


MOLECULAR DOCKING STUDY OF SOME HETEROCYCLICS ON STAPHYLOCOCCUS
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ABSTRACT: In this work, we collected the three dimensional structure of Peptidoglycon which plays an important role in staphylococcus pathway. The protein structures were collected from PDB data bank. From the 3D structures of the proteins, the targeted derivatives were designed. Docking studies was performed with designed ligands from the drug. The drug derivatives docked to the protein by hydrogen bonding interactions and these interactions play an important role in the binding studies. Docking results showed the best compounds among the derivatives.

Key words: Antibacterial activity, docking studies, Peptidoglycon

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INTRODUCTION

Bacteria, including Staphylococcus aureus, can bind opsonically active C3b molecules through activation of the classical as well as the alternative complement (C) pathways. In either case, the fixation of C3 to the surface of the bacteria seems to be critical for rapid ingestion by phagocytes (Gigli & Nelson, 1968; Stossel et al 1975). In normal human serum S. aureus opsonization is rapidly mediated via the IgG-dependent classical C pathway. In the absence of antibody, however, bacterial opsonization proceeds at a significantly slower rate via the alternative C pathway (Verbrugh, van Dijk et al 1979). Recently attention was focused on peptidoglycan (PG) as being the S. aureus cell-wall component of key importance in promoting bacterial opsonization (Kronvall et al 1970). Purified PG has been found to remove opsonic factors from serum (Humphreys et al 1974; PRYJMA et al 1976), to become rapidly ingested by human polymorphonuclear leucocytes when opsonized (Peterson et al 1978) and to consume C3-9 via the classical as well as the alternative C pathways and in the absence of immunoglobulins (Peterson et al., 1978; Wilkinson et al 1978). Teichoic acid (TA) and protein A (PA), the other two major polymers of the cell wall of S. aureus, were not included in these studies and their importance in bacterial opsonization was thus assessed only indirectly (Archibald et al 1972). Furthermore, the consumption of C by staphylococci has not been correlated with the process of bacterial opsonization.

METHODOLOGY

The structures of the compounds (Figure 1) were constructed and optimized using chemsketch software.

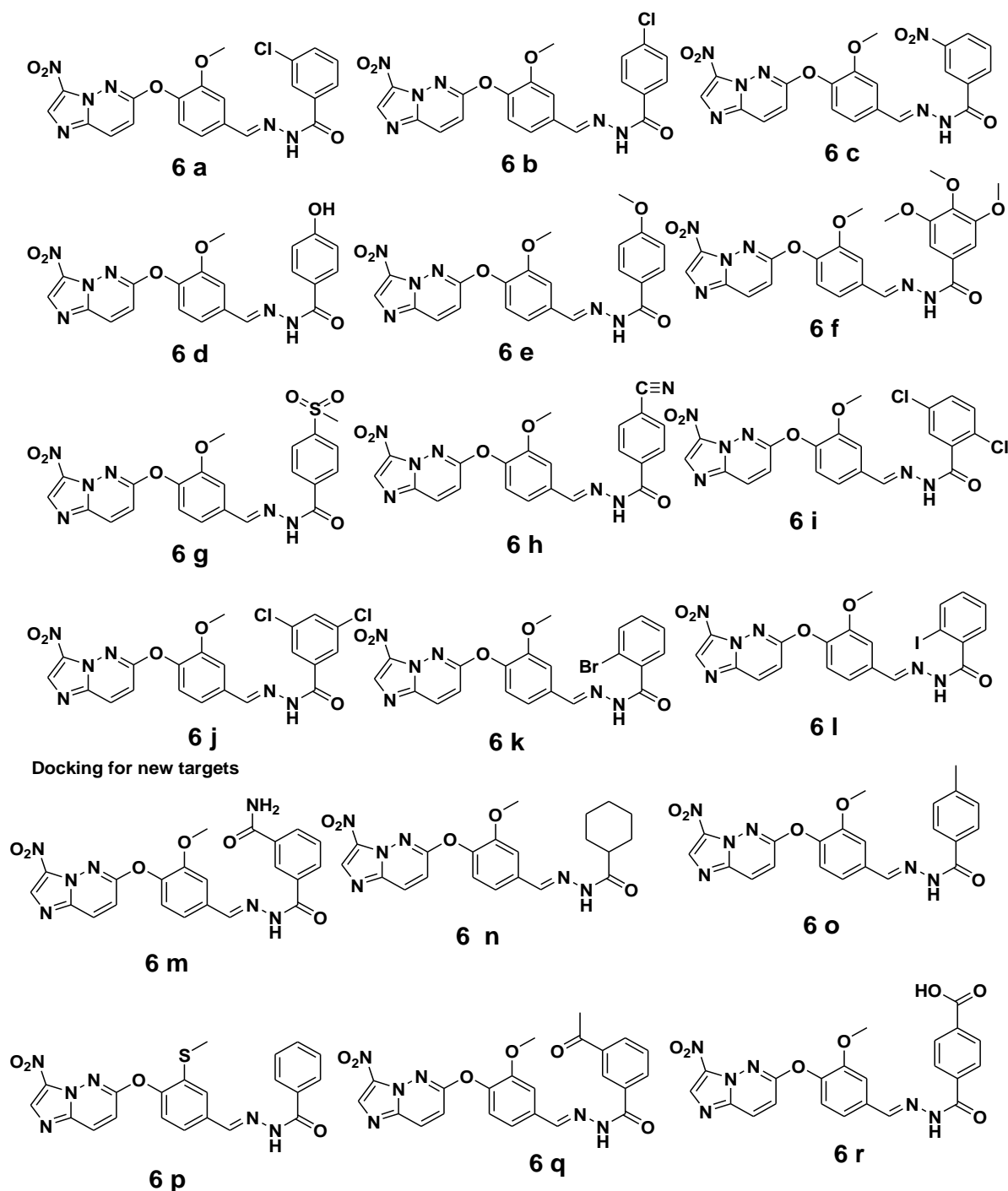


Fig 1: compounds used for docking studies

To prepare the Peptidoglycan from *Staphylococcus*, the crystal structure was taken from the Protein Data Bank (PDB_ID: 3NB6) (Figure 2).



Fig 2: Peptido glycon

Hetero atoms were removed from the binding site and the chain A was selected for docking studies. Hydrogen atoms were added to the enzyme. The molecular docking method was performed using the Gold version 3.0.1 program to study the binding orientation of compounds into the Sortase structure. The docking experiments were performed using the binding site of Sortase. The binding site identification was carried out using Cast P server. A new program, CAST, for automatically locating and measuring protein pockets and cavities, is based on precise computational geometry methods, including alpha shape and discrete flow theory. CAST identifies and measures pockets and pocket mouth openings, as well as cavities. The program specifies the atoms lining pockets, pocket openings, and buried cavities; the volume and area of pockets and cavities; and the area and circumference of mouth openings.

Docking method

Docking was carried out using GOLD (Genetic Optimization of Ligand Docking) software which is based on genetic algorithm (GA). This method allows as partial flexibility of protein and full flexibility of ligand. The compounds are docked to the active site of the protein. The interaction of these compounds with the active site residues are thoroughly studied using molecular mechanics calculations. The parameters used for GA were population size (100), selection pressure (1.1), number of operations (10,000), number of island (1) and niche size (2). Operator parameters for crossover, mutation and migration were set to 100, 100 and 10 respectively. Default cutoff values of 3.0 Å° (dH-X) for hydrogen bonds and 6.0 Å° for vanderwaals were employed. During docking, the default algorithm speed was selected and the ligand binding site in the alpha glucosidase was defined within a 10 Å° radius with the centroid as CE atom of ASP175. The number of poses for each inhibitor was set 100, and early termination was allowed if the top three bound conformations of a ligand were within 1.5Å° RMSD. After docking, the individual binding poses of each ligand were observed and their interactions with the protein were studied. The best and most energetically favorable conformation of each ligand was selected.

Gold Score fitness function:

Gold Score performs a force field based scoring function and is made up of four components: 1. Protein-ligand hydrogen bond energy (external H-bond); 2. Protein-ligand vander Waals energy (external vdw); 3. Ligand internal vander Waals energy (internal vdw); 4. Ligand intramolecular hydrogen bond energy (internal- H- bond). The external vdw score is multiplied by a factor of 1.375 when the total fitness score is computed. This is an empirical correction to encourage protein-ligand hydrophobic contact. The fitness function has been optimized for the prediction of ligand binding positions.

GoldScore = S (hb_ext) + S (vdw_ext) + S (hb_int) + S (vdw_int)

Where S (hb_ext) is the protein-ligand hydrogen bond score, S (vdw_ext) is the protein-ligand van der Waals score, S (hb_int) is the score from intramolecular hydrogen bond in the ligand and S (vdw_int) is the score from intramolecular strain in the ligand.

RESULTS AND DISCUSSION

After collecting the crystal structures, the possible binding sites of 3NB6 was searched with CASTP server as shown in Figure 3. The residues included in active site were ILE 110, ASP 111, SER 112, PRO 113, THR 115, TYR 116, GLY 121, TYR 122, LYS 123, ALA 137, LEU 138, PRO 139, PRO 140, VAL 141, ALA 142, CYS 145, GLY 150, LEU 352, GLN 355, GLN 356, PHE 357, GLN 358, and LYS 518.

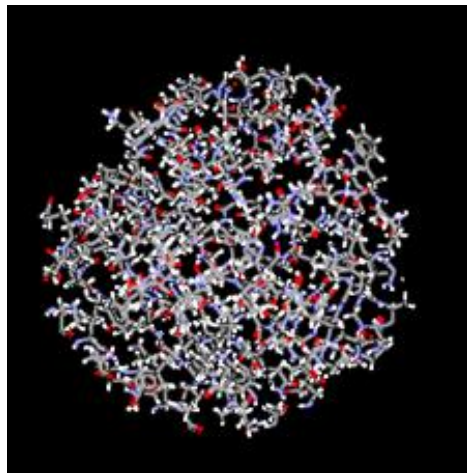
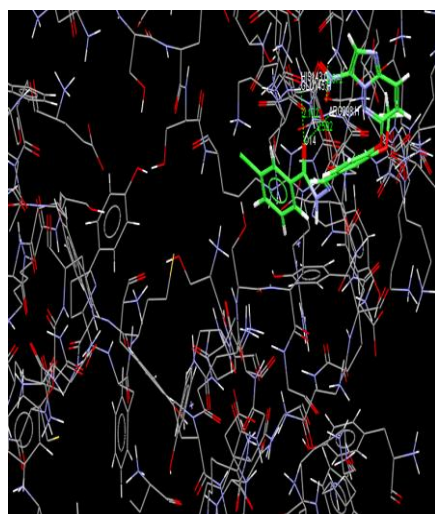


Fig 3: Active site

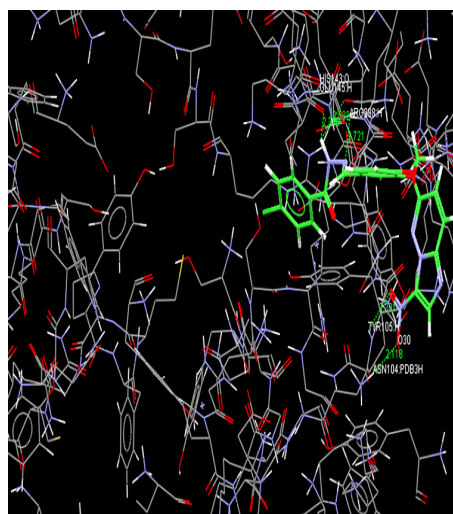
From the binding site analysis of 3NB6 we identified that, the binding pockets are identical in all chains and the largest binding pocket was taken for further docking studies. The crystal structure of 3NB6 was similar hence we have taken as representative structure for docking studies. The docking of drugs into the active site of 3NB6 was performed using the GOLD software and the docking evaluations were made on the basis of GoldScore fitness functions. We preferred Gold fitness score than Chemscore fitness as Gold fitness score is marginally better than Chemscore fitness function.

Molecular docking study

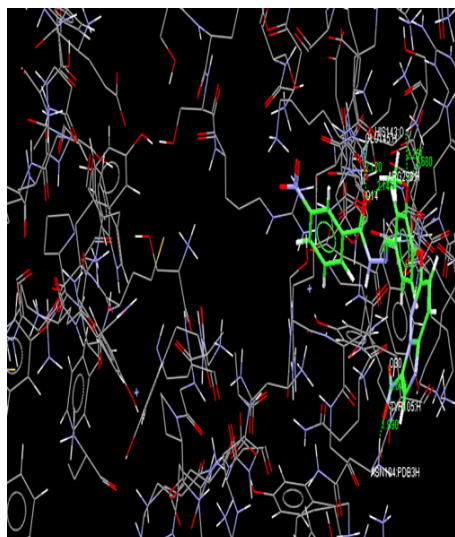
Structure-based drug design begins with the identification of a molecular target such as a protein such as 3NB6 in this study. This structure is then used as a blueprint for the drug design of a lead compound. The compounds are modelled for their fit in the active site of the target, considering both steric aspects (*i.e.*, geometric shape) and functional group interactions, such as hydrogen bonding and hydrophobic interactions. The selected docked conformations of analogues into the 3NB6 binding site are shown in Figure 4.



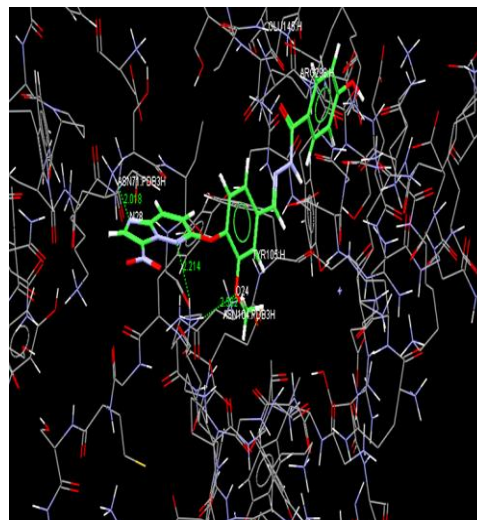
4a) Docking of 6a



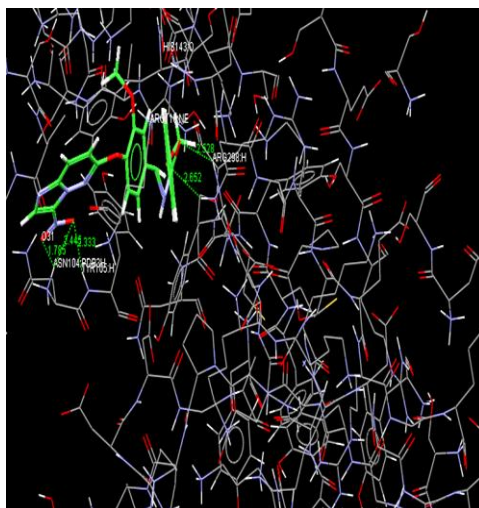
4b) Docking of 6b



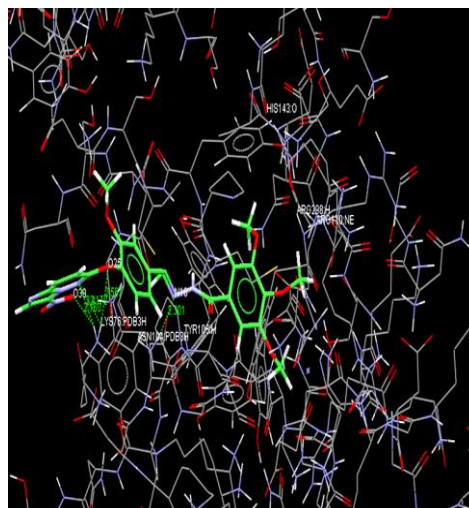
4c) Docking of 6c



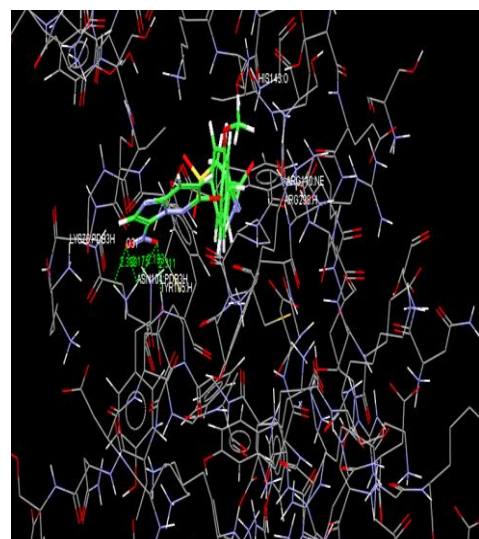
4d) Docking of 6d



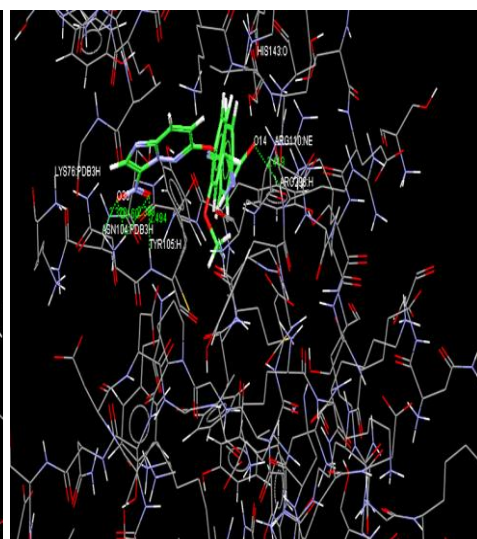
4e) Docking of 6e



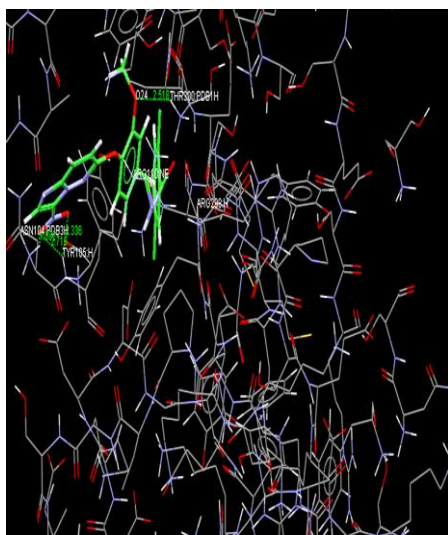
4f) Docking of 6f



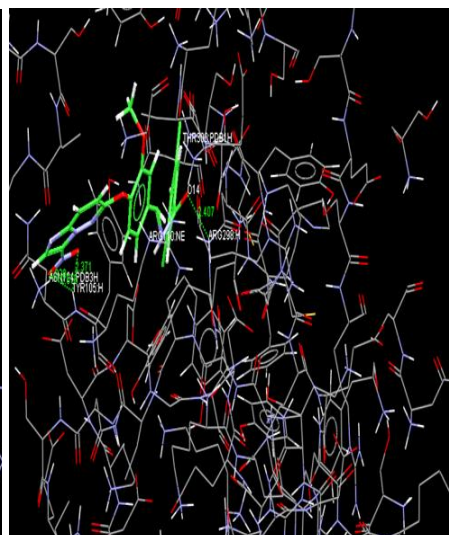
4g) Docking of 6g



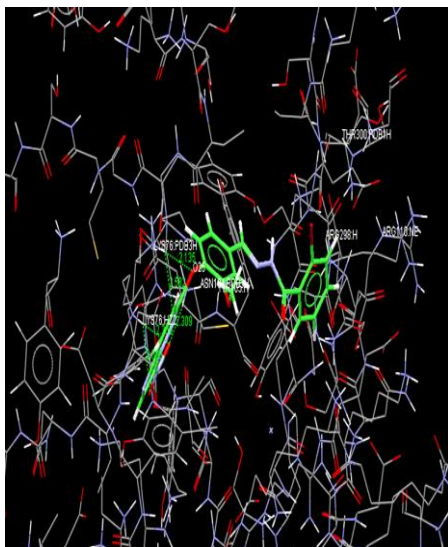
4h) Docking of 6h



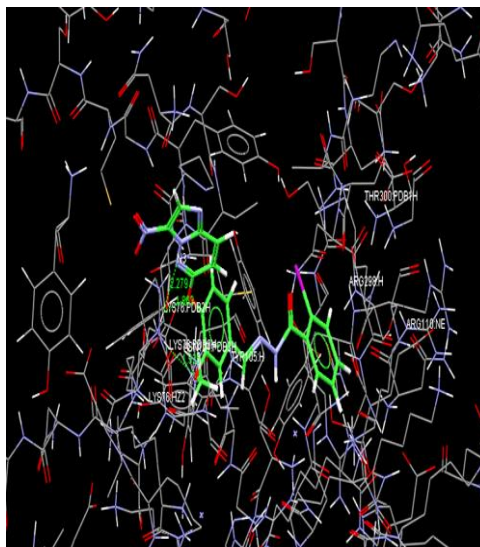
4i) Docking of 6i



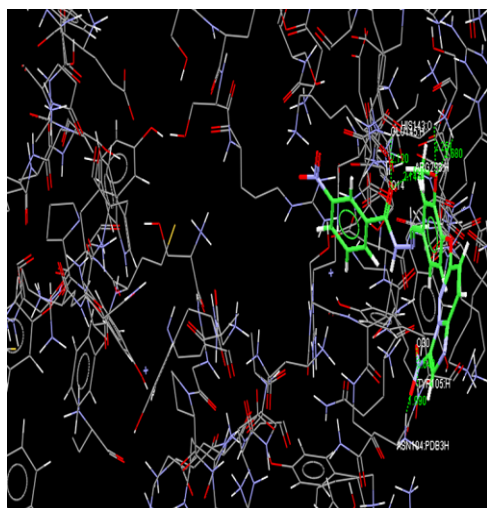
4j) Docking of 6j



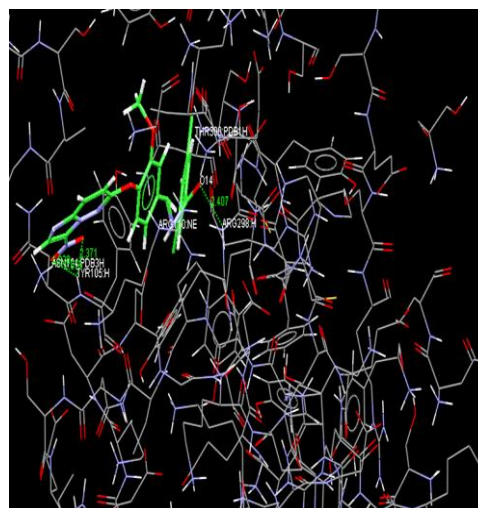
4k) Docking of 6k



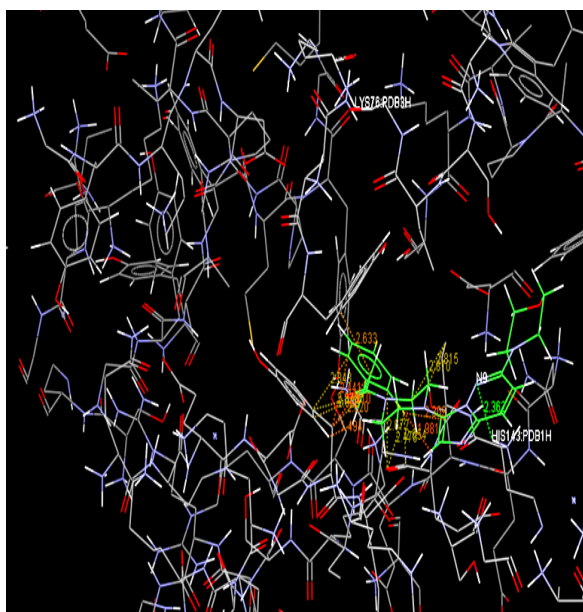
4l) Docking of 6l



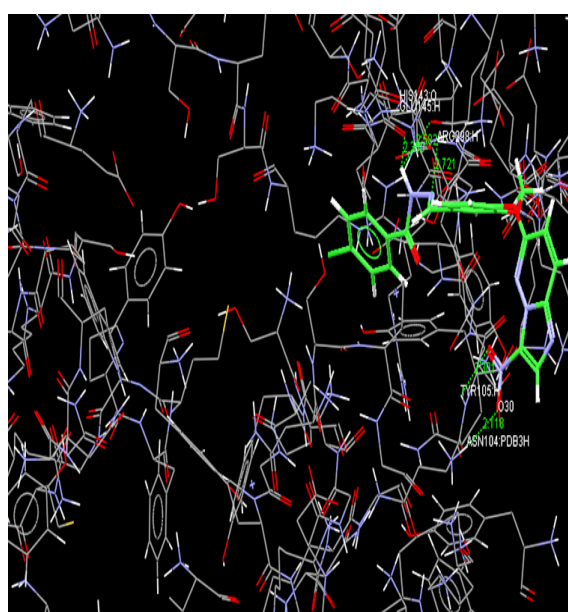
Docking of 6m



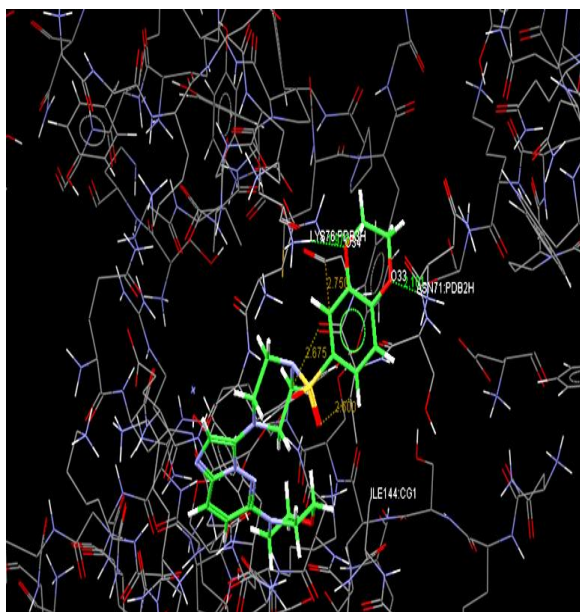
Docking of 6n



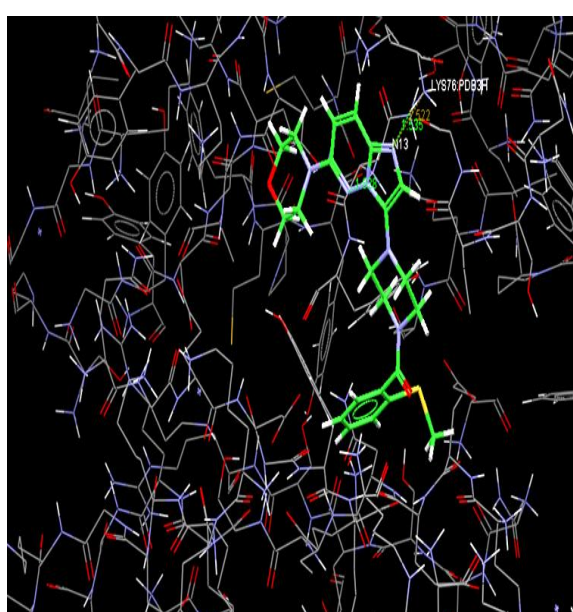
Docking of 6o



Docking of 6p



Docking of 6q



Docking of 6r

Fig 4a-4l: Docking of compounds

The docked conformations revealed that all molecules were located in the hydrophobic binding pocket. In this study, all docked drugs were found to have some interactions between an oxygen atom of the drugs and target proteins. Moreover, these docked conformations also formed an H-bonding interaction within the active site (Table 1).

In the binding pocket, common H-bonding interactions were formed between all docked drugs and GLY 121, TYR 122, LYS 123, ALA 137. In order to explain the binding of these compounds, the H-bonding interactions with the other surrounding residues in the hydrophobic binding pocket were also investigated. In Figure 4, strong H-bonding interactions between the hydroxyl group (H12) of analogue and an oxygen atom of GLU31 and another hydrogen bond between hydroxyl group (H11) and oxygen atom of ASP111. Two H-bonding interactions were also formed between Oxygen atom (O4) of 5-FU, ASP29 and CYS 44 of 3NB6.

Table 1: Fitness score

| Fitness | S(hb_ext) | S(vdw_ext) | S(hb_int) | S(int) | Ligand name |
|---------|-----------|------------|-----------|--------|-------------|
| 36.24 | 0.00 | 33.77 | 0.00 | -10.18 | 6a |
| 38.40 | 6.00 | 30.48 | 0.00 | -9.51 | 6b |
| 42.21 | 3.94 | 40.79 | 0.00 | -17.81 | 6c |
| 43.68 | 9.26 | 30.36 | 0.00 | -7.32 | 6d |
| 41.94 | 6.20 | 39.23 | 0.00 | -18.21 | 6e |
| 35.21 | 8.99 | 33.74 | 0.00 | -20.16 | 6f |
| 38.68 | 8.00 | 39.02 | 0.00 | -22.97 | 6g |
| 41.20 | 5.95 | 39.37 | 0.00 | -18.88 | 6h |
| 44.02 | 4.42 | 40.72 | 0.00 | -16.38 | 6i |
| 44.46 | 4.45 | 40.68 | 0.00 | -15.92 | 6j |
| 41.63 | 18.00 | 23.48 | 0.00 | -8.66 | 6k |
| 41.51 | 9.17 | 28.38 | 0.00 | -6.68 | 6l |
| 21.68 | 5.03 | 32.60 | 0.00 | -28.19 | 6m |
| 15.69 | 4.74 | 24.46 | 0.00 | -22.68 | 6n |
| 18.71 | 0.63 | 33.58 | 0.00 | -28.10 | 6o |
| 31.21 | 8.99 | 33.74 | 0.00 | -20.16 | 6p |
| 33.57 | 8.00 | 39.02 | 0.00 | -22.97 | 6q |
| 40.12 | 5.95 | 39.37 | 0.00 | -18.88 | 6r |

CONCLUSION

The docking results agreed well with the observed *in vitro* data, in which the anti-microbial activity of the analogues was higher than other drugs and formed five hydrogen bonds. The docking study revealed the binding orientation of compounds in the 3NB6 binding pocket surrounding the active site, which resulted in inhibition of enzyme activity. From these results we can conclude that compound 6j is one of the good inhibitory compounds of 3NB6. The application of computational sciences to pharmaceutical research is a discipline, which is phenomenal.

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