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Nelfinavir was predicted to be a potential inhibitor of 2019-nCov main protease by an integrative approach combining homology modelling, molecular docking and binding free energy calculation

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Abstract

2019-nCov has caused more than 80 deaths as of 27 January 2020 in China, and infection cases have been reported in more than 10 countries. However, there is no approved drug to treat the disease. 2019-nCov M^{pro} is a potential drug target to combat the virus. We built homology models based on SARS M^{pro} structures, and docked 1903 small molecule drugs to the models. Based on the docking score and the 3D similarity of the binding mode to the known M^{pro} ligands, 4 drugs were selected for binding free energy calculations. Both MM/GBSA and SIE methods voted for nelfinavir, with the binding free energy of -24.69 ± 0.52 kcal/mol and -9.42 ± 0.04 kcal/mol, respectively. Therefore, we suggested that nelfinavir might be a potential inhibitor against 2019-nCov M^{pro}.

1. Introduction

In December 2019, cluster of patients with pneumonia were reported in Wuhan, Hubei Province, China.^{1, 2} Shortly, a new coronavirus, temporally named 2019-nCov, was identified to be the cause of the disease, which is the seventh member of the family betacoronavirus.¹ More than 2,700 infection cases and 80 deaths were reported as of 27 January 2020 from China. In addition, infection cases have been reported from Thailand, Australia, Malaysia, Singapore, France, Japan, South Korea, the United States, Vietnam, Canada and Nepal, indicating that the disease is a potential threat to the global health.³ Sadly, the number is still growing rapidly and no drug has been approved to be effective. Therefore, it is urgent to discover and develop drugs to cure the disease.

Based on its function, the main protease (M^{pro}) or chymotrypsin-like protease ($3CL^{pro}$)⁴ is suggested to be a potential drug target to combat 2019-nCov, which is highly conservable among coronaviruses. Sequence alignment revealed that the M^{pro} of 2019-nCov shares 96% similarity with that of SARS (severe acute respiratory syndrome) (Figure 1). Studies for identifying the inhibitors of 2019-nCov M^{pro} were quickly performed for discovering and developing drugs against the disease. For example, Hualiang Jiang and collaborators identified 30 drugs and compounds as the M^{pro} inhibitors via protein modelling and virtual screening, which is a rapid progress in the way to cope with the crisis. In addition, one of the 30 drugs/compounds, remdesivir, was also suggested to be potential inhibitor against 2019-nCov by Liu et al,⁵ who also suggested 3 other possible inhibitors.



Figure 1. Sequence alignment of 2019-nCov M^{pro} and SARS M^{pro}.

For finding more potential drugs as inhibitors of the protein, we modelled the 2019-nCov M^{pro} structures using SARS M^{pro} (PDB ID: 2GTB) as template and docked 1903 approved drugs against the model in this study. Fifteen drugs were selected based on the docking score and three dimensional (3D) similarity to available M^{pro} inhibitors against other coronavirus. For validation, we modelled 10 additional new models of 2019-nCov M^{pro} and docked the 15 drugs against the new models, which revealed that 6 drugs (nelfinavir, praziquantel, pitavastatin, perampanel, eszopiclone, and zopiclone) have good binding modes with the new models. Binding free energy calculation were then performed for 4 of the 6 drugs using MM/GBSA and SIE methods. Nelfinavir was identified as the best one with predicted binding free energies of -24.69 ± 0.52 kcal/mol by MM/GBSA and -9.42 ± 0.04 kcal/mol by SIE, respectively. Taking into account its high 3D similarity of binding mode to the known M^{pro} inhibitor, we proposed that nelfinavir should be a potential inhibitor against 2019-nCov M^{pro}.

2. Materials and methods

2.1 Homology modelling. 43 M^{pro} complexes with ligands were downloaded from protein data bank⁶ (PDB IDs: 1WOF, 2A5I, 2A5K, 2ALV, 2AMD, 2GTB, 2GX4, 2GZ7, 2GZ8, 2OP9, 2QIQ, 2V6N, 2ZU4, 2ZU5, 3SN8, 3SND, 3SZN, 3TIT, 3TIU, 3TNS, 3TNT, 3V3M, 4F49, 4MDS, 4TWW, 4TWY, 4WY3, 4YLU, 4YOG, 4YOI, 4YOJ, 4ZRO, 5C5N, 5C5O, 5EU8, 5N5O, 5N19, 5NH0, 5WKJ, 5WKK, 5WKL, 5WKM,

6FV1) and aligned to 2GTB in PyMOL.⁷ 11 complexes (PDB IDs: 2A5K, 2GTB, 2GX4, 3SND, 3TNS, 3V3M, 4F49, 4YLU, 5NH0, 5WKJ, 5WKM) were served as templates to build 11 2019-nCov M^{pro} models in SWISS-MODEL server by “user template” mode.⁸

2.2 Approved Drugs. 1905 approved small molecule drugs with 3D coordinates were downloaded from DrugBank release version 5.1.5,⁹ while 1903 drugs could be converted to pdbqt format by `prepare_ligand4.py` script in MGLToos version 1.5.6.¹⁰

2.3 Molecular Docking. 1903 approved drugs in pdbqt format were docked to 2019-nCov M^{pro} model (template: 2GTB) by `smina`,¹¹ which is a fork of AutoDock Vina¹² with improving scoring and minimization. The hydrogens were added to 2019-nCov M^{pro} model by `pdb2pqr` (`--ff=amber --ffout=amber --chain --with-ph=7`).¹³ Then the model was converted to pdbqt format by `prepare_receptor4.py` script in MGLToos version 1.5.6.¹⁰ The ligand in 2GTB was used to define the grid and the buffer space was set to 6.0 Å (`autobox_add`). The random seed was explicitly set to 0 (`seed`). The exhaustiveness of the global search was set to 32 (`exhaustiveness`) and at most 1 binding mode was generated for each drug (`num_modes`). MolShaCS, which utilized Gaussian-based description of molecular shape and charge distribution, was used to calculate the 3D similarities between approved drugs and available M^{pro} inhibitors.¹⁴

2.4 Molecular dynamics simulation. Each simulation system was immersed in a cubic box of TIP3P water that was extended by 9 Å from the solute, with a rational number of counter ions of Na⁺ or Cl⁻ to neutralize the system. General Amber force field (GAFF)¹⁵ and Amber ff03 force field¹⁶ were used to parameterize the ligand and protein, respectively. 10,000 steps of minimization with constraints (10 kcal/mol/Å²) on heavy atoms of complex, including 5,000 steps of steepest descent minimization and 5,000 steps of conjugate gradient minimization, was used to optimize each system. Then each system was heated to 300 K within 0.2 ns followed by 0.1 ns equilibration in NPT ensemble. Finally, 5 ns MD simulation on each system at 300 K was performed. The minimization, heating and equilibrium are performed with *sander* program in Amber16. The 5 ns production run was performed with *pmemd.cuda*.

2.5 Binding free energy calculation. Based on the 5 ns MD simulation trajectory, bind

free energy (ΔG) was calculated with MM/GBSA^{17, 18} and SIE¹⁹ approaches. In the MM/GBSA, the ΔG was calculated according to equation (1),

$$\Delta G = \Delta H - T\Delta S = \Delta E_{ele} + \Delta E_{VDW} + \Delta G_{gb} + \Delta G_{np} - T\Delta S \quad (1)$$

where ΔE_{ele} and ΔE_{VDW} refer to electrostatic and van der Waals energy terms, respectively. ΔG_{gb} and ΔG_{np} refer to polar and non-polar solvation free energies, respectively. Conformational entropy ($T\Delta S$) was calculated by *nmode* module in Amber16. The dielectric constants for solvent and solute were set to 80.0 and 1.0, respectively, and OBC solvation model ($igb = 5$ and $PBradii = 5$)²⁰ was used in this study. Other parameters are set to default values.

In the SIE, the ΔG was calculated based on equation (2),

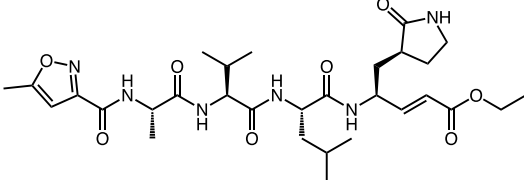
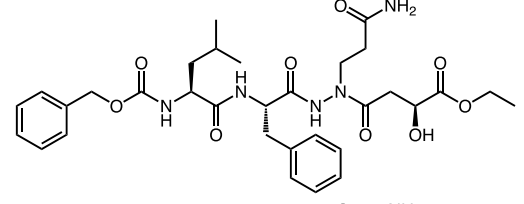
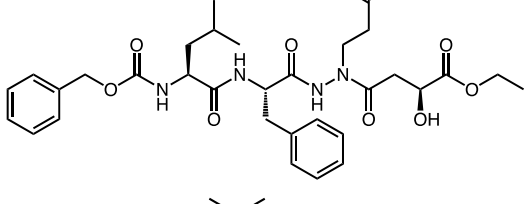
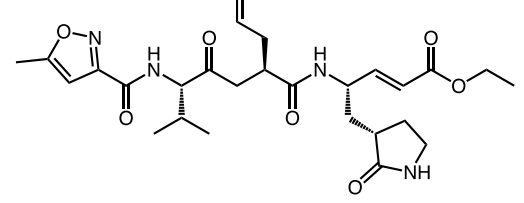
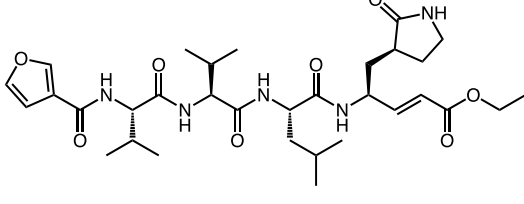
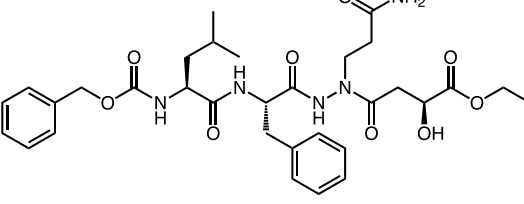
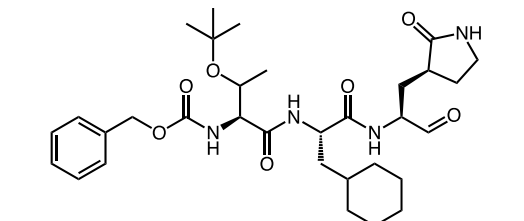
$$\Delta G_{bind} = \alpha[E_C(D_{in}) + \Delta G_{bind}^R(\rho, D_{in}) + E_{VDW} + \gamma\Delta MSA(\rho)] + C \quad (2)$$

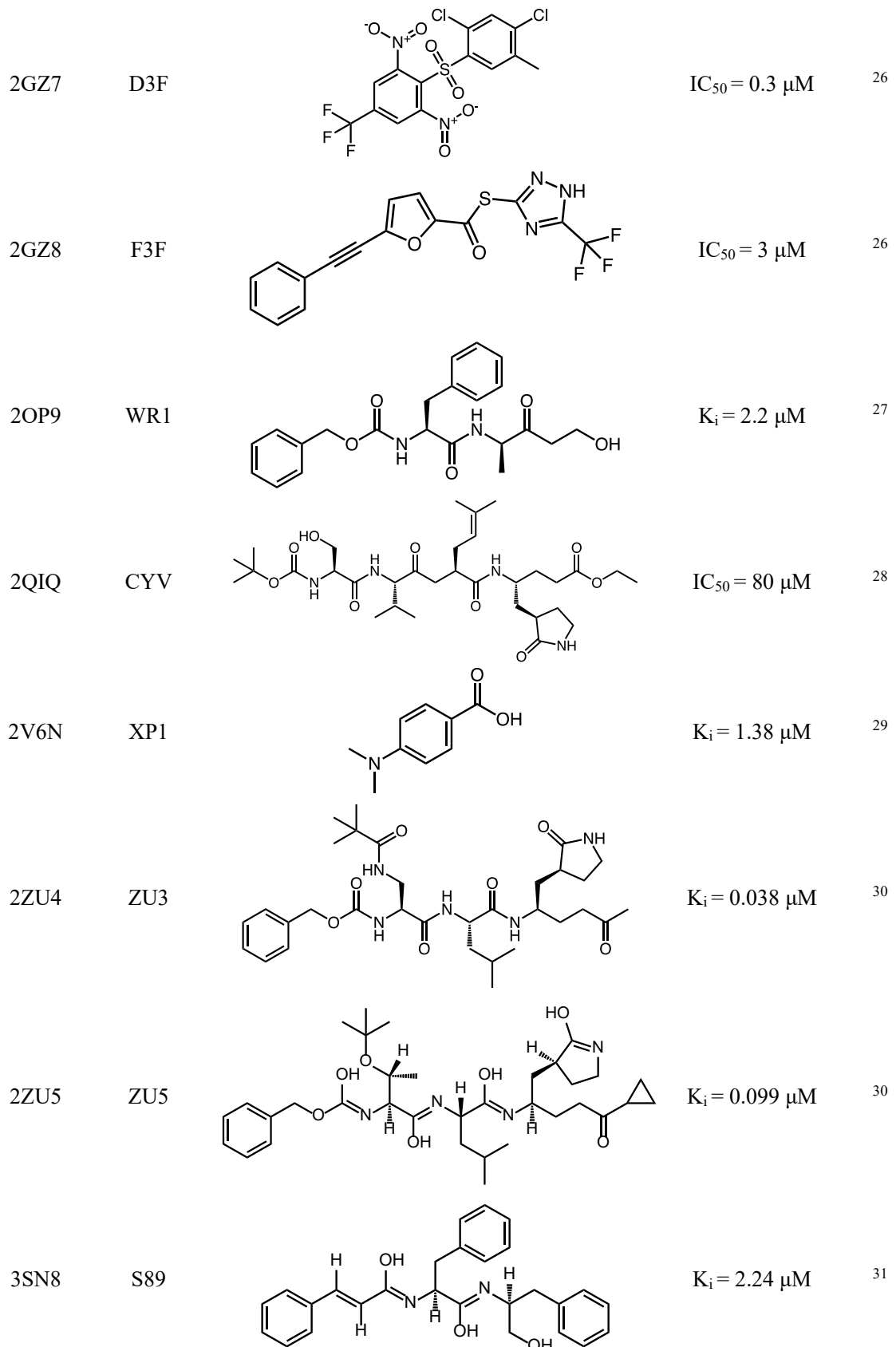
where E_C and E_{VDW} refer to the sum of intermolecular Coulomb and van der Waals interaction energies, respectively. ΔG_{bind}^R and ΔMSA refer to the changes of reaction field energy and molecular surface area upon ligand binding, respectively. Default values of the global proportionality coefficient ($\alpha = 0.1048$), the solute interior dielectric constant ($D_{in} = 2.25$), the van der Waals radii linear scaling coefficient ($\rho = 1.1$), the molecular surface area coefficient ($\gamma = 0.0129 \text{ kcal/mol/\AA}^2$), and the constant ($C = -2.89 \text{ kcal/mol}$) are used in this study.

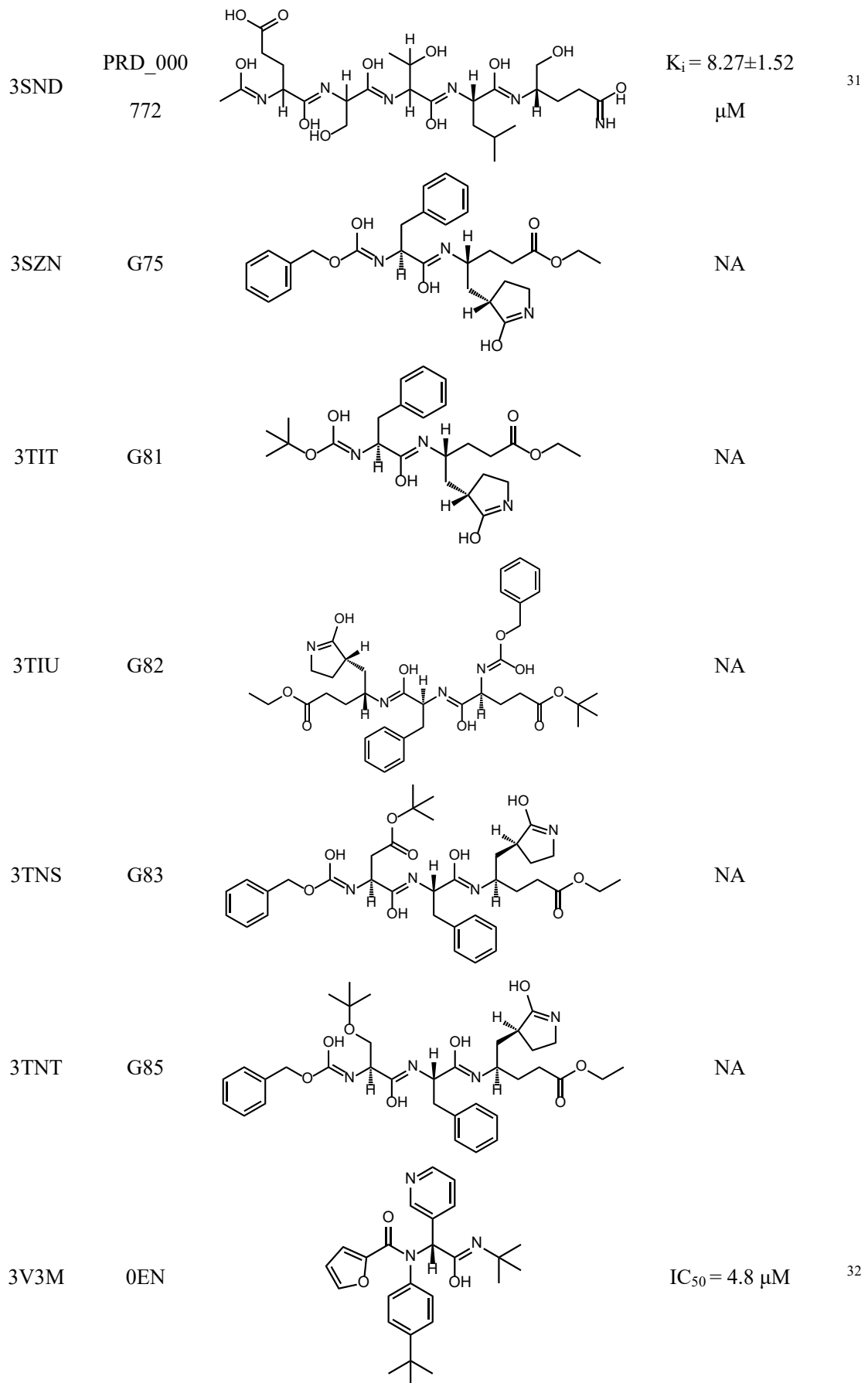
3. Results and discussion

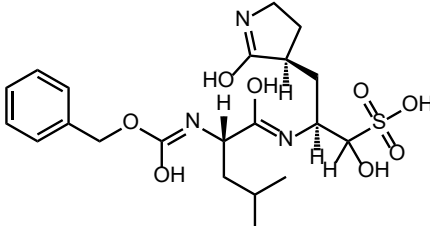
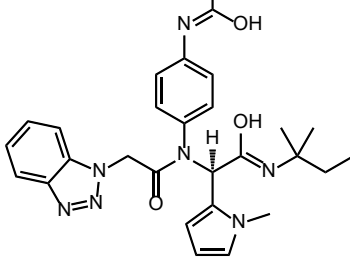
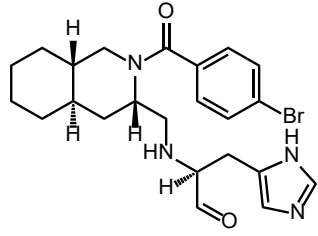
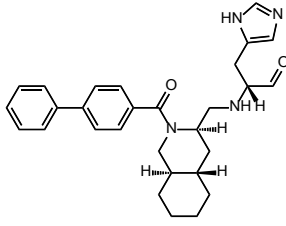
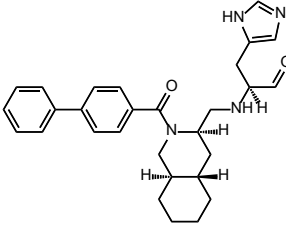
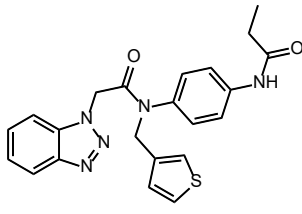
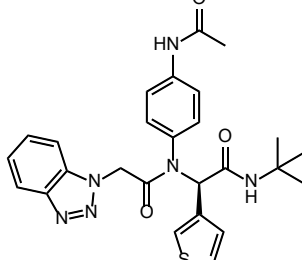
3.1 Docking results of 1903 approved drugs against M^{pro} models. 1903 approved small molecule drugs were docked to a M^{pro} homology model built by SWISS-MODEL using 2GTB as template. There are 672 drugs with the docking score better than -7.0 kcal/mol. The binding modes are essential to the activities. If a compound shares a similar binding mode to a known ligand, it is more likely to have a similar activity to the ligand. Therefore, we calculated the 3D similarities of the binding mode of the docked drugs to the available binders against M^{pro}. 39 binders in 44 complexes were chosen as the references to calculate the 3D similarities (Table 1). 159 drugs have a 3D similarity larger than 65.0% with at least one known M^{pro} binder.

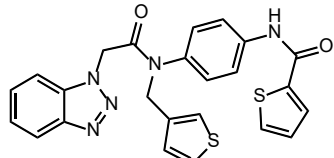
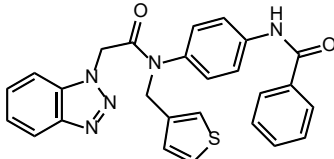
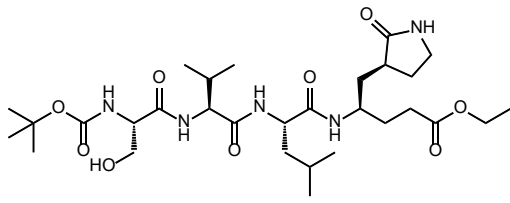
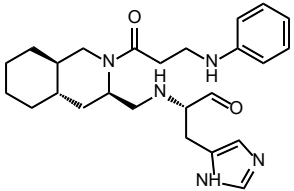
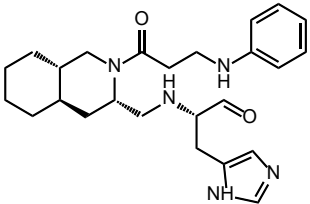
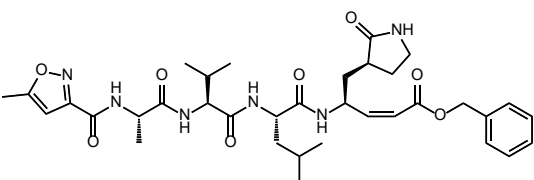
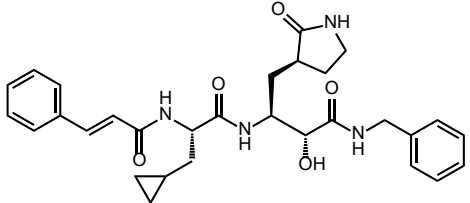
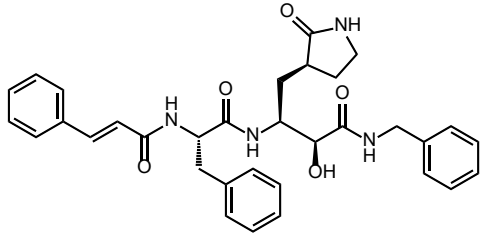
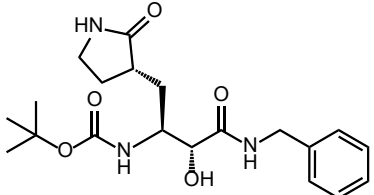
Table 1. Known M^{pro} binders as references to calculate the 3D similarity of binding modes.

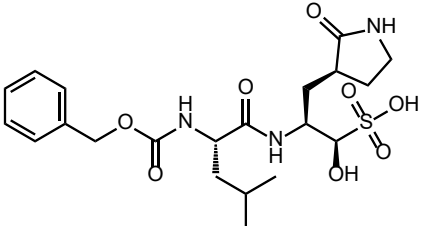
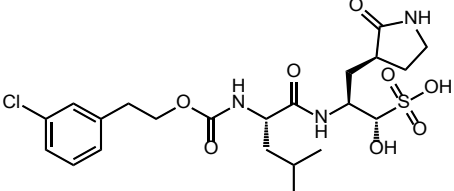
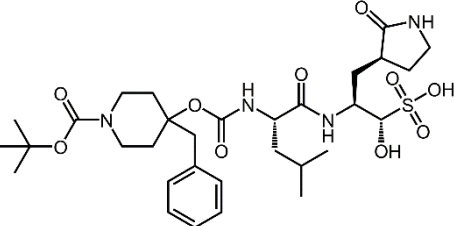
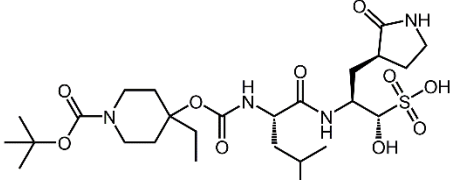
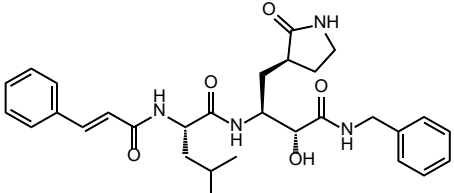
PDB ID	Ligand ID	Ligand Structure	Activity	Ref
1WOF	I12		$K_i = 10.7 \mu\text{M}$	21
2A5I	AZP		$K_i = 18 \mu\text{M}$	22
2A5K	AZP		$K_i = 18 \mu\text{M}$	22
2ALV	CY6		$IC_{50} = 70 \mu\text{M}$	23
2AMD	9IN		$K_i = 6.7 \mu\text{M}$	21
2GTB	AZP		$kinact/K_i = 1900 \pm 400 \text{ M}^{-1} \text{ s}^{-1}$	24
2GX4	NOL		$K_i = 0.053 \mu\text{M}$	25





4F49	K36		$IC_{50} = 0.82 \mu M$	33
4MDS	23H		$IC_{50} = 6.2 \mu M$	34
4TWW	3A7		$IC_{50} = 63 \mu M$	35
4TWY	3BL		$IC_{50} = 108 \mu M$	35
4WY3	3X5		$IC_{50} = 240 \mu M$	35
4YLU	R30		$IC_{50} > 100 \mu M$	36
4YOG	4F5		$IC_{50} = 1.8 \mu M$	37

4YOI	4F4		$IC_{50} = 0.33 \mu M$	37
4YOJ	RFM		$IC_{50} = 0.41 \mu M$	37
4ZRO	PRD_002 174		$IC_{50} = 0.59 \mu M$	38
5C5N	SLH		NA	
5C5O	SDJ		NA	
5EU8	PRD_002 214		NA	
5N5O	8O5		NA	
5N19	D03		NA	
5NH0	8X8		NA	

5WKJ	B1S		$EC_{50} = 0.9 \mu\text{M}^*$	39
5WKK	AW4		$EC_{50} = 0.5 \mu\text{M}^*$	39
5WKL	B3J		$IC_{50} = 0.8 \mu\text{M}$	39
5WKM	N02		$IC_{50} = 6.1 \mu\text{M}$	39
6FV1	E8E		NA	

* Racemic activity; NA: Not Available.

After visualizing the docked complexes carefully, we selected 15 drugs (Table 2) for further analysis. There are some differences between the conformations of the protein in the 3D similarity reference and 2GTB model. Therefore, we modelled 10 new homology models using the proteins in 3D similarity reference as templates and we re-docked the 15 drugs to the 10 new homology models. 6 drugs (nelfinavir, pitavastatin, perampanel, praziquantel, zopiclone, and eszopiclone) show good docking scores and binding modes (Table 3). Because eszopiclone and zopiclone was used to treat insomnia in a low dosage which may not be suitable to treat pneumonia, we carried out further binding free energy calculation for the rest 4 drugs.

Table 2. 15 drugs selected from 2GTB model.

DrugBank ID	Docking Score against 2GTB model (kcal/mol)	3D Similarity Reference	3D Similarity
DB00328	-7.16	3V3M	71.3%
DB13680	-7.13	3SND	70.8%
DB01165	-7.31	3V3M	70.2%
DB01198	-7.81	5NH0	68.9%
DB08934	-8.65	2GX4	68.6%
DB01165	-7.31	5WKM	68.3%
DB08860	-8.06	2GTB	68.1%
DB00402	-7.75	5NH0	67.8%
DB08883	-8.17	5NH0	67.1%
DB01288	-7.96	5WKM	67.1%
DB08860	-8.06	2A5K	66.9%
DB00972	-7.71	5NH0	66.7%
DB00482	-8.46	4YLU	66.7%
DB00220	-8.91	2GX4	66.4%
DB01058	-7.90	3V3M	65.7%
DB00328	-7.16	4F49	65.6%
DB08934	-8.65	3TNS	65.5%
DB00904	-7.45	5WKM	65.4%
DB11951	-8.27	4F49	65.3%
DB11951	-8.27	5WKJ	65.2%

Table 3. 6 drugs selected from 10 new homology models.

DrugBank ID	Name	Homology Template	Docking Score (kcal/mol)	3D Similarity
DB00220	nelfinavir	2GX4	-9.18	70.2%
DB08860	pitavastatin	2GTB	-8.06	68.1%
DB08883	perampanel	5NH0	-8.63	66.9%
DB01058	praziquantel	3V3M	-7.38	65.8%
DB01198	zopiclone	5NH0	-8.46	63.5%
DB00402	eszopiclone	5NH0	-8.46	63.5%

3.2 Binding Free Energy calculated by MM/GBSA and SIE. The 4 docked complex structures, i.e., nelfinavir-2GX4, pitavastatin-2GTB, perampanel-5NH0 and praziquantel-3V3M, were subjected to 5 ns molecular dynamics simulations using

Amber 16. To provide insight into their binding mechanisms, the binding free energies were calculated by MM/GBSA and SIE approaches. In results of the MM/GBSA approach, the calculated binding free energies of nelfinavir-2GX4, pitavastatin-2GTB, perampanel-5NH0, and praziquantel-3V3M, are -24.69 ± 0.52 , -12.70 ± 0.38 , -14.98 ± 0.34 , and -6.51 ± 0.21 kcal/mol, respectively (Table 4), which highlight nelfinavir as the most active one. In pitavastatin-2GTB, perampanel-5NH0, and praziquantel-3V3M, the van der Waals interaction (E_{vdw}) makes a more significant contribution than the electrostatic interaction (E_{ele}) (Table 4), indicating that van der Waals interaction is the main driving force for the 3 drugs binding. However, the E_{ele} interaction of nelfinavir-2GX4 is very strong, suggesting that the electrostatic interaction also play an important role in the binding of nelfinavir. Furthermore, in results of the SIE approach, the rank of the calculated binding free energies for the 4 docking complexes is consistent with that of the MM/GBSA approach (Table 5) that the nelfinavir has the strongest binding free energy (-9.42 ± 0.04 kcal/mol), indicating the reliability of the current binding free energy analysis.

Table 4. Components of the Binding Free Energy (kcal/mol) Calculated by MM/GBSA Approach*.

Energy term	nelfinavir	pitavastatin	perampanel	praziquantel
E_{vdw}	-58.36 ± 0.32	-42.77 ± 0.30	-44.33 ± 0.25	-34.92 ± 0.32
E_{ele}	-117.95 ± 0.81	37.95 ± 0.86	0.05 ± 0.03	-8.64 ± 0.35
G_{gb}	134.49 ± 0.83	-22.40 ± 1.63	14.76 ± 0.29	21.53 ± 0.32
G_{np}	-6.69 ± 0.03	-5.60 ± 0.03	-5.43 ± 0.03	-4.34 ± 0.03
ΔH	-48.50 ± 0.36	-32.82 ± 0.38	-34.96 ± 0.27	-26.37 ± 0.30
$T\Delta S$	-23.81 ± 0.67	-20.12 ± 0.38	-19.98 ± 0.42	-19.86 ± 0.12
ΔG_{cal}	-24.69 ± 0.52	-12.70 ± 0.38	-14.98 ± 0.34	-6.51 ± 0.21

*: The statistical error was estimated based on 0.5-5 ns MD simulation trajectory. 500 snapshots evenly extracted from the 0.5-5 ns MD trajectory of complex were used for MM/GBSA calculations and 50 snapshots for the entropy term calculations.

Table 5. Components of the Binding Free Energy (kcal/mol) Calculated by SIE

Approach*.

Energy term	nelfinavir	pitavastatin	perampanel	praziquantel
E_{vdw}	-58.36±0.32	-42.77±0.30	-44.33±0.25	-34.92±0.32
E_C	-52.44±0.36	16.87±0.82	0.02±0.01	-3.84±0.16
$\Delta G_{\text{bind}}^{\text{R}}$	58.44±0.30	-10.57±0.67	8.06±0.15	12.05±0.16
E_{cavity}	-9.93±0.05	-7.79±0.04	-8.19±0.04	-6.66±0.06
ΔG_{cal}	-9.42±0.04	-7.53±0.04	-7.55±0.03	-6.39±0.04

*: The binding free energy was calculated with the equation: $\Delta G_{\text{bind}} = 0.1048 * (E_C + E_{\text{vdw}} + \Delta G_{\text{bind}}^{\text{R}} + E_{\text{cavity}}) - 2.89$. $E_{\text{cavity}} = \gamma \Delta \text{MSA}$. The statistical error was estimated based on 0.5-5 ns MD simulation trajectory. 500 snapshots evenly extracted from the 0.5-5 ns MD trajectory were used.

3.3 Binding modes of nelfinavir against 2019-nCov M^{pro}.

As shown in Figure 2, the binding model of nelfinavir in its docking complex turned out to be very similar with that of the original ligand (TG-0205221)²⁵ of 2GX4 (Figure 2a&b), which is an inhibitor of SARS M^{pro} with in vitro K_i of 0.053 μM and IC_{50} of 0.6 μM in the Vero-E6 cells.²⁵ In its crystal structure, TG-0205221 is able to form hydrogen bonds with HIS-163, GLU-166 and GLN-189, and a very weak hydrogen bond with PHE-140 (Figure 2c). Our docking results showed that three of the hydrogen bonds involving GLU-166 and GLN-189 maintained upon the binding of nelfinavir and 2019-nCov M^{pro}, with additionally the possible formation of π - π stacking interaction with HIS-41 (Figure 2d). These observations further demonstrated that nelfinavir would interact with key residues of 2019-nCov M^{pro} in a similar way to that of the existing inhibitors against coronavirus M^{pro}.

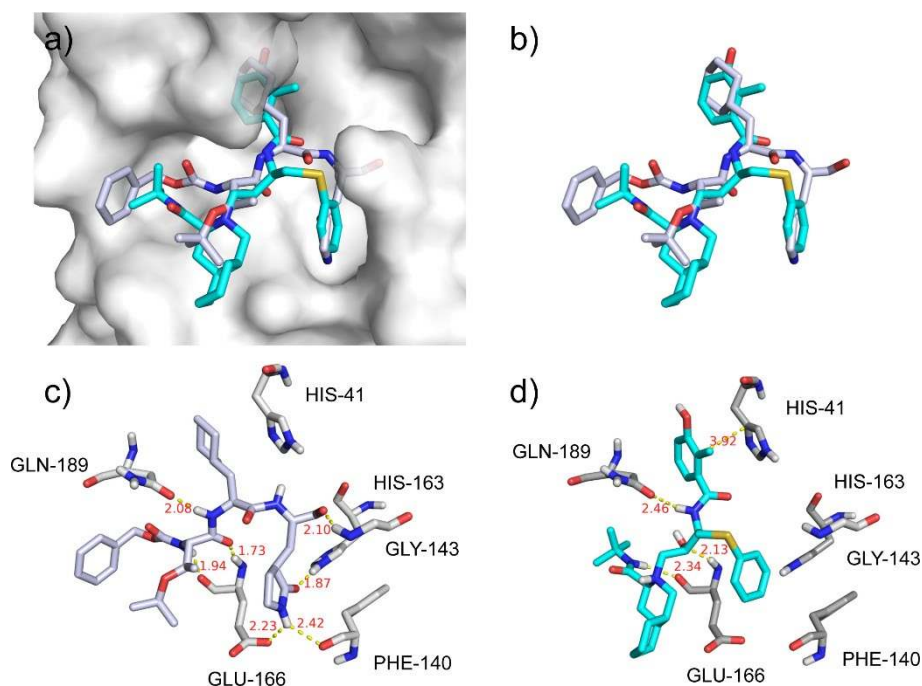


Figure 2. The binding model of nelfinavir against 2019-CoV M^{pro}. (a) Binding models of the original ligand (TG-0205221, white) in 2GX4 and nelfinavir (cyan) in the 2019-nCoV M^{pro} protein pocket (white surface); (b) Superposition of TG-0205221 (white) and nelfinavir (cyan) in their binding conformations; (c) Interactions between TG-0205221 and associated residues in the crystal structure (2GX4) of SARS M^{pro}; (d) Interactions between nelfinavir and associated residues in the homology model of 2019-nCoV M^{pro}. The data in red is the interaction distance (Å).

4. Conclusions

2019-nCoV caused more than 80 deaths in China as of 27 January 2020 and is a potential threat to the global health. However, there is no approved drug to treat the disease. 2019-nCoV M^{pro} is a potential drug target to combat the virus, which shares 96% sequence similarity with the corresponding one in SARS. We built 11 homology models of 2019-nCoV M^{pro} and docked 1903 approved small molecule drugs to the 2GTB model. Based on the docking score and the 3D similarity of binding mode to 39 known M^{pro} binders, 15 drugs were selected for further evaluation. The 15 drugs were then docked to all the 11 homology models, leading to 4 drugs for binding energy calculations. Both MM/GBSA and SIE calculations voted for nelfinavir, a HIV-1 protease inhibitor to treat HIV. Therefore, we suggested that nelfinavir might be active against 2019-nCoV M^{pro}. In addition, pitavastatin, perampanel, and praziquantel might

also have moderate activities against 2019-nCov.

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