



Cite this: *RSC Adv.*, 2020, **10**, 19790

Nature as a treasure trove of potential anti-SARS-CoV drug leads: a structural/mechanistic rationale†

Ahmed M. Sayed,^a Amira R. Khattab,^b Asmaa M. AboulMagd,^b Hossam M. Hassan,^b Mostafa E. Rateb,^f Hala Zaid^g and Usama Ramadan Abdelmohsen^b              <img alt="ORCID iD icon" data-bbox="7645 228 7665 24

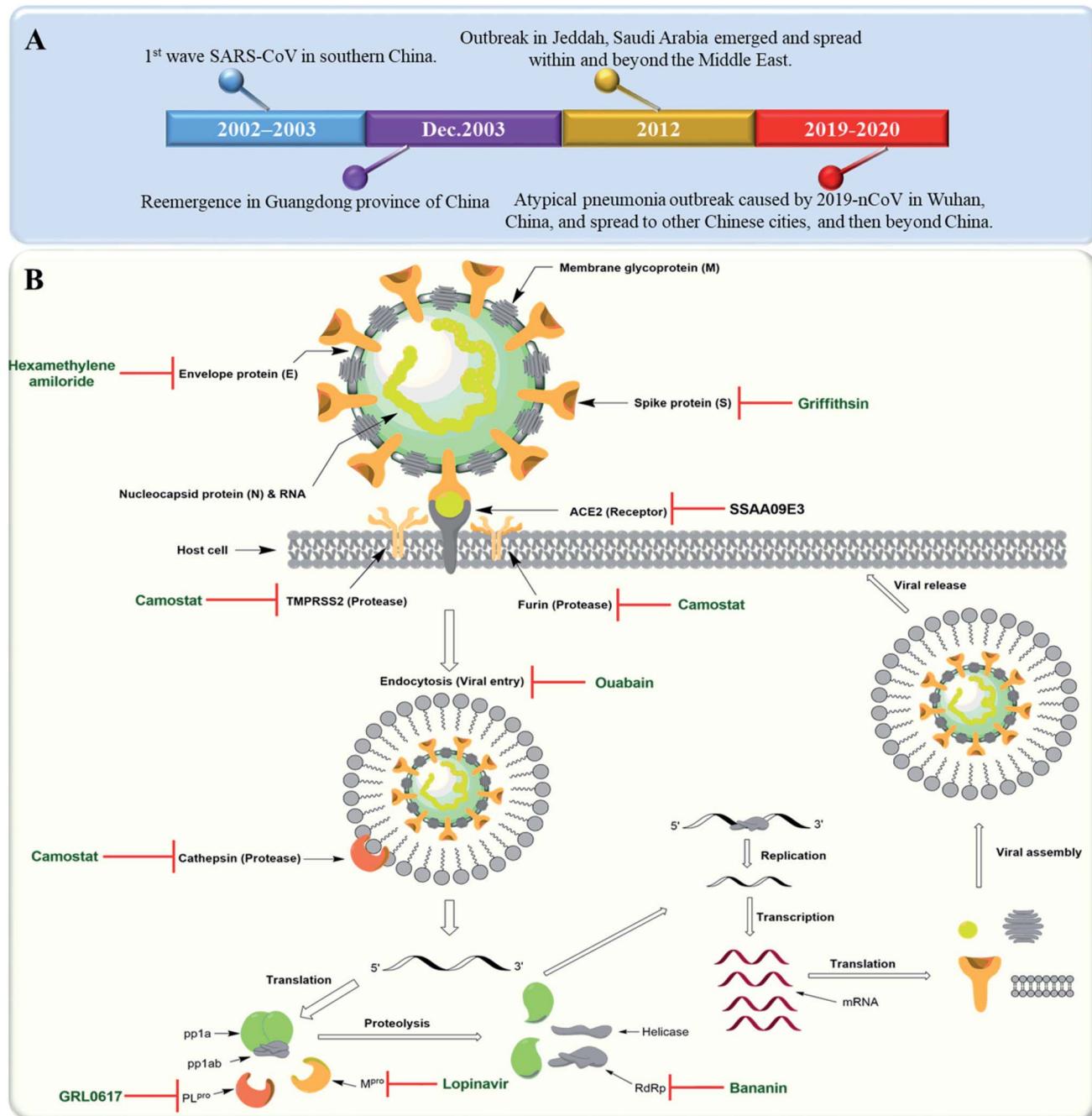


Fig. 1 (A) Timeline of SARS-CoV outbreaks during the last two decades. (B) Coronavirus structure and its life cycle, illustrating the main molecular targets and their inhibitors.

products against SARS-CoV". Cross-referencing and "related articles" functions were used to expand the search criteria. Searches were further supplemented with publicly available information and reports from US Centers for Disease Control and Prevention and WHO websites. Furthermore, we carried out an *in silico* exploration of a library of phenolic natural products (496 compounds) *via* a computer-aided virtual screening against the active site of the SARS-CoV Main protease (M^{Pro}) and a drug-likeness computation of the compounds reported in this review (75 compounds).

2. Anti-SARS CoV therapeutic drugs

2.1. Virus-based targets and their therapeutics

Many plant derived therapeutics exhibited an *in vitro* efficacy against multiple SARS viral targets *viz.* viral entry into host cells, viral processing (viral protease), viral replication (transcription and translation), and viral release from infected cells (Fig. 1B).

SARS-CoV replicase gene has been revealed to encode a number of enzymatic functions.⁵ These include M^{Pro} (nsp5), a papain-like protease (PL^{Pro} , nsp3), an RNA-dependent RNA



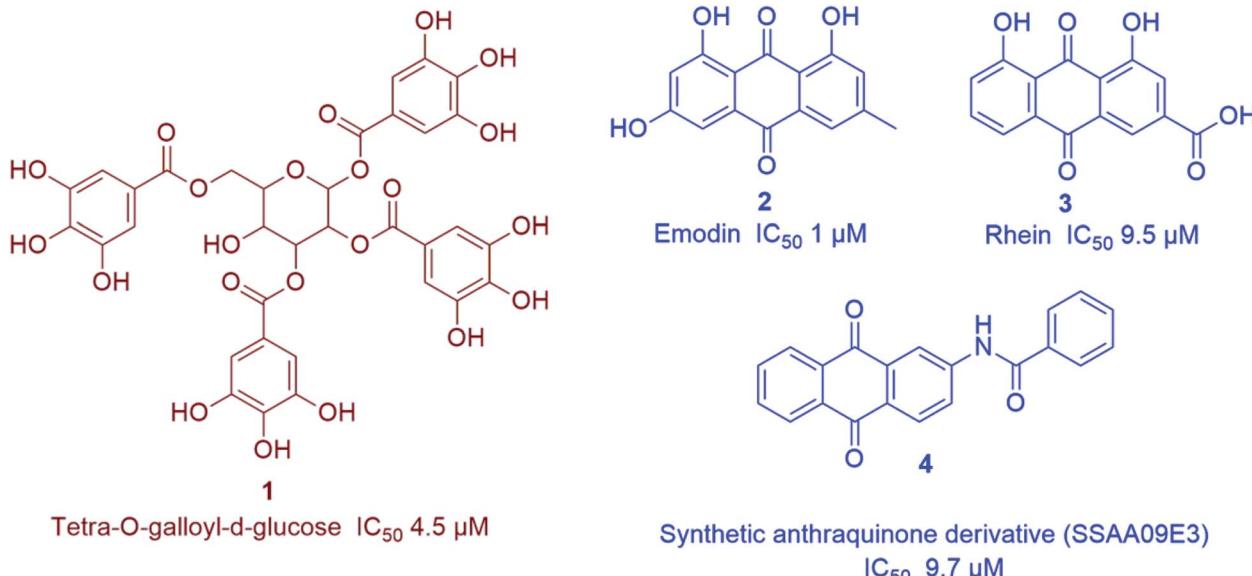


Fig. 2 Representative natural products with SARS-CoV S-protein-ACE-II binding inhibitory action (*in vitro*) viz. anthraquinones (blue) and tannins (brown).

polymerase (RdRp, nsp12), and a helicase (HEL1, nsp13). Such essential functional proteins have been selected as potential targets for the development of new anti-SARS-CoV therapeutics during the last two decades. Both viral M^{PRO} and PL^{PRO} cleave two large replicase polyproteins, *viz.* pp1a and pp1ab, that are encoded by the open reading frame 1a/b (ORF1a/b) to produce a number of non-structural proteins (nsp 1–15) such as RNA-dependent RNA polymerases (RdRp) and helicases that initiate the viral replication process. Such proteases can be considered key factors in the viral life cycle.⁶ Accordingly, their *in vitro* and *in vivo* inhibition has exhibited anti-SARS-CoV potential (Fig. 1B).

Several small molecules inhibitors for these two enzymes have been reported since 2002 (outlined in Fig. 1B), of which both lopinavir and ritonavir showed a broad spectrum *in vitro* M^{PRO} inhibitory activity⁷ as well as clinical efficacy in patients infected by COVID-19.⁸ Similarly, a number of specific and selective inhibitors have been developed against PL^{PRO}, and two of them with a common naphthalene benzamide scaffold are under clinical trials now.^{9,10} Broad-spectrum RdRp and helicase inhibitors like ribavirin and bananin, respectively have shown some clinical efficacy against SARS-CoV and MERS-CoV.^{11,12} The CoV small hydrophobic envelope protein (E-protein) was found to contain cation-selective ion channels similar to the mouse hepatitis virus (MHV) and human immunodeficiency virus (HIV). Inhibitors for such ion channels like hexamethylene amiloride have a well-proven *in vitro* and *in vivo* anti-CoV efficacy.¹³

2.2. Host-based targets and their therapeutics

ACE-II enzyme was found to be a functional and specific receptor for the viral spike-glycoproteins (S-protein). Hence, its antibodies were able to interfere with the viral entry to the host

cells, and thus blocking viral replication.¹⁴ The spike glycoproteins (S1 and S2 subunits) were also reported to be potential targets for neutralizing antibodies and oligosaccharide-specific binding proteins (*e.g.* Griffithsin, a natural product derived from the red algae *Griffithsia*), in addition to several small molecules fusion inhibitors (S2 subunit inhibitors).¹⁵ Other host proteases that could be targeted for drug development include the endosomal cysteine protease *i.e.* cathepsins that facilitate endosomal cell entry of CoV,¹⁶ and both the transmembrane protease serine 2 (TMPRSS2) and furin, that cleave and activate spike proteins into the S1 and S2 subunits to mediate the non-endosomal virus entry at the plasma membrane.¹⁷ The broad-spectrum serine protease inhibitor camostat possessed *in vitro* inhibitory activity toward cathepsins, furin and TMPRSS2 and significant *in vivo* efficacy. Moreover, viral entry *via* endocytosis was also found to be inhibited by cardiotonic steroids such as ouabain and bufalin at nanomolar concentrations¹⁸ (Fig. 1B).

3. Anti-SARS-CoV natural therapeutics

3.1. Viral entry inhibitors

Natural therapeutic agents that target viral entry are considered an important class of antiviral therapeutics because of their potential to block the first step of viral infection and hence, preventing its propagation and ability to evolve and acquire drug resistance. SARS-CoV virus entry is initiated when the spike glycoprotein (S) efficiently bind with a metallopeptidase, ACE-II on host cell.¹⁹

3.1.1. S-protein-ACE-II binding inhibitors. Several natural products that target CoV S-protein to prevent its binding with ACE-II, have been reported. In 2004, a group of Chinese



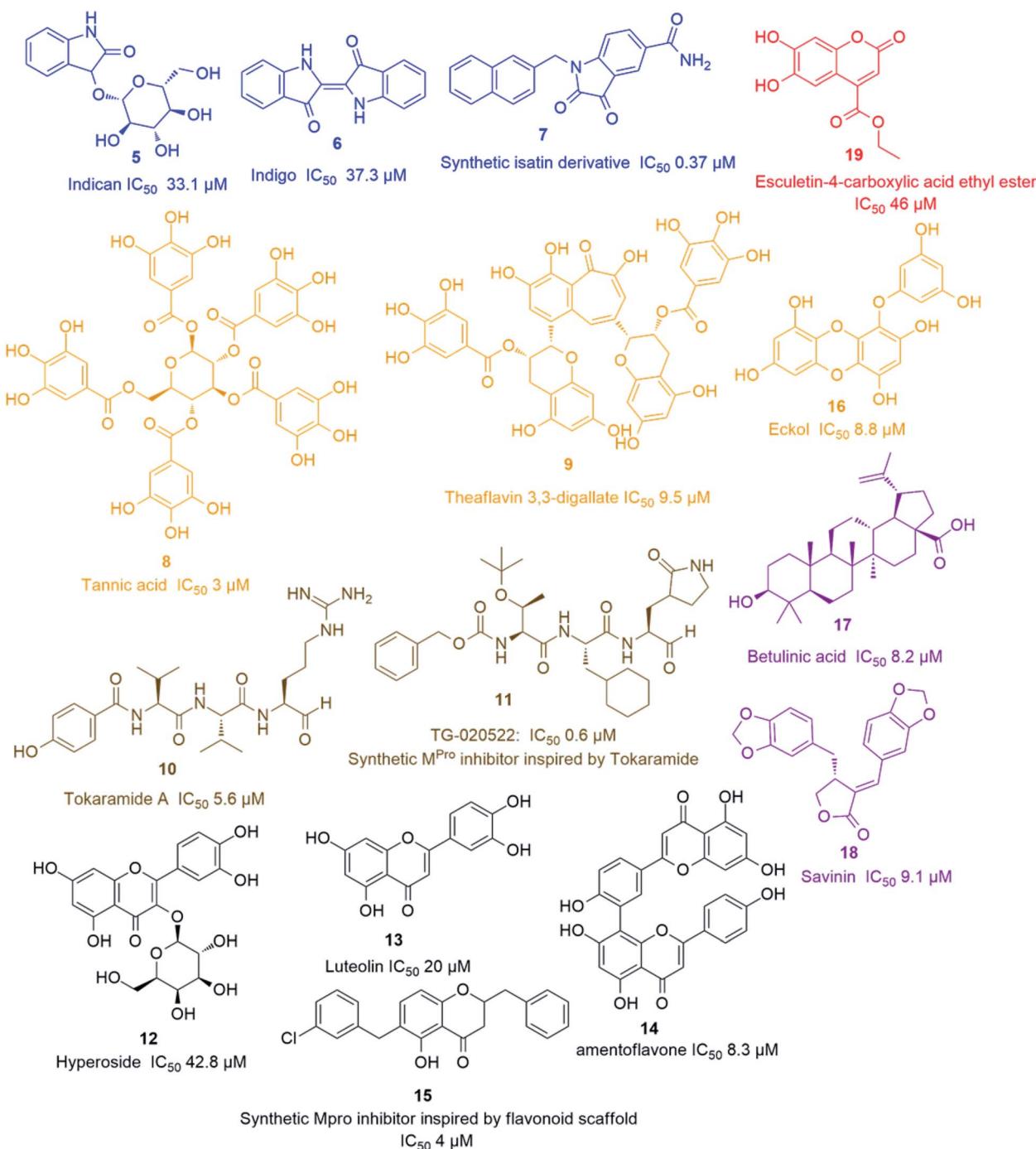


Fig. 3 Representative natural products with SARS-CoV M^{pro} *in vitro* inhibitory action viz. alkaloids (blue), coumarins (red), tannins (orange), peptides (brown), terpenes (purple) and flavonoids (black).

researchers screened their traditional herbal medicine (Traditional Chinese Medicine, TCM) for potential natural products that can bind with S-protein. Their extensive virtual screening followed by *in vitro* validation enabled them to identify a number of phenolic compounds, among which tetra-O-galloyl- β -D-glucose (1) (Fig. 2) was reported as a potent S-protein-ACE-II binding inhibitor (IC₅₀ 4.5 μ M).²⁰ Three years later, another Chinese research group screened a large number of medicinal plants belonging to Polygonaceae family, *i.e.* *Rheum officinale*

and *Polygonum multiflorum*, for the possible discovery of anti-CoV natural therapeutics. They found that emodin (2) and its related anthraquinone derivative, *i.e.* rhein (3) significantly blocked the ACE-II and S-protein interaction in a dose-dependent manner with IC₅₀ of 1–10 μ M (Fig. 2).²¹ Thereafter, this class of compounds inspired the synthesis of an anthraquinone-based derivative such as SSAA09E3 (4) (IC₅₀ 9.7 μ M) (Fig. 2) which exhibited interesting pharmacodynamic and pharmacokinetic properties and hence, it was selected to be

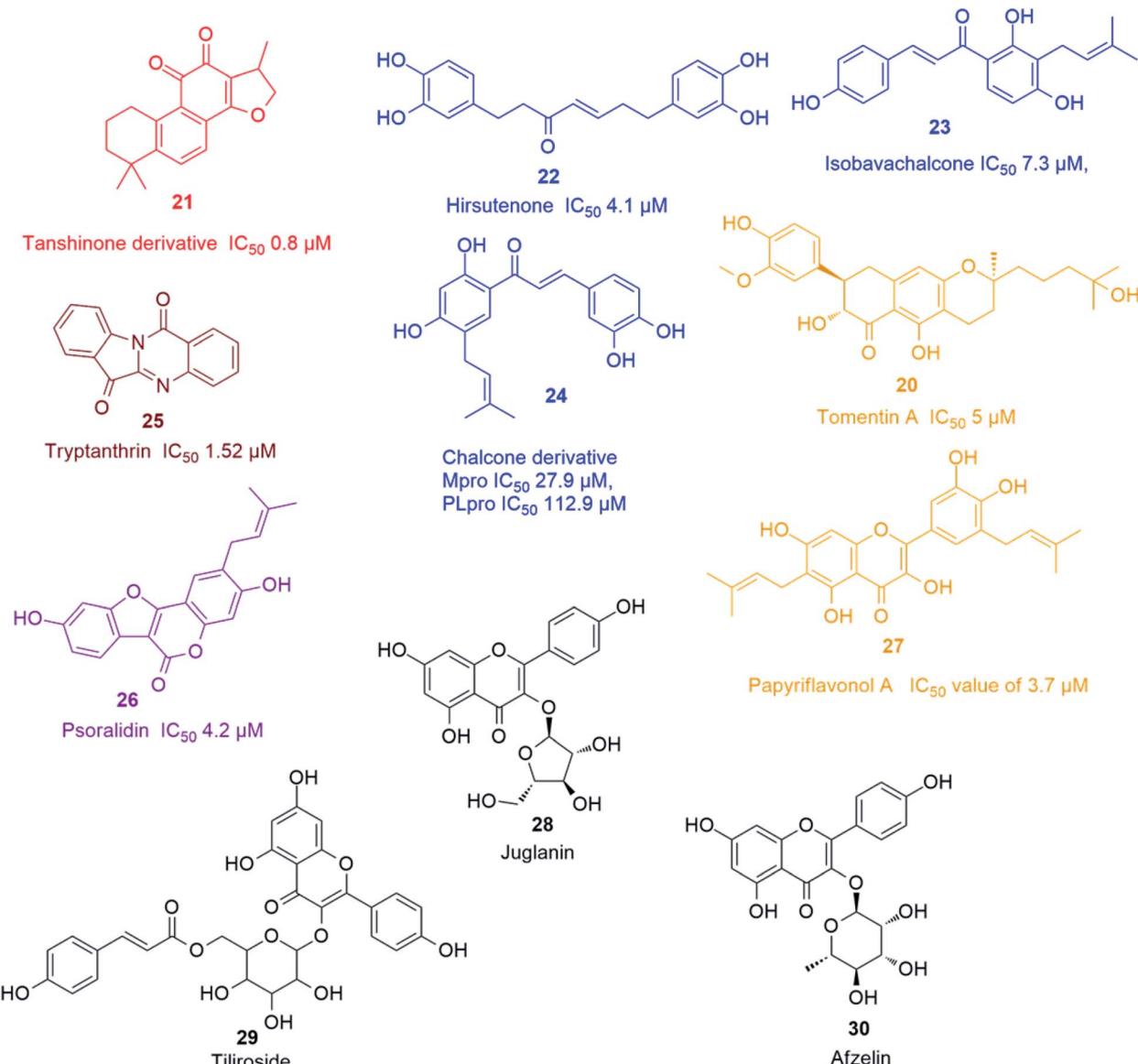


Fig. 4 Representative natural products with SARS CoV PL^{pro} *in vitro* inhibitory action viz. chalcones (blue), terpenes (Red), flavonoids (orange), alkaloids (brown), coumarins (purple), and viral release inhibitors viz. flavonoid glycosides (black).

further subjected for preclinical evaluation as a CoV entry inhibitor.²²

3.2. Inhibitors of viral replication

Being one of the most prominent molecular targets in COVID-19, M^{pro} and PL^{pro} inhibitors were selected for the development of potential anti-SARS CoV therapeutics. Several small molecules have been developed during and after the first and second CoV spread waves. Among which, many edible plants-derived natural products and their related synthetic derivatives have offered a number of interesting potential inhibitors with well-proven safety and efficacy.

3.2.1. M^{pro} inhibitors. *Isatis indigotica* was one of the first medicinal plants that have been reported to possess anti-SARS CoV potential. A deeper investigation into its metabolites that

could be attributable to the observed antiviral action led to the characterization of several isatin-derived alkaloids indican (5) and indigo (6) (Fig. 3) and other phenolic compounds with M^{pro} *in vitro* inhibitory action.²³ A year later, the isatin scaffold was utilized for developing more potent semi-synthetic derivative (7) that possessed M^{pro} inhibitory action at nanomolar concentrations with IC₅₀ of 0.37 μ M (Fig. 3).²⁴

The rapid development of computer-aided drug design (CADD) used for the *in silico* molecular modelling along with the growing small molecules libraries derived by the aid of combinatorial chemistry have dramatically improved the drug discovery and development process. During the last coronavirus waves, such computer-based approaches have contributed to finding a wide array of new and potent drug leads. Natural product libraries and databases have also been utilized to find



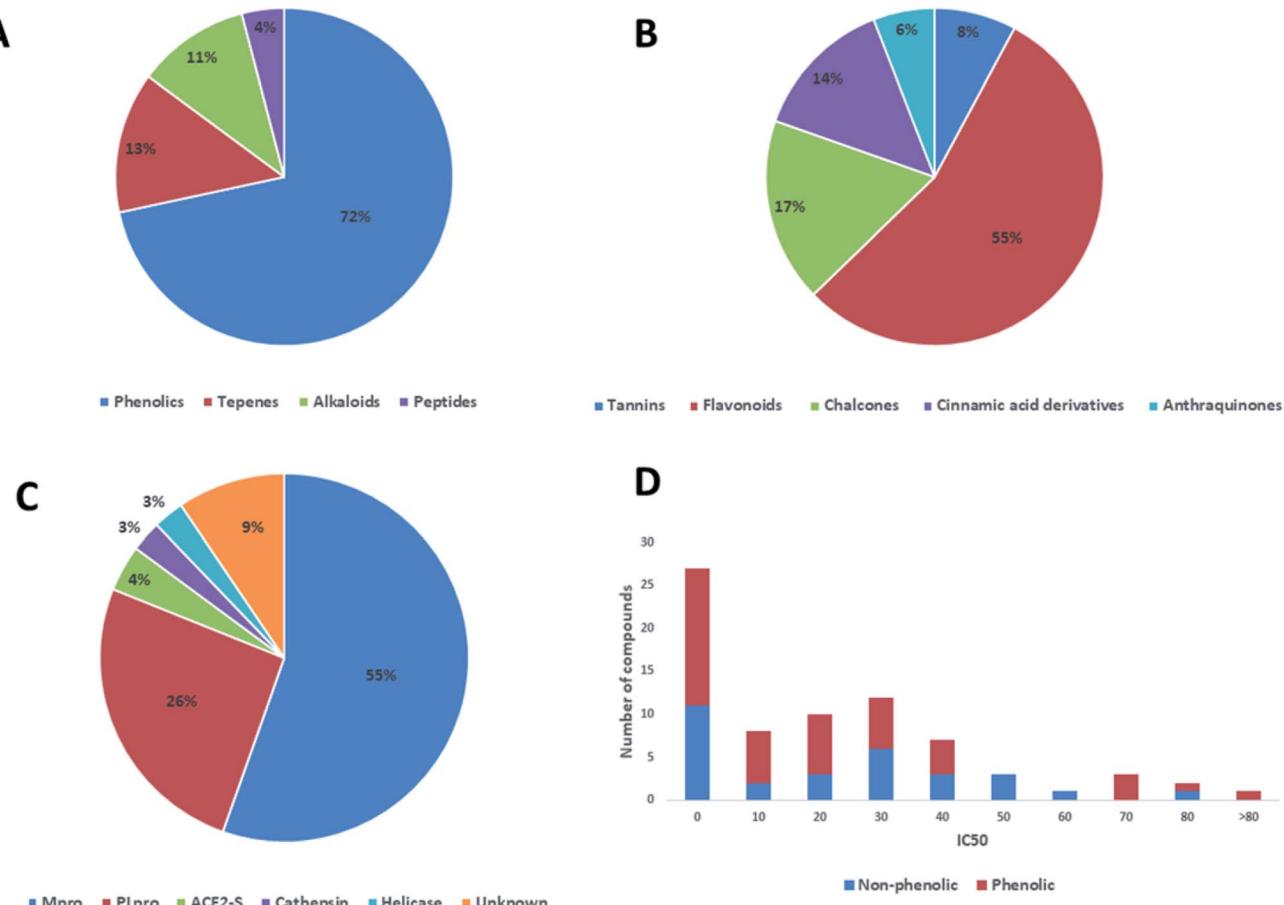


Fig. 5 Pie charts indicating the reported anti-SARS CoV natural therapeutic classes (A & B), and their common viral targets (C). A bar chart depicting activity range of the reported natural products illustrating phenolic compounds as being the most prominent class of compounds (D).

potential anti-SARS CoV therapeutics. In 2004, shortly after the first SARS-CoV spread, a number of polyphenolic metabolites of which tannic acid (8), and theaflavin-3,3'-digallate (9) (Fig. 3) were reported as potent viral M^{pro} inhibitors (IC₅₀ 3 and 9.5 μ M, respectively). These compounds were filtered out from a library of phenolic compounds consisted of 720 compounds using an *in silico* approach called structure-based virtual screening.²⁵

Later on, a number of natural peptides *i.e.* tokaramide A (10) (Fig. 3) derived from the marine organism *Theonella mirabilis*^{26,27} inspired another research group to develop a semi-synthetic peptide derivative *i.e.* TG-0205221 (11) with a nanomolar M^{pro} inhibition activity (IC₅₀ 0.6 μ M), where they exhibited a non-competitive inhibition through the formation of covalent bonds with their binding sites (Fig. 3).²⁸ Following the screening of natural product databases, another research group reported for hyperoside (quercetin-3- β -galactoside) (12) as being a promising M^{pro} inhibitor with IC₅₀ of 42.79 μ M. They also studied the structure-activity relationship of this flavonoid in comparison with a number of related derivatives and found that a hydroxyl group at C-3' is an essential structural requirement for hyperoside competitive inhibitory action (Fig. 3).²⁹

As part of the ongoing investigation of flavonoids-based antiviral candidates, Ryu and co-workers in 2010, reported the

M^{pro} inhibitory activity (IC₅₀ 20–200 μ M) of some common flavonoids *i.e.* quercetin, apigenin and luteolin (13) and bioflavonoids such as amentoflavone (14), with the latter possessing better *in vitro* inhibitory activity (IC₅₀ 8.3 μ M). These findings revealed that larger polyhydroxylated compounds are more preferred to develop potent M^{pro} noncompetitive inhibitors (Fig. 3).³⁰ Such interesting activity together with the simplicity of flavonoids-based inhibitors encouraged another researcher group to develop potent inhibitors analogs such as compound (15) based on the flavonoids scaffold which were subjected for a preclinical trial (Fig. 3).³¹ In 2013, eight phlorotannins, isolated from an edible brown algae *Ecklonia cava*, namely eckol (16), dieckol, triphloretol A, dioxinodehydroeckol, 2-phloroekkol, 7-phloroekkol, fucodiphloroethol G and phlorofucuroeckol A, with the first two compounds (Fig. 3) possessing the most potent SARS-CoV M^{pro} inhibitory activities (IC₅₀ 8.8 and 2.7 μ M, respectively).³² Epigallocatechin gallate and gallicatechin gallate showed good M^{pro} inhibition activity (IC₅₀ 73 and 47 μ M, respectively) with their galloyl moiety at 3-OH position required for the antiviral action.³³

Herbacetin, rhoifolin and pectolinarin showed anti-SARS-CoV M^{pro} action (IC₅₀ 33.17, 27.45 and 37.78 μ M, respectively) using a tryptophan-based fluorescence method.³⁴



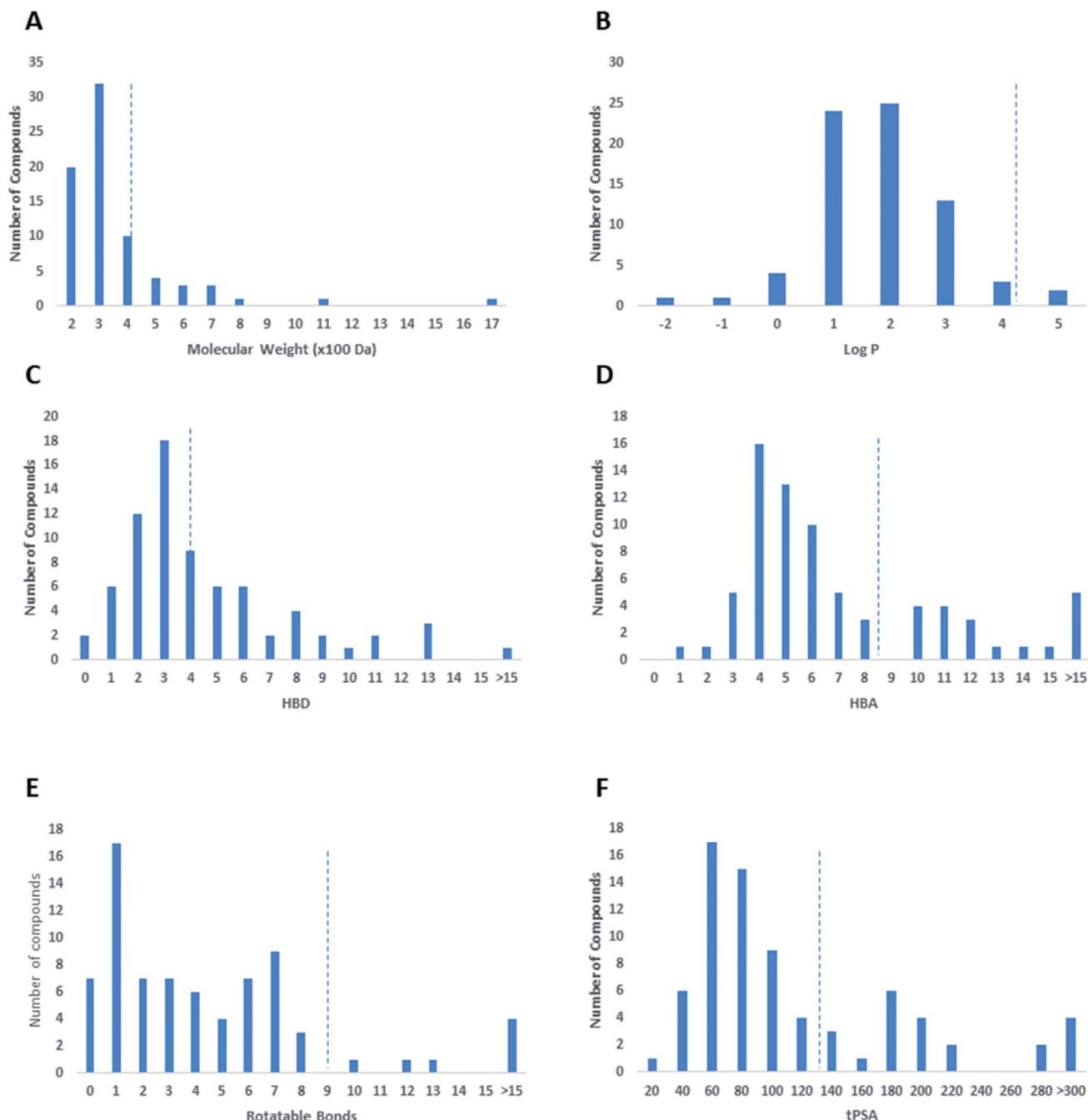


Fig. 6 Analysis of the physicochemical properties of anti-SARS CoV natural products displayed in terms of molecular weight (A), calculated $\log P$ (B), HBD (C), HBA (D), number of rotatable bonds (E), and tPSA (F).

Scutellarein, myricetin, quercetagetin, and robinetin were reported in a Chinese patent to inhibit SARS M^{pro} activity in a fluorescent assay.² The former two compounds also were proved to inhibit the SARS-CoV helicase protein *in vitro* by affecting the ATPase activity.³⁵ High throughput screening of natural product libraries led to the discovery of two interesting M^{pro} competitive inhibitors *i.e.* betulinic acid (17) and savinin (18) (IC_{50} 8.2 and 9.1 μ M, respectively). Both compounds were able to bind the enzyme's active site constituting S1 and S2 subsites *via* multiple hydrogen bonds with Asn 142, Gln 189 and

Cys 145 amino acid residues. Thus, they could be promising starting points for developing more potent derivatives, particularly, betulinic acid (17) (Fig. 3) which is abundant in many edible and medicinal plants.³⁶

Four quinone-methide triterpenes were reported to exhibit M^{pro} inhibitory actions namely celastrol, pristimerin, tingenone and iguesterin (IC_{50} 10.3, 5.5, 9.9, and 2.6 μ M, respectively), which were isolated from the bark of *Tripterygium regelii* (F. Celastraceae), a woody vine used in traditional Chinese

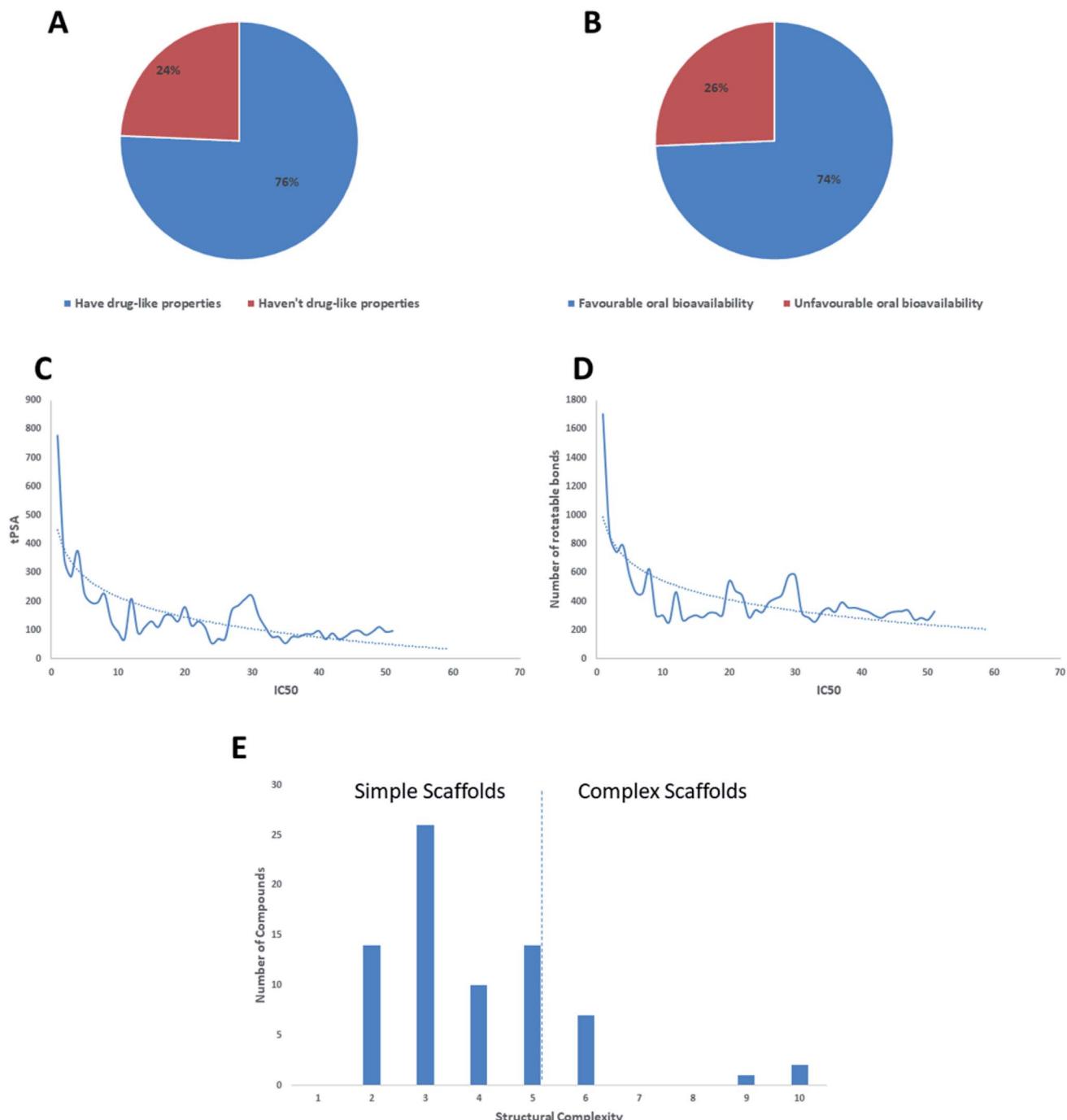


Fig. 7 The compliance of the tested natural products with drug-likeness rules Lipinski's and Veber's rules (A & B, respectively). The relationship between the IC₅₀ and both tPSA and the number of rotatable bond (C & D, respectively). Distribution of the reported anti-SARS CoV natural products depending on their scaffold's complexity (E).

medicine in the treatment of inflammatory and autoimmune diseases.³⁷

A coumarin compound named esculetin-4-carboxylic acid ethyl ester (**19**) (Fig. 3), isolated from tropical marine sponge *Axinella corrugata*, was reported to possess M^{pro} inhibitory action with IC₅₀ value of 46 μ M.³⁸ Isolinoleic acid (at a concentration of 50 μ M) possessed M^{pro} inhibitory potential as reported in another Chinese patent.²

3.2.2. PL^{pro} inhibitors. *In vitro* activity against CoV PL^{pro} was reported by a number of natural products such as geranyl-ated flavonoids (Fig. 4), *i.e.* tomentin A (**20**), tomentin B, tomentin C, tomentin D, tomentin E, 3'-O-methyldiplacol, 4'-O-methyldiplacol, 3'-O-methyldiplacone, 4'-O-methyldiplacone, mimulone, diplacone, and 6-geranyl-4',5,7-trihydroxy-3',5'-dimethoxyflavanone, isolated from *Paulownia tomentosa* fruits.³⁹ Tanshinones isolated from *Salvia miltiorrhiza*,⁴⁰ and



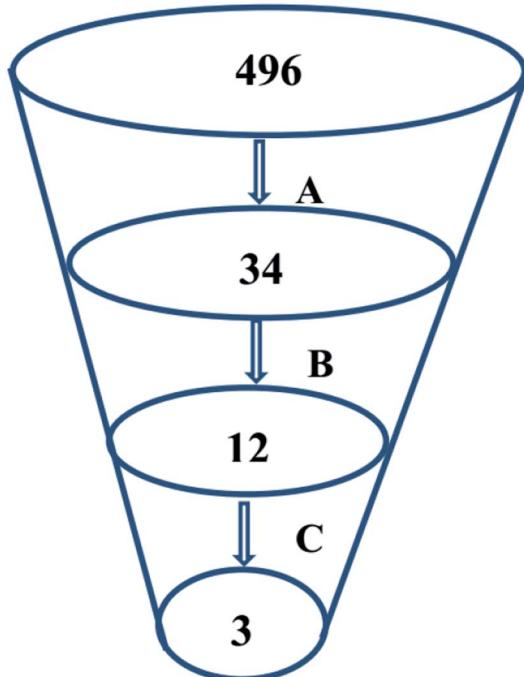


Fig. 8 An outline of the employed virtual screening methodology that comprised of (A) phenolic compounds library virtual screening via molecular docking, (B) selection of the docking poses maintaining the essential binding interactions and (C) selection of top phenolic hits with the most acceptable drug-likeness properties.

diarylheptanoid hirsuteneone derived from the stem bark of *Alnus japonica*.⁴¹ Park and co-workers identified a number of tanshinone derivatives as being potent PL^{pro} inhibitors. One of them (21) was found to be the most potent noncompetitive inhibitor (IC_{50} 0.8 μ M), that was able to bind to the enzyme's active site covalently. In parallel work, the same research group characterized a diarylheptanoid derivative hirsuteneone (22) with an α,β -unsaturated carbonyl group that was also able to form a covalent bond with a cysteine residue (Cys-145) in the enzyme's active site leading to a significant *in vitro* inhibition (IC_{50} 4.1 μ M). They also reported a novel geranylated flavonoid derivative with a dihydro-2H-pyran moiety with promising inhibitory activity (IC_{50} value of 5 μ M).⁴²

Recently, Park and his co-workers identified several polyhydroxylated chalcone derivatives, *i.e.* isobavachalcone (23), 4-hydroxyderricin, xanthoangelol, xanthoangelol F, xanthoangelol D, xanthoangelol E, xanthoangelol B, xanthoangelol G and xanthoangelol A, and coumarins from *Angelica keiskei*, *i.e.* psoralen, bergapten, xanthotoxin and isopimpinellin (Fig. 4), as being potent competitive inhibitors of both M^{pro} and PL^{pro} with IC_{50} values ranging from 5.8 to 11.9 μ M. Xanthoangelol E, containing the perhydroxyl group, exhibited the most potent M^{pro} and PL^{pro} inhibitory activity with IC_{50} values of 11.4 and 1.2 μ M, respectively.⁴³ Another group of polyphenolic compounds, were isolated from the medicinal plant *Broussonetia papyrifera*, of which a chalconoid derivative (24) showed the best inhibitory potential against both M^{pro} and PL^{pro} (IC_{50} 27.9 and 112.9 μ M, respectively).⁴⁴ Trypanthrin (25) isolated

from *Strobilanthes cusia* leaf inhibited the cleavage activity of PL^{pro} and the post-entry stage of viral replication with IC_{50} of 1.52 μ M.⁴⁵ Six cinnamic amides *i.e.* *N*-trans-caffeooyltyramine, *N*-trans-coumarooyltyramine, *N*-trans-feruloyltyramine, terrestriamide and *N*-trans-feruloyloctopamine, isolated from *Tribulus terrestris* fruits, showed significant PL^{pro} inhibition (IC_{50} values ranging from 15.8 to 70.1 μ M). The latter compound possessed the most potent inhibition with IC_{50} of 15.8 μ M.⁴⁶ Six phenolic compounds with good PL^{pro} inhibitory actions (IC_{50} ranging between 4.2 and 38.4 μ M), namely bavachinin, neobavaisoflavone, 4'-O-methylbavachalcone, corylifol A, isobavachalcone and psoralidin (26) were isolated from *Psoralea corylifolia* seeds. The latter two compounds showed the most potent actions with IC_{50} values of 7.3 and 4.2 μ M, respectively.⁴⁷ The inhibitory potential of ten polyphenols derived from *Broussonetia papyrifera* roots, *i.e.* broussochalcone A, broussochalcone B, 4-hydroxyisolonchocarpin, papyriflavonol A (27), 3'-(3-methylbut-2-enyl)-3',4,7-trihydroxyflavane, kazinol A, kazinol B, broussoflavan A, kazinol F and kazinol J, were tested against the two SARS CoV proteases with a more potent inhibition recorded against PL^{pro} than that of $3CL^{pro}$. The most potent PL^{pro} inhibition was exhibited by the prenylated flavone derivative *viz.* papyriflavonol A (27) with IC_{50} value of 3.7 μ M, exceeding the inhibitory potential of non-prenylated flavone derivatives *viz.* quercetin and kaempferol (IC_{50} 8.6 and 16.3 μ M, respectively). This signified the crucial role of the prenyl group in forming stronger hydrophobic interactions with the enzyme as well as the increase in the hydroxylation in the flavone backbone.⁴⁸

3.3. Viral release inhibitors

Several kaempferol derivatives (Fig. 4) were reported to block the 3a channel of Coronavirus, which are formed by the ORF 3a-coded proteins, thus counteracting the viral production and release from the host cells. This gives the body a chance to adjust its immune system to counteract the viral attack. Kaempferol glycosides were proved to possess more potent inhibitory effect than kaempferol and this pointed out to the significance of sugar residues for the antiviral activity. Juglanin (kaempferol-3-*O*- α -L-arabinofuranoside) (28) was the most effective glycoside with an IC_{50} value of 2.3 μ M. Other kaempferol glycosides *i.e.* tiliroside (kaempferol-3-*O*-(6-*p*-coumaroyl)-glucoside) (29) and afzelin (kaempferol-3-*O*- α -L-rhamnoside) (30) were less potent than juglanin (28) but showed similar activity to that of emodin (2).⁴⁹

4. Analysis of drug-likeness properties

The druggability of any molecule is determined depending on its pharmacokinetic properties as well as its excellent pharmacodynamics. Calculating the physicochemical properties (*e.g.* $\log P$ and molecular weight) of a bioactive compound can predict its possible pharmacokinetics (*e.g.* absorption and bioavailability). Lipinski's rule of five considers a given bioactive compound as a drug candidate if it possess the following



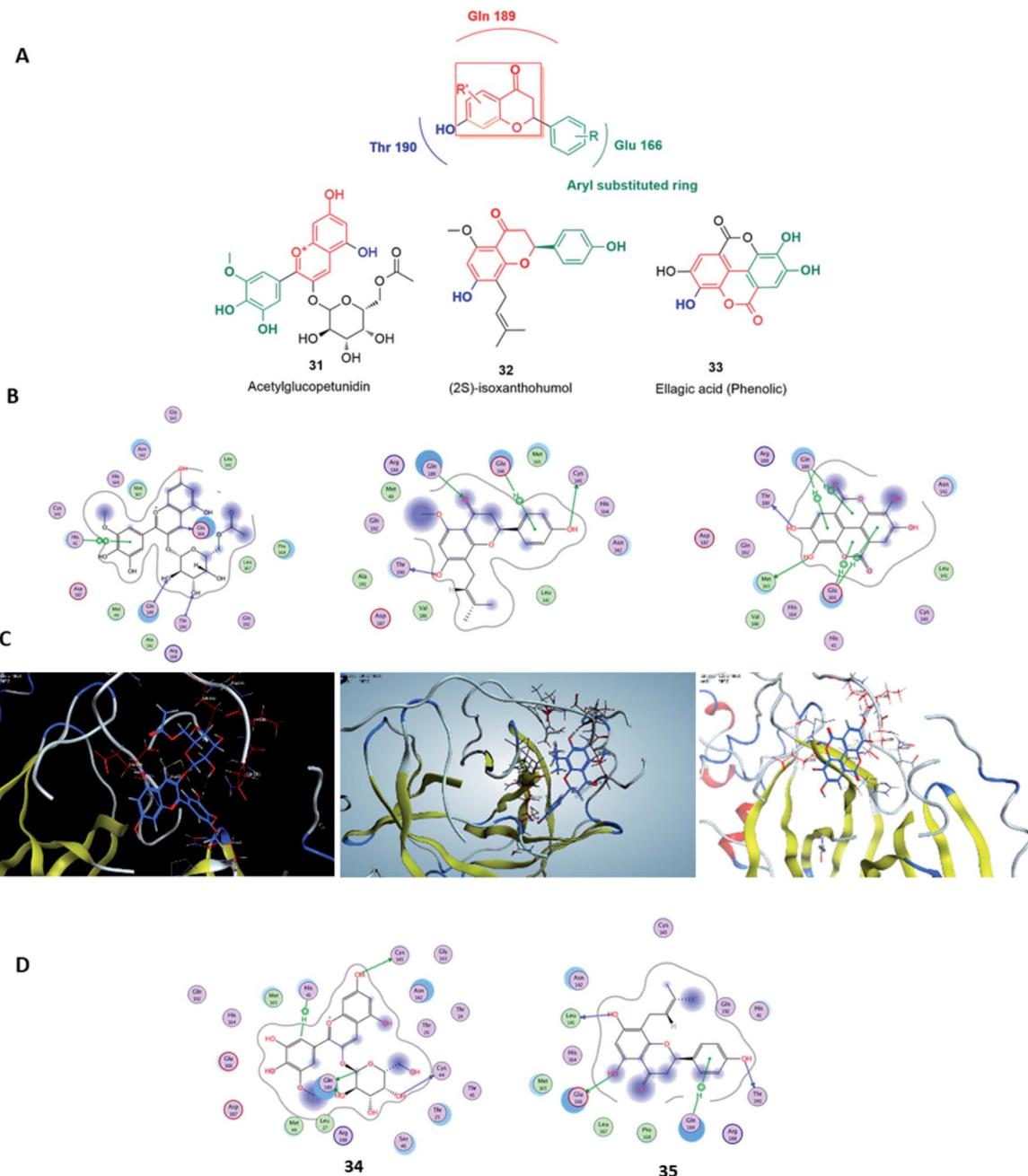


Fig. 9 The main pharmacophore of the top three hits (A), along with their binding modes inside SARS CoV M^{Pro} active sites (B & C). The binding modes of the hydrolysis metabolites derived from compounds 31 and 32 (D).

physicochemical parameters *i.e.* $\log P \leq 5$, molecular weight ≤ 500 , hydrogen bond donor [HBD] ≤ 5 , and hydrogen bond acceptor [HBA] ≤ 10 . Despite, 90% of orally active drugs that have reached phase 2 clinical status⁵⁰ follow these parameters range, however, meeting the rule of five does not guarantee that a molecule is a drug-like.⁵¹ Drug's cellular permeability and in turn its distribution and excretion are associated with its topological polar surface area (tPSA) and molecular flexibility (*i.e.* the number of rotatable bonds). Hence, bioactive molecules with tPSA of 140 Å or less and rotatable bonds of ten or fewer

(*i.e.* Veber's oral bioavailability) can be considered as potential drug candidates.⁵¹

In this review, the 74 anti-CoV natural products (Table S1, ESI†) were found to belong to four classes of compounds *viz.* phenolics, terpenoids, alkaloids and peptides, where phenolics were the most prominent compounds of which flavonoids were the major class (72%) (Fig. 5). The majority of compounds were found targeting M^{Pro} and PL^{Pro} (55% and 26%, respectively, Fig. 5) indicating that they are the most susceptible targets towards natural products, particularly the phenolics. The anti-viral activity of these compounds possessed activity ranging

from potent to moderate (IC_{50} ranging from 0.0025 to 90 μM , average IC_{50} 20.13 μM), where phenolics were also the most active ones (Fig. 5). The physicochemical characters, *viz.* molecular weight, $\log P$, HBD, HBA, tPSA and rotatable bonds, of the 74 anti-SARS CoV were calculated using LigandScout 4.3 (Vienna, Austria) and projected onto a drug-likeness cut-off.⁵²

Regarding the drug-likeness properties, 76% of the reviewed molecules followed Lipinski's and Veber's rules (Fig. 7), indicating the promising potential of natural products to serve as drug candidates against the newly emerged anti-SARS CoV. The molecular weights of the 74 compounds distributed between 248.24 and 1701.2 Da (average of 138.248 Da), and most of them were between 200 and 400 Da (Fig. 6). Both tannic acid and β -aescin were the largest molecules. However, they showed potent anti-SARS CoV (IC_{50} 3 and 6 μM , respectively). The $\log P$ of the compounds under study followed a normal distribution with the major was around 2, and 79% of the compounds were of favourable $\log P$ values (≤ 5) (Fig. 6). Regarding HBD and HBA parameters (average of 3 and 4, respectively, Fig. 6), they showed a distribution pattern similar to that of $\log P$ (Fig. 6) with more than 80% of compounds were of acceptable numbers. Additionally, 74% of compounds can demonstrate good oral bioavailability (Fig. 6). Being the most predominant and active compounds, phenolic's tPSA and to some extent of flexibility (number of rotatable bonds) which are interestingly linked to their activity. The compounds with higher tPSA and number of rotatable bonds (*i.e.* flexible) were found to exhibit higher IC_{50} values (Fig. 7). Such observation was also found in the peptides class, as they were among the most active compounds (IC_{50} 0.0025 to 6.8 μM), and also have high tPSA ($>140 \text{ \AA}$) and high flexibility (rotatable bonds 12 to 27). Natural products are not always available to supply the drug industry due to their limited natural occurrence for some ecological considerations. Accordingly, combinatorial chemistry can aid in providing an alternative source for the natural product itself or its analogs. In this context, we examined the complexity and the synthetic accessibility of our reviewed natural products. As illustrated in Fig. 7, most of the compounds ($>85\%$) got a score ranging from 2 to 5, indicating that they or their analogs could be synthesized if the natural product itself cannot be readily accessible.

5. *In silico* analysis of potential anti SARS-CoV 2 agents

Virtual screening (VS) proved to be a helpful tool for drug hit/lead identification. Many studies have recently employed *in silico* exploration of large sets of natural products with the aid of online databases in attempt to discover new hits against potential molecular targets. Among which, a study by ul Qamar M. T. reported for structural insights into anti-viral phytochemicals/traditional Chinese medicinal compounds with *in silico* screening of 32 297 natural products for their SARS-CoV-2 3CL^{pro} inhibitory action.⁵³ As shown in our *in silico* analysis, phenolic derivatives were the most prominent and active compounds against previous CoVs (Fig. 5). Thus, we subjected a library of phenolic metabolites,⁵⁴ to a computer-

aided virtual screening against the active site of the recently reported COVID-19 M^{pro} (PDB code: 5R7Z) based on high-throughput flexible docking technique for the sake of finding possible candidate acting as inhibitors for the newly emerged SARS-CoV2. Thirty four compounds were ranked by highest energy score and the poses that maintained the essential interactions. Twelve compounds were retrieved as the top-hits that fit into the M^{pro} target pocket (Table S2, ESI†). Only three of these top candidates obey both Lipinski's and Veber's rules of drug-likeness (Fig. 8 and 9). The structure–activity relationship analyses of the final top-hits presented in Fig. 9(A) revealed that they shared convergent interactions with the active site's amino acid residues that are comparable with the previous CoV-main proteases inhibitors. All top hits exhibited hydrogen bonding interaction with Thr 190 *via* the hydroxyl phenolic group and the carbonyl oxygen of Thr 190 residue. Additional hydrogen bonding interactions were found to exist between ellagic acid (33) and Met-165 or isoxanthohumol (32) and Cys-145. Regarding van der Waal interactions, all the hits showed interaction with Gln-189, which is assumed to be essential for the activity (Fig. 9). Additionally, the presence of Gln-191, Glu-166 and Cys-145 with some top-hits to form the hydrophobic subsite in other CoV M^{pro} which may create an extra van der Waal interactions. These essential interactions were found to be missing in low-scoring compounds. The selected hits with drug-like properties are well reported to be abundant in many edible fruits such as raspberry and pomegranate. Hence, they are readily available for the *in vitro* investigation to validate them as potential anti-SARS CoV2 drug leads (Fig. 9). Regarding the possibility of the metabolic instability of acetylglucopetunidin (31) and (2S)-isoxanthohumol (32), both compounds show the presence of acetylester or methoxy groups that could be easily hydrolyzed inside the human body. Accordingly, we redocked their respective metabolites (34 & 35) to explore their inhibitory potential. Both metabolites were found to reserve the same hydrophobic and hydrogen bond interactions with His-41 and Gln-189 amino acid residues, respectively however, they gained an additional hydrogen bond interaction with Gln-189. Moreover, both metabolites bind with Cys-44 and Cys-145 *via* two hydrogen bond donors with binding affinity of $-7.147 \text{ kcal mol}^{-1}$. The metabolite of (2S)-isoxanthohumol (35) was found to bind with the same three amino acid residues *i.e.* Gln-189, Thr-190, and Glu-166, with a binding affinity of $-6.021 \text{ kcal mol}^{-1}$, which was comparable to its parent compound (32), however, an extra hydrogen bond donor with Leu 141 was recorded (Fig. 9(D)).

6. Concluding remarks

The rapid identification of effective therapeutic interventions against COVID-19 is a major challenging issue nowadays. Taking into consideration that the development of new synthetic drugs is a time-consuming process, so repurposing the already known medications is the most convenient choice under the current situation. Hence, natural products can provide a valuable source for the rapid drug discovery of effective as well as safe anti-SARS CoV therapeutics. As has been



illustrated in the current review, and other recently published reviews, a wide array of chemically-diverse natural products have proven to be excellent candidates as anti-SARS CoV, among which phenolic derivatives, *viz.* flavonoids were the most reported active agents, due to their promising pharmacokinetic profiles.⁵⁵

Being the most susceptible targets in CoV, both M^{pro} and PL^{pro} of SARS-CoV in addition to a number of host-based proteases *viz.* cathepsins and furin, can be utilized for the sake of developing broad-spectrum anti-SARS CoV therapeutic agents that target multiple viral and non-viral proteins. Some phenolics such as tannins and flavonoids have exhibited a broad-spectrum activity not only against SARS-CoV but also other viruses⁵⁶. Based on our findings, tPSA and molecular flexibility are the most essential descriptors that should be taken into consideration for selecting active and bioavailable phenolic compounds. The reported IC₅₀ values of SARS natural products-inhibitors were mostly recorded in a micromolar range. However, more potent synthetic or semi-synthetic derivatives can be developed from the current scaffolds as they showed low to moderate molecular complexity. With the tremendous ongoing efforts to develop broad-spectrum anti-viral therapeutics to combat coronaviruses and stop the rapid spread of COVID-19 worldwide, we hope that the current outbreak to settle down in a few months, as has been witnessed previously with SARS and MERS pandemics.

Contributors

AMS, AMA and ARK initiated the review. URA, HMH and MER developed the scope of the manuscript. AMS, AMA and ARK did the literature search and prepared the first draft. AMS designed the figures and interpreted the findings. URA, HZ, HMH and MER critically reviewed the data and draft. All authors subsequently modified the manuscript jointly. URA is the guarantor of the final version, which was read and approved by all the authors.

Conflicts of interests

We declare that we have no conflicts of interest.

Acknowledgements

The authors thank Nahda University in Beni Suef (NUB), Minia, Beni Suef Universities and Arab Academy for Science, Technology and Maritime Transport for supporting this work. We thank Jörn Piel (Institute of Microbiology, ETH Zürich, Zürich, Switzerland), Prof. Hossam Eldin Elmalahi (Nahda University, Beni Suef, Egypt) and Prof. Emad Hamdi Ghoz (Deraya University, New Minia, Egypt) for their support and fruitful discussion.

References

- E. De Wit, N. Van Doremalen, D. Falzarano and V. J. Munster, *Nat. Rev. Microbiol.*, 2016, **14**, 523–534.
- V. Kumar, Y. S. Jung and P. H. Liang, *Expert Opin. Ther. Pat.*, 2013, **23**, 1337–1348.
- V. Thiel, K. A. Ivanov, Á. Putics, T. Hertzog, B. Schelle, S. Bayer, B. Weißbrich, E. J. Snijder, H. Rabenau, H. W. Doerr, A. E. Gorbalyena and J. Ziebuhr, *J. Gen. Virol.*, 2003, **84**, 2305–2315.
- K. Anand, J. Ziebuhr, P. Wadhwani, J. R. Mesters and R. Hilgenfeld, *Science*, 2003, **300**, 1763–1767.
- E. J. Snijder, P. J. Bredenbeek, J. C. Dobbe, V. Thiel, J. Ziebuhr, L. L. M. Poon, Y. Guan, M. Rozanov, W. J. M. Spaan and A. E. Gorbalyena, *J. Mol. Biol.*, 2003, **331**, 991–1004.
- S. van Boheemen, M. de Graaf, C. Lauber, T. M. Bestebroer, V. S. Raj, A. M. Zaki, A. D. M. E. Osterhaus, B. L. Haagmans, A. E. Gorbalyena, E. J. Snijder and R. A. M. Fouchier, *mBio*, 2012, **3**, 1–9.
- K. S. Chan, S. T. Lai, C. M. Chu, E. Tsui, C. Y. Tam, M. M. L. Wong, M. W. Tse, T. L. Que, J. S. M. Peiris, J. Sung, V. C. W. Wong and K. Y. Yuen, *Hong Kong Med. J.*, 2003, **9**, 399–406.
- B. Cao, Y. Wang, D. Wen, W. Liu, J. Wang, G. Fan, L. Ruan, B. Song, Y. Cai, M. Wei, X. Li, J. Xia, N. Chen, J. Xiang, T. Yu, T. Bai, X. Xie, L. Zhang, C. Li, Y. Yuan, H. Chen, H. Li, H. Huang, S. Tu, F. Gong, Y. Liu, Y. Wei, C. Dong, F. Zhou, X. Gu, J. Xu, Z. Liu, Y. Zhang, H. Li, L. Shang, K. Wang, K. Li, X. Zhou, X. Dong, Z. Qu, S. Lu, X. Hu, S. Ruan, S. Luo, J. Wu, L. Peng, F. Cheng, L. Pan, J. Zou, C. Jia, J. Wang, X. Liu, S. Wang, X. Wu, Q. Ge, J. He, H. Zhan, F. Qiu, L. Guo, C. Huang, T. Jaki, F. G. Hayden, P. W. Horby, D. Zhang and C. Wang, *N. Engl. J. Med.*, 2020, **382**(19), 1787–1799.
- Y. M. Báez-Santos, S. J. Barraza, M. W. Wilson, M. P. Agius, A. M. Mielech, N. M. Davis, S. C. Baker, S. D. Larsen and A. D. Mesecar, *J. Med. Chem.*, 2014, **57**, 2393–2412.
- K. Ratia, S. Pegan, J. Takayama, K. Sleeman, M. Coughlin, S. Baliji, R. Chaudhuri, W. Fu, B. S. Prabhakar, M. E. Johnson, S. C. Baker, A. K. Ghosh and A. D. Mesecar, *Proc. Natl. Acad. Sci. U. S. A.*, 2008, **105**, 16119–16124.
- C. M. Booth, L. M. Matukas, G. A. Tomlinson, A. R. Rachlis, D. B. Rose, H. A. Dwosh, S. L. Walmsley, T. Mazzulli, M. Avendano, P. Derkach and I. E. Ephtimios, *J. Am. Med. Assoc.*, 2003, **289**, 2801–2809.
- J. A. Tanner, B. J. Zheng, J. Zhou, R. M. Watt, J. Q. Jiang, K. L. Wong, Y. P. Lin, L. Y. Lu, M. L. He, H. F. Kung, A. J. Kesel and J. D. Huang, *Chem. Biol.*, 2005, **12**, 303–311.
- L. Wilson, P. Gage and G. Ewart, *Virology*, 2006, **353**, 294–306.
- W. Li, M. J. Moore, N. Vasllieva, J. Sui, S. K. Wong, M. A. Berne, M. Somasundaran, J. L. Sullivan, K. Luzuriaga, T. C. Greeneugh, H. Choe and M. Farzan, *Nature*, 2003, **426**, 450–454.
- J. Gao, G. Lu, J. Qi, Y. Li, Y. Wu, Y. Deng, H. Geng, H. Li, Q. Wang, H. Xiao, W. Tan, J. Yan and G. F. Gao, *J. Virol.*, 2013, **87**, 13134–13140.
- Z. Qian, S. R. Dominguez and K. V. Holmes, *PLoS One*, 2013, **8**, e76469.



17 K. Shirato, M. Kawase and S. Matsuyama, *J. Virol.*, 2013, **87**, 12552–12561.

18 C. Burkard, M. H. Verheij, B. L. Haagmans, F. J. van Kuppeveld, P. J. M. Rottier, B.-J. Bosch and C. A. M. de Haan, *J. Virol.*, 2015, **89**, 4434–4448.

19 A. K. Ghosh, K. Xi, M. E. Johnson, S. C. Baker and A. D. Mesecar, *Annu. Rep. Med. Chem.*, 2006, **41**, 183–196.

20 L. Yi, Z. Li, K. Yuan, X. Qu, J. Chen, G. Wang, H. Zhang, H. Luo, L. Zhu, P. Jiang, L. Chen, Y. Shen, M. Luo, G. Zuo, J. Hu, D. Duan, Y. Nie, X. Shi, W. Wang, Y. Han, T. Li, Y. Liu, M. Ding, H. Deng and X. Xu, *J. Virol.*, 2004, **78**, 11334–11339.

21 T. Y. Ho, S. L. Wu, J. C. Chen, C. C. Li and C. Y. Hsiang, *Antiviral Res.*, 2007, **74**, 92–101.

22 A. O. Adedeji, W. Severson, C. Jonsson, K. Singh, S. R. Weiss and S. G. Sarafianos, *J. Virol.*, 2013, **87**, 8017–8028.

23 C. W. Lin, F. J. Tsai, C. H. Tsai, C. C. Lai, L. Wan, T. Y. Ho, C. C. Hsieh and P. D. L. Chao, *Antiviral Res.*, 2005, **68**, 36–42.

24 L. Zhou, Y. Liu, W. Zhang, P. Wei, C. Huang, J. Pei, Y. Yuan and L. Lai, *J. Med. Chem.*, 2006, **49**, 3440–3443.

25 C. N. Chen, C. P. C. Lin, K. K. Huang, W. C. Chen, H. P. Hsieh, P. H. Liang and J. T. A. Hsu, *Evid. base Compl. Alternative Med.*, 2005, **2**, 209–215.

26 N. Fusetani, M. Fujita, Y. Nakao, S. Matsunaga and R. W. Van Soest, *Bioorg. Med. Chem. Lett.*, 1999, **9**, 3397–3402.

27 Y. Nakao, M. Fujita, K. Warabi, S. Matsunaga, N. Fusetani and A. Miraziridine, *J. Am. Chem. Soc.*, 2000, **122**, 10462–10463.

28 S. Yang, S. J. Chen, M. F. Hsu, J. D. Wu, C. T. K. Tseng, Y. F. Liu and W. C. Chen, *J. Med. Chem.*, 2006, **49**, 4971–4980.

29 L. Chen, J. Li, C. Luo, H. Liu, W. Xu, G. Chen and H. Jiang, *Bioorg. Med. Chem.*, 2006, **14**, 8295–8306.

30 Y. B. Ryu, H. J. Jeong, J. H. Kim, Y. M. Kim, J. Y. Park, D. Kim and M. C. Rho, *Bioorg. Med. Chem.*, 2010, **18**, 7940–7947.

31 A. O. Adedeji and S. G. Sarafianos, *Curr. Opin. Virol.*, 2014, **8**, 45–53.

32 J. Y. Park, J. H. Kim, J. M. Kwon, H. J. Kwon, H. J. Jeong, Y. M. Kim and Y. B. Ryu, *Bioorg. Med. Chem.*, 2013, **21**, 3730–3737.

33 T. T. H. Nguyen, H. J. Woo, H. K. Kang, V. D. Nguyen, Y. M. Kim, D. W. Kim and D. Kim, *Biotechnol. Lett.*, 2012, **34**, 831–838.

34 Y. Yang, M. S. Islam, J. Wang, Y. Li and X. Chen, *Int. J. Biol. Sci.*, 2020, **16**, 1708.

35 M. S. Yu, J. Lee, J. M. Lee, Y. Kim, Y. W. Chin, J. G. Jee and Y. J. Jeong, *Bioorg. Med. Chem. Lett.*, 2012, **22**, 4049–4054.

36 C. C. Wen, Y. H. Kuo, J. T. Jan, P. H. Liang, S. Y. Wang, H. G. Liu and C. C. Hou, *J. Med. Chem.*, 2007, **50**, 4087–4095.

37 Y. B. Ryu, S. J. Park, Y. M. Kim, J. Y. Lee, W. D. Seo, J. S. Chang and W. S. Lee, *Bioorg. Med. Chem. Lett.*, 2010, **20**, 1873–1876.

38 P. Hamill, D. Hudson, R. Y. Kao, P. Chow, M. Raj, H. Xu and F. Jean, *Biol. Chem.*, 2006, **387**, 1063–1074.

39 J. K. Cho, M. J. Curtis-Long, K. H. Lee, D. W. Kim, H. W. Ryu, H. J. Yuk and K. H. Park, *Bioorg. Med. Chem.*, 2013, **21**, 3051–3057.

40 J. Y. Park, J. H. Kim, Y. M. Kim, H. J. Jeong, D. W. Kim, H. H. Park and Y. B. Ryu, *Bioorg. Med. Chem.*, 2012a, **20**, 5928–5935.

41 J. Y. Park, H. J. Jeong, J. H. Kim, Y. M. Kim, S. J. Park, D. Kim and Y. B. Ryu, *Biol. Pharm. Bull.*, 2012, **35**, 2036–2042.

42 H. Wang, S. Xue, H. Yang and C. Chen, *Virol. Sin.*, 2016, **31**, 24–30.

43 J. Y. Park, J. A. Ko, D. W. Kim, Y. M. Kim, H. J. Kwon, H. J. Jeong and Y. B. Ryu, *J. Enzyme Inhib. Med. Chem.*, 2016, **31**, 23–30.

44 J. Y. Park, H. J. Yuk, H. W. Ryu, S. H. Lim, K. S. Kim, K. H. Park, Y. B. Ryu and W. S. Lee, *J. Enzyme Inhib. Med. Chem.*, 2017, **32**, 504–512.

45 Y. C. Tsai, C. L. Lee, H. R. Yen, Y. S. Chang, Y. P. Lin, S. H. Huang and C. W. Lin, *Biomolecules*, 2020, **10**, 366.

46 Y. H. Song, D. W. Kim, M. J. Curtis-Long, H. J. Yuk, Y. Wang, N. Zhuang and K. H. Park, *Biol. Pharm. Bull.*, 2014, **37**, 1021–1028.

47 D. W. Kim, K. H. Seo, M. J. Curtis-Long, K. Y. Oh, J. W. Oh, J. K. Cho and K. H. Park, *J. Enzyme Inhib. Med. Chem.*, 2014, **29**, 59–63.

48 J. Y. Park, H. J. Yuk, H. W. Ryu, S. H. Lim, K. S. Kim, K. H. Park and W. S. Lee, *J. Enzyme Inhib. Med. Chem.*, 2017, **32**, 504–512.

49 S. Schwarz, D. Sauter, K. Wang, R. Zhang, B. Sun, A. Karioti and W. Schwarz, *Planta Med.*, 2014, **80**, 177–182.

50 C. A. Lipinski, *Drug Discov. Today Technol.*, 2004, **1**, 337–341.

51 D. F. Veber, S. R. Johnson, H. Y. Cheng, B. R. Smith, K. W. Ward and K. D. Kopple, *J. Med. Chem.*, 2002, **45**, 2615–2623.

52 G. Wolber and T. Langer, *J. Chem. Inf. Model.*, 2005, **45**, 160–169.

53 M. T. ul Qamar, S. M. Alqahtani, M. A. Alamri and L. L. Chen, *J. Pharm. Anal.*, 2020, DOI: 10.1016/j.jpha.2020.03.009.

54 V. Neveu, J. Perez-Jiménez, F. Vos, V. Crespy, L. du Chaffaut, L. Mennen and A. Scalbert, *Database*, 2010, **2010**, bap024.

55 M. T. Islam, C. Sarkar, D. M. El-Kersh, S. Jamaddar, S. J. Uddin, J. A. Shilpi and M. S. Mubarak, *Phytother Res.*, 2020, 1–22.

56 O. S. Martins, I. A. Santos, D. M. Oliveira, R. V. Grosche and A. C. G. Jardim, *Viruses*, 2020, **12**, 272.

