

Source Apportionment of PM_{2.5} During Haze and Non-haze in Kuala Lumpur Urban Environment

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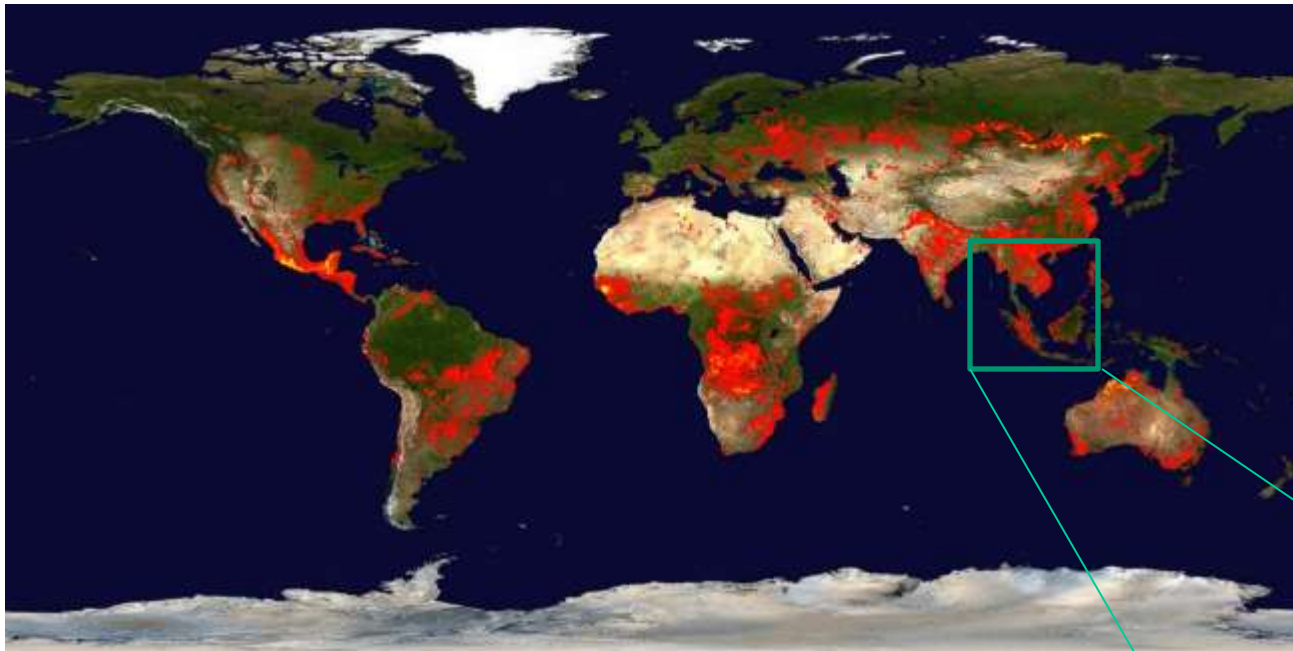
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Introduction

Biomass burning in Southeast Asia



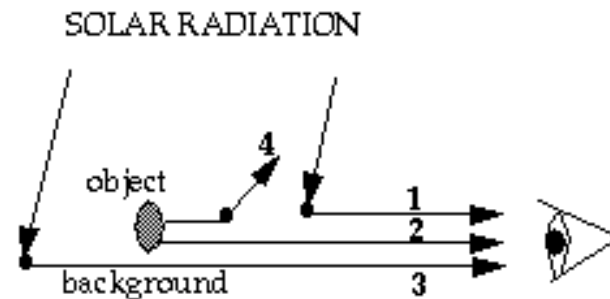
Southeast Asia

Ichoku and Kahn (2012)

Kuala Lumpur Hazy Condition



- $PM_{2.5} \geq 35 \mu\text{g m}^{-3}$
- Visibility $< 10 \text{ km}$
- Dry/Low humidity



Trigger to Biomass Burning

- In South East Asia biomass burning has become a traditional method of **clearing land** in the practice of shifting cultivation, which involves field rotation and the **slashing and burning** of a new plot of land once the existing plot has lost its fertility





Biomass Burning from Peat Soil Combustion





Haze: Bad new folks! Sumatra hotspots double to 118 on Saturday

JUNE 23, 2013 BY ADMIN [LEAVE A COMMENT](#)

Smouldering
High amount of
soot and smoke !!

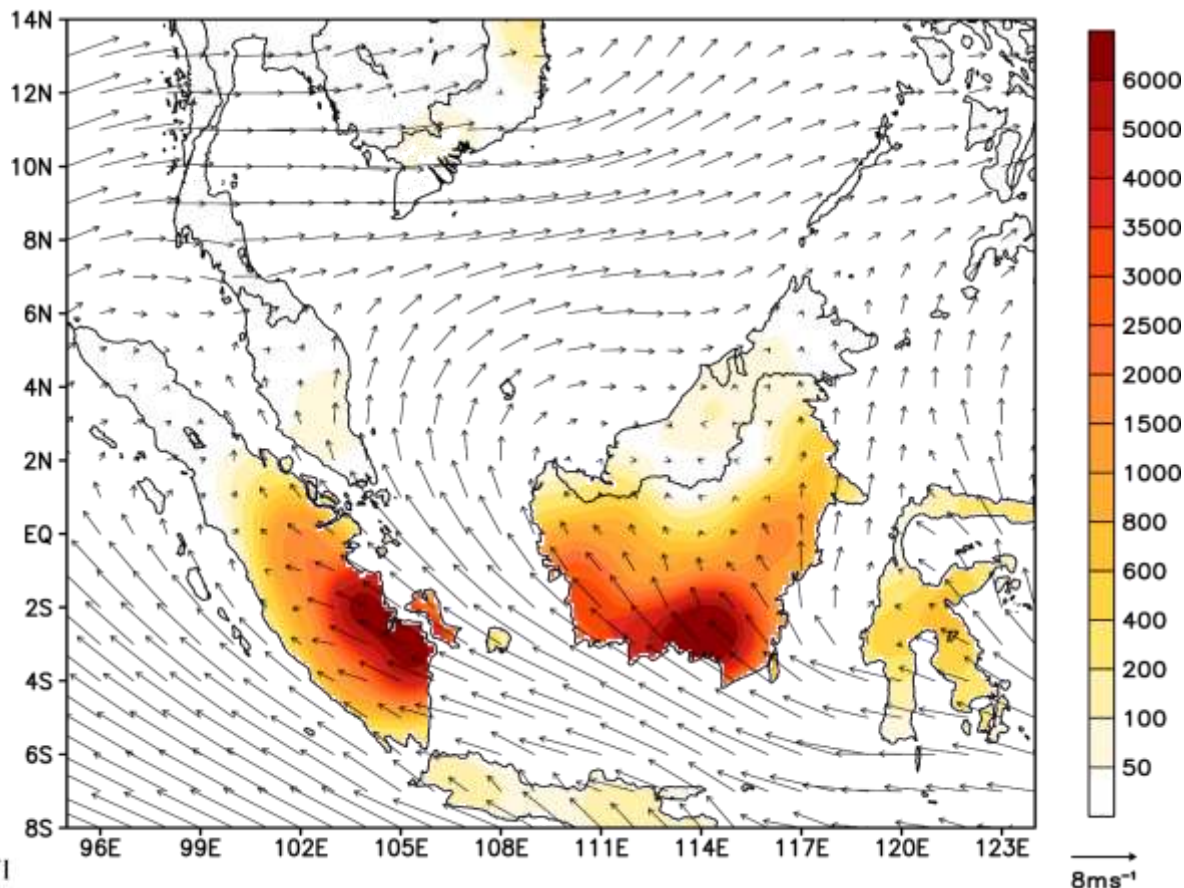


Flaming

Smouldering

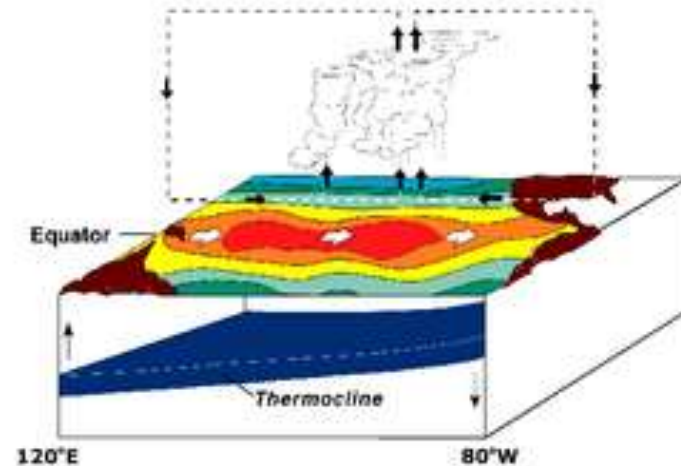


Wind Direction – Southwest Monsoon



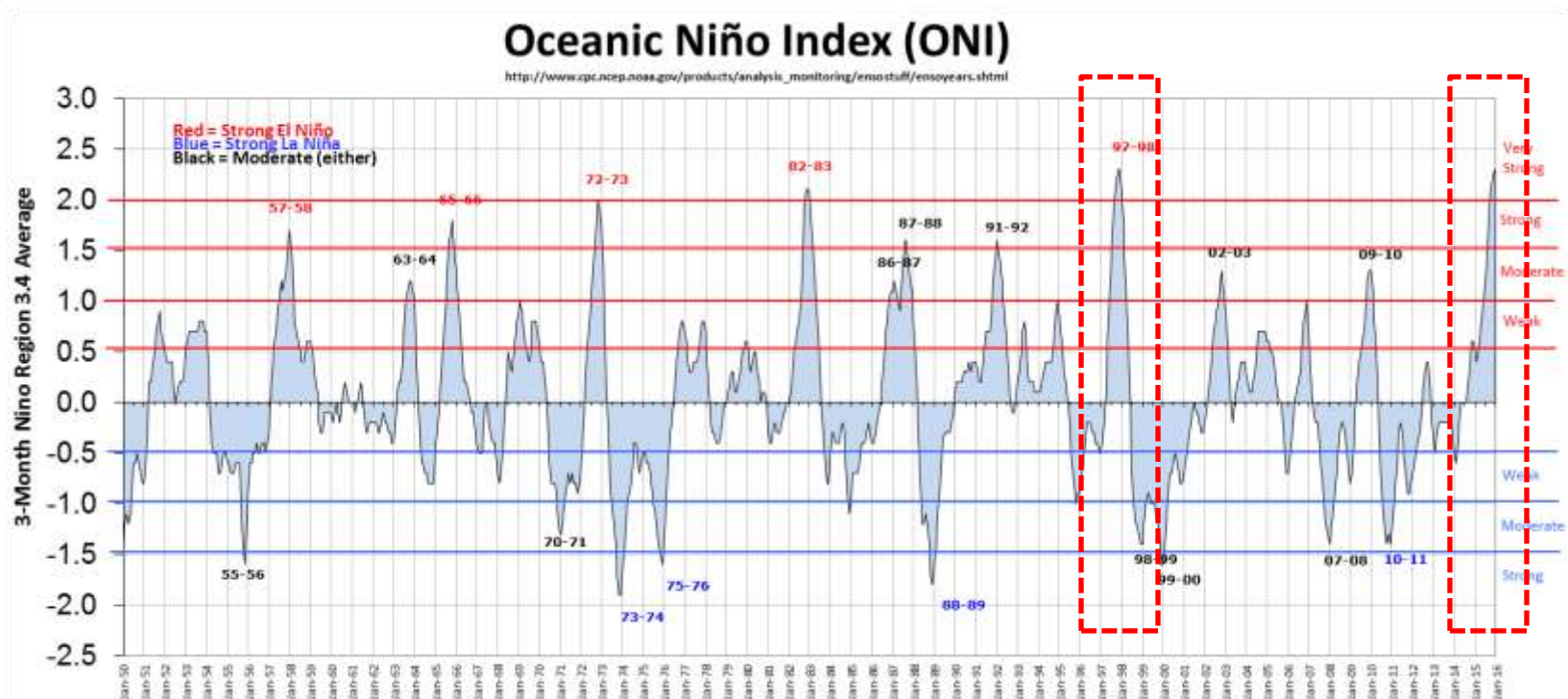
Trigger to Biomass Burning

- **El Niño–Southern Oscillation (ENSO)** event have repeatedly created conditions that make even rainforest susceptible to wildfires.





El-Niño and La-Niña

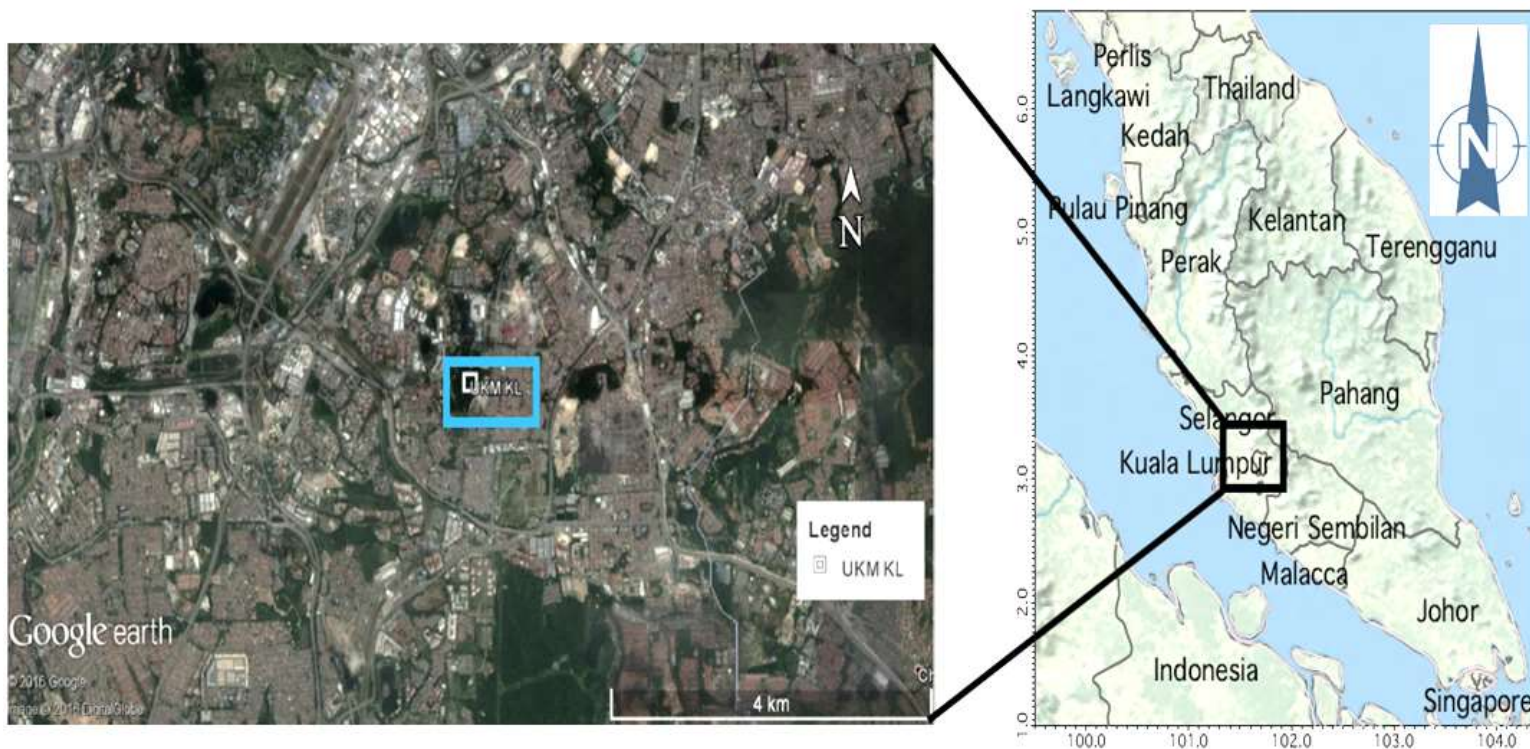


Research Objectives

- To determine the concentrations of $PM_{2.5}$ in Kuala Lumpur and its inorganic compositions during pre-haze, haze and post-haze periods.
- To predict the $PM_{2.5}$ concentration transported towards Kuala Lumpur using Numerical Atmospheric-dispersion Modelling Environment (NAME) together with the Global Fire Assimilation System (GFAS).
- To apportion possible sources of $PM_{2.5}$ using Positive Matrix Factorisation (PMF).
- To estimate the carcinogenic and non-carcinogenic health risks among specific age groups during pre-haze, haze and post-haze episodes.



Study Location



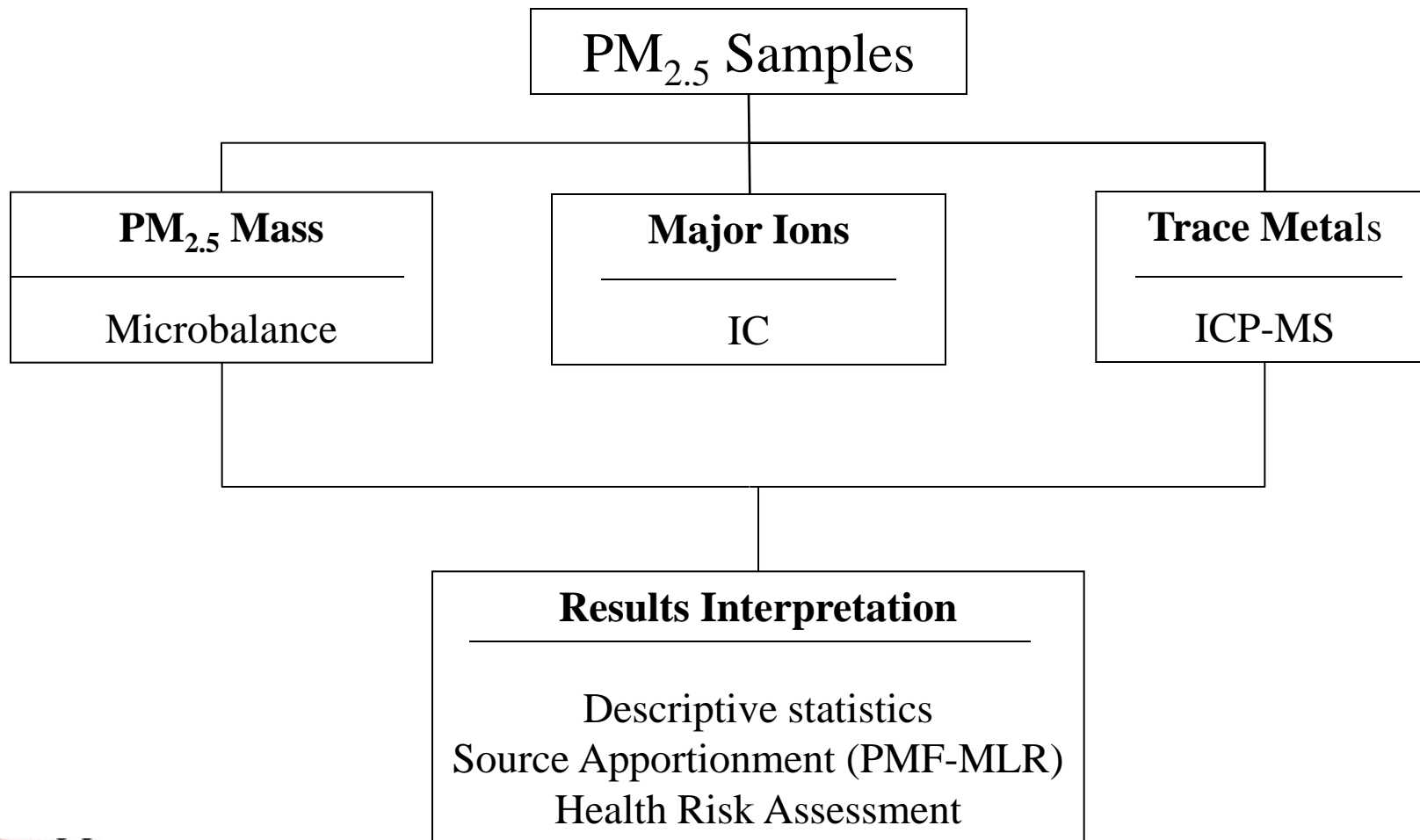
Sampling Site

- Kuala Lumpur located within the Klang Valley and situated in the middle of the west coast of the Malaysian Peninsular.
- In 2015, Kuala Lumpur had an estimated population of 1.78 million people in an area of just 243 km² with an average annual population growth rate of 2.4% (DOSM, 2016).
- The PM_{2.5} aerosol sampling was conducted on the rooftop (30 m above ground level) of a building within the compound of Universiti Kebangsaan Malaysia Kuala Lumpur's campus.

PM_{2.5} Sampling



- Tisch HVS PM_{2.5}
- Flowrate of 1.13 m³ min⁻¹
- 24 h sampling/filter
- Quartz filter [Whatman QM-A; 8' X 10']
- June 2015– January 2016
- Nine samples per month



Backward Trajectories

- Numerical Atmospheric-dispersion Modelling Environment (NAME), a Lagrangian particle dispersion model, produced by the United Kingdom's Met Office's.
- The backward trajectories started at the latitude-longitude coordinates of the measurement site in Kuala Lumpur within an altitude range of 0 – 100 m and ran for 5 days.

Modelled PM_{2.5} concentration

- Global Fire Assimilation System (GFAS) PM_{2.5} emissions data.
- This dataset relies on fire radiative power (FRP) observations obtained from the Terra and Aqua satellites - Moderate Resolution Imaging Spectroradiometer (MODIS) instruments.
- Combination of emission sensitivities derived from NAME with the GFAS emissions - modelled PM_{2.5} concentration

Modelled PM_{2.5} concentration

- Combination of emission sensitivities derived from NAME with the GFAS emissions - modelled PM_{2.5} concentration

Mass/cubic meter ($\mu\text{g m}^{-3}$)

= Emission sensitivities (s m^{-1}) x GFAS emissions ($\text{g m}^{-2} \text{s}^{-1}$) x 10^6 ($\mu\text{g g}^{-1}$)

Source Apportionment of PM_{2.5}

- Positive Matrix Factorization (PMF) 5.0 model from the United States Environmental Protection Agency (US EPA).

$$X_{ij} = \sum_{k=1}^p g_{ik} f_{kj} + e_{ij}$$

Normalized data

Source profile

Source contribution

Measurement error

Other Air Pollutants and Meteorological Data

- Temperature, rainfall, relative humidity, visibility, wind speed and wind direction (MetMalaysia Petaling Jaya station, 9 km from the sampling station)
- Air Pollution Index (API) data was collected from Malaysian DOE (Batu Muda Station, 6 km from the sampling site)

Carcinogenic Metal Health Risk

$$\text{Lifetime average daily dose (LADD)} \ (\mu\text{gkg}^{-1}\text{day}^{-1}) = \frac{C \times IR \times ED \times EF}{BW \times AT}$$

C = Concentration of the contaminant in the atmosphere (ng m^{-3}),

IR = inhalation rate ($\text{m}^3 \text{day}^{-1}$)

ED = exposure duration (years)

EF = exposure frequency (day year^{-1})

BW = body weight (kg)

AT = averaging time (70 years x 365 days).

Excess lifetime cancer risk (ELCR Inhalation)

= LADD \times inhalation unit risk ($\mu\text{g m}^{-3}$)⁻¹

Non-Carcinogenic Metal Health Risk

$$\text{Average daily dose (mgkg}^{-1}\text{day}^{-1}) = \frac{C \times IR \times ED \times EF}{BW \times AT}$$

$$\text{Hazard quotients (HQ)} = \text{ADD}/\text{RfC}$$

ADD = average daily dose

RfC = reference concentration

$$\text{HI} = \sum \text{HQ}$$

If the $\text{HQ} \leq 1$, it is believed that there is no risk of developing non-cancer health effects.

Non-cancer effects may occur if the $\text{HQ} > 1$.

An adverse effect is deemed more likely to occur if the HQ value is larger.

Exposure factors for calculating the exposure dose in health risk assessment

EP	Unit	Infant 0-<1year	Toddler 1-<6years	Children 6-<12years	Adolescent 12-<18years	Adult 18-<70years
IR	m ³ day ⁻¹	5.4	9	12	15.7	15.7
ED	years	1	5	6	6	52
EF	days year ⁻¹	60 ^a , 305 ^b	60, 305	60, 305	60, 305	60, 305
BW*	kg	7	15	31.2	38	66
AT	years	70 ^{**} , 1 ^{***}	70, 5	70, 6	70, 6	70, 52

* Adapted from National Health and Morbidity Survey III 2006 (NHMS III, 2008)

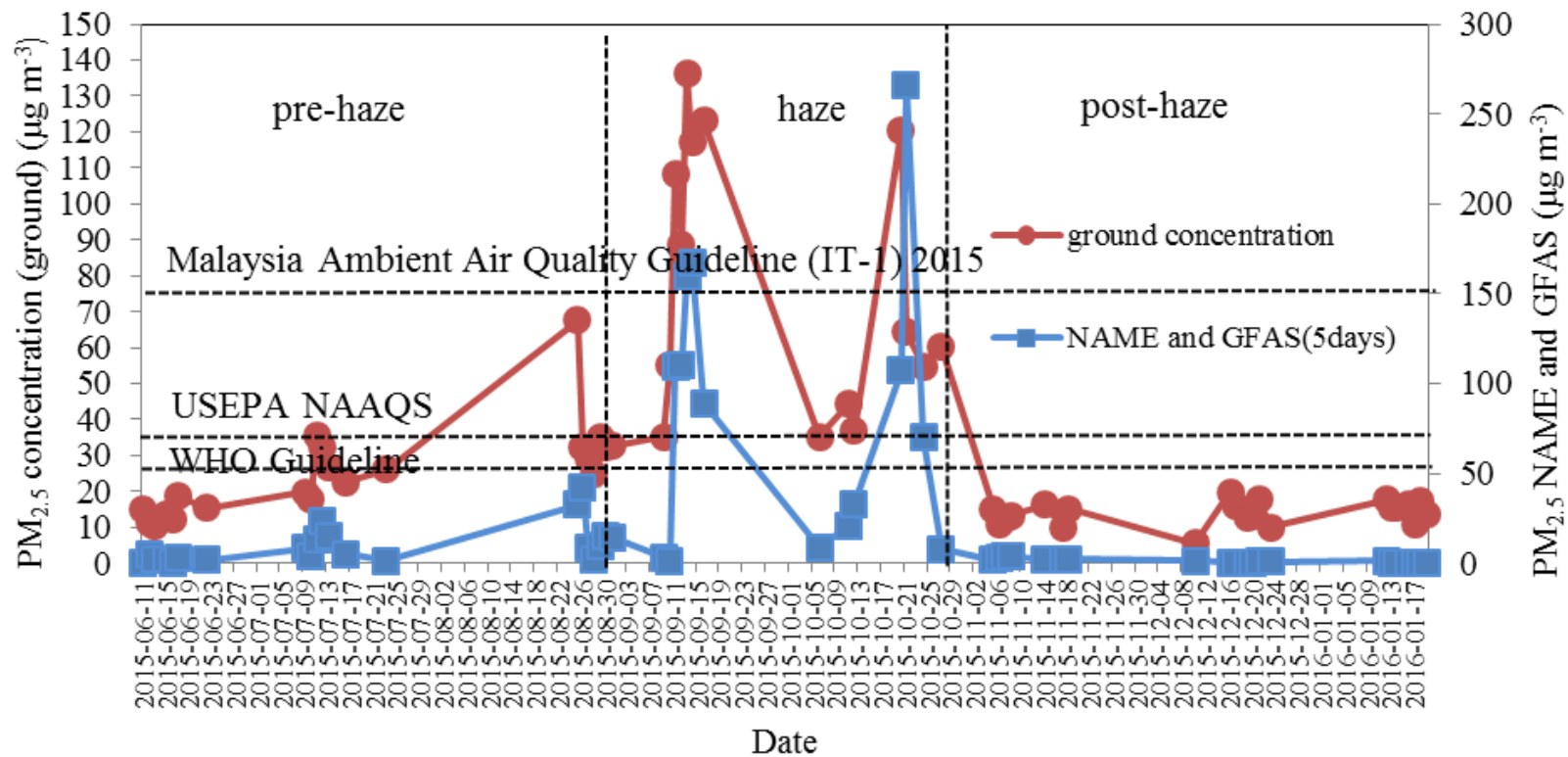
a exposure factors of haze episode

b exposure factors of non-haze episode

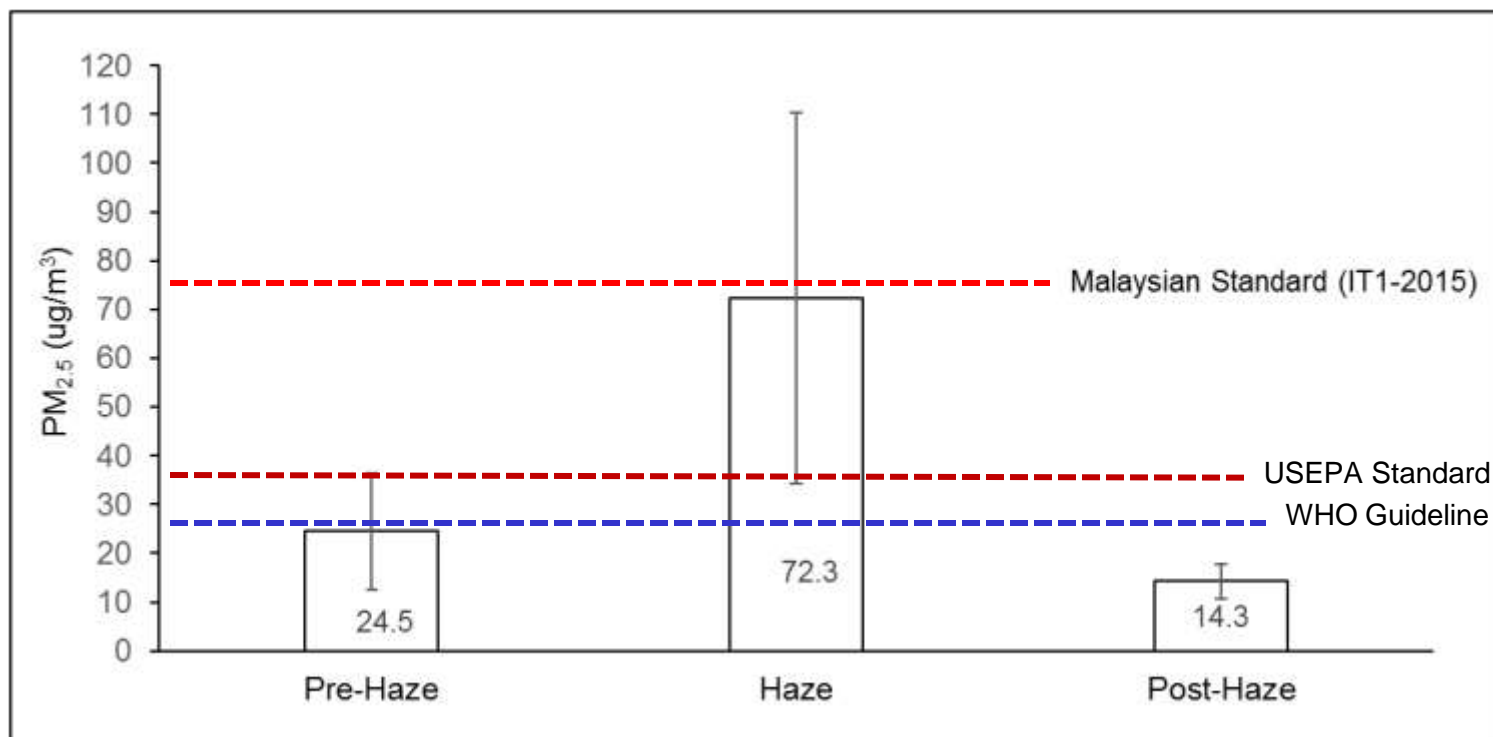
** AT for carcinogens (fixed at 70 years of exposure)

*** AT for non-carcinogens (average years of exposure)

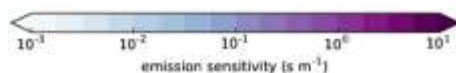
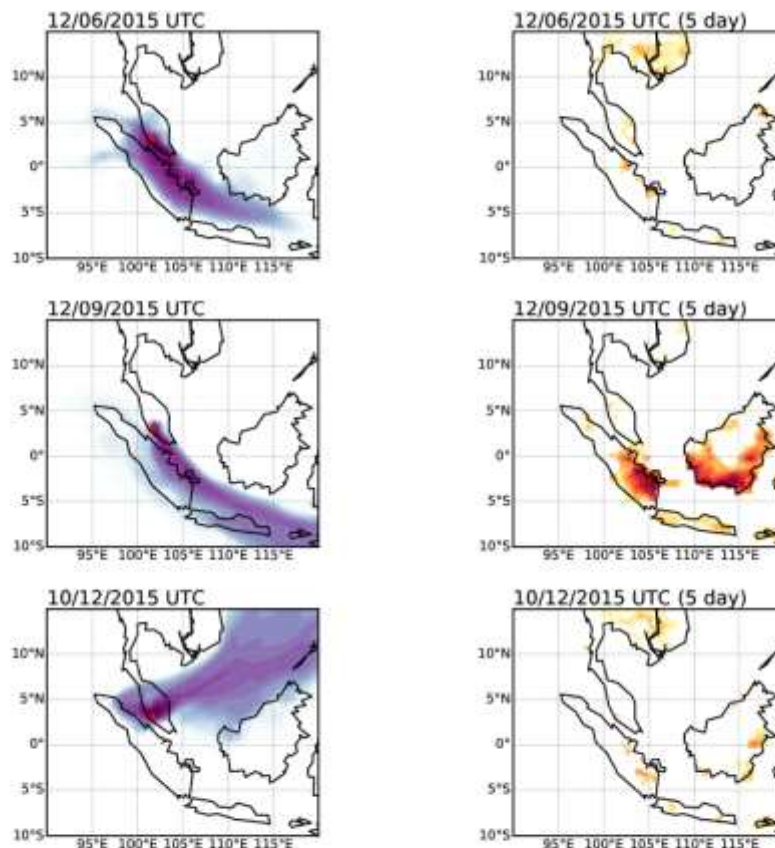
Measured HVS PM_{2.5} vs Predicted NAME-GFAS PM_{2.5}



Average PM_{2.5} Concentration

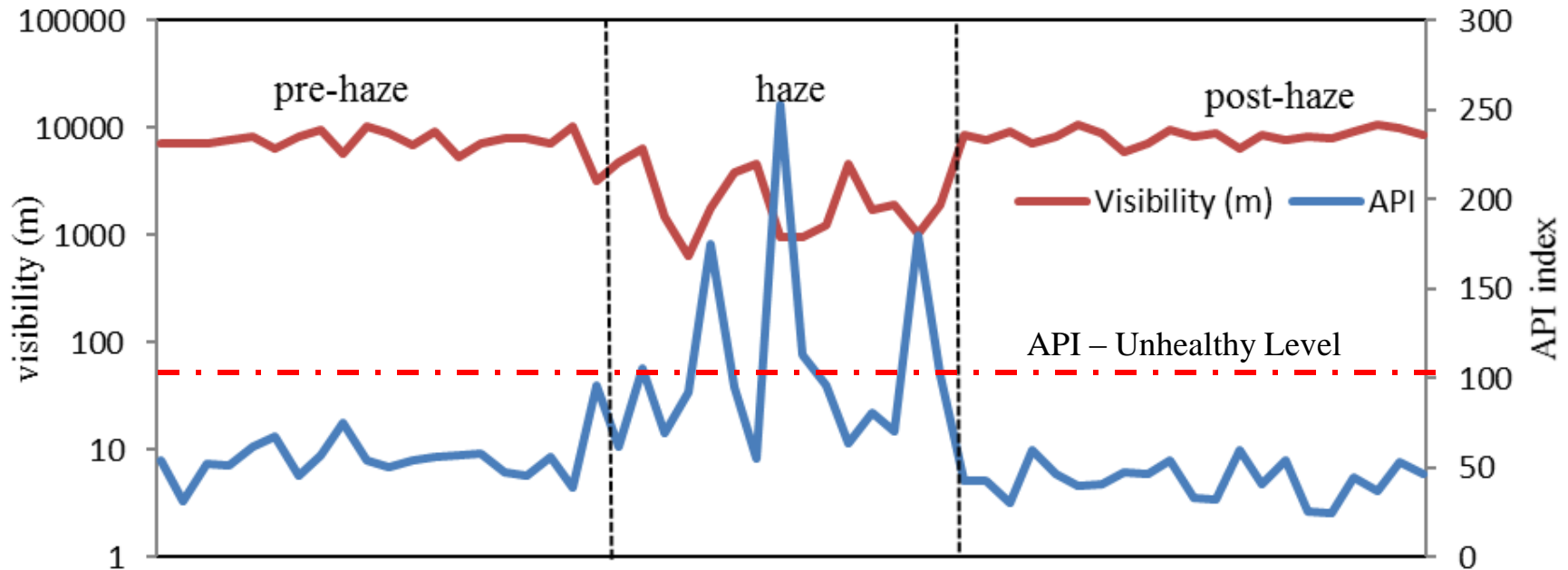


Emission Sensitivity and Emission



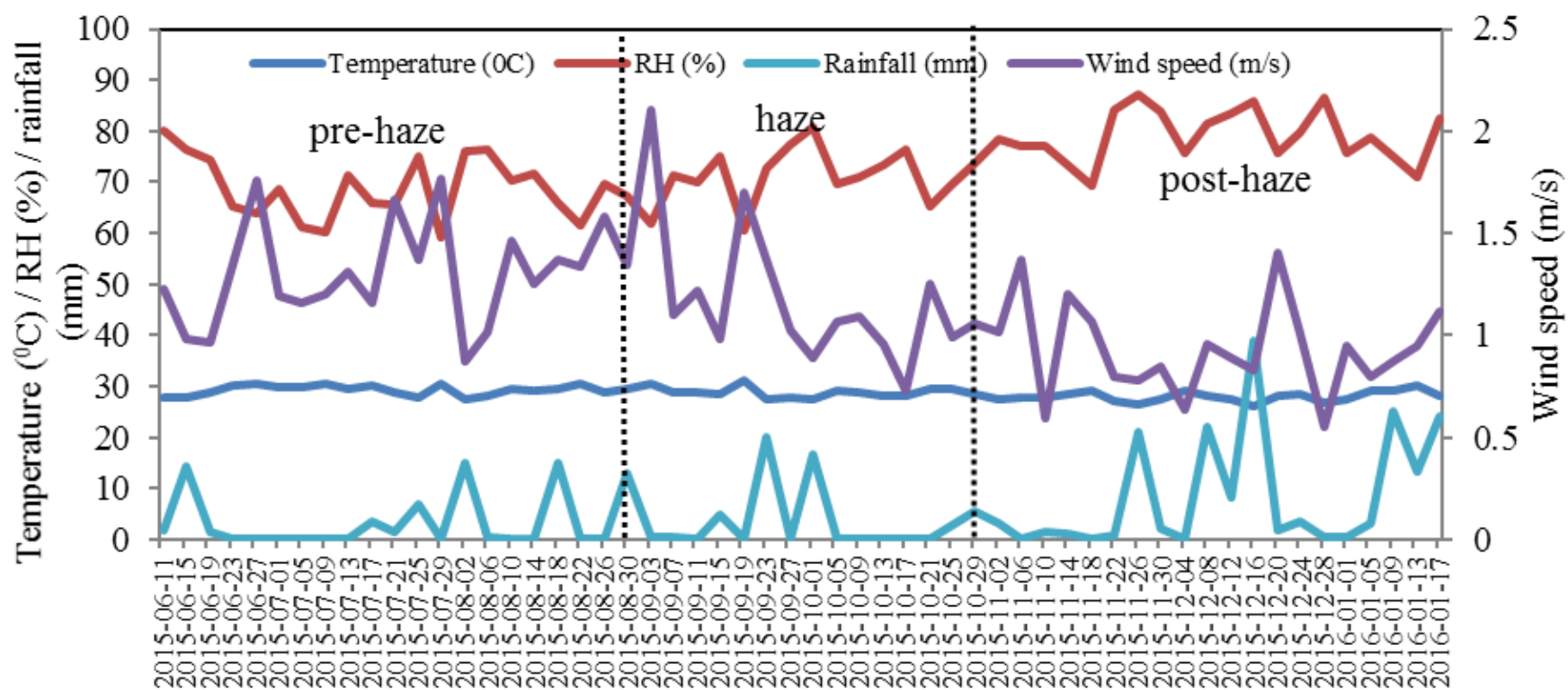


Visibility and API Index



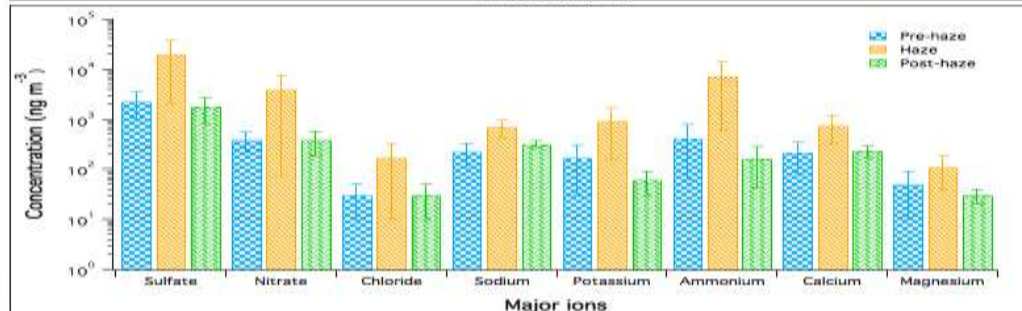
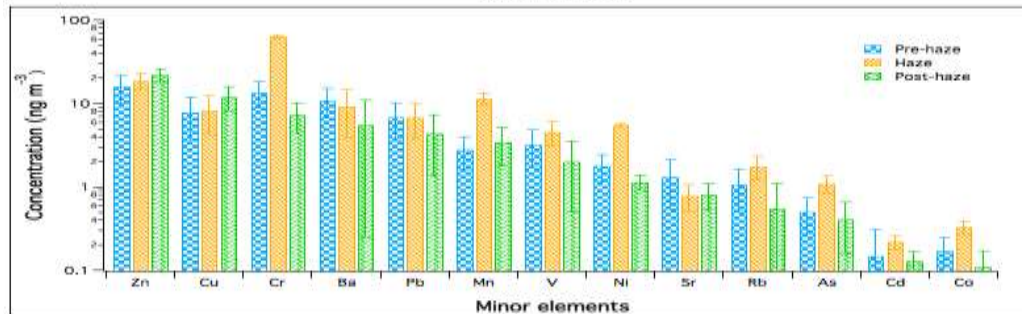
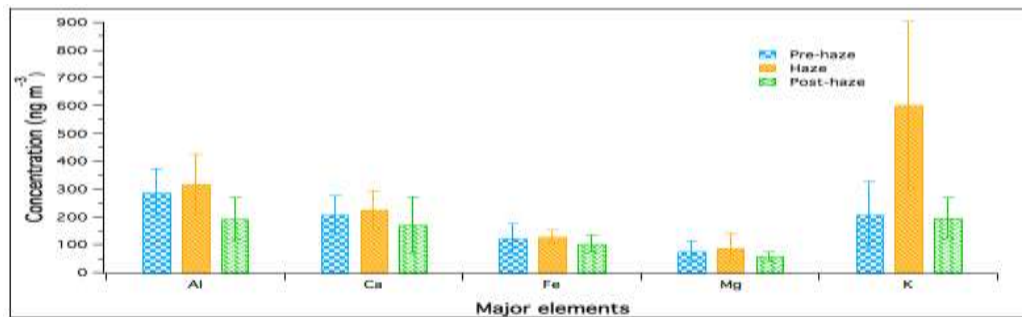


Other Meteorological Data

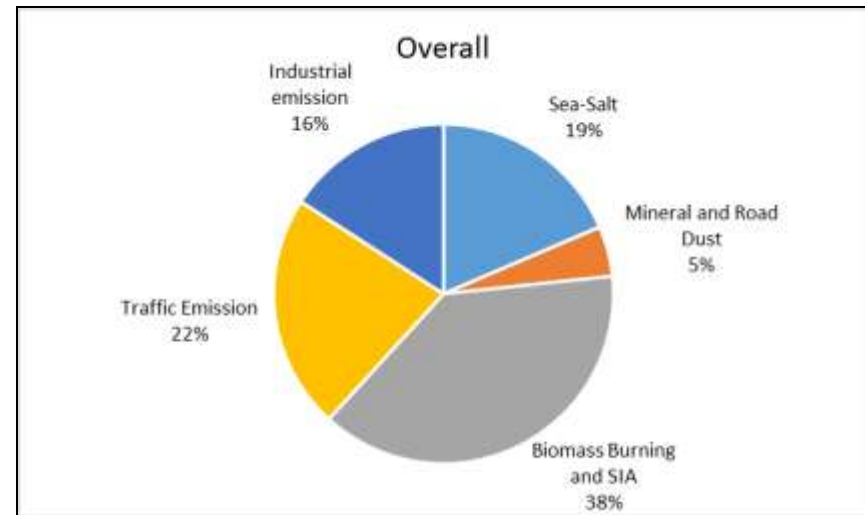
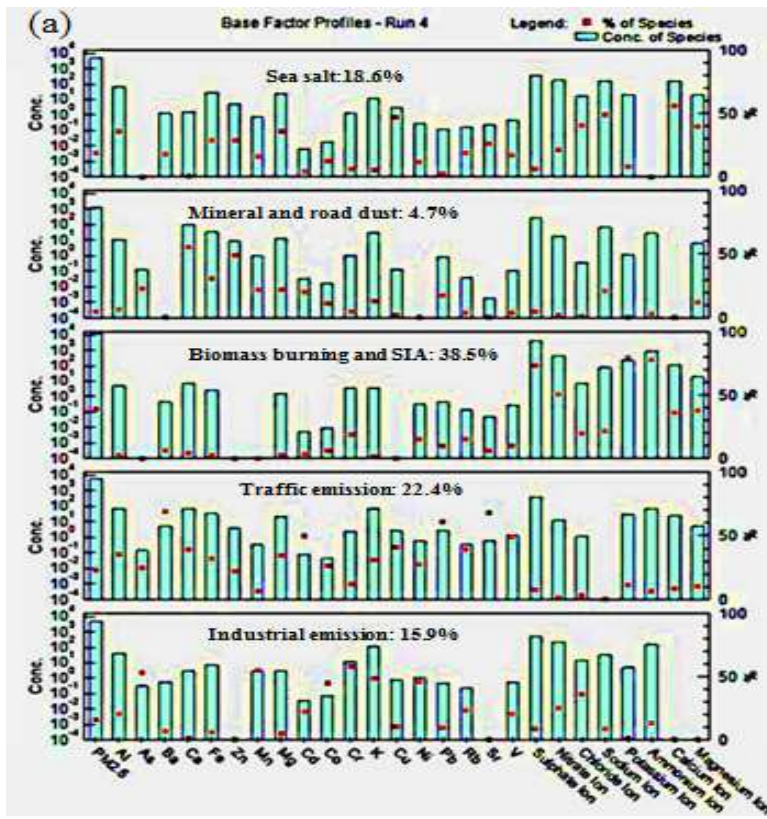




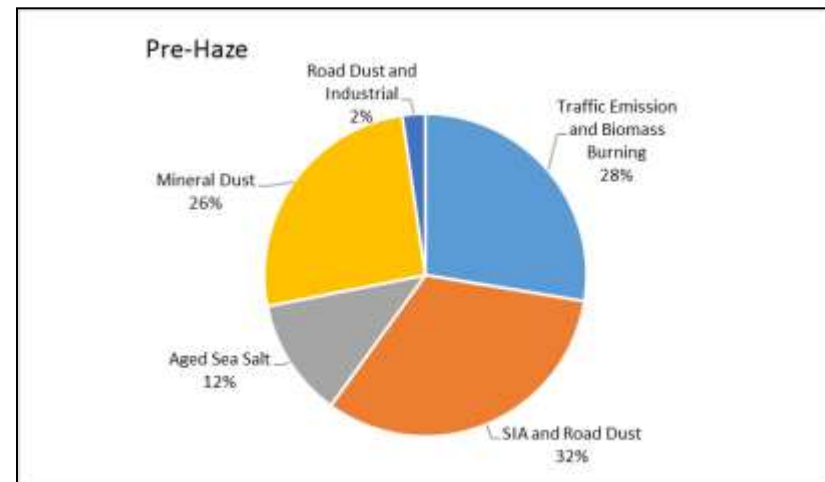
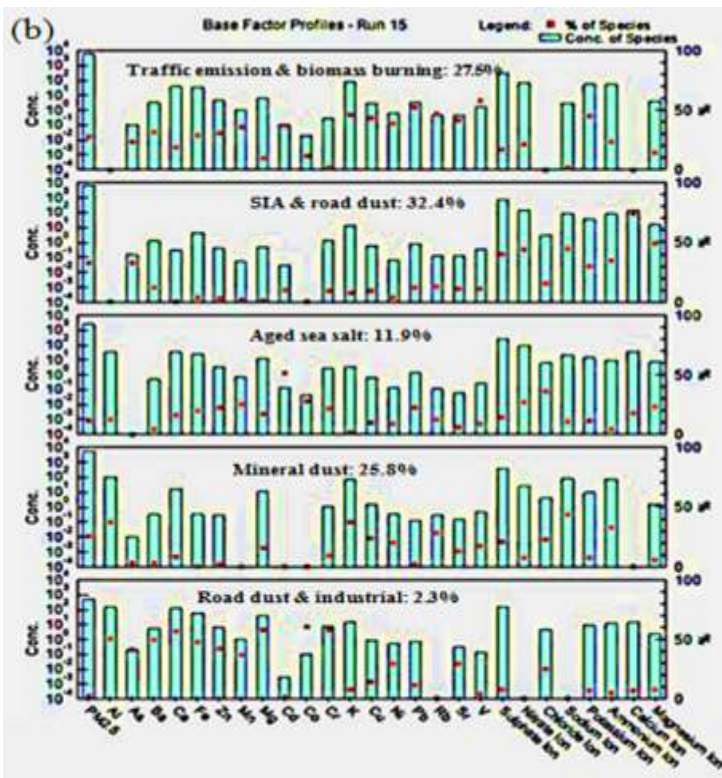
PM_{2.5} Inorganic Composition



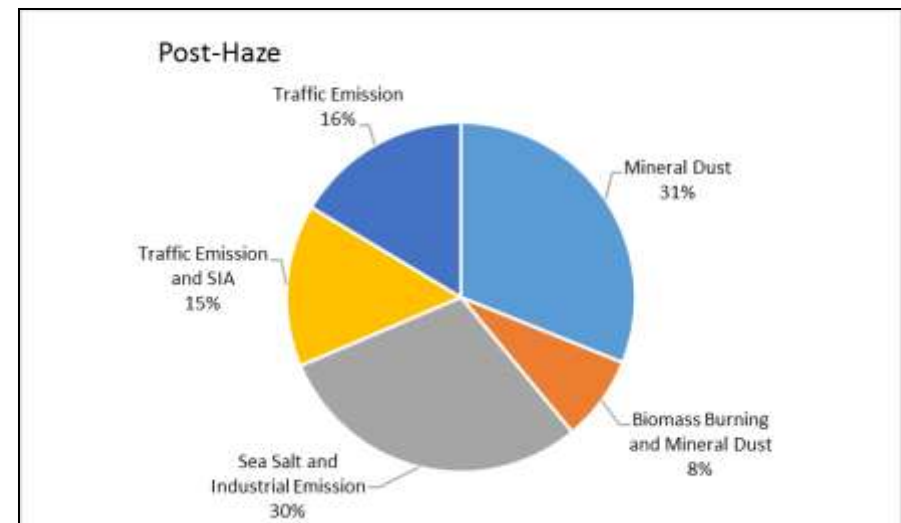
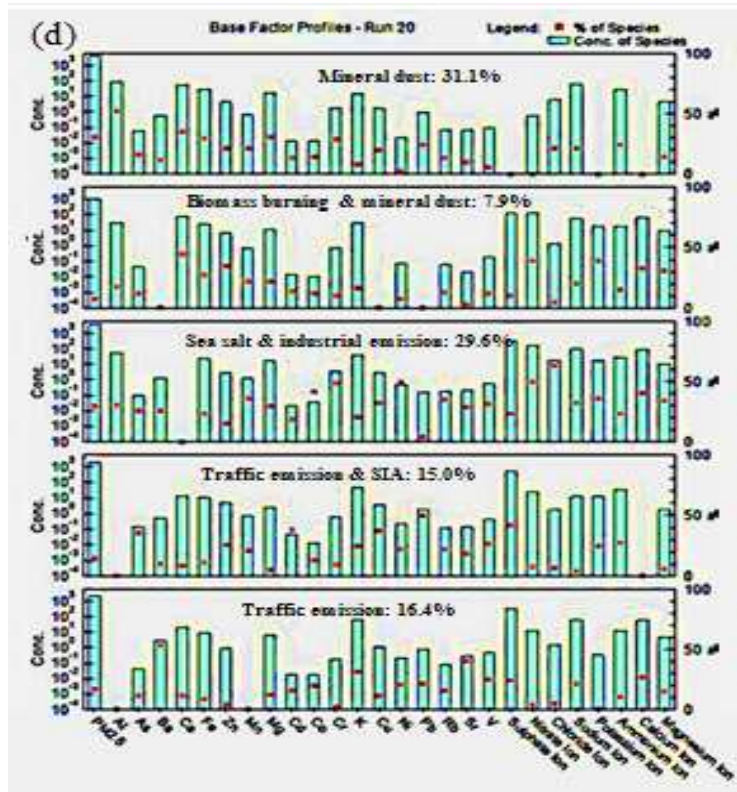
Source Apportionment -Overall



Source Apportionment –Pre-haze



Source Apportionment – Post-Haze



Hazard index (HI) and hazard quotient (HQ)

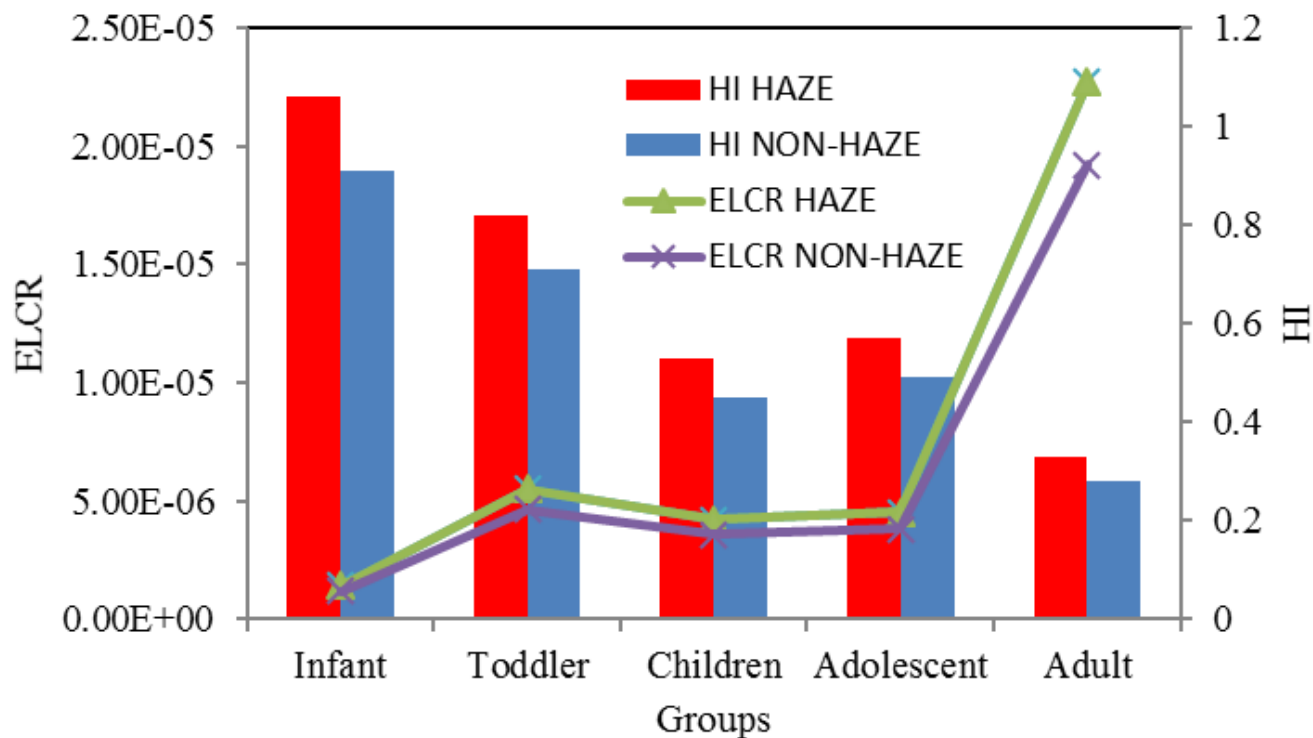
Elements	RfC (mg m ⁻³)	HQ				
		Infant 0-<1year	Toddler 1-<6years	Children 6-<12years	Adolescent 12-<18years	Adult 18-<70years
Pre-haze						
Cr	8.00×10^{-6}	1.10×10^0	8.59×10^{-1}	5.50×10^{-1}	5.91×10^{-1}	3.40×10^{-1}
Mn	5.00×10^{-5}	3.60×10^{-2}	2.80×10^{-2}	1.80×10^{-2}	1.93×10^{-2}	1.11×10^{-2}
Ni	2.00×10^{-4}	5.70×10^{-3}	4.46×10^{-3}	2.86×10^{-3}	3.07×10^{-3}	1.77×10^{-3}
Cd	1.00×10^{-5}	9.67×10^{-3}	7.52×10^{-3}	4.82×10^{-3}	5.18×10^{-3}	2.98×10^{-3}
As	5.00×10^{-5}	6.57×10^{-3}	5.11×10^{-3}	3.28×10^{-3}	3.52×10^{-3}	2.03×10^{-3}
∑HI		1.16	0.90	0.58	0.62	0.36
Haze						
Cr	8.00×10^{-6}	1.02×10^0	7.93×10^{-1}	5.08×10^{-1}	5.46×10^{-1}	3.14×10^{-1}
Mn	5.00×10^{-5}	2.94×10^{-2}	2.29×10^{-2}	1.47×10^{-2}	1.58×10^{-2}	9.07×10^{-3}
Ni	2.00×10^{-4}	3.51×10^{-3}	2.73×10^{-3}	1.75×10^{-3}	1.88×10^{-3}	1.08×10^{-3}
Cd	1.00×10^{-5}	2.79×10^{-3}	2.17×10^{-3}	1.39×10^{-3}	1.49×10^{-3}	8.60×10^{-4}
As	5.00×10^{-5}	2.81×10^{-3}	2.19×10^{-3}	1.40×10^{-3}	1.51×10^{-3}	8.68×10^{-4}
∑HI		1.06	0.82	0.53	0.57	0.33
Post-haze						
Cr	8.00×10^{-6}	5.87×10^{-1}	4.57×10^{-1}	2.93×10^{-1}	3.15×10^{-1}	1.81×10^{-1}
Mn	5.00×10^{-5}	4.53×10^{-2}	3.52×10^{-2}	2.26×10^{-2}	2.42×10^{-2}	1.40×10^{-2}
Ni	2.00×10^{-4}	3.67×10^{-3}	2.86×10^{-3}	1.83×10^{-3}	1.97×10^{-3}	1.13×10^{-3}
Cd	1.00×10^{-5}	8.38×10^{-3}	6.52×10^{-3}	4.18×10^{-3}	4.49×10^{-3}	2.58×10^{-3}
As	5.00×10^{-5}	5.29×10^{-3}	4.11×10^{-3}	2.63×10^{-3}	2.83×10^{-3}	1.63×10^{-3}
∑HI		0.65	0.51	0.32	0.35	0.20
∑HI	Haze	1.06	0.82	0.53	0.57	0.33
∑HI	Non-haze*	0.91	0.71	0.45	0.49	0.28

Excess lifetime cancer risks (ELCR) of carcinogenic

Elements	IUR ($\mu\text{g m}^{-3}$) ⁻¹	Excess lifetime cancer risk ($\mu\text{g m}^{-3}$) ⁻¹				
		Infant 0-<1year	Toddler 1-<6years	Children 6-<12years	Adolescent 12-<18years	Adult 18-<70years
Pre-haze						
Pb	1.20×10^{-5}	7.53×10^{-10}	2.93×10^{-9}	2.25×10^{-9}	2.42×10^{-9}	1.21×10^{-8}
Cd	1.80×10^{-3}	2.49×10^{-9}	9.67×10^{-9}	7.44×10^{-9}	8.00×10^{-9}	4.00×10^{-8}
Cr	1.20×10^{-2}	1.51×10^{-6}	5.89×10^{-6}	4.53×10^{-6}	4.86×10^{-6}	2.43×10^{-5}
Ni	2.40×10^{-4}	3.93×10^{-9}	1.53×10^{-8}	1.18×10^{-8}	1.26×10^{-8}	6.31×10^{-8}
As	4.30×10^{-3}	2.02×10^{-8}	7.85×10^{-8}	6.04×10^{-8}	6.49×10^{-8}	3.24×10^{-7}
Co	9.00×10^{-3}	1.41×10^{-8}	5.48×10^{-8}	4.22×10^{-8}	4.53×10^{-8}	2.26×10^{-7}
Σ		1.56×10^{-6}	6.05×10^{-6}	4.65×10^{-6}	5.00×10^{-6}	2.49×10^{-5}
Haze						
Pb	1.20×10^{-5}	1.50×10^{-10}	5.83×10^{-10}	4.49×10^{-10}	4.82×10^{-10}	2.41×10^{-9}
Cd	1.80×10^{-3}	7.17×10^{-10}	2.79×10^{-9}	2.15×10^{-9}	2.31×10^{-9}	1.15×10^{-8}
Cr	1.20×10^{-2}	1.40×10^{-6}	5.44×10^{-6}	4.18×10^{-6}	4.49×10^{-6}	2.24×10^{-5}
Ni	2.40×10^{-4}	2.41×10^{-9}	9.37×10^{-9}	7.21×10^{-9}	7.74×10^{-9}	3.86×10^{-8}
As	4.30×10^{-3}	8.65×10^{-9}	3.36×10^{-8}	2.59×10^{-8}	2.78×10^{-8}	1.39×10^{-7}
Co	9.00×10^{-3}	5.38×10^{-9}	2.09×10^{-8}	1.61×10^{-8}	1.73×10^{-8}	8.63×10^{-8}
Σ		1.42×10^{-6}	5.50×10^{-6}	4.23×10^{-6}	4.55×10^{-6}	2.27×10^{-5}
Post-haze						
Pb	1.20×10^{-5}	4.91×10^{-10}	1.91×10^{-9}	1.47×10^{-9}	1.58×10^{-9}	7.87×10^{-9}
Cd	1.80×10^{-3}	2.15×10^{-9}	8.38×10^{-9}	6.45×10^{-9}	6.92×10^{-9}	3.46×10^{-8}
Cr	1.20×10^{-2}	8.06×10^{-7}	3.13×10^{-6}	2.41×10^{-6}	2.59×10^{-6}	1.29×10^{-5}
Ni	2.40×10^{-4}	2.52×10^{-9}	9.80×10^{-9}	7.54×10^{-9}	8.10×10^{-9}	4.04×10^{-8}
As	4.30×10^{-3}	1.62×10^{-8}	6.31×10^{-8}	4.86×10^{-8}	5.22×10^{-8}	2.60×10^{-7}
Co	9.00×10^{-3}	8.29×10^{-9}	3.22×10^{-8}	2.48×10^{-8}	2.66×10^{-8}	1.33×10^{-7}
Σ		8.35×10^{-7}	3.25×10^{-6}	2.50×10^{-6}	2.68×10^{-6}	1.34×10^{-5}
Σ ELCR	Haze	1.42×10^{-6}	5.50×10^{-6}	4.23×10^{-6}	4.55×10^{-6}	2.27×10^{-5}
Σ ELCR	Non-haze*	1.20×10^{-6}	4.65×10^{-6}	3.58×10^{-6}	3.84×10^{-6}	1.92×10^{-5}



Atmosphere Investigation



Conclusion

- $\text{PM}_{2.5}$ mass collected during pre-haze ($24.5 \pm 12.0 \mu\text{g m}^{-3}$), haze ($72.3 \pm 38.0 \mu\text{g m}^{-3}$), and post-haze ($14.3 \pm 3.58 \mu\text{g m}^{-3}$) events in Kuala Lumpur were significantly different ($p < 0.005$)
- The highest concentration of $\text{PM}_{2.5}$ during haze episode - 5 times higher than WHO guidelines, 3.9 times higher than the US EPA standards and 1.8 times higher than the Malaysian Ambient Air Quality Standards 2015 (IT-1)
- The concentration of $\text{PM}_{2.5}$ recorded during the haze episode had a good correlation with the Malaysian Air Pollutants Index (API) ($r = 0.466$; $p < 0.05$) and significantly reduce the visibility ($r = -0.631$; $p = 0.005$).

Conclusion

- The SIA (SO_4^{2-} , NO_3^- and NH_4^+) dominated the composition of $\text{PM}_{2.5}$ - contribute to 43% inorganic composition of $\text{PM}_{2.5}$ mass during haze compared to pre-haze and post-haze, where they only contributed 12% and 16%, respectively.
- The overall dominant sources of $\text{PM}_{2.5}$ in Kuala Lumpur urban environment were SIA and biomass burning (38.5%); and traffic emission (22.4%).

Conclusion

- The non-carcinogenic health risk assessment - infant group faced more significant health risk than the other age groups during haze (HI = 1.06).
- The carcinogenic health risk assessment - adult group is the most affected group for haze exposure (ECLR= 2.27×10^{-5})
- The lowest ELCR estimation was posed by the infant group during non-haze (1.20×10^{-6}) indicating that 1 – 2 individuals in 1,000,000 are likely to develop cancer in their lifetime due to exposure of urban PM_{2.5} aerosols.

Further reading....

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Source apportionment and health risk assessment among specific age groups during haze and non-haze episodes in Kuala Lumpur, Malaysia

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ABSTRACT

This study aims to determine $PM_{2.5}$ concentrations and their composition during haze and non-haze episodes in Kuala Lumpur. In order to investigate the origin of the measured air masses, the Numerical Atmospheric-dispersion Modelling Environment (NAME) and Global Fire Assimilation System (GFAS) were applied. Source apportionment of $PM_{2.5}$ was determined using Positive Matrix Factorization (PMF). The carcinogenic and non-carcinogenic health risks were estimated using the United State Environmental Protection Agency (USEPA) method. $PM_{2.5}$ samples were collected from the centre of the city using a high-volume air sampler (HVS). The results showed that the mean $PM_{2.5}$ concentrations collected during non-haze, haze and



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THANK YOU