


## RESEARCH ARTICLE

# Clinical characteristics of 50 466 hospitalized patients with 2019-nCoV infection

Pengfei Sun MD<sup>1</sup> | Shuyan Qie MD<sup>2</sup> | Zongjian Liu MD<sup>2</sup> | Jizhen Ren MD<sup>3</sup> |  
Kun Li MD<sup>4</sup> | Jianing Xi MD<sup>5</sup> 

<sup>1</sup>Department of Plastic Surgery, Zibo Central Hospital, Zibo, China

<sup>2</sup>Department of Rehabilitation, Beijing Rehabilitation Hospital of Capital Medical University, Beijing, China

<sup>3</sup>Department of Plastic Surgery, Affiliated Hospital of Qingdao University, Qingdao, China

<sup>4</sup>Department of Plastic Surgery, Qingdao Eighth People's Hospital, Qingdao, China

<sup>5</sup>Department of Respiratory Rehabilitation, Beijing Rehabilitation Hospital of Capital Medical University, Beijing, China

## Correspondence

Jianing Xi, MD, Beijing Rehabilitation Hospital of Capital Medical University, Bada Road, Shijingshan District, Beijing 100044, China.  
Email: [zbzbspf@163.com](mailto:zbzbspf@163.com)

## Abstract

We aim to summarize reliable evidence of evidence-based medicine for the treatment and prevention of the 2019 novel coronavirus (2019-nCoV) by analyzing all the published studies on the clinical characteristics of patients with 2019-nCoV. PubMed, Cochrane Library, Embase, and other databases were searched. Several studies on the clinical characteristics of 2019-nCoV infection were collected for meta-analysis. Ten studies were included in the meta-analysis, including a total number of 50 466 patients with 2019-nCoV infection. Meta-analysis shows that among these patients, the incidence of fever was 89.1%, the incidence of cough was 72.2%, and the incidence of muscle soreness or fatigue was 42.5%. The incidence of acute respiratory distress syndrome (ARDS) was 14.8%, the incidence of abnormal chest computer tomography (CT) was 96.6%, the percentage of severe cases in all infected cases was 18.1%, and the case fatality rate of patients with 2019-nCoV infection was 4.3%. Fever and cough are the most common symptoms in patients with 2019-nCoV infection, and most of these patients have abnormal chest CT examination. Several people have muscle soreness or fatigue as well as ARDS. Diarrhea, hemoptysis, headache, sore throat, shock, and other symptoms only occur in a small number of patients. The case fatality rate of patients with 2019-nCoV infection is lower than that of the severe acute respiratory syndrome and Middle East respiratory syndrome.

## KEYWORDS

2019-nCoV, clinical symptoms, coronavirus, meta-analysis

## 1 | INTRODUCTION

Since December 2019, the epidemic of 2019 novel coronavirus (2019-nCoV) infectious pneumonia in Wuhan, China. The Chinese government and researchers have taken rapid measures to control the epidemic.<sup>1</sup> On January 30 2020, WHO declared that the epidemic of 2019-nCoV was a public health emergency of international concern (PHEIC). At present, the number of patients with 2019-nCoV infection is still rising, and its harm to human beings has exceeded the outbreak of severe acute respiratory syndrome (SARS) in China, 2002.<sup>2</sup> The clinical

characteristics and the case fatality rate after 2019-nCoV infection have always been concerned by people, and it is also the focus of medical workers' research at present. However, due to the different design of different clinical studies and insufficient sample size, the conclusions of the published studies were different. To acquire more accurate conclusions on the clinical characteristics and mortality of patients with 2019-nCoV infection, we searched the relevant literatures and carried out single-arm meta-analysis.<sup>3</sup> Our findings provide important guidance for current clinical work on the prevention and treatment of 2019-nCoV infection.

## 2 | MATERIAL AND METHODS

### 2.1 | Search strategy

Three popular medical databases including PubMed, Cochrane Library, and Embase databases were searched for related literature, using the following keywords: "2019-nCoV," "Coronavirus," "COVID-19," "SARS-CoV-2," and "Wuhan Coronavirus." Articles reviewed were dated up to February 2020. In this meta-analysis, there was no language restriction. Only available data from published articles were collected. Data from unpublished papers were not included.

### 2.2 | The inclusive and exclusive criteria

#### 2.2.1 | Inclusive criteria

Studies that include randomized controlled trials, nonrandomized controlled trials, case-control studies, cohort studies, cross-sectional studies, and also case reports on the clinical characteristics of

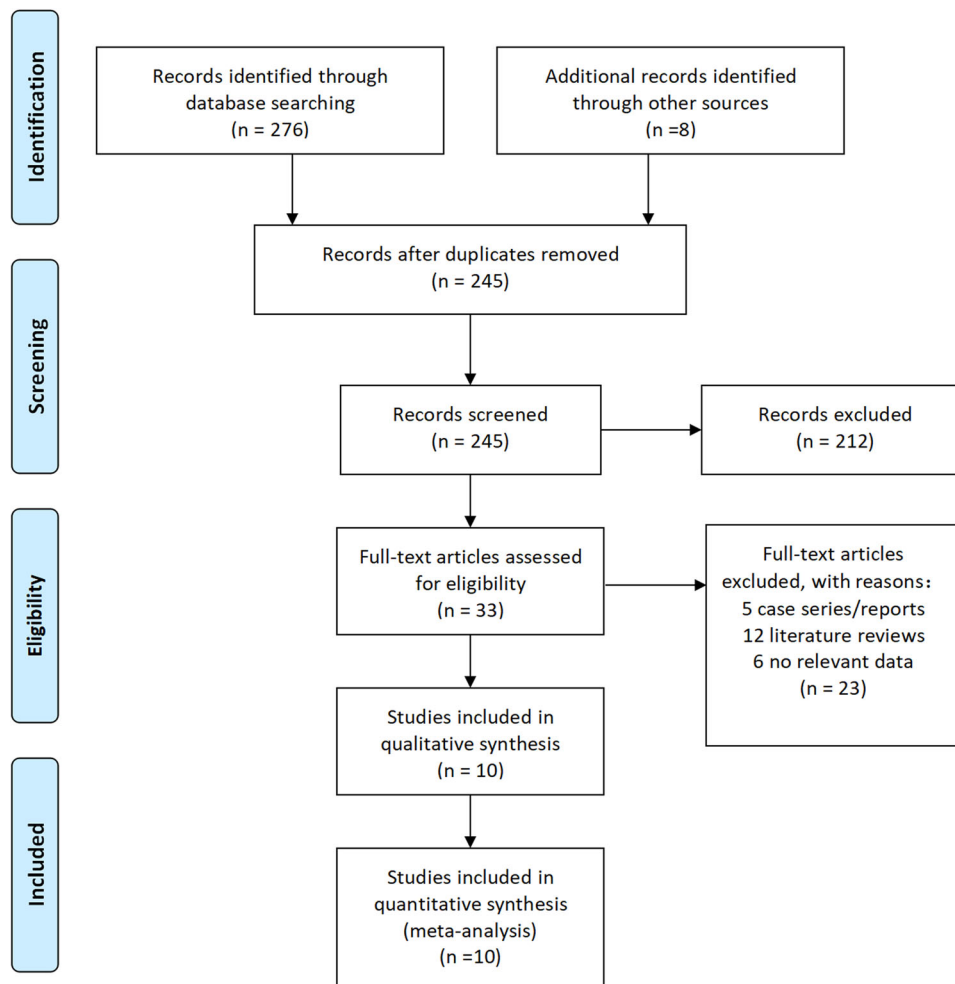
patients with 2019-nCoV infection. Clinical characteristics of patients with 2019-nCoV infection include fever (temperature  $\geq 37.3^{\circ}\text{C}$ ), cough, muscle soreness or fatigue, acute respiratory distress syndrome (ARDS), abnormal chest computer tomography (CT) detection, patients in critical condition (severe cases are those who need to be sent to the intensive care unit (ICU) for treatment), as well as death due to 2019-nCoV infection.

#### 2.2.2 | Exclusive criteria

Articles were published repeatedly; studies did not include the research indicators needed for meta-analysis; research data were missing.

### 2.3 | Data extraction and paper quality evaluation

First of all, we screened the literature according to the literature abstract, excluding the articles that obviously do not meet the

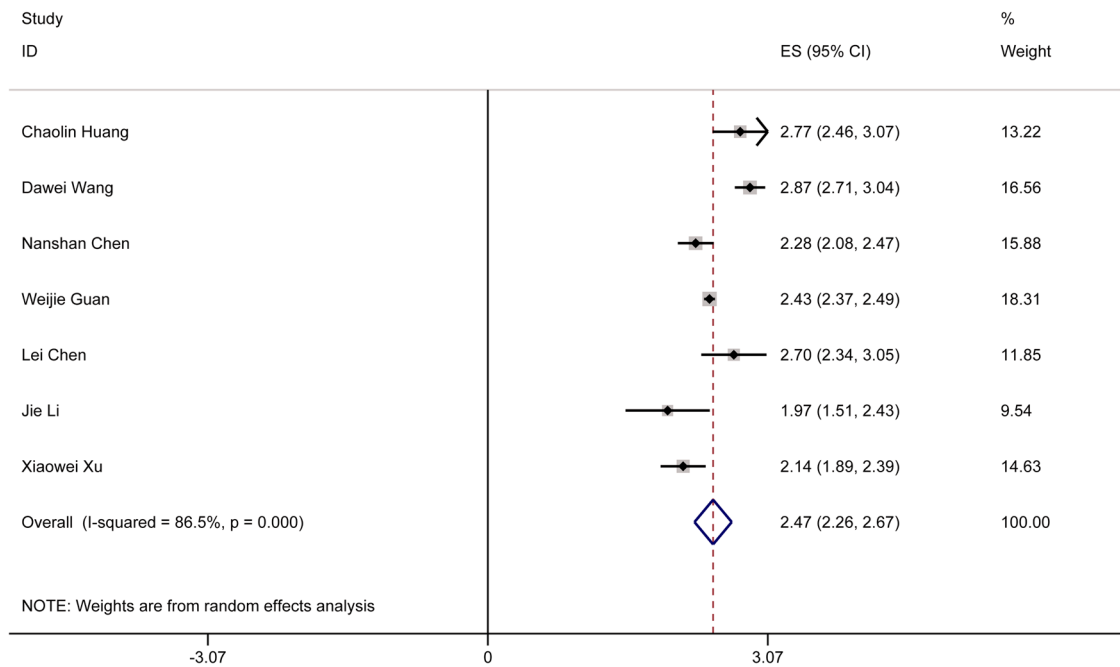


**FIGURE 1** PRISMA flow diagram

**TABLE 1** The characteristics of the literature

First author	Year	Country	Follow-up (days)	No. patients	Sex	Average age	Research type	Literature quality	Clinical symptom
Huang et al <sup>5</sup>	2020	China	18	41	Male: 30 Female: 11	49	Retrospective study	7	Fever ARDS Muscle soreness or fatigue Cough Abnormal chest CT Patient in critical condition Death of patient
Wang et al <sup>6</sup>	2020	China	34	138	Male: 75 Female: 63	56	Retrospective study	7	Fever ARDS Muscle soreness or fatigue Cough Abnormal chest CT Patient in critical condition Death of patient
Chen et al <sup>7</sup>	2020	China	25	99	Male: 67 Female: 32	55.5	Retrospective study	6	Fever ARDS Muscle soreness or fatigue Cough Abnormal chest CT Death of patient
Guan W-j <sup>8</sup>	2020	China	28	1099	Male: 640 Female: 459	47	Retrospective study	8	Fever ARDS Muscle soreness or fatigue Cough Abnormal chest CT Patient in critical condition Death of patient
Chen et al <sup>9</sup>	2020	China	15	29	Male: 21 Female: 8	56	Retrospective study	6	Fever Muscle soreness or fatigue Cough Patient in critical condition Death of patient
Sun et al <sup>10</sup>	2020	America	42	288	Male: 182 Female: 106	44	Retrospective study	5	Death of patient
Yang et al <sup>11</sup>	2020	China	51	4021	Male: 2211 Female: 1810	49	Retrospective study	5	Patient in critical condition Death of patient
Li et al <sup>12</sup>	2020	China	21	17	Male: 9 Female: 8	45	Retrospective study	6	Fever Muscle soreness or fatigue Cough Abnormal chest CT
China CDC <sup>13</sup>	2020	China	43	44 672	Male: 22 981 Female: 21 691	-	Retrospective study	6	Patient in critical condition Death of patient
Xu <sup>14</sup>	2020	China	16	62	Male: 36 Female: 26	41	Retrospective study	6	Fever Muscle soreness or fatigue Cough Abnormal chest CT Patient in critical condition Death of patient

Abbreviation: China CDC, Chinese Center for Disease Control and Prevention.



**FIGURE 2** Meta-analysis of the incidence of fever

inclusive criteria, and then read the full article for rescreening. If any disagreement on the choice of the literature exists, a third evaluator will join to make the decision. All included literature were evaluated using the Newcastle-Ottawa Scale (NOS).<sup>4</sup> The highest quality of the literature was 9 stars and the lowest 0 stars.

## 2.4 | Statistical analysis

The statistical software Stata version 12.0 was used to carry out the single-arm meta-analysis. Original data included in the literature were first transformed by double arcsine method to make them conform to the normal distribution and then analyzed in Stata. The initial conclusion obtained by meta-analysis was then restored using formula  $(P = (\sin(tp/2))^2)$  to reach final conclusion. To objectively evaluate the publication bias of the included literature, the Egger test with  $P < .05$  as the existence of publication bias was performed, the values larger than which were considered as no publication bias.

## 3 | RESULTS

### 3.1 | Literature inclusion

A total of 284 articles were retrieved, among which 39 papers were removed due to repeated retrieval, 212 papers were removed after reading abstracts, and 23 were eliminated after reading the full text. At the end, a total of 10 articles<sup>5-14</sup> were included in this meta-analysis, including data from 50 466 patients. The specific operation

flow is shown in Figure 1. The characteristics of the literature are shown in Table 1.

### 3.2 | Meta-analysis results

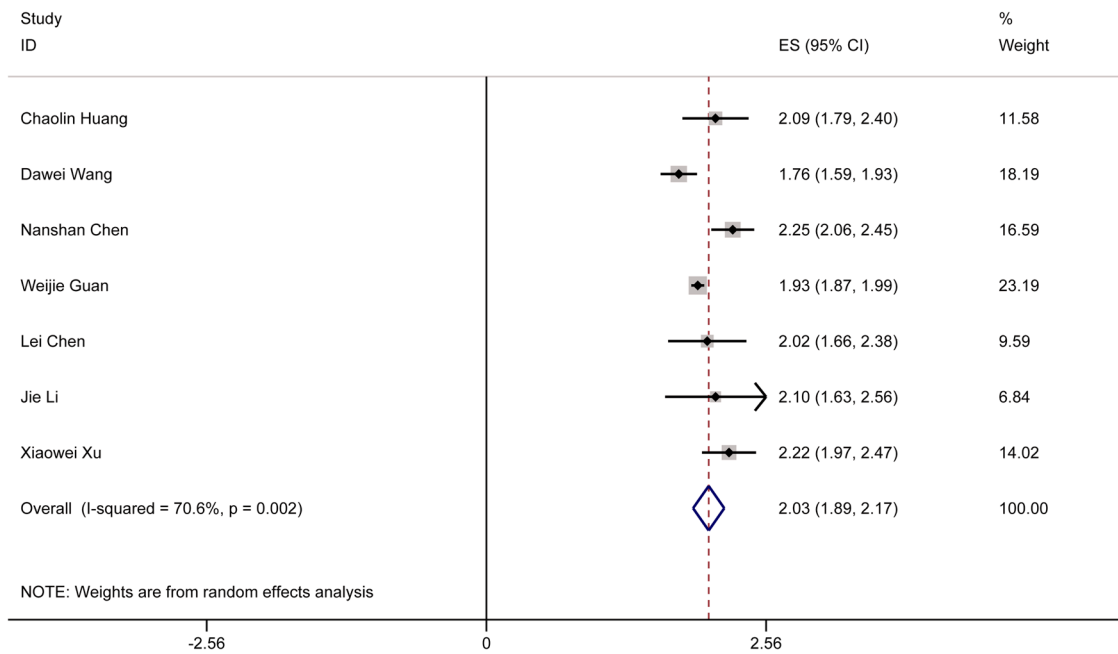
Through meta-analysis, we found that among all the clinical characteristics of patients with 2019-nCoV infection, the incidence of fever was 89.1% (Figure 2), that of cough was 72.2% (Figure 3), and that of muscle soreness or fatigue was 42.5% (Figure 4). The incidence of ARDS was 14.8% (Figure 5), correspondingly that of abnormal chest CT detection was 96.6% (Figure 6), severe cases in all infected cases occupied a percentage of 18.1% (Figure 7). The case fatality rate of patients with 2019-nCoV infection was 4.3% (Figure 8). Detailed results of the meta-analysis are shown in Table 2.

### 3.3 | Publication bias detection

The results of the Egger test are shown in Table 3. There was a publication bias in the meta-analysis of the ARDS group ( $P = .008$ ).

## 4 | DISCUSSION

2019-nCoV is one type of coronaviruses that belong to the  $\beta$ -coronavirus cluster. It causes the third kind of zoonotic coronavirus disease after SARS and Middle East respiratory syndrome (MERS). Wei Ji et al<sup>15</sup> found that 2019-nCoV seemed to be a recombinant virus between bat coronavirus and coronavirus of another unknown

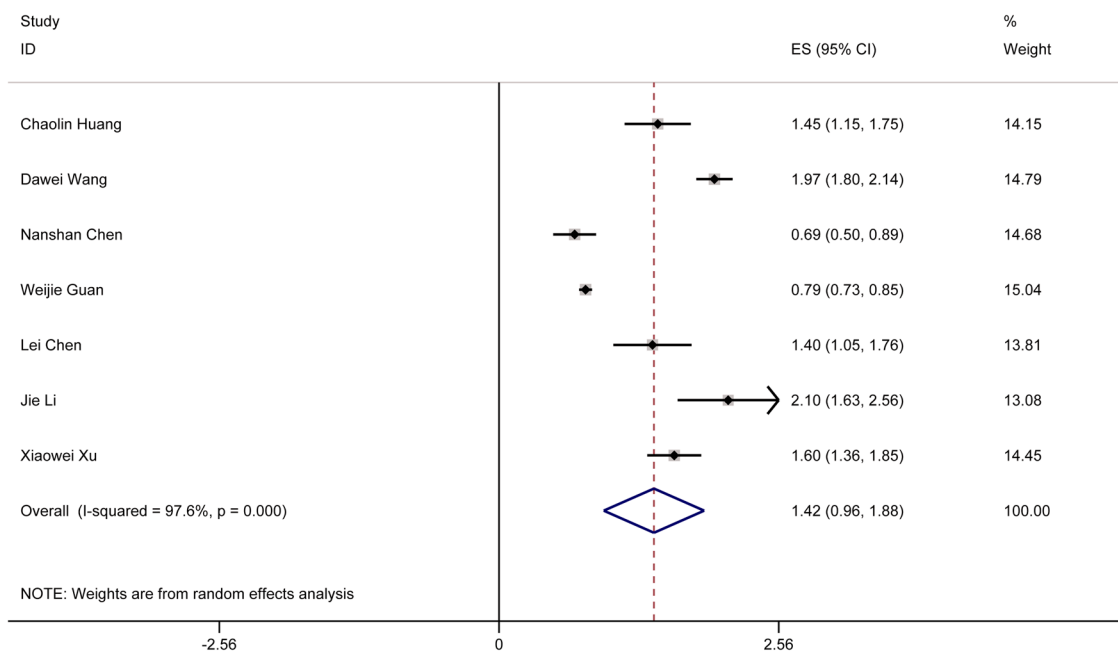


**FIGURE 3** Meta-analysis of the incidence of cough

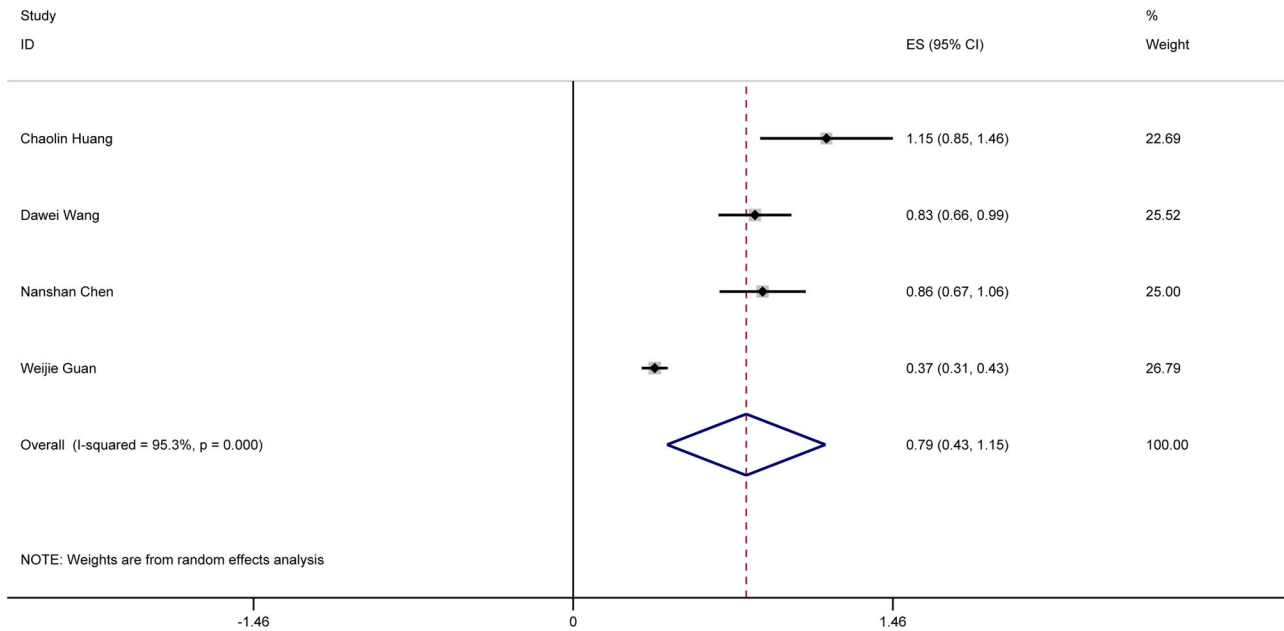
origin. Xiao et al<sup>16</sup> confirmed that Malayan pangolins were the most likely intermediate host of 2019-nCoV. The research of Benvenuto et al<sup>17</sup> found that 2019-nCoV was only closely related to the coronavirus isolated from the chrysanthemum-headed bat in 2015. Their research suggests that bats may be the reservoir host for 2019-nCoV. According to the research by Zhou et al<sup>18</sup> and Wu et al,<sup>19</sup> it was found that the sequence homology between 2019-nCoV and

SARS-CoV was 79.5%. The homology between 2019-nCoV and bat coronavirus at the genetic level was 96%. Therefore, very likely it can be confirmed that 2019-nCoV comes from bats.

According to the results of meta-analysis, we find that the most common symptoms of patients with 2019-nCoV infection were fever and cough, and most of the patients had abnormal chest CT. Some patients with 2019-nCoV infection had muscle soreness or fatigue,



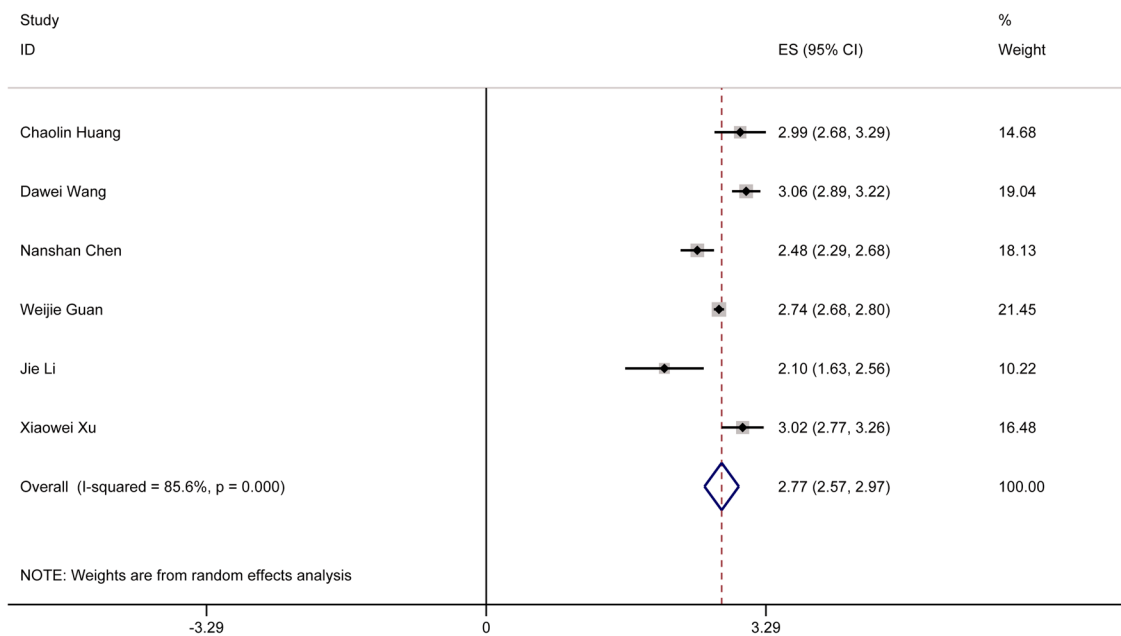
**FIGURE 4** Meta-analysis of the incidence of muscle soreness or fatigue



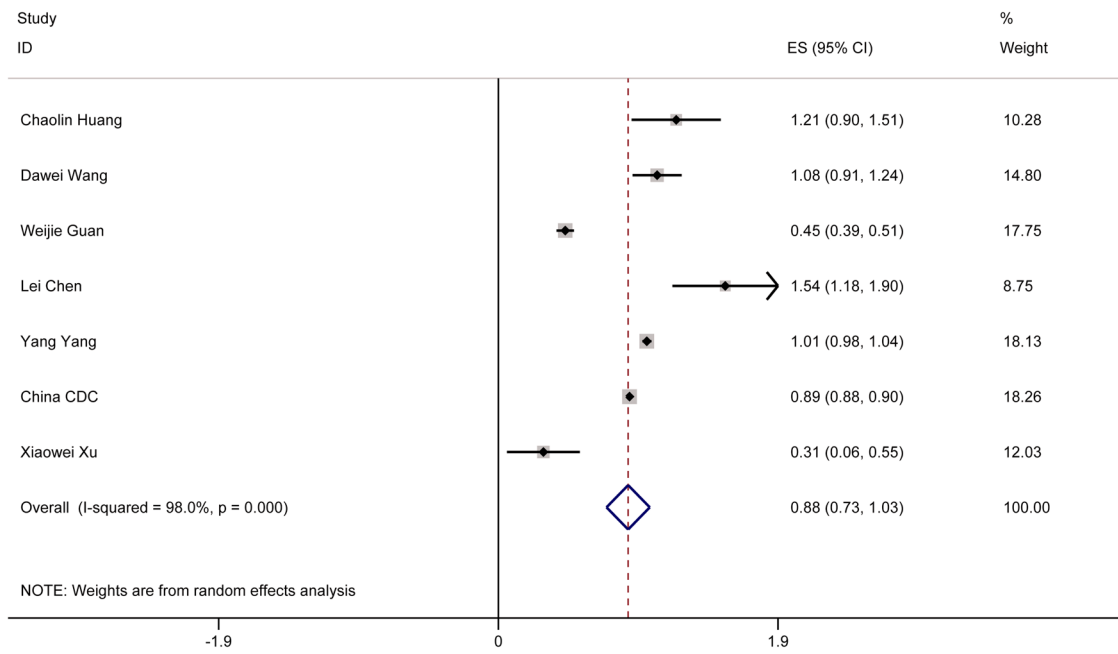
**FIGURE 5** Meta-analysis of the incidence of ARDS. ARDS, acute respiratory distress syndrome

ARDS, and other symptoms. Diarrhea, hemoptysis, headache, sore throat, shock, and other symptoms only occurred among a small number of patients. The proportion of severe cases in all infected cases was 18.1%, and the case fatality rate was 4.3%. At present, the Chinese government has announced that the case fatality rate of patients with 2019-nCoV infection in China is 3.36%, which is lower than that of another two widely contagious zoonotic coronavirus diseases, SARS and MERS. Infection mortality rates of SARS and MERS are 9.6%<sup>20</sup> and 35%,<sup>21</sup> respectively.

Sheng et al<sup>22</sup> have shown that viral infection could increase the risk of pulmonary fibrosis. Xie et al<sup>23</sup> found that 45% of patients showed signs of pulmonary fibrosis within 1 month after being infected with SARS-CoV. Hui<sup>24</sup> revealed that 36% and 30% of patients infected with SARS-CoV developed pulmonary fibrosis 3 and 6 months after infection. These studies consistently suggest that pulmonary fibrosis will become one of the serious complications in patients with 2019-nCoV infection. How to prevent and reduce the occurrence of pulmonary fibrosis in patients with 2019-nCoV



**FIGURE 6** Meta-analysis of the incidence of abnormal chest CT. CT, computer tomography

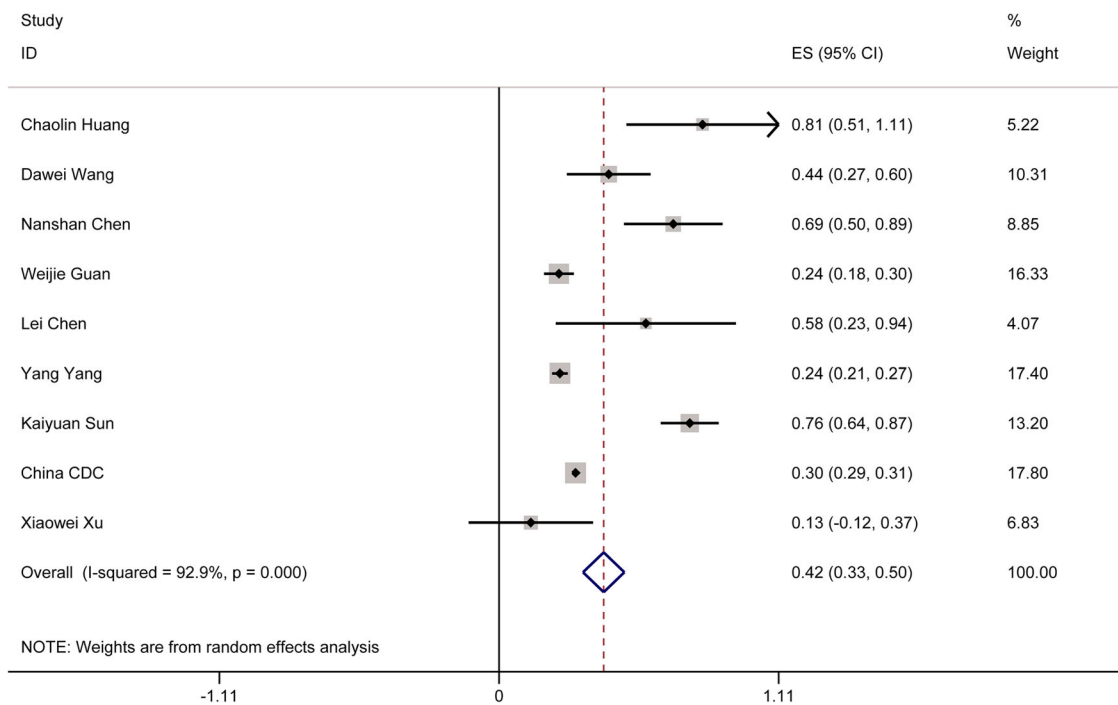


**FIGURE 7** Meta-analysis of the proportion of severe cases in all infected cases

infection are urgent problems for medical workers in the treatment of 2019-nCoV.

Zhao et al<sup>25</sup> found that angiotensin-converting enzyme 2 (ACE2) was the receptor of 2019-nCoV. In normal lung tissue, ACE2 is mainly expressed by type I and type II alveolar epithelial cells. It was reported that 83% of II type alveolar cells expressed ACE2. Therefore, 2019-nCoV infection causes damages to most II

type alveolar cells. In addition, the use of mechanical ventilation in the treatment of patients with 2019-nCoV infection can also aggravate the injury of alveolar cells. After alveolar cell injury, transforming growth factor- $\beta$  (TGF- $\beta$ ) released in tissue promotes lung repair. Virus infection often leads to excessive activation of TGF- $\beta$  pathway, which leads to the occurrence of pulmonary fibrosis.



**FIGURE 8** Meta-analysis of the mortality rate of patients with 2019-nCoV infection. nCoV, novel coronavirus

Clinical symptom	Results of meta-analysis	Adjusted results
Fever	2.47(2.26,2.67)	0.891(0.818,0.945)
Cough	2.03(1.89,2.17)	0.722(0.657,0.782)
Muscle soreness or fatigue	1.42(0.96,1.88)	0.425(0.213,0.652)
ARDS	0.79(0.43,1.15)	0.148(0.046,0.296)
Abnormal chest CT	2.77(2.57,2.97)	0.966(0.921,0.993)
Patient in critical condition	0.88(0.73,1.03)	0.181(0.127,0.243)
Death of patient	0.42(0.33,0.50)	0.043(0.027,0.061)

Abbreviations: ARDS, acute respiratory distress syndrome; CT, computer tomography.

**TABLE 2** Meta-analysis results

**TABLE 3** Results of Egger test

Groups	Fever group	Cough group	Muscle soreness or fatigue group	ARDS group	Abnormal chest CT group	Severe cases group	Death cases group
P	.866	.278	.090	.008	.908	.826	.258

Abbreviations: ARDS, acute respiratory distress syndrome; CT, computer tomography.

Based on the mechanism of pulmonary fibrosis after virus infection, we can formulate corresponding prevention and treatment proposals. The first is the treatment of 2019-nCoV infection. At present, a variety of drugs have been found to inhibit 2019-nCoV. Holshue et al<sup>26</sup> achieved remarkable results in the treatment of patients with 2019-nCoV infection with remdesivir. This drug is currently in clinical trials. Second, it is necessary to inhibit the inflammatory reaction and reduce exudation. In this direction, antibiotics not only have traditional antibacterial effects but also have immunomodulatory and anti-inflammatory properties.<sup>27</sup> Macrolide antibiotics such as clarithromycin, azithromycin, and erythromycin prevent the production of proinflammatory cytokines and immune mediators.<sup>28</sup> The third is to inhibit the fibrogenic effect of TGF- $\beta$ . Luo et al<sup>29</sup> found that arsenic trioxide could inhibit the signal transduction of TGF- $\beta$  and play a role in antifibrosis. The fourth is the rational use of ventilator to avoid unnecessary lung damages. In addition, pirfenidone and nintedanib as drugs for the treatment of idiopathic pulmonary fibrosis can also be used to prevent and treat pulmonary fibrosis in patients with 2019-nCoV infection.<sup>30</sup>

In all, a total of 10 articles were covered in the meta-analysis, including 50 466 patients with 2019-nCoV infection. By far, it is the first meta-analysis with the largest sample size. The quality of the literature included in this study is high, the analysis is rigorous, and the conclusions drawn by the study are highly credible. However, this meta-analysis also has some limitations: (a) all studies included in this meta-analysis are retrospective studies with large heterogeneity; (b) most patients in our meta-analysis are Chinese, and we aimed to use the conclusions of this study to predict patients in general, including other countries and races; (c) there was publication bias in the meta-analysis of the ARDS group. The analytical conclusion of the ARDS group may be influenced by publication bias; and (d) the study subjects were inpatients who have been diagnosed with 2019-nCoV infection. So, the incidence of severe pneumonia or the fatality rate in

the total population infected with 2019-nCoV would be much lower than the results shown in this study.

Therefore, based on the above-mentioned limitations, the conclusions of this meta-analysis still need to be verified by more relevant studies with more careful design, more rigorous execution, and larger sample size.

#### ACKNOWLEDGMENT

We would like to acknowledge TopEdit LLC for the linguistic editing and proofreading during the preparation of this manuscript.

#### CONFLICT OF INTERESTS

The authors declare that there are no conflict of interests.

#### AUTHOR CONTRIBUTIONS

JX and PS developed the idea for and designed the study and had full access to all data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis. PS and SQ contributed to the writing of the report. JX contributed to the critical revision of the report. SQ, ZL, JR, and KL contributed to the statistical analysis. All authors contributed to data acquisition, data analysis, or data interpretation, and reviewed and approved the final version.

#### ORCID

Jianing Xi  <http://orcid.org/0000-0002-8405-2355>

#### REFERENCES

1. Lancet T. Emerging understandings of 2019-nCoV. *Lancet*. 2020; 395(10221):311.
2. Lu H, Stratton CW, Tang YW. Outbreak of pneumonia of unknown etiology in Wuhan China: the mystery and the miracle. *J Med Virol*. 2020;92:401-402. <https://doi.org/10.1002/jmv.25678>



3. Moher D, Liberati A, Tetzlaff J, Altman DG. Preferred reporting items for systematic reviews and meta-analyses: the PRISMA statement. *J Clin Epidemiol*. 2009;62(10):1006-1012.
4. Stang A. Critical evaluation of the Newcastle-Ottawa scale for the assessment of the quality of nonrandomized studies in meta-analyses. *Eur J Epidemiol*. 2010;25(9):603-605.
5. Huang C, Wang Y, Li X. Clinical features of patients infected with 2019 novel coronavirus in Wuhan, China. *Lancet*. 2020;395:497-506.
6. Wang D, Hu B, Hu C, et al. Clinical characteristics of 138 hospitalized patients with 2019 novel coronavirus-Infected Pneumonia in Wuhan, China. *JAMA*. 2020. <https://doi.org/10.1001/jama.2020.1585>
7. Chen N, Zhou M, Dong X, et al. Epidemiological and clinical characteristics of 99 cases of 2019 novel coronavirus pneumonia in Wuhan, China: a descriptive study. *Lancet*. 2020;395:507-513.
8. Guan W-j, Ni Z-y, Hu Y, et al. Clinical characteristics of 2019 novel coronavirus infection in China. *medRxiv*. 2020. <https://doi.org/10.1101/2020.02.06.20020974>
9. Chen J, Liu HG, Liu W. Analysis of clinical features of 29 patients with 2019 novel coronavirus pneumonia. *Zhonghua Jie He He Hu Xi Za Zhi*. 2020;43:E005.
10. Sun K, Chen J, Viboud C. Early epidemiological analysis of the 2019-nCoV outbreak based on a crowdsourced data. *medRxiv*. 2020. <https://doi.org/10.1101/2020.01.31.20019935>
11. Yang Yang, Lu Qingbin, Liu Mingjin, et al. Epidemiological and clinical features of the 2019 novel coronavirus outbreak in China. *medRxiv*. 2020. <https://doi.org/10.1101/2020.02.10.20021675>
12. Li Jie, Li Shilin, Cai Yurui, et al. Epidemiological and clinical characteristics of 17 hospitalized patients with 2019 novel coronavirus infections outside Wuhan, China. *medRxiv*. 2020. <https://doi.org/10.1101/2020.02.11.20022053>
13. The Noval Coronavirus Pneumonia Emergency Response Epidemiology Team, Chinese Center for Disease Control and Prevention. The epidemiological characteristics of an outbreak of 2019 novel coronavirus diseases (COVID-19) in China. *Zhonghua Liu Xing Bing Xue Za Zhi*. 2020;41:145-151.
14. Xu XW, Wu XX, Jiang XG, et al. Clinical findings in a group of patients infected with the 2019 novel coronavirus (SARS-Cov-2) outside of Wuhan, China: retrospective case series. *BMJ*. 2020;368:m606.
15. Ji W, Wang W, Zhao X, Zai J, Li X. Homologous recombination within the spike glycoprotein of the newly identified coronavirus may boost cross-species transmission from snake to human. *J Med Virol*. 2020;92:433-440. <https://doi.org/10.1002/jmv.25682>
16. Xiao Kangpeng, Zhai Junqiong, Feng Yaoyu, et al. Isolation and characterization of 2019-nCoV-like coronavirus from Malayan Pangolins. *medRxiv*. 2020. <https://doi.org/10.1101/2020.02.17.951335>
17. Benvenuto D, Giovanetti M, Ciccozzi A, Spoto S, Angeletti S, Ciccozzi M. The 2019-new coronavirus epidemic: evidence for virus evolution. *J Med Virol*. 2020;92:455-459. <https://doi.org/10.1002/jmv.25688>
18. Zhou P, Yang XL, Wang XG, et al. A pneumonia outbreak associated with a new coronavirus of probable bat origin. *Nature*. 2020. <https://doi.org/10.1038/s41586-020-2012-7>
19. Wu F, Zhao S, Yu B, et al. A new coronavirus associated with human respiratory disease in China. *Nature*. 2020. <https://doi.org/10.1038/s41586-020-2008-3>
20. Hui DSC, Zumla A. Severe acute respiratory syndrome: historical, epidemiologic, and clinical features. *Infect Dis Clin North Am*. 2019;33(4):869-889.
21. Azhar EI, Hui DSC, Memish ZA, Drosten C, Zumla A. The Middle East respiratory syndrome (MERS). *Infect Dis Clin North Am*. 2019;33(4):891-905.
22. Sheng G, Chen P, Wei Y, et al. Viral infection increases the risk of idiopathic pulmonary fibrosis: a meta-analysis. *Chest*. 2019. <https://doi.org/10.1016/j.chest.2019.10.032>
23. Xie L, Liu Y, Xiao Y, et al. Follow-up study on pulmonary function and lung radiographic changes in rehabilitating severe acute respiratory syndrome patients after discharge. *Chest*. 2005;127(6):2119-2124.
24. Hui DS. Impact of severe acute respiratory syndrome (SARS) on pulmonary function, functional capacity and quality of life in a cohort of survivors. *Thorax*. 2005;60(5):401-409.
25. Zhao Y, Zhao Z, Wang Y, et al. Single-cell RNA expression profiling of ACE2, the putative receptor of Wuhan 2019-nCoV. *bioRxiv*. 2020. <https://doi.org/10.1101/2020.01.26.919985>
26. Holshue ML, DeBolt C, Lindquist S, et al. First case of 2019 novel coronavirus in the United States. *N Engl J Med*. 2020. <https://doi.org/10.1056/NEJMoa2001191>
27. Buret AG. Immuno-modulation and anti-inflammatory benefits of antibiotics: the example of tilmicosin. *Can J Vet Res*. 2010;74(1):1-10.
28. Wuyts WA, Willems S, Vos R, et al. Azithromycin reduces pulmonary fibrosis in a bleomycin mouse model. *Exp Lung Res*. 2010;36(10):602-614.
29. Luo F, Zhuang Y, Sides MD, et al. Arsenic trioxide inhibits transforming growth factor- $\beta$ 1-induced fibroblast to myofibroblast differentiation in vitro and bleomycin induced lung fibrosis in vivo. *Respir Res*. 2014;15:51.
30. Chunbin Wang, Shuang Zhang, Qian Zhang, et al. Research progress of two new antifibrotic drugs for idiopathic pulmonary fibrosis. *Chin Pharm J*. 2019;09:682-686.

**How to cite this article:** Sun P, Qie S, Liu Z, Ren J, Li K, Xi J. Clinical characteristics of 50 466 hospitalized patients with 2019-nCoV infection. *J Med Virol*. 2020;1-9. <https://doi.org/10.1002/jmv.25735>