Molecular Docking of *Glycyrrhiza glabra* against the Conserved Target M1, NA and NS1 Proteins of Influenza A Viral Strains Identified through Pangenome Analysis

Aishwarya. S *, Evangeline Shantha, Nantha Devi. E, Sagaya Jansi R

Department of Bioinformatics, Stella Maris College, Chennai, Tamilnadu, India.

Corresponding author – Aishwarya S*

s.aishwaryabiotech@gmail.com

Abstract

Influenza viruses that infect humans are known to swiftly evolve over time. Influenza A virus has a negative single-stranded RNA genome in eight segments. Pangenome analysis of twelve strains of Influenza A viruses H1N1, H1N2, H2N2, H3N2, H5N1, H5N6, H7N2, H7N3, H7N7, H7N9, H9N2, and H10N8 gave insight on the core genes that are conserved and accessory genes that are specific for the strains. The proteins Neuraminidase, Matrix M1 and Nonstructural protein 1 were encoded by the core genes of segments 6, 7, and 8 respectively which proves that they are conserved in almost all the strains of influenza. The 3Dimensional structures of the core genes were interpreted by homology modeling and compared with corresponding Protein Data Bank structures (4MWQ, IEA3, 2GX9). Among several anti-viral phytocompounds that were virtually screened against the modeled and PDB target proteins, three molecules of Indian plant *Glycyrrhiza glabra* had high scores and interactions. Compounds 2,4,4' Trihydrochalcone, Davidigenin and Licoflavone B docked well with the Neuraminidase, Matrix protein M1 and Nonstructural Protein NS1 respectively with good scores, minimized energy and interacted with the active sites. The compounds obeyed Lipinski's Rule of five and exhibited drugability as well. Thus the present study focused on the drugable lead compounds from *glabra* that has inhibitory activities against the viral attachment, replication and matrix structure.

Keywords: Pan Genome Analysis, Influenza A virus, Homology Modeling, Molecular Docking, Core Genome Fragments

Introduction

The most common cause of respiratory infections in humans is due to seasonal or epidemic outbreak of influenza viruses and recently there are many different strains identified (Aishwarya *et. al.*, 2019). Influenza viruses belonging to the family *Orthomyxoviridae* are enveloped, with negative- single-stranded RNA genomes found in seven to eight gene segments (Miller, MS *et. al* 2013). Among the types A, B and C, Influenza A is more predominant in affecting humans and is mutated to form multiple strains. Influenza A viruses are subdivided by antigenic characterization of the Hemagglutinin (HA) and Neuraminidase (NA), a surface glycoprotein and occur in eighteen HA (H1–H16) and eleven NA (N1–9) subtypes (Hay AJ *et. al*, 2001). These subtypes are further divided into two or three groups: group 1 HA (H1, H2, H5, H6, H8, H9, H11, H12, H13, H16, H17, and H18) and group 2 HA (H3, H4, H7, H10, H14, and H15), or group 1 NA (N1, N4, N5, and N8), group 2 NA (N2, N3, N6, N7, and N9) and group 3 NA (N10 and N11) (Zhang *et. al.*,2014). Currently available treatment options like vaccines and anti-influenza drugs have a short protection plan and they become less effective over time due to the aberrant mutation rates of the viruses (ShinJ *et. al.*, 2019).

Pangenome analysis of viruses enabled the identification of core genes that are conserved in multiple strains and enables identification of the conserved targets from the core genes (Darricarrere *et. al*, 2018). The conserved targets identified were Nonstructural protein 1 that is involved in viral-host interaction (Mok W *et. al*, 201), Matrix protein M1 is responsible for the viral envelope structure (Couch RB *et. al* 2014) and Neuraminidase enhances viral replication (Benton *et. al*, 2012). Though there are neuraminidase inhibitors available, their efficacy is debated since there are varied NA groups in different strains of the virus. Molecular docking is a computational approach that had accelerated drug design research with the efficient prediction of lead compounds against the 3-dimensional protein targets (Shanti N.,2011)

The current study aimed at identifying the core genes of multiple strains of influenza and determining the protein targets which were then studied for rigid molecular docking with the available antiviral plant compounds.

Methodology

Retrieval of complete genomes

The complete genomes of 12 different strains of Influenza A virus infecting humans were retrieved from NCBI Genome sequence portal (www.ncbi.nlm.nih.gov/)

Pan-genome analysis

Pan-genome analysis of the retrieved segmented genes was performed using panseq (<https://lfz.corefacility.ca/panseq/>) to predict the core genes of the eight segments. The open reading frame was identified from the core genes and the translated protein segments were validated using BLAST –Basic Local Alignment Search Tool (www.blast.ncbi.nlm.nih.gov/).

Comparative modeling of the translated protein sequences

Comparative modeling also called homology modeling is the best way to predict the structures of the proteins. The translated core genes of segments 6, 7 and 8 were modeled with suitable templates using the web server Phyre2 (<http://www.sbg.bio.ic.ac.uk/phyre2>).

Library preparation

The entry point for any chemistry program within drug discovery research is generally the identification of specifically acting low-molecular-weight modulators with an adequate activity in a suitable target assay. Around 657 antiviral compounds were identified and retrieved from IMPPAT database. The library thus generated was optimized with an OPLS3 force field and stereoisomers were predicted using the ligprep module of Schrödinger software.

Screening of the druggable molecules

The prepared library compounds were screened based on the Lipinski's rule of 5 that describes the drugability of the molecules based on the following four criteria

1. The molecular weight of the molecule is less than or equal to 500 Daltons
2. LogP (Ratio of Octanol to Water) is less than or equal to 5
3. Number of Hydrogen bond donor is less than or equal to 5
4. Number of Hydrogen bond acceptors is less than or equal to 10

Target structure retrieval from PDB

Identifying and selecting the most appropriate drug target or receptor is the initial step in the drug designing procedure. Mostly proteins act as good targets for the drugs but in some cases, enzymes can also serve as excellent drug targets. Three viral targets namely Neuraminidase

(PDB ID-4MWQ), Nonstructural protein NS-1 (PDB ID- 2GX9) and Matrix protein M1 (PDB ID- 1EA3) were selected and their 3D structure was retrieved from PDB (<https://www.rcsb.org/>). The available protein structures will enable the comparison of the modeled protein and its docking results.

Protein Preparation and Receptor grid generation

The modeled proteins and the retrieved 3D protein structures were energy minimized and optimized using the Protein Preparation wizard of the Schrödinger suite. The binding pocket of the modeled proteins was identified using the Computed Atlas of Surface Topography of Proteins (CASTP server - <http://sts.bioe.uic.edu/castp/>) and the all the six structures were subjected to grid generation through the receptor grid generation package of Schrödinger suite for drug discovery.

Molecular docking and interpretation of results

The prepared and filtered library compounds were subjected to rigid docking against the three sets of target protein structures using the Ligand Docking module of Schrödinger. Standard precision mode screened out better-docked compounds and the next Xtra Precision mode yielded the best-docked compounds. The results were interpreted based on the minimized energy, docking score, and interacting residues

Fig 1 Flowchart represents the exact protocol that was followed in the current study

Results and Discussion

Retrieval of complete genomes

There were 12 different strains of Influenza A viral sequences available in the NCBI that were identified and sequenced at different time scales and periods. The complete genome sequences for the entire study were retrieved from NCBI and their accession numbers are listed in the table 1a and 1b.

Table 1.a List of Genomes of Influenza A strains

Year	2016	2009	2018	2013	2012	1962
Strains	H1N1	H1N2	H3N2	H5N1	H7N3	H2N2
Segment 1	KY509794	JQ340008	MH701027	KF918495	CY125725	AY209940
Segment 2	KY509798	JQ340007	MH701026	KF918496	CY125726	AY210014
Segment 3	KY509802	JQ340006	MH701025	KF918497	CY125727	AY209996
Segment 4	KY509806	JQ340003	MH701020	KF918498	CY125728	AY209959
Segment 5	KY509811	JQ340005	MH701023	KF918499	CY125729	AY210079
Segment 6	KY509814	JQ340004	MH701022	KF918500	CY125730	AY209908
Segment 7	KY509818	JQ340009	MH701021	KF918501	CY125731	AY210037
Segment 8	KY509822	JQ340010	MH701024	KF918502	CY125732	AY210164

Table 1.b List of Genomes of Influenza A strains

Year	2015	5-p 2003	2013	2017	2017	2013
Strains	H5N6	H7N2	H7N7	H7N9	H9N2	H10N8
Segment 1	KT245150	EU783920	KF918334	MF455313	MF440743	KJ406534
Segment 2	KT245149	EU587369	KF918335	MF455314	MF440742	KJ406538
Segment 3	KT245148	EU587370	KF918336	MF455315	MF440741	KJ406542
Segment 4	KT245143	EU587368	KF918337	MF455316	MF440736	KJ406546
Segment 5	KT245146	EU587371	KF918338	MF455317	MF440739	KJ406550
Segment 6	KT245145	EU587372	KF918339	MF455318	MF440738	KJ406554
Segment 7	KT245144	EU587373	KF918340	MF455319	MF440737	KJ406558
Segment 8	KT245147	EU587374	KF918341	MF455320	MF440740	KJ406562

Pangenome analysis

Pan-Genome Analysis was performed for all the 12 different strains of Influenza A virus infecting humans using Panseq. There were multiple core fragments and the most predominant core genes with less mutation were selected. The ORFs of the resulted Core fragments were predicted using ORF Finder. Table 2 represents ORFs of segments 6, 7 and 8 by ORF finder.

333 **Table 2- ORFs of Core Fragments**

Segment Number	Open Reading Frames	Strand	Start	Stop	Length	Translated Protein
Segment 8	ORF 1	+	<1	714	237	Nonstructural Protein NS1
	ORF 2	-	767	117	216	
	ORF 3	+	518	>838	106	
Segment 7	ORF 1	+	<1	759	252	Matrix Protein M1
	ORF 4	+	692	>982	96	
	ORF 11	-	278	>3	91	
	ORF 5	-	958	752	68	
	ORF 9	-	330	157	57	
Segment 6	ORF 2	+	221	>910	229	Neuraminidase NA
	ORF 3	+	54	218	54	
	ORF 6	+	567	692	49	
	ORF 7	-	270	157	37	

Validation of the core genes

The validation of the Predicted core fragments was performed using Pfam sequence search. The segments 6,7 and 8 coded for the proteins neuraminidase, Matrix M1 and Nonstructural protein NS1 respectively and are represented in the figures 2.a, 2.b, and 2.c. Pfam results thus validated the protein-coding core genes of all 12 strains.

Fig 2.a Segment 6

Fig 2.b Segment 7

Fig 2.c. Segment 8

Homology modeling

The ORF sequences were subjected to Blast against a nonredundant Protein database to identify the most similar sequence that was chosen as a template. The sequences that shared 100% identity and maximum query coverage was modeled comparatively and a 3 D structure of each protein was obtained.

Table 3 BLAST results of the ORFs of segment 6,7 and 8

Protein Name	Maximum score	Total score	Query coverage	E value	Identity %	UniProt ID
Neuraminidase	467	467	99%	2.00E-165	98.69%	Q6XUA7
Matrix protein 1	524	524	100%	0	100.00%	Q6XT42
<i>Nonstructural</i> protein 1	478	478	100%	6.00E-173	97.47%	P69277

The sequences were modeled using Phyre2 and models were generated. Predicted models were validated using rampage and the best model was retained for each segment and are represented in the figures 3.a, 3.b, and 3.c. Figures 4a, 4b and 4c represent the Ramachandran plot for the modeled proteins M1, NS1 and Neuraminidase respectively. The favored and allowed regions of residues are represented in table 4. The results thus show that the modeled protein structures are valid to be used for docking.

Figure 3.a -3.c Phyre2 Modeled proteins of the segments 6,7 and 8

Fig 3. a. Matrix M1 Fig. 3.b NonStructural Protein NS1 Fig. 3.c. Neuraminidase

Figure 4.a – 4.c Ramachandran plots for validating the modeled protein

Fig 4a. Matrix M1

Fig. 4.b Non-Structural Protein NS1 Fig. 4.c. Neuraminidase

Table 4 Results of Ramachandran plot for the three modeled proteins

Protein	# residues in the Favoured region	# Residues in the Allowed region	# Residues in the Outlier region
Matrix Protein 1	154 (98.7%)	2 (1.3%)	0 (0.0%)
Nonstructural Protein 1	378 (95.5%)	12 (3.0%)	6 (1.5%)
Neuraminidase	372 (96.4%)	13 (3.4%)	1 (0.3%)

Retrieval of structures from PDB

The known structures of the proteins coding the segment 6, 7 and 8 were retrieved from Protein Data Bank based on the resolution. The Ramachandran plot analysis was performed for the retrieved protein structures whose values are mentioned in table 5.

Table 5 Results of Ramachandran Plot for the three proteins retrieved from PDB

Protein	PDB ID	# residues in the Favoured region	# residues in the Allowed region	#residues in the Outlier region
Matrix Protein 1	1EA3	269 (86.8%)	26 (8.4%)	15 (4.8%)

NonStructural Protein 1	2GX9	237 (96.7%)	8 (3.3%)	0 (0.0%)
Neuraminidase	4MWQ	372 (96.4%)	14 (3.6%)	0 (0.0%)

Fig 5. Ramachandran plot analysis for PDB retrieved proteins

5.a Matrix Protein(1EA3) **5.b** Neuraminidase Protein(4MWQ) **5.c** NS Protein (2GX9)

Active site prediction

The prediction of Active site was performed using the CASTp server for modeled protein and for the PDB proteins using sitemap module of Schrödinger. The number of binding sites and the residues involved are represented in table 6.

Table 6 – Active site residues predicted using CastP web server

Protein	Binding site for modeled protein using Castp		Binding site for PDB proteins using Sitemap	
	# Binding Sites	Residues	# Binding Sites	Residues
Matrix Protein 1	1	81, 84, 85, 93, 96, 126, 130, 133, 135	1	13-16, 47-57, 79, 82-89, 92, 93, 121, 122, 125, 126, 154, 157, 158
Nonstructural Protein 1	1	13, 17, 20, 21, 24, 44, 58, 62, 66	1	89, 91, 92, 94-97, 99, 131-133, 145-148, 150
Neuraminidase	1	118, 119, 134, 151, 152, 156, 178, 179, 221, 222, 224, 225, 227, 240-242, 246, 252, 275-278, 292, 371, 406	1	120, 135, 152, 153, 157, 180, 181, 223, 224, 226-229, 242-244, 246-248, 277-280, 294, 296, 406

						(OH..O) TRP 180	2.09
			XP	-3.649	-34.408	(OH..O) TRP 180	2.81
						(OH..O) GLU 279	2.63

Fig 6 a: 2,4,4' Trihydroxychalcone binding with the modeled neuraminidase protein

Fig 6 b: 2,4,4' Trihydroxychalcone binding with the PDB structure of neuraminidase protein (4MWQ)

Among the 392 virtually screened phytochemicals, 2,4,4' Trihydroxychalcone from *Glycyrrhiza glabra* was found to exhibit good glide score and minimal energy against the neuraminidase protein. The interactions were also good with the active site residues of both the modeled protein and PDB structures. The compound also obeyed Lipinski rule of five with the molecular weight of 256.29g/mol, LogP value of 2.6, 3 Hydrogen bond donors and 4 hydrogen bond acceptors. Percentage of human oral absorption was 92% which makes the compound an effective lead for drug development.

Table 8 Docking results of Matrix protein

Modeled / PDB structure	Compound Name & PubChem ID	Plant Name	SP/ XP	Glide Score	Glide energy	Interactions	Bond length (Å°)
Modeled	Davidigenin 442342	Glycyrrhiza glabra	SP	-6.885	-36.249	(OH..O) SER 126	2.64
						(OH..O) ASN133	2.81
						(OH..O) MET135	2.92
			XP	-5.71	-36.108	(OH..O) SER 126	2.64
						(OH..O) MET 135	2.95
						(OH..O) ASN 133	2.8
PDB IEA3			SP	-6.012	-33.074	lig(OH..O) ASP 89	2.71
						lig(OH..O) ASN 85	2.21
			XP	-4.993	-32.676	(OH..O) ASN 85	2.12
						(OH..O) SER 157	2.9

Fig 7.a Davidigenin docking with modeled Matrix M1

Fig 7.b Davidigenin docking with PDB Matrix M1 (IEA3)

Among the 392 virtually screened phytochemicals, Davidigenin from *Glycyrrhiza glabra* was found to exhibit good glide score and minimal energy against the matrix protein M1. The interactions were also good with the active site residues of both the modeled protein and PDB structures. The compound also obeyed Lipinski rule of five with the molecular weight of 258.23g/mol, LogP value of 3, with 3 Hydrogen bond donors and 4 hydrogen bond acceptors. Percentage of human oral absorption was 89% which makes the compound an effective lead for drug development.

Table 9: Docking results of the Neuraminidase protein

Modeled / PDB structure	Compound Name & PubChem ID	Plant Name	SP/ XP	Glide Score	Glide energy	Interactions	Bond length (Å°)
Modeled	Licoflavone B 11349817	Glycyrrhiza glabra	SP	-5.31	-37.367	(OH..O) TRP 102	2.970
						SER 161(OH..O)	2.960
			XP	-3.7	-30.894	LYS 20 (NH2..O)	2.95
						(OH..O) ASP 24	2.68
PDB 2GX9			SP	-4.37	-34.324	(OH..O) MSE 98	2.18
						(OH..O) TRP 102	2.79
			ARG 148(NH2..O)			2.49	
			SER 161(OH..O)			2.07	
			XP	-4.37	-34.324	(OH..O) MSE 98	2.18
						(OH..O) TRP 102	1.79

whole genomes of 12 different strains of Influenza A virus which has been recorded to infect humans that included H1N1, H1N2, H3N3, H5N1, H7N3, H2N2, H5N6, H7N2, H7N7, H7N9, H9N2, H10N8. The Influenza virus is 8 segmented single-stranded negative RNA virus where each segment codes for a protein. The pan-genome analysis revealed that though the strains were continuously mutating over time, there are a set of genes and proteins that are strongly conserved. The segments 6, 7, and 8 were included in this study because they were less targeted. Virtually screening of antiviral compounds led to the identification of novel lead compounds, 2,4,4' Trihydroxychalcone, davidigenin and licofalvone B from the well known Indian Plant *Glycyrrhiza glabra*. The current study is an extensive computational approach starting from comparative genome analysis, validating the core genes, identifying the acceptable targets and predicting leads that inhibit the targets. The study requires an investigational invitro and invivo tests done subsequently to be able to prove but still the results suggested here is valid and recommends proceeding for further research. If successful we will be able to target multiple strains of influenza viruses at the shortest time span.

Acknowledgment

Deep gratitude to DST-FIST grants which had enabled us to procure the state of art drug development Schrödinger suite.

Conflicts of Interests

No conflicts of interest among the authors.

Abbreviations used

BLAST	- Basic Local Alignment Search Tool
BLASTP	- Basic Local Alignment Search Tool of Proteins
ORF	- Open Reading Frames
CASTp	- Computed Atlas of Surface Topography of Proteins
ADME	- Absorption, Distribution, Metabolism, and Excretion
SP	- Standard Precision
XP	- Extra Precision
PDB	- Protein Data Bank

References

1. Aishwarya Sekar, Nargis Begum, and Anita Margret, An in silico study of lignans as selective estrogen receptor modulators to treat viral infections, International Research Journal of Biological Sciences, 2019: 8(3): 19-25.

2. Benton DJ, Wharton SA, Martin SR, McCauley JW. Role of Neuraminidase in Influenza A(H7N9) Virus Receptor Binding. Williams BRG, ed. *J Virol*. 2017;91(11):e02293-16. doi:10.1128/JVI.02293-16
3. Couch RB, Bayas JM, Caso C, *et al*. Superior antigen-specific CD4+T-cell response with AS03-adjuvantation of a trivalent influenza vaccine in a randomized trial of adults aged 65 and older. *BMC Infect Dis*. 2014;14(1):425. doi:10.1186/1471-2334-14-425
4. Darricarrère N, Pougatcheva S, Duan X, *et al*. Development of a Pan-H1 Influenza Vaccine. Schultz-Cherry S, ed. *J Virol*. 2018;92(22):e01349-18. doi:10.1128/JVI.01349-18
5. J. HA, Victoria G, R. DA, Pu LY. The evolution of human influenza viruses. *Philos Trans R Soc London Ser B Biol Sci*. 2001;356(1416):1861-1870. doi:10.1098/rstb.2001.0999
6. Miller MS, Gardner TJ, Krammer F, *et al*. Neutralizing Antibodies Against Previously Encountered Influenza Virus Strains Increase over Time: A Longitudinal Analysis. *Sci Transl Med*. 2013;5(198):198ra107 LP-198ra107. doi:10.1126/scitranslmed.3006637
7. Mok BW-Y, Song W, Wang P, *et al*. The NS1 Protein of Influenza A Virus Interacts with Cellular Processing Bodies and Stress Granules through RNA-Associated Protein 55 (RAP55) during Virus Infection. *J Virol*. 2012;86(23):12695 LP-12707. doi:10.1128/JVI.00647-12
8. Shin W-J, Seong BL. Novel antiviral drug discovery strategies to tackle drug-resistant mutants of influenza virus strains. *Expert Opin Drug Discov*. 2019;14(2):153-168. doi:10.1080/17460441.2019.1560261
9. Santhi N, Aishwarya S. Insights from the molecular docking of withanolide derivatives to the target protein PknG from Mycobacterium tuberculosis. *Bioinformation*. 2011;7(1):1–4.
10. Zhang T, Bi Y, Tian H, *et al*. Human infection with influenza virus A(H10N8) from live poultry markets, China, 2014. *Emerg Infect Dis*. 2014;20(12):2076–2079. doi:10.3201/eid2012.14091