**RESEARCH ARTICLE**



# **Modifcation efects of ambient temperature on ozone‑mortality relationships in Chengdu, China**

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#### **Abstract**

A multitude of epidemiological studies have demonstrated that both ambient temperatures and air pollution are closely related to health outcomes. However, whether temperature has modifcation efects on the association between ozone and health outcomes is still debated. In this study, three parallel time-series Poisson generalized additive models (GAMs) were used to examine the efects of modifying ambient temperatures on the association between ozone and mortality (including non-accidental, respiratory, and cardiovascular mortality) in Chengdu, China, from 2014 to 2016. The results confrmed that the ambient high temperatures strongly amplifed the adverse efects of ozone on human mortality; specifcally, the ozone effects were most pronounced at > 28 °C. Without temperature stratification conditions, a 10-μg/m<sup>3</sup> increase in the maximum 8-h average ozone ( $O_{3-8hmax}$ ) level at lag01 was associated with increases of 0.40% (95% confidence interval [CI] 0.15%, 0.65%), 0.61% (95% CI 0.27%, 0.95%), and 0.69% (95% CI 0.34%, 1.04%) in non-accidental, respiratory, and cardiovascular mortality, respectively. On days during which the temperature exceeded 28 °C, a 10- $\mu$ g/m<sup>3</sup> increase in O<sub>3–8hmax</sub> led to increases of 2.22% (95% CI 1.21%, 3.23%), 2.67% (95% CI 0.57%, 4.76%), and 4.13% (95% CI 2.34%, 5.92%) in non-accidental, respiratory, and cardiovascular mortality, respectively. Our fndings validated that high temperature could further aggravate the health risks of  $O_{3-8hmax}$ ; thus, mitigating ozone exposure will be brought into the limelight especially under the context of changing climate.

Keywords Temperature · Modification effect · Ozone · Mortality · Chengdu



## **Introduction**

It is universally acknowledged that air pollution has adverse efects on human health; among all air pollutants, particles with aerodynamic diameters less than 2.5  $\mu$ m (PM<sub>2.5</sub>) and ground-level ozone are considered to be most extraordinarily associated with morbidity and mortality (Dimakopoulou et al. [2017](#page-7-0); Vicedo-Cabrera et al. [2020;](#page-8-0) Zhang et al. [2019a,](#page-8-1) [2020b;](#page-8-2) Cohen et al. [2015\)](#page-7-1). China, the largest developing country, is currently facing a serious situation regarding air pollution, and  $PM_{2.5}$  and ozone are the first and second major pollutants, respectively (Kan et al. [2012](#page-7-2); Wang [2021](#page-8-3)). Over the past decades, most studies have focused on the health risks of  $PM_{2.5}$ ; subsequently, a series of corresponding policies regarding  $PM<sub>2.5</sub>$  control and intervention measures have been implemented based on these research results, leading to the  $PM<sub>2.5</sub>$  level being effectively controlled in China (Lu et al. [2019](#page-8-4); Maji et al. [2020](#page-8-5)). However, evidence concerning the adverse effects of ozone on health outcomes has been limited due to a lack of ozone data availability. No data

were published concerning ozone before 2008. The Ministry of Ecology and Environment of the People's Republic of China set the subsequent ozone air quality standards, and ground-level ozone data have been available online since 2013 [\(https://www.mee.gov.cn/](https://www.mee.gov.cn/)). Compared with other countries, it is difficult to evaluate ozone standards in China due to the lack of sufficient evidence regarding the health effects of ozone. Recently, environmental monitoring observations have indicated that the concentration of  $PM<sub>2.5</sub>$  in the Sichuan Basin of China is decreasing yearly, whereas the concentration of surface ozone is increasing (Ning et al. [2017\)](#page-8-6); these results aroused our concern regarding the health risks caused by ozone in this area.

The ambient temperature is another important health risk factor. Substantial epidemiological and toxicological literature has been published showing a clear and consistent association between exposure to adverse ambient temperature, especially in cases of hypothermia and hyperpyrexia, both of which can cause a series of acute health efects, including respiratory tract injury, chronic cardiovascular conditions, systemic infammation, and premature mortality (Wang [2021](#page-8-3); Dimakopoulou et al. [2017;](#page-7-0) Qian et al. [2020\)](#page-8-7). At the same time, the spatiotemporal distribution of ambient pollutants is afected by meteorological conditions, especially the ambient temperature (Zhang et al. [2019b\)](#page-8-8); hence, ambient temperature and air pollution are generally highly correlated in many places and may symmetrically interact to afect health outcomes (Bae et al. [2020](#page-7-3); Chen et al. [2017](#page-7-4)). Nonetheless, related studies about the modifcation efects of the ambient temperature on ozone-mortality relationships are rare, and published results have been inconsistent: some researchers have claimed a strong enhancement of ozone risks on health outcomes only under high temperature levels, whereas others have found outstandingly increases only under low temperature conditions, and some have validated relatively high ozone risks for both high and low temperatures (Iny et al. [2014](#page-7-5); Chen et al. [2018b,](#page-7-6) [a](#page-7-7); Ren et al. [2007](#page-8-9)). These inconsistencies may correspond to the diferent climate and topographic conditions, ozone distribution characteristics, demographic compositions, people's lifestyles, and education levels of diverse areas (Li et al. [2018](#page-7-8)). As the capital city of Sichuan province, the population of Chengdu exceeded 16.04 million as of June 2018, and the city sufers serious ozone pollution, especially in summer (Yang et al. [2021](#page-8-10)). The modulatory effects of ambient temperatures on ozone-mortality relationships in this area are still unclear.

In the present study, we assessed ozone mortality risks on non-accidental, respiratory, and cardiovascular mortality in Chengdu, China, from 2014 to 2016. In addition, we explored whether the associations between ozone and nonaccidental mortality as well as cause-specifc mortality were modifed by ambient temperature. To achieve this aim, three parallel time-series Poisson generalized additive models (GAMs) were used to estimate how the air temperature modulates the health risks of ozone on the three analyzed kinds of mortality.

### **Data and methods**

#### **Data collection**

The daily cause-specifc mortality count data for each district/county of Chengdu (including Jinjiang District, Qingyang District, Jinniu District, Wuhou District, Chenghua District, Longquanyi District, Qingbaijiang District, Xindu District, Wenjiang District, Shuangliu District, and Xinjin County) during 2014 to 2016 were obtained from the National Center for Chronic and Non-communicable Disease Control and Prevention (NCNCD) of the Chinese Centers for Disease Control and Prevention (China CDC). The death data come from 421 hospitals with diferent grades in Chengdu. According to the 10th version of the International Classifcation of Diseases (ICD-10), data on three types of deaths were collected: non-accidental causes (ICD-10 codes A00-R99), cardiovascular diseases (ICD-10 codes I00–I99), and respiratory diseases (ICD-10 codes J00–J99). The total number of non-accidental, respiratory, and cardiovascular cases was 243,135, 76,721, and 59,676, respectively.

The air pollution data were retrieved from the Chengdu Environmental Monitoring Center. There are six monitoring stations in Chengdu including three urban environmental monitoring stations (Jinquan Lianghe, Sanwayao, and Shahepu), two traffic pollution monitoring stations (Shilidian, Liangjiaxiang), and one suburban environmental monitoring station (Lingyansi). Previous studies have indicated that the daily maximum 8-h average ozone  $(O_{3-8hmax})$  concentration is more strongly associated with health outcomes than other metrics, such as the 1-h maximum ozone concentration or the daily average ozone concentration (Yang et al. [2012](#page-8-11)). Consistent with the preceding studies, we therefore chose the daily  $O_{3-8hmax}$  as the ozone concentration indicator. We collected the daily 24-h mean concentrations of  $PM_{10}$ ,  $PM_{2.5}$ ,  $SO_2$ , and  $NO_2$  in Chengdu. To calculate the daily concentrations of the diferent pollutants at a single station, at least 75% of the hourly values had to be available on a particular day for each pollutant. Moreover, to calculate the 8-h maximum ozone concentration, an extra restrictive condition was that at least 6 h of data had to be available in each 8-h period within a day. The daily concentrations were the mean value of the data from the available monitoring results of six monitoring stations.

Daily meteorological data recorded during the same time period were retrieved from the China Meteorological Data Sharing Service System (<http://data.cma.cn/>). The daily surface meteorological data of Chengdu were obtained through station averages, mainly including the daily average temperature (°C), daily average relative humidity (*RH*) (%), and daily average wind speed (m/s).

#### **Statistical methods**

We utilized three GAMs to assess how the air temperature modulates the health risks of  $O_{3-8hmax}$  on health outcomes: an independent model, a nonparametric bivariate response surface model, and a stratifcation parametric model (Zhang et al. [2020b](#page-8-2)). First, we used an independent GAM to investigate the adverse health effects of  $O_{3-8hmax}$  on non-accidental mortality, as well as cause-specifc (respiratory and cardiovascular) mortality at diferent lag days. Model 1 can be expressed as follows:

Temperatures were categorized into high and low temperature levels and then to determine whether the  $O_{3-8hmax}$  risks varied across diferent temperature strata. However, there is by no means uniform standard for choosing temperature cutoff points so far (Zhang et al.  $2020b$ ). According to previous research results, people feel more comfortable when the daily average temperature is near approximately 24℃ and begin to feel uncomfortable due to heat when the daily average is  $>$  28 °C (Ssl et al., [2019\)](#page-8-12). Therefore, we chosen temperature cutoff points corresponding to the transition from comfort to discomfort with increasing heat (i.e., 24 °C, 26 °C, and 28 °C), as outlined by Zhang et al. ([2020b](#page-8-2)). Finally, we divided the temperature data into two strata, including high temperatures (above the cutofs) and low temperatures (below the cutoffs). Then, we assessed



where subscript *t* is the day of the observation;  $E(Y_t|X)$  indicates the expected death counts on day *t*; and *NS*(∙) denotes the natural cubic spline function. According to the preceding study (Yang et al. [2012](#page-8-11)), the *Time* variable was controlled with 7 degrees of freedom (*df*) per year to express a long-term trend. At the same time, (*df*=3) was also used to control the current day's average relative humidity (*RH*) and mean wind speed (*Wind*), as well as the moving average daily mean temperature of the current and the previous day (lag01) (Zhang et al. [2020b](#page-8-2)). *DOW* and *Holiday* are two categorical variables that represent the day of the week and the presence of a public holiday, respectively. The subscript *i* represents diferent lag days. We explored the adverse health effects of  $O_{3-8hmax}$  on three types of mortality at diferent lag days which include both a single-day lag (from lag0 to lag4) and cumulative lags (using moving averages of the current day and the previous 1, 2, 3, or 4 days [lag01 to lag04]). $\alpha$  represents the intercept. In addition, *COVs* is all other covariates.

Second, a nonparametric bivariate response surface model was adopted to visually examine the combing efects of both the ambient average temperature and  $O_{3-8hmax}$  on three types of mortality. The model is described as follows:

$$
log[E(Yt|X)] = ST(Temp, O3-8hmax) + COVs
$$
 (2)

where *ST*(∙) indicates the thin-plate spline functions. Statistics revealed that the adverse health risks of  $O_{3-8hmax}$  on three types of mortality were strongest between the current and previous day (lag01) (as is described later in this study). Therefore, the daily  $O_{3-8hmax}$  at lag01 was used in the followup research to efectively capture the overall efects. The *COVs* are the same as those used in model 1.

Finally, we adopted a temperature-stratifed parametric model to examine the heterogeneity of  $O_{3-8hmax}$  risks across diferent temperature ambient temperature strata.

how the  $O_{3-8h\text{max}}$  mortality risks varied among the different temperature levels using varied temperature cutoff points. Model 3 is given as follows:

$$
\log \left[ E(Y_t|X) \right] = \beta_1 \mathcal{O}_{3-8hmax} + \beta_2 Temp_k + \beta_3 (\mathcal{O}_{3-8hmax} : Temp_k) + \text{COVs}
$$
 (3)

where  $Temp_k$  is the *k*-th temperature strata;  $\beta_1$  and  $\beta_2$  denote the main independent effects of  $O_{3-8hmax}$  levels and daily average temperature, respectively; and  $\beta_3$  refers to a vector of coefficients reflecting the conjunction effects between  $O_{3-8hmax}$ and daily average temperatures; this vector was also adjusted for temperature  $(Temp_k)$  within each temperature strata. The COVs are the same as those used in model 1.

To test the robustness of the models, we performed several sensitivity analyses. First, we changed the *df* of time from 4 to 10 per year. Second, we changed the *df* of Temp, RH, and Wind from 3 to 5, respectively. Third, we applied diferent maximum lags for temperature (1, 3, 5, and 7 days, respectively). Moreover, we used co-pollutant models of other pollutants, such as  $PM_{2.5}$ ,  $PM_{10}$ ,  $SO_2$ ,  $NO_2$ , and CO to account for potential confounding efects from multiple exposures. All of the above analyses were conducted when the temperature levels were stratifed by using 24℃.

All statistical analyses were conducted with R 4.1.2. The estimated modulating effects and corresponding 95% confdence interval (CI) were showed as percentage changes in health outcomes with each  $10$ - $\mu$ g/m<sup>3</sup> increment in the  $O_{3-8hmax}$  mass concentration.

#### **Results**

Table [1](#page-3-0) summarizes the distributions of three types of mortality, meteorological factors, and  $O_{3-8h\text{max}}$  concentrations during the study period. There were considerable

variations in the three mortality types, ranging from 144 to 430 non-accidental mortalities, 35 to 136 cardiovascular mortalities, and 23 to 121 respiratory mortalities. During the study period, the daily mean temperature range between − 1.9 and 29.8℃ and there were no any heat waves (Fig S1). According to statistical analysis, the average temperature in Chengdu was  $16.6 \pm 7.2$  °C, the average *RH* was  $81.7 \pm 8.2\%$ , and the average wind speed was  $1.3 \pm 0.5$  m/s. Notably, the average daily  $O_{3-8hmax}$  concentration was  $119.9 \text{ µg/m}^3$  that quite higher than those reported in some developed countries (Tao et al. [2016](#page-8-13); Nyssanbayeva et al. [2019](#page-8-14); Winiewski et al. [2021\)](#page-8-15) and in other Chinese cities (Sui, et al. [2021;](#page-8-16) Zhang et al. [2006](#page-8-17)). According to the National Ambient Air Quality Standard (GB3095-2012), the  $O_{3-8hmax}$  concentrations exceeded the primary (100  $\mu$ g/m<sup>3</sup>) and secondary (160  $\mu$ g/m<sup>3</sup>) standard limits in 611 days and 280 days, respectively, and the corresponding over standard rates were 55.75% and 25.55%, respectively.

Figure [1](#page-3-1) illustrates the efects of percentage changes in  $O_{3-8h\text{max}}$  on three types of mortality at different lags. The most significant effects of  $O_{3-8hmax}$  on the three types of mortality all appeared at a cumulative lag of one day (lag01).

Therefore, lag01  $O_{3-8hmax}$  was used as the research object in subsequent studies. After the calculations, a  $10$ -μg/m<sup>3</sup> increase in  $O_{3-8hmax}$  was found to lead to 0.40% (95% CI 0.15%, 0.65%), 0.61% (95% CI 0.27%, 0.95%), and 0.69% (95% CI 0.34%, 1.04%) increases in non-accidental, respiratory, and cardiovascular mortality, respectively.

Figure [2](#page-4-0) graphically depicts the combined efects of the daily average temperature and  $O_{3-8hmax}$  on non-accidental, cardiovascular, and respiratory mortality using three-dimensional visualization graphs. It is apparent that the combined efects were extremely complex. It is interesting to note that the non-accidental, cardiovascular, and respiratory mortality all reached their maxima when high-temperature and highconcentration  $O_{3-8hmax}$  coexisted, thus indicating that the high temperature exacerbated/amplifed the mortality risks of  $O_{3-8hmax}$ .

Table [2](#page-4-1) summarizes the modulating efects of low/high temperatures on  $O_{3-8hmax}$ -mortality relationships using varied temperature cutoff points. It should also be noted that the modulation effects of temperature on  $O_{3-8hmax}$ -mortality were more pronounced in the high-temperature section than in the low-temperature section, and these efects were stronger than those obtained in the independent effect model.



*SD*, standard deviation;  $O_{3-8hmax}$ , 8-h maximum ozone concentration

<span id="page-3-1"></span>**Fig. 1** Percentage changes (%) in mortality for every 10-μg/  $m<sup>3</sup>$  increase in O<sub>3–8hmax</sub> at both single lag times of  $0 \sim 4$  days and cumulative lag times of  $01 \sim 04$  days

<span id="page-3-0"></span>**Table 1** Summary statistics of three types of mortality, meteorological factors, and  $O_{3-8h\text{max}}$  concentrations in Chengdu, China, from 2014 to

2016



<span id="page-4-0"></span>**Fig. 2** Bivariate response

O3–8hmax on health outcomes



Furthermore, the higher the temperature cutoff points were, the greater the health risks of  $O_{3-8hmax}$  were on the same kind of mortality at a high temperature level, indicating that high temperatures signifcantly aggravated the health risk of  $O_{3-8hmax}$  on mortality compared to low temperatures.

The sensitivity analyses turned out that the O3–8hmax-mortality risks kept robust to changing the *df* of the temporal smoothness per year (Fig. [3\)](#page-5-0). The percentage changes derived per 10- $\mu$ g/m<sup>3</sup> increment in O<sub>3-8hmax</sub> signifcantly increased from 0.35 (95% CI 0.24%, 0.46%) to 0.40% (95% CI 0.27%, 0.53%) for non-accidental mortality, 0.48 (95% CI 0.26%, 0.70%) to 0.61% (95% CI 0.35%, 0.87%) for respiratory mortality, and 0.62 (95% CI 0.39%, 0.85%) to 0.69% (95% CI 0.46%, 0.92%) for cardiovascular mortality when the *df* was changed from 4 to 10 per year. The changes of *df* for *Temp*, *RH*, and *Wind*, and the maximum lag days of mean temperature did not substantially afect the magnitude of the estimates (see Table S1, S2), and the same patterns were found in the co-pollutant models with the inclusion of  $PM_{2.5}$ ,  $PM_{10}$ ,  $SO_2$ ,  $NO_2$ , and CO (see Table S3), suggesting that the ozone-mortality relationship was not confounded by other pollutants.

## **Discussion**

Ground-level ozone has become a compelling environmental problem that has drawn substantial attention worldwide (Stocker et al. [2013\)](#page-8-18). Assessing ground-level ozone health efects could provide additional evidences for policymaking

<span id="page-4-1"></span>



 $\binom{a}{P}$  < 0.05



<span id="page-5-0"></span>



on the topic of ozone control measures, particularly under the background of climate change (Madaniyazi et al., [2016](#page-8-19)). Our findings proved that exposure to ground-level  $O_{3-8hmax}$ were positively associated with non-accidental mortality as well as cardiovascular and respiratory mortality in Chengdu, China, during the study period. Furthermore, our study further validated that high temperature signifcantly amplifed  $O_{3-8hmax}$ -mortality risks on the three analyzed mortality types. In particular, there existed a consistent pattern of increasing  $O_{3-8hmax}$ -mortality risks as we progressively adopted higher cutofs for high-temperature category.

It is worth noting that the average daily  $O_{3-8hmax}$  concentration was 119.9  $\mu$ g/m<sup>3</sup> and has a high ozone exceeding standard rate in Chengdu during the study period. From the perspective of air-pollution meteorology, there exist two key factors leading to air pollution: one is the excessive emission of air pollutants and secondary transformation, and the other is the dilution and difusion of air pollutants by the unfavorable meteorological conditions (Cai et al., [2017](#page-7-9)). As we all known, ozone, as a secondary pollutant, is widespread in the atmospheric troposphere and mainly produced by photochemical reactions of precursors (nitrogen oxides and volatile organic compounds (VOCs)); the concentration of ground-level ozone are infuenced by anthropogenic and natural emissions and by chemical, physical, and biological processes. The anthropogenic VOCs mainly come from incomplete combustion in motor vehicle exhaust, the volatilization of oil and gas coatings, and industrial emissions (Dang et al. [2021\)](#page-7-10). As of June 2018, motor vehicle ownership had exceeded 3.89 million in Chengdu, and these vehicles produce plenty of nitrogen oxides and VOCs, which are conducive to the formation of ozone. On the other hand, Chengdu is located in the Sichuan Basin and is thus afected by the topography of the Qinghai-Tibetan Plateau; the average wind speed in the Sichuan Basin is low year-round, and the frequency of static and stable weather is high. These conditions are unfavorable to the difusion or dilution of groundlevel ozone (Zhang et al. [2019b\)](#page-8-8). These high precursor concentrations and poor air difusion conditions ultimately

synergistically lead to high ozone pollution concentrations in Chengdu. Therefore, Chengdu should strengthen its air quality control, reduce its emission of ozone precursors, and formulate corresponding motor vehicle control and dispatching policies according to the changing meteorological conditions.

The temperature stratifcation results showed that the health risks of  $O_{3-8hmax}$  were more prominent at high-temperature levels than at low-temperature levels. For instance, 1 day where temperatures exceeded 24 °C, a 10-μg/m<sup>3</sup> increment in  $O_{3-8hmax}$  increased mortality risks of non-accidental, respiratory, and cardiovascular by 0.52%, 0.65%, and 1.15%, respectively. The corresponding risks were 0.17%, 0.18%, and 0.22%, respectively, under low-temperature conditions  $(< 24 °C)$ . These results keep consistent with some previous findings that  $O_{3-8hmax}$ -mortality risks were more prominent in warm season (or summer half year) than in cold season (or winter half year) (Gryparis et al. [2004](#page-7-11); Sun et al. [2018](#page-8-20)). Nevertheless, several previous studies (Cheng and Kan [2012\)](#page-7-12) suggested that the most pronounced health risks of ozone on mortality were always observed in the cold season in southern coastal cities of China, which were distinctly opposite from our results. There are several reasons for the inconsistency between previous studies and our fndings. First, the exposure pattern is an important factor affecting the results (Bell and Michelle [2004](#page-7-13)). In southern coastal cities of China, it is hot during summer and people prefer to stay indoors and mild in winter and people prefer to stay outdoors (Cheng and Kan [2012\)](#page-7-12). Chengdu city is located in southwestern China and has a subtropical climate. In Chengdu, the warm season is relatively mild, and few extreme weather events occur. For instance, the average warm-season temperature is 20.93 °C. People therefore have passion for staying outdoors and open windows in these mild temperatures, which might increase the exposure of the population to ambient ozone (Wong et al. [2001](#page-8-21)). In contrast, people prefer to stay at home rather than go out in cold season, especially in winter, due to the bitter cold outdoor temperatures and poor air quality, ultimately reducing human exposure to ambient ozone in this season.

Second, diference in the adjustment of the daily mean temperature might be possible reasons underlying these diferences (Chen et al. [2018b](#page-7-6), [a](#page-7-7)). In the temperature-mortality association, the efect of a high temperature was immediate, while the efects of cold remained signifcant over long time lags (Zhang et al. [2020a](#page-8-22)). Previous studies have chosen different temperature lags of one day to a maximum lag of 21 days (Chen et al. [2018b,](#page-7-6) [a;](#page-7-7) Jhun et al. [2014](#page-7-14)). Diferent from previous studies, considering the characteristics of ozone (short-lived gas), we pay more attention to the shortterm temperature modifcation efects on ozone-mortality in this study. This discrepancy could cause inconsistent modification patterns.

Although the independent health risks of adverse temperatures or  $O_{3-8hmax}$  on human health have been studied extensively and expounded in numerous studies, the interactions between temperature and  $O_{3-8hmax}$  have been explored only in fragments, and the results remain controversial (Ren et al. [2007](#page-8-9); Shi et al. [2020](#page-8-23); Rainham and Smoyer-Tomic [2003](#page-8-24)). Only some studies have found interactive efects, while others have not. These discrepancies mainly result from environmental and climatic conditions, acclimatization, education attainment, infrastructures, etc. (Zhang et al. [2020b](#page-8-2)). Furthermore, the analytical methods used in various studies would lead to the inconsistency of results. Compared with previous studies, we divided temperatures into two levels (low and high temperatures) by using diferent temperature thresholds corresponding to comfort and discomfort. Our fndings further support the notion that high-concentration ozone and high temperatures mutually interact to afect public health. The evidence from our study indicated the higher the temperature cutoff points were, the greater the health risks of  $O_{3-8hmax}$  were on the same kind of mortality at a high-temperature level. Therefore, heat exposure may exacerbate physiological responses to short-term ozone exposure. For instance, each 10- $\mu$ g/m<sup>3</sup> increase in O<sub>3-8hmax</sub> concentration increased mortality risks by 0.74%, 0.81%, and 1.30% in non-accidental, respiratory, and cardiovascular mortality under high-temperature  $(>26 \degree C)$  conditions; the corresponding risks were 2.22%, 2.67%, and 4.13% when chosen  $28^{\circ}$ C as the temperature cutoff, as a warmer climate will likely increase individual susceptibility to ambient ozone exposure. As a result, it will become even more important to mitigate ozone exposure in the future (Tao et al. [2021](#page-8-25); Vicedo-Cabrera et al. [2020](#page-8-0)).

The mechanisms by which the ambient temperature causes modulation efects on the relationships of ozone on human health remain unclear. There are several possible underlying mechanisms that explain this phenomenon. High temperatures are a well-known cause of heat-related mortality and can thus afect the physiological and psychological stress of the human body and aggravate many pre-existing diseases (Rainham and Smoyer-Tomic [2003](#page-8-24)). Furthermore,

high temperatures are a necessary meteorological condition for ozone generation. Extreme high temperatures may further aggravate the generation rate of ozone and subsequently increase the health risks posed by ozone to the population. Ozone is a potent oxidant capable of generating reactive oxygen species/free radicals in lung cells, thus leading to the promotion of oxidative stress, inducing acute airway infammation and damaging biomolecules (Lodovici et al., [2011](#page-7-15); Ahmad et al. [2005](#page-7-16)). The infammation of pulmonary tissues could further induce a spectrum of mediators and alter cardiac functions or the irritant receptor-mediated stimulation of parasympathetic pathways (Watkinson et al. [2001\)](#page-8-26), making people more vulnerable to the efects of ozone variability. Therefore, both high-temperature and high-concentration ozone may interact to synergistically afect people health.

The study has several advantages should be acknowledged. First, as far as I am concerned, this is the frst time to study the modification effects of temperature on  $O_{3-8hmax}$ mortality in Chengdu, China. Second, diferent from the traditional studies, we divided the temperature data into two strata (including high and low temperatures) by using different temperature cutoff points from the perspective of human comfort, which is more in line with the actual situation. Third, our study further validated that high temperature significantly amplified  $O_{3-8hmax}$ -mortality risks on the three analyzed mortality types. This interaction pattern remained consistent when diferent cutofs for temperature were applied. All these fndings suggest that diferent environmental governance and disease control strategies should be considered in Chengdu. Within a context of the changing climate, our results suggested that health damage due to ozone pollution may be infuenced by the impact of increasing temperatures, and it might become much more important to mitigate ozone exposure in Chengdu of China to further reduce the public health burden.

Some limitations of this study should be acknowledged. First, we utilized mortality data from only a 3-year period, and the statistical power was thus reduced. Second, we had no access to sub-categorical mortality characteristics, such as age, sex, educational background, work status, or the air conditioning utilization rate. Iny et al. ([2014](#page-7-5)) proved that air conditioning can mitigate the mortality risks caused by ozone exposure in 97 US cities, especially during the warm season. Unfortunately, we did not collect the relevant data mentioned above, and this limited our ability to link potentially sensitive subpopulations. Third, similar to most previous time-series studies (Bae et al. [2020](#page-7-3); Shin et al. [2020\)](#page-8-27), we only collected available outdoor monitoring data to represent personal exposure to ambient ozone, but not collected ozone concentration information in the indoor environments where people spend more time, and this omission could have biased the assessment accuracy obtained for ozone risks, resulting in a large exposure measurement error (Maji and Namdeo [2021\)](#page-8-28). Fourth, previous studies (Shin et al. [2020](#page-8-27); Zhang et al. [2020a](#page-8-22)) have demonstrated that both  $PM_{2.5}$  and ozone have adverse efects on mortality. It should be noted that we tried to include multiple interaction efects (including temperature and  $PM_{2.5}$ , temperature, and  $O_{3-8hmax}$ , as well as  $PM_{2.5}$  and  $O_{3-8hmax}$ ) on mortality in the model at initially. However, taking such complications into account did not signifcantly improve goodness of ft and lead to huge computation burden. In addition, previous study (Murase et al., [2009\)](#page-8-29) has pointed out that if multiple interactions are considered simultaneously in the model, collinearity efects are inevitable and further lead to overftting of the model, resulting in a large error. Therefore, we just pay more attention to the temperature modifcation efects on the relationship between  $O_{3-8hmax}$  and mortality. We hope to do further research and overcome these problems in the future.

## **Conclusions**

In conclusion, high temperatures strongly amplifed the adverse health risks of  $O_{3-8hmax}$  on non-accidental mortality as well as cause-specifc mortality (including respiratory and cardiovascular mortality) in Chengdu, China. The results validated that reducing  $O_{3-8hmax}$  emissions, especially in hot weather, would beneft public health. These fndings improve our cognition for the short-term health risks of ozone and offer substantial reference information for policymaking regarding ground-level ozone control and adaptation strategies with the aim of protecting public health.

**Supplementary Information** The online version contains supplementary material available at<https://doi.org/10.1007/s11356-022-20843-5>.

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### **Declarations**

**Ethics approval** Not applicable.

**Consent to participate** Not applicable.

**Consent for publication** Not applicable.

**Conflict of interest** The authors declare no competing interests.

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