1	Measurement Report: Rapid changes of chemical characteristics and health risks for high
2	time-resolved trace elements in PM _{2.5} in a typical industrial city in response to stringent
3	clean air actions
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14	Abstract
15	Atmospheric trace metals entail significant damages in human health and ecosystem safety, and thus
16	a series of clean air actions have been implemented to decrease the ambient element concentrations.
17	Unfortunately, the impact of these emission control measures on element concentrations in fine
18	particles remains poorly understood. In our study, the random forest (RF) model was applied to
19	distinguish the effects of emission and meteorology to trace elements in $\text{PM}_{2.5}$ in a typical industrial
20	city named Tangshan based on a three-year (2017-2020) hourly field observation. The result
21	suggested that the clean air actions have facilitated the dramatic decreases of the deweathered
22	concentrations of Ga, Co, Pb, Zn, and As by 72%, 67%, 62%, 59%, and 54%, respectively. It is
23	attributable to the strict implementation of "coal to gas" strategies and optimization of industrial
24	structure and layout. However, the deweathered levels of Ca (8.3%), Cr (18.5%), and Fe (23%) only
25	displayed minor decreases, indicating that the emission control measures for ferrous metal smelting
26	and vehicle emission were not very effective. The positive matrix factorization (PMF) results
27	suggested that the contribution ratios of biomass burning, non-ferrous metal smelting, coal

28 combustion, ferrous metal smelting, heavy oil combustion, and traffic-related dust changed from 29 33%, 11%, 15%, 13%, 3%, and 25% to 33%, 8%, 8%, 13%, 4%, and 33%, respectively. To date, 30 no significant noncarcinogenic and carcinogenic risks were observed for all of the elements, while 31 both of As and Pb still showed relatively high health damages. It was proposed to further cut down the combustion-related emissions (e.g., As and Pb) because it showed the highest marginal health 32 33 benefits. Besides, the control of traffic-related emissions might be a key abatement strategy to 34 facilitate the reduction of elements in fine particles.

35 Keywords: hourly trace elements; chemical characteristics; health risks; clean air actions; Tangshan

36 1. Introduction

37 Along with the rapid economic development and accelerated urbanization, the energy 38 consumption and output of various industrial products worldwide displayed persistent increases, 39 thereby leading to massive emissions of elements especially trace metals into the atmosphere (Tian 40 et al., 2015; Zhu et al., 2020). These elements injected into the atmosphere could pose great threat 41 to the terrestrial and aquatic ecosystem via dry/wet deposition and then endanger human health 42 through the physicochemical transfer and bioaccumulation in food chains (Fernandez et al., 2000; 43 Harmens et al., 2010; Storelli, 2008). For instance, some toxic trace metals including cadmium (Cd), lead (Pb), and mercury (Hg) were often regarded as human carcinogens even in trace amounts 44 45 (Micheline et al., 2019; Olujimi et al., 2015). Besides, the excessive accumulation of some 46 biological essential elements such as copper (Cu), iron (Fe), and zinc (Zn) could initiate activation of inflammatory cascades in tissues and the induction of biochemical synthesis pathways by 47 catalyzing the generation of reactive oxygen species (ROS) (Alies et al., 2013; Lopez-Cruz et al., 48 2016; Saffari et al., 2014), though minor enrichment of these elements was beneficial to the human 49 50 health and plant growth (Oldani et al., 2017). Apart from the health impacts, some transition metals 51 (e.g., Ni, Zn) could catalyze some chemical reactions such as particle-phase sulfate generation and 52 heterogeneous production and removal of gas-phase hydrogen oxide radicals (HOx) to aggravate the 53 haze formation (Clements et al., 2013; Guo et al., 2014). Therefore, it is highly imperative to 54 recognize the pollution status of elements in the atmosphere, to identify the major sources and then to propose effective control measures to alleviate their negative effects on air pollution and human 55 56

health especially in some developing countries.

57 In the past decades, hundreds of studies investigated the pollution levels of elements and 58 revealed their sources in various study regions including urban (Das et al., 2018; Duan and Tan, 59 2013; Lyu et al., 2017; Grivas et al., 2018; Clements et al., 2014), marine (Shi et al., 2015; Witt et al., 2006), mountainous (Kang et al., 2016). Most of these studies used filter sampling (one sample 60 or two samples each day) coupled with offline analysis using inductively coupled plasma mass 61 spectrometry (ICP-MS) or inductively coupled plasma-atomic emission spectrometry (ICP-AES) to 62 63 determine the element concentrations in the atmosphere (Ao et al., 2019; Lin et al., 2016). Although 64 these studies have obtained much valuable information about the occurrence levels and key sources 65 of ambient elements, the low time-resolution data cannot accurately reflect the dynamic transformation and evolution of ambient elements. It was well known that atmospheric emissions, 66 transport and deposition significantly relied on rapidly evolving meteorological conditions (Holden 67 68 et al., 2016; Rasmussen, 1998), and thus the offline samples inevitably ignored the impacts of 69 environmental shifts with rapid temporality on the atmospheric element concentrations. Moreover, most of current source apportionment studies employed receptor models (positive matrix 70 71 factorization (PMF)) to determine the potential sources of elements (Jeong et al., 2016; Lyu et al., 72 2017), and the accuracy of these models was strongly dependent on the sample size and time 73 resolution (Liu et al., 2018b). In this regard, the high time-resolved observation of atmospheric 74 elements provided an unprecedented opportunity to characterize the occurrence levels, identify their 75 major sources, and assess the health impacts. 76 To date, only a few studies applied the high-resolution devices to capture the hourly variability

of ambient elements. Prati et al. (2000) firstly used Particle Induced X-ray Emission (PIXE) 77 measurements to measure hourly trace elements in Genoa in Italy. Following this work, 78 79 D'Alessandro et al. (2003) and Dall'Osto et al. (2013) also employed the same technique to 80 determine the trace metals in Italian towns and Barcelona, respectively. Later on, Jeong et al. (2016) 81 used the Xact metals monitor to reveal the temporal variability of atmospheric elements in Toronto, 82 Canada in summer and winter during 2013-2014. Recently, the Xact metals monitor has begun to 83 be employed in China due to the higher accuracy and convenience. Chang et al. (2018) firstly used 84 the online multi-element analyzer to achieve a one-year near real-time observation of ambient elements in China and found that traffic, nonferrous metal smelting and coal combustion were major 85

sources of atmospheric trace metals. Afterwards, Cui et al. (2019) applied the analyzer to monitor atmospheric elements during a full year, and demonstrated that dust, industry, and biomass burning were the dominant sources of most trace elements in Beijing, accounting for 36%, 10.7%, and 27% of total PM_{2.5} concentration, respectively. Up to date, continuous hourly element observation was only performed less than one year in most of the previous studies, and the long-term temporal variability of absolute concentrations and key pollution sources of atmospheric elements cannot be fully revealed.

93 Since 2013, Chinese government proposed a strict Air Pollution Prevention and Control Action 94 Plan (the Action Plan) across China and the emissions of multiple gaseous pollutants showed 95 significant decreases (Ma et al., 2019; Li et al., 2021a). In turn, the absolute concentrations and 96 health effects of air pollutants also experienced the rapid changes due to these stringent control 97 measures. Zhang et al. (2019) reported that the population-weighted annual mean PM2.5 98 concentration decreased from 62 to 42 µg/m3 during 2013-2017 and reduced PM2.5-attributable premature deaths by 0.4 million due to the impact of the Action Plan. Shortly after that, Geng et al. 99 100 (2019) estimated that the population-weighted mean concentrations of SO₄²⁻, NO₃⁻, and NH₄⁺ in 101 PM2.5 decreased from 11.1, 13.8, and 7.4 µg/m3 to 6.7, 13.1, and 5.8 µg/m3, respectively, during the 102 same period. Nevertheless, the impact of the Action Plan on trace elements in fine particles still 103 remained poorly understood. Especially, the knowledge about the variation of source apportionment 104 and health risks for trace elements response to the Action Plan was extremely limited. Moreover, most of the previous studies only utilized the original concentrations to analyze the impact of the 105 clean air policy (He et al., 2021; Xiao et al., 2020). It was well known that the pollutant 106 concentrations in the atmosphere were affected by meteorology and anthropogenic emissions 107 108 simultaneously (Li et al., 2021b), and the use of original element concentrations alone cannot assess 109 the unique contribution of emission reduction to the air pollutants. Thus, it is urgently needed to 110 remove the effect of meteorology and accurately capture the independent influence of the Action 111 Plan on the chemical characteristics, source apportionment, and health risks of trace elements. Such 112 knowledge is critical to design effective air pollution mitigation strategies in the near future.

113 As a heavily industrialized city located in the North China Plain (NCP), Tangshan possesses

114 many energy-intensive industries including coal-fired power plants, non-ferrous smelting industries,

115 textiles, building materials, chemical engineering, and papermaking industries (Ren et al., 2011). 116 Intensifying industrial development and urbanization aggravated local air quality. Previous studies 117 performed in Tangshan focused on the trace metals in soils and dusts (Cui et al., 2020; Song et al., 118 2011), whereas no study analyzed the long-term and high-resolution variabilities of atmospheric 119 elements. Since 2013, many emission control measures such as the establishment of desulfurization 120 and denitration facilities for the coal-fired power sector have been strictly implemented in Tangshan 121 (Ma et al., 2019). Especially after 2017, the coal to gas project has started to be implemented in 122 Tangshan and the energy structure underwent significant change (Wang et al., 2020b). In response 123 to substantial pollution control policies, the chemical compositions and major sources of trace elements might show corresponding change. Here, we conducted a near real-time measurement of 124 125 atmospheric elements in PM_{2.5} using a Xact multi-metals analyzer in Tangshan, China, during 126 September 2017 to August 2020. The primary objectives of our study were to (1) determine the 127 occurrence levels of elements in PM2.5 of Tangshan; (2) to analyze the seasonal and intra-week variations of atmospheric elements and to distinguish the separate contributions of emission and 128 129 meteorology to these species; (3) to quantify the changes of major sources for atmospheric elements 130 during this period; (4) to assess the changes of health risks in response to these pollution control 131 measures.

132 2. Material and methods

133 2.1 Sampling site

134 The sampling site (39.66°N, 118.18°E) is situated on the rooftop (~20 m above the ground) of a building in the urban district of Tangshan and no high buildings spread around within 100 m range. 135 136 The sampling site is close to some major roads including the Airport Road, Huayan North Road, 137 and Changhong Road. A large number of commercial streets and recreation facility surround the 138 site. Although no big industrial point source was closely adjacent to the sampling site, many 139 potential pollution sources were located more than 15 km away from the site. For instance, the Beihu 140 industrial region is located about 15 km in the eastern direction of the site. Some large iron steel 141 industries and nonferrous/ferrous smelting industries were located on the north of sampling site (more than 30 km). Besides, most of the large petrochemical industries, coal-fired power plants, and 142 143 shipping industries focus on the Caofeidian and Haigang developing zones, both of which were 144 located about 50 km in the South area of the sampling site. The detailed location is depicted in

145 Figure 1.

146 2.2 Instrumentation

147 Hourly mass concentrations of 22 elements, including Ag, As, Au, Ca, Co, Cu, Cd, Cr, Fe, Ga, 148 Hg, K, Mn, Ni, Pb, Pd, Sb, Se, Sn, Tl, V, and Zn in PM2.5 were determined continuously by an online 149 multi-element analyzer (Model Xact 625, Cooper Environment Service, USA) (Table S1). The 150 sample air is drawn through a small spot on the tape where the PM2.5 was collected at a flow rate of 151 16.7 L min⁻¹ during September 2017-August 2020. An internal Pd pod is utilized as an internal 152 standard to determine the stability of the instrument. TI was removed from the datasets because over 95% of their concentrations were below the limit of detection (LOD) (Table S2). Au, Cd, Sn and Sb 153 154 were also excluded from the datasets because over 50% of the concentrations for these metals were 155 below the LOD. To validate the reliability of the online multi-element analyzer, many previous 156 studies used the filter sampling coupled with ICP-MS and ICP-AES to determine the daily concentrations of elements and confirmed that the online device showed good agreement with the 157 filter sampling (Furger et al., 2017; Tianxue et al., 2006). Hourly averaged meteorological 158 159 parameters including air temperature (T), relative humidity (RH), air pressure (P), wind direction (WD), and wind speed (WS) during the sampling period were measured by a weather station with 160 161 sonic anemometer (150WX, Airmar, Milford, NH, USA). The hourly mass concentration of PM2.5 was determined by a particulate monitor (Thermo, FH62C-14). The routine procedures, including 162 the daily zero or standard calibration, span and range check, station environmental control and staff 163 certification, followed the Technical Guideline of Automatic Stations of Ambient Air Quality in 164 Tangshan based on the national specification HJ/T193-2005, which was revised based on the 165 166 technical guidance established by the US EPA. Quality Assurance and Quality Control (QA/QC) for 167 the Xact measurements was implemented throughout the sampling period. The internal Pd upscale 168 value was recorded after daily programmed test for the instrument. 169 2.3 Deweathered model development

170 The concentrations of air pollutants were affected by meteorological parameters and emissions

171 simultaneously. In order to separate the contributions of emissions, the impacts of meteorological

172 conditions must be eliminated. In this study, a typical machine-learning model named random forest

173 (RF) approach was applied to distinguish the effects of emissions and meteorological conditions 174 (Chen et al., 2018). All trace elements in PM2.5 were treated as the dependent variables. The time 175 predictors (year, day of year (DOY), day of week (DOW), hour of day (HOY)) and meteorological factors including air temperature (T), relative humidity (RH), wind speed (WS), wind direction 176 (WD), and air pressure (P) were regarded as the predictors (Figure S1). The original dataset was 177 178 randomly classified into a training dataset (80% of input dataset) for developing the RF model and 179 the remained remaining one was treated as the test dataset. After the building of the RF model, the 180 deweathered technique was employed to predict the concentrations of trace elements at a specific 181 time point. The deweathered element concentrations served as the concentrations contributed by 182 emission alone. The differences between the original element concentrations and the deweathered 183 element concentrations were regarded as the concentrations contributed by meteorology. Many 184 statistical indicators including R² value, root-mean-square error (RMSE), and mean absolute error 185 (MAE) were regarded as the major indicators to evaluate the RF modelling accuracy. The RF model with the 5-fold cross-validation R² value less than 0.5 was considered to be an unconvincing result 186 187 and cannot reflect the impacts of emission and meteorology on air pollutants accurately because 188 more than 50% variability of the training model cannot be appropriately explained. After the model 189 evaluation, only the trace elements with the cross-validation R² values larger than 0.5 were selected 190 to estimate the respective contributions of emission and meteorology to the total element 191 concentrations.

192 2.4 PMF model

As a typical receptor model applied to source apportionment, the PMF 5.0 version was used to 193 194 identify the major origins of the atmospheric elements and to determine the contribution ratio of 195 each source to these elements (Norris et al., 2014). The objective of PMF is to solve the issues of 196 chemical mass balance between the measured concentration of each element and its source 197 contributions by decomposing the input matrix into factor contributions and factor profiles. The 198 detailed equation is shown in Eq. (1). Besides, the contribution of each source for an individual 199 element must be non-negative because no sample has a negative source contribution. In brief, the basic principle of PMF is to calculate the least object function Q when the gik must be a positive-200 201 definite matrix based on Eq. (2) (Paatero and Tapper, 1994; Reff et al., 2007).

202
$$x_{ij} = \sum_{k=1}^{p} g_{ik} f_{kj} + e_{ij} \quad (1)$$

203
$$Q = \sum_{i=1}^{n} \sum_{j=1}^{m} \left[\frac{x_{ij} - \sum_{k=1}^{p} g_{ik} f_{kj}}{u_{ij}} \right]^{2}$$
(2)

where x_{ij} and e_{ij} denote the concentration and uncertainty of the jth element, respectively. g_{ik} 204 205 represents the contribution ratio of the kth source to the ith sample, fkj represents the ratio of the jth 206 element in the kth source, and eii indicates the residual of the jth element in the ith sample. The 207 uncertainties associated with factor profiles were evaluated using three error calculation methods 208 including the bootstrap (BS) method, displacement (DISP) analysis, and the combination method of DISP and BS (BS-DISP). For the BS method, 100 runs were performed and the result has been 209 210 believed to be valid since all of the factors showed a mapping of above 90%. DISP analysis also 211 confirmed that the solution was considered to be stable because the observed drop in the Q value 212 was less than 0.1% and no factor swap occurred. For the BS-DISP analysis, the solution has been 213 verified to be useful because the observed drop in the Q value was less than 0.5%. Furthermore, both of the results from BS and BS-DISP did not suggest any asymmetry or rotational ambiguity 214 215 for all of the factors (Manousakas et al., 2017; Taghvaee et al., 2018).

216 2.5 Health risk assessment of trace metals in PM_{2.5}

As a typical industrial city, Tangshan possesses a large number of residents and poor air quality. 217 218 Therefore, the residents in Tangshan might suffer from severe exposure risks of trace metals. In our 219 work, the carcinogenic and non-carcinogenic risks of trace metals in PM2.5 were evaluated based on 220 some statistical threshold proposed by the International Agency for Research on Cancer (IARC). 221 Based on the criterion of the IARC, As, Ni, Cr, and Pb were considered to be carcinogenic to humans. 222 The carcinogenic and non-carcinogenic risks induced by metal exposure for adults and children 223 were evaluated based on the carcinogenic risks (CR) and hazard quotient (HQ). The formulas for 224 calculating ADD, CR, and HQ are as follows: ADD=(C×InhR×EF×ED)/(BW×AT) 225 (3) 226 HQ=ADD/RfD (4) 227 $CR = ADD \times CSF$ (5)

228 where C (mg m⁻³) denotes the concentration of the corresponding trace metal in PM_{2.5}; InhR is the

respiratory rate $(m_1^3 d_1^{-1})$; EF represents the annual exposure frequency (d y⁻¹); ED is the exposure duration (year); BW is the average body weight (kg); AT denotes the average exposure time (d); ADD means the daily intake (mg/kg/day) of trace metals; RfD represents the reference dose (mg kg⁻¹ d⁻¹), calculated with reference concentrations; CSF is the cancer slope factor (kg d mg⁻¹). The potential non-carcinogenic risk of the trace metal might be high when HQ was above 1.0, whereas the health risk is not obvious when HQ is below 1.0. The carcinogenic risk of each trace metal is evaluated based on whether CR is higher than 10⁻⁴.

236 3. Results and discussion

237 3.1 Occurrence levels and inter-annual variations of original element concentrations

238 The total mass concentrations of 16 elements in PM2.5 of Tangshan varied between 230 ng/m3 239 to 20000 ng/m³, with the average value (±standard deviation) of 3100 ± 900 ng/m³. The total 240 element concentrations in Tangshan accounted for 5.7% of the total mass concentrations of PM2.5, 241 which was slightly higher than those in Beijing (4.7%) and Qingdao (4.0%), and significantly higher 242 than that in Shanghai (1.80%) (Chang et al., 2018; Cui et al., 2019). As depicted in Figure 2, the 243 average concentrations (± standard deviation) concentrations of these elements followed the order 244 of K (1400 \pm 950 ng/m³) > Fe (880 \pm 590 ng/m³) > Ca (330 \pm 270 ng/m³) > Zn (320 \pm 160 ng/m³) > 245 Pb $(58 \pm 36 \text{ ng/m}^3) > \text{Mn} (54 \pm 32 \text{ ng/m}^3) > \text{Cu} (22 \pm 19 \text{ ng/m}^3) > \text{As} (15.3 \pm 11.0 \text{ ng/m}^3) > \text{Se} (6.5 \pm 10.0 \text{ ng/m}^3) > \text{Se} ($ 246 $\pm 5.3 \text{ ng/m}^3$) > V (4.0 $\pm 3.6 \text{ ng/m}^3$) > Cr (2.8 $\pm 2.2 \text{ ng/m}^3$) > Ag (2.8 $\pm 2.1 \text{ ng/m}^3$) > Ni (2.2 $\pm 1.8 \text{ ng/m}^3$) = Ni (2.2 \pm 1.8 \text{ ng/m}^3) = Ni (2.2 \pm 1.8 \text{ ng/m} 247 ng/m^3) > Hg (1.5 ± 0.8 ng/m³) > Ga (0.9 ± 0.7 ng/m³) > Co (0.7 ± 0.2 ng/m³). Among all of these elements, K, Fe, Zn, and Ca were the most abundant species, accounting for 95% of the total 248 249 elements in PM2.5. The remaining element concentrations only accounted for less than 6% of the 250 total element concentrations, which was similar to in previous studies (Chang et al., 2018; Cui et al., 251 2019). Nearly all of the trace elements in Tangshan, Beijing, Qingdao, and Shanghai were 252 significantly lower than those in Zibo during 2006-2007 (Table S3). It suggested that the trace 253 elements in China experienced marked decreases in the past decades (Zhang et al., 2018). Compared 254 with some cities in some developed countries, all of the trace element concentrations were 255 significantly higher than those in London and Toronto. Moreover, the concentrations of K, Ca, V, 256 Cr, Mn, and Fe in Tangshan were higher than those in Venice, Italy.

257 Due to the higher exposure risk and great threat to human health, it is necessary to compare the

设置了格式: 上标 **设置了格式:** 上标 258 trace metal concentrations with the risk threshold proposed by many organizations or countries. As 259 shown in Table 1, we have collected many risk thresholds in different countries and found that the Hg ($1.5 \pm 0.8 \text{ ng/m}^3$), Ni ($2.2 \pm 1.8 \text{ ng/m}^3$), and Pb concentrations ($58 \pm 36 \text{ ng/m}^3$) in Tangshan were 260 261 significantly lower than the thresholds of the Chinese Ambient Air Quality Standard (CAAQS) (Hg: 50 ng/m³), World Health Organization (WHO) (Hg: 1000 ng/m³, Ni: 25 ng/m³, and Pb: 1000 ng/m³), 262 European Union (EU) (Ni: 20 ng/m³), and the United States (Pb: 150 ng/m³). However, both of the 263 264 As $(15 \pm 11 \text{ ng/m}^3)$ and Cr concentrations $(2.8 \pm 2.2 \text{ ng/m}^3)$ in PM_{2.5} of Tangshan were much higher 265 than the standard values of the CAAQS (As: 6.0 ng/m^3 and Cr: 0.03 ng/m^3), WHO (As: 6.6 ng/m^3 266 and Cr: 0.25 ng/m³), and EU (As: 6.0 ng/m³).

267 The inter-annual variation of the original concentrations of the trace elements in PM2.5 are 268 depicted in Figure 3 and S2-S3. The original concentrations of all the trace elements exhibited 269 decreasing trends. The Cu, Co, Zn, Pb, As, and Ga concentrations showed dramatic decreases from 270 37 to 12 ng/m³ (68%), 1.21 to 0.4 ng/m³ (66%), 400 to 190 ng/m³ (53%), 71 to 40 ng/m³ (44%), 20 to 11 ng/m3(44%), and 1.09 to 0.6 ng/m3(42%), respectively. Following these species, the K (40%), 271 272 Ag (39%), V (39%), Ni (36%), Ca (33%), Mn (29%), Se (29%), Fe (27%), and Cr (21%) 273 concentrations showed moderate decreasing ratios. The observed Hg level exhibited the lowest 274 decreasing ratio from 1.59 to 1.43 ng/m³ (9.9%).

275 3.2 Impact of emission reduction on trace element concentrations

276 Although the original concentrations of the trace elements could be utilized to analyze the impact 277 of the clean air policy, the role of emission reduction on the element concentration might not be 278 clearly clarified because the meteorological factors were also important variables affecting the air 279 quality. In order to accurately reflect the response of the element concentrations to the emission 280 reduction alone during 2017-2020, the meteorological conditions were eliminated by the RF model 281 in our study. Based on the results in Figure S4, the RF models for all of the species showed better 282 performance because their R² values were higher than 0.50, and the slopes of all of the fitting curves 283 were also close to the R² values. The result suggested that the separation of meteorology and emission of trace elements based on the RF model was reliable. During 2017-2020, the deweathered 284 concentrations of Ga, Co, Pb, Zn, and As showed the rapid decreases from 1.52 to 0.4 ng/m³ (72%), 285 1.31 to 0.4 ng/m³ (67%), 92 to 35 ng/m³ (62%), 410 to 170 ng/m³ (59%), and 21 to 10 ng/m³ (54%), 286

287 respectively (Figure 3). It was well known that As, Co, and Pb were typical marker elements for 288 coal combustion and the "coal-to-gas" and "coal-to-electricity" strategies have been widely 289 performed in Tangshan (Fang et al., 2020; Li et al., 2017). Wang et al. (2020a) have estimated that 290 these effective control measures have contributed to around 60% of the total PM_{2.5} reductions. 291 Meanwhile, the upgradation and optimization of the industrial structure/layout and the shutdown of 292 high-pollution industries were also strictly implemented in Tangshan, and thus led to the dramatic 293 decreases of Ga and Zn concentrations because Ga and Zn were common forms of nonferrous metal 294 smelting (Tian et al., 2015). In contrast, the deweathered Ca level displayed the lowest decrease 295 ratio (8.3%) from 2017 to 2020, indicating that clean air actions cannot significantly reduce the fugitive emissions. In addition, the deweathered Fe (23%) and Cr (18.5%) also suffered from 296 297 relatively low decrease ratios. It was well documented that Fe and Cr originated from metallurgical 298 industry such as steel production and ferrous metal smelting (Tian et al., 2015), and the slight 299 decreases of the deweathered Fe and Cr levels during the sampling period suggested that the 300 emission control measures for ferrous metal smelting should be strengthened in the future.

301 In addition, the decreasing ratios of the deweathered concentrations for each species displayed 302 different seasonal characteristics. The deweathered concentrations of some elements related with 303 industrial activities (e.g., Ga, Zn, and Cr) showed rapid decreases in autumn and winter compared 304 with other seasons during 2017-2020 (Figure 4), indicating that the optimization of the industrial 305 layout and shutdown of outdated industries were effective to decrease these element emissions 306 especially in the high-pollution season. Some elements derived from biomass burning including K 307 (66%) and Se (50%) also exhibited the most dramatic decreasing ratios in autumn. It was assumed 308 that enhanced crop residual burning occurred frequently during the autumn harvest season. Ke et al. 309 (2019) confirmed that the number of fire spots in October-November was even higher than that in 310 June and the burned area in the harvest season was highest during 2013-2017. However, the control 311 on open biomass burning has been implemented strictly in recent years, largely reducing the K and 312 Se emissions in autumn. It should be noted that the deweathered Pb (46%), Co (65%), and As (45%) 313 concentrations in winter did not display high decreasing ratios though the annual mean deweathered 314 Pb, Co and As levels experienced dramatic decreases. The result revealed that it was still difficult 315 to reduce the Pb, Co, and As emissions during the heating season because increased coal consumption for domestic heating largely offset the contributions of emission control measures(Zhu et al., 2018; Zhu et al., 2020).

318 Apart from the seasonal difference of each species, the decreasing ratios of these elements also 319 suffered from distinctly intra-weekly variations. The deweathered concentrations of most elements 320 except Ca, Cu, Ni, and V exhibited higher decreasing ratios at the weekends than on the weekdays 321 (Figure 5). Cui et al. (2020) have demonstrated that the weaker supervision on industrial enterprises 322 on weekends could lead to the higher concentrations of non-traffic elements such as K, As, Se, and 323 Cr in some cities. Fortunately, grid monitoring has been widely performed in Tangshan recently 324 (http://hbepb.hebei.gov.cn/hbhjt/xwzx/jicengfengcai/101624062321621.html), and many low-cost 325 sensors were installed at some energy-intensive industries, which could decrease the stealing 326 emissions of some elements. Nonetheless, the decreasing ratios of Ca, Cu, V, and Ni did not show 327 the regular intra-weekly characteristics. In recent years, Tangshan adopted strict traffic management 328 regulation and the nonlocal light duty vehicles were restricted to drive inside the urban area one day 329 per week based on the end number of the license plates (Westerdahl et al., 2009; Wu et al., 2011), 330 whereas the restrictions were not valid at weekends (Liu et al., 2007). Theoretically, the traffic 331 control should result in marked decreases of traffic-related element concentrations on weekdays 332 compared with weekends. However, in our study, some traffic-related elements such as Ca and Cu 333 did not show similar characteristics. Meanwhile, as the important tracer of vehicle emission, the 334 NOx concentration in Tangshan did not show a regular intra-weekly pattern. It was supposed that the vehicle volume in Tangshan has increased from 2.0 to 2.4 million (http://tjj.hebei.gov.cn/), 335 which largely offset the benefits of traffic controls. The shipping-related elements including V and 336 337 Ni also did not show regular intra-weekly variation because no heavy metal emission control 338 measures for shipping were performed.

339 3.3 The role of meteorology on the year-to-year variations of element concentrations

The difference between the original and <u>the</u> deweathered element concentrations could be regarded as the concentrations contributed by meteorological parameters. The positive impacts of meteorological parameters on the trace elements suggested that the meteorological conditions were unfavorable to the pollutant diffusion, while the negative impacts of meteorological indicators meant the favorable condition to trace elements. In our study, the roles of meteorological conditions 345 on Ca (-25%), V (-10%), Cr (-2.5%), Mn (-0.7%), Fe (-4.6%), Ni (-7.6%), and Cu (-21%) during 346 2017-2020 were negative (Figure S5), while the roles of meteorological parameters on other 347 elements were positive. The result suggested that those elements derived from vehicle emission (Ca, 348 Cu, and Fe), ferrous metal smelting (Cr and Mn), and heavy oil combustion (V and Ni) were less 349 sensitive to the emission reduction actions compared with other elements and the meteorological 350 conditions were much beneficial to the diffusion of these elements. In order to further reveal the key 351 meteorological factors for these elements, we used the RF model to calculate the variable 352 importance of all of these meteorological parameters including P, RH, T, WD, and WS. The result 353 suggested that Ca, Fe, and Cu were mainly influenced by T, whereas V, Ni, Cr, and Mn were often associated with WD and WS (Figure 6). During the spring and summer in 2017-2020, the average 354 air temperature decreased from 8.9 and 27 to 7.2 and 26°C, respectively. The decreased air 355 356 temperature led to a higher water content in the soil and a lower tendency of dust suspension, and 357 might decrease the concentrations of Ca, Fe, and Cu (Manju et al., 2018; Yang et al., 2017; Lyu et 358 al., 2016). Although the annual average wind speed in Tangshan decreased from 1.70 to 1.45 m/s, 359 the mean wind speed from the southeastern direction displayed a slight increase from 1.34 to 1.50 360 m/s. Zhao et al. (2013) verified that V and Ni were usually emitted from heavy oil combustion of 361 ocean-going ship engines. Many coastal ports and ferrous metal smelting industries were located on 362 the southeastern direction of the sampling site, and thus the enhanced WS might promote the 363 dilution and dispersion of trace elements (Figures S6-S8). As shown in Figure S8, both of-V and Ni 364 showed the higher concentrations in the southeastern part of Tangshan and the concentrations 365 displayed gradual decreases along the Southeast-Northwest transect, which also demonstrated that 366 both of V and Ni in the sampling site could be derived from coastal shipping emission. 367 3.4 The impact of clean air policy on the source apportionment of the trace elements

368 Although the major sources of elements could be determined based on some important tracers 369 (e.g., K, V), the contributions of the major sources to each element still remained unknown. 370 Therefore, Positive matrix factorization (version PMF 5.0) was applied to identify more source 371 information of the elements in $PM_{2.5}$ during 2017-2020 based on the deweathered levels. After 20 372 runs, more than 26000 samples were trained to determine the optimal six factors with the lowest 373 values of Q (robust) and Q (true). The BS, DISP, and BS-DISP methods confirmed that the most reliable solution was obtained with six factors. The detailed information of the PMF analysis and
error diagnostics is summarized in Tables S4-S6.

376 As shown in Figure S9, the trace elements in PM2.5 during 2017-2020 showed similar 377 characteristics. Factor 1 possesses high loadings of K (55%) and Se (42%). K and Se were often 378 regarded as the major tracers of biomass burning. Due to the increasing usage of biomass fuels for 379 domestic heating during the heating season, K and Se in PM2.5 of Tangshan showed higher values 380 in winter, suggesting that these metals in fine particles could originate from the combustion of 381 biomass fuels. Except for the domestic heating, we found some episodes during the harvesting 382 season in late summer (2500 and 11.2 ng/m3) and early autumn (2600 and 9.5 ng/m3) also showed 383 extremely high concentrations of K, which might be linked with local biomass burning (Chen et al., 384 2017). Based on the map of fire points and backward air masses trajectories (Figures S6-S8), the 385 metals released from biomass burning in the NCP could be transported to the sampling site by the 386 dominant southerly wind, which further proved the impacts of biomass burning (Chen et al., 2017). The abundant elements in factor 2 included Ag (53%), Zn (51%), and Cu (36%). Owing to the 387 388 higher temperatures during the roasting, sintering and smelting processes for the extraction of Cu, 389 and Zn from ores, some metals such as Ag in nonferrous metal ores could be vaporized as a byproduct and released into the flue gas (Pacyna and Pacyna, 2001; Wu et al., 2012). Therefore, 390 391 factor 2 was interpreted as the non-ferrous metal smelting source. 392 Factor 3 was characterized by a large mass fraction of Co (81%), Pb (61%), Hg (57%), and As (39%). After the phase-out of leaded gasoline since 1980s, the contribution from coal combustion 393 394 to Pb showed rapid increase and accounted for the major fraction of particulate Pb (Das et al., 2018).

Meanwhile, Co and Hg were also treated as important byproducts released from coal burning and the Co and Hg concentrations often increased significantly with the elevation of the burning temperature (Tang et al., 2018). Tian et al. (2015) estimated that 73% of As, 56 % of Pb, and 47 % of Hg were found to be emitted from coal combustion in China. Coal consumption in South China was mainly driven by coal-fired power plants, while the coal-based heating was the major sector for the coal consumption in the NCP. In our study, As, Co and Pb showed the higher concentrations in

401 winter (heating season) (18.7, 0.9, and 76 ng/m³) compared with other seasons (14, 0.6, and 51

402 ng/m³). The markedly seasonal discrepancies of As, Co and Pb strongly supported the impact of the

403 coal combustion for domestic heating on the enhancement of As, Co and Pb in the fine particles.

404 Factor 4 was distinguished by high loadings of Cr (78%) and Mn (39%), respectively. Cr and 405 Mn mainly originated from the metallurgical industry such as steel production and ferrous metal smelting (Liu et al., 2018a; Tian et al., 2015; Zhu et al., 2018). China was responsible for more than 406 407 49% of the world steel production in 2017 (approximate 830 million tons), and 60% of the large 408 steel producers were located in China (Chang et al., 2018). Tangshan possesses many large steel 409 production industries such as Tangshan Steel, Qian'an Steel, and Guofeng Steel. Besides, some 410 industries of Capital Steel have been migrated into Tangshan (Li et al., 2019), which might increase 411 the Cr and Mn emissions.

412 Factor 5 explained 10.1% of the total species and it was characterized by high loadings of V 413 (88%) and Ni (51%). It was well documented that V was a key fingerprint of heavy oil combustion, 414 which was generally emitted from shipping emission and petrochemical refining (Shafer et al., 415 2012). Ni was widely utilized as a tracer of fuel oil combustion in industries (Zhu et al., 2018). 416 Many oil-fired power plants were located in Tangshan for central heating (Yu et al., 2013). Based 417 on the backward trajectory and wind direction (Figures S6-S8), we found that high concentrations 418 of V and Ni might be derived from the southeastern air masses especially in summer and autumn, 419 indicating the impacts of coastal port and petroleum refinery industry. In addition, the V and Ni 420 concentrations displayed the gradual decreases along the Southeast-Northwest transect, indicating 421 the potential sources were located on Southeast Tangshan (Figure S8). Gathering evidence 422 suggested that the V/Ni ratio in petroleum coke with a low-sulfur content and fuel oil usage ranged from 1 to 3 (Moreno et al., 2010). The annual mean ratios of V and Ni in our study reached 1.2 423 424 during the sampling period, which was in the range of this interval. The result also revealed that 425 petrochemical refining and heavy oil combustion derived from coastal shipping emission might be 426 an important source of V and Ni in the fine particles.

Factor 6 was characterized by high loadings of Ca (78%), Cu (32%), and Fe (33%), and moderate loadings of Mn (31%) and Zn (29%). Some previous studies have demonstrated that Cu, Fe, and Zn were released from tyre and brake wear because they were the necessary materials for brake pads and the agents in brake linings (Dall'Osto et al., 2013; Hjortenkrans et al., 2007). Ca probably originated from the road fugitive dust because it was one of the most abundant elements in the upper soil (Alves et al., 2015; Liu et al., 2018a). Moreover, we have found <u>that</u> Fe, Ca, and Zn displayed
remarkably high values during the morning rush hours and a small peak during the sunset (Figure
S10), which was coincident to the diurnal variation of the traffic volume. Thus, the factor 6 was
identified as the traffic-related dust source.

436 Although six similar sources were revealed during 2017-2020, the contribution concentrations 437 and ratios of these sources varied greatly in these years. As shown in Figures 7 and 8, the 438 contribution concentrations of biomass burning, non-ferrous metal smelting, coal combustion, and 439 ferrous metal smelting to trace elements decreased from 1460, 480, 640, and 570 ng/m³ to 900, 230, 440 230, and 350 ng/m3, respectively. However, the contribution concentrations of heavy oil combustion 441 and traffic-related dust displayed a slight increase during 2017-2019, while they decreased rapidly 442 after 2019. The contribution concentrations for nearly all sources to the trace elements suffered 443 fromshowed decreases during 2017-2020 because the total deweathered levels of trace elements 444 experienced decreases in the past three years. However, the contribution ratios of these sources to 445 the trace elements did not show similar characteristics. For instance, the contribution ratio of the traffic-related dust increased from 25% to 33%. In contrast, the contributions of non-ferrous metal 446 447 smelting and coal combustion decreased from 11% to 8% and 15% to 8%, respectively. The 448 contributions of ferrous metal smelting, heavy oil combustion, and biomass burning remained 449 relatively stable during this period.

450 Due to the strict implementation of the clean air policy, many outdated industrial capacities were 451 shut down and cleaner technologies have been upgradedwere implemented, which facilitated the production decreases of pig iron and coal-fired power plants (Ma et al., 2019). Hence, the 452 453 contribution concentrations and ratios of non-ferrous metal smelting and coal combustion 454 experienced dramatic decreases. Although the open biomass burning has been strictly restricted in 455 Tangshan (Chang et al., 2018), the contribution ratios of biomass burning to the trace elements in 456 PM2.5 remained relatively stable, which might be attributable to the rapid decreases of the 457 contributions derived from coal combustion and non-ferrous metal smelting. In addition, the biofuel 458 combustion was still widespread in some rural and suburb areas (Kamal et al., 2015; Li et al., 2020), 459 which might offset the decreases in the contributions of open biomass burning. Although the 460 contribution concentrations of traffic-related dust to trace elements also showed a slight decrease,

the contribution ratios of traffic-related dust to some trace elements exhibited marked increases (8%) during 2017-2020 because the contribution ratios of metal smelting and coal combustion displayed substantial decreases. The result also demonstrated that the implementation of coal to gas project facilitated the decreases of trace element concentrations. In addition, the source variation trend also suggested that the formulation of many new quality standards for non-road diesel fuels cannot fully decrease the element emissions (Cui et al., 2017), and thus the control of traffic-related dust should be enhanced in the future.

468 3.5 Health risk assessment of trace metals in PM_{2.5}

Although the trace metals only accounted for a minor fraction of the total mass concentration of PM_{2.5}, it might pose a great threat to the human health because most of these metals were bioavailable and non-degradable (Rai et al., 2019; Yi et al., 2011). Unfortunately, previous studies mainly used filter sampling techniques to determine the concentrations of trace metals and then assess their health risks (Cui et al., 2018; Huang et al., 2016). These low-resolution data might not accurately reflect the real health risks triggered by metal exposure. In our study, we employed online data to assess the health risks derived from metal exposure.

476 The health risks of trace metals could be classified into two types including carcinogenic and 477 non-carcinogenic risk. Based on the major parameters summarized in Tables S7 and S8, we 478 estimated both of the carcinogenic and non-carcinogenic risks of the major metals. To evaluate the 479 impacts of emission control measures on the element concentrations, both the health risks based on 480 the original element levels and the deweathered element concentrations were calculated. The mean 481 CR values based on the original concentrations were in the order of Pb $(2.3 \times 10^{-6} (adult) and 1.4 \times 10^{-6} (adult))$ 6 (child)) > As (1.9×10⁻⁶ and 1.1×10⁻⁶) > Cr (0.11×10⁻⁶ and 0.07×10⁻⁶). The total CR values for adults 482 483 and children reached 4.3×10⁻⁶ and 2.6×10⁻⁶ (Table 2), respectively. The total CR values were located 484 in the range of the acceptable (10⁻⁶) and least stringent risk levels (10⁻⁴), which suggested that 485 Tangshan suffered from a slight metal carcinogenic risk. Among all of these metals, Pb and As 486 displayed the higher CR values. It was assumed that the coal combustion for domestic heating might 487 be the dominant factor for the higher risks of Pb and As in Tangshan because both Pb and As in PM2.5 were mainly derived from coal combustion. With regard to the non-carcinogenic risks of the 488 trace metals, the HQ of As $(1.2 \times 10^{-2} \text{ and } 2.9 \times 10^{-2})$ and Pb $(6.8 \times 10^{-4} \text{ and } 17 \times 10^{-4})$ showed higher 489

values compared with other elements. The result indicated that nearly all elements did not display
potential non-carcinogenic risk because the HQ values of all the metals were less than 1. The total
HQ value of these metals was also lower than 1, indicating that the trace elements in Tangshan did
not show a significant non-carcinogenic risk.

494 By removing the impact of the meteorological conditions, we can isolate the impact of the clean 495 air policy on health risks associated with metal exposure alone. The decreased ratios of the CR 496 values based on the deweathered As and Pb concentrations during 2017-2020 were 54% and 62%, 497 respectively (Table 3). However, the decreased ratios of the CR values based on the original As and 498 Pb levels only reached 44%. The result suggested that the clean air policy in recent years 499 significantly decreased the As and Pb emissions. Additionally, the decreased ratios of the HQ values 500 for the original Cu (41%) and Zn (53%) were much lesslower than those for the deweathered ones 501 (Cu: 47% and Zn: 59%). Nevertheless, some other elements did not show similar characteristics. 502 For instance, the decreased ratios of the HQ values for the original Cr (21%) and Fe (27%) were even slightly higher than those for the deweathered ones (Cr: 19% and Fe: 23%). It was assumed 503 504 that the clean air policy in recent years facilitated the emission reduction of non-ferrous metal 505 smelting and coal combustion efficiently. However, the concentrations of the elements derived from 506 ferrous metal smelting and vehicle emission did not show marked decreases, which was in good 507 agreement with the source apportionment result in section 3.3. Thus, in the future work, it is highly 508 imperative to further reduce the industrial/traffic-related emissions in order to alleviate potential 509 health risks.

510 3.6 Limitations and uncertainties

It should be noted that our work is still subject to some limitations. At first, some elements such as Cr (0.5) and Ga (0.5) showed relatively lower CV R^2 values though they were generally higher than 0.5. These elements might show the higher uncertainties during the meteorology-normalization compared with other elements such as Cu (0.85) and K (0.85). Besides, few variables were applied to deweather the element concentrations, which might be responsible for the lower CV R^2 value for some elements. Due to the lack of available hourly emission inventory of each element, we only used the time variable to train the model. This method still suffered from some uncertainties, which

should be improved by the establishment of near-time emission database.

519 4. Conclusions and implications

520	Three-year continuous hourly observation of elements in PM2.5 was conducted in Tangshan
521	during September 2017-August 2020. The effect of the clean air policy on the element
522	concentrations in $PM_{2.5}$ was quantified. The main conclusions were drawn-as follows:
523	(1) The deweathered concentrations of Ga, Co, Pb, Zn, and As showed rapid decreases from 1.52
524	to 0.42 ng/m ³ (72%), 1.31 to 0.44 ng/m ³ (67%), 92 to 35 ng/m ³ (62%), 411 to 170 ng/m ³ (59%),
525	and 21 to 10 ng/m ³ (54%), respectively. The clean air actions played the important role on the
526	emission reduction of coal combustion and non-ferrous metal smelting.

- 527 (2) The deweathered levels of Ca (8.3%), Cr (19%), and Fe (23%) displayed relatively low
 528 decreases compared with other elements, indicating that the vehicle emission and ferrous529 smelting industries might not be sensitive to the air clean policy.
- 530 (3) The deweathered levels of some elements related with industrial activities (e.g., Ga, Zn, and Cr)
- 531 exhibited rapid decreases in autumn and winter compared with other seasons during 2017-2020,
- 532 while the combustion-related elements such as Pb and As did not show high decreasing ratios
- in winter. The enhanced coal consumption during the heating season offsets the benefits derivedfrom strict emission control measures.
- (4) The favorable meteorological conditions promoted the decreases of Ca (-25%), V (-10%), Cr (2.6%), Mn (-0.68%), Fe (-4.6%), Ni (-7.6%), and Cu (-21%) concentrations.
- (5) The contribution ratios of biomass burning, non-ferrous metal smelting, coal combustion,
 ferrous metal smelting, heavy oil combustion, and traffic-related dust changed from 33%, 11%,
- 539 15%, 13%, 3%, and 25% to 33%, 8%, 8%, 13%, 4%, and 33%, respectively.
- (6) All elements did not show significant noncarcinogenic and carcinogenic risks, while both Asand Pb still displayed relatively high health damages.
- 542 Our study presented detailed information about the impact of clean air policy on the chemical 543 compositions and source apportionment of trace elements in PM_{2.5} in Tangshan, and provided new 544 insights for the scientific community and policymakers. Many targeted measures could be 545 undertaken to alleviate the air pollution and further to reduce avoided premature health risks.
- 546 However, this study still suffered some limitations and more steps will be taken toward thoroughly
- 547 addressing these problems. First of all, the PMF model still showed some uncertainties, and thus

- 548 characterizing the isotopic signatures of the elements is of great significance. In addition, a Sunset
- 549 OC/EC analyzer, a Monitoring of Aerosols and Gases (MARGA) platform, and other on-line
- 550 measurements should be collocated to probe into the synergistic effect of emission reduction and
- 551 meteorology on air quality.
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- 554 Data availability
- 555 The boundary layer height dataset was obtained from the website of https://www.ecmwf.int/. The
- 556 dataset is archived in https://zenodo.org/record/7031975#.Ywys8cjfmfU (Li et al., 2022).
- 557 Author contributions
- 558 LR wrote the manuscript. LR, PM, ZWD, and HJM contributed to the conceptualization of the study.
- 559 LR, GYN, CYB, and PM conducted the research, and visualized the results. WGH revised the
- 560 manuscript.
- 561 Competing interests
- 562 The contact authors have declared that neither they nor their co-authors have any competing
- 563 interests.

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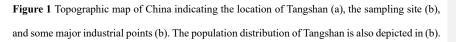
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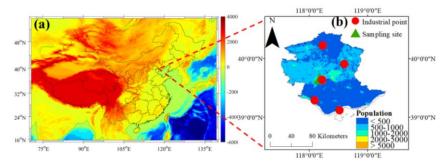


Figure 2 Bar chart of the concentrations of 16 trace elements including K, Fe, Ca, Zn, Pb, Mn, Fe, As, Se, V, Cr, Ag, Ni, Hg, Ga, and Co. The bars and black lines represent mean values and associated standard deviations, respectively.

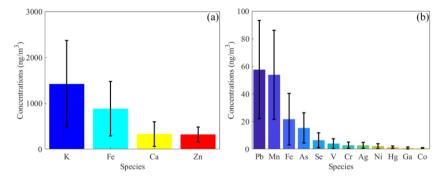


Figure 3 Inter-annual variations of the original (red) and deweathered (manganese purple) element concentrations (ng/m³) in PM_{2.5} in Tangshan. The dark, nattier blue, and nattier yellow backgrounds represent the species during 2017-2018 (from September in 2017 to August in 2018), 2018-2019 (from September in 2018 to August in 2019), and 2019-2020 (from September in 2019 to August in 2020). The bars and black lines represent mean values and associated standard deviations, respectively.

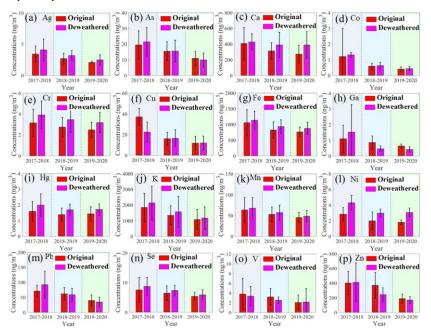


Figure 4 Original (red and orange) and deweathered (green and blue) element concentrations (ng/m³) in PM_{2.5} in Tangshan in four seasons during 2017-2018, 2018-2019, and 2019-2020. S1, U1, A1, and W1 represent the spring, summer, autumn, and winter during 2017-2018. S2, U2, A2, and W2 denote the spring, summer, autumn, and winter during 2018-2019. S3, U3, A3, and W3 are the spring, summer, autumn, and winter during 2019-2020. The points and shaded areas represent mean values and associated standard deviations, respectively.

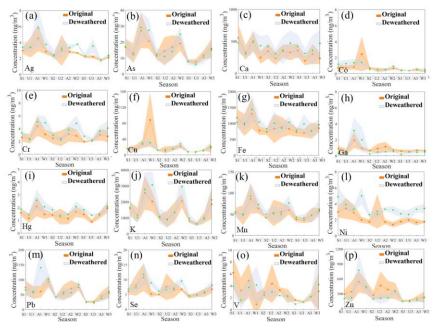
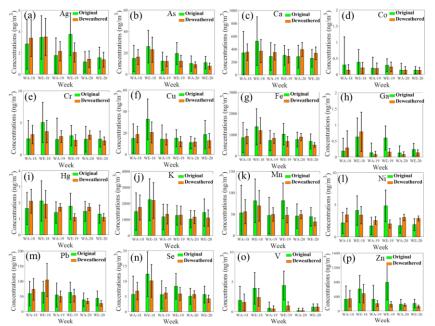
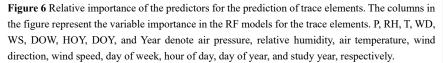


Figure 5 Weekly variations of the original (green) and deweathered (orange) element concentrations (ng/m³) in PM_{2.5} in Tangshan. The green and dark backgrounds denote the error bars of the original and deweathered elements, respectively. The bars and black lines represent mean values and associated standard deviations, respectively.





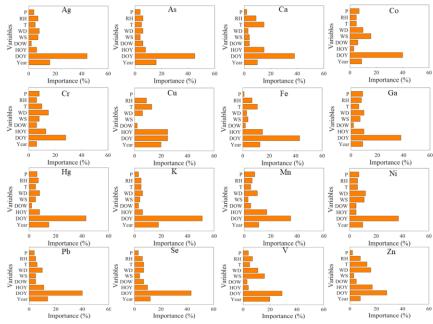
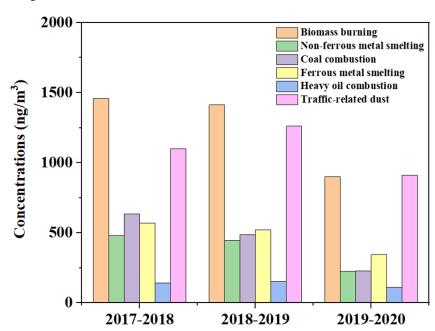
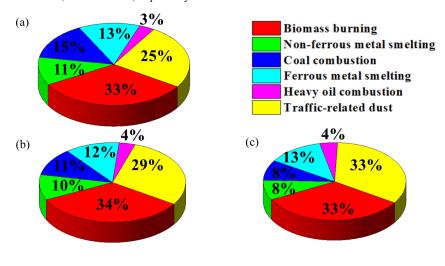


Figure 7 Deweathered mean concentrations of trace elements derived from six sources in Tangshan



during 2017-2020.

Figure 8 Average contributions of the six sources to the deweathered concentrations of the elements in PM_{2.5} based on the PMF model. The red panel means the biomass burning; the green panel denotes the non-ferrous metal smelting; the blue one represents the coal combustion; the cyan one is the ferrous metal smelting; the pink one represents the heavy oil combustion; and the yellow one denotes the traffic-related dust. (a), (b), and (c) represent the source contributions during 2017-2018, 2018-2019, and 2019-2020, respectively.



<u> </u>					
Elements	Mean±SD	CAAQS	WHO	EU	United States
Со	0.7±0.2				
Ga	0.9±0.7				
Hg	$1.47 {\pm} 0.8$	50	1000		
Ni	2.2±1.8		25	20	
Ag	2.8±2.1				
Cr	2.8±2.2	0.03	0.3		
v	4.0±3.6				
Se	6.5±5.3				
As	15.3±11.0	6	6.6	6	
Cu	22±19				
Mn	54±32				
РЬ	58±36		1000		150
Zn	320±160				
Ca	330±270				
Fe	880 ± 590				
К	1420±950				

Table 1 Comparison of the element concentrations in $PM_{2.5}$ of Tangshan and the standard values forthese elements in World Health Organization (WHO), China, Europe, and the United States (Unit:

		0		0			0			2.0	
Age	Year	Indicator	Cr	Mn	Fe	Co	Ni	Cu	Zn	As	РЬ
Adult	2017-2018	HQ	2.5×10 ⁻⁴	1.07×10 ⁻⁴	3.6×10 ⁻⁴	4.5×10 ⁻⁴	0.3×10 ⁻⁴	1.18×10 ⁻⁴	3.2×10 ⁻⁴	1.53×	8.4×10 ⁻⁴
										10-2	
		CR	0.1×10 ⁻⁶				-	-	-	2.4×10 ⁻⁶	2.9×10 ⁻⁶
	2018-2019	HQ	2.2×10 ⁻⁴	0.9×10 ⁻⁴	2.8×10 ⁻⁴	4.7×10 ⁻⁴	0.2×10 ⁻⁴	1.00×10 ⁻⁴	2.9×10 ⁻⁴	1.21×	7.3×10 ⁻⁴
										10-2	
		CR	0.1×10 ⁻⁶				-			1.87×	2.5×10 ⁻⁶
										10-6	
	2019-2020	HQ	2.0×10 ⁻⁴	0.8×10 ⁻⁴	2.6×10 ⁻⁴	3.3×10 ⁻⁴	0.2×10 ⁻⁴	0.7×10 ⁻⁴	1.49×10 ⁻⁴	0.9×10 ⁻²	4.7×10 ⁻⁴
		CR	0.1×10 ⁻⁶				-	-	-	1.33×	1.61×
										10-6	10-6
Child	2017-2018	HQ	6.0×10 ⁻⁴	2.6×10 ⁻⁴	8.6×10 ⁻⁴	1.11×10 ⁻³	0.8×10 ⁻⁴	2.9×10 ⁻⁴	7.7×10 ⁻⁴	3.7×10 ⁻²	2.0×10 ⁻³
		CR	0. 1×10 ⁻⁶				-			1.44×	1.75×
										10-6	10-6
	2018-2019	HQ	5.3×10 ⁻⁴	2.2×10 ⁻⁴	6.7×10 ⁻⁴	1.10×10 ⁻³	0.6×10 ⁻⁴	2.3×10 ⁻⁴	7.1×10 ⁻⁴	3.0×10 ⁻²	1.77×
											10 ⁻³
		CR	0.1×10 ⁻⁶				-	-	-	1.14×	1.52×
										10-6	10-6
	2019-2020	HQ	4.8×10 ⁻⁴	1.85×10 ⁻⁴	6.3×10 ⁻⁴	7.9×10 ⁻⁴	0.5×10 ⁻⁴	1.70×10 ⁻⁴	3.6×10 ⁻⁴	2.1×10 ⁻²	1.14×
											10 ⁻³
		CR	0.1×10^{-6}				-	-	-	0.8×10^{-6}	1.0×10 ⁻⁶

Table 2 Non-carcinogenic and carcinogenic risks for the original element levels in $\ensuremath{\text{PM}_{2.5.}}$

Age	Year	Indicator	Cr	Mn	Fe	Co	Ni	Cu	Zn	As	Pb
Adult	2017-2018	HQ	3.1×10 ⁻⁴	1.14×10 ⁻⁴	3.8×10 ⁻⁴	1.03×10 ⁻³	0.5×10 ⁻⁴	1.33×10 ⁻⁴	3.2×10 ⁻⁴	1.68×	1.08×
										10-2	10-3
		CR	0.2×10 ⁻⁶				-	-	-	2.6×10 ⁻⁶	3.7×10 ⁻⁶
	2018-2019	HQ	2.7×10 ⁻⁴	1.00×10 ⁻⁴	3.1×10 ⁻⁴	4.9×10 ⁻⁴	0.4×10 ⁻⁴	1.00×10 ⁻⁴	2.9×10 ⁻⁴	1.22×	6.9×10 ⁻⁴
										10-2	
		CR	0.1×10 ⁻⁶				-			1.89×	2.4×10 ⁻⁶
										10-6	
	2019-2020	HQ	2.5×10 ⁻⁴	0.8×10 ⁻⁴	3.0×10 ⁻⁴	3.5×10 ⁻⁴	0.4×10 ⁻⁴	0.7×10 ⁻⁴	1.49×10 ⁻⁴	0.8×10 ⁻²	4.1×10 ⁻⁴
		CR	0.1×10 ⁻⁶				-			1.20×	1.40×
										10-6	10-6
Child	2017-2018	HQ	7.5×10 ⁻⁴	2.8×10 ⁻⁴	9.3×10 ⁻⁴	1.17×10 ⁻³	1.23×10 ⁻⁴	3.2×10 ⁻⁴	3.2×10 ⁻⁴	4.1×10 ⁻²	2.6×10 ⁻³
		CR	0. 1×10 ⁻⁶				-			1.58×	2.3×10 ⁻⁶
										10-6	
	2018-2019	HQ	6.6×10 ⁻⁴	2.3×10 ⁻⁴	7.6×10 ⁻⁴	1.19×10 ⁻³	0.9×10 ⁻⁴	2.4×10 ⁻⁴	1.91×10 ⁻⁴	3.0×10 ⁻²	1.68×
											10-3
		CR	0.1×10 ⁻⁶				-			1.15×	1.44×
										10-6	10-6
	2019-2020	HQ	6.1×10 ⁻⁴	1.99×10 ⁻⁴	7.2×10 ⁻⁴	8.4×10 ⁻⁴	0.9×10 ⁻⁴	1.71×10 ⁻⁴	1.34×10 ⁻⁴	1.89×	9.9×10 ⁻⁴
										10-2	

 $\textbf{Table 3} \text{ Non-carcinogenic and carcinogenic risks for the deweathered element levels in PM_{2.5}.}$