

Article

# Identification of an Antiviral Compound from the Pandemic Response Box that Efficiently Inhibits SARS-CoV-2 Infection In Vitro

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**Abstract:** With over 50 million currently confirmed cases worldwide, including more than 1.3 million deaths, the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) pandemic has a major impact on the economy and health care system. Currently, limited prophylactic or therapeutic intervention options are available against SARS-CoV-2. In this study, 400 compounds from the antimicrobial “pandemic response box” library were screened for inhibiting properties against SARS-CoV-2. An initial screen on Vero E6 cells identified five compounds that inhibited SARS-CoV-2 replication. However, validation of the selected hits in a human lung cell line highlighted that only a single compound, namely Retro-2.1, efficiently inhibited SARS-CoV-2 replication. Additional analysis revealed that the antiviral activity of Retro-2.1 occurs at a post-entry stage of the viral replication cycle. Combined, these data demonstrate that stringent in vitro screening of preselected compounds in multiple cell lines refines the rapid identification of new potential antiviral candidate drugs targeting SARS-CoV-2.

**Keywords:** pandemic response box; drug repurposing; compound screen; SARS-CoV-2; Retro-2.1; remdesivir; Vero E6; Calu-3

## 1. Introduction

In December 2019, a new zoonotic coronavirus emerged in Wuhan, Hubei province, China, named severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), which is the etiological agent of coronavirus disease 2019 (COVID-19) [1–3]. The clinical features of SARS-CoV-2-infected patients range from mild cold-like symptoms to severe illness ultimately leading to acute respiratory distress syndrome [2,4]. Patients at an older age and with underlying comorbidities are at higher risk of developing severe courses of COVID-19 [5]. Despite unprecedented international public health response measures to contain SARS-CoV-2 transmissions, the viral outbreak is currently categorized as a pandemic with over 50 million confirmed laboratory cases reported worldwide, including over 1.3 million deaths as of November 2020 [6]. At present, and despite earlier outbreaks of SARS-CoV and Middle East respiratory syndrome (MERS)-CoV, there are limited approved antiviral treatment

options, such as antiviral drugs, vaccines and immuno-prophylaxis, that can be used prophylactically or therapeutically to halt the current SARS-CoV-2 infections.

Vaccine development is a long process to reach approval for clinical use, and although SARS-CoV-2 vaccine developments are currently expedited, the eventual worldwide vaccine distribution may be delayed for additional months [7]. Moreover, while vaccines are used prophylactically, antiviral drugs can be employed both prophylactically and therapeutically. For SARS-CoV-2, several antiviral compounds have been evaluated, such as the nucleoside analogue remdesivir, the transmembrane serine protease 2 (TMPRSS2) inhibitor camostat mesylate, and the antimalaria drug (hydroxy-) chloroquine, targeting different stages of the viral replication cycle [8–10]. All three antiviral drugs have recently been tested in large patient cohorts, whereby remdesivir has been shown to reduce the hospitalization time, but proved to exert only a marginal benefit for patients with severe COVID-19 disease [11]. Several clinical trials with either hydroxychloroquine or chloroquine did not reveal any benefit in the clinical outcome [11–15]. Additionally, RNA viruses, including coronaviruses, are known to rapidly evade antiviral drug inhibition by developing resistance mutations and subsequent selection of drug-resistant viral populations [16–18]. Therefore, the use of multiple drug regimens as well as expanding the repertoire of available antiviral treatment options are of crucial importance to combat the SARS-CoV-2 pandemic.

Since the beginning of the 21st century, multiple epidemics have been caused by a viral or bacterial agent, such as the Ebola-, measles-, Zika-viruses and cholera [19]. Some epidemics have even reached pandemic proportions, such as the influenza A/H1N1 virus and the currently circulating SARS-CoV-2 [20,21]. As a rapid response to these virulent agents, the Medicines for Malaria Venture and Drugs for Neglected Diseases initiatives developed the pandemic response box (PRB), a compound library containing 400 compounds with antibacterial, antifungal and antiviral properties. This compound library allows rapid evaluation of potential drug repurposing against newly emerging pathogens.

To this end, we performed an *in vitro*-based screen of 400 preselected compounds with antibacterial, antifungal, and antiviral properties contained in the PRB and assessed their antiviral activity against SARS-CoV-2. A stringent large-scale screen in Vero E6 cells highlighted sixteen compounds that prevented virus-induced cytopathogenic effects (CPE) while displaying low cytotoxicity and no detrimental effect on cell viability. Further validation using Vero E6 and Calu-3 cells revealed that only one compound, Retro-2.1, showed substantial SARS-CoV-2 inhibition while the antiviral effects of the other compounds were only observed in Vero E6 cells. Time-of-addition analysis indicated that Retro-2.1 targets SARS-CoV-2 at a post-entry stage of the viral replication cycle. Together, these data demonstrate that stringent *in vitro* screening of preselected compounds on different cell lines refines the rapid identification of a new potential antiviral candidate drugs targeting SARS-CoV-2.

## 2. Materials and Methods

### 2.1. Cell Lines and Viruses

Vero E6 cells (kindly provided by M. Müller and C. Drosten, Charité, Berlin, Germany) and Calu-3 cells (HTB-55, American Type Culture Collection (ATCC), Manassas, VA, USA) were propagated in Dulbecco's modified Eagle Medium–GlutaMAX, 10% (*v/v*) heat-inactivated fetal bovine serum, 100 mg/mL streptomycin, 100 IU/mL penicillin, 1% (*w/v*) non-essential amino acids and 15 mM 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES, Gibco, Gaithersburg, MD, USA). Cells were maintained at 37 °C in a humidified incubator with 5% CO<sub>2</sub>. SARS-CoV-2 (SARS-CoV-2/München-1.1/2020/929 [22]) stocks were produced on Vero E6 cells, aliquoted and stored at –80 °C. Viral titers were determined by tissue culture infectious dose 50 (TCID<sub>50</sub>) on Vero E6 cells after 72 h incubation at 37 °C in a humidified incubator with 5% CO<sub>2</sub>.

## 2.2. Compound Preparation of the Pandemic Response Box

All 400 compounds (purity of >90%) distributed in 96-well plates were dissolved and diluted in dimethyl sulfoxide (DMSO, Sigma-Aldrich, St. Louis, MO, USA) and aliquoted at a concentration of 1 mM in 96-well plates (Techno Plastic Products AG, Trasadingen, Switzerland) and were kept at  $-20^{\circ}\text{C}$  until further use. Compounds were diluted at the indicated concentration in cell culture medium. The control compound remdesivir (MedChem Express, NJ, USA, HY-104077) was diluted to a 20 mM working stock in DMSO and stored at  $-80^{\circ}\text{C}$ , while K22 was prepared and stored as described previously [17].

## 2.3. Antiviral Activity Screening

Vero E6 cells were seeded in 96-well clear bottom, black wall plates (Costar, Tewksbury, MA, USA), at a density of 20,000 cells per well, one day prior to the experiment. Cells were pretreated for 2 h with 1  $\mu\text{M}$  of each compound contained in the PRB. Remdesivir [8], K22 [17], and vehicle controls (DMSO) were included in each plate. Subsequently, cells were infected with SARS-CoV-2 at a multiplicity of infection (MOI) of 0.01 in compound-containing medium and incubated at  $37^{\circ}\text{C}$  in a humidified incubator with 5%  $\text{CO}_2$ . Uninfected (mock) controls were included in each plate. At 48 h post-infection (hpi), cells were fixed with 4% (*v/v*) neutral buffered formalin (Formafix AG, Hittnau, Switzerland) and stained with crystal violet. Cell viability and cytotoxicity were assessed in parallel, in identically treated, uninfected plates. Two independent experiments were performed, each including a technical duplicate. Wells containing an intact cell layer without apparent CPE after infection and displaying high cell viability and low cytotoxicity were considered as hits.

## 2.4. Cell Cytotoxicity and Cell Viability

Cell cytotoxicity and viability were assessed using CellTox™ green cytotoxicity assay (Promega, Dübendorf, Switzerland) and CellTiter-Glo®2.0 assay (Promega), respectively, according to manufacturer's protocols. Readout was performed on a Cytation 5 Cell Imaging Multi-Mode Reader (BioTek, Sursee, Switzerland). The mean raw values were standardized by calculating the z-scores and individual cell cytotoxicity and viability scores of compounds exceeding a confidence interval of 95% (z-score of -2 or +2) were regarded as nonviable hits.

## 2.5. Half Maximal Effective Concentration ( $\text{EC}_{50}$ ) Determination of Selected Compounds

Vero E6 and Calu-3 cells were seeded in 96-well clear bottom, black wall plates (Costar), at a density of 20,000 or 40,000 cells per well, respectively, one day prior to the experiment. Cells were pretreated for 2 h with 2-fold serial dilutions of selected compounds, ranging from 4  $\mu\text{M}$  to 0.062  $\mu\text{M}$ . Cells were infected with SARS-CoV-2 (MOI of 0.01) in compound-containing medium and incubated at  $37^{\circ}\text{C}$  in a humidified incubator with 5%  $\text{CO}_2$ . Cells were fixed with 4% (*v/v*) neutral buffered formalin at 24 and 48 (Vero E6) or 48 (Calu-3) hours post-infection and processed for immunofluorescence analysis. Briefly, cells were permeabilized with 0.1% (*v/v*) Triton X-100 (Merck, Kenilworth, NJ, USA) for 5 min and blocked in phosphate-buffered saline (PBS), supplemented with 50 mM  $\text{NH}_4\text{Cl}$ , 0.1% (*w/v*) saponin (Sigma) and 2% (*w/v*) bovine serum albumin (Immunoglobulin G (IgG)-free, Jackson Immunoresearch, Westgrove, PA, USA). SARS-CoV-2 antigen-positive cells were detected using a rabbit polyclonal anti-SARS-CoV nucleocapsid protein (Rockland, Limerick, PA, USA 200-401-A50) and a secondary Alexa Fluor® 488-labeled donkey anti-rabbit IgG (H + L) or Alexa Fluor® 647-labeled donkey anti-rabbit IgG (H + L) (Jackson Immunoresearch). Samples were counterstained using 4',6-diamidino-2-phenylindole (DAPI, Thermo Fisher Scientific, Darmstadt, Germany) to visualize the nuclei and finally washed with PBS.

Images were acquired on a Cytation 5 Cell Imaging Multi-Mode Reader (BioTek) equipped with a 4 $\times$  air objective (numerical aperture: 0.13). Four images per well were acquired to cover the entire surface of the well and processed and stitched using the Gen5 ImagePrime software package

(v. 3.08.01). The percentage of virus-infected cells in each well was calculated by dividing the number of cells with a cytoplasmic green fluorescent protein (GFP) signal (SARS-CoV nucleocapsid) by the total number of detected cells (DAPI). Nuclei were segmented using a primary mask with dynamic thresholding and a secondary mask encompassing the cytoplasm was based on the primary mask, dilated by 5  $\mu\text{m}$ . The half maximal effective concentration ( $\text{EC}_{50}$ ) was calculated in GraphPad Prism (San Diego, CA, USA, v. 9.0.0) using the non-linear variable slope with four parameters option, the corresponding  $\text{EC}_{90}$ -value was calculated based on the previously calculated  $\text{EC}_{50}$ -value and the hill slope values.

#### 2.6. Half-Maximum Cytotoxicity Concentration ( $\text{CC}_{50}$ ) Determination

To determine the cytotoxicity of the compounds and for later calculation of the selectivity index (SI), a dilution series ranging from 100  $\mu\text{M}$  to 0.19  $\mu\text{M}$  for both Vero E6 cells and Calu-3 cells was prepared in cell culture medium and incubated for 48 h at 37 °C in a humidified incubator with 5% CO<sub>2</sub>. Cytotoxicity and cell viability were both assessed as described earlier, whereby the  $\text{CC}_{50}$  was calculated in GraphPad Prism (v. 9.0.0) using the non-linear variable slope with four parameters option.

#### 2.7. Time-of-Addition Experiment

Vero E6 cells were seeded in 96-well, clear bottom, black wall plates (Costar) at a density of 20,000 cells per well. Retro-2.1 and the vehicle control (DMSO) were diluted in cell culture medium at a concentration of 10  $\mu\text{M}$  or using an equal volume of DMSO. Cells were (pre-) treated and infected with a MOI of 0.01. Following each incubation step, cells were washed three times with PBS to remove any residual compound or virus. At 24 h post-infection, supernatant was collected for viral titration, while cells were fixed and processed for immunofluorescence analysis as described before using an Alexa Fluor® 647-labeled donkey anti-rabbit IgG (H + L) as a secondary antibody (Jackson ImmunoResearch).

#### 2.8. Data Representation

Graphs were generated using GraphPad Prism software (v. 9.0.0) and the final figures were assembled in Adobe Illustrator CS6 (v. 16.0.0). Brightness and contrast of microscopy pictures were minimally adjusted and processed identically to their corresponding control using FIJI (v. 1.53c). Images were assembled using the FigureJ plugin in FIJI [23].

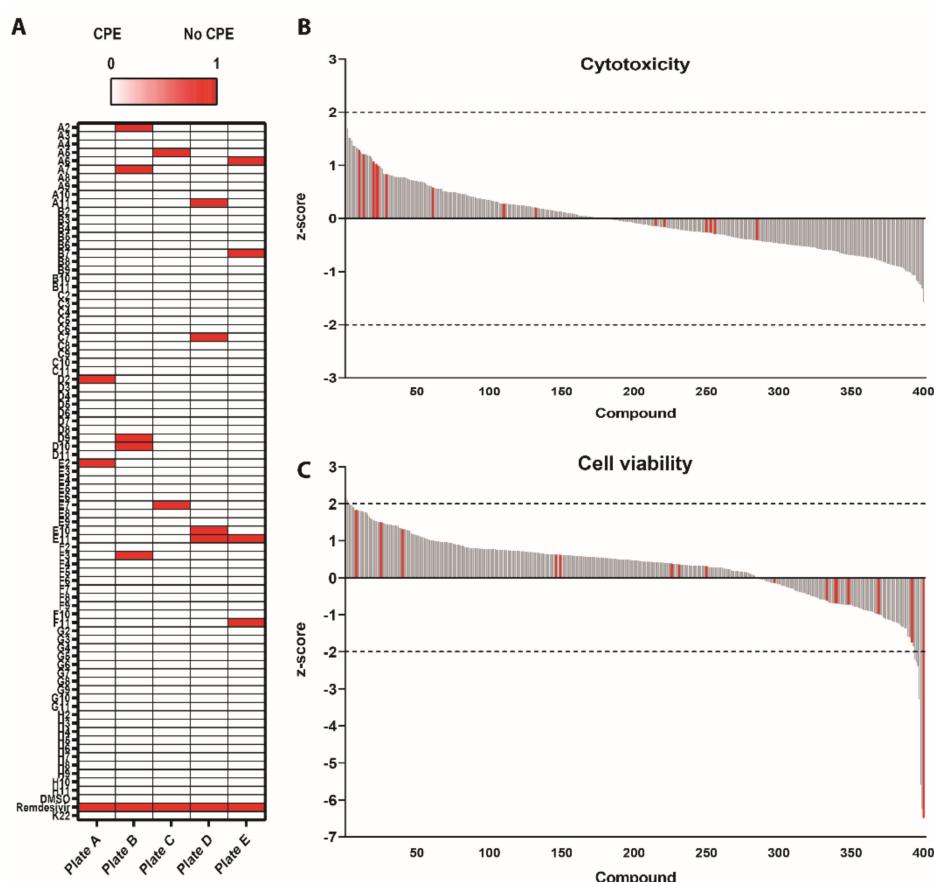
### 3. Results

#### 3.1. Survival Screen with Compounds Included in the Pandemic Response Box against SARS-CoV-2

To identify potential antiviral compounds against SARS-CoV-2 replication, 201 antibacterial, 46 antifungal and 153 antiviral molecules included in the PRB were screened using a conservative concentration of 1  $\mu\text{M}$ . Based on the documented inhibition of coronavirus replication, remdesivir was included as a positive control [8]. The compound K22 was also included in the analysis since it has been shown to inhibit a broad range of viruses including coronaviruses, arteriviruses, toroviruses as well as different members of the family Flaviviridae [17,24,25]. Vero E6 cells were pretreated for 2 h and subsequently infected with SARS-CoV-2 (MOI of 0.01) for 48 h in drug-containing medium. Cell survival was scored from 0 (virus-induced cytopathogenic effects; CPE) to 1 (no CPE), upon evaluation of SARS-CoV-2-induced CPE using crystal violet staining (Figure 1A). The cytotoxicity and cell viability were assessed on uninfected plates processed in parallel, to exclude detrimental effects of the compounds on the cells.

This survival screen resulted in a total of seventeen compounds that inhibited SARS-CoV-2-induced CPE (red). In parallel, cell viability and cell cytotoxicity assays were performed to exclude detrimental effects of each compound on the cells. The mean raw values were standardized by calculating the individual cell cytotoxicity and viability z-scores. Compounds exceeding a 95% confidence interval

(z-score of  $-2$  or  $+2$ ) were regarded as nonviable hits. This showed that only a single compound caused significant cytotoxicity (Plate D, G11; z-score 3.12) (Figure 1B). In contrast, 10 compounds exceeded the cell viability cutoff, including the previously identified compound in Plate B, D10 (z-score:  $-6.498$ ) (Figure 1C). Of note, this compound was also observed as a positive hit in the survival screen, but due to its impaired cell viability, it was excluded from further analysis. The sixteen remaining hit compounds are categorized as antifungal (three), antibacterial (six) and antiviral (seven) compounds (Appendix A), which, similarly to their vehicle control (DMSO) and remdesivir, did not influence cell cytotoxicity and cell viability. Interestingly, K22 did not show any inhibitory activity against SARS-CoV-2 at a concentration of  $1 \mu\text{M}$ . These results highlight the relevance of a conservative and rapid screening of libraries containing compounds that could potentially inhibit SARS-CoV-2.

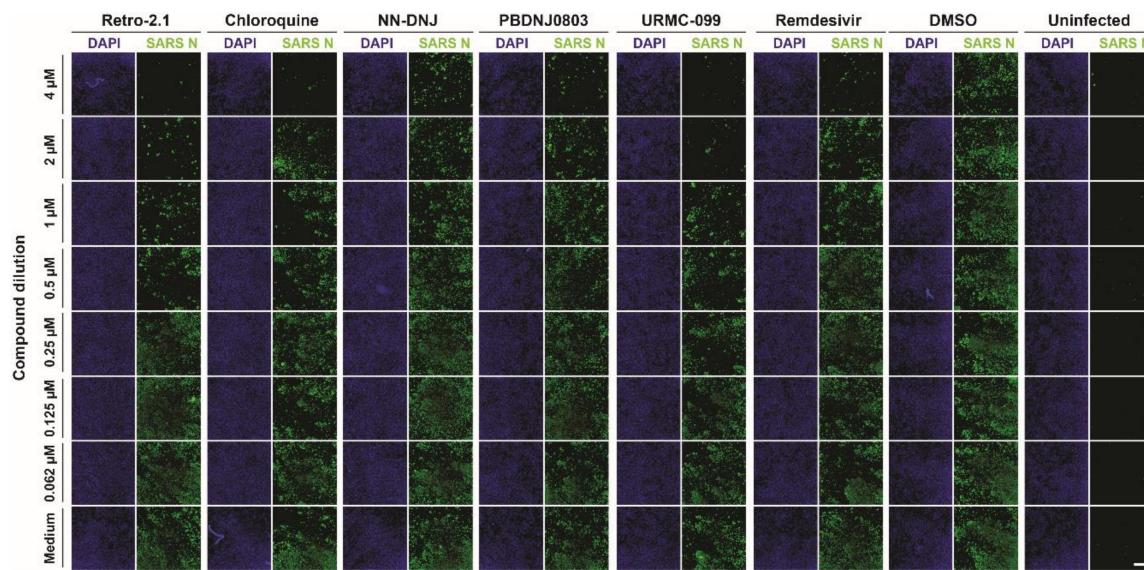


**Figure 1.** Cell survival upon infection with severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) (A), cytotoxicity (B) and viability (C) after incubation with pandemic response box compounds. The survival, cytotoxicity and viability were measured after 48 h of incubation on Vero-E6 cells at  $37^\circ\text{C}$  in a humidified incubator with  $5\% \text{CO}_2$ . The heatmap shows the position on the plates where survival was observed with a multiplicity of infection (MOI) of 0.01, which was scored from 0 to 1 (0 = cytopathogenic effects (CPE) and 1 = no CPE) (A). Calculated z-scores of each individual compound for cell cytotoxicity (B) and cell viability (C). A confidence interval of 95% (z-score of  $-2$  or  $+2$ ) was used as cutoff values, which are indicated with gray dashed lines. Each bar represents an individual compound sorted from high to low z-scores (*x*-axis). The red bars indicate the compounds that showed inhibition of SARS-CoV-2 during the survival screen. Results are shown as mean of two individual experiments performed in two technical replicates.

### 3.2. Antiviral Efficacy against SARS-CoV-2

To further confirm and evaluate the extent of antiviral activity of the previously highlighted sixteen compounds, cells were pretreated with the selected compounds at concentrations ranging

from 4  $\mu\text{M}$  to 0.062  $\mu\text{M}$  and infected with SARS-CoV-2 (MOI of 0.01). After 24 h of infection, cells were fixed and processed for immunofluorescence analysis using the anti-SARS-CoV nucleocapsid protein antibody and DAPI (Figure 2).

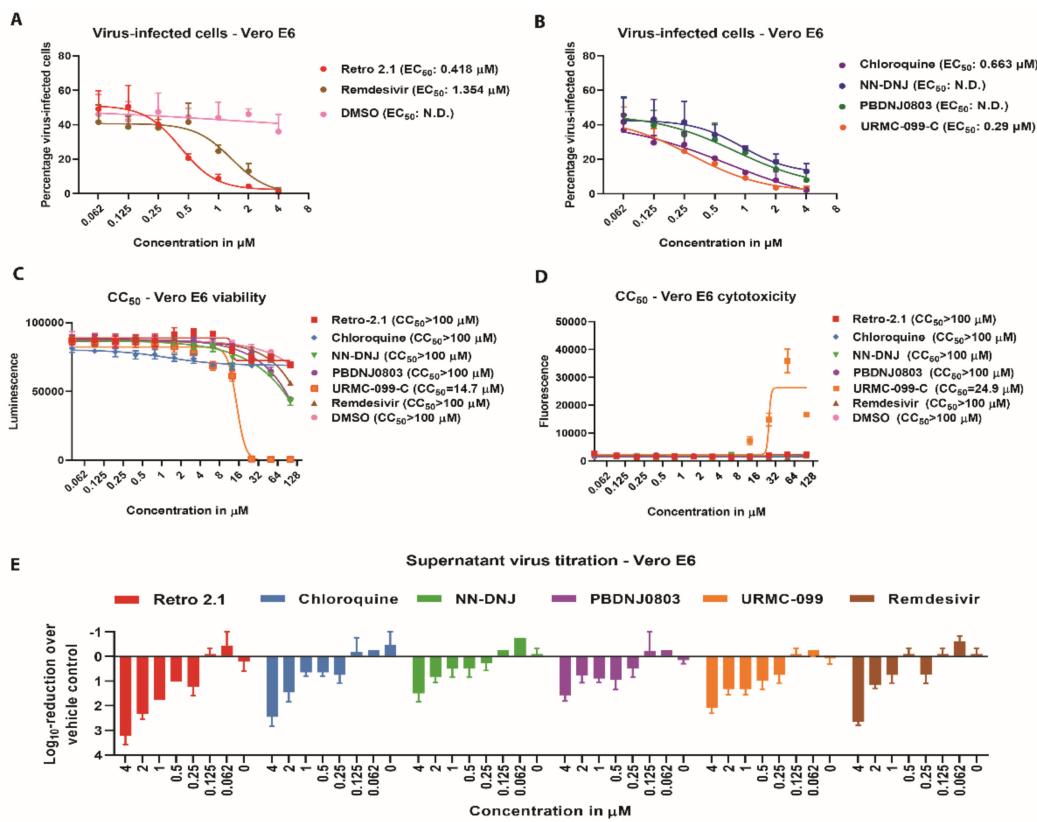


**Figure 2.** Assessment of the antiviral efficiency by immunofluorescence. To determine the efficiency of inhibition of each specific compound from the survival screen, a two-fold dilution series ranging from 4  $\mu\text{M}$  to 0.062  $\mu\text{M}$  was prepared following infection with SARS-CoV-2. Vero E6 cells were pretreated with the compound 2 h prior to infection with SARS-CoV-2 (MOI of 0.01) at 37 °C in a humidified incubator with 5% CO<sub>2</sub>. Cells were fixed 24 h post-infection, followed by immunostaining with the cross-reactive SARS-CoV nucleoprotein antigen (SARS N) and 4',6-diamidino-2-phenylindole (DAPI). Remdesivir was included as a positive control. The images are representative of the results of three individual experiments. Scale bar: 1000  $\mu\text{m}$ .

The efficiency of the selected compounds to inhibit SARS-CoV-2 (Figure 3A,B), as well as their individual effects on cell viability (Figure 3C), cytotoxicity (Figure 3D), and viral titer were assessed (Figure 3E). The EC<sub>50</sub> values for each compound were inferred by calculating the percentage of virus-infected cells. This indicated that five out of sixteen candidate compounds, *n*-nonyldeoxynojirimycin (NN-DNJ), PDNJ0803, chloroquine, Retro-2.1 and URMC-099-C, inhibited SARS-CoV-2 in a dose-dependent manner (Figure 3C,D). Moreover, these five compounds showed EC<sub>50</sub> values ranging from 0.29  $\mu\text{M}$  to 0.63  $\mu\text{M}$ , without increased cytotoxicity or decreased cell viability (Figure 3A–D). These results were corroborated by the dose-dependent reduction in viral titer, where Retro-2.1 displayed the strongest reduction (Figure 3E). Processing the samples at 48 h post-infection rather than 24 h post-infection showed that the inhibitory effect of all compounds is reduced at a later phase during infection (Figure S1). The remaining eleven compounds showed little or no inhibition at lower concentrations and were therefore excluded from further analysis (Figure S2).

In parallel to the inhibition efficiency, the half-maximum cytotoxicity concentration (CC<sub>50</sub>) was determined for each compound at concentrations ranging from 0.04  $\mu\text{M}$  to 100  $\mu\text{M}$  on Vero E6 cells. This demonstrated that the previously tested compounds were all well tolerated at concentrations up to 100-fold higher than the one used in the initial screen (1  $\mu\text{M}$ ) (Figure 3C,D, Appendix B). In contrast, only URMC-099-C displayed moderate cell cytotoxicity, resulting in a CC<sub>50</sub> of 14.7  $\mu\text{M}$  based on viability and 24.9  $\mu\text{M}$  based on cytotoxicity, while all other compounds had a CC<sub>50</sub> above 100  $\mu\text{M}$  (Figure 3C,D, Appendix B). The resulting selectivity indexes (SI, the CC<sub>50</sub>-score divided by the EC<sub>50</sub>-score) showed that Retro-2.1 (SI = 239.2) is the most efficient and least cytotoxic inhibitor, followed by chloroquine (SI = 150.8) and URMC-099-C (SI = 50.7). The positive control remdesivir showed a SI of 73.8, while no SI values were calculated for NN-DNJ and PBDNJ0803 as the EC<sub>50</sub>-value could not be determined.

(Figure 3B, Appendix B). Combined, these results demonstrate the identification of five compounds of the pandemic response box library that inhibited SARS-CoV-2 in Vero E6 cells.

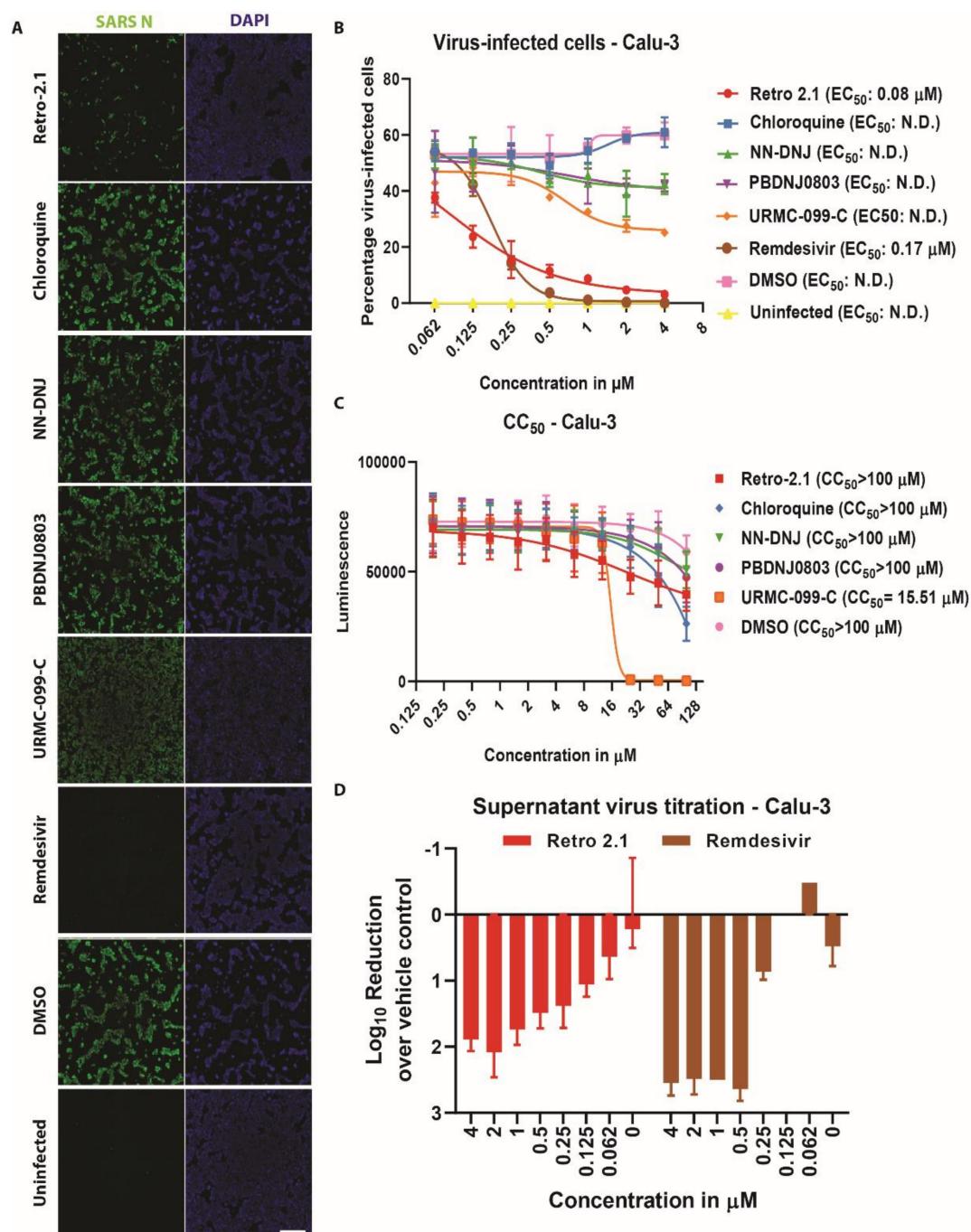


**Figure 3.** Half maximal effective concentration ( $\text{EC}_{50}$ ) determination of the five hit compounds showing inhibition against SARS-CoV-2. Vero E6 cells were pre-treated for 2 h with the indicated compound concentrations prior to SARS-CoV-2 infection (MOI 0.01) at  $37^\circ\text{C}$  in a humidified incubator with 5%  $\text{CO}_2$ . Following infection, cells were fixed and processed for immunofluorescence analysis. To determine the reduction in the percentage of virus-infected cells, the number of cells with a green fluorescent protein (GFP)-positive cytoplasmic signal (infected cells) was divided by the total number of cells (DAPI, nuclei). The percentage of virus-infected cells of Retro-2.1 was compared to remdesivir, DMSO (A) and chloroquine, *n*-nonyldeoxyojirimycin (NN-DNJ), PBDNJ0803 and URMC-099 (B). Abbreviations: N.D.: not determined. Results are displayed as means and SD of three independent experiments. The results of the half-maximum cytotoxicity concentration ( $\text{CC}_{50}$ ) based on viability (C) or cytotoxicity (D) are shown as means and SD from two individual experiments performed in two technical replicates. Corresponding supernatants were titrated by tissue culture infectious dose 50 (TCID<sub>50</sub>) to quantify the  $\log_{10}$  reduction in SARS-CoV-2 titer (E). Results are displayed as means and SD of two independent experiments.

### 3.3. Inhibition Analysis of SARS-CoV-2 Infection on Calu-3 Cells

To exclude potential cell-specific biases, the previous experiments were additionally performed using a human lung adenocarcinoma cell line (Calu-3), which recapitulates important biological aspects of the natural site of infection [26,27]. Interestingly, inhibition of SARS-CoV-2 replication was only observed upon treatment with Retro-2.1 ( $\text{EC}_{50}: 0.08 \mu\text{M}$ ) or with remdesivir ( $\text{EC}_{50}: 0.17 \mu\text{M}$ ), based on the percentage of virus-infected cells (Figure 4A,B). The other four compounds showed no viral inhibition in Calu-3 cells. Determination of the  $\text{CC}_{50}$  showed that most compounds were well tolerated by the Calu-3 cells with values  $>100 \mu\text{M}$ , except for URMC-099-C ( $\text{CC}_{50} = 15.51 \mu\text{M}$ ) (Figure 4C). Consistently, the percentage of SARS-CoV-2-infected Calu-3 cells treated with Retro-2.1 (SI: 1250) or remdesivir (SI: 588) correlated with viral titers (Figure 4D). This highlights that from the

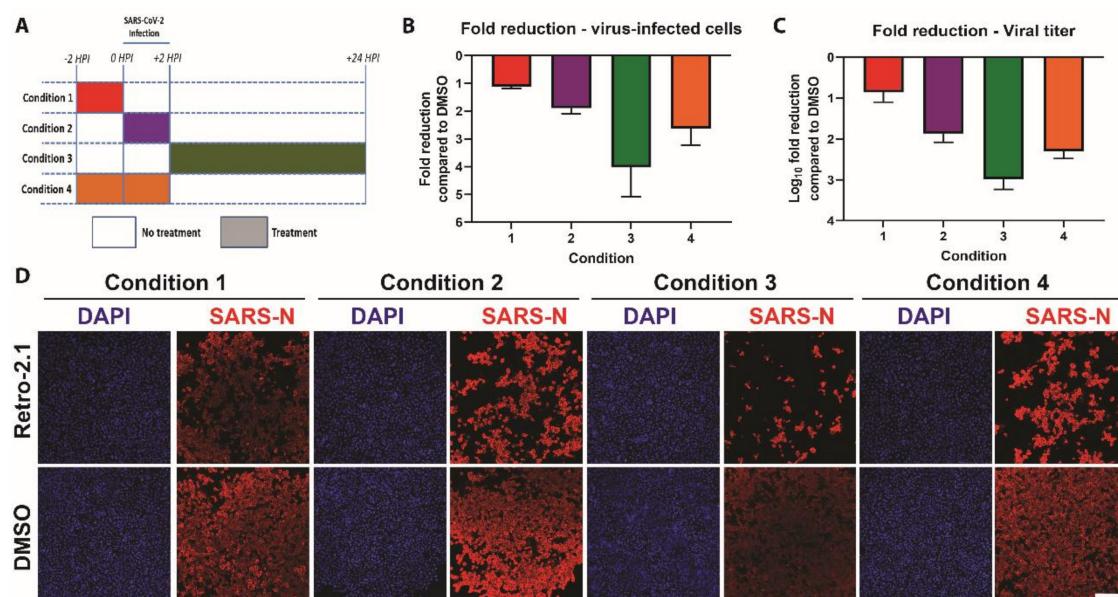
five antiviral compounds identified using Vero E6 cells, only Retro-2.1 showed efficient inhibition of SARS-CoV-2 replication in both Vero E6 and Calu-3 cells.



**Figure 4.** Analysis of the antiviral efficiency on Calu-3 cells. Calu-3 cells were pre-treated 2 h at the indicated concentration prior to SARS-CoV-2 infection (MOI 0.02) at 37 °C in a humidified incubator with 5% CO<sub>2</sub>. Cells were fixed 48 h post-infection, followed by immunostaining with the cross-reactive SARS-CoV nucleoprotein antibody (SARS N) and DAPI (A). The images are representative of the results of three individual experiments (only treatments with 4  $\mu\text{M}$  are shown). Scale bar: 1000  $\mu\text{m}$ . The half maximum inhibitory concentration ( $\text{EC}_{50}$ ) was determined using the percentage of infected cells and half maximum cytotoxicity concentration ( $\text{CC}_{50}$ ) was determined using cell viability (B, C). Results are displayed as means and SD of three individual experiments. Abbreviation: N.D.: not determined. Corresponding log<sub>10</sub>-reduction in SARS-CoV-2 viral titer for Retro 2.1 and remdesivir treatments (D).

### 3.4. Time-of-Addition

To determine which step of the viral replication cycle is affected by Retro-2.1, we performed a time-of-addition experiment. Confluent Vero E6 cells were either pre-treated with 10  $\mu$ M of Retro-2.1, or an equal volume of vehicle control (DMSO), either two hours prior to infection, during infection, or two hours after infection. Alternatively, Retro-2.1 treatment was performed during both the pretreatment and virus infection steps (Figure 5A). At 24 h post-infection, the supernatant from each condition was titrated to determine the infectious viral titer, and the cells were processed for immunofluorescence analysis as described earlier.



**Figure 5.** Time-of-addition of Retro-2.1. Time-of-addition was performed by Retro-2.1 treatments pre-, during and/or post-infection (A). Confluent Vero E6 cells were fixed 24 h post-infection (hpi) followed by processing for immunostaining with the cross-reactive SARS-CoV nucleoprotein antibody (SARS-N) and DAPI. The percentage of SARS-CoV-2-infected cells (B) and viral titers in the supernatant (C) were determined at 24 h post-infection. Graphs show means and SD from three individual experiments, performed in two technical replicates. (D) Representative images of the indicated conditions. Scale bar: 300  $\mu$ m.

The most pronounced inhibitory effect was observed when cells were treated with Retro-2.1 after SARS-CoV-2 infection (condition 3) (Figure 5B,D). In contrast, when cells were treated with Retro-2.1 prior to infection (condition 1), only a marginal reduction was observed. Treatment of cells during infection (condition 2) showed a more pronounced inhibition, which was comparable to condition 4, which combines treatment with Retro-2.1 prior and during infection. These results were corroborated by the viral titers analyzed at 24 h post-infection, demonstrating a >1000-fold reduction in viral titer in condition 3 (Figure 5C). These data indicate that Retro-2.1 likely interferes with the viral replication cycle of SARS-CoV-2 during a post-entry step.

## 4. Discussion

In this study, we have demonstrated that a conservative in vitro screening of 400 compounds from the PRB using two different cell lines refines the identification of effective antiviral candidate drugs targeting SARS-CoV-2. The first stage of the screening using Vero E6 cells led to the identification of five compounds displaying effective antiviral activity against SARS-CoV-2. This included the anti-malaria drug chloroquine and antibacterial Retro-2.1, as well as the antiviral compounds NN-DNJ, PDNJ0803, and URM-C-099-C. These compounds showed a dose-dependent inhibition of SARS-CoV-2

infection. However, validation of these selected hits on the human lung cell line Calu-3 revealed that only Retro-2.1 efficiently inhibited SARS-CoV-2 replication. Time-of-addition analysis showed that Retro-2.1 likely impairs SARS-CoV-2 replication during a post-entry step of the viral replication cycle. Combined, these results demonstrate that robust evaluation of novel antiviral candidate drugs should be performed in different cell lines to exclude potential cell-dependent artefacts. K22, which exerts a well-documented inhibition of several coronaviruses including SARS-CoV and MERS-CoV, was, to our knowledge, not tested against SARS-CoV-2 [17,24,25]. Here, we show that K22 did not inhibit SARS-CoV-2 infection in Vero E6 cells, further emphasizing the requirements of rigorous screening conditions and robust controls during drug library screening assays.

In the light of the current pandemic, it has become evident that drug repurposing is a promising strategy that could rapidly help to mitigate SARS-CoV-2 infections. In our study, we employed a stringent survival screening approach using a library of 400 preselected compounds at a concentration of 1  $\mu$ M. Other recent studies have employed different types of libraries and screening conditions, such as incubation time, cell density and MOI, but alike our study, most employed Vero E6 cells as an initial screening platform and used remdesivir as a positive control [28,29]. In line with these studies, we also identified chloroquine, which is approved for clinical use in the context of malaria treatment, as a potent antiviral compound against SARS-CoV-2, showing the reproducibility of these screens. Although the evaluation of chloroquine treatments against SARS-CoV-2 infections in clinical settings has been rapidly undertaken, it has been demonstrated that its antiviral activity on viral entry is restricted to Vero E6 cells [26]. Concordantly, in our study, we also observed that the impairment of SARS-CoV-2 replication by chloroquine was restricted to Vero E6 cells. Additionally, limited inhibition of SARS-CoV-2 by chloroquine was more recently confirmed by experimental infections of Syrian golden hamsters and rhesus macaques [30]. Altogether, these results emphasize the importance of the rigorous testing of antiviral compounds in multiple cell lines to exclude potential cell-based artefacts prior to initiating clinical trials.

In this study, we identified Retro-2.1 as a novel potent compound that inhibited SARS-CoV-2 replication in both Vero E6 and the human lung Calu-3 cell lines. Retro-2.1 has been previously shown to inhibit a broad range of viruses including enteroviruses, filoviruses, herpes simplex virus, vaccinia virus and polyomaviruses [31–36]. It has been suggested that Retro-2.1 remodels the intracellular distribution of syntaxins, which consequently alters vesicular retrograde transport between endosomes and the Golgi apparatus [37,38]. Moreover, convergent data have reported the strong dependence on cellular endomembranes and cellular trafficking pathways for efficient coronavirus replication. Several syntaxins (stxbp1, stxbp3, stx5a, and stx18) were identified in close proximity of the coronavirus replication and transcription complex during infection, some of which were further suggested to assist viral replication [39]. Consistent with these studies, our time-of-addition analysis revealed that Retro-2.1 substantially impairs SARS-CoV-2 infection during a post-entry stage of the viral replication cycle. Further investigations are required to define the precise molecular targets as well as the mode of action of Retro-2.1.

In summary, the stringent screening of 400 compounds identified Retro-2.1 as a potent antiviral compound against SARS-CoV-2 in vitro in different cell lines. This vindicates further efficacy and safety testing in other biologically relevant pre-clinical in vitro and in vivo models as novel intervention strategies against SARS-CoV-2 and other coronaviruses.

**Supplementary Materials:** The following are available online at <http://www.mdpi.com/2076-2607/8/12/1872/s1>. Figure S1: The EC<sub>50</sub> determination of the five top compounds after 48 h of incubation. Figure S2: EC<sub>50</sub> determination of eleven compound hits that showed partial inhibition against SARS-CoV-2.

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**Conflicts of Interest:** The authors declare no conflict of interest.

## Appendix A

**Table A1.** The sixteen compounds that showed inhibition of SARS-CoV-2 after the survival screen. Remdesivir was included as a reference drug control.

Plate Position:	Section	MVV-Number	CHEMBL ID	Reference	Name	Clinical Stage According CHEMBL	Chemical Formula
Plate A, D2	Antifungals	MMV1634386	CHEMBL3311228	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL3311228/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL3311228/</a>	Otesconazole	3 (Phase III)	C23 H16 F7 N5 O2
Plate A, E2	Antifungals	MMV637528	CHEMBL64391	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL64391/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL64391/</a>	Itraconazole	4 (approved)	C35 H38 Cl2 N8 O4
Plate B, A2	Antibacterials	MMV1483032	CHEMBL243644	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL243644/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL243644/</a>	AC1MTT7I	0 (research)	C22 H18 N4 O4
Plate B, A7	Antibacterials	MMV637945	CHEMBL403	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL403/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL403/</a>	Sulbactam	4 (approved)	C8 H11 N O5 S
Plate B, D9	Antibacterials	MMV1582492	CHEMBL3109593	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL3109593/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL3109593/</a>	Retro-2.1	0 (research)	C23 H18 F N3 O S2
Plate B, F3	Antibacterials	MMV1582487	CHEMBL198796	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL198796/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL198796/</a>	Decylphosphinate	0 (research)	C13 H28 N O4 P
Plate C, A5	Antibacterials	MMV1578576	CHEMBL1568820	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL1568820/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL1568820/</a>	-	0 (research)	C15 H12 F N3 O
Plate C, E7	Antibacterials	MMV000008	CHEMBL76	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL76/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL76/</a>	Chloroquine	4 (approved)	C18 H26 Cl N3
Plate D, A11	Antivirals	MMV1593513	CHEMBL408500	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL408500/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL408500/</a>	N-Nonyl Deoxynojirimycin (NN-DNJ)	0 (research)	C15 H31 N O4
Plate D, C7	Antivirals	MMV690621	-	Patent: <a href="#">WO2006118607A2</a>	NA for racemic	-	C18 H16 Cl N3 O
Plate D, E10	Antivirals	MMV1634401	CHEMBL1652119	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL1652119/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL1652119/</a>	PBDNJ0803	0 (research)	C20 H33 N O5
Plate D, E11	Antivirals	MMV002780	CHEMBL402487	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL402487/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL402487/</a>	Noscapine	0 (research)	C22 H23 N O7
Plate E, A6	Antivirals	MMV1580482	CHEMBL2436978	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL2436978/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL2436978/</a> Patent: <a href="#">WO 2014085795 A1</a>	URMC-099-C	0 (research)	C27 H27 N5
Plate E, B7	Antivirals	MMV1593544	CHEMBL3752642	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL3752642/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL3752642/</a>	-	0 (research)	C36 H43 N3 O5
Plate E, E11	Antifungals	MMV002350	CHEMBL561	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL561/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL561/</a>	Lomefloxacin	4 (approved)	C17H19F2N3O3
Plate E, F11	Antivirals	MMV1580483	CHEMBL3960662	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL3960662/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL3960662/</a>	AZD0156	0 (research)	C26 H31 N5 O3
-	Control	-	CHEMBL4065616	<a href="https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL4065616/">https://www.ebi.ac.uk/chembl/compound_report_card/CHEMBL4065616/</a>	Remdesivir	2 (Phase II)	C27 H35 N6 O8 P

## Appendix B

**Table A2.** The EC<sub>50</sub>, EC<sub>90</sub> and CC<sub>50</sub> values of the five compounds that showed inhibition of SARS-CoV-2 during the compound dilution series on VeroE6-cells, including references to the mode of action of these compounds against other viruses. Remdesivir is included as a reference drug control. Abbreviations: N.D.: not determined.

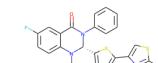
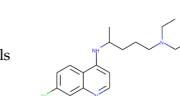
Compound	EC <sub>50</sub> in $\mu\text{M}$	EC <sub>90</sub> in $\mu\text{M}$	CC <sub>50</sub> in $\mu\text{M}$	Selective Index (SI)	Class	Examples of Susceptible Viruses	Literature Reference	Mode of Action	Administration	Tested on Cells or Organism	Molecular Structure
Retro-2.1	0.418	1.03	100	239.2	Retrograde transport inhibitor	<ul style="list-style-type: none"> <li>• Enterovirus</li> <li>• Herpes simplex virus</li> <li>• Filoviruses</li> <li>• Vaccinia virus</li> <li>• Polyomavirus</li> </ul>	[31–36]	<ul style="list-style-type: none"> <li>• Modulate intracellular vesicle transport (Enterovirus 71),</li> <li>• Blocks entry (Herpes Simplex Virus 2)</li> <li>• Blocks entry that follows by glycoprotein proteolysis (Filoviruses)</li> <li>• Localization towards the ER (Polyomavirus)</li> <li>• endosome-to-Golgi apparatus trafficking (Vaccinia virus)</li> <li>• induce cellular STX5 displacement from the Golgi upon treatment, leading to an inhibition of retrograde transport (Cytomegalovirus)</li> </ul>	<ul style="list-style-type: none"> <li>• Prophylactic</li> <li>• Therapeutic</li> </ul>	<ul style="list-style-type: none"> <li>• Vero</li> <li>• HeLa</li> <li>• 293S</li> <li>• Mice</li> </ul>	
Chloroquine	0.663	10.44	100	150.82	<ul style="list-style-type: none"> <li>• Autophagic proteolysis inhibitor</li> <li>• Endosomal acidification inhibitor</li> </ul>	<ul style="list-style-type: none"> <li>• Herpes simplex virus</li> <li>• Picornaviruses</li> <li>• Poliovirus</li> <li>• SARS-CoV</li> <li>• HIV-1</li> <li>• Hepatitis A virus</li> <li>• Hepatitis B virus</li> <li>• Hepatitis C virus</li> <li>• Influenza A virus</li> <li>• Influenza B virus</li> <li>• Bovine viral diarrhea virus</li> <li>• 229E-CoV</li> <li>• OC43-CoV</li> <li>• Chikungunya virus</li> <li>• Dengue virus</li> <li>• Crimean-Congo Hemorrhagic fever</li> <li>• Ebola virus</li> <li>• Nipah virus</li> <li>• Hendra virus</li> <li>• Lassa virus</li> <li>• Rabies virus</li> </ul>	[40–62]	<ul style="list-style-type: none"> <li>• Raise endosomal pH (Picornavirus)</li> <li>• Terminal glycosylation of the ACE2-receptor (SARS-CoV)</li> <li>• Alter the glycosylation of the glycoprotein (HIV-1)</li> <li>• Interference with endosomal acidification (Hepatitis C virus)</li> <li>• Endosomal acidification (BVDV)</li> <li>• P38 MAPK and ERK activation (229E)</li> </ul>	<ul style="list-style-type: none"> <li>• Prophylactic</li> <li>• Therapeutic</li> </ul>	<ul style="list-style-type: none"> <li>• VeroE6</li> <li>• Vero 76</li> <li>• L132</li> <li>• MDBK</li> <li>• Huh7</li> <li>• NS20 murine neuroblastoma cells</li> <li>• B-SC-1</li> <li>• A549</li> <li>• Humans</li> <li>• U937</li> <li>• MRC5</li> <li>• Guinea pigs</li> </ul>	

Table A2. Cont.

Compound	EC <sub>50</sub> in $\mu\text{M}$	EC <sub>90</sub> in $\mu\text{M}$	CC <sub>50</sub> in $\mu\text{M}$	Selective Index (SI)	Class	Examples of Susceptible Viruses	Literature Reference	Mode of Action	Administration	Tested on Cells or Organism	Molecular Structure	
N-Nonyl Deoxyx Jirimycin (NNDNJ)	N.D.	N.D.	100	N.D.	ER $\alpha$ -glucosidase inhibitors	<ul style="list-style-type: none"> <li>Bovine viral diarrhea virus</li> <li>West nile virus</li> <li>Dengue virus</li> <li>Hepatitis B virus</li> <li>Hepatitis C virus</li> <li>Flaviviruses</li> <li>Woodchuck hepatitis virus</li> <li>Ebola virus</li> <li>Tacaribe virus</li> <li>Junin virus</li> <li>SARS-CoV</li> </ul>	[63–71] US patent: 9040488	<ul style="list-style-type: none"> <li>Protein folding (Hepatitis B virus, Bovine viral diarrhea virus)</li> <li>Protein trafficking (Woodchuck hepatitis virus)</li> </ul>	<ul style="list-style-type: none"> <li>Prophylactic</li> <li>Therapeutic</li> </ul>	<ul style="list-style-type: none"> <li>MDBK</li> <li>BHK (-21)</li> <li>HepG2</li> <li>Huh7.5</li> <li>Woodchucks</li> </ul>		
PBDNJ0803	N.D.	N.D.	100	N.D.	ER $\alpha$ -glucosidase inhibitors	<ul style="list-style-type: none"> <li>Dengue virus</li> <li>Bovine viral diarrhea virus</li> <li>West nile virus</li> <li>Ebola virus</li> <li>Tacaribe virus</li> <li>Junin virus</li> </ul>	[69,70] US Patent: 9040488	<ul style="list-style-type: none"> <li>Protein folding (Bovine viral diarrhea virus)</li> </ul>	<ul style="list-style-type: none"> <li>Therapeutic</li> </ul>	<ul style="list-style-type: none"> <li>MDBK</li> <li>BHK</li> <li>Vero</li> </ul>		
URMC-099-C	0.29	1.83	14.7	50.69	Mixed-lineage kinase 3 inhibitor	<ul style="list-style-type: none"> <li>HIV-1</li> <li>Zika virus</li> </ul>	[72–75]	<ul style="list-style-type: none"> <li>Anti-inflammatory</li> <li>Blocks phosphorylation of MLK3</li> <li>Inhibits viral maturation in the endosome</li> <li>pro-viral effect in replication</li> </ul>	<ul style="list-style-type: none"> <li>Prophylactic</li> <li>Therapeutic</li> </ul>	<ul style="list-style-type: none"> <li>BV-2 microglial</li> <li>Mice</li> <li>Macrophages</li> <li>SNB-19</li> </ul>		
Remdesivir	1.354	3.39	100	73.85	Nucleoside analogue	<ul style="list-style-type: none"> <li>Ebola virus</li> <li>Marburg virus</li> <li>SARS-CoV</li> <li>MERS-CoV</li> <li>Respiratory Syncytial Virus</li> <li>Junin virus</li> <li>Lassa virus</li> <li>Hendra virus</li> <li>Nipah virus</li> <li>Rift Valley Fever Virus</li> <li>Mumps virus</li> <li>Measles virus</li> <li>Coronaviruses</li> <li>Murine hepatitis virus</li> </ul>	[18,76–80]	<ul style="list-style-type: none"> <li>Interferes with the RNA dependent RNA polymerase activity</li> </ul>	<ul style="list-style-type: none"> <li>Prophylactic</li> <li>Therapeutic</li> </ul>	<ul style="list-style-type: none"> <li>Primary macrophages</li> <li>HeLa</li> <li>HFF-1</li> <li>HMVEC-TERT</li> <li>Rhesus monkeys</li> <li>Huh7</li> <li>Hep-2</li> <li>U2OS</li> <li>Vero</li> <li>Mice</li> <li>Human airway epithelial</li> </ul>		

## References

- Wang, C.; Horby, P.W.; Hayden, F.G.; Gao, G.F. A novel coronavirus outbreak of global health concern. *Lancet* **2020**, *395*, 470–473. [[CrossRef](#)]
- Huang, C.; Wang, Y.; Li, X.; Ren, L.; Zhao, J.; Hu, Y.; Zhang, L.; Fan, G.; Xu, J.; Gu, X.; et al. Clinical features of patients infected with 2019 novel coronavirus in Wuhan, China. *Lancet* **2020**, *395*, 497–506. [[CrossRef](#)]
- Zhu, N.; Zhang, D.; Wang, W.; Li, X.; Yang, B.; Song, J.; Zhao, X.; Huang, B.; Shi, W.; Lu, R.; et al. A novel coronavirus from patients with pneumonia in China, 2019. *N. Engl. J. Med.* **2020**, *382*, 727–733. [[CrossRef](#)] [[PubMed](#)]
- Zhou, F.; Yu, T.; Du, R.; Fan, G.; Liu, Y.; Liu, Z.; Xiang, J.; Wang, Y.; Song, B.; Gu, X.; et al. Clinical course and risk factors for mortality of adult inpatients with COVID-19 in Wuhan, China: A retrospective cohort study. *Lancet* **2020**, *395*, 1054–1062. [[CrossRef](#)]
- Yang, J.; Zheng, Y.; Gou, X.; Pu, K.; Chen, Z.; Guo, Q.; Ji, R.; Wang, H.; Wang, Y.; Zhou, Y. Prevalence of comorbidities in the novel Wuhan coronavirus (COVID-19) infection: A systematic review and meta-analysis. *Int. J. Infect. Dis.* **2020**, *94*, 91–95. [[CrossRef](#)] [[PubMed](#)]
- Coronavirus Disease (COVID-19) Situation Reports. Available online: <https://www.who.int/emergencies/diseases/novel-coronavirus-2019/situation-reports> (accessed on 18 November 2020).
- Lurie, N.; Saville, M.; Hatchett, R.; Halton, J. Developing Covid-19 Vaccines at Pandemic Speed. *N. Engl. J. Med.* **2020**, *382*, 1969–1973. [[CrossRef](#)] [[PubMed](#)]
- Wang, M.; Cao, R.; Zhang, L.; Yang, X.; Liu, J.; Xu, M.; Shi, Z.; Hu, Z.; Zhong, W.; Xiao, G. Remdesivir and chloroquine effectively inhibit the recently emerged novel coronavirus (2019-nCoV) in vitro. *Cell Res.* **2020**, *30*, 269–271. [[CrossRef](#)] [[PubMed](#)]
- Hoffmann, M.; Kleine-Weber, H.; Schroeder, S.; Krüger, N.; Herrler, T.; Erichsen, S.; Schiergens, T.S.; Herrler, G.; Wu, N.-H.; Nitsche, A.; et al. SARS-CoV-2 Cell Entry Depends on ACE2 and TMPRSS2 and Is Blocked by a Clinically Proven Protease Inhibitor. *Cell* **2020**, *181*, 271–280.e8. [[CrossRef](#)]
- Home—ClinicalTrials.gov. Available online: <https://clinicaltrials.gov/> (accessed on 28 October 2020).
- Beigel, J.H.; Tomashek, K.M.; Dodd, L.E.; Mehta, A.K.; Zingman, B.S.; Kalil, A.C.; Hohmann, E.; Chu, H.Y.; Luetkemeyer, A.; Kline, S.; et al. Remdesivir for the Treatment of Covid-19—Final Report. *N. Engl. J. Med.* **2020**. [[CrossRef](#)]
- Boulware, D.R.; Pullen, M.F.; Bangdiwala, A.S.; Pastick, K.A.; Lofgren, S.M.; Okafor, E.C.; Skipper, C.P.; Nascene, A.A.; Nicol, M.R.; Abassi, M.; et al. A Randomized Trial of Hydroxychloroquine as Postexposure Prophylaxis for Covid-19. *N. Engl. J. Med.* **2020**, *383*, 517–525. [[CrossRef](#)]
- Geleris, J.; Sun, Y.; Platt, J.; Zucker, J.; Baldwin, M.; Hripcak, G.; Labella, A.; Manson, D.K.; Kubin, C.; Barr, R.G.; et al. Observational study of hydroxychloroquine in hospitalized patients with COVID-19. *N. Engl. J. Med.* **2020**, *382*, 2411–2418. [[CrossRef](#)] [[PubMed](#)]
- Wang, Y.; Zhang, D.; Du, G.; Du, R.; Zhao, J.; Jin, Y.; Fu, S.; Gao, L.; Cheng, Z.; Lu, Q.; et al. Remdesivir in adults with severe COVID-19: A randomised, double-blind, placebo-controlled, multicentre trial. *Lancet* **2020**, *395*, 1569–1578. [[CrossRef](#)]
- Pan, H.; Peto, R.; Karim, Q.A.; Alejandria, M.; Henao-Restrepo, A.M.; García, C.H.; Kieny, M.-P.; Malekzadeh, R.; Murthy, S.; Preziosi, M.-P.; et al. Repurposed antiviral drugs for COVID-19 –interim WHO SOLIDARITY trial results. *MedRxiv* **2020**. [[CrossRef](#)]
- Smith, E.C.; Blanc, H.; Vignuzzi, M.; Denison, M.R. Coronaviruses Lacking Exoribonuclease Activity Are Susceptible to Lethal Mutagenesis: Evidence for Proofreading and Potential Therapeutics. *PLoS Pathog.* **2013**, *9*, e1003565. [[CrossRef](#)]
- Lundin, A.; Dijkman, R.; Bergström, T.; Kann, N.; Adamiak, B.; Hannoun, C.; Kindler, E.; Jónsdóttir, H.R.; Muth, D.; Kint, J.; et al. Targeting Membrane-Bound Viral RNA Synthesis Reveals Potent Inhibition of Diverse Coronaviruses Including the Middle East Respiratory Syndrome Virus. *PLoS Pathog.* **2014**, *10*, e1004166. [[CrossRef](#)]
- Agostini, M.L.; Andres, E.L.; Sims, A.C.; Graham, R.L.; Sheahan, T.P.; Lu, X.; Smith, E.C.; Case, J.B.; Feng, J.Y.; Jordan, R.; et al. Coronavirus susceptibility to the antiviral remdesivir (GS-5734) is mediated by the viral polymerase and the proofreading exoribonuclease. *MBio* **2018**, *9*, 1–15. [[CrossRef](#)]
- Bloom, D.E.; Cadarette, D. Infectious Disease Threats in the Twenty-First Century: Strengthening the Global Response. *Front. Immunol.* **2019**, *10*, 549. [[CrossRef](#)]

20. WHO Director-General's Opening Remarks at the Media Briefing on COVID-19—11 March 2020. Available online: <https://www.who.int/dg/speeches/detail/who-director-general-s-opening-remarks-at-the-media-briefing-on-covid-19---11-march-2020> (accessed on 18 November 2020).
21. WHO | World Now at the Start of 2009 Influenza Pandemic. Available online: [https://www.who.int/mediacentre/news/statements/2009/h1n1\\_pandemic\\_phase6\\_20090611/en/](https://www.who.int/mediacentre/news/statements/2009/h1n1_pandemic_phase6_20090611/en/) (accessed on 18 November 2020).
22. Thao, T.T.N.; Labroussaa, F.; Ebert, N.; V'kovski, P.; Stalder, H.; Portmann, J.; Kelly, J.; Steiner, S.; Holwerda, M.; Kratze, A.; et al. 9 Rapid reconstruction of SARS-CoV-2 using a synthetic genomics platform. *Nature* **2020**. [[CrossRef](#)]
23. Mutterer, J.; Zinck, E. Quick-and-clean article figures with FigureJ. *J. Microsc.* **2013**, *252*, 89–91. [[CrossRef](#)]
24. Rappe, J.C.F.; de Wilde, A.; Di, H.; Müller, C.; Stalder, H.; V'kovski, P.; Snijder, E.; Brinton, M.A.; Ziebuhr, J.; Ruggli, N.; et al. Antiviral activity of K22 against members of the order Nidovirales. *Virus Res.* **2018**, *246*, 28–34. [[CrossRef](#)]
25. García-Nicolás, O.; V'kovski, P.; Vielle, N.J.; Ebert, N.; Züst, R.; Portmann, J.; Stalder, H.; Gaschen, V.; Vieyres, G.; Stoffel, M.; et al. The small-compound inhibitor K22 displays broad antiviral activity against different members of the family Flaviviridae and offers potential as a panviral inhibitor. *Antimicrob. Agents Chemother.* **2018**, *62*, 1–17. [[CrossRef](#)] [[PubMed](#)]
26. Hoffmann, M.; Mösbauer, K.; Hofmann-Winkler, H.; Kaul, A.; Kleine-Weber, H.; Krüger, N.; Gassen, N.C.; Müller, M.A.; Drosten, C.; Pöhlmann, S. Chloroquine does not inhibit infection of human lung cells with SARS-CoV-2. *Nature* **2020**, *585*, 588–590. [[CrossRef](#)] [[PubMed](#)]
27. Pruijssers, A.J.; George, A.S.; Schäfer, A.; Leist, S.R.; Gralinski, L.E.; Dinnon, K.H.; Yount, B.L.; Agostini, M.L.; Stevens, L.J.; Chappell, J.D.; et al. Remdesivir Inhibits SARS-CoV-2 in Human Lung Cells and Chimeric SARS-CoV Expressing the SARS-CoV-2 RNA Polymerase in Mice. *Cell Rep.* **2020**, *32*, 107940. [[CrossRef](#)] [[PubMed](#)]
28. Touret, F.; Gilles, M.; Barral, K.; Nougairède, A.; van Helden, J.; Decroly, E.; de Lamballerie, X.; Coutard, B. In vitro screening of a FDA approved chemical library reveals potential inhibitors of SARS-CoV-2 replication. *Sci. Rep.* **2020**, *10*, 1–8. [[CrossRef](#)] [[PubMed](#)]
29. Riva, L.; Yuan, S.; Yin, X.; Martin-Sancho, L.; Matsunaga, N.; Pache, L.; Burgstaller-Muehlbacher, S.; De Jesus, P.D.; Teriete, P.; Hull, M.V.; et al. Discovery of SARS-CoV-2 antiviral drugs through large-scale compound repurposing. *Nature* **2020**, *586*, 113–119. [[CrossRef](#)]
30. Rosenke, K.; Jarvis, M.; Feldmann, F.; Schwarz, B.; Okumura, A.; Lovaglio, J.; Saturday, G.; Hanley, P.; Meade-White, K.; Williamson, B.; et al. Hydroxychloroquine Proves Ineffective in Hamsters and Macaques Infected with SARS-CoV-2. *BioRxiv Prepr. Serv. Biol.* **2020**. [[CrossRef](#)]
31. Dai, W.; Wu, Y.; Bi, J.; Lu, X.; Hou, A.; Zhou, Y.; Sun, B.; Kong, W.; Barbier, J.; Cintrat, J.C.; et al. Antiviral effects of Retro-2cycl and Retro-2.1 against Enterovirus 71 in vitro and in vivo. *Antivir. Res.* **2017**, *144*, 311–321. [[CrossRef](#)]
32. Shtanko, O.; Sakurai, Y.; Reyes, A.N.; Noël, R.; Cintrat, J.C.; Gillet, D.; Barbier, J.; Davey, R.A. Retro-2 and its dihydroquinazolinone derivatives inhibit filovirus infection. *Antivir. Res.* **2018**, *149*, 154–163. [[CrossRef](#)]
33. Dai, W.; Wu, Y.; Bi, J.; Wang, J.; Wang, S.; Kong, W.; Barbier, J.; Cintrat, J.C.; Gao, F.; Jiang, Z.; et al. Antiviral effect of Retro-2.1 against herpes simplex Virus type 2 in Vitro. *J. Microbiol. Biotechnol.* **2018**, *28*, 849–859. [[CrossRef](#)]
34. Harrison, K.; Haga, I.R.; Pechenick Jowers, T.; Jasim, S.; Cintrat, J.-C.; Gillet, D.; Schmitt-John, T.; Digard, P.; Beard, P.M. Vaccinia Virus Uses Retromer-Independent Cellular Retrograde Transport Pathways To Facilitate the Wrapping of Intracellular Mature Virions during Virus Morphogenesis. *J. Virol.* **2016**, *90*, 10120–10132. [[CrossRef](#)]
35. Maru, S.; Jin, G.; Desai, D.; Amin, S.; Shwetank; Lauver, M.D.; Lukacher, A.E. Inhibition of Retrograde Transport Limits Polyomavirus Infection In Vivo. *mSphere* **2017**, *2*. [[CrossRef](#)] [[PubMed](#)]
36. Desai, D.; Lauver, M.; Ostman, A.; Cruz, L.; Ferguson, K.; Jin, G.; Roper, B.; Brosius, D.; Lukacher, A.; Amin, S.; et al. Inhibition of diverse opportunistic viruses by structurally optimized retrograde trafficking inhibitors. *Bioorganic Med. Chem.* **2019**, *27*, 1795–1803. [[CrossRef](#)] [[PubMed](#)]
37. Gupta, N.; Pons, V.; Noël, R.; Buisson, D.A.; Michau, A.; Johannes, L.; Gillet, D.; Barbier, J.; Cintrat, J.C. (S)-N-methyldihydroquinazolinones are the active enantiomers of retro-2 derived compounds against toxins. *ACS Med. Chem. Lett.* **2014**, *5*, 94–97. [[CrossRef](#)] [[PubMed](#)]

38. Stechmann, B.; Bai, S.K.; Gobbo, E.; Lopez, R.; Merer, G.; Pinchard, S.; Panigai, L.; Tenza, D.; Raposo, G.; Beaumelle, B.; et al. Inhibition of retrograde transport protects mice from lethal ricin challenge. *Cell* **2010**, *141*, 231–242. [CrossRef] [PubMed]
39. V'kovski, P.; Gerber, M.; Kelly, J.; Pfaender, S.; Ebert, N.; Braga Lagache, S.; Simillion, C.; Portmann, J.; Stalder, H.; Gaschen, V.; et al. Determination of host proteins composing the microenvironment of coronavirus replicase complexes by proximity-labeling. *Elife* **2019**, *8*, 1–30. [CrossRef] [PubMed]
40. Dai, W.; Wu, Y.; Bi, J.; Wang, S.; Li, F.; Kong, W.; Barbier, J.; Cintrat, J.C.; Gao, F.; Gillet, D.; et al. Antiviral effects of abma against herpes simplex virus type 2 in vitro and in vivo. *Viruses* **2018**, *10*, 1–15. [CrossRef]
41. Barnard, D.L.; Day, C.W.; Bailey, K.; Heiner, M.; Montgomery, R.; Lauridsen, L.; Chan, P.K.S.; Sidwell, R.W. Evaluation of immunomodulators, interferons and known in vitro SARS-CoV inhibitors for inhibition of SARS-CoV replication in BALB/c mice. *Antivir. Chem. Chemother.* **2006**, *17*, 275–284. [CrossRef]
42. Mizui, T.; Yamashina, S.; Tanida, I.; Takei, Y.; Ueno, T.; Sakamoto, N.; Ikejima, K.; Kitamura, T.; Enomoto, N.; Sakai, T.; et al. Inhibition of hepatitis C virus replication by chloroquine targeting virus-associated autophagy. *J. Gastroenterol.* **2010**, *45*, 195–203. [CrossRef]
43. Eng, E.O.; Chew, J.S.W.; Jin, P.L.; Chua, R.C.S. In vitro inhibition of human influenza A virus replication by chloroquine. *Virol. J.* **2006**, *3*, 3–5.
44. Yan, Y.; Zou, Z.; Sun, Y.; Li, X.; Xu, K.F.; Wei, Y.; Jin, N.; Jiang, C. Anti-malaria drug chloroquine is highly effective in treating avian influenza A H5N1 virus infection in an animal model. *Cell Res.* **2013**, *23*, 300–302. [CrossRef]
45. Paton, N.I.; Lee, L.; Xu, Y.; Ooi, E.E.; Cheung, Y.B.; Archuleta, S.; Wong, G.; Smith, A.W. Chloroquine for influenza prevention: A randomised, double-blind, placebo controlled trial. *Lancet Infect. Dis.* **2011**, *11*, 677–683. [CrossRef]
46. De Lamballerie, X.; Boisson, V.; Reynier, J.C.; Enault, S.; Charrel, R.N.; Flahault, A.; Roques, P.; Grand, R. Le On chikungunya acute infection and chloroquine treatment. *Vector Borne Zoonotic Dis.* **2008**, *8*, 837–839. [CrossRef] [PubMed]
47. Khan, M.; Santhosh, S.R.; Tiwari, M.; Lakshmana Rao, P.V.; Parida, M. Assessment of in vitro prophylactic and therapeutic efficacy of chloroquine against Chikungunya virus in Vero cells. *J. Med. Viro.* **2010**, *82*, 817–824. [CrossRef] [PubMed]
48. Farias, K.J.S.; Machado, P.R.L.; de Almeida Junior, R.F.; de Aquino, A.A.; da Fonseca, B.A.L. Chloroquine interferes with dengue-2 virus replication in U937 cells. *Microbiol. Immunol.* **2014**, *58*, 318–326. [CrossRef] [PubMed]
49. Delvecchio, R.; Higa, L.M.; Pezzuto, P.; Valadão, A.L.; Garcez, P.P.; Monteiro, F.L.; Loiola, E.C.; Dias, A.A.; Silva, F.J.M.; Aliota, M.T.; et al. Chloroquine, an endocytosis blocking agent, inhibits zika virus infection in different cell models. *Viruses* **2016**, *8*, 1–15. [CrossRef] [PubMed]
50. Porotto, M.; Orefice, G.; Yokoyama, C.C.; Mungall, B.A.; Realubit, R.; Sganga, M.L.; Aljofan, M.; Whitt, M.; Glickman, F.; Moscona, A. Simulating Henipavirus Multicycle Replication in a Screening Assay Leads to Identification of a Promising Candidate for Therapy. *J. Virol.* **2009**, *83*, 5148–5155. [CrossRef]
51. Freiberg, A.N.; Worthy, M.N.; Lee, B.; Holbrook, M.R. Combined chloroquine and ribavirin treatment does not prevent death in a hamster model of Nipah and Hendra virus infection. *J. Gen. Virol.* **2010**, *91*, 765–772. [CrossRef]
52. Kono, M.; Tatsumi, K.; Imai, A.M.; Saito, K.; Kuriyama, T.; Shirasawa, H. Inhibition of human coronavirus 229E infection in human epithelial lung cells (L132) by chloroquine: Involvement of p38 MAPK and ERK. *Antivir. Res.* **2008**, *77*, 150–152. [CrossRef]
53. Ferraris, O.; Moroso, M.; Pernet, O.; Emonet, S.; Ferrier Rembert, A.; Paranhos-Baccalà, G.; Peyrefitte, C.N. Evaluation of Crimean-Congo hemorrhagic fever virus in vitro inhibition by chloroquine and chlorpromazine, two FDA approved molecules. *Antivir. Res.* **2015**, *118*, 75–81. [CrossRef]
54. Dowall, S.D.; Bosworth, A.; Watson, R.; Bewley, K.; Taylor, I.; Rayner, E.; Hunter, L.; Pearson, G.; Easterbrook, L.; Pitman, J.; et al. Chloroquine inhibited ebola virus replication in vitro but failed to protect against infection and disease in the in vivo guinea pig model. *J. Gen. Virol.* **2015**, *96*, 3484–3492. [CrossRef]
55. Feng, Z.; Hensley, L.; McKnight, K.L.; Hu, F.; Madden, V.; Ping, L.; Jeong, S.H.; Walker, C.; Lanford, R.E.; Lemon, S.M. A pathogenic picornavirus acquires an envelope by hijacking cellular membranes. *Nature* **2013**, *496*, 367–371. [CrossRef] [PubMed]
56. Lecot, S.; Belouzard, S.; Dubuisson, J.; Rouillé, Y. Bovine Viral Diarrhea Virus Entry is Dependent on Clathrin-Mediated Endocytosis. *J. Virol.* **2005**, *79*, 10826–10829. [CrossRef] [PubMed]

57. Vincent, M.J.; Bergeron, E.; Benjannet, S.; Erickson, B.R.; Rollin, P.E.; Ksiazek, T.G.; Seidah, N.G.; Nichol, S.T. Chloroquine is a potent inhibitor of SARS coronavirus infection and spread. *Virol. J.* **2005**, *2*, 1–10. [CrossRef] [PubMed]
58. Blanchard, E.; Belouzard, S.; Goueslain, L.; Wakita, T.; Dubuisson, J.; Wychowski, C.; Rouillé, Y. Hepatitis C Virus Entry Depends on Clathrin-Mediated Endocytosis. *J. Virol.* **2006**, *80*, 6964–6972. [CrossRef]
59. Tsiang, H.; Superti, F. Ammonium chloride and chloroquine inhibit rabies virus infection in neuroblastoma cells. *Arch. Virol.* **1984**, *81*, 377–382. [CrossRef]
60. Kronenberger, P.; Vrijen, R.; Boeyé, A. Chloroquine induces empty capsid formation during poliovirus eclipse. *J. Virol.* **1991**, *65*, 7008–7011. [CrossRef]
61. Romanello, F.; Smith, K.; Hoven, A. Chloroquine and Hydroxychloroquine as Inhibitors of Human Immunodeficiency Virus (HIV-1) Activity. *Curr. Pharm. Des.* **2005**, *10*, 2643–2648. [CrossRef]
62. Bishop, N.E. Examination of potential inhibitors of hepatitis A virus uncoating. *Intervirology* **1998**, *41*, 261–271. [CrossRef]
63. Mehta, A.; Ouzounov, S.; Jordan, R.; Simsek, E.; Lu, X.; Moriarty, R.M.; Jacob, G.; Dwek, R.A.; Block, T.M. Imino sugars that are less toxic but more potent as antivirals, in vitro, compared with N-n-nonyl DNJ. *Antivir. Chem. Chemother.* **2002**, *13*, 299–304. [CrossRef]
64. Wu, S.-F.; Lee, C.-J.; Liao, C.-L.; Dwek, R.A.; Zitzmann, N.; Lin, Y.-L. Antiviral Effects of an Iminosugar Derivative on Flavivirus Infections. *J. Virol.* **2002**, *76*, 3596–3604. [CrossRef]
65. Zitzmann, N.; Mehta, A.S.; Carrouée, S.; Butters, T.D.; Platt, F.M.; McCauley, J.; Blumberg, B.S.; Dwek, R.A.; Block, T.M. Imino sugars inhibit the formation and secretion of bovine viral diarrhea virus, a pestivirus model of hepatitis C virus: Implications for the development of broad spectrum anti-hepatitis virus agents. *Proc. Natl. Acad. Sci. USA* **1999**, *96*, 11878–11882. [CrossRef] [PubMed]
66. Steinmann, E.; Whitfield, T.; Kallis, S.; Dwek, R.A.; Zitzmann, N.; Pietschmann, T.; Bartenschlager, R. Antiviral effects of amantadine and iminosugar derivatives against hepatitis C virus. *Hepatology* **2007**, *46*, 330–338. [CrossRef] [PubMed]
67. Block, T.M.; Lu, X.; Mehta, A.S.; Blumberg, B.S.; Tennant, B.; Ebling, M.; Korba, B.; Lansky, D.M.; Jacob, G.S.; Dwek, R.A. Treatment of chronic hepadnavirus infection in a woodchuck animal model with an inhibitor of protein folding and trafficking. *Nat. Med.* **1998**, *4*, 610–614. [CrossRef] [PubMed]
68. Gu, B.; Mason, P.; Wang, L.; Norton, P.; Bourne, N.; Moriarty, R.; Mehta, A.; Despande, M.; Shah, R.; Block, T. Antiviral profiles of novel iminocyclitol compounds against bovine viral diarrhea virus, West Nile virus, dengue virus and hepatitis B virus. *Antivir. Chem. Chemother.* **2007**, *18*, 49–59. [CrossRef] [PubMed]
69. Chang, J.; Wang, L.; Ma, D.; Qu, X.; Guo, H.; Xu, X.; Mason, P.M.; Bourne, N.; Moriarty, R.; Gu, B.; et al. Novel imino sugar derivatives demonstrate potent antiviral activity against flaviviruses. *Antimicrob. Agents Chemother.* **2009**, *53*, 1501–1508. [CrossRef] [PubMed]
70. Qu, X.; Pan, X.; Weidner, J.; Yu, W.; Alonzi, D.; Xu, X.; Butters, T.; Block, T.; Guo, J.T.; Chang, J. Inhibitors of endoplasmic reticulum  $\alpha$ -glucosidases potently suppress hepatitis C virus virion assembly and release. *Antimicrob. Agents Chemother.* **2011**, *55*, 1036–1044. [CrossRef]
71. Fukushi, M.; Yoshinaka, Y.; Matsuoka, Y.; Hatakeyama, S.; Ishizaka, Y.; Kirikae, T.; Sasazuki, T.; Miyoshi-Akiyama, T. Monitoring of S Protein Maturation in the Endoplasmic Reticulum by Calnexin is Important for the Infectivity of Severe Acute Respiratory Syndrome Coronavirus. *J. Virol.* **2012**, *86*, 11745–11753. [CrossRef]
72. Goodfellow, V.S.; Loweth, C.J.; Ravula, S.B.; Wiemann, T.; Nguyen, T.; Xu, Y.; Todd, D.E.; Sheppard, D.; Pollack, S.; Polesskaya, O.; et al. Discovery, synthesis, and characterization of an orally bioavailable, brain penetrant inhibitor of mixed lineage kinase 3. *J. Med. Chem.* **2013**, *56*, 8032–8048. [CrossRef]
73. Marker, D.F.; Tremblay, M.È.; Puccini, J.M.; Barbieri, J.; Gantz Marker, M.A.; Loweth, C.J.; Chris Muly, E.; Lu, S.M.; Goodfellow, V.S.; Dewhurst, S.; et al. The new small-molecule mixed-lineage kinase 3 inhibitor URMC-099 is neuroprotective and anti-inflammatory in models of human immunodeficiency virus-associated neurocognitive disorders. *J. Neurosci.* **2013**, *33*, 9998–10010. [CrossRef]
74. Zhang, G.; Guo, D.; Dash, P.K.; Araíga, M.; Wiederin, J.L.; Haverland, N.A.; Knibbe-Hollinger, J.; Martinez-Skinner, A.; Ciborowski, P.; Goodfellow, V.S.; et al. The mixed lineage kinase-3 inhibitor URMC-099 improves therapeutic outcomes for long-acting antiretroviral therapy. *Nanomed. Nanotechnol. Biol. Med.* **2016**, *12*, 109–122. [CrossRef]

75. Xu, H.; Cheng, M.; Chi, X.; Liu, X.; Zhou, J.; Lin, T.; Yang, W. High-Throughput Screening Identifies Mixed-Lineage Kinase 3 as a Key Host Regulatory Factor in Zika Virus Infection. *J. Virol.* **2019**, *93*, 1–16. [[CrossRef](#)]
76. Warren, T.K.; Jordan, R.; Lo, M.K.; Ray, A.S.; Mackman, R.L.; Soloveva, V.; Siegel, D.; Perron, M.; Bannister, R.; Hui, H.C.; et al. Therapeutic efficacy of the small molecule GS-5734 against Ebola virus in rhesus monkeys. *Nature* **2016**, *531*, 381–385. [[CrossRef](#)] [[PubMed](#)]
77. Lo, M.K.; Jordan, R.; Arvey, A.; Sudhamsu, J.; Shrivastava-Ranjan, P.; Hotard, A.L.; Flint, M.; McMullan, L.K.; Siegel, D.; Clarke, M.O.; et al. GS-5734 and its parent nucleoside analog inhibit Filo-, Pneumo-, and Paramyxoviruses. *Sci. Rep.* **2017**, *7*, 1–7. [[CrossRef](#)] [[PubMed](#)]
78. Sheahan, T.P.; Sims, A.C.; Leist, S.R.; Schäfer, A.; Won, J.; Brown, A.J.; Montgomery, S.A.; Hogg, A.; Babusis, D.; Clarke, M.O.; et al. Comparative therapeutic efficacy of remdesivir and combination lopinavir, ritonavir, and interferon beta against MERS-CoV. *Nat. Commun.* **2020**, *11*, 1–14. [[CrossRef](#)] [[PubMed](#)]
79. Sheahan, T.P.; Sims, A.C.; Zhou, S.; Graham, R.L.; Pruijssers, A.J.; Agostini, M.L.; Leist, S.R.; Schäfer, A.; Dinnon, K.H.; Stevens, L.J.; et al. An orally bioavailable broad-spectrum antiviral inhibits SARS-CoV-2 in human airway epithelial cell cultures and multiple coronaviruses in mice. *Sci. Transl. Med.* **2020**, *12*. [[CrossRef](#)]
80. Sheahan, T.P.; Sims, A.C.; Graham, R.L.; Menachery, V.D.; Gralinski, L.E.; Case, J.B.; Leist, S.R.; Pyrc, K.; Feng, J.Y.; Trantcheva, I.; et al. Broad-spectrum antiviral GS-5734 inhibits both epidemic and zoonotic coronaviruses. *Sci. Transl. Med.* **2017**, *9*. [[CrossRef](#)]

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