B-Cell Epitope Mapping from Eight Antigens of *Candida albicans* to Design a Novel Diagnostic Kit: An Immunoinformatics Approach

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Abstract

Invasive candidiasis is an emerging fungal infection and a leading cause of morbidity in health care facilities. Despite advances in antifungal therapy, increased antifungal drug resistance in *Candida albicans* has enhanced patient fatality. The most common method for *Candida albicans* diagnosing is blood culture, which has low sensitivity. Therefore, there is an urgent need to establish a valid diagnostic method. Our study aimed to use the bioinformatics approach to design a diagnostic kit for detecting *Candida albicans* with high sensitivity and specificity. Eight antigenic proteins of *Candida albicans* (HYR1, HWP1, ECE1, ALS, EAP1, SAP1, BGL2, and MET6) were selected. Next, a construct containing different immunodominant B-cell epitopes was derived from the antigens and connected using a suitable linker. Different properties of the final construct, such as physicochemical properties, were evaluated. Moreover, the designed construct underwent 3D modeling, reverse translation, and codon optimization. The results confirmed that the designed construct could identify *Candida albicans* with high sensitivity and specificity in serum samples of patients with invasive candidiasis. However, experimental studies are needed for final confirmation.

Keywords *Candida albicans* · Epitope mapping · B-cell epitope · Diagnostic kit · Antigens

Introduction

*Candida* is a significant cause of fungal infections, with *Candida albicans* responsible for 70–90% of candidiasis infections worldwide (Bhakt et al. 2018; Erwig and Gow 2016). As the natural flora of the human body, *Candida albicans* is found on different parts of the body, such as mucosal surfaces, skin, mouth, gastrointestinal tract, and genitals (Antinori et al. 2016; León et al. 2014). The pathogenicity of *Candida albicans* is due to the transformation from single-celled yeast to long multicellular fibers called hyphae. The filamentous form can bind and penetrate the host tissue faster than the single-cell form. Therefore, the production of hyphae indicates the beginning of infection (Nobile et al. 2012; Sachivkina et al. 2020; Tsui et al. 2016). Today, the prevalence of invasive candidiasis as an opportunistic clinical infection is increasing for different reasons (Chen et al. 2020; Lewis and Williams 2017; Thomson et al. 2015). Due to pregnancy and the use of contraceptive estrogens, women are at high risk of fungal infection. *Candida albicans* have been isolated from vaginal biopsies of 20–50% of asymptomatic women of gestational age (Brown et al. 2018).
2019; Gharaghani et al. 2018). Azole antifungal agents such as miconazole, ketoconazole, and fluconazole are effective treatments for *Candida albicans* infections. Nevertheless, increasing drug resistance has made this treatment problematic. Therefore, the early detection of fungal infections is impressive for treatment, prevention of drugs overuse, and reduction of costs (Thomson et al. 2015).

Various identification methods such as culture on Corn Mill Agar-Tween 80, sugar uptake test by UX 20C API kit, and Germ tube test have been used to identify *Candida* species (Souza et al. 2015). Though blood culture is the standard method for diagnosing candidiasis, for the low sensitivity of the method (about 50%), positive results appear only in the advanced stages of infection, which significantly impairs the treatment process. Although molecular methods such as Polymerase Chain Reaction (PCR), cell wall, and cytoplasm antigens identification are used to early detect the disease, none of these methods has been widely approved (He et al. 2015). Immunological methods like Immunofluorescence Antibody Test (IFAT), Enzyme-linked Immunoassay (ELISA), and immunoblotting are more sensitive than molecular methods and help assess the host’s immune response (Buonfrate et al. 2015; Levenhagen and Costa-Cruz 2014). Considering the production of recombinant antigens and synthetic peptides that have immunogenic epitope domains, desirable improvement has been achieved in serological methods (Señido et al. 2013). Extensive efforts have been made to establish a valid serological test to diagnose invasive candidiasis. Detecting specific antibodies against yeast antigens in patient samples is one of the documented methods (Calandra et al. 2016; Ma et al. 2013). Based on the results of several studies, various proteins from *Candida albicans* were regarded as antigens. Proteins that stimulate the immune system and produce antibodies are found in two categories. The first group contains proteins that appear specifically in the invasive phase of the disease, such as HYR1, HWP1, ECE1, and ALS. The expression of the second group, including EAP1, SAP1, BGL2, and MET6, increases in the pathogenic stage (Araújo et al. 2017; Lain et al. 2008; Mayer et al. 2013; Wang et al. 2015).

The last decade has witnessed a considerable increase in the processing speed of computers and the storage capacity of information. On the one hand, sequencing the genomes of many prokaryotes, eukaryotes, and many proteins gives new insights into biological data analysis (Bansal 2005). In addition, the design, production, and purification of each of the essential antigens for the diagnosis of infectious diseases are time-consuming, requiring high costs and controlling different processes. Thus, combining these various antigenic proteins as a chimeric protein in a vector can be a valuable way to reduce the time and cost of processing. This study aimed to bioinformatically design and clone the recombinant *Candida albicans* multi-epitope antigen to diagnose invasive candidiasis using the ELISA system.

**Methods**

**Study Design**

The current study was done in three following steps. In the first step, according to the results of *Candida albicans* proteomic microarray and literature review on *Candida albicans* antigens, the antigenic proteins of *Candida albicans* were chosen. Invasive phase-specific proteins, including HWP1, ALS, ECE1, and HYR1, and proteins associated with incremental expressions of the pathogenic stage, such as BGL2, EAP1, MET6, and SAP1, were the select antigens. In the next step, the above-mentioned antigens from different *Candida albicans* strains were aligned via CLC Genomics Workbench 20 to select the consensus regions. Next, using immunoinformatics servers, the antigenic regions and B cell epitopes of each protein were defined, and then the final epitopes were joined by an appropriate linker. Afterward, the structural analysis of the final construct, such as 3D modeling, physicochemical properties, and validation, was performed. In the ultimate step, the protein sequence was reverse-translated to gene sequence and optimized for expression in the *Escherichia coli* (*E. coli*) host.

**Sequence Retrieval**

The amino acid sequences of BGL2, EAP1, MET6, SAP1, HWP1, ALS, ECE1, and HYR1 were retrieved from the Protein knowledgebase (UniProtKB) database (www.uniprot.org) in the FASTA format. Also, CLC Genomics Workbench 20 was applied for antigenic protein aligning. The reviewed sequences of ALS protein were aligned to achieve a protected sequence of this protein.

**Immunoinformatics Analyses**

**Antigenicity Prediction**

The EMBOSS server 6.6.0.0 (https://www.bioinformatics.nl/cgi-bin/emboss/antigenic) was used to define antigenic regions of the proteins. The performance of this server is based on the Kolaskar and Tongaonkar technique, which forecasts protein antigenic regions using physicochemical properties of amino acid residues. In the present study, initial parameters such as antigenic region with a minimum length of 6 and output report format EMBOSS motif were utilized to establish the antigenic region of the target proteins (Rice et al. 2000).
Prediction of Linear B Cell Epitopes

B cell epitopes considerably affect antibody production. For this purpose, in this study, B cell epitopes were determined using three servers to increase the prediction accuracy. In the beginning, for predicting linear B cell epitopes, ABCpred at (https://webs.iiitd.edu.in/raghava/abcpred/ABC_submission.html) was utilized to predict linear B cell epitopes in the antigenic sequence using an artificial neural network algorithm. In addition, ABCpred is the first server developed according to the recurrent neural network by fixed-length samples with about 65.93% accuracy (Chatterjee et al. 2021; Saha and Raghava 2006).

The BCPRED 2.0 server (https://webs.iiitd.edu.in/raghava/bcepred/bcepred_submission.html) was also applied to predict linear B cell epitopes. This server predicts B cell considering the physicochemical properties, including accessibility, hydrophilicity, polarity, flexibility/mobility, exposed surface, turns, or a combination of these features. Based on different physicochemical properties, the server can predict epitopes with various accuracies from 52.92 to 57.53%. In the case of using flexibility, hydrophilicity, polarity, and surface features simultaneously at a threshold of 2.38, this server can predict epitopes with 58.7% accuracy (Saha and Raghava 2004).

BepiPred-2.0 (http://www.cbs.dtu.dk/services/BepiPred/) is another server that predicts B cell epitopes based on epitope information from solved 3D structures and a wide range of linear epitopes downloaded from the IEDB database. About 50 sequences can be submitted simultaneously, with a length of 10–6000 amino acids. The prediction for each sequence usually takes a few minutes (Jespersen et al. 2017).

Multi-epitope Candidate Construction

The B cells epitope candidates, which were selected by relevant servers and got the highest scores, were ligated using a suitable amino acid linker. Eventually, an amino acid structure with multiple epitope characteristics was constructed.

Physicochemical Properties Assessment

The ProtParam tool at the EXPASY server (http://web.expasy.org/protparam/), which is the proteomics server of the Swiss Institute of Bioinformatics, was utilized for calculating different physicochemical features of proteins. The parameters consisted of theoretical pI (isoelectric point), molecular weight (kilodalton), instability index, estimated aliphatic index, and grand average of hydropathicity (GRAVY) (Gasteiger et al. 2005).

Tertiary Structure Prediction of Designed Construct

Galaxy TBM (http://galaxy.seoklab.org/cgi-bin/submit.cgi?type=TBM) was applied to predict the tertiary protein structures. Galaxy TBM performs homology modeling using comparative analysis of the amino acid sequence with those of proteins whose structures are experimentally determined to predict the structure of the regions of proteins that do not have a high degree of homology with other proteins (Ko et al. 2012).

Tertiary Structure Validation

The ProSA-web server (https://prosa.services.came.sbg.ac.at/prosa.php) is one of the servers used for evaluating the 3D structure of the protein. The software mainly revolves around the alpha carbon of amino acid residues to determine the atomic interactions of the protein query. The energy level of the query structure, depicted via a color code system, determines the z-score, which illustrates the quality of the protein structure. Natively folded proteins lie within a defined z-score value, which is used as the reference range with which query proteins are evaluated. The ProSA-web server is accessible elsewhere (Wiederstein and Sippl 2007).

The conformation of proteins is defined by the $\phi$ and $\psi$ values, which are the allowable freedom of rotation in Cα-N and Cα-C of peptide bonds, respectively. The allowable torsion angles between atoms are those that do not cause any steric hindrance; thus, the spherical shape and van der Waals radii were meticulously analyzed. The models were categorized into three groups using the above
parameters: Favored, allowed, and outlier regions. The plot was generated using PROCHECK online software (https://saves.mbi.ucla.edu/) (Lovell et al. 2003).

The ERRAT online software (http://services.mbi.ucla.edu/ERRAT/) was employed to verify the 3D structure of proteins. The program uses non-covalently bonded atom–atom interactions and determines whether regions of the target proteins are correctly folded or not. Six types of atomic interactions, including C–C, C–N, C–O, N–N, and N–O, are evaluated where C, N, and O denote carbon, nitrogen, and oxygen, respectively. The software posits that the distance between the aforementioned non-bonded interaction should not surpass 3.5 Å, and the covalently bonded atoms within the same polypeptide strand should be excluded from the analysis (Colovos and Yeates 1993).

Molecular Dynamics (MD) Simulations
Understanding macromolecular functioning requires dynamic systems simulations with biologically relevant sizes and time scales. As an alternative to atomistic simulations, coarse-grained representations combined with Normal Mode Analysis (NMA) have been established. Internal Coordinates (IC) can significantly improve the versatility and efficiency of current approaches that are usually based on Cartesian coordinates. The iMODS server (https://imods.iqfr.csic.es/) was employed to explain protein collective motion (López-Blanco et al. 2011). To investigate the motions of proteins using NMA in internal coordinates, one can simply submit the PDB-ID or the atomic coordinates in PDB format.

In Silico Cloning
Codon optimization and back-translation of the final construct were done by the OPTIMIZER server (http://genomes.urv.es/OPTIMIZER/) for the high-level expression of the multi-epitopic protein structure in E. coli. Afterward, the different factors of DNA sequence such as Codon with a Frequency Distribution (CFD), overall GC content, and no negative CIS elements and repeat sequences were evaluated by GenScript Rare Codon Analysis Tool (https://www.genscript.com/tools/rare-codon-analysis). In the next step, the pET-23a+ expression vector was used to simulate in silico cloning of the DNA sequence of the construct.

Results
Sequence Retrieval
The UniprotKB database was used to obtain the amino acid sequences of eight pathogenic Candida albicans proteins. All sequences of MET6, two sequences of BGL2, three sequences of EAP, 17 sequences of SAP1, 24 sequences of HWP1, 44 sequences of ALS, four sequences of ECE1, and 21 sequences of HYR1 were retrieved from UniProtKB (Table 1).

Immunoinformatics Analyses
Antigenicity Prediction
The EMBOSS server was used to detect the antigenic sequence with the highest score for antigenicity. Approximately 13 antigenic regions of BGL2 protein, 27 antigenic regions of EAP1 protein, 32 antigenic regions of MET6 protein, 18 antigenic regions of SAP1 protein, 25 antigenic regions of HWP1 protein, 23 antigenic regions of ALS protein, 13 antigenic regions of ECE1 protein, and 32 antigenic regions of HYR1 protein were identified. All the results obtained from this server were found in Supplementary Tables (S1a–h).

Determining Linear B Cell Epitopes
The ABCpred, BCPRED, and BepiPred servers were used to detect linear B-cell epitopes from all eight proteins based on their amino acid sequences. The number of linear epitopes and their scores are listed in Table 2.

The shared sequences predicted by the two mentioned servers were selected by Microsoft Excel software, and the best immunogenic epitopes were chosen. The finally selected regions of BGL2, EAP1, MET6, SAP1, HWP1, ALS, ECE1, and HYR1 proteins are presented in Table 3.

Antigenicity and Allergenicity of Designed Construct’s Epitopes
The antigenicity of each epitope was estimated separately using the VaxiJen 2.0 server and a fungal model. Epitopes were identified as antigens or potential antigens based on the findings. The scores obtained from the VaxiJen 2.0 server are as follows: BGL2 is 1.35, EAP1 is 1.26, MET6 is 0.5, SAP1 is 0.76, HWP1 is 0.96, ALS is 0.93, ECE1 is 0.56, and HYR1 is 1.67. Allergenicity prediction by the AllerTOP server showed that all epitopes in the designed structure were non-allergenic.

Assembling Epitopes
The select B cell epitopes of BGL2, EAP1, MET6, SAP1, HWP1, ALS, ECE1, and HYR1 proteins, displayed in Table 3, were connected using an EAAAK linker. Linkers are short peptides to connect different protein domains. Building a multi-epitope diagnostic kit without a proper linker can exacerbate protein folding issues. The EAAAK
A rigid linker was used to assemble the designed diagnostic kit. The EAAAK linker not only maintains proper spacing between epitopes and fragments of the designed vaccine but also eliminates unwanted interactions. Also, a 6xHis
A tag was added to the C-terminus to facilitate the purification process of multi-epitope protein structures (Fig. 1).

**Assessment of Physicochemical Properties**

Different physicochemical properties of the designed multi-epitope construct were determined via the ProtParam server. The construct’s number of amino acids and molecular weight (MW) were 255 and 27.3 kDa, respectively. The theoretical PI of the construct to indicate the basic condition of the protein was 6.80. The estimated half-life was 2.8 h in mammalian reticulocytes, 3 min in yeast, and 2 min in *E. coli*. The aliphatic index was 71.37, and the grand average of hydropathicity (GRAVY) was −0.289, which showed that the protein was hydrophobic and could better interact with neighboring water molecules. The instability index was computed as 32.48, which classified the protein as stable.

**Tertiary Structure Modeling**

The 3D model of the structure was predicted by the GalaxyTBM server, resulting in five initial models (Fig. 2a–e).

**Tertiary Structure Validation**

The ERRAT (Fig. 3a–e), ProSA-web (Fig. 4a–e), and Ramachandran plot servers (Fig. 5a–e) were used to evaluate the quality of the predicted models to select the best model. The ProSA-web server shows the overall quality score of protein (z-score) obtained by comparing the 3D model of the target protein with similar proteins of the same size deposited in PDB. The validation results of the five modeled structures by the ERRAT, ProSA-Web and Ramachandran servers are presented in Table 4. After comparing all the obtained results from the servers mentioned above, model 4 was selected as the best structure. In this model, the ERRAT quality and ProSA z-score were 78.59 and −3.33, respectively (Table 4). Moreover, the Ramachandran plot showed that 86.6%, 11.5%, and 1.5% of the amino acid residues were located in the favored region, allowed region, and generously allowed regions, respectively.

**Molecular Dynamics (MD) Simulations**

Understanding the macromolecular function of the protein complex was performed using the internal coordinates of the iMODS server (Fig. 6a). The complex’s eigenvalue was

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**Table 2** Linear B cells epitopes analysis of BGL2, EAP1, MET6, SAP1, HWP1, ALS, ECE1 and HYR1 proteins by ABCpred, BCPRED and BepiPred servers

| Protein | Server | Number of linear B cell epitopes | Maximum score |
|---------|--------|----------------------------------|---------------|
| BGL2    | ABCpred| 29                               | 0.91          |
| BGL2    | BCPRED| 6                                | 0.999         |
| BGL2    | BepiPred| 11                             | 0.66          |
| EAP1    | ABCpred| 84                               | 0.95          |
| EAP1    | BCPRED| 26                               | 1             |
| EAP1    | BepiPred| 1                              | 0.66          |
| MET6    | ABCpred| 78                               | 0.97          |
| MET6    | BCPRED| 10                               | 0.999         |
| MET6    | BepiPred| 28                             | 0.67          |
| SAP1    | ABCpred| 33                               | 0.94          |
| SAP1    | BCPRED| 7                                | 0.997         |
| SAP1    | BepiPred| 11                             | 0.7           |
| HWP1    | ABCpred| 65                               | 0.97          |
| HWP1    | BCPRED| 26                               | 1             |
| HWP1    | BepiPred| 1                              | 0.71          |
| ALS     | ABCpred| 62                               | 0.95          |
| ALS     | BCPRED| 19                               | 1             |
| ALS     | BepiPred| 15                             | 0.65          |
| ECE1    | ABCpred| 24                               | 0.90          |
| ECE1    | BCPRED| 3                                | 1             |
| ECE1    | BepiPred| 7                              | 0.65          |
| HYR1    | ABCpred| 92                               | 0.92          |
| HYR1    | BCPRED| 27                               | 1             |
| HYR1    | BepiPred| 8                              | 0.7           |

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**Table 3** The final selected B cell epitopes regions of BGL2, EAP1, MET6, SAP1, HWP1, ALS, ECE1 and HYR1 proteins

| Protein | Start–end position | Sequence | Length of sequence |
|---------|--------------------|----------|--------------------|
| BGL2    | 276–306            | WKPDTSGTSSVEKHWGVWQSDKTLKYSIDCK | 31 |
| EAP1    | 91–114             | STSYYTDTAYTTTVTVCDDGGS          | 24 |
| MET6    | 77–110             | NAPIERYTFLAPIPYLVFLAMGRGLQKKATETQA | 34 |
| SAP1    | 22–53              | KRSPGFVTLDFVKTVPVQDQEGKVRQAI   | 32 |
| HWP1    | 376–408            | TTVITVTCSSESSCETSEVTVGTVVTSEETVY | 33 |
| ALS     | 313–354            | TGYRNSDAGSNGIVIVATTTRTVTSTTAATTLTVPFDPRNKTK | 42 |
| ECE1    | 79–103             | QIIMSVKAFKGNKREDIDSVVAGII      | 25 |
| HYR1    | 831–863            | ANPVTSTESDTTISVTSTYMTGFDKGPKP  | 33 |

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2.744204e40 (Fig. 6b). Arrows indicated the direction of each residue in the 3D model, and the length of the line represented the extent of mobility. The B-factor values derived from normal mode analysis were comparable to RMS (Fig. 6c). Hinges in the chain indicated the complex’s probable deformability as measured by the contortion of each individual residue (Fig. 6d). Each normal mode’s variance was inversely related to its eigenvalue. The covariance matrix explained the link between pairs of residues by representing correlated, uncorrelated, or anti-correlated motions with red, white, and blue colors, respectively (Fig. 6e). As a result, an elastic network model (Fig. 6f) was generated, which identified the pairs of atoms connected by springs. Each dot in the diagram represented one spring between the corresponding pair of atoms and was colored according to its stiffness.

In Silico Cloning of the Final Construct

The OPTIMIZER online server was used for back-translation and codon optimization of the multi-epitope construct for this study. The gene sequence’s CAI, CFD, and GC content were 0.88, 0%, and 53.3%, respectively (Fig. 7a–c). A CAI of > 0.8 is considered good for expression in a selected host. The optimal percentage range of GC content is between 30 and 70%. Codons with a frequency distribution (CFD) < 30% are likely to impede transcripational and translational efficiency. All the above-mentioned factors of our sequence were in the ideal range. This indicated that the final multi-epitope construct with optimized sequence would be efficiently expressed in *E. coli*. Next, an optimized nucleotide sequence was cloned into the pET-23a+ expression vector. For this purpose, *NdeI* and *XhoI* restriction sites were added to the N and C-terminus of the sequence, respectively (Fig. 8).

**Discussion**

The occurrence of invasive candidiasis was increased by 50% globally over the past decade, varying from 2.4 per 100,000 to 15 per 100,000, according to the clinical and national conditions. Despite considerable advances in antifungal remedies, candidiasis spreads with high mortality (40–60%). The timely use of appropriate antifungal agents is necessary to increase survival. However, yeast blood cultures are much less available and require long-term (24 h or longer) incubation (Poissy et al. 2020). ELISA and concomitant immunological methods are used in most detection approaches due to desirable features such as high sensitivity, convenience, and relatively low cost (Khatami et al. 2020b). Recombinant antigenic proteins are the best options for detecting antibodies against invasive *Candida* antigenic proteins in the blood. Also, as an efficient serological diagnostic method, ELISA is one of the most effective procedures for detecting antibodies in invasive candidiasis. Using the complete sequence of an antigen to detect antibodies can cause nonspecific binding to the target antibody due to the similarity and overlap with similar sequences of other living antigenic proteins. In general, this method reduces the degree of specificity (Dey et al. 2021; Laín et al. 2007b). Each antigen’s design, production, and purification in diagnosing infectious diseases are time-consuming, requiring high costs and controlling different processes. Thus, a combination of different antigenic proteins as a chimeric protein can be a valuable way to reduce the time and cost of the process. In recent years, bioinformatics knowledge has increased in various scientific fields, e.g., the production of recombinant products and proteins (Khatami et al. 2020a; Mahapatra et al. 2020; Sabetian et al. 2019; Shahbazi et al. 2016; Tehrani et al. 2020; Zarei et al. 2019). The present study used various bioinformatics strategies to design the *Candida albicans* diagnostic kit based on linear B cell epitopes of BGL2, EAP1, MET6, SAP1, HWP1, ALS, ECE1 and HYR1 antigens. Zheng et al. conducted serological assays for rapid diagnosis of invasive candidiasis. In this regard, recombinant BGL2 protein of *Candida albicans* cell wall was produced, and an indirect ELISA method was developed to detect IgG antibodies. The sensitivity, specificity, positivity, and negative predictive values were 80.8%, 90.0%, 89.4%, and 81.8%, respectively for anti-BGL2 detection. The results showed that recombinant BGL2 antigen from *Candida albicans* reacted to the sera of proven IC patients infected with *Candida albicans*. 
The EAP1 protein, expressed by the EAP1 gene, binds to glycosylphosphatidylinositol in the *Candida albicans* cell wall. The EAP1 protein is involved in surface adhesion. Different studies have shown that mutations in this protein reduce *Candida albicans* adhesion to epithelial cells, indicating the importance of this protein in the binding of yeast cells. The EAP1 protein is also effective in forming and developing *Candida albicans* biofilms (Li et al. 2007). Recently, it was discovered that EAP1 plays a role in macrophage regulation and lymphocyte activation. Furthermore, EAP1 suppressed CD69 expression, implying that EAP1 is involved in CD4T-cell activation (Ramalho-Oliveira et al. 2019). Secreted Aspartic Proteinases (SAPs) are a group of hydrolytic enzymes secreted by *Candida albicans* to penetrate host cell membranes. There is an unequal tendency for SAPs proteins to penetrate various cell types. Therefore, in different stages of yeast infection, they could be expressed and secreted into both intracellular and extracellular environments. In this regard, the SAP1 protein is mainly observed in vaginal infections caused by *Candida albicans* (Oliver et al. 2019; Singh et al. 2019). According to studies, the increase in Th1/Th2/Th17 cytokine levels

![Fig. 2](image-url) In order to predict the third structure of the designed multi-epitope structure, GalaxyTBM server was used. As a result, five different models of designed multi-epitope protein by GalaxyTBM server, a model 1, b model 2, c model 3, d model 4, e model 5.
observed in SAP2-vaccinated mice indicates that SAP2 has immunomodulatory properties. SAP2 immunization also significantly increases total CD45+ leukocytes in the spleen, preventing a significant decrease in their number following fungal infection. Furthermore, SAP2 immunization increases plasma cell counts and the proportion of fungal-binding B cells in the spleens of immunized mice (Shukla and Rohatgi 2020). Leighann et al. showed that using a defined panel of genes during the initial growth of biofilm may be very significant for diagnosing *Candida albicans*. Panels of genes such as SAP, HWP1, and EAP1 are examined sequentially to identify different biomarkers for diagnosing invasive candidiasis in immunocompromised patients (Sherry et al. 2014). Methionine synthase (MET6) is a protein with an essential role in methionine biosynthesis during *Candida albicans* morphogenesis (Sáez-Rosón et al. 2014). Patients with disseminated candidiasis have a better prognosis when MET6-specific IgG serum antibodies are produced (Adams et al. 2021). Hyphal Wall Protein 1 (HWP1) facilitates *Candida albicans* attachment to host epithelial cells. In addition, HWP1 encodes a principal protein of *Candida albicans* that is involved in cell wall proliferation and intracellular signaling (Orsi et al. 2014). Agglutinin-Like Sequence (ALS) proteins are important proteins in the *Candida albicans* cell wall, which encode large cell surface glycoproteins and play a key role in adhesion to different surfaces (Lombardi et al. 2019). Generally, a valid diagnostic method is used to detect antibodies against ALS, HWP1, and MET6 *Candida albicans* proteins. The results obtained by ELISA showed that the sensitivity to detect the desired antibodies is more than 66%. Also, in immunocompromised patients, anti-MET6 antibodies were named the best biomarkers (Díez et al. 2021). ECEIp is a large protein with 271 aa that is processed in the Endoplasmic Reticulum (ER) and the Golgi complex. Candidalysin toxin is also derived from the Ece1p protein (Richardson et al. 2018). Antifungal drugs have been shown to eliminate various components of fungi required for their virulence. They also influence the expression of genes involved in hyphal growth, such as Hwp1 and Ecel, in the epithelium. As a result, Hwp1 reduced IL-8 production, inhibited aspartyl proteinase expression, and reduced the inflammatory response (Rodríguez-Cerdeira et al. 2020). A Hyphally Regulated Gene (HYR1) is expressed specifically during hyphal development. Hyr1p contributes to *Candida albicans* virulence by resisting phagocyte killing (a major host defense mechanism against candidiasis) via an unknown mechanism. (Uppuluri et al. 2018). Numerous studies have investigated the recombinant *Candida albicans* antigens for their potential use in diagnosing invasive candidiasis. In this context, SAP, HWP1, HYR1, ECE1, and ALS proteins as *Candida albicans* recombinant antigens have been tested for serodiagnosis of invasive candidiasis (Lain et al. 2007a, 2008).

In the current study, the proteins sequences were obtained from the UniProt databases at www.uniprot.org. Prediction of B cell epitopes plays a vital role in developing diagnostic kits based on multi-epitope proteins. Due to the subject’s significance, three different servers were used to identify the
Fig. 4 ProSA-Web z-score, indicate the validation of protein 3D models. The ProSA Z-score, based on X-ray crystallography and NMR spectroscopy, of Models a 1, b 2, c 3, d 4, e 5, are −3.33, −3.23, −2.9, −2.96, and −3.4, respectively
linear 16-mer B cell epitopes regions. The final epitopes were selected and then connected by the EAAAK linker. Selecting a suitable linker is crucial for the appropriate binding of protein fragments in a multi-epitope structure. The linkers are responsible for connecting protein segments and have an essential role in maintaining the biological activity and increasing the

Fig. 5 The PROCHECK’s Ramachandran plot validate the 3D models of selected multi-epitopic construct. Favoured regions in models was shown; a model 1 (87.0%), b model 2 (87.4%), c model 3 (86.6%), d models 4 (85.9%), e model 5 (86.3%)
expression of the structure (Chen et al. 2013; Li et al. 2016; Mahapatra et al. 2021). The EAAAK (Glu–Ala–Ala–Ala–Lys) sequence is a tough and helix-forming linker. Lee et al. conducted a study to enhance the antibacterial activity of an attacin-coleoptericin hybrid protein fused with a linker. They used helical (Glu–Ala–Ala–Ala–Lys) and non-helical linkers (Gly–Gly–Gly–Gly–Ser) to compare them in binding the two antimicrobial proteins. The results showed that the fused proteins with the EAAAK linker had more antibacterial properties (Lee et al. 2013). Our study selected and ligated antigenic regions of different proteins using linkers. The three-dimensional structures of the proteins were modeled and evaluated by several servers. In this way, the Galaxy-TBM server developed five models for the desired structure. Then, the obtained models were evaluated by the ERRAT, ProSA-Web, and Ramachandran plot servers, and finally, model 4 was selected.

## Conclusion

In this present study, to provide a diagnostic kit for *Candida albicans*, we used immunodominant linear B cell epitopes from eight pathogenic *Candida albicans* antigens (BGL2, EAP1, MET6, SAP1, HWP1, ALS, ECE1, and HYR1) to model and design a multi-epitope protein using suitable linkers. The structural and physicochemical evaluations of the designed protein construct showed that our recombinant protein was thermostable, hydrophilic, and basic. Finally, it was found that the designed chimeric protein could be significantly expressed in *E. coli*. The multi-epitope-designed protein can be used to construct a high-performance serological test for the diagnosis of candidiasis infections.
Fig. 6 Molecular dynamics simulation of multi-epitope diagnostic kit; stability of the protein complex was investigated through mobility, b eigenvalue, c B-factor, d deformability, e covariance and f elastic network analysis.

Fig. 7 Designed structural analysis to express high levels of multi-epitope protein in E. coli host. a Codon frequency distribution (CFD) value of gene sequence is 91–100, 81–90, 71–80, 61–70, 51–60 in the gene is 61%, 7%, 10%, 13%, and 5%, respectively. b The average GC content of our sequence is 53.3%. c CAI of gene sequence is 0.88; a CAI of 40.8 is considered as good for expression in selected host.
Fig. 8 Schematic representation of cloning process. The gene sequence of multi-epitope protein with the name of Vahedi opt, and the length of 882 nucleotides shown in the designed structure. The Vahedi opt gene sequence in the Vahedi cloned structure is distinct between the NdeI and XhoI cleavage sites.

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Data Availability  All data generated or analysed during this study are included in this published article, and its supplementary information files.

Declarations

Conflict of interest  All the authors declare that they have none conflict of interest.

Research Involving Human and/or Animal Participants  This article does not contain any studies with human participants or animals performed by any of the authors.
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