

# Polycyclic aromatic hydrocarbons in atmospheric dust deposition from Anqing, China: pollution characteristics, sources and health risk assessment

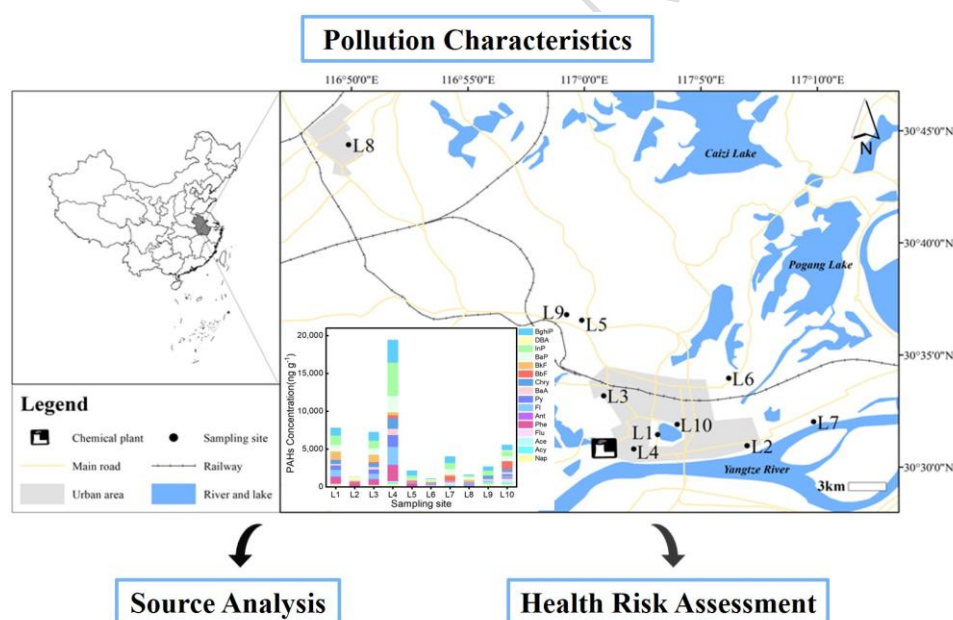
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## Graphical abstract



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## Abstract

This study systematically analyzed the pollution characteristics, source apportionment, and health risks of 16 polycyclic aromatic hydrocarbons (PAHs) in atmospheric dust deposition, using Anqing City—a typical industrial city in China—as the research subject. Results indicate that the concentration range of  $\Sigma_{16}\text{PAHs}$  in atmospheric dust in Anqing City is 85.22 – 21,351.03 ng g<sup>-1</sup>, with

an average of 5,301.21 ng g<sup>-1</sup>, placing the pollution level in the lower-middle range nationally. Spatial distribution revealed significantly higher PAH concentrations in industrial clusters of Daguan District compared to other areas, indicating industrial emissions as a major point source of PAHs. The composition was dominated by high-ring (5–6 ring) PAHs (average proportion: 58.90%), indicating primary origin from high-temperature combustion processes. Integrated source apportionment using ring number distribution, characteristic ratio analysis, and positive matrix factorization (PMF) models identified fossil fuel combustion and industrial activities as the primary PAH sources. Health risk assessments revealed skin contact as the predominant exposure pathway for both adults and children, with carcinogenic risks approaching or exceeding acceptable thresholds under certain high-exposure scenarios.

**Keywords:** Polycyclic aromatic hydrocarbons; Atmospheric dust deposition; Health risk assessment; Anqing

## 1. Introduction

The presence of environmental pollutants represents a significant threat to human health. Among these pollutants, polycyclic aromatic hydrocarbons (PAHs) are of significant concern due to their toxicity, carcinogenicity, and teratogenicity (Barbosa Jr et al., 2023). PAHs constitute a diverse group of organic compounds found extensively in environmental matrices like road dust, soil, and sediments (Ma et al., 2017). In light of the pervasive presence of these compounds in the environment and their potential implications for human and ecological health, the United States Environmental Protection Agency (USEPA) has identified 16 PAHs as priority pollutants (Samburova et al., 2017).

Globally, a significant quantity of PAHs are released into the atmosphere on an annual basis, and since PAHs are mostly products of incomplete combustion, most of them primarily manifest as particulate matter, which will migrate and diffuse with the atmosphere, and finally land on the ground in the form of dust (Yang et al., 2021). In addition to direct harm to the human body, atmospheric dust

fall pollution will also produce some damage to humans and the environment through various environmental media(Kothiyal et al., 2022).

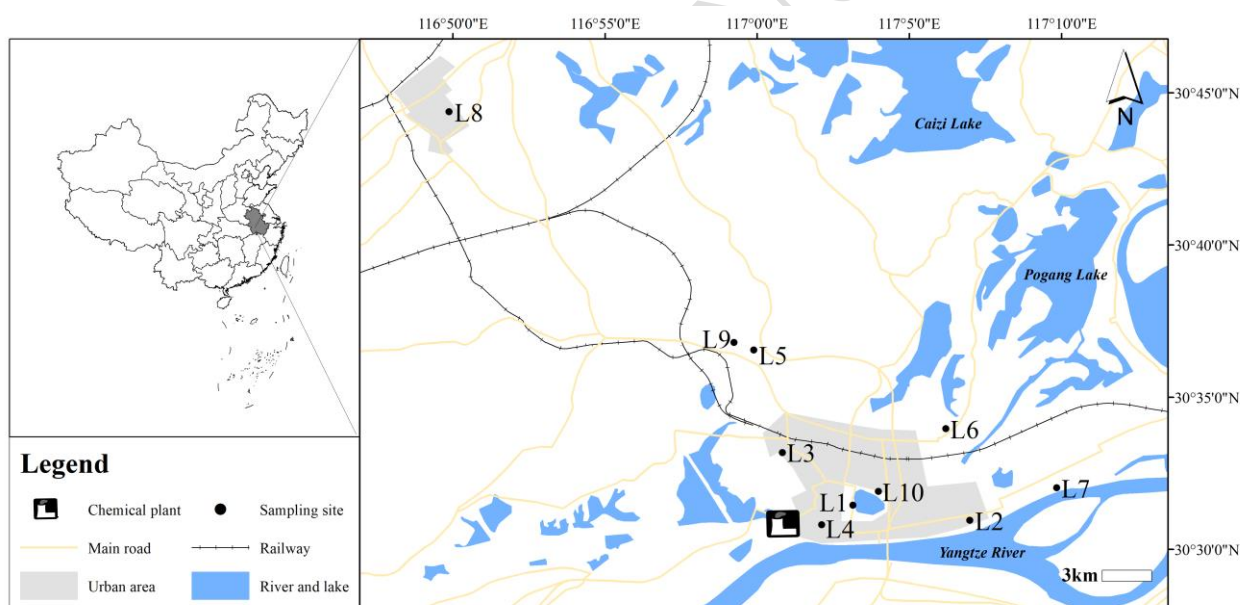
In recent years, people have gradually realised that PAHs contamination will constitute a great threat to public health, so the research on PAHs has become a hot spot. The research on PAHs in foreign countries started earlier, but the research on PAHs in China started later, and the cities that have carried out the research on PAHs in urban air are Shenzhen, Guangzhou, Tianjin, Xi'an and Beijing and other large cities(Fu et al., 2023, Jiang et al., 2023, Wu et al., 2022, Zhang et al., 2025, Li et al., 2022). Anqing, as a representative medium-sized industrial city in the Yangtze River Economic Belt, hosts a dense cluster of chemical, thermal power, steel, and building material industries(Sun et al., 2022). These sectors are characterized by intensive fossil fuel consumption and high-temperature processes, which are significant PAHs emission sources. Despite the proliferation of PAHs studies in megacities, systematic investigations in medium-sized industrial cities—particularly those with mixed industrial profiles like Anqing—remain scarce. This study aims to fill this gap by providing a comprehensive assessment of PAHs pollution characteristics, sources, and health risks in Anqing, thereby offering a scientific basis for targeted pollution control in similar urban contexts.

In this study, atmospheric dust deposition samples were collected from 10 different areas in Anqing City. The samples were extracted and concentrated using ultrasonic extraction before being quantitatively analysed for PAHs using gas chromatography-mass spectrometry (GC-MS). The primary objectives of this paper are to: (1) reveal the concentration and compositional characteristics of PAHs in atmospheric dust deposition in Anqing City; (2) identify the main origins of PAHs in atmospheric dust deposition; (3) assess the human health hazards associated with PAHs in atmospheric dust deposition.

## **2. Materials and Methods**

### **2.1. Study area and sampling**

In the last few years, the Anqing area of China has been designated a hub for chemical-based industries, boasting over 400 chemical companies, including Anqing Petrochemical and numerous others(Sun et al., 2022). With the concomitant increase in population and rapid economic growth, the issue of environmental pollution has become increasingly salient. In this study, 10 dust samples were gathered from diverse sites in Anqing City, with the sampling points distributed across the primary urban areas of Anqing City and Huaining County (Figure 1), covering industrial, commercial, residential, and suburban areas. During sampling, approximately 5 grams of dust is collected from outdoor window sills, etc., using a brush impregnated with acetone solution according to a strict procedure, and placed in a polythene sample bag labelled with the appropriate collection site. Subsequent to collection, the samples are returned to the laboratory to remove impurities, sieved through a 0.15 mm screen and stored at low temperatures.



**Figure 1.** Schematic diagram of atmospheric dust deposition sampling points

## 2.2. Analysis of PAHs in atmospheric dust deposition

The study analyzed the sixteen priority PAHs as identified by the USEPA, which include naphthalene (Nap), acenaphthylene (Acy), acenaphthene (Ace), fluoranthene (Fl), phenanthrene (Phe), anthracene (Ant), fluorene (Flu), pyrene (Py), benzo[a]anthracene (BaA), chrysene (Chry), benzo[a]pyrene (BaP), benzo[b]fluoranthene (BbF), benzo[k]fluoranthene (BkF), indeno[1,2,3-

86 cd]pyrene (InP), dibenzo[a,h]anthracene (DBA), and benzo[g,h,i]perylene (BghiP). Ultrasonic  
 87 extraction method (USA EPA 3550B) was used. About 2 g of sediment was weighed, 2 g of copper  
 88 powder was added, and the extract was extracted by ultrasonication with 15 mL of 1:1 (V:V) n-  
 89 hexane/acetone solution for 30 min, centrifuged and transfer the supernatant, and the above  
 90 ultrasonication was repeated for 2 times and the temperature was controlled to be below 40 °C. The  
 91 extracts were combined, rotary evaporated to 2 mL, and the extracts were passed through a  
 92 chromatography column (1 cm anhydrous sodium sulfate, 6 cm 3% deactivated alumina, and 12 cm  
 93 3% deactivated silica gel from top to bottom). The column was pre-eluted with 15 mL of n-hexane,  
 94 the hexane eluent was discarded, and then eluted with 70 mL of a (3:7 dichloromethane/n-hexane)  
 95 mixture, all of which was received. The received solution was rotary evaporated and fixed to 1.0 mL  
 96 with n-hexane, and stored at low temperature for measurement. The analysis of PAHs was conducted  
 97 utilizing a GC-MS (Agilent 5975C/7890A) with ionization in electron impact (EI) mode, and data  
 98 acquisition took place under the selective ion monitoring (SIM) mode.

### 99 2.3. Health risk assessment

100 The toxicity of PAHs present in atmospheric dust deposition is determined by means of the toxicity  
 101 equivalency factor (TEF) of these compounds(Zhang et al., 2022). The most toxic PAH-BaP had a  
 102 TEF value of 1, and the TEF values for the other PAHs were derived by comparing their  
 103 carcinogenicity levels with those of BaP(Škrbić et al., 2019). CS denotes the total concentration of  
 104 PAH congeners calculated using the TEF method (Equation (1))(Wang et al., 2011). Each PAH  
 105 congener's concentration is denoted by  $C_n$  ( $m^3 \text{ kg}^{-1}$ ), and the corresponding TEF for each PAH is  
 106 denoted by  $TEF_n$  (Table 2).

$$107 \quad CS = \sum (C_n \times TEF_n) \quad (1)$$

108 The incremental lifetime cancer risk (ILCR) associated with exposure to PAHs was determined  
 109 based on previous research(Wu et al., 2020). The ILCRs for dermal contact, inhalation, and ingestion  
 110 were quantified for each area dust sample in Anqing according to the following Equation (2) - (5):

$$111 \quad \text{ILCR}_{\text{Ingestion}} = \frac{\text{CS} \times (\text{CSF}_{\text{Ingestion}} \times \sqrt[3]{\text{BW}/70}) \times \text{IR}_{\text{Ingestion}} \times \text{EF} \times \text{ED}}{\text{BW} \times \text{AT} \times 10^6} \quad (2)$$

$$112 \quad \text{ILCR}_{\text{Inhalation}} = \frac{\text{CS} \times (\text{CSF}_{\text{Inhalation}} \times \sqrt[3]{\text{BW}/70}) \times \text{IR}_{\text{Inhalation}} \times \text{EF} \times \text{ED}}{\text{BW} \times \text{AT} \times \text{PEF}} \quad (3)$$

$$113 \quad \text{ILCR}_{\text{Dermal}} = \frac{\text{CS} \times (\text{CSF}_{\text{Dermal}} \times \sqrt[3]{\text{BW}/70}) \times \text{SA} \times \text{AF} \times \text{ABS} \times \text{EF} \times \text{ED}}{\text{BW} \times \text{AT} \times 10^6} \quad (4)$$

$$114 \quad \text{CR} = \text{ILCR}_{\text{Ingestion}} + \text{ILCR}_{\text{Inhalation}} + \text{ILCR}_{\text{Dermal}} \quad (5)$$

115 To minimise computational uncertainty, a Monte Carlo simulation (Oracle Crystal Ball, USA) was  
 116 employed. This simulation was based on parameters from previous researchs (Kong et al., 2025,  
 117 Tarafdar and Sinha, 2019)(Table 1) research and comprised 20,000 iterations.

118 **Table 1.** Exposure parameters for probabilistic risk assessment.

Exposure variable	Distribution types <sup>a</sup>	Adult	Child	Unit
Averaging life span (AT)	Point	70×365=25,550	70×365=25,550	day
Body weight (BW)	Normal	[52.10, 6.50]	[16.30, 2.40]	kg
Dermal exposure area (SA)	Lognormal	[4,619.11, 1.77]	[4,619.11, 1.77]	cm <sup>2</sup>
Dermal adherence factor (AF)	Lognormal	[0.04, 3.41]	[0.04, 3.41]	mg cm <sup>-2</sup>
Dermal adsorption fraction (ABS)	Lognormal	[0.13, 1.26]	[0.13, 1.26]	unitless
Exposure duration (ED)	Uniform	[0, 24]	[0, 6]	year
Exposure frequency (EF)	Triangular	345 [180, 365]	345 [180, 365]	day year <sup>-1</sup>
Ingestion rate (IR <sub>ingestion</sub> )	Point	100	200	mg day <sup>-1</sup>
Inhalation rate (IR <sub>inhalation</sub> )	Lognormal	[13.9, 1.07]	[7.19, 1.62]	m <sup>3</sup> day <sup>-1</sup>
Particle emission factor (PEF)	Point	1.36×10 <sup>9</sup>	1.36×10 <sup>9</sup>	m <sup>3</sup> kg <sup>-1</sup>
Ingestion carcinogenic slope factor (CSF <sub>ingestion</sub> )	Lognormal	[7.30, 1.15]	[7.30, 1.15]	mg kg <sup>-1</sup> day <sup>-1</sup>
Inhalation carcinogenic slope factor (CSF <sub>inhalation</sub> )	Lognormal	[3.14, 1.80]	[3.14, 1.80]	mg kg <sup>-1</sup> day <sup>-1</sup>
Dermal carcinogenic slope factor (CSF <sub>dermal</sub> )	Point	25	25	mg kg <sup>-1</sup> day <sup>-1</sup>

119 <sup>a</sup> For normal, point, uniform and lognormal distributions, the values in parentheses represent the  
120 arithmetic mean and standard deviation, the fixed values, the minimum and maximum, and the  
121 geometric mean and geometric standard deviation, respectively.

## 122 2.4. Positive matrix factorization (PMF)

123 Source apportionment was performed using the US EPA PMF 5.0 model. The model decomposes  
124 the original data matrix  $X$  ( $n$  samples  $\times$   $m$  species) into factor contributions ( $G$ ) and factor profiles  
125 ( $F$ ), minimizing the objective function  $Q$ :

$$126 \quad X = G \times F + E \quad (6)$$

$$127 \quad Q = \sum_{i=1}^n \sum_{j=1}^m \left( \frac{e_{ij}}{u_{ij}} \right)^2 \quad (7)$$

128 where  $E$  is the residual matrix and  $u_{ij}$  is the uncertainty of species  $j$  in sample  $i$ .

129 Uncertainties ( $u_{ij}$ ) were calculated following (Guo et al., 2025b):

$$130 \quad u_{ij} = \begin{cases} 0.2 \times C_{ij} + \frac{MDL}{3}, & \text{if } C_{ij} \leq MDL \\ 0.1 \times C_{ij} + \frac{MDL}{3}, & \text{if } C_{ij} > MDL \end{cases} \quad (8)$$

131 where  $C_{ij}$  is the concentration and MDL is the method detection limit.

132 Model robustness was evaluated using multiple criteria. The optimal number of factors (three) was  
133 determined by examining the change in  $Q$  ( $Q_{\text{robust}}/Q_{\text{true}}$ ) with increasing factor numbers, ensuring  
134 physical interpretability of profiles. Bootstrap analysis (100 runs) was performed to assess factor  
135 stability and uncertainty. Key goodness-of-fit metrics, including the coefficient of determination ( $R^2$ )  
136 between observed and modeled concentrations for key species, are reported to validate model  
137 performance.

## 138 2.5. Quality assurance/quality control

139 To ensure the representativeness, accuracy, and comparability of the data, stringent quality control  
140 measures were implemented. Results for reagent blanks, laboratory blanks, and recovery experiments  
141 fell within acceptable ranges. Each PAH working standard curve exhibited good linearity with  
142 correlation coefficients ( $R^2$ )  $\geq 0.995$ . No PAHs were detected in reagent blanks. Laboratory blanks  
143 showed only trace amounts of Nap and Phe (below 3% of actual sample levels), with final  
144 concentrations adjusted by corresponding blank subtraction. The method's minimum detection limits  
145 ranged from 0.20 to 1.20 ng g<sup>-1</sup>. Except for the relatively low recovery rate of highly volatile



146 naphthalene (65.8%), recovery rates for other compounds ranged from 76 to 110%, with relative  
147 standard deviations < 8.0%. PAH concentrations in the measured dust were not corrected for recovery.

### 148 **3. Results and Discussion**

#### 149 3.1. Distribution of PAHs content in dust

150 In this study, the contents of PAHs in dust samples from 10 sampling sites in urban areas of Anqing  
151 were counted. Table 2 lists the TEF, mean, median and range of 16 PAHs in the dust samples from  
152 the 10 sampling points.

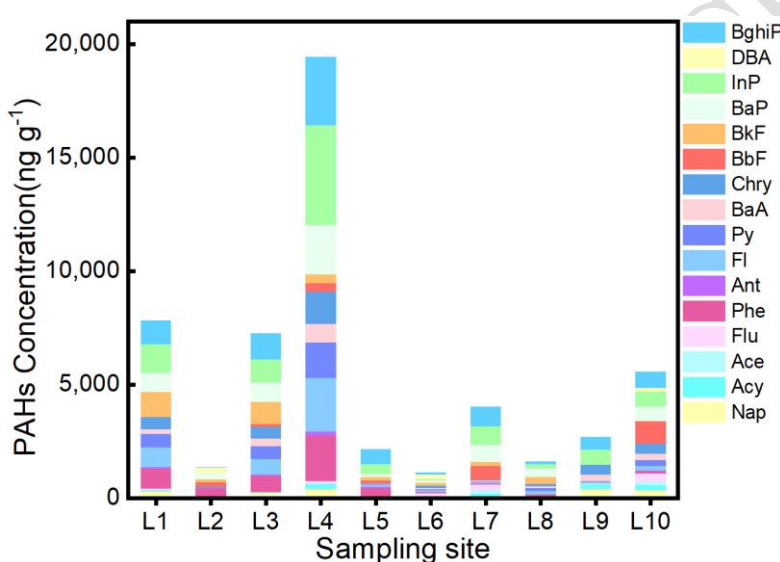
153 **Table 2.** Distribution of PAHs in dust samples from different areas of Anqing city.

Congeners	TEF	Mean( $\text{ng g}^{-1}$ )	Median( $\text{ng g}^{-1}$ )	Concentration( $\text{ng g}^{-1}$ )
Nap	0.001	175.016	127.96	0 - 409.51
Acy	0.001	98.229	42.12	0 - 232.7
Ace	0.001	46.33	19.475	0 - 157.06
Flu	0.001	93.833	28.16	0 - 365.27
Phe	0.001	482.271	273.635	17.11 – 2,005.35
Ant	0.01	38.783	14.335	1.75 - 194.64
Fl	0.001	453.363	116.385	1.64 – 2,343.66
Py	0.001	323.667	80.505	0.96 – 1,567.34
BaA	0.1	211.601	138.275	15.57 - 815.77
Chry	0.01	365.861	266.145	28.29 – 1,408.7
BbF	0.1	245.095	122.935	0 - 998.03
BkF	0.1	334.646	170.54	0 – 1,077.93
BaP	1	616.714	525.87	0 – 2,177.43
InP	0.1	943.586	652.815	7.49 – 4,378.85
DBA	1	59.467	0	0 - 226.64
BghiP	0.01	812.849	679.085	12.41 – 2,992.15
$\Sigma_{16}\text{PAHs}$	—	5,301.21	3,258.29	85.22 – 21,351.03

154 As shown in Table 2, the concentration of  $\Sigma_{16}\text{PAHs}$  ranged from 85.22 - 21,351.03  $\text{ng g}^{-1}$ , with a  
155 mean concentration of 5301.21  $\text{ng g}^{-1}$  and a median concentration of 3,258.29  $\text{ng g}^{-1}$ ; of these, InP  
156 had the highest concentration, with a mean concentration of 943.59  $\text{ng g}^{-1}$ , followed by BghiP and  
157 BaP; Ant had the lowest concentration with an average of 38.79  $\text{ng g}^{-1}$ , while Ace and DBA also had  
158 relatively low concentrations, detected in only 3 sampling sites. The minimum concentrations of the  
159 2-ring PAHs (Nap, Acy, Ace, Flu) were all 0. This may be due to the fact that most of the 2-ring

160 PAHs are mainly present in the gas-phase and are volatile, and therefore their content in the particles  
 161 may be below the detection limit.

162 Combined with Figure 2 we can clearly see the degree of contamination at each sampling point:  
 163  $L4 > L1 > L3 > L10 > L7 > L9 > L5 > L8 > L2 > L6$ . It can be seen that the most serious pollution in the urban  
 164 area of Anqing is Daguan District, Daguan District, there are Sinopec Anqing Petrochemical  
 165 Company, Anqing Changhong Chemical Co., Ltd, Anqing Yicheng Chemical Technology Co., Ltd,  
 166 Anqing Jingyi Fine Chemical Co., Ltd, and other chemical enterprises, and these chemical enterprises  
 167 are very likely to be an important reason for the more serious pollution of PAHs in this area.



**Figure 2.** Distribution of PAHs in dust samples from different areas of Anqing city

170 Compared with other cities and countries, the levels of PAHs in atmospheric dust deposition in  
 171 Anqing were significantly lower than those in Tianjin (average concentration of  $7,990 \text{ ng g}^{-1}$ )(Yu et  
 172 al., 2014), Canada (average concentration of  $24,700 \text{ ng g}^{-1}$ )(Gill et al., 2020), Laiwu (average  
 173 concentration of  $10,892 \text{ ng g}^{-1}$ )(Wei et al., 2021), and Shenzhen (average concentration of  $29,920 \text{ ng}$   
 174  $\text{g}^{-1}$ )(Liu et al., 2016). The difference with Hubei Province (average concentration of  $4,430 \text{ ng g}^{-1}$ ) is  
 175 small(Zhang et al., 2016). It is not difficult to see that the average level of PAHs in atmospheric dust  
 176 deposition in Anqing is lower than that in some large cities in China, which may be due to the high  
 177 population density, motor vehicle emissions and fossil energy demand in large cities, and thus the

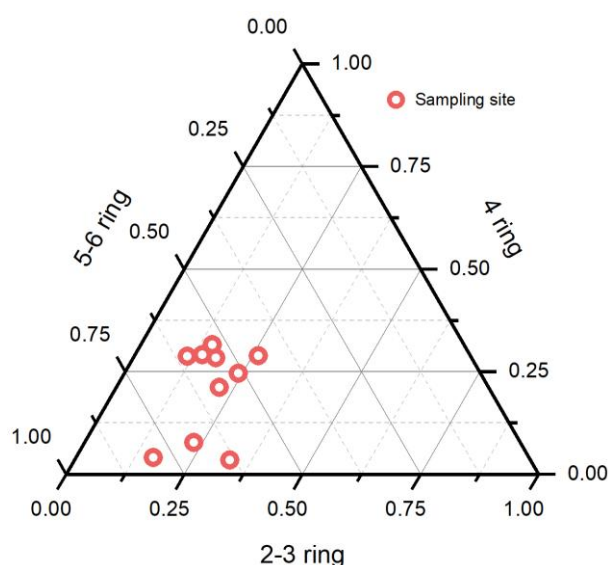
178 PAHs content in dust in large cities is much higher than that in small and medium-sized cities like  
179 Anqing.

180 Although this study's sample size is sufficient for a preliminary spatial assessment, it may limit the  
181 statistical robustness and spatial representativeness of the findings, particularly in heterogeneous  
182 urban environments. This limitation has been identified as a constraint of the study, and caution  
183 should be exercised when interpreting the results in relation to city-wide extrapolation. It should be  
184 noted that this study did not account for seasonal variations or meteorological influences, which may  
185 affect the deposition patterns and source contributions of PAHs. Future research should incorporate  
186 multi-seasonal sampling and atmospheric dispersion modelling to improve temporal  
187 representativeness.

### 188 3.2. Source analysis of PAHs in dust

#### 189 3.2.1. Comparison of ring numbers of PAHs in dust

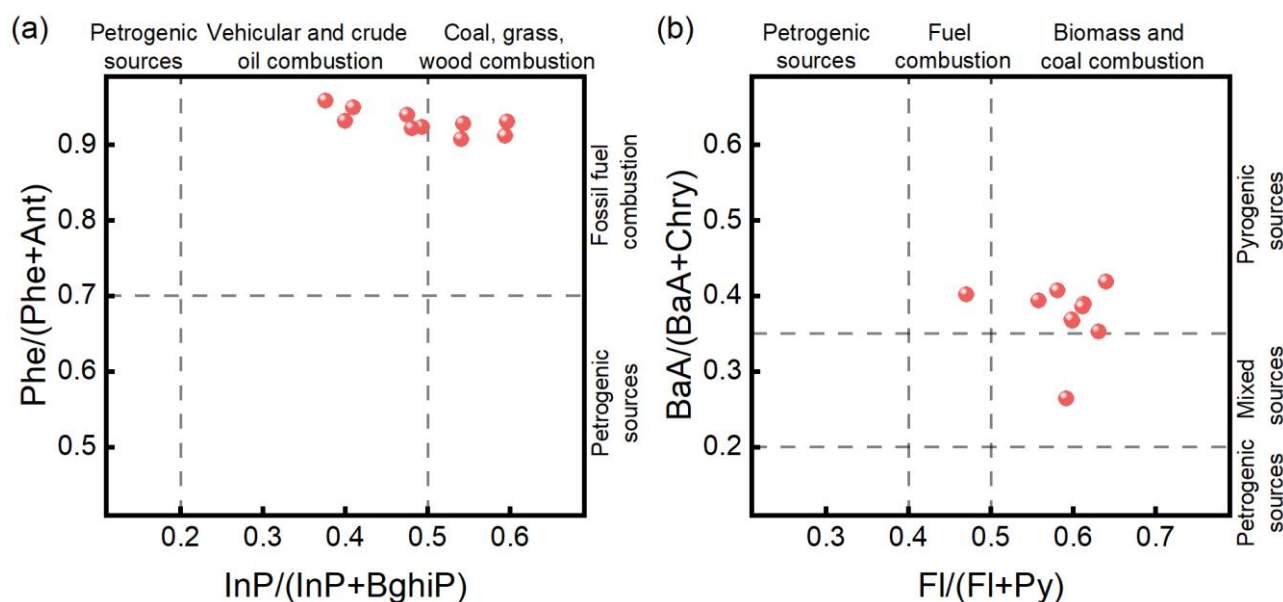
190 The distribution of PAHs with different ring numbers in the 10 dust samples collected in this  
191 experiment is shown in Figure 3, and the PAHs in the studied area are mainly 5-ring and 4-ring,  
192 followed by 3-ring. The other 16 PAHs can be divided into three groups according to their ring  
193 numbers: low molecular weight (2-3 ring, LMW), middle molecular weight (4 ring, MMW) and high  
194 molecular weight (5-6 rings, HMW) to discriminate their different sources of contamination. In  
195 general, LMW PAHs primarily originate from crude oil contamination and the combustion of  
196 materials such as wood and coal at temperatures below moderate levels(Zhang et al., 2020). MWM  
197 PAHs mainly stem from fuel combustion, while HWM PAHs primarily result from the high-  
198 temperature combustion of fossil fuels. The proportion of LMW PAHs in the dust samples in the  
199 studied area ranged from 11.30 - 32.95%, with a mean value of 20.32%, the proportion of MMW  
200 PAHs ranged from 3.43 - 31.59%, with a mean value of 20.78%, and the proportion of HMW PAHs  
201 ranged from 44.83 - 79.42%, with a mean value of 58.90%. This indicates that PAHs in atmospheric  
202 dust deposition in Anqing City primarily originate from high-temperature combustion processes, such  
203 as coal combustion and motor vehicle exhaust emissions.



**Figure 3.** Ternary plot showing comparative contribution of 2–3 ring, 4 ring and 5–6 ring PAHs in samples

### 3.2.2. Characteristic ratio method of PAHs in dust

In order to gain a more detailed understanding of the emission origins of PAHs in atmospheric dust deposition, a diagnostic comparison of PAHs was performed, the results of which are shown in Figure 4(Li et al., 2017). The majority of the samples exhibited  $Fl/(Fl + Py)$  ratios greater than 0.5, suggesting a mixture of fuel combustion sources(Han et al., 2021). The ratios of  $BaA/(BaA + Chry)$  and  $InP/(InP + BghiP)$  both exceeded 0.2, indicating that the primary origins of PAHs in atmospheric dust deposition were emissions from coal, wood, fuel combustion, and vehicular exhaust(Wang et al., 2022). The predominance of PAHs at most road atmospheric samples, as indicated by ratios of  $Phe/(Phe + Ant) > 0.7$ , was attributed to fossil fuel combustion(Zhang et al., 2024).



**Figure 4.** Characteristic ratios of PAHs in dust in different areas of Anqing City

In summary, the PAHs in atmospheric dust deposition in Anqing City primarily originate from a combination of pollution sources, including fossil fuel combustion, coal combustion, and motor vehicle emissions. The pollution source structure was similar across all sampling sites, with no significant spatial variation observed.

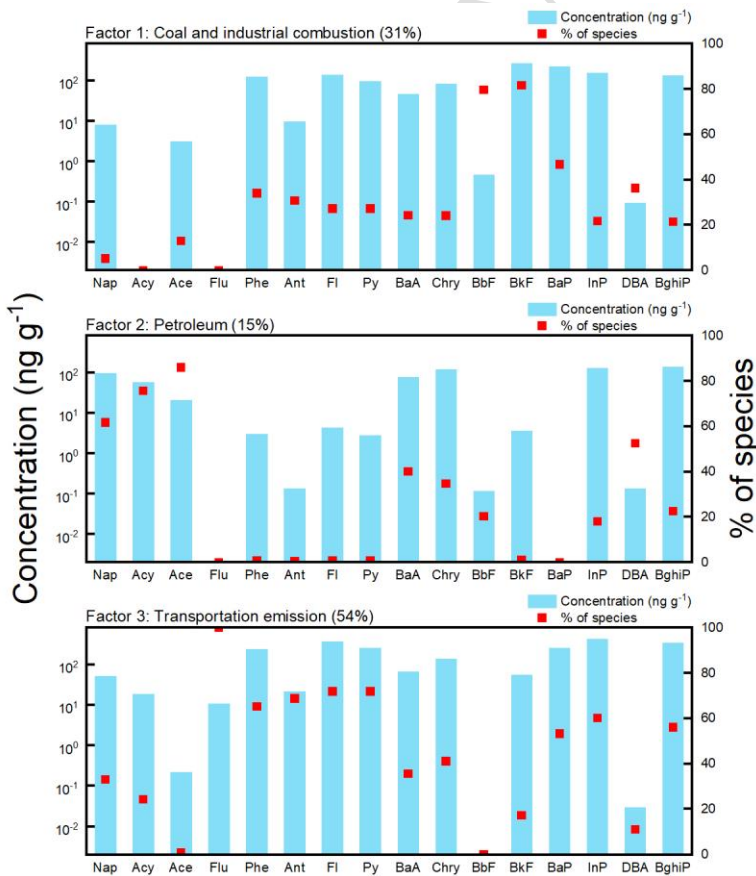
### 3.2.3. PMF of PAHs in dust

To identify the primary sources of PAHs in atmospheric dust deposition, this study employed a positive matrix factorization model for source apportionment of 16 PAHs. Based on model goodness-of-fit and factor independence, three factors were ultimately determined as the optimal solution. The loadings of PAHs across each factor and their contributions to total concentration are shown in Figure 5.

Factor 1 explained 30.66% of the total PAH concentration, primarily contributed by high-molecular-weight PAHs such as BkF (81.65%), BaP (46.77%), and InP (21.77%). These compounds are typically associated with high-temperature combustion processes, particularly coal combustion and industrial high-temperature operations (Wang et al., 2025). Additionally, medium-ring PAHs like Phe, Fla, and Pyr constitute a significant proportion within this factor, further supporting its combustion-derived characteristics. Thus, factor 1 is identified as originating from coal and industrial combustion sources.

Factor 2 contributed 15.49% of the total concentration, primarily loaded onto low-ring PAHs such as Acy (75.75%), Ace (86.07%), and BaA (40.11%). These compounds are frequently associated with incomplete combustion of biomass or volatilization of petroleum substances(Guo et al., 2025a). Notably, Acy and Ace are widely recognized as tracers of biomass combustion. Therefore, factor 2 is classified as a biomass and petroleum volatiles source.

Factor 3 is the largest contributing source, accounting for 53.85% of total PAH concentration. It primarily includes medium-to-low ring PAHs such as Flu (100%), Phe (65.20%), Fl (71.84%), and Py (71.94%). These compounds are commonly found in transportation emissions, particularly gasoline vehicle exhaust(Ma et al., 2025). Additionally, InP and BghiP constitute a significant proportion within this factor, further supporting its association with motor vehicle emissions. Consequently, factor 3 is identified as a transportation emission source.



**Figure 5.** Analysis of sources of PAHs based on PMF

The contribution order of the three sources to total PAHs was: traffic emissions (53.85%) > coal and industrial combustion (30.66%) > biomass and petroleum volatiles (15.49%). Coal and industrial

250 combustion contributed more than in Shanghai (14.7 %)(Feng et al., 2022) but less than in Anshan  
251 (32.6 %)(Wang et al., 2020). This source profile aligns with Anqing's status as an industrial city with  
252 growing vehicular activity. Comparing with other Chinese cities, the contribution of traffic sources  
253 here (53.85%) is higher than in some heavily industrial cities where coal combustion dominates,  
254 highlighting the need for targeted vehicular emission controls alongside industrial regulations.

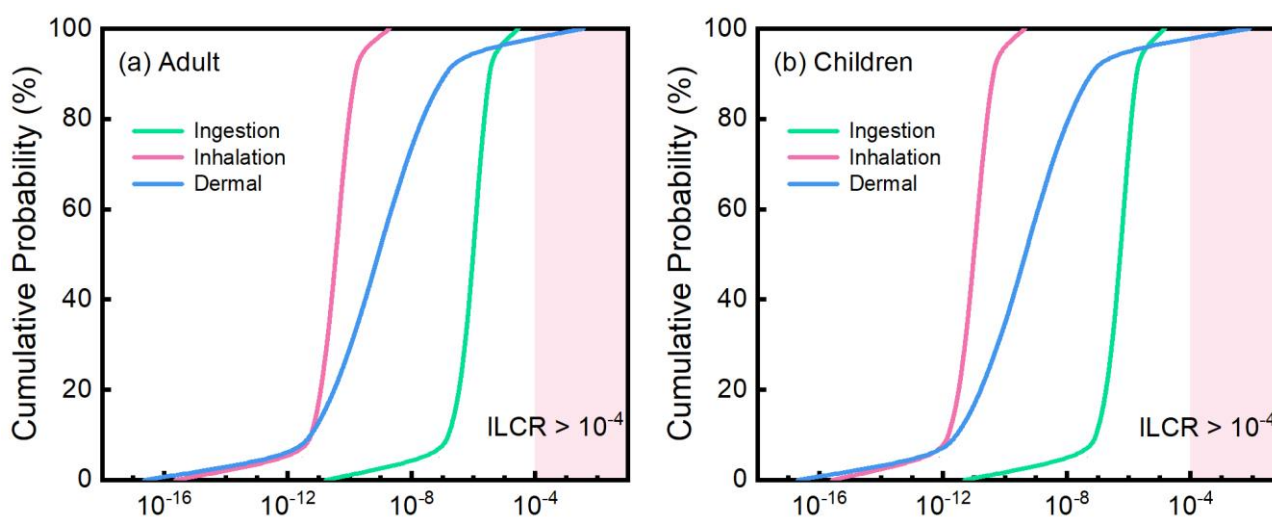
### 255 3.3. Health risk assessment of PAHs

256 This study employed Monte Carlo simulations to assess the ILCR of PAHs in atmospheric dust  
257 deposition for adults and children(Bharali et al., 2025). Overall, dermal contact and oral ingestion  
258 represent the predominant exposure pathways, while inhalation risks are negligible.

259 As shown in the ILCR probability distribution in Figure 6, both adults and children exhibit  
260 significantly right-skewed risk distributions for skin contact and oral ingestion, with pronounced  
261 high-risk “tails.” Notably, in the skin contact pathway, the rightmost tail of the distribution curve  
262 approaches or even exceeds  $1 \times 10^{-4}$ , indicating that skin contact may pose unacceptable carcinogenic  
263 risks under extreme exposure scenarios(Sun et al., 2025).

264 Although the median and mean ILCR values for skin contact are below  $10^{-4}$ , the high-end quantiles  
265 (e.g., 95th percentile or higher) suggest that for a portion of highly exposed individuals  
266 (approximately 5% or less), the carcinogenic risk via this route may have reached levels warranting  
267 intervention. This finding aligns with the high contribution of the AF in sensitivity analyses,  
268 indicating that individual behavioral variations and uncertainties in exposure levels are key drivers of  
269 the high-risk tail.

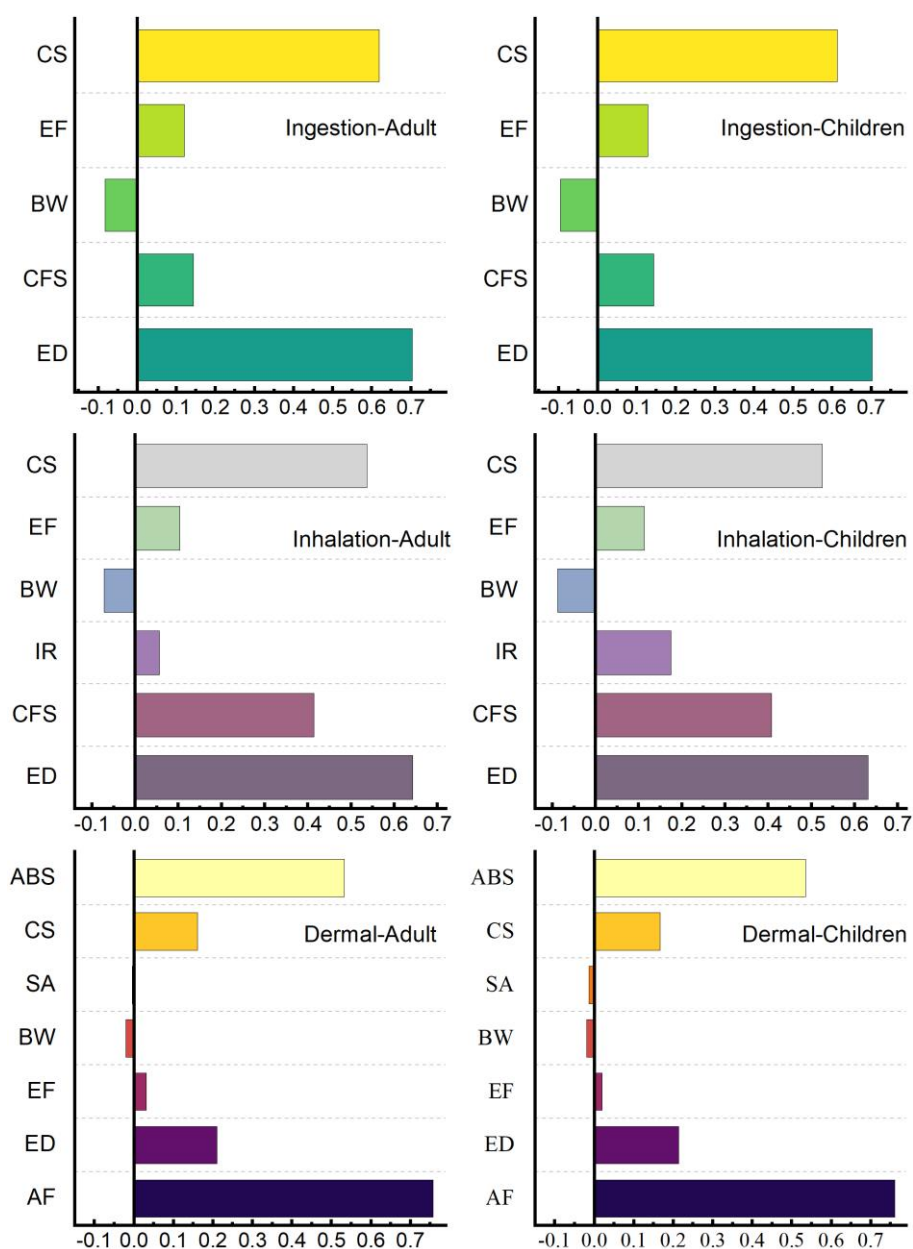




**Figure 6.** Probability distribution for carcinogenic risks of PAHs for (a) adults and (b) children

As shown in Figure 7, ED and BaP equivalent toxicity concentration (CS) are the most sensitive parameters across all exposure pathways. For the dermal exposure pathway, the AF is also a critical influencing factor (Khalili et al., 2021). This indicates that, in addition to controlling the toxic equivalent concentration of dust, reducing the frequency of skin contact with dust and the amount of dust adhering to the skin—particularly through protective measures in high-risk environments such as frequent ground activities and cleaning operations—is especially crucial for controlling the high tail risk associated with the dermal pathway.

In summary, although the average carcinogenic risk from PAHs in indoor dust is generally manageable, skin contact exposure under extreme conditions may pose unacceptable health risks ( $ILCR > 10^{-4}$ ). This finding underscores that risk assessment and management should prioritize high-risk populations and extreme exposure scenarios, beyond merely monitoring average risk levels. This underscores that, in addition to reducing environmental PAHs concentrations, behavioral and exposure mitigation measures—such as improved hygiene practices, use of protective gear for outdoor workers, and reducing children's direct contact with dust—are crucial for managing population risk, especially for vulnerable subgroups.



**Figure 7.** Sensitivity analysis of ILCR for PAHs

#### 4. Conclusions

This study systematically examined the pollution levels, composition profiles, sources, and health risks of 16 priority PAHs in atmospheric dust deposition across different functional zones in Anqing City, China. The concentration range of  $\Sigma_{16}\text{PAHs}$  in atmospheric dust in Anqing City was relatively wide (85.22 – 21,351.03 ng g<sup>-1</sup>), with an average concentration of 5,301.21 ng g<sup>-1</sup>, indicating a pollution level in the lower-middle range nationally. Spatial distribution revealed significantly higher concentrations in industrial clusters like Daguan District (L1, L3, L4) compared to other areas, indicating local industrial emissions as a major point source of PAHs. Composition analysis showed

high-ring (5–6 ring) PAHs dominated (average 58.90%), revealing that PAHs in the study area primarily originate from high-temperature combustion processes. Comprehensive source apportionment using ring number distribution, characteristic ratio method, and PMF models yielded mutually corroborating results, confirming that Anqing's PAH pollution stems from a hybrid contamination pattern involving fossil fuel combustion and industrial activities. Probabilistic health risk assessment was conducted using a Monte Carlo simulation-based ILCR model. Results indicate that for the general population of Anqing City, the overall carcinogenic risk from PAH exposure via particulate matter deposition is at an acceptable level. Skin contact is the primary exposure pathway, with its high-risk tail (e.g., 95th percentile) approaching or reaching  $10^{-4}$ , warranting attention. Children's total risk is slightly higher than adults', which is associated with their behavioral patterns (frequent hand-to-mouth contact).

## **5. Environmental implications**

The findings underscore that in industrial cities like Anqing, control strategies must adopt a multi-target approach. Prioritizing traffic emission reductions (e.g., upgrading vehicle fleets, promoting EVs) is essential, given its dominant contribution. Simultaneously, stringent controls on industrial and coal combustion emissions are needed, especially in identified hotspot areas like Dagan District. Public health measures should focus on reducing dermal exposure for high-risk groups (children, outdoor workers). This study establishes a critical baseline and a methodological framework (integrating PMF and probabilistic risk assessment) that can be applied to other industrial cities for evidence-based environmental management.

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## **Conflict of Interest**

The authors declare no conflict of interest.

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