Exposure to ambient gaseous pollutant and daily hospitalizations for Sjögren’s syndrome in Hefei: A time-series study

Tian-Ping Zhang1, Li-Jun Wang2, Shan Wang3, Ping Wang4, Xiao-Hui Zhou5, Li Wang1, Chun-Mei Yang6* and Xiao-Mei Li1*

1Department of Rheumatology and Immunology, The First Affiliated Hospital of USTC, Division of Life Sciences and Medicine, University of Science and Technology of China, Hefei, China, 2Department of Infectious Diseases, The First Affiliated Hospital of Anhui Medical University, Hefei, China, 3Department of Rheumatology, The First People’s Hospital of Hefei (Binhu Hospital), Hefei, China, 4Department of Rheumatology and Immunology, the Third People’s Hospital of Hefei, Hefei, China, 5Department of Scientific Research, The First Affiliated Hospital of USTC, Division of Life Sciences and Medicine, University of Science and Technology of China, Hefei, China

Objective: Increasing evidence suggested that gaseous pollutants were associated with the development of autoimmune diseases, while there were few studies on the association between gaseous pollutants and Sjögren’s syndrome (SS). This study sought to assess the relationship between exposure to several gaseous pollutants and the hospitalizations for SS.

Methods: The data regarding SS hospitalizations, gaseous pollutants, and meteorological factors in Hefei from 2016 to 2021 were collected. A distributed lag non-linear model combined with a generalized linear model were adopted to analyze the association between gaseous pollutants and SS hospitalizations, and stratified analyses were also conducted.

Results: We detected significant associations between gaseous pollutants (NO2, SO2, O3, CO) and SS hospitalizations. Exposure to NO2 was linked with the elevated risk of hospitalizations for SS (RR=1.026, lag1 day). A positive correlation between CO exposure and hospitalizations for SS was found (RR=1.144, lag2 day). In contrast, exposure to SO2, O3 was respectively related to the decreased risk of hospitalizations for SS (SO2: RR=0.897, lag14 day; O3: RR=0.992, lag9 day). Stratified analyses found that female patients were more vulnerable to these gaseous pollutants. SS patients ≥ 65 years were more susceptible to NO2, CO exposure, and younger patients were more vulnerable to O3 exposure. In addition, exposure to O3, CO in cold season were more likely to affect hospitalizations for SS.
Introduction

Sjögren’s syndrome (SS) is a chronic, systemic autoimmune disease characterized by the impaired function of the exocrine glands, which lead to several clinical symptoms such as xerostomia and dry eye (1). Moreover, this disease could affect other systems in the body, primarily the nervous system, musculoskeletal system, kidneys, skin, lungs, and blood vessels (2). SS could be classified as primary SS (pSS) alone or secondary SS (sSS) according to the presence or absence of other autoimmune diseases, such as systemic lupus erythematosus (SLE), rheumatoid arthritis (RA), systemic sclerosis (SSc) (3). It is generally considered that pSS is the most common autoimmune disease after RA, with a prevalence of 0.1-4.8% among different populations (4). In addition to dry eyes and mouth, most patients with SS also experience pain, fatigue, anxiety, and other symptoms, which could seriously affect people’s physical health (5). At present, the etiology and pathogenesis of SS are not completely clear, and studies have shown that genetic variation and environmental factors greatly affect the occurrence and development of SS (6). It is important to note that the identified genetic risk variants only convey modestly increased risk, further suggesting that other factors were involved in SS development. For example, SS patients often point out the influence of the weather on their symptoms, and believe that wet weather, extreme cold could aggravate the symptoms (7, 8).

More attention had been focused on air pollution because of its harmful effects on health. Some gaseous pollutant, such as carbon monoxide (CO), sulphur dioxide (SO2) and nitrogen dioxide (NO2), might be involved in regulating many important biological processes, including the increase of free radicals in the body, activation of the immune system and triggering inflammation (9, 10). Previous studies showed that air pollutants could increase the risks of atopic dermatitis, diabetes mellitus, and multiple sclerosis (11–13). In addition, Zhao et al. found that exposure to high levels of NO2, SO2 could affect the risk of first admissions for SLE (14). Another study also reported that short-term exposure to NO2 and CO had some effect on acute gout hospitalizations (15). These studies suggested that air pollution had some effects on autoimmune disease.

Current studies confirmed a more important role of environmental risk factors in the pathogenesis of SS. For example, one study by Xin et al. (16) suggested that extreme environmental conditions exposure could increase the risk of the outpatient visits for SS. Nevertheless, the role of other environmental factors, such as gaseous pollutants, in the pathogenesis of SS also required further analysis. For all we know, available study on the association between gaseous pollutants and SS were still scarce in humid subtropical regions. We performed this time-series study to investigate the effect of several ambient gaseous pollutants (NO2, SO2, O3, CO) on daily hospitalizations for SS in Hefei City, and identified the susceptible subpopulations and season.

Material and methods

Study area and study population

This time-series study was a retrospective, ecological study, and carried out in Hefei (31°52′N, 117°17′E), which was located in the central part of Anhui Province. Hefei featured a subtropical humid monsoon climate with moderate rainfall, and covered a total area of 11,445.1 km² with 9.465 million population by the end of 2021.

The SS hospitalizations of this study were obtained from three hospitals which most SS patients sought medical help from, including the First Affiliated Hospital of University of Science and Technology of China, the First People’s Hospital of Hefei, and the Third People’s Hospital of Hefei, from 1 January 2016 to 31 December 2021 in Hefei. We first obtained the electronic medical records of all hospitalized patients from the hospital information system (HIS) of these hospitals, and identified SS patients based on their diagnosis in the electronic medical records. Moreover, all SS patients were diagnosed by senior rheumatologists according to the 2002 American European Consensus Group (AECG) classification criteria (17) and the following information of SS patients was collected, including age, gender, residential address, and date of hospitalization. Meanwhile, the SS patients who lived outside Hefei, had incomplete demographic information, and were

Conclusion: Our results demonstrated a significant association between exposure to NO2, CO and elevated risk of hospitalizations for SS, and SO2, O3 exposure might be linked to reduced risk of SS hospitalizations.

KEYWORDS
gaseous pollutants, Sjögren’s syndrome, time-series study, autoimmune disease, air pollutant
hospitalized in other departments other than the department of rheumatology and immunology were excluded from this study. The ethics approval was granted by the Ethics Committee of The First Affiliated Hospital of University of Science and Technology of China (2022-RE-037).

Pollutants and meteorological data

In this study, we used the air pollutants concentration of the residential address of SS patient to reflect their amount of exposure to air pollutants. Air pollution data (24-h for NO2, SO2, CO, and 8-h for O3) from 2016 to 2021 were collected through the environmental monitoring network (including 10 air quality monitoring stations) established by Hefei Environmental Monitoring Center. The daily air pollutant data was an average of 10 monitoring stations. Data on daily relative humidity (RH, %) and mean temperature (°C) during this study period were obtained from the China Meteorological Data Service Center (http://data.cma.cn/).

Statistical analysis

The effect of gaseous pollutants on hospitalizations for SS was investigated with a distributed lag non-linear model (DLNM) combined with a generalized linear model in this study. These two models were respectively used to describe the additional lag-response correlation and the traditional exposure-response correlation. Since the daily admissions for SS was considered as a small probability event, thus, the DLNM with a quasi-Poisson distribution was adopted to investigate the association between gaseous pollutant and SS hospitalizations (15). Spearman analysis was used to analyze the correlation between each covariate, and two variables whose correlation coefficient less than 0.7 could not enter in the same model for avoiding multicollinearity. Finally, the model for NO2, SO2, CO, O3 were shown as below:

\[
Y_t \sim \text{quasipoisson}(\mu_t)
\]

\[
\log(\mu_{t1})_{\text{NO2}} = \alpha_{t1} + \beta_{1,24} \text{NO2}_{t1} + \text{ns}(\text{SO2}, 3) + \text{ns}(\text{RH}, 3) + \text{ns}(\text{time}, 6^6) + \gamma_{1, \text{DOW}} + \gamma_{1, \text{Holiday}}
\]

\[
\log(\mu_{t2})_{\text{SO2}} = \alpha_{t2} + \beta_{2,24} \text{SO2}_{t1} + \text{ns}(\text{NO2}, 3) + \text{ns}(\text{MT}, 3) + \text{ns}(\text{RH}, 3) + \text{ns}(\text{time}, 8^6) + \gamma_{2, \text{DOW}} + \gamma_{2, \text{Holiday}}
\]

\[
\log(\mu_{t3})_{\text{O3}} = \alpha_{t3} + \beta_{3,24} \text{O3}_{t1} + \text{ns}(\text{SO2}, 4) + \text{ns}(\text{MT}, 4) + \text{ns}(\text{RH}, 4) + \text{ns}(\text{time}, 6^6) + \gamma_{3, \text{DOW}} + \gamma_{3, \text{Holiday}}
\]

\[
\log(\mu_{t4})_{\text{CO}} = \alpha_{t4} + \beta_{4,24} \text{CO}_{t1} + \text{ns}(\text{O3}, 3) + \text{ns}(\text{RH}, 3) + \text{ns}(\text{time}, 7^6) + \gamma_{4, \text{DOW}} + \gamma_{4, \text{Holiday}}
\]

The subscript t referred to the day of observation, Yt and \( \mu_t \) were the actual and expected SS hospitalizations on day t, respectively. In above models, \( \alpha \) represented the intercept of the model. In NO2 model, \( NO2_t \) was the dlnm cross basis matrix of NO2, \( I \) represented the lag day, \( \beta \) was the vector of NO2, ns() means natural cubic spline. A natural cubic spline curve of time with 6 dfs/year was adopted to control the seasonality and long-term trend (18). DOW was the dummy variable of the day of week and the two-category variable \( \text{Holiday} \) was used to control the effect of holidays. Quasi-Poisson Akaike Information Criterion (Q-AIC) was preferred to identify the optimal dfs and determined the final model parameters. In order to identify the susceptible populations, further stratified analyses were conducted according to different age (<65 years vs ≥65 years), gender (male vs female). In addition, the relationship between gaseous pollutants and SS hospitalizations was respectively analyzed among hot season (April-September) and cold season (October-March). The statistical analyses were conducted using R software version 3.6.1 (http://www.R-project.org) with “dlnm” and “splines” packages, and \( P<0.05 \) (two-sided) was considered as statistically significant.

Sensitivity analyses

Sensitivity analyses were performed by varying the dfs in the ns function for gaseous pollutants (3-5 dfs), meteorological variables (3-5 dfs) and time (6-8 dfs per year) in turn in the model to verify the robustness of our models.

Results

Descriptive analysis

A total of 1119 SS hospitalizations were collected in Hefei from 1 January 2016 to 31 December 2021, and the characteristics of these SS patients, ambient gaseous pollutant and meteorological factors were shown in Table 1. Among the SS patients, 1061 cases were female (94.8%), 350 cases (31.3%) were aged 65 years or older, and there were more hospital admissions for SS in hot season (580, 51.8%). The daily mean values for NO2, CO, SO2, and O3 (24-h for NO2, CO, SO2, and 8-h for O3) were 42.09 μg/m3 (range: 9 μg/m3 – 137 μg/m3), 0.80 mg/m3 (range: 0.3 mg/m3 – 2.8 mg/m3), 8.97 μg/m3 (range: 2 μg/m3 – 58 μg/m3) and 94.57 μg/m3 (4 μg/m3 – 269 μg/m3). The number of daily hospital admissions for SS ranged from 0 to 6 during the study period. Figure 1 presented the temporal trends of gaseous pollutants, mean temperature, and hospitalizations for SS from 2016 to 2021 in Hefei. The results of Spearman rank correlation analysis and scatter plot were arranged in Figure S1, and the correlation coefficients between any two variables are less than 0.7.

The association between gaseous pollutants and SS hospitalizations

Overall effects

The exposure-response relationships between SS hospitalizations and gaseous pollutants (NO2, SO2, O3, and CO)
in different lag days were presented in Figure 2. The results indicated a meaningful positive correlation between high concentrations of NO₂ and CO (reference concentrations of 38 \( \mu \text{g/m}^3 \) and 0.7 mg/m³, respectively) and the elevated risk of SS hospitalizations, while exposure to high concentration of SO₂ and O₃ (reference concentrations of 8 \( \mu \text{g/m}^3 \) and 90 \( \mu \text{g/m}^3 \), respectively) could reduce the risk of SS hospitalization. In addition, the concentration-response relationships between NO₂, SO₂, O₃, CO and SS hospitalizations were shown in Figure S2.

**Effects of NO₂ on SS hospitalizations**

There was a significant single-day effect between NO₂ and SS hospitalizations with every 10 \( \mu \text{g/m}^3 \) increase in NO₂ concentration from lag1 (RR: 1.026, 95% CI: 1.004-1.048) to lag3 (RR:1.018, 95% CI: 1.002-1.034), and the highest RR of SS admissions was at lag1 (Figure 3; Table S1). Moreover, the meaningful estimated risk effects were found from lag0-2 (RR: 1.079, 95% CI: 1.009-1.153) to lag0-8 (RR:1.094, 95% CI: 1.005-1.191) in the cumulative lag structure, and the highest RR was found at lag5 (RR: 1.118, 95% CI: 1.041-1.201) (Figure 4; Table S1). When stratified by gender, age, season, the results suggested and NO₂ exposure was positively correlated with the number of SS hospitalizations in female (RR:1.027, 95% CI: 1.005-1.050, lag1), the patients \( \geq 65 \) years (RR: 1.055, 95% CI: 1.031-1.080, lag2), hot seasons (RR:1.031, 95% CI: 1.003-1.060, lag1), and cold seasons (RR: 1.014, 95% CI: 1.000-1.027, lag2) (Figure 5; Table S2).

![Figure 1](image1.png)

**FIGURE 1**
The time series of gaseous pollutants, mean temperature, and hospitalizations for SS from 2016 to 2021 in Hefei.
Effects of SO2 on SS hospitalizations

For every 10 μg/m³ increase in SO2 concentration, the association between SO2 and the daily hospitalizations for SS was statistically significant from lag9 (RR: 0.944, 95% CI: 0.896-0.994) to lag14 (RR: 0.897, 95% CI: 0.807-0.998) in the single-day lag structure, and the lowest RR was found at lag14 (Figure 3; Table S3). In multi-day cumulative effect, the risk of SS hospitalization decreased with increasing SO2 over the lag0-11 (RR: 0.536, 95% CI: 0.302-0.952) to lag0-14 (RR: 0.400 95% CI: 0.199-0.803), and the lowest RR was also at lag0-14 (Figure 4; Table S3). In addition, the effects of SO2 exposure on SS hospitalizations were statistically significant in female (RR: 0.925, 95% CI: 0.864-0.991, lag12), the patients < 65 years (RR: 0.917, 95% CI: 0.852-0.983, lag4), the patients ≥ 65 years
(RR: 0.771, 95% CI: 0.636-0.934, lag14), hot seasons (RR: 0.824, 95% CI: 0.711-0.955, lag14), and cold seasons (RR: 0.947, 95% CI: 0.902-0.994, lag11) (Figure 6; Table S4).

Effects of O₃ on SS hospitalizations
An inverse single-day single day lag association was found between O₃ and the risk of SS hospitalizations (lag8, RR: 0.993, 95% CI: 0.988-0.999, and lag9, RR: 0.992, 95% CI: 0.985-0.999, per 10 μg/m³ increase in O₃ concentration) (Figure 3; Table S5).

In the cumulative lag structure, the estimated risk effects decrease from lag0-8 (RR: 0.956, 95% CI: 0.919-0.995) to lag0-10 (RR: 0.940, 95% CI: 0.900-0.981) (Figure 4; Table S5). The results of the subgroup analyses also revealed that the relation between O₃ exposure and a decreased risk of SS hospitalizations was consistently significant in female (RR: 0.992, 95% CI: 0.984-0.999, lag9), the patients < 65 years (RR: 0.991, 95% CI: 0.984-0.998, lag7), and cold seasons (RR: 0.965, 95% CI: 0.934-0.997, lag1) (Figure 7; Table S6).
Effects of CO on SS hospitalizations
The single-day lag association between CO and SS hospitalizations was statistically significant at lag2 (RR: 1.144, 95% CI: 1.023-1.278) with every 1 mg/m³ increase in CO concentration (Figure 3; Table S7), and the multi-day cumulative risk was significant at lag0-4 (RR: 1.607, 95% CI: 1.006-2.566) (Figure 4; Table S7). Interestingly, this study also observed a significant relationship between CO and an increased SS hospitalizations in female (RR: 1.160, 95% CI: 1.035-1.299, lag2), the patients ≥ 65 years (RR: 1.227, 95% CI: 1.007-1.496, lag2), and cold seasons (RR: 1.160, 95% CI: 1.040-1.293, lag3) (Figure 8; Table S8).

Sensitivity analyses
The sensitivity analyses results suggested that the effects of gaseous pollutants on SS hospitalizations were stable under df variations for time, gaseous pollutants, meteorological factors. Hence, these models were considered to be reliable and robust (Figures S3–S6).

Discussions
Air pollution, including gaseous pollutants, had become an important public problem threatening human health. Several
studies had suggested that there was a close association between exposure to gaseous pollutants and the risk of a variety of autoimmune diseases (14, 15, 19), which provided important clues for further research on the pathogenesis of autoimmune diseases. The results of a recent study showed that cold, damp and long sunshine duration might associated with the number of SS patients visiting hospitals, and the effects were influenced by age, sex (16). This suggested that the role of environmental factors in the pathogenesis of SS deserved more attention. In this study, we identified significant associations between exposure to gaseous pollutants and the hospitalizations for SS. Exposure to NO2, CO was positively related to the risk of hospitalizations for SS, while exposure to SO2, O3 was negatively related to the risk of hospitalizations for SS.

As common gaseous pollutants produced by traffic, exposure to high concentrations of NO2, CO might cause various adverse health effects. One cohort study showed that the average exposure levels of O2, CO in 5-year prior to RA diagnosis were significantly associated with the high incidence of RA (20). A time-series study by Wu et al. found that exposure to high-concentration NO2 was significantly related to an increased risk of RA readmissions (21). Severe dryness of the mucosal surface in eyes and oral cavity were the primary clinical manifestation among SS patients, and the patients with SS who were exposed to air pollution was associated with more severe abnormalities of the ocular surface and eye irritation (22). In addition, environmental NO2 concentration was proved to be associated with dry eye syndrome (23). This might support our results that exposure to NO2 was associated with an increased risk of daily hospitalizations for SS. Moreover, we found a promoting effect of CO exposure on the risk of hospitalization for SS. This effect could also be observed in another autoimmune disease, where adults exposed to CO for one year had a higher risk of developing RA in a population-based cohort study (24). At present, the biological mechanism by which NO2 and CO affect the risk of SS remained unclear. It was possible that exposure to these pollutants might cause oxidative stress and increase the release of proinflammatory mediators within the lung and systemically. Chronic inflammation of the lungs might promote the susceptibility to diseases characterized by inflammation, which was considered a suitable biomarker of exposure to relevant gaseous pollutants, and might contribute to the occurrence of rheumatic diseases (20). We hypothesized that exposure to high concentrations of CO and NO2 might be involved in some inflammatory processes and promoted the onset of inflammation in SS patients.

Among the gaseous pollutants, the lower level of O3 and high concentrations of nitrogen oxides often appeared in proximity to high-traffic areas (25). Several studies had suggested significant associations between O3 exposure and the risk of human diseases. One previous study found that O3 exposure could reduce the risk of interstitial lung disease in pSS patients (26). In this study, our results showed the protective effect of O3 exposure against the risk of SS hospitalizations. Exposure to SO2 were also negatively related to the risk of SS hospitalizations in the present study. Similarly, the study by Jung et al. found that exposure to increased O3 and SO2 were negatively associated with the increased risk of SLE (27). These results suggested that SO2 and O3 exposures had certain protective effect on the risk of medical visits for autoimmune diseases included SS, while the specific mechanism was still unclear due to the lack of relevant studies. Intriguingly, O3 was involved in the high Th2 response in airway cells by enhancing the type 2 innate lymphoid cell (ILC)-related pathways (28). Therefore, the O3-associated expansion of the Th2 pathway through ILCs might partially explain the potential protective effect of O3 on SS development by ameliorating Th17/Th1-related signaling pathways. In addition, ground-level O3 was
a secondary pollutant produced by photochemical reactions between traffic-related air pollutants, including NO2 and volatile organic compounds (26). Our results also implied that the primary pollutants (NO3 and CO) had greater importance on the SS hospitalizations in comparison to the secondary pollutants (O2) produced by photochemical processes.

The stratified analyses showed that the effects of NO2, SO2, O3, and CO exposure on SS hospitalizations remained significant in female SS patients but not in male patients. Women tend to exhibit higher immune reactivity, with different numbers or reactivity of cells constituting the immune responses and different resistance to target organ damage, which might result in different effects of air pollutants on the risk of autoimmune diseases among male and female patients (29). NO2, CO exposure were associated with higher risk of SS hospitalizations in elderly patients, while O3 exposure was linked to lower risk of SS hospitalizations in young patients. This might be caused by aging leading to the alterations in the function of Toll like receptors (30), which were potent inflammation mediators in SS and played an important function in salivary tissue. We also found a statistically significant association between CO exposure and the risk of SS hospitalizations during the cold season. This might be associated with the dry environment in cold season, and exposure to dry environment could result in the deterioration of tear function unit in SS patients with dry eye symptom through promoting inflammatory activity (31).

There were several strengths in this study. First, as we all know, this was the first time-series study to examine the association between exposure to gaseous pollutants and hospitalizations for SS. Second, stratified analyses by gender, age and season were conducted to identify the populations and seasons that were particularly vulnerable to gaseous pollutants. In addition, we acknowledged that some limitations existed in this study. First, we only collected SS hospitalizations data from three hospitals in Hefei, which might not be able to represent SS inpatients in Hefei. Another limitation was that the design of this study had an ecological fallacy, which might restrict the ability of our study to explore the causal relationships. Finally, the association between SS activity, complications and gaseous pollutants was not analyzed because of missing data. In order to analyze the precise mechanisms responsible for the associations between gaseous pollutants and SS, further studies based on epidemiological investigations and functional investigations should be conducted.

In summary, our study provided key evidence that exposure to NO2, CO was positively related to the increased risk of hospitalizations for SS in Hefei. While, a negative association was identified between exposure to SO2, O3 and SS hospitalizations. Stratified analyses found that the effect of these gaseous pollutants on risk of SS hospitalizations could be modified by gender. The effect of NO2, CO, O3 exposure on SS hospitalizations was affected by age, and the relationship between CO, O3 exposure and SS hospitalizations remained significant in cold season. This study was of great significance for further explore the role of gaseous pollutants in the pathogenesis of SS.

Data availability statement

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

Ethics statement

This study was approved by the Ethical Committee of the First Affiliated Hospital of USTC (Hefei, Anhui, China).

Author contributions

X-ML, C-MY and T-PZ designed the study. SW, PW, X-HZ and LW participated in the data collection. L-JW performed data processing. T-PZ conducted the data analysis and drafted the manuscript. X-ML and C-MY contributed to manuscript revision. All authors contributed to the article and approved the submitted version.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fimmu.2022.1028893/full#supplementary-material
