once the first phase of the job is finished, the initial phylogenetic tree is displayed to the user using a strongly updated adaptation of PhyloTree library (Figure 3A) [39], together with the table of removed sequences (Figure 3B). By clicking on the individual leaves of the phylogenetic tree, the user can exclude selected sequences from future calculations. Furthermore, whole subtrees can be removed by choosing this option in the menu of the selected ancestral node. The MSA of the homologous sequences can be also visualized by switching to the multiple sequence alignment tab. This mode is intended for the expert users with the greater knowledge of the system of interest as it allows for the removal of the noise and outliers from the initial set of homolog sequences. If the expert mode is utilized, it is recommended to exclude the sequences that do not share the function similar to the query protein or that cause a significant disturbance in the MSA.

Web server output

The calculation's progress can be tracked in the 'calculation browser' similarly to the selection step. Once finished, users can either download the results in the zipped archive directly from the calculation page or navigate to the 'Result page' for further analysis. The 'Result page' is organized into several panels allowing users to interactively visualize and design ancestral enzymes.

Protein visualization

The homology model of the query protein predicted by ProMod3 is interactively visualized in the web browser using the JSmol

applet [40] (Figure 3D). Users can switch between different visualization styles such as backbone, wireframe or cartoon and change the quality of the visualized structure. It is also possible to visualize the differences between the query and the selected ancestral sequence on the modelled protein structure: substitutions and deletions are shown in blue and red, respectively, while insertions are indicated by regions between red and yellow residues.

Ancestral tree panel

The 'ancestral panel' shows the final phylogenetic tree constructed by RAxML [17] along with further information about the precalculated ancestral sequences (Figure 3E). By selecting any of the ancestral nodes, it is possible to either (i) visualize the differences between a wild-type protein and the selected ancestor node on the protein structure or (ii) open a new window providing an overview of the posterior probabilities for individual amino acids in the sequence of the selected ancestor (Figure 3G). Posterior probabilities are shown in the bar-styled sequence logo together with the percentages for each considered amino acid, and each bar is expanded with information about the charge and hydrophobicity of the most probable amino acids. The bar representation was in part derived from the SequenceLogo library [41]. The user can edit the ancestral sequence and store it as a new user-defined ancestor (Figure 3F). This option is useful for the experts with more in-depth knowledge of the system of interest and allows to force some specific mutations, e.g., the mutations with the previously known effect on proteins stability, into the constructed ancestral sequence. It can also be used to bring some biological insight into the positions with noisy posterior probabilities. Furthermore, the ancestral sequences' MSA can be visualized in the multiple sequence alignment tab for further analysis.

Sequence designer

The 'Sequence designer' panel allows users to manage and edit user-defined ancestral sequences. Additionally, new sequences can be created by modifying existing custom ancestors (Figure 3C). Differences between the query sequence and custom ancestors can also be visualized on the protein structure in this panel. All prepared designs can be downloaded in one zipped archive together with the original ancestors and the structure prepared by homology modelling.

Web server experimental validation

In one of our previous studies, we have presented experimental characterizations of six inferred ancestral proteins from haloalkane dehalogenase subfamily II [10]. Relative to their contemporary counterparts, these ancestral proteins exhibited higher thermal stability (by 8–24°C), improved yields and broadened substrate specificity. Those ancestral sequences were reconstructed by clustering an initial set of homologous sequences that was reduced by inspection in the sequence-editing program BioEdit [42]. A multiple sequence alignment was then manually curated using a structure-guided alignment of eight proteins from HLD-II and poorly conserved regions were removed from the alignment. The topology of the phylogenetic tree was optimized by subtree pruning and re-grafting, and the tree's root was established using outgroup selected on the basis of expert judgement. Finally, the ancestral sequences and positioning of gaps were refined by manual inspection.

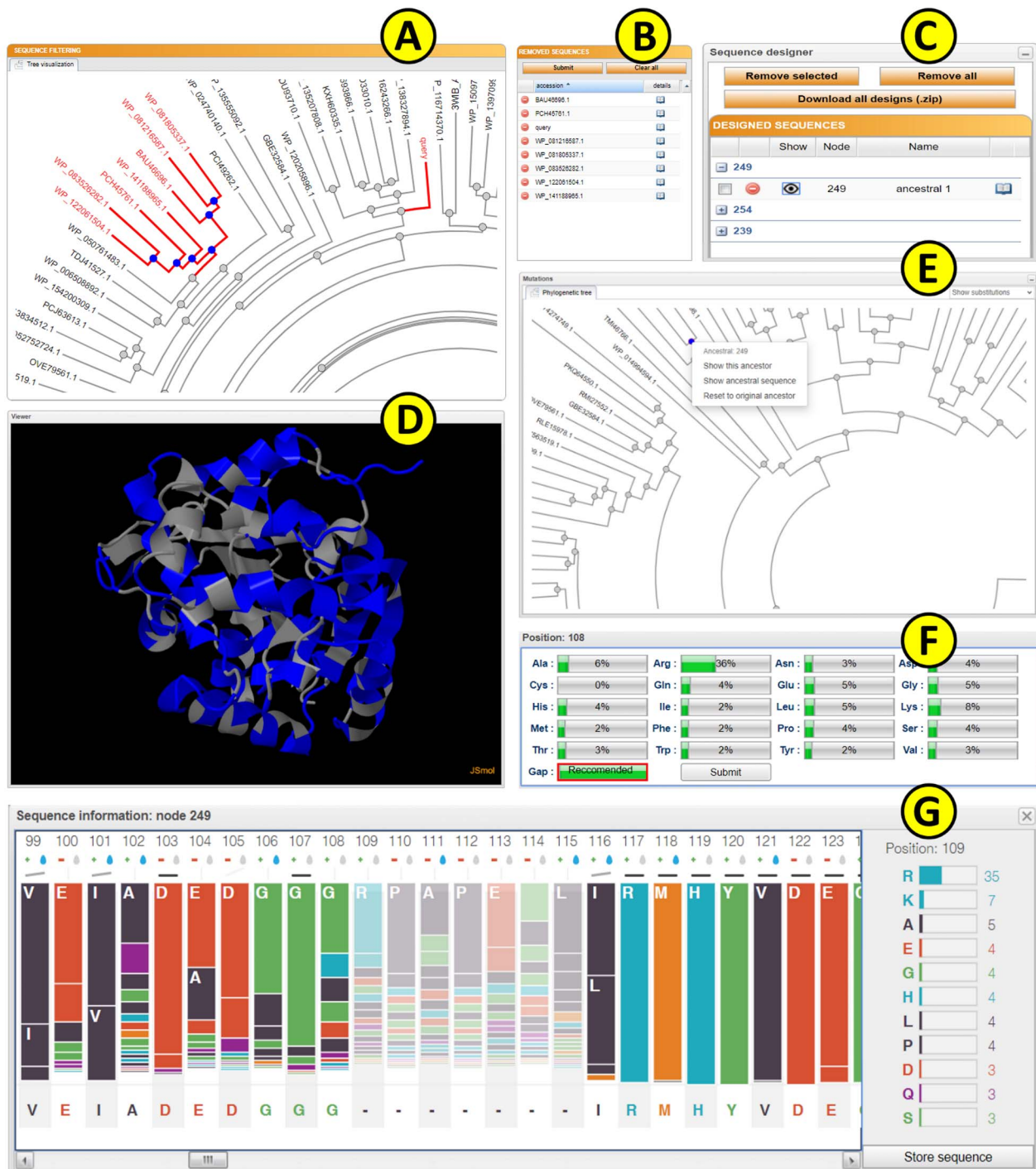


Figure 3. The FireProt^{ASR} graphical user interface showing results obtained for the haloalkane dehalogenase DhaA (UniProt ID P0A3G2, PDB ID 4E46). (A) The sequence-filtering panel allows users to exclude selected branches from the calculation. (B) The reduction table shows the list of removed sequences. (C) The sequence designer allows users to download and edit ancestral sequences. (D) The JSmol viewer provides interactive protein visualization. (E) The mutations panel contains all designed ancestral sequences in the ancestral tree. (F) The edit window enables amino acid substitutions at individual positions. (G) The sequence information window shows detailed information on selected ancestral sequences.

As part of the validation of FireProt^{ASR}, we tried to replicate these results by using the sequence of haloalkane dehalogenase DhaA (UniProt ID P0A3G2) as the only input query. All steps of the calculation, including homologous sequence selection, multiple sequence alignment construction, phylogenetic rooting

and ancestral reconstruction were carried out automatically. Three pairs of ancestral sequences were selected, each pair containing one 'global' and one 'local' ancestral node (Figure 4A). Global ancestor (Glob) represents ancestral sequence obtained directly from the fully automated workflow, while local ancestor

Table 1. Characteristics of reconstructed and experimentally characterized ancestral haloalkane dehalogenases

Protein code	Expression (% of total protein)	Solubility (%)	Yield (mg/l)	T _m (°C)	HLD act. (μmol/mg-s)	LUC act. (RLU/mg-s)
DhaA wt	17	83.1	91.1	50.56 ± 2.4	0.032 ± 0.0059	n.a.
DhaA 172Loc	23	85.5	74.9	71.60 ± 0.7	0.038 ± 0.0002	1.41 ± 0.26
DhaA 172Glob	21	65.2	88.2	70.04 ± 1.5	0.061 ± 0.0045	n.a.
DhaA 230Loc	20	n.d.	n.d.	n.d.	n.d.	n.d.
DhaA 230Glob	23	84.8	108.5	72.14 ± 0.4	0.061 ± 0.0118	n.a.
DhaA 238Loc	23	63.2	74.9	70.36 ± 0.6	0.014 ± 0.0021	353.5 ± 14.58
DhaA 238Glob	19	83.3	94.4	76.19 ± 0.2	0.030 ± 0.0012	3.18 ± 0.33

Notes: DhaA, haloalkane dehalogenase from *Rhodococcus rhodochrous* NCIMB 13064; wt, wild type; Loc, ancestral protein inferred from local alignment; Glob, ancestral protein inferred from global alignment; T_m, melting temperature; HLD act., haloalkane dehalogenases activity; LUC act., luciferase activity; n.d., not determined due to poor solubility of this protein; n.a., not active under tested conditions.

catalytic activity. Moreover, inference based on both haloalkane dehalogenases and luciferases led to the discovery of the very interesting enzyme ancHLD-Rluc, which exhibits dual dehalogenase and monooxygenase activity. This experimental validation provides direct experimental evidence of the good functionality and reliability of the fully automated version of FireProt^{ASR}.

Additionally, results obtained using FireProt^{ASR} were thoroughly and quantitatively compared to three previously published experimental studies. For this purpose, Euclidean distance [43], and the Subtree prune and regraft distance [44] were calculated to compare the trees obtained from the FireProt^{ASR} and published literature. The two trees were also graphically compared using the Jaccard index utilizing ColorBrewer [45] scheme. Detailed comparison of all three experimental studies with the results produced by FireProt^{ASR} server is attached in [Supplementary Data 1–3](#), available online at <https://academic.oup.com/bib>. Finally, the robustness and reliability of the FireProt^{ASR} server was tested using 60 diverse proteins from various protein families (see [Supplementary Data 4](#) available online at <https://academic.oup.com/bib>).

Discussion

ASR has been shown to be a very effective strategy for the protein thermostability engineering and as such was implemented in various computational tools using maximum-likelihood (FastML [46], RaxML [17], Ancestors [47]) or Bayesian inference (HandAlign [48], MrBayes [18]) methods. However, a significant limitation of those methods is that they require complex input data to be uploaded by the users. Those requirements are reaching from a simple set of homolog sequences to the MSA or even rooted phylogenetic tree, leaving the most crucial and laborious parts of the calculation in the hands of the users. Non-expert users without the deep knowledge of the bioinformatics tools and the system of interest are therefore hindered from the successful use of the ASR method.

FireProt^{ASR} is a web server that aims to provide users with one-stop-shop solution for the ancestral sequence reconstruction. FireProt^{ASR} requires minimal input from the users, and the whole calculation can be processed from a single protein sequence, set of homolog sequences, MSA and phylogenetic tree. All steps of the calculation, including the search for biologically relevant homolog sequences, dataset reduction and the ancestral reconstruction are automated. Moreover, a novel algorithm based on localized weighted back-to-consensus analysis is implemented to resolve an issue with ancestral gap reconstruction. FireProt^{ASR} web server is also complemented by an easy-to-use web interface that allows users to interactively analyze

sequences of the individual ancestral nodes together with the ability to design their own ancestral sequences based on the posterior probabilities of the existing nodes.

The robustness and reliability of the results produced by the FireProt^{ASR} workflow was evaluated by experimental characterization of six ancestral sequences of haloalkane dehalogenase from HLD-II subfamily. With the exception of the local variant of the ancestral node 230, all designed ancestral sequences are soluble and also retain high expressibility and yields on the levels comparable to the DhaA wild type. However, the thermal stability has increased by over 20°C and global variants 172 and 230 have also increased the HLD activity by two-fold. Increase in HLD activity cannot be observed in the constructed local variants that utilize smaller subsets of homolog sequences, and thus only a limited amount of evolutionary information. This would encourage the usage of the global variants for the design of highly stable and active proteins. However, more focused view using a localized variants of the ancestral nodes can provide some useful results as can be observed in the local variant of the node 238 that shows both dehalogenase and monooxygenase activity. High thermal stabilization was also achieved in those variants.

Finally, the results provided by the FireProt^{ASR} web server are consistent with the designs presented in the published literature as the fully automatized designs obtained by FireProt^{ASR} method maintain high sequence similarity (>90%) with the manually designed and curated ancestors. Finally, the comprehensive analysis of approximately 60 different proteins from various protein families have proven the robustness and reliability of the presented method.

The full automation of the FireProt^{ASR} method eliminates the need to select, install and evaluate individual tools, optimize their parameters and interpret intermediate results. Together with its general applicability for a wide range of protein families, FireProt^{ASR} makes the procedure of ancestral reconstruction accessible to the users without any prior expertise in bioinformatics, and the intuitive web interface allows for a further analysis utilizing both sequence and structural information.

Key Points

- FireProt^{ASR} is a web service for a fully automated design of stable proteins using ancestral sequence reconstruction and is accompanied by an interactive and easy-to-use interface.

- FireProt^{ASR} allows users to utilize ancestral reconstruction without prior knowledge of the necessary bioinformatics tools and the biological system.
- The robustness and reliability of the FireProt^{ASR} method were thoroughly tested by both laboratory experiments and by comparing predictions with the results published in scientific literature.
- Laboratory characterization of the ancestral designs showed up to 26°C improvement in thermostability and some of the proteins poses even dual catalytic activity.

Data availability

All data validating the robustness and accuracy of our service are available in the Supplementary materials 1-4. Web service and tutorials are freely available at <https://loschmidt.chemi.muni.cz/fireprotasr/>.

Supplementary Data

Supplementary data are available online at *Briefings in Bioinformatics*.

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