High-Throughput Screening and Identification of Potent Broad-Spectrum Inhibitors of Coronaviruses

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ABSTRACT Coronaviruses (CoVs) act as cross-species viruses and have the potential to spread rapidly into new host species and cause epidemic diseases. Despite the severe public health threat of severe acute respiratory syndrome coronavirus and Middle East respiratory syndrome CoV (MERS-CoV), there are currently no drugs available for their treatment; therefore, broad-spectrum inhibitors of emerging and endemic CoVs are urgently needed. To search for effective inhibitory agents, we performed high-throughput screening (HTS) of a 2,000-compound library of approved drugs and pharmacologically active compounds using the established genetically engineered human CoV OC43 (HCoV-OC43) strain expressing Renilla luciferase (rOC43-ns2Del-Rluc) and validated the inhibitors using multiple genetically distinct CoVs in vitro. We screened 56 hits from the HTS data and validated 36 compounds in vitro using wild-type HCoV-OC43. Furthermore, we identified seven compounds (lycorine, emetine, monensin sodium, mycophenolate mofetil, mycophenolic acid, phenazo-pyridine, and pyrvinium pamoate) as broad-spectrum inhibitors according to their strong inhibition of replication by four CoVs in vitro at low-micromolar concentrations. Additionally, we found that emetine blocked MERS-CoV entry according to pseudovirus entry assays and that lycorine protected BALB/c mice against HCoV-OC43-induced lethality by decreasing viral load in the central nervous system. This represents the first demonstration of in vivo real-time bioluminescence imaging to monitor the effect of lycorine on the spread and distribution of HCoV-OC43 in a mouse model. These results offer critical information supporting the development of an effective therapeutic strategy against CoV infection.

IMPORTANCE Currently, there is no approved therapy to treat coronavirus infection; therefore, broad-spectrum inhibitors of emerging and endemic CoVs are needed. Based on our high-throughput screening assay using a compound library, we identified seven compounds with broad-spectrum efficacy against the replication of four CoVs in vitro. Additionally, one compound (lycorine) was found to protect BALB/c mice against HCoV-OC43-induced lethality by decreasing viral load in the central nervous system. This inhibitor might offer promising therapeutic possibilities for combatting novel CoV infections in the future.

KEYWORDS Coronaviruses, bioluminescence imaging, broad-spectrum, high-throughput screening, inhibitor, mice

Emerging viruses are difficult to control, because they periodically cycle in and out of humans and livestock; therefore, effective vaccines and antivirals are urgently needed. Coronaviruses (CoVs) represent a group of enveloped, positive-sense, single-stranded viruses with large genomes (27 to 33 kb) and capable of causing respiratory, enteric, hepatic, and neurological diseases of differing severities in diverse animal
species, including humans. All CoVs have a similar genome organization: approximately two-thirds of the 5’-proximal genome contains the ORF1a/b replicase gene, and the remainder encodes the spike, envelope, membrane, and nucleocapsid structural proteins along with several accessory proteins. CoVs belong to the family Coronaviridae in the order Nidovirales (1) and are divided into four genera: alpha-, beta-, gamma-, and delta-CoVs. Only alpha- and beta-CoVs can infect humans, with four CoVs currently known to be prevalent: human CoV 229E (HCoV-229E), HCoV-OC43, HCoV-HKU1, and HCoV-NL63. Severe acute respiratory syndrome CoV (SARS-CoV) and Middle East respiratory syndrome CoV (MERS-CoV) (2, 3) are considered the most emergent CoVs.

CoV infections are difficult to prevent and cure. Although CoV replication machinery exhibits substantial proofreading activity, estimates of the nucleotide mutation rate in CoVs are moderate to high relative to that of other single-stranded RNA viruses. Additionally, the large RNA genome in CoVs allows for extra plasticity in genome modification by recombination (4–6). Moreover, many animal CoVs cause long-term or persistent enzootic infections, which increase the probability of infecting a new host species. SARS-CoV and MERS-CoV are recent examples of newly emergent CoVs that cause severe human diseases (7, 8). Several drugs, such as ribavirin, lopinavir-ritonavir, interferon, and corticosteroids, have been used to treat patients infected with SARS-CoV or MERS-CoV (9–12). However, contradictory findings on their efficacy and concerns over tolerability and clinical benefit have limited the use of antiviral therapeutics for CoVs. Although substantial effort has focused on identifying antivirals for CoV treatment, no approved therapeutic (drug or biological agent) is currently available for the prophylaxis or treatment of CoV-related disease. Treatments for emerging CoV diseases rely upon supportive care and the judicious use of limited quantities of experimental therapeutics (13). Moreover, the lack of effective drugs, the high morbidity and mortality rates caused by the virus, and the potential of epidemic spread highlight the need for new broad-spectrum anti-CoV drugs, especially given the likelihood of infection by novel CoVs (13).

Several recent studies highlighted potential broad-spectrum inhibitors against CoVs (13–17). de Wilde et al. (14) identified numerous potent MERS-CoV inhibitors through screening of a U.S. Food and Drug Administration (FDA)-approved drug library. Interestingly, all of the screened compounds were also capable of inhibiting the replication of SARS-CoV and HCoV-229E. Dyall et al. (15) also screened 27 compounds with activity against both MERS-CoV and SARS-CoV from a 290-compound library; however, the half-maximal effective concentration (EC50) values of most of these drugs were relatively high in vitro but were not assessed in vivo, making their clinical utility questionable. Müller et al. (16) found that silvestrol was a potent and nontoxic inhibitor of cap-dependent viral mRNA translation in CoV-infected human primary cells, with EC50 values of 1.3 nM and 3 nM for MERS-CoV and HCoV-229E, respectively. Notably, Sheahan et al. (17) showed that a nucleotide prodruk (GS-5734) could inhibit SARS-CoV and MERS-CoV replication in multiple in vitro systems at submicromolar half-maximal inhibitory concentration (IC50) values. Furthermore, the prophylactic and early therapeutic administration of GS-5734 significantly reduced the lung viral load and improved clinical signs of disease, as well as respiratory function, in a mouse model of SARS-CoV pathogenesis, further supporting the development of GS-5734 as a broad-spectrum therapeutic to protect against CoVs.

HCoV-OC43, SARS-CoV, and MERS-CoV all belong to beta-CoVs and show a high degree of conservation of essential functional domains, especially within 3CLpro, RdRp, and the RNA helicase, which might represent potential targets for broad-spectrum anti-CoV drugs. We recently reported that a genetically engineered CoV strain (HCoV-OC43) expressing Renilla luciferase (Rluc; rOC43-ns2Del-Rluc) facilitates high-throughput screening (HTS) for broad-spectrum anti-CoV agents and quantitative analysis of CoV replication (18). In the present study, we performed HTS of a 2,000-compound library containing FDA-approved drugs and pharmacologically active compounds and assessed broad-spectrum anti-CoV activity in vitro and in vivo in an experimental infection mouse model. This comprehensive screening and assessment
provided new candidate inhibitors to effectively treat infections by existing CoVs, as well as those by emergent strains in the future.

RESULTS

HTS of anti-HCoV-OC43 compounds. Optimal screening conditions were established using the rOC43-ns2Del-Rluc reporter virus to infect BHK-21 cells in 96-well plates (multiplicity of infection [MOI] = 0.01; 10,000 cells/well). Under this condition, the coefficient of variation and Z factor were 2.9% and 0.86, respectively, demonstrating that the assay was robust and suitable for HTS.

A schematic of the HTS strategy is depicted in Fig. 1A. In the primary screening from the 2,000-compound library under a concentration of 10 μM, 56 hits were found to significantly inhibit rOC43-ns2Del-Rluc replication (Fig. 1B, red and yellow squares), with ≥70% reduced Rluc activity and ≤80% cytotoxicity, including 12 FDA-approved drugs. To obtain more potent inhibitors and exclude the possibility that the observed antiviral activity was specific to rOC43-ns2Del-Rluc, we confirmed the antiviral activity of the 56 hits against wild-type HCoV-OC43 (HCoV-OC43-WT) by quantitative reverse transcription (qRT)-PCR under a lower concentration (5 μM), which confirmed the antiviral activity of 36 compounds (Fig. 1B, yellow squares).

Identification of broad-spectrum anti-CoV inhibitors in vitro. Because only alpha- and beta-CoVs infect humans, we focused on three other CoVs (MERS-CoV [beta-CoV], mouse hepatitis virus A59 [MHV-A59] [beta-CoV], and HCoV-NL63 [alpha-CoV]) to assess the broad-spectrum antiviral activity of the 36 compounds by using an eight-point dose-response confirmation (15). We identified 17 compounds that inhib-
TABLE 1 Properties and antiviral activities of 36 compounds against four CoVs

<table>
<thead>
<tr>
<th>Compound name*</th>
<th>CAS no.</th>
<th>Formula</th>
<th>Bioactivity*</th>
<th>HCoV-OC43 EC50, C50</th>
<th>HCoV-NL63 EC50, C50</th>
<th>MERS-CoV EC50, C50</th>
<th>MHV-A59 EC50, C50</th>
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<tbody>
<tr>
<td>Lycorine</td>
<td>476-28-8</td>
<td>C16H17NO4</td>
<td>Inhibits cell division, antineoplastic, antiviral</td>
<td>0.15, 4.37</td>
<td>0.47, 3.81</td>
<td>1.63, 3.14</td>
<td>0.31, 3.51</td>
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<tr>
<td>Emetine</td>
<td>483-18-1</td>
<td>C20H26ClN4O4</td>
<td>Inhibits RNA, DNA, and protein synthesis</td>
<td>0.30, 2.69</td>
<td>1.43, 3.63</td>
<td>0.34, 3.08</td>
<td>0.12, 3.51</td>
</tr>
<tr>
<td>Mycophenate moefitil</td>
<td>115007-34-6</td>
<td>C23H26N2O4</td>
<td>Immune suppressant, antineoplastic, antiviral</td>
<td>1.58, 3.43</td>
<td>0.23, 3.01</td>
<td>1.54, 3.17</td>
<td>0.27, 3.33</td>
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<tr>
<td>Phenazopyridine</td>
<td>94-78-0</td>
<td>C10H12N4</td>
<td>Analgesic</td>
<td>1.90, &gt;20</td>
<td>2.02, &gt;20</td>
<td>1.93, &gt;20</td>
<td>0.77, &gt;20</td>
</tr>
<tr>
<td>Mycophenic acid</td>
<td>24280-93-1</td>
<td>C16H18O4</td>
<td>Immune suppressant, antineoplastic, antiviral</td>
<td>1.95, 3.55</td>
<td>0.18, 3.44</td>
<td>1.95, 3.21</td>
<td>0.17, 4.18</td>
</tr>
<tr>
<td>Pyrvinium pamoate</td>
<td>3546-41-6</td>
<td>C24H24N2O4</td>
<td>Analgesic</td>
<td>0.32,</td>
<td>2.34,</td>
<td>6.05, 4.24</td>
<td>1.98, 14.5</td>
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<tr>
<td>Menonisin sodium</td>
<td>22373-78-0</td>
<td>C24H28N2O10</td>
<td>Antibacterial</td>
<td>0.45, 3.12</td>
<td>2.64, 3.24</td>
<td>2.56, 2.96</td>
<td>5.21, 3.19</td>
</tr>
<tr>
<td>Cetylpyridinium chloride</td>
<td>6004-24-6</td>
<td>C26H22ClN2</td>
<td>Protein synthesis inhibitor</td>
<td>4.31, 8.23</td>
<td>1.24, 8.52</td>
<td>0.69, 8.14</td>
<td>7.86, 8.19</td>
</tr>
<tr>
<td>Oligomycin</td>
<td>1404-19-9</td>
<td>C10H12N4</td>
<td>Antimicrobial, antifungal</td>
<td>0.19, 6.56</td>
<td>2.63, 4.26</td>
<td>0.21, 5.16</td>
<td>6.43, 6.78</td>
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<tr>
<td>Promazine</td>
<td>58-40-2</td>
<td>C21H20N4S</td>
<td>Antiinfective</td>
<td>0.41, 20</td>
<td>1.37, 20</td>
<td>13.72, 20</td>
<td>0.51, &gt;20</td>
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<tr>
<td>Diperoxon</td>
<td>537-12-2</td>
<td>C22H24N2O4</td>
<td>Antimicrobial, anesthetic</td>
<td>1.71, 14.3</td>
<td>4.91, 13.6</td>
<td>8.77, 14.2</td>
<td>1.98, 14.5</td>
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<tr>
<td>Valinomycin</td>
<td>2001-95-8</td>
<td>C36H30N2O18</td>
<td>Antimicrobial</td>
<td>0.29,</td>
<td>1.97,</td>
<td>5.16, 3.84</td>
<td>5.16, 2.45</td>
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<tr>
<td>Phenylmercuric acetate</td>
<td>62-38-4</td>
<td>C13H16O2</td>
<td>Antifungal</td>
<td>2.17, 5.35</td>
<td>6.79, 5.47</td>
<td>4.64, 5.39</td>
<td>6.81, 5.97</td>
</tr>
<tr>
<td>Hycanthone</td>
<td>3105-97-3</td>
<td>C22H26N6O2</td>
<td>Antimicrobial, hepatotoxic</td>
<td>0.16, 3.58</td>
<td>5.76, 3.68</td>
<td>5.11, 4.32</td>
<td>5.78, 4.19</td>
</tr>
<tr>
<td>Ticlopidine</td>
<td>53885-35-1</td>
<td>C17H10N2O8</td>
<td>PAH inhibitor</td>
<td>1.41, &gt;20</td>
<td>15.65, &gt;20</td>
<td>11.25, &gt;20</td>
<td>14.28, &gt;20</td>
</tr>
<tr>
<td>Papaverine</td>
<td>61-25-6</td>
<td>C21H20N2O4</td>
<td>Muscle relaxant, cerebral vasodilator</td>
<td>1.61, 12.11</td>
<td>7.32, 11.71</td>
<td>9.45, 11.98</td>
<td>11.64, 12.44</td>
</tr>
<tr>
<td>Propanolol</td>
<td>318-98-9</td>
<td>C14H18N2O4</td>
<td>Antitumor, antitumor, antineoplastic</td>
<td>0.48, &gt;20</td>
<td>8.11, &gt;20</td>
<td>11.01, &gt;20</td>
<td>13.54, &gt;20</td>
</tr>
<tr>
<td>Tilorone</td>
<td>27591-69-1</td>
<td>C20H25N3O2</td>
<td>Antiviral</td>
<td>0.32, &gt;20</td>
<td>6.89, &gt;20</td>
<td>10.56, &gt;20</td>
<td>16.11, &gt;20</td>
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<tr>
<td>Antimycin A</td>
<td>1397-94-0</td>
<td>C20H25N3O2</td>
<td>Antifungal, antitumor, interferes in cytochrome oxidation</td>
<td>1.65, 3.62</td>
<td>6.05, 4.21</td>
<td>6.89, 4.32</td>
<td>5.42, 3.98</td>
</tr>
<tr>
<td>Salinomycin sodium</td>
<td>53003-10-4</td>
<td>C20H24NaO11</td>
<td>Antibacterial</td>
<td>0.29, 1.97</td>
<td>5.71, 2.41</td>
<td>5.49, 3.84</td>
<td>5.16, 2.45</td>
</tr>
<tr>
<td>Exalamide</td>
<td>53370-90-4</td>
<td>C18H21N2O2</td>
<td>Antifungal</td>
<td>1.48, &gt;20</td>
<td>17.49, &gt;20</td>
<td>15.91, &gt;20</td>
<td>16.39, &gt;20</td>
</tr>
</tbody>
</table>

*The compounds in bold inhibited the replication of all CoVs with EC50 values of <5 μM.

†CNS, central nervous system; PAH, platelet-activating factor.

**The replication of the CoVs was done at 10 C50 values with EC50 values of <5 μM.

**The compounds indicated in Table 1 were used to test the replication of four CoVs in vitro.

**The seven broad-spectrum inhibitors suppressed the replication of all CoVs in a dose-dependent manner and with low EC50 values of <5 μM.
FIG 2 Dose-response curves for seven broad-spectrum inhibitors of four types of CoVs in vitro. BHK-21, Vero E6, LLC-MK2, or DBT cells were infected with HCoV-OC43-WT, MERS-CoV, HCoV-NL63, or MHV-A59 at an MOI of 0.01, respectively, and treated for 72 h with eight doses (0.1, 0.25, 0.5, 1, 2, 5, 10, and 20 μM) of each inhibitor. The inhibition of viral replication was measured by qRT-PCR. The EC\textsubscript{50} and CC\textsubscript{50} values were calculated using the GraphPad Prism software. The insets show the SI (EC\textsubscript{50} / CC\textsubscript{50}) for each dose. (Continued on next page)
anti-HCoV-OC43 activity (Fig. 2A). Emetine is an active principle of ipecac and inhibits the replication of both DNA and RNA viruses. Additionally, emetine displayed potent anti-CoV activity and the strongest anti-MERS-CoV activity among the top seven inhibitors, with an EC<sub>50</sub> value of 0.34 μM and an SI of 9.06 (Fig. 2B). Mycophenolic acid (an immunosuppressant) exerted a significant inhibitory effect on HCoV-OC43 replication, with an EC<sub>50</sub> of 1.95 μM, and showed stronger anti-HCoV-NL63 activity than the others (EC<sub>50</sub> of 0.18 μM and SI of 19.11) (Fig. 2E). Mycophenolate mofetil, a derivative of mycophenolic acid, showed an antiviral effect on the four CoVs similar to that of mycophenolic acid, suggesting that the two drugs might harbor similar core structures and antiviral mechanisms (Fig. 2C and E). Moreover, phenazopyridine, a widely used urinary analgesic, also displayed strong broad-spectrum anti-CoV activity for the first time, especially against MHV-A59 (EC<sub>50</sub> of 0.77 μM and SI of >25.97) (Fig. 2D). Pyrvinium pamoate is an FDA-approved anthelminthic drug and a potent inhibitor of WNT signaling, suggested to occur through direct activation of protein kinase CK1ε (24). Pyrvinium pamoate inhibited the replication of all CoVs and displayed low toxicity (50% cytotoxic concentration [CC<sub>50</sub>], >19 μM) (Fig. 2F). Finally, monensin sodium, previously shown to inhibit the formation of gamma-CoV infectious bronchitis virus (IBV), inhibited all CoVs at low EC<sub>50</sub> values and displayed low toxicity (Fig. 2G). Although the specific antiviral mechanisms of these seven inhibitors against CoVs are unknown, they showed potential as new antivirals for the treatment of infections caused by a range of CoVs.

Validation of anti-CoV activity. We verified the antiviral activity of the seven inhibitors against HCoV-OC43 by indirect immunofluorescence assay (IFA) and Western blot analysis. As shown in Fig. 3A, all seven inhibitors significantly suppressed HCoV-OC43 replication, compared with that of the control (dimethyl sulfoxide [DMSO]), and with a >90% inhibitory effect for most compounds, except for monensin sodium. Additionally, we observed inhibitory activity when the cells were treated with the inhibitors after viral infection, resulting in significantly reduced levels of HCoV-OC43 nucleocapsid protein (Fig. 3B). Although the inhibitory effect of mycophenolic acid differed according to IFA and Western blot results, the seven inhibitors clearly suppressed HCoV-OC43 replication.

Emetine inhibited MERS-CoV entry. Viral entry is an essential step of the viral life cycle and is thus an attractive target for therapy. Inhibition of this step can block viral propagation at an early stage of infection, thereby minimizing the chance for the virus to evolve and acquire drug resistance. Therefore, we tested the effect of the seven screened inhibitors on CoV entry using a pseudotype virus with a human immunodeficiency virus type 1 (HIV-1) backbone but expressing the spike protein of MERS-CoV in order to generate dose-response curves. Measurement of the inhibition percentage showed that only emetine was an entry inhibitor that blocked MERS-CoV S-mediated infection, with luciferase activity reduced 50-fold compared with that of the control and an EC<sub>50</sub> value of 0.16 μM (Fig. 4).

Antiviral activity of lycorine against lethal HCoV-OC43 infection in vivo. HCoV-OC43 infects neurons and causes encephalitis in mice, with this model previously used for anti-CoV drug evaluation (25). Moreover, this model was convenient based on the lack of need for three biological facilities, in contrast to experiments involving SARS-CoV or MERS-CoV. Therefore, we used this model to evaluate the in vivo antiviral activity of the seven inhibitors. Intracerebral or intranasal inoculation of HCoV-OC43 results in acute-onset severe neurological illness and causes death, with high levels of viral replication in the brain (titer, >10<sup>5</sup> 50% tissue culture infective doses [TCID<sub>50</sub>/ml) at 3

FIG 2 Legend (Continued)
2, 5, 10, or 20 μM of lycorine (A), emetine (B), mycophenolate mofetil (C), phenazopyridine (D), mycophenolic acid (E), pyrvinium pamoate (F), or monensin sodium (G). At 72 h postinfection, the cell culture supernatants were subjected to a viral load assay, and cell lysates were assessed for cytotoxicity. Percent inhibition was calculated as follows: inhibition of viral load (%) = 100% – [viral load (titers or copies) of each CoV in the compound-treated cells/viral load of DMSO-treated control cells]. Inhibition is shown in red, and cytotoxicity is shown in blue. EC<sub>50</sub> values and SI (CC<sub>50/EC<sub>50</sub></sub>) are shown. Data represent the mean ± standard deviation (error bars) of the results of three independent experiments.
to 5 days after infection (26–29). Briefly, female BALB/c mice (12 days old) were inoculated via the intranasal route with 100 TCID$_{50}$ of HCoV-OC43-WT and treated with the seven inhibitors for 14 days, and their survival was monitored for up to 20 days. The inhibitor doses and regimens were selected based on acute-toxicity assessments. Emetine was used at 5 mg/kg, and chloroquine was used as the positive control, which showed antiviral activity at 30 mg/kg. All mice in the phosphate-buffered saline (PBS)-DMSO-treated group died within 6 days after HCoV-OC43-WT challenge (Fig. 5A). In contrast, 83.3% of mice in the lycorine-treated group were still alive at 20 days postinoculation ($P < 0.001$), similar to the survival rate of the chloroquine-treated group (Fig. 5A). Additionally, viral loads in the brain and spinal cord were under the limit of detection in the lycorine-treated group (Fig. 5B), and immunohistochemistry (IHC) of mouse brain coronal sections showed that HCoV-OC43 nucleocapsid protein was present only in the PBS-DMSO-treated group, not in the lycorine-treated group (Fig. 5C).

**Lycorine blocked the spread of rOC43-ns2Del-Rluc in the mouse brain.** To more closely monitor the effect of lycorine on the spread and replication of HCoV-OC43 in the mouse central nervous system in real time, we used bioluminescence imaging (BLI) based on its important advantages of an intrinsically low background signal and very high sensitivity for monitoring light emission in vivo. As expected, PBS-DMSO-treated mice showed gradually increasing signal intensity postinoculation, whereas no signal was detected in the brains of lycorine-treated mice (Fig. 6A). These observations were confirmed by the $\sim 2 \times 10^5$-fold-higher Rluc activity in the PBS-DMSO-treated mice

![Image of inhibitors and molecular weights](image-url)
than in the lycorine-treated mice. Therefore, lycorine showed promise as an anti-CoV agent (Fig. 6B).

DISCUSSION
The SARS epidemic in 2003 and the ongoing MERS-CoV outbreak highlight the inadequacy of available treatments for life-threatening zoonotic CoV infections in humans. Indeed, no specific antiviral agent or vaccine is currently available for human or zoonotic CoV infections, despite the extensive research efforts triggered by the SARS outbreak (30–32). Several FDA-approved drugs (ritonavir, lopinavir, nelfinavir, mycophenolic acid, and ribavirin) inhibit the entry and/or replication of MERS-CoV, SARS-CoV, or other human CoVs in multiple cell lines; however, their antiviral efficacy in vivo remains unknown (16, 32–37). To date, few small molecules have demonstrated anti-CoV activity in animal models of CoV infection (17, 38, 39), and most of the available anti-CoV drugs that target structural proteins might not be effective against other CoVs. For example, the human monoclonal antibodies and antiviral peptides that block virus-host cell binding typically have a limited breadth of protection due to the antigenic diversity in the CoV spike glycoprotein (40, 41). It is worth mentioning that combining antiviral peptides with beta interferon (IFN-β) (40) or combination therapy with different humanized or human monoclonal antibodies targeting non-cross-resistant epitopes (42) can enhance antiviral therapeutic effects. There are also several reports of the enhanced therapeutic effects of combinations of other antiviral agents for the treatment of MERS-CoV. These results indicate that the combined use of different antiviral agents might be synergistic in their treatment of CoV infection.

Here, we identified seven potent broad-spectrum anti-CoV agents, among which lycorine was confirmed as showing strong anti-CoV activity in vivo. Our results suggest that FDA-approved drugs can be used for the prophylaxis of severe or lethal CoV...
infections, thereby greatly facilitating the rapid and rational development of anti-CoV agents with desirable pharmacokinetic and biodistribution properties.

The criteria for the inhibition rate associated with the signals in the antiviral HTS assay ranged from ~30% to ~90% according to different references, with more primary hits but fewer efficacy hits screened when the inhibition rate was set to the lower threshold. In our experiment, we focused on screening hits with higher potency and, therefore, so established the inhibition threshold at 70%. Furthermore, we identified several effective hits with a high degree of cytotoxicity under 10 μM in our primary screening, which was why the cytotoxicity value was set to 80%. Finally, we identified 36 compounds exhibiting anti-HCoV-OC43 activity from a library of compounds, two of which (chloroquine and loperamide) were previously reported as exhibiting broad-spectrum anti-CoV activity. Nine compounds (phenazopyridine, lycorine, pyrvinium pamoate, monensin sodium, cetylpyridinium chloride, oligomycin, loperamide, harmine, and conessine) have not previously been demonstrated as exhibiting antiviral activity against MERS-CoV, thereby offering new therapeutic possibilities for this severe CoV. Furthermore, seven compounds (lycorine, emetine, monensin sodium, mycophenolate mofetil, mycophenolic acid, phenazopyridine, and pyrvinium pamoate) showed inhib-
itory activity against multiple genetically distinct CoVs in vitro at low-micromolar concentrations (EC$_{50}$ values ranging from 0.12 to 3.81 μM). Among these, lycorine is an alkaloid isolated from Amaryllidaceae plants and reportedly exhibits anticancer, antiplasmodial, antitrypanosomal, anti-inflammatory, analgesic, and emetic activities (43–45). Lycorine inhibits the replication of poliomyelitis virus, herpes simplex virus 1, Bunyamwera virus, West Nile virus, dengue virus, and SARS-CoV in vitro, although the mechanism remains to be elucidated (46–49). Liu et al. (50) demonstrated the ability of lycorine to reduce the mortality of human enterovirus 71-infected mice by inhibiting viral replication. Furthermore, Guo et al. (51) evaluated a total of 32 lycorine derivatives, demonstrating that 1-acetyllycorine suppressed enterovirus 71 and hepatitis C virus replication in various cells. Moreover, drug resistance analysis revealed that 1-acetyllycorine targeted a phenylalanine (F76) in the viral proteases. Lycorine also exhibits strong activities against influenza A virus H5H1 in vitro and delays the export of nucleoprotein from the nucleus to the cytoplasm during replication (52). However, the potential mechanism of lycorine against CoVs requires further exploration, and its potential selection for drug-resistant strains must be assessed. We found no reports concerning whether its use in combination enhances the antiviral efficacy of lycorine. Additionally, emetine is a drug used mainly as both an antiprotozoal and an emetic (53), with a number of groups recently reporting new antiviral roles for emetine. Emetine is reportedly a potent inhibitor that suppresses the replication of several RNA viruses (dengue virus and HIV) without generating drug-resistant variant viruses (54–56). We noted that emetine inhibits human cytomegalovirus (HCMV) replication by disrupting the HCMV-induced interaction between p53 and the E3-ubiquitin ligase MDM2 (57). Moreover, a recent study revealed emetine as an inhibitory modulator of rabies virus
axonal transport via a mechanism independent of protein synthesis inhibition (58). Furthermore, Yang et al. (59) demonstrated that emetine inhibits Zika virus (ZIKV) replication via inhibition of ZIKV NS5 polymerase activity and disruption of lysosomal function. In the present study, our preliminary result indicated that emetine inhibited MERS-CoV entry. Interestingly, the immunosuppressant mycophenolic acid and its derivative mycophenolate mofetil, which suppress MERS-CoV replication (33), also inhibited HCoV-OC43 replication (EC₅₀ values of 1.95 and 1.58 μM, respectively). However, a previous study showed that MERS-CoV-infected common marmosets treated with mycophenolate mofetil had more severe forms of disease and higher lung viral loads than did untreated animals, and renal transplant recipients on maintenance mycophenolate mofetil therapy reportedly developed severe or fatal MERS (60). Therefore, these two inhibitors are unlikely to be useful against CoV infections. Monensin sodium, which affects IBV and MHV-A59 assembly (61,62), was also identified as an inhibitor of other CoVs (MERS-CoV, HCoV-OC43, and HCoV-NL63) in this study.

Lycorine showed strong antiviral activity against multiple genetically distinct CoVs in vitro and protected mice against lethal HCoV-OC43 infection in vivo. To our knowledge, this is the first report of the use of BLI to evaluate the effects of antiviral agents on CoV replication and dissemination in vivo without the need for sacrificing animals to quantify viral titers and establish the complete pattern of virus dissemination in the central nervous system. However, most of the data and conclusions concerning the antiviral effects of the inhibitors were derived only from one cell line and should be tested in different CoV replication cell lines in order to exclude the influence of host cell factors, especially for MERS-CoV, for which there are a larger number of susceptible cell lines (e.g., Huh 7 or Calu3/2B4 cells). Therefore, further in vitro and in vivo studies are warranted to determine the potential antiviral mechanisms associated with the seven compounds, as well as their clinical efficacy. Additionally, the efficacy of lycorine-interferon combinations should be explored in animal models.

Following SARS and MERS, future emerging CoVs will likely pose a threat to public health. Therefore, identification of broad-spectrum inhibitors of SARS-CoV, MERS-CoV, and future emerging CoVs is a research priority. From this perspective, the potent broad-spectrum inhibitors (lycorine and emetine) identified in this study might be effective against CoV infections either as single agents or in combination.

**MATERIALS AND METHODS**

**Cell lines, viruses, compounds, and antibodies.** BHK-21, Vero-E6, LLC-MK2, DBT, 293FT, DPP4-expressing Huh7.5 cells, and 17Cl-1 cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM; Gibco, Thermo Fisher Scientific, Waltham, MA, USA) supplemented with 10% fetal bovine serum (FBS; Gibco) and incubated at 37°C in an atmosphere containing 5% CO₂.

HCoV-OC43 (GenBank accession number [AY391777.1](https://www.ncbi.nlm.nih.gov/nuccore/AY391777)) expressing the Rluc gene (rOC43-n2Del-Rluc) and derived from an infectious cDNA clone (17) was used for HTS in HBK-21 cells. HCoV-NL63 strain Amsterdam 1 was used to infect monolayers of LLC-MK2 cells at an MOI of 0.01. The MERS-CoV strain EMC was used to infect monolayers of Vero-E6 cells at an MOI of 0.01. The MHV strain A59 was propagated in 17Cl-1 cells, and a plaque assay was performed in monolayers of DBT cells infected with MHV at an MOI of 0.01.

A 2,000-compound library containing FDA-approved drugs and pharmacologically active compounds was purchased from MicroSource Discovery Systems, Inc. (Gaylordsville, CT, USA) (see Table S1 in the supplemental material). The anti-HCoV-OC43 nucleocapsid protein mouse monoclonal antibody was made in-house. Anti-β-actin (13E5) rabbit monoclonal antibody was obtained from Cell Signaling Technology (Danvers, MA, USA). Infrared IRDye 800CW-labeled goat anti-mouse IgG and 680RD-labeled goat anti-rabbit IgG were purchased from LI-COR Biosciences (Lincoln, NE, USA).

**Viral load assays.** MHV titers were quantified by a plaque assay, as described previously (63). Viral genomic RNA from HCoV-OC43, HCoV-NL63, and MERS-CoV was extracted from 50 μl of cell culture supernatants using a QiAamp viral RNA minikit (Qiagen, Valencia, CA, USA) and quantified by real-time RT-PCR, as described previously (64, 65). The primer and probe sequences were as follows: q-OC43-F, 5’-GCT CAG GAA GGT CTG CTC C-3’; q-OC43-R, 5’-TCC TGc ACT AGA GGC TCT GC-3’; q-OC43-probe, 5’-FAM-TAC TTC GGG CAC ATC C-TAMRA (6-carboxytetramethylrhodamine)-3’; q-NL63-F, 5’-AGG ACC TTA AAT TCA GAC AAC GTT CT-3’; q-NL63-R, 5’-GAT TAC GTT TGC GAT TCA GAC G-3’; q-NL63-probe, 5’-FAM-TAA CAG TTT TAG CAC TTC CTG CTA CCC AAA CA-TAMRA-3’; q-MERS-F, 5’-GGC ACT GAG GAC CCA CGT T-3’; q-MERS-R, 5’-TTG CGA CAT ACC CAT AAA AGC A-3’; and q-MERS-probe, 5’-FAM-CCC CAA ATT GCT GAG CTT CCT GCT ACA-TAMRA-3’.
Primary screening assay and secondary confirmation assay. HCoV-OC43 can replicate efficiently in BHK-21 cells, with this cell line commonly used for virus isolation or antiviral screening assays. Therefore, the primary screening assay was conducted using 10 μM each compound in BHK-21 cells. We observed no difference in the average Fluc signal between mock controls (only reporter virus added) and DMSO controls. The average control (mock or DMSO control) signal was ~6.3 × 10^5 luciferase light units, and that for the background (only BHK-21 cells) was ~25 luciferase light units.

Briefly, ~4,000 BHK-21 cells were seeded on 96-well plates in DMEM supplemented with 10% FBS. After overnight incubation in a 5% CO₂ atmosphere at 37°C, each well contained ~10,000 cells. The medium was then replaced with 94 μL DMEM supplemented with 3% FBS, and 1 μL of each compound (diluted in DMSO) was added to the plates at a final concentration of 10 μM (one compound per well) in triplicate. An equal volume of DMSO alone was added to the DMSO control wells (Fig. 1A). After incubation for 60 min, 5 μL of diluted viral suspension was added to each well (MOI = 0.01) for a final total screening volume of 100 μL/well. The plates were incubated at 37°C for 72 h, and luciferase activity was determined using the Renilla-Glo luciferase assay system (Promega, Madison, WI, USA) according to the manufacturer’s instructions. Antiviral activity was calculated as follows: inhibition of Fluc activity (%) = 100% − (relative luminescence units of compound-treated cells/relative luminescence units of DMSO-treated control cells). Cytotoxicity was calculated as follows: inhibition of BHK-21 proliferation (%) = 100% − (cell viability of compound-treated cells/cell viability of DMSO-treated control cells). The EC₅₀ value and the compound-specific toxicity (CC₅₀) were calculated with GraphPad Prism 5 software (GraphPad, Inc., La Jolla, CA, USA) using the nonlinear regression model. The Z factor, an assessment of the quality of screening assays, was determined, and compounds were considered effective if they reduced Fluc activity by ≥70% and cytotoxicity by ≤80%.

A confirmation assay was performed using HCoV-OC43-WT treated with the compounds at two concentrations (10 and 5 μM), and viral RNA load was determined by qRT-PCR. To screen hits with higher potency and narrow the screening scope, a compound was considered effective only if the EC₅₀ value was ≤5 μM.

Western blot analysis. BHK-21 cells were lysed by incubation in 0.5% NP-40 buffer for 30 min at 4°C. Lysates were separated by sodium dodecyl sulfate-polyacrylamide gel electrophoresis and transferred to nitrocellulose membranes, which were blocked with 5% skim milk in PBS for 1 h and incubated with the primary antibody overnight at 4°C. After washing with PBS plus Tween 20 buffer, the membranes were incubated for 1 h with the appropriate secondary antibody and scanned using the Odyssey Infrared imaging system (LI-COR Biosciences).

IFN. BHK-21 cells in 96-well plates were infected with HCoV-OC43-WT (MOI = 0.01) in the presence of 10 μM of the indicated inhibitors, with chloroquine and DMSO used as the positive and negative controls, respectively. At 72 h postinfection, cells were fixed in 4% formaldehyde, permeabilized in 0.5% Triton X-100, blocked in 5% bovine serum albumin in PBS, and then probed with primary antibodies (anti-HCoV-OC43 nucleocapsid protein) for 1 h at room temperature. The cells were washed three times with PBS and then incubated with fluorescein isothiocyanate-labeled goat anti-mouse IgG (Sigma-Aldrich, St. Louis, MO, USA) at a dilution of 1:100 for 1 h. The cells were then washed and stained with 4,6-diamidino-2-phenylindole (DAPI; Invitrogen, Carlsbad, CA, USA) to detect nuclei. Fluorescence images were obtained and analyzed using a fluorescence microscope (TE2000U; Nikon, Melville, NY, USA) with a video documentation system.

Cell viability assay. Cell viability was assessed by a methyl-thiazolyl-tetrazolium (MTT) assay. After 72 h, MTT was added to a final concentration of 0.5 mg/mL, and cells were incubated for 3 h in a humidified 5% CO₂ incubator at 37°C. The plates were then centrifuged (500 × g, 10 min), and the supernatant was removed from each well by aspiration with a micropipette. Subsequently, 100 μL of DMSO was added per well, and the plates were gently shaken. The absorbance at 580 nm was detected using a MicroELISA Auto Reader MRS80 spectrometer (Dynatech Laboratories, Inc., Charlottesville, VA, USA).

MERS-CoV entry inhibition assay. The inhibitory activity of the selected inhibitors on CoV entry was determined using pseudoviruses, as described previously (41). Briefly, 293FT cells were cotransfected with plasmids PNL4-3.luc.RE- and pVRC-MERS-S, and the culture supernatant containing sufficient pseudotyped MERS-CoV was collected at 72 h posttransfection. Upon reaching a density of 5,000 cells/well in a 96-well plate, dipeptidyl peptidase 4 (DPP4)-expressing HuH7.5 cells were infected with 200 TCID₅₀ of the pseudovirus MERS-CoV in the presence of each inhibitor at different concentrations. The culture medium was renewed with fresh medium containing 2% FBS, and luciferase activity was determined after an additional 48 h of incubation using a Promega GloMax 96 plate luminometer (Promega).

Mice and infection. Twelve-day-old female BALB/c mice were inoculated via the intranasal route with 100 TCID₅₀ HCoV-OC43-WT or rOC43-ns2Del-Rluc. After 2 h, the mice were treated with the test compounds in 1× DMSO and PBS buffer. The animals were injected intraperitoneally daily with 10 μL of the compounds (emetine was used at 5 mg/kg, chloroquine was used at 30 mg/kg, and other inhibitors were used at 15 mg/kg), and survival was monitored for 20 days postinoculation. All procedures involving animals were performed in compliance with the Guide for the Care and Use of Laboratory Animals of the People’s Republic of China. The study protocol was approved by the Committee on the Ethics of Animal Experiments of the Chinese Center for Disease Control and Prevention.

BLI. BALB/c mice were infected with rOC43-ns2Del-Rluc and treated with lycorine or DMSO-PBS, followed by immediate anesthetization with 2% isoflurane and intraperitoneal administration of ViviRen in vivo Renilla luciferase substrate (20 μg/g; Promega) at 2, 3, and 4 days postinoculation. The mice were positioned in a specially designed box and placed onto the stage inside the light-tight camera box.
were imaged within 5 min postinjection of the substrate, and photon flux was quantitated using Living Image software (PerkinElmer, Waltham, MA, USA).

ICH. Mouse brains were removed at 3 days postinfection and fixed in formalin, and the tissues were processed for IHC, as described previously (66).

Statistical analysis. Differences between groups were examined for statistical significance using Student’s t test. A P of <0.05 was considered statistically significant.

SUPPLEMENTAL MATERIAL

Supplemental material for this article may be found at https://doi.org/10.1128/JVI.00023-19.

SUPPLEMENTAL FILE 1, XLS file, 0.5 MB.

ACKNOWLEDGMENTS

We thank P. J. Talbot (INRS-Institute Armand-Frappier, Université du Québec, Laval, Québec, Canada) for providing the infectious clone of HCoV-OC43 (WT), Bart L. Haagmans and Ron A. M. Fouchier (Erasmus MC, Rotterdam, the Netherlands) for providing MERS-CoV (isolate hCoV-EMC/2012), Lia van der Hoek (Center for Infection and Immunity Amsterdam, Academic Medical Center, University of Amsterdam, the Netherlands) for providing HCoV-NL63 strain Amsterdam I, and Hongyu Deng (CAS Key Laboratory of Infection and Immunity, Institute of Biophysics, Chinese Academy of Sciences) for providing MHV strain A59.

This work was supported by the National Key Research and Development Program of China (grant 2016YFD0500300 to W.T. and grant 2016YFC1200200 to B.H.) and by the Megaproject for Infectious Disease Research of China (grant 2016ZX10004001-003 to W.T.).

The funders had no role in the study design, data collection and analysis, decision to publish, or preparation of the manuscript.

We have submitted the ICMJE Form for Disclosure of Potential Conflicts of Interest. Conflicts that the editors consider relevant to the content of the manuscript have been disclosed.

L.S. and W.T. designed the study and wrote the first draft of the manuscript. L.S., J.N., C.W., B.H., W.W., N.Z., Y.D., H.W., and F.Y. conducted the experiments. L.S. conducted the statistical analysis. L.S., S.C., and W.T. performed data interpretation. B.H. and W.T. supervised the project.

The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the China CDC.

REFERENCES


36. Shen et al. Journal of Virology

June 2019 Volume 93 Issue 12 e00023-19

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