Discovery and Nanosized Preparations of (S,R)-Tylophorine Malate as Novel anti-SARS-CoV-2 Agents

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ABSTRACT: The coronavirus disease 2019 (COVID-19) pandemic caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) has stimulated the search for effective drugs for its prevention and treatment. Natural products are an important source for new drug discovery. Here, we report that NK007 (S,R), a tylophorine malate, displays high antiviral activity against SARS-CoV-2 with an EC\textsubscript{50} 0.03 \textmu M \textit{in vitro}, which is substantially lower than that of remdesivir (EC\textsubscript{50}: 0.8 \textmu M \textit{in vitro}), the only authorized drug to date. The histopathological research revealed that NK007(S,R) (5 mg/kg/dose) displayed a protection effect in lung injury induced by SARS-CoV-2, which is better than remdesivir (25 mg/kg/dose). We also prepared two nanosized preparations of NK007(S,R), which also showed good efficacy (EC\textsubscript{50}: NP-NK007, 0.007 \textmu M \textit{in vitro}; LP-NK007, 0.014 \textmu M \textit{in vitro}). Our findings suggest that tylophorine alkaloids, isolated from the traditional Chinese medicine Cynanchum komarovii AL, offer a new skeleton for the development of anticoronavirus drug candidate.

KEYWORDS: Coronavirus, COVID-19, SARS-CoV-2, Traditional Chinese medicine, Tylophorine, Nanosized preparation

As of September 1, 2021, there have been 217,558,771 reported cases of coronavirus disease 2019 (COVID-19), which is caused by severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2), and the disease has led to 4,517,240 deaths in more than 200 countries.\textsuperscript{1,2} Kissler and co-workers reported that one-off blocking measures cannot prevent the spread of SARS-CoV-2, and the United States may need to continue social isolation measures until 2022.\textsuperscript{3} So far, there are currently no specific drugs or vaccines available for the treatment of SARS-CoV-2-infected patients, the only authorized drug by the United States Food and Drug Administration is the nucleotide analogue remdesivir.\textsuperscript{4,5} Related studies show that remdesivir has a strong resistance to SARS-CoV-2 \textit{in vitro}.\textsuperscript{6} The COVID-19 pandemic has stimulated the search for effective drugs for prevention and treatment of the disease.

The response to a viral pandemic requires the simultaneous implementation of short-term measures and long-term planning. The use of existing drugs is the focus of short-term measures, and indeed a series of first-line drugs that have been found to have potential utility for treatment of COVID-19 are currently being explored, including ritonavir and lopinavir,\textsuperscript{7,8} ribavirin,\textsuperscript{9} chloroquine and hydroxychloroquine,\textsuperscript{10} and favipiravir and remdesivir.\textsuperscript{11} Long-term planning focuses on the identification of druggable targets and the discovery and development of new drug molecules. Progress toward target-based design of new drugs against COVID-19 will undoubtedly be facilitated by the recent determination of the structures of the main protease (M\textsubscript{pro})\textsuperscript{12} and the RNA-dependent RNA polymerase\textsuperscript{13} from SARS-CoV-2. The advantages of target-based drug design are the clear target and mechanism and high selectivity, but this approach is subject to the development of drug resistance.

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An alternative approach is to design new drugs based on natural products, which are excellent sources of molecules with novel chemical structures and unique mechanisms of action. Natural products often act on multiple targets, which tends to hinder the development of drug resistance. However, this approach is relatively labor-intensive, and the determining the mechanism of action of natural products can be difficult and time-consuming. Nevertheless, they have been an important source of new drugs14−16 and may be useful for the identification of molecules with activity against SARS-CoV-2.

SARS-CoV-2 is an enveloped single-stranded RNA virus. The spike protein on the viral envelope binds to the ACE2 receptors of host cells, and a viral protease can assist virus invasion. After the virus enters the cell, its genes are released, the viral replicase and transcriptase are synthesized, and RNA replication and transcription are accomplished by an RNA-dependent RNA polymerase. Then structural proteins are synthesized, and finally new virus particles are assembled and released. All these steps of the virus life cycle are potential targets for drug therapy,6,17 and a number of small-molecule compounds showed that they have good anticancer,28 anti-HIV,29 and immunomodulatory activities.30,31 As an optimized approach is relatively labor-intensive, and the determining the mechanism of action of natural products can be difficult and time-consuming. Nevertheless, they have been an important source of new drugs14−16 and may be useful for the identification of molecules with activity against SARS-CoV-2.

NK007(S,R) was further used to study the therapy outcome in the COVID-19 rat model. A COVID-19 golden hamster rat model was generated according to a recently published paper.32 Briefly, golden hamster rats were randomly divided into four groups. Rats were then infected intranasally with SARS-CoV-2 (1 × 105 PFU). One hour after SARS-CoV-2 infection, rats in the NS (normal saline, 0.9% NaCl) group, the remdesivir group, and the NK007(S,R) group received normal saline, remdesivir, or NK007(S,R) intranasally. The Mock group is mock-infected rats. After virus infection at day 0, rats received NK007(S,R) or remdesivir or NS treatment 3 times as shown in Figure 4A. All rats were euthanized at day 4, and several parameters were measured, including body weight loss, SARS-CoV-2 RNA copies in the lungs, and pathological change in lung tissues.

NS-treated COVID-19 hamsters had a significant body weight loss compared with Mock hamsters on day 4 post infection (p value = 0.0093), while NK007(S,R) or remdesivir treated hamsters were with no significant body weight loss when compared to the Mock hamsters (Figure 4B). We found that the average number of viral RNA copies in the lungs of NK007(S,R) (∼10^5) or remdesivir (∼10^4) treated rats have

**Figure 1.** Discovery of NK007(S,R).

**Figure 2.** Determination of the EC_{50} values of remdesivir and NK007(S,R).

**Figure 3.** Determination of the CC_{50} value of NK007(S,R).

**Figure 4A.** Safety test of NK007(S,R) in COVID-19 hamsters. (A) NK007(S,R) or remdesivir or NS treated rats have significant body weight loss compared with Mock hamsters on day 4 post infection (p value = 0.0093), while NK007(S,R) or remdesivir treated hamsters were with no significant body weight loss when compared to the Mock hamsters (Figure 4B). We found that the average number of viral RNA copies in the lungs of NK007(S,R) (∼10^5) or remdesivir (∼10^4) treated rats have
no significant difference with NS-treated infected rats ($\sim 10^{3.5}$) (Figure 4C). And the histopathological changes in rat lung tissues are a key index to assess the therapy effects of NK007(S,R) or remdesivir in COVID-19 golden hamster rats. The lungs of rats were assessed by grading the injuries in accordance with the International Harmonization of Nomenclature and Diagnostic Criteria (INHAND) scoring standard. As shown in Figure 4D,E, the average histopathological score of virus-infected rats in NS group was approximately 3, along with an infiltration of lymphocytes and neutrophils, and the alveolar septum, bronchus, and perivascular interstitium were significantly widened. For rats infected by SARS-CoV-2, treatment of remdesivir at a dose of 25 mg/kg got histopathological scores, approximately 2.7 where the alveolar septum, bronchus, and perivascular interstitium were obviously widened, along with an infiltration of some lymphocytes and neutrophils. Treatment of NK007(S,R) at a dose of 5 mg/kg got pathological scores, approximately 2.0. NK007(S,R) treatment significantly decreased lung inflammation in SARS-CoV-2 infected rats. The local alveolar septum, bronchi, and
perivascular interstitial widening were significantly decreased in the remdesivir treatment group compared with NS-treated COVID-19 hamsters. And the NK007(S,R) treated group showed the lowest lung inflammation. According to histopathological results, NK007(S,R) shows protection from lung injury induced by SARS-CoV-2 infection, and the therapy outcome of NK007(S,R) at a dose of 5 mg/kg is better than that of remdesivir at a dose of 25 mg/kg. The good pharmacological activity of NK007(S,R) lays a foundation for its application in the treatment of COVID-19.

We further developed two nanosized preparations of NK007(S,R) and evaluated their anti-SARS-CoV-2 activities in vitro. The self-assembled PEG−PLGA nanoparticle loaded with NK007(S,R) (NP-NK007) was prepared using a double-emulsion (W1/O/W2) solvent evaporation method (Figure 5A). The mean particle size of NP-NK007 was 145.8 ± 7.4 nm with a PDI of 0.067 ± 0.032 (Figure 5B) and a slightly positive zeta potential of 1.83 ± 0.77 mV (Figure S4). The encapsulation efficiency and drug loading of nanoparticles was 87.47% ± 1.70 and 13.10% ± 0.61 (n = 3), respectively. The release profile of NK007(S,R) out of NP-NK007 was performed (Figure 5C). NP-NK007 released NK-007(S,R) gradually. Within a 48 h period, only 66.51% ± 2.00% of NK007(S,R) was released from NP-NK007. All the results illustrated that NK007(S,R) was efficiently entrapped in the nanoparticles and might achieve a sustained release in vivo. The biodistribution of nanoparticles was studied in healthy C57BL/6 mice. As shown in Figure 5D, NP-DiD, the self-assembled
PEG–PLGA nanoparticle loaded with a lipophilic fluorescent dye DiD prepared in the same method of NP-NK007, displayed enhanced distribution in the lung compared to the free DiD. It indicated that NP-NK007 might have the potential to achieve higher antivirus activity in vivo than NK-007(S,R) because of the improved accumulation of NK-007(S,R) delivered by the nanoparticles in the lung.

The liposome loaded with NK007(S,R) (LP-NK007) was prepared by reverse-phase evaporation method. As presented in Figure 6A, we designed another platform for lung targeted delivery of NK007(S,R) encapsulating into a lipid bilayer envelope. First, the in vivo distribution test was used to screen the lipid with the best lung targeting ability. Similarly, DiD was employed as the fluorescence probe to characterize the in vivo distribution profiles of different liposomes. As seen in Figure 6B, the relative fluorescence intensity of LP12 (DOTAP-based LP-DiD) was the strongest in the lung and was similar to free DiD in the spleen and liver. This implied that NK007(S,R) could be enriched in the lung at most when delivered by LP12 and had the least impact on other organs. Therefore, LP-NK007 containing DOTAP was selected for the subsequent experiments. The characteristics of the optimal LP-NK007 were shown in Figure 6C including EE%, DL%, particle size, and PDI. LP-NK007 had average diameters around 75 nm with narrow PDI. The encapsulation efficiency and drug loading of LP-NK007 was 62.4% and 36.7%, respectively.

Under our test conditions, the in vitro EC50 values of NP-NK007 and LP-NK007 were 0.007 and 0.014 μM respectively, and the CC50 values of NP-NK007 and LP-NK007 were 20 and 1 μM respectively, which indicated that these two nanosized preparations could be used as drug sustained-release agents.

Starting from a traditional Chinese medicine, we obtained an anticoronavirus drug candidate with a novel skeleton. The compound, a tylophorine malate designated NK007(S,R), showed substantially higher activity (EC50: 0.03 μM) against SARS-CoV-2 than remdesivir (EC50: 0.8 μM). NK007(S,R) showed protection in lung injury induced by SARS-CoV-2 infection, and the therapy outcome of NK007(S,R) at a dose of 5 mg/kg was better than that of remdesivir at a dose of 25 mg/kg. We further developed two nanosized preparations of NK007(S,R) and evaluated their anti-SARS-CoV-2 activities, which indicated that these two nanosized preparations could be used as drug sustained-release agents. The findings reported herein are encouraging, and we hope they will contribute to the ongoing fight against the COVID-19 pandemic.

### ASSOCIATED CONTENT

[Supporting Information](https://pubs.acs.org/doi/10.1021/acsmedchemlett.1c00481)

Experimental and synthetic methods, spectral characterization of compounds, tables and figures of EC50 and CC50 values, size distribution and ζ potential graphs (PDF)

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W., W.T., and X.S. conceived the project. Z.W., H.S., Y. Liu, Y. Li, and Y.Z. are responsible for proposal, compound design, synthesis and structure determination. F.Y., L.Z., B.H., and W.W. are responsible for the test of activity. Y.F. and W.X. are responsible for the preparation of two nanosized preparations of NK007(S.R). Y.D. is responsible for providing sample.

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Notes

The authors declare no competing financial interest.

ABBREVIATIONS

COVID-19, the coronavirus disease 2019; SARS-CoV-2, severe acute respiratory syndrome coronavirus; ACE2, angiotensin-converting enzyme 2; INHAND, International Harmonization of Nomenclature and Diagnostic Criteria

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