Sex differences in clinical characteristics and risk factors for disease severity of hospitalized patients with COVID-19

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ORIGINAL ARTICLE

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Abstract
Recent studies reported sex differences in patients with coronavirus disease-2019 (COVID-19). We aim to analyze sex differences in clinical characteristics and risk factors for disease severity of hospitalized patients with COVID-19 in Beijing. All adults (185 cases) diagnosed with COVID-19 and admitted to Beijing Ditan Hospital, Capital Medical University were included in samples. The median age of all patients was 41 years. The mean body mass index (BMI) of males was relatively higher compared to females (p < 0.001). The proportion of male patients with coronary heart disease (CHD), nonalcoholic fatty liver disease (NAFLD), history of smoking and drinking was higher than females. Male patients developed more clinical symptoms, obtained more abnormal laboratory test results, while they were less aware of care-seeking than female patients. There were no significant differences in clinical complications and outcomes between two groups. Age (odds ratio [OR]: 1.082; 95% confidence interval [CI]: 1.034–1.132; p = 0.001) and BMI (OR: 1.237; 95% CI: 1.041–1.47; p = 0.016) were considered risk factors for refractory pneumonia in multivariate regression analysis. The findings of the current study showed that SARS-CoV-2 was more likely to affect older males with comorbidities. Further researches into factors underlying obesity and disease severity may provide mechanistic insight into COVID-19 development.

KEYWORDS
COVID-19, disease severity, risk factors, SARS-CoV-2, sex differences

Abbreviations: AKI, acute kidney injury; ARDS, acute respiratory distress syndrome; AST, aspartate aminotransferase; BMI, body mass index; CHD, coronary heart disease; CI, confidence interval; CK, creatine kinase; COVID-19, coronavirus disease-2019; CRP, c-reactive protein; CT, computerized tomography; DBP, diastolic blood pressure; DIC, diffuse intravascular coagulation; EMR, electronic medical record; FIB, fibrinogen; HT, hypertension; MERS, Middle East respiratory syndrome; NAFLD, nonalcoholic fatty liver disease; OR, odds ratio; SARS, severe acute respiratory syndrome; SARS-CoV-2, severe acute respiratory syndrome coronavirus 2; SBP, systolic blood pressure; T2DM, type 2 diabetes mellitus

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INTRODUCTION

Coronavirus disease-2019 (COVID-19) is an emerging respiratory infectious disease caused by the severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) and triggered a global pandemic which has not been under control yet. Even the situation has been curbed in some countries as of July 11, there are still 12,322,395 cases diagnosed and 556,335 deaths confirmed around the world. Previous reports have reported that there existed a sex imbalance based on confirmed cases and case fatality rate of COVID-19. Notably, the roles and significance of sex are often neglected in studies of infectious disease. As studied, hormone difference alters the host response to infection disease, and sex-biasing differences influence host pathogen interactions. Previous studies have reported that European and American had higher percentage of male patients with COVID-19 were far worse than females. More than that, male sex, older age, underlying disease such as type 2 diabetes mellitus (T2DM), hypertension (HT), obesity, coronary heart disease (CHD) are risk factors associated with worse outcomes. The health status of Beijing residents has reached the level of high-income countries and regions, and the health status of women is better than that of men. Based on recent reports, the prevalence of overweight was the highest in Beijing among Chinese adults published in 2019, the prevalence of overweight was the highest in Beijing. The differences in the health status of men and women might contribute to the sex differences in COVID-19, and the characteristics of COVID-19 may be different from those of other cities as well. However, sex differences associated with clinical characteristics, severity and mortality of COVID-19 in Beijing have not been well described yet. We summarized the clinical characteristics, comorbid conditions, severity and outcomes of 185 hospitalized patients in order to investigate whether sex bias is associated with the clinical characteristics and early outcomes of the patients who were diagnosed with COVID-19 in Beijing and to assess potential risk factors on patients with severe COVID-19 at admission.

METHODS

2.1 Study design and participants

From January 13, 2019 to March 19, 2020, a total of 185 adults with COVID-19 were diagnosed and admitted if tested positive for respiratory symptoms, showed typical chest imaging findings, and tested positive SARS-CoV-2 test results of pharyngeal swab specimens in Beijing Ditan Hospital. Data collected through the electronic medical record system included baseline demographic information (age, sex, body mass index [BMI], smoking history, and alcohol consumption), underlying disease (CHD, HT, T2DM, respiratory disease, and nonalcoholic fatty liver disease [NAFLD]), symptoms (fever, headache, fatigue, cough, sore throat, cough, expectoration, shortness of breath, nausea/vomiting, myalgia/arthritis, diarrhea), vital signs (indoor air oxygen saturation, the highest temperature, heart rate [HR], systolic blood pressure [SBP], diastolic blood pressure [DBP]), laboratory data of the first 24-h hospital stay (leukocyte count, neutrophil count, lymphocyte count, neutrophil/lymphocyte ratio (NLR), hemoglobin, platelet count, creatinine, AST, alanine aminotransferase (ALT), troponin T, bilirubin, albumin, creatine kinase (CK), lactate dehydrogenase, d-dimer, fibrinogen (FIB), CRP, procalcitonin, serum amyloid A (SAA)), computerized tomography (CT), clinical complications including acute respiratory distress syndrome (ARDS), acute kidney injury (AKI) or diffuse intravascular coagulation (DIC), septic shock, co-infection, myocardial damage and final diagnosis including remained in hospital, discharge, death, and readmission. All data were analyzed independently by three researchers and obtained in accordance with standard biosafety and institutional safety procedures during the observation period of this study. The severity of the patient with COVID-19 is determined based on the diagnostic and treatment guideline for SARS-CoV-2 issued by Chinese National Health Committee (version 7).

2.2 Statistical analysis

The normality of continuous variables’ distribution was tested by one-sample Kolmogorov-Smirnov test. Continuous variables with normal distribution were presented as mean ± standard deviation (SD). Mean of two continuous normally distributed variables were compared using independent samples Student’s test; non-normal variables were reported as median (interquartile range [IQR]). Mann-Whitney U test and Kruskal-Wallis test were used, respectively, to compare means of non-normal distributions. Categorical variables were summarized as numbers and percentages. We made comparisons via analysis of chi-square for categorical variables between males and females. Correlations between severity of COVID-19 and underlying disease (CHD, HT, T2DM, respiratory disease, NAFLD) were analyzed in terms of the Spearman Pearson correlation coefficient. Multivariable binary logistic regression analyses were used to assess the association among age, sex, BMI, underlying comorbidity (CHD, HT, T2DM, respiratory disease, NAFLD), history of smoking, alcohol consumption and the dependent variable of
severity of disease (general group and refractory group). Results of logistic regression are given as the odds ratio (OR) with the 95% confidence interval (CI). A p value < 0.05 was considered statistically significant using two-sided tests. The data were analyzed by SPSS version 16.0 and Graph Pad Prism 8.0.

3 | RESULTS

3.1 Baseline characteristics for different sex groups

A total of 185 patients were admitted and initially diagnosed with COVID-19, among whom 95 (51.4%) were men. The age of all patients ranged from 18 to 92 years with a median of 41 years, furthermore, there is no differences in age between males and females (p = 0.914). Based on data collected, the mean BMI was higher in males (25.45 vs. 22.29, p < 0.001), and the mean duration from symptoms initiation to hospital admission was longer for men than that for women (6 vs. 4, p = 0.042). Considering previous living habits of patients, the number of patients with smoking and drinking history was significantly higher in male patients (specifically smoking: 18.9% vs. 2.2% and drinking: 28.4% vs. 6.7%, p < 0.001). Among all patients, 77 cases (41.6%) had one or more following comorbidities including but not limited to HT (22.7%), NAFLD (14.6%), respiratory diseases (9.2%), T2DM (7.6%), CHD (3.2%). More specifically, the percentages of males with CHD (6.3% vs. 0.0%) and NAFLD (20.0% vs. 8.9%) were significantly higher than females (p < 0.05) (Table 1).

3.2 Vital signs and laboratory parameters

All vital signs and laboratory parameters were collected on the day of hospital admission for all patients. The peak temperature measured prior to hospital admission was higher in male patients than females (38.20°C vs. 37.65°C, p = 0.008). Moreover, the medians of SBP, DBP in males were higher than in females (p < 0.05). There was no significant difference in HR and respiratory rate between male and female patients. The most common symptoms observed onset were fever (74.6%), cough (56.2%), fatigue (35.7%), expectoration (29.7%), myalgia/arthritis (28.1%), sore throat (20.0%), and chills (19.5%). Furthermore, several less common symptoms such as headache (18.4%), diarrhea (9.2%), nausea, and vomiting (5.4%) were also observed. Of the 185 patients, 138 patients (74.6%) had two or more symptoms concurrently. Among all the listed symptoms, the percentages of patients with fever (T ≥ 38°C, 60.0% vs. 44.4%), cough (65.3% vs. 46.7%) and chills (25.3% vs. 13.3%) were significantly higher in male group than females (p < 0.05) (Table 1). Although leukocyte count, neutrophil count, lymphocyte count did not differ between two groups (p > 0.05), the platelet count was significantly lower in male group than that in female group (p < 0.001). However, no sex differences were observed in lymphopenia and thrombocytopenia (p > 0.05). In addition, there were numerous factors showed significant higher proportion in males than females including ALT, CK, CRP (p < 0.001, p = 0.006, p = 0.008, respectively). Of note, FIB, another acute response protein, was also higher in male patients than that in female patients (p = 0.033) (Table 2).

3.3 Chest computed tomographic images

We analyzed the chest CT imaging of all 185 patients at the first examination after admission. All data analyzed from CT were divided into two categories: normal and abnormal. We performed a R × C chi-square test to compare chest CT imaging between two groups. Higher percentages of CT abnormalities were discovered in male patients (89.47% vs. 77.78%, p = 0.031) (Figure 1).

3.4 Severity assessment and clinical complications on admission

As described in the diagnostic and treatment guideline for SARS-CoV-2 issued by the Chinese National Health Committee, all patients were divided into general and refractory groups according to the clinical efficacy after hospitalization. By comparing the severity of disease in two groups, the rate of patients diagnosed as general and refractory illness and refractory patients in the male group were more than that in female group (p = 0.014) (Figure 2). There is no significant difference between males and females in complications, such as ARDS, acute kidney insufficiency (AKI) or acute exacerbation of chronic kidney insufficiency, diffuse DIC, acute myocardial injury, co-infection and septic shock (p > 0.05) (Table 3).

3.5 Outcomes of patients with COVID-19

As of March 23, 2020, a total of two patients died during this period, both of whom were males. The mortality rate of COVID-19 in this study was 1.1%, which did not differ
### TABLE 1  Demographics and baseline characteristics of patients infected with SARS-CoV-2 according to sex

| Variables                                 | All patients (n = 185) | Men (n = 95) | Women (n = 90) | p values |
|-------------------------------------------|------------------------|--------------|----------------|----------|
| Age                                       | 41 (32.57)             | 42 (32.54)   | 40.5 (29.75, 57.25) | 0.914    |
| BMI                                       | 23.94 ± 4.41 (n = 132) | 25.45 ± 4.40 (n = 69) | 22.29 ± 3.81 (n = 63) | <0.001   |
| From symptom to diagnosis"                | 5 (3, 8)               | 6 (3, 8)     | 4 (3, 8)       | 0.042    |
| History of smoking                        | 20 (10.80%)            | 18 (18.90%)  | 2 (2.20%)      | <0.001   |
| Alcohol consumption                       | 33 (17.80%)            | 27 (28.40%)  | 6 (6.70%)      | <0.001   |
| Coronary heart disease                    | 6 (3.20%)              | 6 (6.30%)    | 0 (0.00%)      | 0.029    |
| Hypertension                              | 42 (22.70%)            | 24 (25.30%)  | 18 (20.00%)    | 0.393    |
| Non-alcoholic fatty liver disease         | 27 (14.60%)            | 19 (20.00%)  | 8 (8.90%)      | 0.032    |
| Respiratory disease                       | 17 (9.20%)             | 7 (7.40%)    | 10 (11.10%)    | 0.378    |
| Diabetes                                  | 14 (7.60%)             | 9 (9.50%)    | 5 (5.60%)      | 0.314    |
| Peak temperature,"°C"                     | 38.0 (36.50, 38.50)    | 38.20 (37.50, 38.60) | 37.65 (36.50, 38.30) | 0.008    |
| Fever ≥ 38°C                               | 97 (52.40%)            | 57 (60.00%)  | 40 (44.40%)    | 0.034    |
| HR                                        | 88 (80.99)             | 89 (81.98)   | 86 (78.99)     | 0.620    |
| RR                                        | 20 (17.20)             | 20 (17.21)   | 19 (17.20)     | 0.559    |
| SBP                                       | 128 (117, 140)         | 130 (121, 143) | 126 (112, 137.25) | 0.004    |
| DBP                                       | 83 (75, 90)            | 85 (76, 91)  | 80 (74, 89)    | 0.030    |
| O2 Saturation < 93%                       | 19 (10.30%)            | 10 (10.50%)  | 9 (10.00%)     | 0.906    |
| Headache                                  | 34 (18.40%)            | 16 (16.80%)  | 18 (20.00%)    | 0.579    |
| Fever                                     | 138 (74.60%)           | 74 (77.90%)  | 64 (71.10%)    | 0.289    |
| Fatigue                                   | 66 (35.70%)            | 36 (37.90%)  | 30 (33.30%)    | 0.517    |
| Cough                                     | 104 (56.20%)           | 62 (65.30%)  | 42 (46.70%)    | 0.011    |
| Sore throat                               | 37 (20.00%)            | 21 (22.10%)  | 16 (17.80%)    | 0.462    |
| Expectoration                             | 55 (29.70%)            | 29 (30.50%)  | 26 (28.90%)    | 0.808    |
| Shortness of breath                       | 30 (16.20%)            | 13 (13.70%)  | 17 (18.90%)    | 0.337    |
| Nausea/vomiting                           | 10 (5.40%)             | 5 (3.30%)    | 5 (3.30%)      | 0.930    |
| Diarrhea                                  | 17 (9.20%)             | 11 (11.60%)  | 6 (6.70%)      | 0.248    |
| Myalgia/arthritis                         | 52 (28.10%)            | 29 (30.50%)  | 23 (25.60%)    | 0.452    |
| Chills                                    | 36 (19.50%)            | 24 (25.30%)  | 12 (13.30%)    | 0.041    |
| Signs or symptoms ≥ two                   | 138 (74.60%)           | 75 (78.90%)  | 63 (70.00%)    | 0.162    |

Note: The normality of continuous variables’ distribution was tested by one-sample Kolmogorov-Smirnov test. Continuous variables with normal distribution were presented as mean ± standard deviation (SD); Mean of two continuous normally distributed variables was compared using independent samples Student’s test; non-normal variables were reported as median (interquartile range [IQR]). Mann-Whitney U test and Kruskal-Wallis test were used, respectively, to compare means of non-normal distributions. Categorical variables were summarized as numbers and percentages. “From symptom to diagnosis” indicates the days. * The peak temperature was measured prior to hospital admission.

Between two groups (p = 0.501) (Table 3). Furthermore, there was no statistical difference in the number of patients remained in hospital, discharged, re-admitted caused by reactivation of SARS-CoV-2 after discharge (p > 0.05) (Table 3).

### 3.6 Risk factors analysis of COVID-19 severity

Patients were classified in two groups based on severity of COVID-19 as described previously. Correlations between severity of COVID-19 and underlying disease (CHD, HT, T2DM, respiratory disease, and NAFLD) were analyzed in terms of the Spearman correlation coefficient, but weak correlations or no correlation were found with CHD (p = 0.007, r = 0.196), HT (p < 0.001, r = 0.270), T2DM (p < 0.001, r = 0.290), respiratory disease (p < 0.001, r = 0.281), NAFLD (p = 0.132, r = 0.111) and disease severity, respectively. Due to the limited sample size of CHD, the data could not be counted accurately and effectively in multivariate regression, only age (analyzed in quartiles, OR: 1.082; 95% CI: 1.034–1.132; p = 0.001) and BMI (OR: 1.237; 95% CI: 1.041–1.47; p = 0.016) could be considered risk factors for refractory pneumonia in multivariate regression analysis (Figure 3). We added Spearman correlation analysis between male sex and BMI and found it to be significant (p < 0.001, r = 0.486).
TABLE 2  Laboratory findings on admission to hospital of patients infected with SARS-CoV-2

| Variables                          | All patients (n = 185) | Men (n = 95) | Women (n = 90) | p values |
|------------------------------------|------------------------|--------------|----------------|----------|
| Leukocyte, ×10^9/L                 | 5.36 ± 2.36            | 5.40 ± 2.10  | 5.32 ± 2.60    | 0.805    |
| Lymphocyte count, ×10^9/L          | 1.37 ± 0.61            | 1.31 ± 0.57  | 1.44 ± 0.65    | 0.169    |
| NLR                               | 2.50 (1.63, 3.58)      | 2.62 (1.78, 4.07) | 2.41 (1.37, 3.36) | 0.110 |
| Haemoglobin, g/L                   | 140.61 ± 17.22         | 149.85 ± 15.24 | 130.86 ± 13.41 | <0.001  |
| Lymphopenia*                       | 123 (66.50%)           | 67 (70.50%)  | 56 (62.20%)    | 0.232    |
| Thrombocytopenia*                  | 39 (21.10%)            | 25 (26.30%)  | 14 (15.60%)    | 0.073    |
| blood platelet count, ×10^9/L      | 197 (154, 242)         | 176 (147, 225) | 223 (172, 263.25) | <0.001  |
| Sodium, mmol/L                     | 139.1 (137.0, 140.6)   | 138.1 (136.1, 140.5) | 139.9 (138.65, 141.15) | 0.001 |
| Potassium, mmol/L                  | 3.71 (3.52, 4.00)      | 3.8 (3.58, 4.00) | 3.7 (3.45, 3.95) | 0.027    |
| Creatinine ≥ 133, µmol/L           | 3 (1.60%)              | 1 (1.10%)    | 2 (2.20%)      | 0.613    |
| Aspartate aminotransferase > 40 U/L (n = 176) | 28 (15.90%) | 19 (20.40%) | 9 (10.80%) | 0.083 |
| Alanine aminotransferase > 40 U/L  | 36 (19.50%)            | 29 (30.50%)  | 7 (7.80%)      | <0.001   |
| Total bilirubin ≥ 17.1, µmol/L (n = 175) | 22 (12.50%) | 13 (14.00%) | 9 (10.80%) | 0.530   |
| Albumin, g/L                       | 42.4 (38.75, 45.45)    | 42.4 (38, 46.30) | 42.2 (39.5, 45.2) | 0.661 |
| Creatinine kinase ≥ 200 U/L        | 23 (12.40%)            | 18 (18.90%)  | 5 (5.60%)      | 0.006    |
| C-reactive protein level ≥ 10 mg/L (n = 182) | 74 (40.70%) | 47 (50.00%) | 27 (30.70%) | 0.008 |
| Serum amyloid A ≥ 10 mg/L (n = 134) | 82 (61.20%) | 48 (61.60%) | 34 (53.10%) | 0.067 |
| Lactose dehydrogenase ≥ 250 U/L (n = 173) | 55 (31.80%) | 34 (37.40%) | 21 (25.60%) | 0.097 |
| D-dimer ≥ 0.5 (n = 171)            | 61 (35.70%)            | 33 (36.30%)  | 28 (35.00%)    | 0.863    |
| Fibrinogen, µg/mL (n = 184)        | 274 (216, 365)         | 306 (214, 386) | 247 (219, 249) | 0.033    |

Note: Continuous variables with normal distribution were present as mean ± standard deviation (SD). Mean of two continuous normally distributed variables was compared using independent samples Student's test; non-normal variables were reported as median (interquartile range [IQR]). Mann-Whitney U test and Kruskal-Wallis test were used, respectively, to compare means of non-normal distributions. Categorical variables were summarized as numbers and percentages.

*A lymphopenia denoted the lymphocyte count of less than 1500 per cubic millimeter. #Thrombocytopenia denoted the platelet count of less than 150,000 per cubic millimeter; $NLR$, neutrophil lymphocyte ratio.

FIGURE 1  Chest computed tomographic (CT) images. (A) Chest CT images of a male patient infected with SARS-CoV-2 were partially absorbed after nasal high-flow oxygen inhalation treatment. (B) Chest CT images of a female patient infected with SARS-CoV-2 showed patchy ground glass shadows and diffuse pulmonary fibrosis. (C) Higher percentages of CT abnormalities were discovered in male patients at the first examination after admission.
TABLE 3 Clinical complications and outcome of patients infected on admission

| Variables          | All patients (n = 185) | Men (n = 95)    | Women (n = 90) | p values |
|--------------------|------------------------|----------------|----------------|----------|
| ARDS               | 18 (9.70%)             | 10 (10.50%)    | 8 (8.90%)      | 0.707    |
| Acute renal injury | 1 (0.50%)              | 1 (1.10%)      | 0 (0.00%)      | >0.99    |
| Septic shock       | 4 (2.20%)              | 3 (3.20%)      | 1 (1.10%)      | 0.339    |
| Co-infections      | 57 (30.80%)            | 32 (33.70%)    | 25 (27.80%)    | 0.384    |
| Myocardial damage  | 6 (3.20%)              | 4 (4.40%)      | 2 (2.10%)      | 0.629    |
| DIC                | 0                      | 0              | 0              | 1        |
| ≥ two complications | 61 (33.00%)            | 34 (35.80%)    | 27 (30.00%)    | 0.402    |
| Remained in hospital | 72 (38.90%)         | 35 (36.80%)    | 37 (41.10%)    | 0.552    |
| Recovery           | 96 (51.90%)            | 48 (50.50%)    | 48 (53.30%)    | 0.703    |
| Mortality          | 2 (1.10%)              | 2 (2.10%)      | 0 (0.00%)      | 0.501    |
| Readmission        | 15 (8.10%)             | 10 (10.50%)    | 5 (5.60%)      | 0.216    |

Note: The observation period of this study was from January 13 to March 19, 2020. Discharge criteria: (1) Body temperature returns to normal for more than 3 days; (2) Respiratory symptoms improved significantly; (3) Pulmonary imaging showed significant improvement of acute exudative lesions; (4) Two consecutive negative nucleic acid tests of respiratory tract specimens such as sputum, nose and throat swabs (sampling time interval of at least 24 h). Reason for readmission: reactivation of SARS-CoV-2 after discharge.

FIGURE 2 Severity of patients infected in male and female groups on admission. (A) Sankey diagram for general and refractory illness according to the clinical efficacy after hospitalization between male and female groups. (B) Refractory patients in the male group were more than that in female group

4 | DISCUSSION

As reported previously, male and female may differ in susceptibility and response to certain infectious diseases. For example, men are more susceptible to certain infectious diseases, such as malaria, SARS, Middle East respiratory syndrome (MERS) and influenza than females. As known, infectious diseases may cause inflammatory-mediated immune response, and the differences in immune responses between women and men may be related to sex hormones and specific X-chromosome-encoded genes. Many previous reports and studies have demonstrated the importance of taken sex difference into consideration during disease research. Taken all the facts into consideration, sex might play a critical role in COVID-19 studies as well. In support of this hypothesis, many existing researches focusing on COVID-19 have reported that there were more male patients admitted than females in the same area, other than that, the mortality and severity rates were also higher in male patients. All previous reports suggested possible sex effects during the progression of COVID-19. In this study, by comparing the baseline data of male and female patients, the mean age and sex ratios of the two groups of patients were similar, which might be limited by geographical factors. However, in male patients, the proportions of smokers (18.9% vs. 2.2%), patients with CHD (6.3% vs. 0.0%) and patients with NAFLD (20.00% vs. 8.89%) were higher than those of female patients. Moreover, more male patients developed symptoms such as high fever, cough, and chills. In the related content, the abnormal CT rate in male patients is higher than that in female patient, suggesting the clinical symptoms were more sever and the pulmonary infection progresses faster in male patients. Based upon analyzed data, the refractory illness rate (including critical and severe COVID-19) is higher in the male group than in the female group, which is consistent with the results of recent COVID-19 studies in other regions. Many believes that the sex differences of SARS-CoV-2 is related to the difference in ACE2 content in men and women, furthermore, sex is used as a strong indicator of ACE2 concentration. ACE2 exists not only in lung, but also in other types of tissue such as heart muscle, kidneys, and blood vessel walls and is particularly high in testis.
has been reported that SARS-CoV-2 may infect the male genital urinary system, presumably inhibiting the function of cells and reproduction. In support of this, a retrospective study of serum samples collected from 81 male patients with COVID-19 who were admitted to Leishen Mountain Hospital, Wuhan. They found that luteinizing hormone and prolactin levels were significantly increased, while testosterone and follicle-stimulating hormone ratios were significantly decreased in patients with COVID-19. This might be caused by damage to the interstitial cells. The gene encoding ACE2 is located on the X chromosome. Any X chromosome-related diseases generally showed higher infectious rate in males than in females due to the genetic chromosome differences. As we discovered similar sex differences patterns in COVID-19 disease research, we propose that ACE2 might be related to the sex differences in COVID-19 between males and females as well. Kimberly E. Stelzig demonstrated that estrogen regulates the expression of SARS-CoV-2 receptor ACE2 in differentiated airway epithelial cells in a single female donor of NHBE cells; however more research is needed to confirm this finding. Other than the genetic contribution, different cultural and behavioral habits might act into the sex difference in COVID-19 as well. In this cohort, the percentage of patients who were smokers was significantly higher in male group than that in female group. Long-term smoking might cause underlying lung diseases, which will affect lung ventilation and exacerbate lung disease progression, which played an important part in COVID-19 progressions. Other than that, during the admission process, we found out that men intended to contradict hospital admission and thus were hospitalized longer after onset symptoms observed. This finding suggested that, compared with men, women were more willing and optimistic on seeking medical help.

Inflammatory storms have been mentioned in many studies, and our study also confirmed the inflammatory response. This study indicated that male patients had higher levels of inflammatory markers (including CRP, SAA, FIB), suggesting that bacterial infection is more common in male patients and might aggravate the disease progression. The elevated AST and CK levels suggested that it may be related to the tissue damage mediated by virus, and the male response is more severe than female. Although the mean platelet counts in males were significantly lower than those in females, the mean platelet counts in both groups were within the normal range, and the thrombocytopenia ratios were parallel between the two groups; therefore, there was no clinical value in the decrease of platelet count between male and female groups. Correlations between severity of COVID-19 and underlying disease (CHD, HT, T2DM, respiratory disease, NAFLD) were analyzed in terms of the Spearman correlation coefficient, but weak correlations or no correlation were found between underlying disease and disease severity respectively. This result is different from previous studies, which may be caused by low proportion of patients with underlying diseases or critically ill patients included in our study. After logistic regression analysis between severe and non-severe cases, we found that age and BMI were independent risk factors, there was a positive correlation between male sex and BMI, which might contribute to the fact that male group had more patients with severe pneumonia and testified the research conclusions of Cai et al and Petrakis et al. Torio Emanuele Bianchi conducted a meta-analysis and found that low testosterone levels were associated with high levels of adipocytokines and inflammatory responses. Adipose tissue is a source of many inflammatory factors and

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**FIGURE 3** Risk factors for severity of COVID-19. Multivariable binary logistic regression analyses were used to assess the association among age, sex, BMI, underlying comorbidity, history of smoking, alcohol consumption and the dependent variable of severity of disease (general group and refractory group). Age and BMI were considered risk factors for refractory pneumonia.
may contribute to a more severe inflammatory response in elderly male obese patients. In particular, the decrease in testosterone levels in older men and the increase in pro-inflammatory cytokines in obese patients may be particularly pronounced in older obese men, possibly exacerbating COVID-19 progression.

There were several limitations in the current study. First of all, we observed that there was a correlation between obesity and refractory pneumonia. The mechanism of obesity causing refractory pneumonia needs further research in the future. Second, Liu et al reported that the viral load of SARS-CoV-2 might be a useful marker for assessing disease severity and prognosis, while the viral load of SARS-CoV-2 was not be detected because of the emergency in progress and limited time availability. We will detect the viral load in the samples which are currently retained in the future and produce conclusions regarding to the relationship between viral load and disease characteristics and prognosis. Thirdly, this is a respective study; the findings need to be confirmed by a randomized controlled study in the future.

In this single-center case series study, we found that age and BMI were associated with disease severity of hospitalized COVID-19 patients in Beijing even after adjusting for other related potential confounders. Males with COVID-19 usually had more respiratory symptoms and abnormal laboratory results, such as CRP, FIB, AST, and CK than females while they were less aware of care-seeking than females. The findings of the current study showed that SARS-CoV-2 was more likely to affect older males with comorbidities. Further researches are still needed to explain the exact relationships between BMI and the severity of COVID-19 in male patients.

ACKNOWLEDGMENTS

The authors acknowledge all patients and the medical staffs in Beijing Ditan Hospital, Capital University.

CONFLICT OF INTEREST

Authors have no conflict of interests.

ETHICS APPROVAL AND CONSENT TO PARTICIPATE

This study was performed after approval by the Committee for Ethical Affairs of the Beijing Ditan Hospital (DTZZLX-202009).

AUTHOR CONTRIBUTIONS

Jing-Jing Wang, Yun-Juan Su, and Wen Xie conceived and designed the study. Jing-Jing Wang, Qi Wang, and Ai-Bin Wang extracted information and analyzed the data. Jing-Jing Wang and Yun-Juan Su wrote the manuscript. Ying Cao and Rui Ding reviewed the manuscript. All authors reviewed and approved the final version of the manuscript.

AVAILABILITY OF DATA AND MATERIALS

The datasets used and/or analyzed during the current study are available from the corresponding author.

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REFERENCES

1. Lu R, Zhao X, Li J, et al. Genomic characterisation and epidemiology of 2019 novel coronavirus: implications for virus origins and receptor binding. Lancet. 2020;395(10224):565–574.
2. Zhou P, Yang XL, Wang XG, et al. A pneumonia outbreak associated with a new coronavirus of probable bat origin. Nature. 2020;579:270–273.
3. Goel S, Jain T, Hooda A, et al. Clinical characteristics and in-hospital mortality for COVID-19 across The globe. Cardiol Ther. 2020;9:553–559.
4. van Lunzen J, Altfeld M. Sex differences in infectious diseases common but neglected. Infect Dis. 2014;209:579–580.
5. Kovats S. Estrogen receptors regulate innate immune cells and signaling pathways. Cell Immunol. 2015;294:63–69.
6. Jin JM, Bai P, He W, et al. Gender differences in patients with COVID-19: focus on severity and mortality. Front Public Health. 2020;8:152.
7. Sharma G, Volgman AS, Michos ED. Sex differences in mortality from COVID-19 pandemic: are men vulnerable and women protected?. JACC Case Rep. 2020;2:1407–1410.
8. Zhang L, Wang Z, Wang X, et al. Prevalence of overweight and obesity in China: results from a cross-sectional study of 441 thousand adults, 2012−2015. Obes Res Clin Pract. 2020;14:119–126.
9. Zhang X, Zhang M, Zhao Z, et al. Geographic variation in prevalence of adult obesity in China: results from the 2013–2014 national chronic disease and risk factor surveillance. Ann Intern Med. 2020;172:291–293.
10. National Health Commission. PRC. COVID 19 diagnosis and treatment program (trial version 7) [EB/OL]. 2020. http://www.nhc.gov.cn/yzwj/s7653p/202003/46c9_294a7dfe4ce80dc7f5912eb1989.shtml. Accessed March 03, 2020.
11. Karlberg J, Chong DS, Lai WY. Do men have a higher case fatality rate of severe acute respiratory syndrome than women do?. Am J Epidemiol. 2004;159:229–231.
12. Alghamdi IG, Hussain II, Almalki SS, Alghamdi MS, Alghamdi MM, El-Sheemy MA. The pattern of Middle East respiratory syndrome coronavirus in Saudi Arabia: a descriptive epidemiological analysis of data from the Saudi Ministry of Health. Int J Gen Med. 2014;7:417–423.
13. vom Steeg LG, Klein SL. SeXX matters in infectious disease pathogenesis. PLoS Pathog. 2016;12:e1005374.
14. Huang C, Wang Y, Li X, et al. Clinical features of patients infected with 2019 novel coronavirus in Wuhan, China. Lancet. 2020;395:497–506.
15. 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;323:1061–1069.
16. Guan WJ, Ni ZY, Hu Y, et al. Clinical characteristics of coronavirus disease 2019 in China. N Engl J Med. 2020;382:1708–1720.
17. Mo P, Xing Y, Xiao Y, et al. Clinical characteristics of refractory COVID-19 pneumonia in Wuhan, China. Clin Infect Dis. 2020. https://doi.org/10.1093/cid/ciaa270.
18. Lan J, Ge J, Yu J, et al. Structure of the SARS-CoV-2 spike receptor-binding domain bound to the ACE2 receptor. Nature. 2020;581:215–220.
19. Ma L, Xie W, Li D, et al. Effect of SARS-CoV-2 infection upon male gonadal function: A single center-based study. medRxiv. 2020. https://doi.org/10.1101/2020.03.21.20037267.
20. Kloc M, Ghobrial RM, Kubiak JZ. The role of genetic sex and mitochondria in response to COVID-19 infection. Int Arch Allergy Immunol. 2020;181:629–634.
21. Culebras E, Hernández F. ACE2 is on the X chromosome: could this explain COVID-19 gender differences?. Eur Heart J. 2020;41:3095.
22. Stelzig KE, Canepa-Escaro F, Schilio M, Berdnikovs S, Prakash YS, Chiarella SE. Estrogen regulates the expression of SARS-CoV-2 receptor ACE2 in differentiated airway epithelial cells. Am J Physiol Lung Cell Mol Physiol. 2020;318:L1280–L1281.
23. Walter LA, McGregor AJ. Sex- and Gender-specific observations and implications for COVID-19. West J Emerg Med. 2020;21:507–509.
24. Wei ZY, Qian HY, Huang J, Geng YJ. Pathogenesis and management of myocardial injury in coronavirus disease. Eur J Heart Fail. 2019;22:1994–2006.
25. Zhu X, Ge Y, Wu T, et al. Co-infection with respiratory pathogens among COVID-2019 cases. Virus Res. 2020;285:198005.
26. Levi M, Thachil J, Iba T, Levy JH. Coagulation abnormalities and thrombosis in patients with COVID-19. Lancet Haematol. 2020;7:e438–e440.
27. Cai Q, Chen F, Wang T, et al. Obesity and COVID-19 severity in a designated hospital in Shenzhen, China. Diabetes Care. 2020;43:1392–1398.
28. Petrakis D, Margină D, Tsarouhas K, et al. Obesity a risk factor for increased COVID19 prevalence, severity and lethality (Review). Mol Med Rep. 2020;22:9–19.
29. Bianchi VE. The anti-Inflammatory effects of testosterone. J Endocr Soc. 2018;3:91–107.
30. Spagnolo PA, Manson JE, Joffe H. Sex and gender differences in health: what the COVID-19 pandemic can teach us. Ann Intern Med. 2020;173:385–386.
31. Liu Y, Yan LM, Wan L, et al. Viral dynamics in mild and severe cases of COVID-19. Lancet Infect Dis. 2020;20:656–657.

How to cite this article: Wang J-J, Su Y-J, Wang Q, et al. Sex differences in clinical characteristics and risk factors for disease severity of hospitalized patients with COVID-19. MedComm. 2021;2:247–255. https://doi.org/10.1002/mco2.66