Seasonal characteristics of atmospheric peroxyacetyl nitrate (PAN) in a coastal city of Southeast China: Explanatory factors and photochemical effects

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Abstract:

Peroxyacetyl nitrate (PAN) acting as a typical indicator of photochemical pollution can redistribute NOx and modulate O_3 production. Coupled with the observation-based model (OBM) and a generalized additive model (GAM), the intensive observation campaigns were conducted to reveal the pollution characteristics of PAN and its impact on O_3, the contributions of influencing factors to PAN formation were also quantified in this paper. The F-values of GAM results reflecting the importance of the influencing factors showed that ultraviolet radiation (UV, F-value=60.64), Ox (Ox=NO_2+O_3, 57.65), and air temperature (T, 17.55) were the main contributors in the PAN pollution in spring, while the significant effects of Ox (58.45), total VOCs (TVOCs, 21.63) and T (20.46) were found in autumn. The PAN formation rate in autumn was 1.58 times higher than that in spring, relating to the intense photochemical reaction and meteorological conditions. Model simulations revealed that acetaldehyde oxidation (46±4%) contributed to the dominant formation pathway of PA (hence PAN), followed by methylglyoxal oxidation (28±3%) and radical cycling (19±3%). The PAN formation was highly VOC-sensitive, as surplus NOx (compared with VOCs abundance) prevented NOx from being the limiting factor photochemical formation of secondary pollution. At our site, PAN promoted and inhibited O_3 formation under high and low ROx levels, respectively. The PAN promoting O_3 formation mainly occurred during the periods of 11:00-16:00 (local time) when the favorable meteorological conditions (high UV and T) stimulated the photochemical reactions to offer ROx radicals, which accounted for 17% of the whole monitoring periods in spring and 31% in autumn. The analysis of PAN formation mechanism and its positive or negative

effect on ozone provided scientific insights into photochemical pollution mechanism under various pollution scenarios in coastal areas.

Keywords: PAN formation mechanism; GAM model; OBM-MCM; Sensitivity analysis; Photochemical pollution; Coastal area

1 Introduction

Peroxyacetyl nitrate ($CH_3C(O)O_2NO_2$, PAN) is a key product of photochemical smog (Penkett and Brice, 1986; Li et al., 2019). PAN is generated through photochemical reactions of precursors emitted by human activities only, and the atmospheric PAN is a reliable and scientific indicator of photochemical pollution (Lonneman et al., 1976; Han et al., 2017). In the surface atmosphere, the level of PAN is much lower than that of ozone (O_3), but its biological toxicity is about one or two magnitudes greater than that of O_3 (Temple and Taylor, 1983). Additionally, PAN acts as a temporary reservoir for NOx and radicals, and can transport to remote regions to redistribute NOx and intervene in O_3 production at regional or even global scale (Kleindienst, 1994; Atkinson et al., 2006; Fischer et al., 2010).

The reaction of peroxyacetyl radical ($CH_3C(O)O_2$, PA) with NO_2 is the only formation pathway of PAN (Han et al., 2017; Xue et al., 2014). PAN affects radical chemistry and modulates O_3 production mainly by affecting PA radical, which is one of the most abundant organic peroxy radicals in the troposphere (Tyndall et al., 2001). Only a small group of oxygenated volatile organic compounds (OVOCs) (i.e. acetaldehyde (CH_3CHO), methacrolein (MACR), methyl vinyl ketone (MVK), methyl ethyl ketone (MEK), and methylglyoxal (MGLY)) can directly produce PA radical to generate PAN (Xue et al., 2014; Zhang et al., 2015). A large proportion of these OVOCs (the second-generation precursors of PAN) are mainly transformed by oxidation reactions from some hydrocarbons such as ethane, propene, isoprene, and aromatics (the first-generation precursors of PAN) (Xu et al., 2021; Qian et al., 2019). The main and direct PAN destruction is thermal decomposition, and the indirect sinks of PAN were the reactions of PA with NO, HO_2, and RO_2 (Wolfe et al., 2014; Zeng et al., 2019).

Some studies on the distribution and sources of PAN have been conducted in urban, suburban, and remote regions around the world (Grosjean et al., 2002; Marley et al., 2007; Roberts et al., 2001). The PAN levels in cities are higher than that in rural and remote areas, and that in background areas such as oceans and mountains can be as low as tens of pptv (Gaffney et al., 1999; Moore et al., 2009). Despite the growing concerns about photochemical pollution in China, PAN measurements and analysis of its

photochemical mechanism are still sparse (Zeng et al., 2019). At present, the observations of PAN were mainly distributed in Beijing, Guangzhou, and Hong Kong (Xue et al., 2014; Yuan et al., 2018; Zeng et al., 2019). Xue et al. (2014) reported that anthropogenic VOCs were the most important precursors of PAN in urban areas, and isoprene was the predominant precursor in suburban regions. In Zeng et al. (2019) study, carbonyls were the most significant contributors to PAN production, followed by aromatics and BVOCs. In addition, some researchers found that atmospheric PAN suppressed local O_3 formation in autumn (Zeng et al., 2019). Recently, negative and positive impacts of PAN photochemistry on O_3 production were captured under the low and high NOx conditions, respectively (Zeng et al., 2019; Liu et al., 2021). However, the PAN formation and its influencing mechanism on O_3 production are still complex and unclear (Hu et al., 2020; Zhang et al 2019; Xu et al., 2018). Long-term field measurements and model simulations could help to verify the mechanisms under various pollution scenarios and environmental conditions.

Xiamen is located in the coastal region of Southeast China under the East Asian monsoon control, belonging to the subtropical marine climate (Liu et al., 2020a; Liu et al., 2020b). In spring, north cold airflow and south warm airflow formed the quasistationary front causing atmospheric stagnation. In autumn, under the control of the west pacific subtropical high (WPSH), favorable meteorological conditions enhanced the formation and accumulation of photochemical pollutants (Wu et al., 2020). Our previous studies focused on the occurrence and pollution characteristics of PAN (Hu et al., 2020). In this study, an observation-based model coupled to the Master Chemical Mechanism (OBM-MCM) was used to better understand PAN photochemistry in spring and autumn, and a generalized additive model (GAM) was adopted to quantify the complex nonlinear relationships of PAN with its precursors and environmental factors (Hua et al., 2021). The study aims to explore (1) the PAN formation mechanism and sensitivity analysis, (2) the impacts of PAN on O_3 formation and radical chemistry, (3) the relationship between PAN and influencing factors under different pollution scenarios.

2 Materials and methods

2.1 Observation site

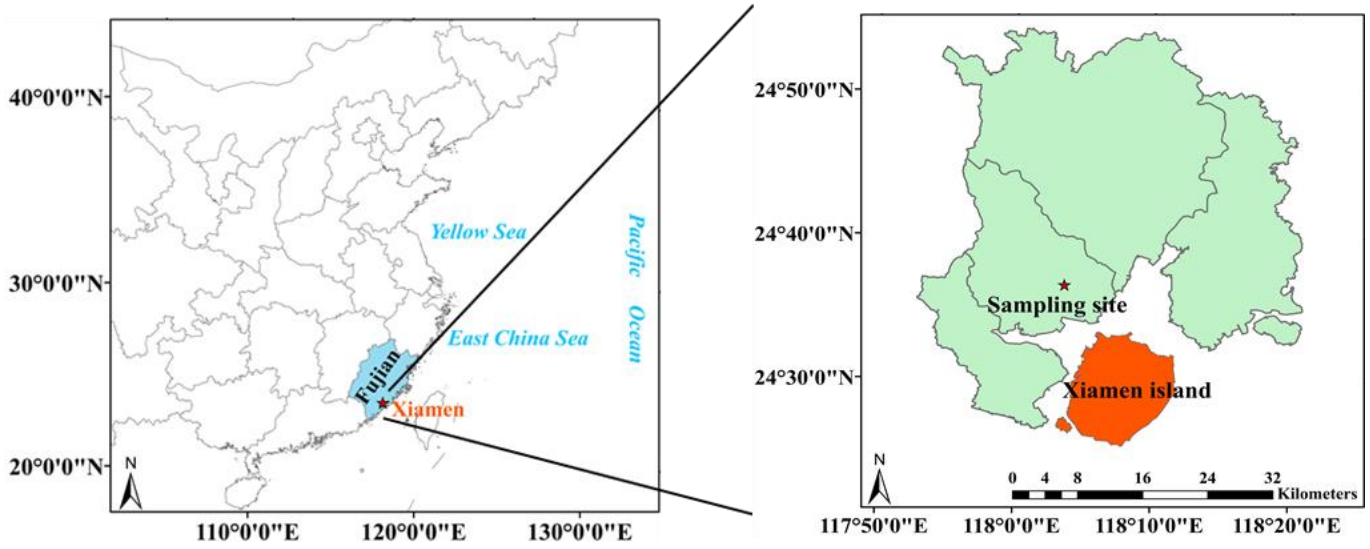


Fig. 1. Location of Xiamen and the observation site.

Observations were carried out at the Atmospheric Environment Observation Supersite (AEOS, 24.61° N, 118.06° E; Fig. 1), located on the rooftop of around a 70 m high building in the Institute of Urban Environment, Chinese Academy of Sciences. The observations site is surrounded by highways, educational institutions, and residential buildings, which was characterized by rapidly urbanizing development area. When the prevailing wind direction was southerly winds, our observation site is downwind of the densely populated downtown (Xiamen island) (Hu et al., 2020; Liu et al., 2022). The field observations were continuously conducted from March 15 to November 4, 2020. The photochemical pollution events mainly appeared during spring and autumn in Xiamen, and we preferred to choose the periods with relatively high O_3 and PAN levels, then the measured data of 53 days in each season was chosen after excluding some special circumstances, such as extreme synoptic situations and instrument calibration.

2.2 Measurement techniques

PAN was monitored using a PAN analyzer (PANs-1000, Focused Photonics Inc., Hangzhou, CN) containing gas chromatography with electron capture detector (GC-ECD). During the observation period, multi-point standard curve calibration was conducted once a month, and single-point calibration was conducted every week, respectively. In the calibration mode of the PAN analyzer, the Mass Flow Controller (MFC) controls the flow rate of NO, acetone and zero gas separately. The PAN standard gas is generated by the reaction of NO and acetone under ultraviolet light irradiation, and the sample is diluted to the required calibration mixing ratio for injection analysis. PAN was detected every 5 min and the detection limit was 50 pptv. The uncertainty and precision of PAN measurement were ±10% and 3%,

respectively.

A gas chromatography-mass spectrometer (GC-FID/MS, TH-300B, Wuhan, CN) was used for monitoring the atmospheric VOCs with a 1-hour time resolution. The instrument conducted sampling with a 30 L/min sampling rate, then samples were pre-concentrated by cooling to -160 °C in a cryogenic trap followed by heating to 100 °C, and subsequently transferred to the secondary trap by high-purity helium (He). The flame ionization detector (FID) detected the low-carbon (C2-C5) hydrocarbons by a PLOT (Al_2O_3/KCl) column (15 m × 0.32 mm × 6.0 μm); the other species were quantified using a DB-624 column (60 m × 0.25 mm × 1.4 μm). The instrument system can quantitatively analyze 106 VOCs in the ambient atmosphere, including 29 alkanes, 11 alkenes, one alkyne, 17 aromatics, 35 halogenated hydrocarbons, and 13 OVOCs. Nine compounds (Acetaldehyde, Propanal, Crotonaldehyde, Methacrolein, n-butanal, Benzaldehyde, Valeraldehyde, m-Tolualdehyde, Hexanal) could not be determined due to lack of aldehyde and ketone calibration gases, and Table S2 showed all VOCs compounds that we used in the OBM model. The single-point calibration was performed every day at 23:00 with the standard mixtures of PAMS and TO15, and multi-point calibration was performed one month. The detection limits of the measured VOCs were in the range of 0.02 ppbv to 0.30 ppbv, and the measurement precision was $\leqslant$10%.

Criteria air pollutants of O_3, CO, SO_2, and NOx, were monitored by using Thermo Instruments TEI 49i, 48i, 43i, and 42i (Thermo Fisher Scientific, Waltham, MA, USA), respectively. HONO was monitored using an analyzer for Monitoring Aerosols and Gases in Ambient Air (MARGA, ADI 2080, Applikon Analytical B.V., the Netherlands). Particulate matters ($PM_{2.5}$) were monitored by oscillating microbalance with tapered element (TEOM1405, Thermo Scientific Corp., MA, US), and the uncertainty of the $PM_{2.5}$ measurement was ±20%, respectively. The meteorological parameters (i.e. wind speed (WS), wind direction (WD), pressure (P), air temperature (T), and relative humidity (RH)) were measured by a weather station with sonic anemometer (150WX, Airmar, USA). Ultraviolet radiation (UV) was determined by a UV radiometer (KIPP & ZONEN, SUV5 Smart UV Radiometer). Photolysis frequencies including $J(O^1D)$, $J(NO_2)$, J(HONO), $J(NO_3)$, J(HCHO), and $J(H_2O_2)$ were analyzed by a photolysis spectrometer (PFS-100, Focused Photonics Inc., Hangzhou, China), and the uncertainty and detection limit of photolysis rates measurement were ±5% and around 1×10^{-5}, respectively.

Table S1 shows the detailed uncertainty and detection limit of instruments for trace gas observation. A schedule was applied to operate and inspect the AEOS monitoring station regularly and strictly to ensure the validity of the data. The detailed applications of the atmospheric monitoring procedure were shown in our previous studies (Wu et al., 2020; Liu et al., 2020a; Liu et al., 2020b; Hu et al., 2020).

2.3 Observation-based model

The OBM-MCM model is successfully used in the simulation of photochemical processes and the quantification of the reaction rates, such as O_3, PAN, and alkyl nitrates ($RONO_2$) (Zeng et al. 2019). In our study, the PAN photochemistry mechanism was simulated using this box model, and the incorporated chemical mechanism was the latest version of MCM-v3.3.1 (http://mcm.leeds.ac.uk/MCM/), which introduced 142 nonmethane VOCs and about 20000 elementary reactions (Jenkin et al., 2003; Saunders et al., 2003). The physical process including dilution effect and dry deposition within the boundary layer height was considered, avoiding the excessive accumulation of pollutants in the model (Li et al., 2018; Liu et al., 2021; Xue et al., 2016). The observed data with a time resolution of 1 h of pollutants (i.e., O_3, CO, NO, NO_2, HONO, SO_2, and VOCs), meteorological parameters (i.e., T, P, and RH), and photolysis rate constants ($J(O^1D)$, $J(NO_2)$, $J(H_2O_2)$, J(HONO), J(HCHO), and $J(NO_3)$), which were mentioned in Section 2.1, were input into the OBM-MCM model as constraints. The photolysis rates of other molecules were driven by solar zenith angle and were scaled by measured JNO_2 (Saunders et al., 2003). Pre-ran for 2 days before running the model to constrain the unmeasured compounds reaching a steady-state (Xue et al., 2014; Liu et al., 2022).

PAN affects atmospheric photochemistry by acting as a temporary source or sinks of PA radical (Xue et al., 2014; Liu et al., 2021), hence the production and sink of PA radical reflecting the PAN formation were discussed in our study. Furthermore, relative incremental reactivity (RIR) was used to analyze the sensitivity of O_3 (Eq. 1) and PAN (Eq. 2) to their precursors, and was calculated as the ratio of the differences in O_3 or PAN net production rate to variety in precursors (Chen et al., 2020; Liu et al., 2021). The production pathways of O_3 include HO_2+NO and RO_2+NO reactions, and the destruction pathways of O_3 involve reactions of O_3 photolysis, O_3+OH, O_3+HO_2, O_3+VOCs, NO_2+OH, and NO_3+VOCs. The net O_3 production rate ($P(O_3)$) is calculated by the difference of O_3 production rate and destruction rate, and the detailed net production rate of O_3 ($P(O_3)$) was introduced in our previous study (Liu et al., 2022). The net production of PAN (P(PAN)) involved the production pathway of PA+NO_2, and the loss of PAN was thermal decomposition and PAN+OH (Zeng et al., 2019).

$$RIR(PAN) = \frac{\Delta P(O_3)/P(O_3)}{\Delta X/X} \quad (1)$$

$$RIR(O_3) = \frac{\Delta P(PAN)/P(PAN)}{\Delta X/X} \quad (2)$$

Here, the $\Delta X/X$ meaning the reduction in the input mixing ratios of each target O_3 and PAN precursor

group was 20% (Liu et al., 2021).

2.4 Generalized additive model

The Generalized Additive Model (GAM) is an extension of the additive model proposed. Different from traditional regression models, GAM is a non-parametric regression model driven by data rather than statistical distribution models (He et al., 2017). GAM does not need to set the parameter model in advance, and it can adjust the functional form of the explained variable according to the specific situation. The Generalized Additive Model (GAM) has been widely used in air pollution research such as O_3 and $PM_{2.5}$, and can effectively deal with the complex nonlinear relationship between air pollutants and influencing factors (Ma et al., 2020; Hua et al., 2021; Guan et al., 2019). It is the first time that the GAM is used to analyze the relationship between PAN and its influencing factors, and the combined effect of multiple influencing factors on the PAN mixing ratio was discussed in our study. Its form is:

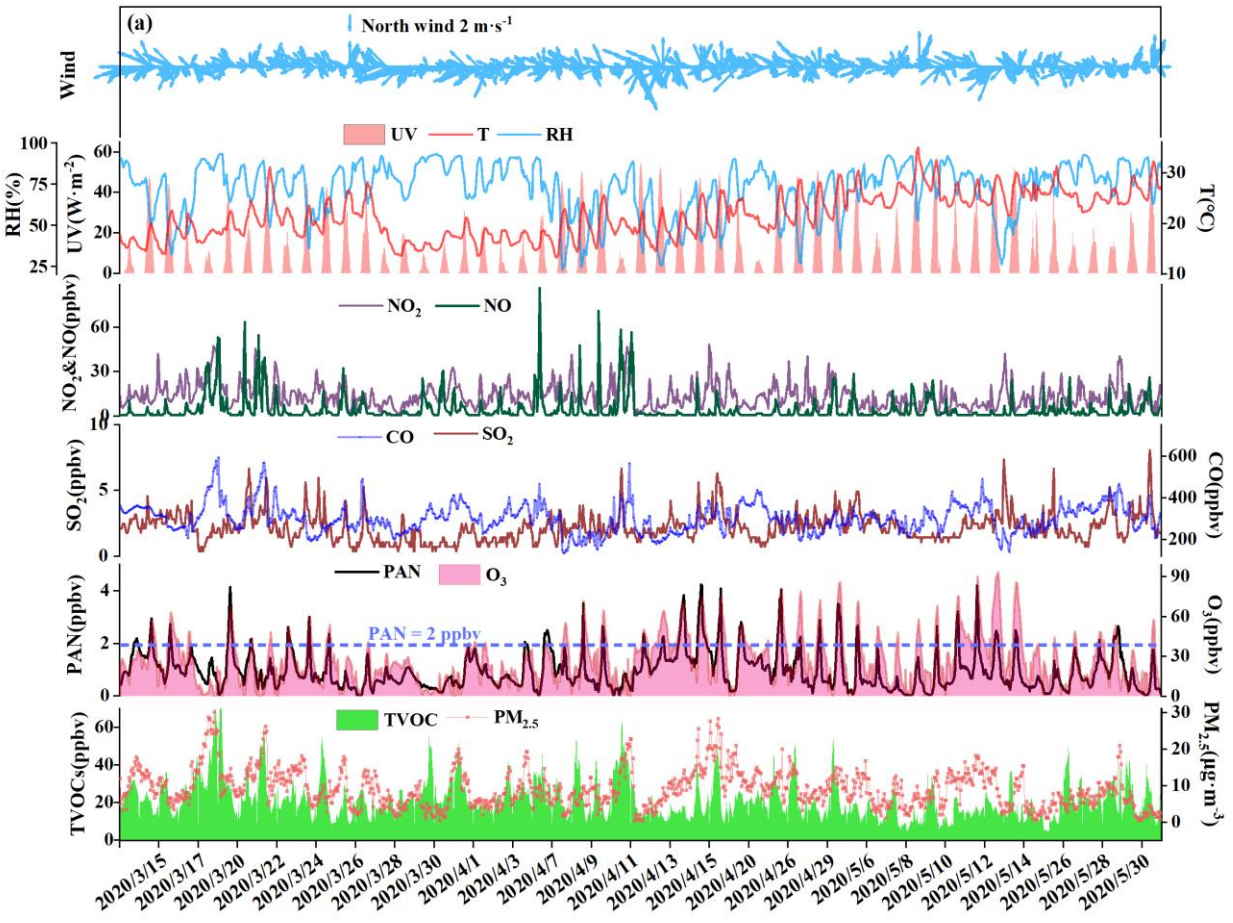
$$g(y)=\beta+f_1(x_1)+f_2(x_2)+\ldots\ldots+f_n(x_n)+\alpha \qquad (3)$$

Where y is the response variable; $g(y)$ is the connection function; x_n, x_i, x_j, x_k, and x_l are the explanatory variables; fn is the non-parametric smoothing functions; β is the intercept; α is the truncation error.

The F-value, P-value, adjust R^2, and deviance explained given by the GAMs model are used to judge the significance of the influencing factors on PAN and the goodness of the model simulation. Among them, a high F-value indicates the great importance of the influencing factor; the P-value is used to judge the significance of the model result; the adjusted R^2 is the value of the regression square ranging from 0 to 1; the deviance explained represents the fitting effect. In addition, when the degree of freedom (edf, ref.df) of the explanatory variable is 1, it indicates that the explanatory variable and the response variable are linear. When the degree>1, it is a non-linear relationship.

3 Results and discussion

3.1. Overview of observation



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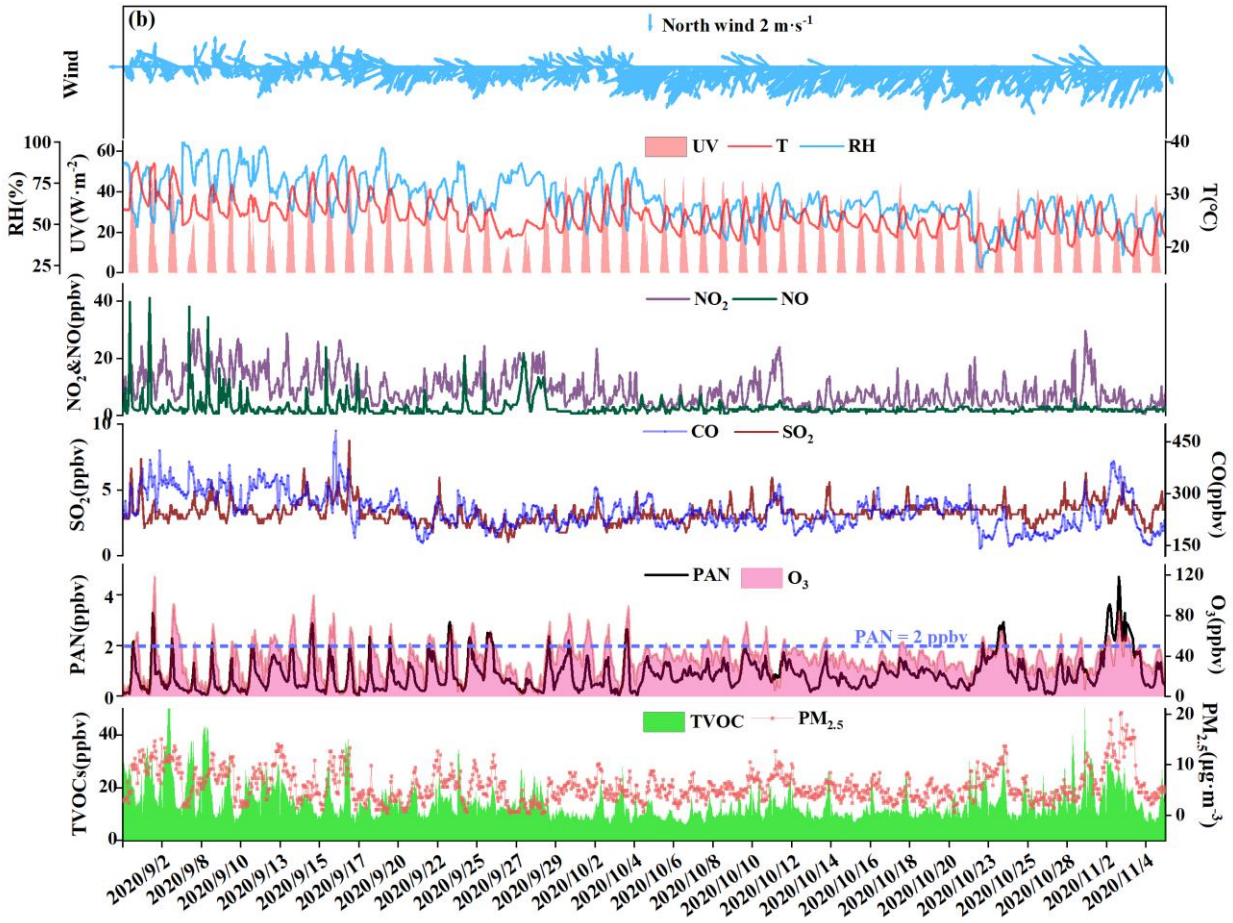

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210 **Fig. 2. Time series of PAN, O_3, NOx, CO, SO_2, TVOCs, $PM_{2.5}$, and meteorological parameters in (a) spring and**

(b) autumn.

The time series of air pollutants and meteorological parameters are shown in Fig. 2. The average levels of PAN in autumn (0.87±0.66 ppbv) were comparable to that in spring (0.96±0.73 ppbv), while O_3 mixing ratios in autumn (37.22±16.89 ppbv) were 1.39 times higher than that in spring (26.73±18.63 ppbv). PAN and O_3 are produced by the photochemical reactions of VOCs and NOx, thus they usually show a relatively close relationship ($R^2 \geq 0.49$, Fig. S3). The PAN level (0.92±0.69 ppbv) in Xiamen was lower than that of megacities such as Beijing (3.79±3.26 ppbv) (Xu et al., 2021), Jinan (2.54 ppbv) (Liu et al. 2018), Santiago (6.4 ppbv) (Rubio et al., 2005) and Chongqing (2.05 ppbv) (Sun et al., 2020), and was comparable to the coastal cities with relatively clean air, including Shenzhen (1.01±0.94 ppbv) (Xia et al., 2021), and Qingdao (0.81 ppbv) (Liu et al., 2021).

The averaged values of PAN and NO, NO_2, CO, TVOCs in spring were 1.70, 1.32, 1.21, and 1.46 times higher than those in autumn, respectively. The details of measured VOCs were provided in Table S2. Alkanes, OVOCs, aromatics, and halocarbons accounted for about 90% of total VOCs, suggesting the impacts of atmospheric oxidation capacity and marine emissions in coastal regions (Liu et al., 2020a; Liu et al., 2020b). During the transition from spring to summer the wind direction fluctuated between northwest and southeast while during the transition from summer to autumn the wind direction fluctuated from southeast to northeast. The wind rose charts showed that the wind direction frequencies with relatively high wind speed (>3 $m \cdot s^{-1}$) in spring and autumn were southeast wind and northeast wind (Fig. S4), respectively. Although the frequency of northwest wind (NNW) also accounted for a certain proportion, the NNW speeds were generally slow, and the direction of the NNW was mainly rural residential and mountainous areas with less anthropogenic emissions, so that it was not the focus of this research. The ultraviolet radiation (UV), WS and T in spring (15.32 $W \cdot m^{-2}$; 1.96 $m \cdot s^{-1}$; 21.51 ℃) were weaker than those in autumn (18.43 $W \cdot m^{-2}$; 3.01 $m \cdot s^{-1}$; 25.85 ℃), and RH and P in spring (73.25 %; 1010.71 hPa) were higher than that in autumn (65.21 %; 1008.71 hPa). These meteorological conditions carried by the WPSH (high T, low RH, and stagnant weather conditions) were conducive to the photochemical reaction and accumulation of air pollutants in autumn (Wu et al., 2019; Xia et al., 2021). High precursor levels of PAN in spring were conducive to the continuous and stable production of PAN, and the high air temperature in autumn accelerated the thermal decomposition of PAN. However, the O_3 levels in autumn were higher than that in spring, attributing to the influence of strong photochemical reaction conditions, regional transport from the Yangtze River Delta region or increased atmospheric

background levels (Monks, 2000). High O_3 values in both seasons were concentrated on the wind direction of southeast and northeast (Fig. S5). High PAN values in spring easily happened in the wind direction of the southeast with low wind speed (<3 $m \cdot s^{-1}$), showing the influence of urban plumes from the downtown of Xiamen island. High PAN values in autumn also appeared in the wind direction of the southeast, as well as the northeast with a relatively high wind speed (from Quanzhou city, an industrial city adjacent to Xiamen). Anymore, PAN lifetimes in our observation site were relatively short due to the high ambient temperature, and the PAN lifetimes in autumn (2.02 hours) were significantly lower than that in spring (6.39 hours), which was not conducive to regional transport (Hu et al., 2020; Liu et al., 2018). Accordingly, O_3 showed obvious characteristics of long-range transport, and PAN pollution was mainly from local production/accumulation in spring and autumn, but short-range transport from adjacent cities might contribute to the high PAN concentrations in autumn to a certain extent.

Based on the above analysis, we found that the photochemical reactions were still intense and even stronger under the low precursor mixing ratios. Although the precursor mixing ratios of PAN and O_3 in spring were significantly higher than those in autumn ($P<0.01$), the PAN mixing ratios in autumn were comparable to those in spring, while the O_3 mixing ratios in autumn were much higher than those in spring. Therefore, it is very necessary to furtherly explore the key influencing factors and their formation mechanisms.

3.2 The influencing factors of PAN using the GAM

PAN levels are not only related to chemical reactions in the boundary layer, but also affected by meteorological conditions (Hu et al., 2020). According to the collinearity analysis (He and Lin, 2017), the meteorological parameters (UV, T, RH, and WS) and other air pollutants (NO, TVOCs, $PM_{2.5}$, and O_x) were considered into the multiple-factor GAM model (Table S3). As shown in Table 1, the adjusted R^2 and deviance explained for the smoothed variables of the multiple-factor GAM model were 0.70 and 72% in spring, 0.60 and 63% in autumn. According to the F-values, the orders of the explanatory variables in spring and autumn were UV (60.64) > Ox (57.65) > T (17.55) > $PM_{2.5}$ (9.94) > TVOCs (9.52) > NO (8.73)> WS (7.42) > RH (3.4) and Ox (58.45) > TVOCs (21.63) > T (20.46) > $PM_{2.5}$ (14.53) > RH (10.99) > UV (7.13) > NO (4.16) >WS (2.55), respectively.

Response curves of the PAN to explanatory factors in the multiple-factor model were presented (Fig. 3 and Fig. S6). Except for UV and T in spring, the degrees of freedom (df) of the explanatory variables were greater than 1, indicating the non-linear relationships between explanatory variables and PAN. The

PAN in both seasons showed a downward trend with the increase of NO. PAN in spring was constant with NO fluctuation between 10 and 23 ppbv, and the confidence interval (CI) of NO concentration was relatively narrow. As we all know, the reaction of PA+NO is one of the most important loss pathways of PA, and the NO_2 production by NO oxidation in the O_3 formation cycle can react with PA radical to produce PAN, suggesting the fact that NO can consume and produce PAN indirectly (Liu et al., 2021). The consumption of NO to PAN was basically equal to the production when the NO levels were relatively high (>10 ppbv), and the consumption of NO to PAN is greater than the production when the NO levels were low in spring. High values of NO mainly happened during rush hour traffic, thus controlling vehicle emissions can effectively alleviate PAN pollution. Ox had a positive correlation with PAN, representing the promotion effects of atmospheric oxidation capacity on PAN formation. The Ox levels <70 ppbv (with narrow CI) played a significant promotion role in PAN formation (Fig. 3(b) and Fig. S4(b)). High Ox >70 ppbv showed little influence on PAN, which could be explained as high Ox with relatively high air temperature leading to intense PAN thermal decomposition. When TVOCs were between 10 and 30 ppbv and $PM_{2.5}$ levels were <17 $\mu g \cdot m^{-3}$, PAN showed an upward trend with narrow CI. According to our previous study (Liu et al., 2022; Hu et al., 2020), the results of sensitivity analysis in Xiamen was VOCs-sensitive; the relatively low $PM_{2.5}$ concentrations in Xiamen showed limited influence on solar radiation through scattering and absorption, but promoted heterogeneous reactions producing radicals to a certain extent. UV and T had significant positive and negative nonlinear correlations with PAN, respectively. When UV changed between 0 and 50 $W \cdot m^{-2}$ and T changed between 15 and 35 $W \cdot m^{-2}$, the CIs barely increased. In addition, when RH was more than 40%, the increase of RH was unfavorable for PAN production in both seasons. Some studies also found that high water vapor content could remove PAN and its precursors (Yan et al., 2018; Ma et al., 2020). Overall, the multiple-factor GAM analysis could better simulate the variations of PAN under real atmospheric conditions and evaluate the contributions of the influence factors to PAN formation.

Table 1 Estimated degree of freedom (Edf), degree of reference freedom (Ref. df), P-value, F-value, deviance explained (%), adjusted R^2, deviance contribution (%) for the smoothed variables in the multiple-factor GAM

model.

Smoothed variables	Spring				Autumn			
	Edf	Ref.df	F-value	P-value	Edf	Ref.df	F-value	P-value
NO (ppbv)	5.21	6.26	8.73	0.00	1.11	1.21	4.16	0.03
Ox (ppbv)	4.73	5.85	57.65	0.00	4.84	5.98	58.45	0.00
TVOCs (ppbv)	7.14	8.19	9.52	0.00	4.08	5.06	21.63	0.00
$PM_{2.5}$ ($\mu g \cdot m^{-3}$)	5.73	6.86	9.94	0.00	1.53	1.90	14.53	0.00
UV ($W \cdot m^{-2}$)	1.00	1.00	60.64	0.00	4.38	5.38	7.13	0.00
T (℃)	1.00	1.00	17.55	0.00	2.73	3.46	20.46	0.00
RH (%)	6.78	7.87	3.40	0.00	6.56	7.68	10.99	0.00
WS ($m \cdot s^{-1}$)	5.22	6.37	7.42	0.00	5.12	6.28	2.55	0.02
	Deviance explained (%)=80%; Adjust R^2=0.79				Deviance explained (%)=72%; Adjust R^2=0.70			

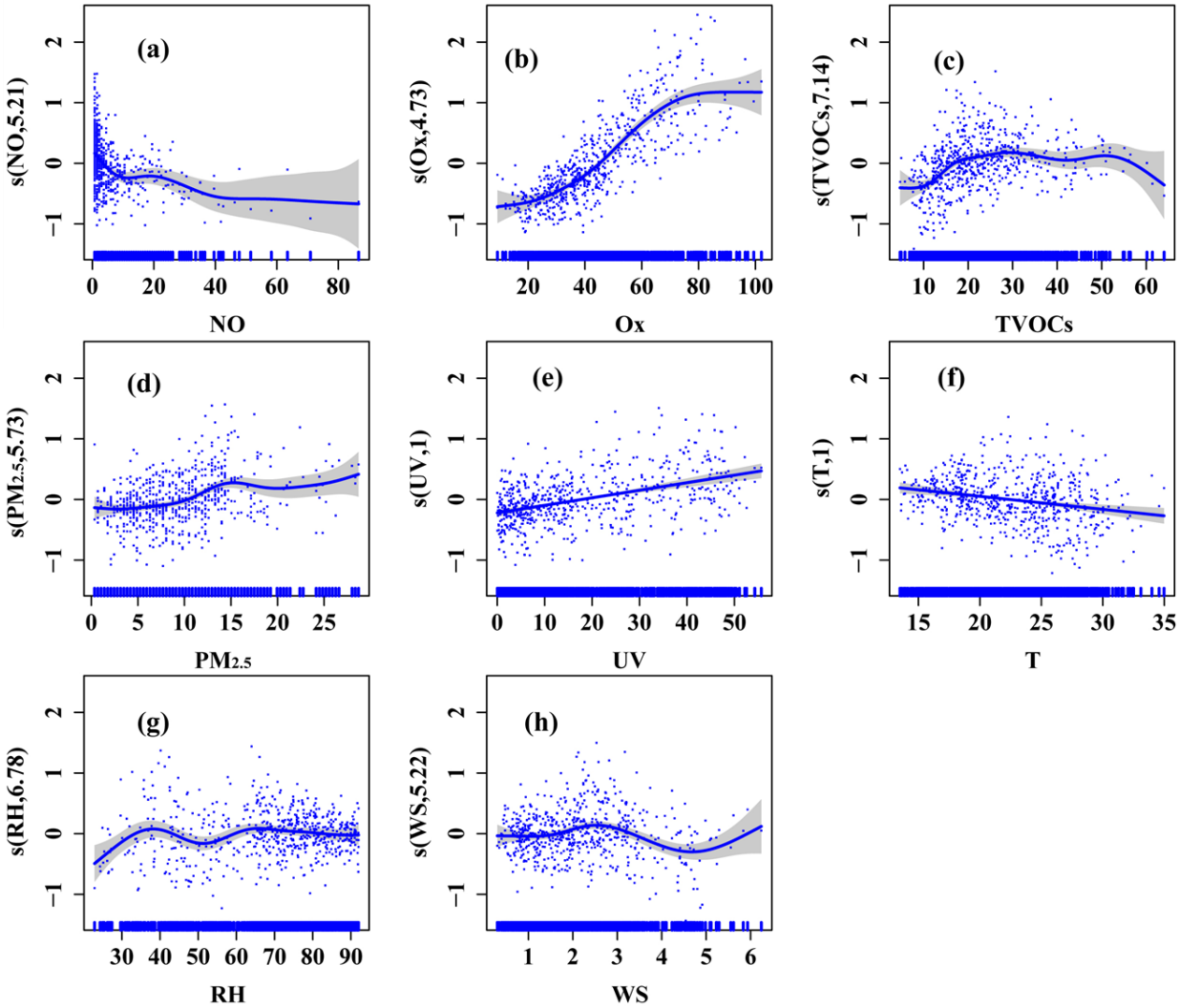

Fig. 3. Response curves (spring) in the multiple-factor model of PAN to changes in (a) NO, (b) Ox (Ox=O_3+NO_2), (c) TVOCs, (d) $PM_{2.5}$, (e) ultraviolet radiation (UV), (f) air temperature (T), (g) relative humidity (RH), and (h) wind speed (WS). The y-axis is the smoothing function values. For example, s(NO, df) shows the trend in PAN when NO changes, and the number of df is the degree of freedom. The x-axis is the influencing factor, and the shaded area around the solid red line indicates the 95% confidence interval of PAN. The blue vertical short lines represent the concentration distribution characteristics of the explanatory variables (units: NO (ppbv), Ox (O_3+NO_2) (ppbv), TVOCs (ppbv), $PM_{2.5}$ ($\mu g \cdot m^{-3}$), UV ($W \cdot m^{-2}$), T (℃), RH (%), WS ($m \cdot s^{-1}$)).

3.3. Formation mechanism of PAN

3.3.1 Diurnal variation during episodes and non-episodes

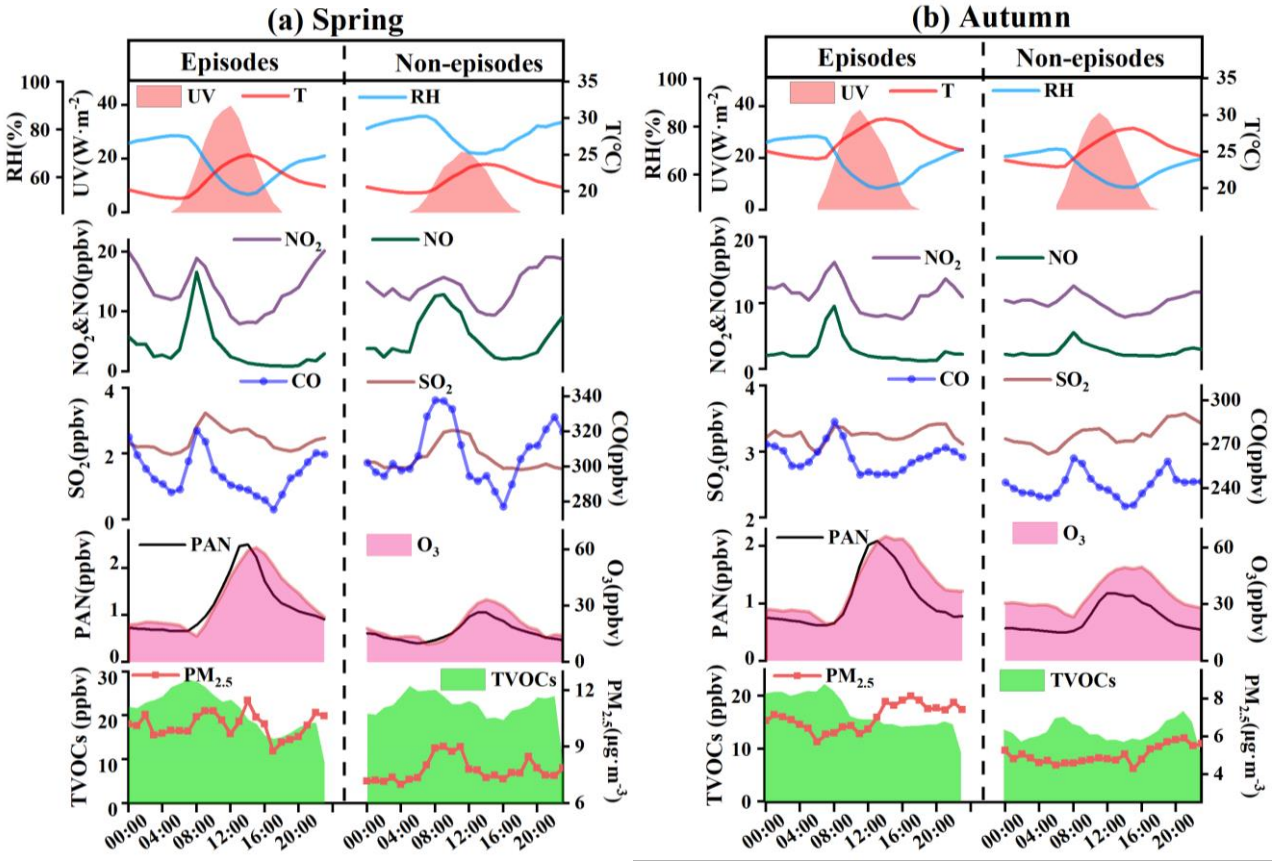

Fig. 4. Diurnal trends of PAN, O_3, TVOCs, $PM_{2.5}$, other trace gases and meteorological parameters during episodes and non-episodes in (a) spring and (b) autumn, respectively.

Throughout the 53-days campaign, 30 and 21 days (i.e., 57% and 40%) with the peak values of PAN exceeding 2 ppbv were observed in spring and autumn, respectively. The scenarios of episodes and non-episodes were classified, according to the previous method (Xue et al., 2014). Diurnal variations of air pollutants and meteorological parameters during episodes and non-episodes are shown in Fig. 4, which could be explained by the evolution of the planetary boundary layer, local emissions, and atmospheric photochemistry. PAN reached a maximum value at 12:00-14:00, then decreased with weak solar radiation and reached the lowest in the early morning. Similar diurnal patterns of PAN and O_3 were observed, indicating the dominance of local photochemistry during the observation period (Zeng et al., 2019). CO, NOx and TVOCs showed highest values in the morning and the lowest values in the afternoon.

In autumn, averaged PAN and O_3 during episodes (PAN: 1.08±0.87 ppbv, and O_3: 40.06±20.27 ppbv) were higher than those during non-episodes (PAN: 0.74± 0.41 ppbv, and O_3: 35.36±13.95 ppbv).

Meanwhile, some air pollutants and meteorological parameters during episodes were 1.03-1.40 times higher than those during non-episodes. The rainfall in Xiamen is more frequent in spring (Hu et al., 2020), leading to the obvious differences in UV and RH levels between episodes and non-episodes. In spring, the precursors (CO, NOx, TVOCs) of PAN during episodes were 1.04-1.49 times lower than those during non-episodes. Moreover, the PAN and O_3 mixing ratios during episodes (PAN: 1.20±0.81 ppbv, and O_3: 32.92±19.81 ppbv) were still significantly higher than those during non-episodes (PAN: 0.64±0.43 ppbv, and O_3: 18.65±13.16 ppbv), attributing to the favorable meteorological conditions of photochemical reactions (strong UV, high T, and low RH). These results further explained that UV, Ox, and T in spring and Ox, TVOCs, T, and $PM_{2.5}$ in autumn played important roles in the formation of PAN based on the GAM analysis.

3.3.2. Formation and loss of PA radical

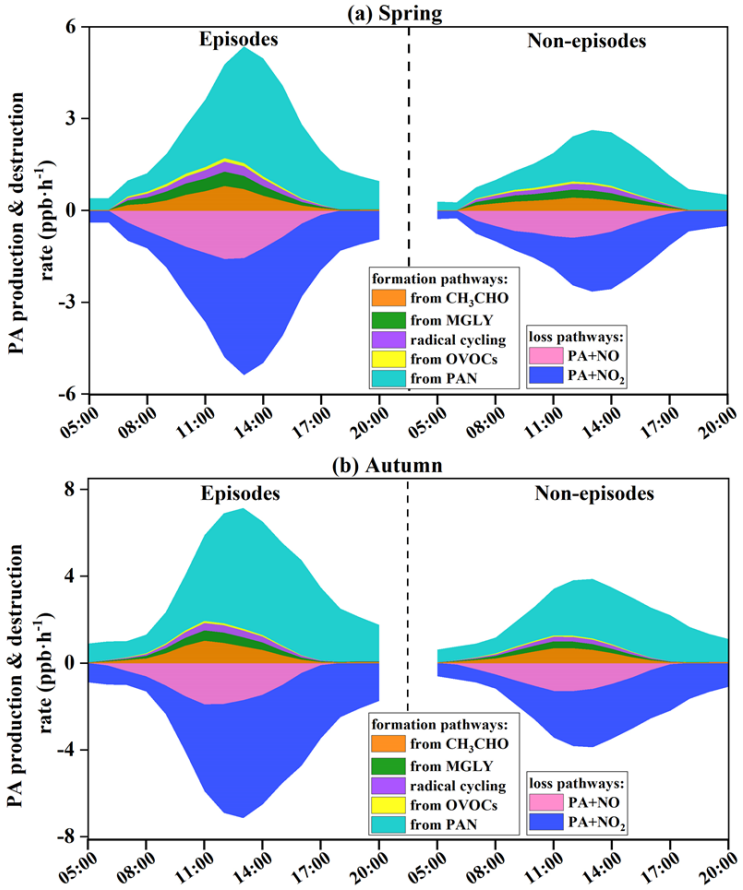

Fig. 5. Formation and destruction rates of PA radical (hence PAN) during episodes and non-episodes in (a) spring and (b) autumn, respectively.

The formation and sink pathways of PA radical were further explored under different pollution scenarios (Fig. 5). Both the PA (hence PAN) production and destruction rates during episodes were 1.80 times higher than those during non-episodes. Combined with the analysis of Section 3.3.1, PA production

rates during the daytime (06:00-17:00 LT) in autumn were 1.58 times higher than that in spring, even though the precursor levels in autumn were much low compared to those in spring. These results indicated favorable meteorological condition was the dominant factor to produce PAN through accelerating its production rate and accumulation. The thermal decomposition of PAN to PA radical in autumn accounted for 77±12% (episodes) and 73±16% (non-episodes) of total PA production, as well as 70±12% (episodes) and 64±15% (non-episodes) in spring, attributing to the relatively high air temperature and UV intensity. The thermal decomposition of PAN peaked at around 13:00~14:00 LT, when the air temperature was the highest in the day, and the pathways without considering the transform between PA and PAN peaked at noontime around 12:00 LT, when the solar radiation was the highest and photochemical reactions became the most intensive.

The average daytime PAN production rate from CH_3CHO by reacting with OH and NO_3 contributed 0.36±0.25 ppb h^{-1} and 0.24±0.13 ppb h^{-1} during episodes and non-episodes in spring. While the rate of 0.46±0.35 ppb h^{-1} and 0.34±0.24 ppb h^{-1} during episodes and non-episodes were observed in autumn. The second production reaction was photolysis and oxidation by OH and NO_3 of MGLY (episode: 0.25±0.15 ppb h^{-1} and non-episodes: 0.17±0.08 ppb h^{-1} in spring; episode: 0.24±0.17 ppb h^{-1} and non-episodes: 0.16±0.11 ppb h^{-1} in autumn). Then, the processes of radical cycling including RO radical decomposition and reactions of acyl peroxy radicals with NO were also the important sources to produce PA, with the contributions of 20±3% and 18±3% in spring and autumn. PA from the other OVOCs (not including CH_3CHO, MGLY, MVK, MACR, and acetone) through reactions of photolysis and oxidation by OH, NO_3, and O_3, accounted for 7±2% and 6±1% in spring and autumn, respectively. Other reactions of acetone, MVK, MACR, MPAN, and isoprene had a minor contribution (around 1% in total) to PA formation. In contrast, the major contributor of PAN destruction rate was PA+NO_2 (69±16% in spring and 73±14% in autumn), followed by PA+NO (31±17% and 27±13%), while the other reactions with NO_3, HO_2, and RO_2 contributed limitedly (around 0.1% of the total).

The second-generation precursors of PAN of CH_3CHO and MGLY have both primary and secondary sources, and the other OVOCs are mainly oxidation products of hydrocarbons (Sinha et al., 2019; Sarkar et al., 2017). Consequently, the contribution and importance of first-generation precursors of PAN are necessary to identify to better control photochemical pollution, which will be discussed in the next section.

3.3.3. Sensitivity of PAN precursors

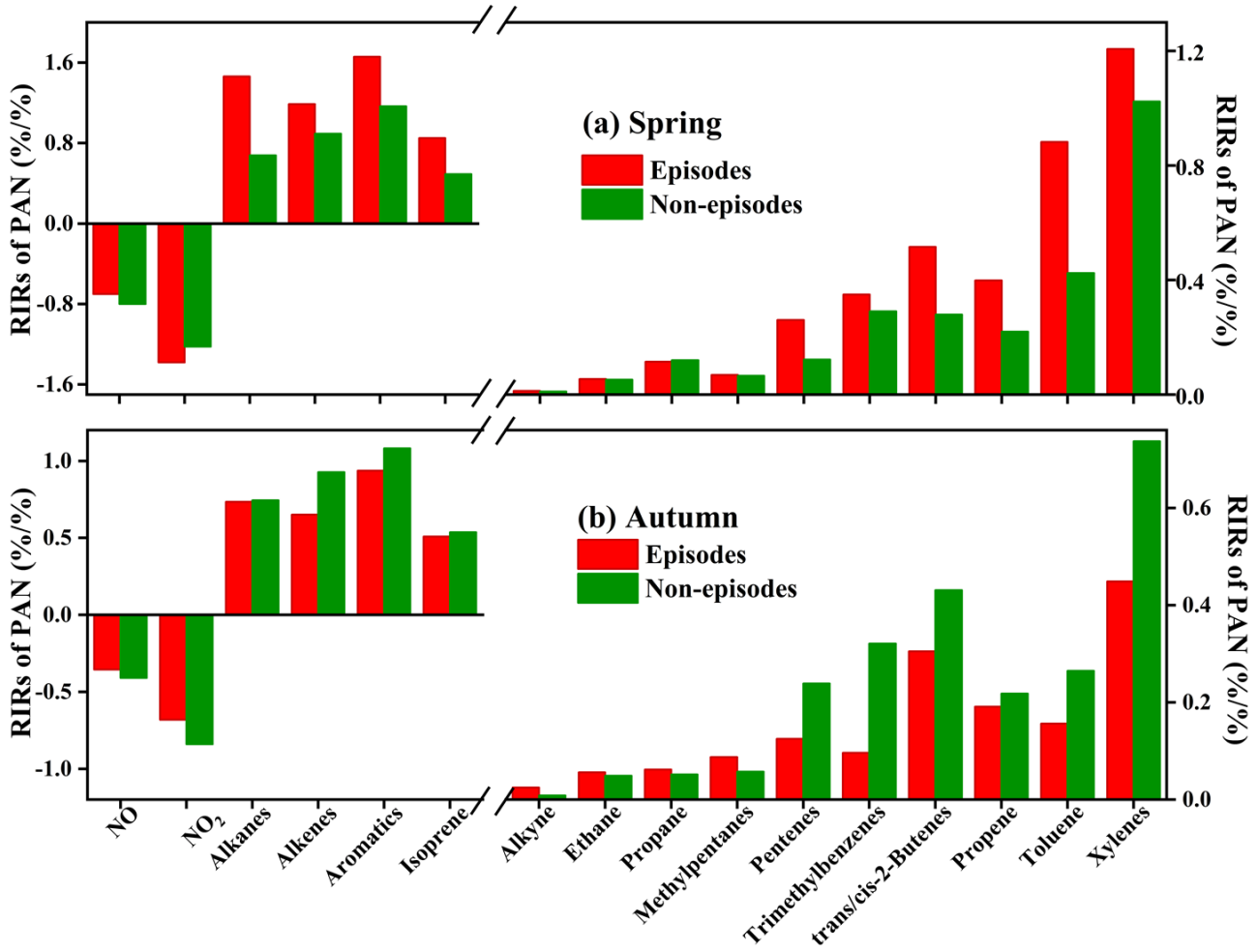

Fig. 6. The OBM-MCM calculated relative incremental reactivity (RIR) for major PAN precursor groups and top 10 specific species in (a) spring and (b) autumn during the daytime (06:00-17:00 LT).

The OBM-MCM model analysis could be used to examine the relationship between PAN and its precursors, and quantify the contribution of first-generation precursors (Liu et al., 2021; Cardelino and Chameides, 1995). During these simulations (except for NO and NO_2), the model was not constrained by the OVOC measurements considering that these first-generation precursors contribute to PAN production through formation of OVOCs. The relative incremental reactivities (RIRs) for O_3 and PAN are shown in Fig. 6 and Fig. S7. The PAN production was highly VOCs-sensitive, while the RIRs of NO and NO_2 were negative ranging from −0.17 to −1.94%/% during the daytime (06:00-17:00 LT). This consisted of the fact that high dense mobiles resulted in the large emissions of vehicle exhausts in Xiamen city. The ratio of VOCs/NOx (1.11±0.32) also convinced NOx was not the limiting factor on the photochemical reaction (Tan et al., 2019). In suburban or rural areas, the transition regime and NOx-sensitive for PAN and O_3 production were usually found (Xue et al., 2014; Liu et al., 2021). Zeng et al. (2019) found NO_2-positive and NO-negative to PAN formation in a suburban of Hong Kong, consisting with the fact that NO_2 directly produced PAN and NO consumed PA radical inhibiting PAN formation.

As shown in Fig.6, aromatics showed the largest RIRs for PAN in spring (1.41%/%) and autumn (1.03%/%), followed by alkanes (1.04%/% in spring and 0.78%/% in autumn), Alkenes (1.04%/% and 0.74%/%), and isoprene (0.67%/% and 0.52%/%). The sensitivities of PAN precursors in spring were 1.37-2.07 times higher than those in autumn, due to the large percentages of PAN decomposition at high air temperatures in autumn. In spring, the weak solar radiation led to poor photochemical reactions, so the RIRs of PAN during non-episodes were lower than that during episodes. However, the PAN sensitivities during episodes were lower than those during non-episodes, attributed to the rapid PAN decomposition in autumn (Liu et al., 2021). In addition, RIRs of VOCs and NOx for PAN were significantly higher than that of O_3 (Fig. S5). For RIRs of VOCs, except for air temperature, the different formation mechanisms of PAN and O_3 should be considered. Only a small part of the VOCs could produce PA to form PAN, thereby, the VOCs were insufficient to produce PAN (Fischer et al., 2014). For RIRs of NOx, O_3 was produced from the NO_2 conversion process, and was also rapidly consumed by NO titration. High levels of VOCs and NOx enhanced the PAN formation, even though a pathway of NO destructed PAN, which was negligible compared to thermal decomposition. For this reason, the RIRs of NOx for PAN were higher than those for O_3.

In addition, the top 10 VOCs species (including xylenes, toluene trans/cis-2-butenes, trimethylbenzenes, propene, pentenes, and methypentanes) governing PAN production were further identified (Fig. 6). The results suggested that the reduction of aromatics, alkenes, and alkanes with ≤5 carbons could effectively decrease PAN pollution.

3.4. Impacts of PAN on O_3 formation

3.4.1 Inhibition and promotion effect of PAN on O_3 formation

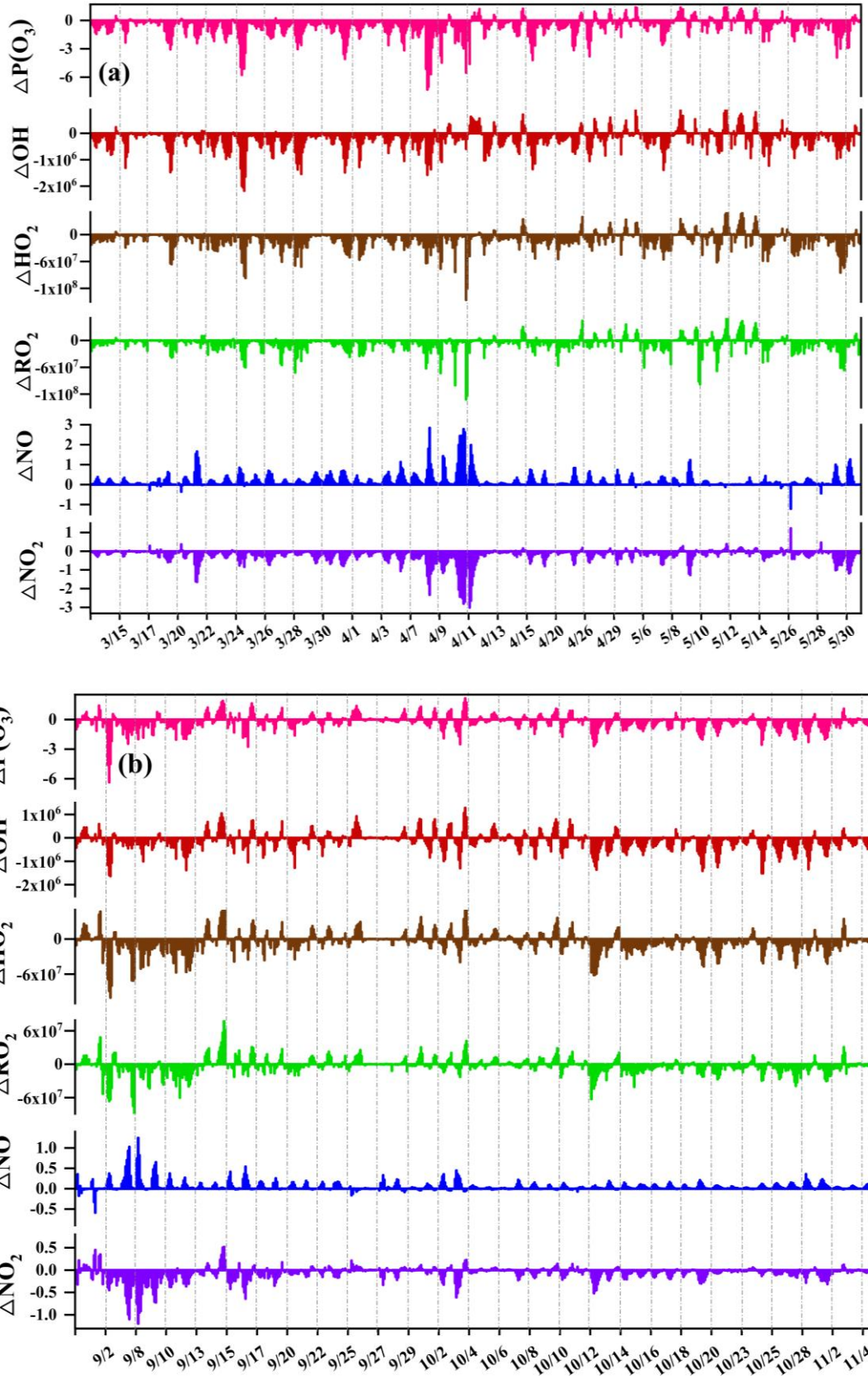

Fig. 7. The differences of O_3 net production $\triangle P(O_3)$, $\triangle OH$, $\triangle HO_2$, $\triangle RO_2$, $\triangle NO$ and $\triangle NO_2$ between the SC1 and the SC2 during the daytime (06:00-17:00) in (a) spring and (b) autumn (Unit: ppbv·h^{-1} for $\triangle P(O_3)$; ppbv for $\triangle NO$ and $\triangle NO_2$; molecules·cm^{-3} for $\triangle OH$, $\triangle HO_2$ and $\triangle RO_2$). The SC1 scenario was the base scenario putting all detected data (i.e. VOCs, trace gases, and meteorological parameters) into the model with all reaction pathways of the MCM mechanism, and the SC2 disabled the PAN chemistry, which is the only difference between SC1 and SC2.

PAN could affect O_3 production by acting as a temporary source of NOx or sink of PA radical to affect precursors and radical chemistry in the troposphere (Xia et al., 2021). To quantify the changes of O_3 in response to PAN chemistry in the coastal city, two parallel scenarios (SC1 and SC2) were conducted based on the OBM model. The SC1 was the base scenario putting all detected data (i.e. VOCs, trace gases, and meteorological parameters) into the model with all reaction pathways (as the description in Section 2.2), and the SC2 disabled the PAN chemistry, which is the only difference between SC2 and SC1. Figure 7 shows the differences of O_3 net production rates $\triangle P(O_3)$, $\triangle OH$, $\triangle HO_2$, $\triangle RO_2$, $\triangle NO$ and $\triangle NO_2$ between the SC1 and the SC2. Negative and positive values represented the inhibition and promotion effects of PAN photochemistry on O_3 formation, respectively. Overall, PAN mostly inhibited the O_3 formation during the observation days. $\triangle P(O_3)$ had significantly positive correlations with $\triangle OH$ (R^2=0.96 in spring and 0.95 in autumn), $\triangle HO_2$ (R^2=0.91 and 0.96), $\triangle RO_2$ (R^2=0.86 and 0.86) and $\triangle NO_2$ (R^2=0.72 and 0.85), and negative correlation with $\triangle NO$ (R^2=-0.63 and -0.65). As shown in Fig. S8, the promotion effects of PAN on O_3 mainly happened during the periods of 11:00-16:00 LT, and most of them concentrated on PAN pollution episodes. The percentage of negative $\triangle P(O_3)$ values were 83% and 69% in spring and autumn, defined as "inhibition effect stages". While the positive $\triangle P(O_3)$ values accounted for 17% and 31% in spring and autumn, defined as "promotion effect stages".

Figure 8 shows the variations of modeled $P(O_3)$, O_3 budgets, and ROx on the inhibition and promotion effect stages in spring and autumn. The abundance of ROx in autumn (2.85×10^8 molecules cm^{-3}) was higher than that in spring (2.08×10^8 molecules cm^{-3}) during inhibition effect stages, while the $P(O_3)$ value in autumn (5.24 ppbv h^{-1}) was higher than that in spring (4.88 ppbv h^{-1}). On the contrary, the level of ROx in spring (4.81×10^8 molecules cm^{-3}) was higher than that in autumn (4.20×10^8 molecules cm^{-3}) during promotion effect stages, and the $P(O_3)$ value (5.95 ppbv h^{-1}) in spring was higher than that in autumn (5.76 ppbv h^{-1}). The results indicated that high ROx concentration was an important factor for the formation of O_3. In the case of closing PAN photochemistry, the $P(O_3)$ increased 1.20 and 1.12 times during inhibition effect stages and decreased 1.09 and 1.08 times during promotion effect stages in spring and autumn, respectively (Fig. 8a). This was consistent with the corresponding changes of ROx radical (Fig. 8b). During the inhibition effect stages, the averaged concentrations of OH, HO_2, and RO_2 increased 1.05, 1.16, and 1.17 times in spring, and increased 1.04, 1.10, and 1.12 times in autumn. During the promotion effect stages, the averaged concentrations of OH, HO_2 and RO_2 decreased 1.02, 1.03, and 1.06 times in spring, and decreased 1.02, 1.04, and 1.05 times in autumn. These results indicated that the changes in ROx dominated the $P(O_3)$ trend without PAN photochemistry. Furthermore, the $P(O_3)$ level

during promotion effect stages (5.95 ppbv h^{-1} in spring, 5.76 ppbv h^{-1} in autumn) was higher than that during inhibition effect stages s (4.88 ppbv h^{-1} in spring, 5.24 ppbv h^{-1} in autumn). For model-simulated $P(O_3)$ and O_3 budgets (Fig. 8a), HO_2+NO (account for 70±4%) and RO_2+NO (30±6%) were the main pathways of O_3 formation, and the main loss reactions were OH+NO_2 (83±12%).

PAN competed with O_3 precursors and terminated the radical chain to suppress O_3 formation by decreasing the ROx production during the inhibition effect stages. During the promotion effect stages, the intensive atmospheric oxidation capacity and photochemical reaction enhance the ROx formation rates from PAN to promote O_3 formation (Fig. 8b).

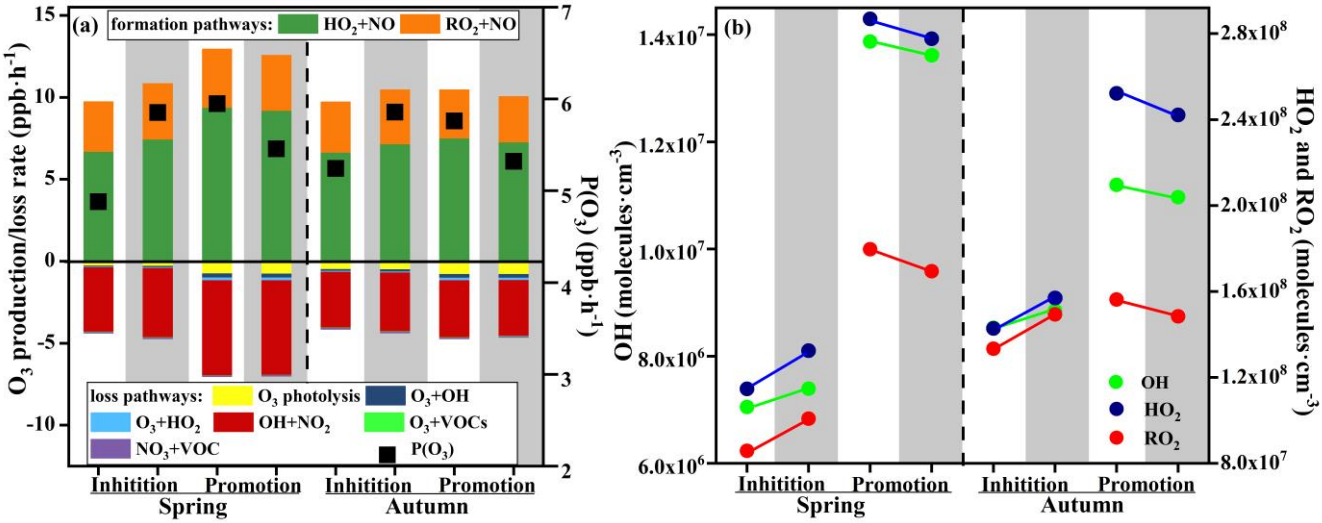

Fig. 8. Model-simulated (a) net O_3 production rate and O_3 budgets, (b) OH, HO_2, and RO_2 on the inhibition effect stages and promotion effect stages. Note: the white background parts represent the SC1 scenarios using the MCM mechanism, and the gray background parts represent the SC2 scenarios using the MCM mechanism with PAN chemistry disabled.

3.4.2 The influencing factors during inhibition and promotion stages

Table S4 showed the air pollutants and meteorological parameters during the inhibition effect stages and promotion effect stages. In detail, the levels of CO and the precursors of O_3 and PAN during the inhibition effect stages were significantly higher than those during the promotion effect stages. However, the $PM_{2.5}$ level during the inhibition effect stages was relatively lower than that during the promotion effect stages, reflecting the influence of heterogeneous reactions on $PM_{2.5}$ by suppling key photochemical oxidants to enhance PAN production (Xu et al., 2021). In addition, SO_2 and wind speed were comparable during the two scenarios. During the promotion effect stages, UV and T were significantly high, while P and RH were significantly low ($P<0.01$). Meanwhile, the PAN (1.89 in spring, 1.58 ppbv in autumn) and O_3 (50.26 ppbv in spring and 53.51 in autumn) under the promotion effects were higher than those under the inhibition effects (PAN: 1.04 and 0.84 ppbv; O_3: 27.32 and 36.42 ppbv in spring and autumn,

494 respectively).

495 In general, ROx radicals dominated the atmospheric oxidative capacity and were the indicators of
496 atmospheric photochemical reaction (Li et al., 2018). According to Section 3.2 of GAM analysis, we
497 chose the factors of NO, TVOCs, $PM_{2.5,}$ UV, T, RH, WS, and △ROx (△ROx=△OH+△HO_2+△RO_2), to
498 discuss the key influencing factor under promotion effect stages. Here, the △P(O_3) rate and the relevant
499 influencing factors were set as the response and explanatory variables, respectively. Table 2 showed the
500 influencing factors on △P(O_3) under promotion effects in spring and autumn. The factors that did not pass
501 the significance test were deleted. As the adjusted model showed, the adjusted R^2 and deviance explained
502 for the smoothed variables in four GAM models ranged from 0.67~0.78 and 70%~80%, verifying the
503 good fitting effect of the multiple-factor GAM model. According to the F-values, the effects of △ROx
504 (21.56 in spring; 45.45 in autumn) and UV (9.66 in spring; 30.55 in autumn) were the main factors leading
505 to the promotion effect in both seasons. Both △ROx and UV had significant positive non-linear
506 relationships with △P(O_3) during promotion effect stages in both seasons (Fig. S9 and S10). The minor
507 influences of WS and T were observed in autumn. The promotion effects easily happened during periods
508 of favorable meteorological conditions for photochemical reactions.

509 Liu et al. (2021) found that PAN photochemistry inhibited O_3 production under low-NOx and low-
510 ROx conditions, and promoted O_3 formation under high-NOx. However, in this study, surplus NOx
511 prevented NOx from being the limiting factor photochemical formation of secondary pollution and the
512 change of NOx could be ignored. Whether PAN photochemistry suppressed or enhanced O_3 production
513 mainly depended on the meteorological conditions of photochemical reaction and the ROx levels.

514

515 **Table 2 Estimated degree (during promotion effect scenarios in spring and autumn) of freedom (Edf), degree of**
516 **reference (Ref. df), P-value, F-value, deviance explained (%), adjusted R^2, deviance contribution (%) for the**
517 **smoothed variables (including NO, △ROx, TVOCs, $PM_{2.5}$, UV, T, RH, and WS) in the multiple-factor GAM model.**

	Incipient				Adjusted			
Smoothed variables	Edf	Ref.df	F-value	P-value	Edf	Ref.df	F-value	P-value
Promotion effect stages in spring								
NO (ppbv)	5.58	6.39	2.09	0.06			Delete	
ROx (molecules·cm^{-3})	5.99	7.06	22.88	0.00	5.72	6.83	21.56	0.00
TVOCs (ppbv)	1.14	1.26	0.60	0.40			Delete	
$PM_{2.5}$ (ppbv)	1.98	2.51	2.62	0.07			Delete	
UV ($W·m^{-2}$)	3.89	4.80	7.40	0.00	2.98	3.73	9.66	0.00
T (℃)	1.00	1.00	1.88	0.17			Delete	
RH (%)	1.00	1.00	0.86	0.36			Delete	
WS ($m·s^{-1}$)	1.41	1.71	3.03	0.13			Delete	
Promotion effect stages in autumn								
NO (ppbv)	1.15	1.28	0.20	0.66			Delete	

ROx (molecules·cm^{-3})	7.10	8.06	41.04	0.00	7.37	8.26	45.45	0.00
TVOCs (ppbv)	1.00	1.00	0.00	0.97			Delete	
$PM_{2.5}$ (μg·m^{-3})	1.00	1.00	0.53	0.47			Delete	
UV (W·m^{-2})	3.11	3.87	28.90	0.00	3.07	3.83	30.55	0.00
T (°C)	2.26	2.87	4.73	0.01	2.28	2.88	7.41	0.00
RH (%)	1.50	1.87	0.58	0.62			Delete	
WS (m·s^{-1})	4.67	5.76	2.73	0.02	4.53	5.60	3.66	0.00

4. Conclusions

Field observation was continuously conducted in spring and autumn in a coastal city of Southeast China. We clarified the seasonal variations of PAN pollution, formation mechanisms, influencing factors, and impacts on O_3 production. The average levels of PAN in autumn were lower than that in spring, while the O_3 showed the opposite characteristics. The multiple-factor GAM model showed that the key factors on PAN mixing ratio were UV, Ox, and T in spring, while Ox, TVOCs, T, and $PM_{2.5}$ played important roles in PAN formation in autumn. The MCM model is an ideal tool to explore PAN photochemical formation and its key precursors at the species level and provides more relevant suggestions for reducing photochemical pollution. The controlling emissions of aromatics and alkenes with ≤5 carbons were benefit for PAN pollution mitigation, and carbonyl compounds especially acetaldehyde were dominant in the PAN production mechanism. PAN presented the inhibition or promotion effects on O_3 under different environmental conditions. The promotion effects of PAN on O_3 mainly happened during the periods of 11:00-16:00 LT, most of which concentrated on PAN pollution episodes. According to the GAM analysis, the levels of ROx and UV were the main factors leading to the promotion effects in both seasons. Overall, PAN stimulated O_3 formation under high levels of UV, T, and ROx in the coastal city. These results indicate that the monitoring of PAN and its precursors and the quantification of its impacts on O_3 formation have significant guidance on photochemical pollution control. The scientific analysis methods used in this study provide a reference for the research on the formation mechanism of PAN and O_3 in other regions.

Authorship Contribution Statement

Taotao Liu performed chemical modeling analyses of OBM-MCM and wrote the paper. Taotao Liu collected the data, contributed to the data analysis. Jinsheng Chen and Youwei Hong designed and revised the manuscript. Jinsheng Chen supported funding of observation and research. Gaojie Chen, Lingling Xu, Mengren Li, Yanting Chen, Xiaoting Ji, Chen Yang, and Yuping Chen contributed to discussions of results. Weiguo Huang, Quanjia Huang and Hong Wang provided part of the data in Xiamen.

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