PM2.5 Carbonaceous species and BC Emitted from Peat Land burning in Riau Sumatera

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Background: Why PM2.5?

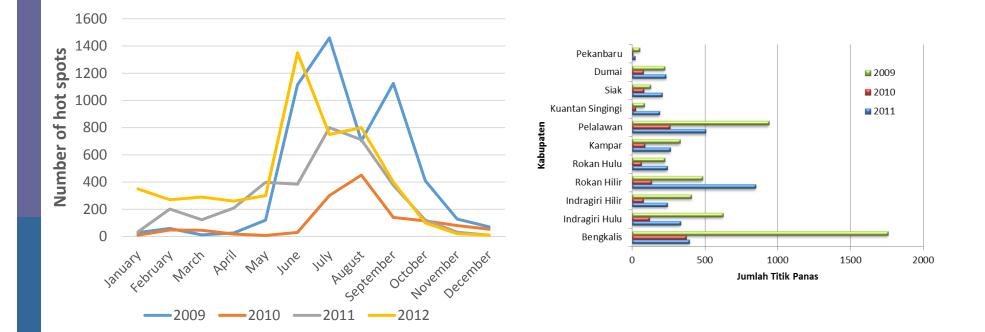
- According to the Global Fire Emissions Database (GFED), average PM2.5 emissions from fire (including deforestation, savanna, forest, agricultural waste, and peat fires) from 1997 to 2010 in Indonesia are accounting for 9.2% of global fire PM2.5 emissions and 62% of Southeast Asian fire emissions
- Peatland fire is a dominant source of PM2.5 emissions, accounting for 55% of all fire sources

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- Riau Province in Sumatra is one of the primary hotspots for peatland fire during the dry season, and the smoke aerosol generated there cause haze in Riau and in neighboring countries such as Malaysia and Singapore
- Limited data exist regarding the chemical characteristics of these
 smoke aerosols
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Background Information

Number of Hot Spot in RiauNumber of Hot Spot in





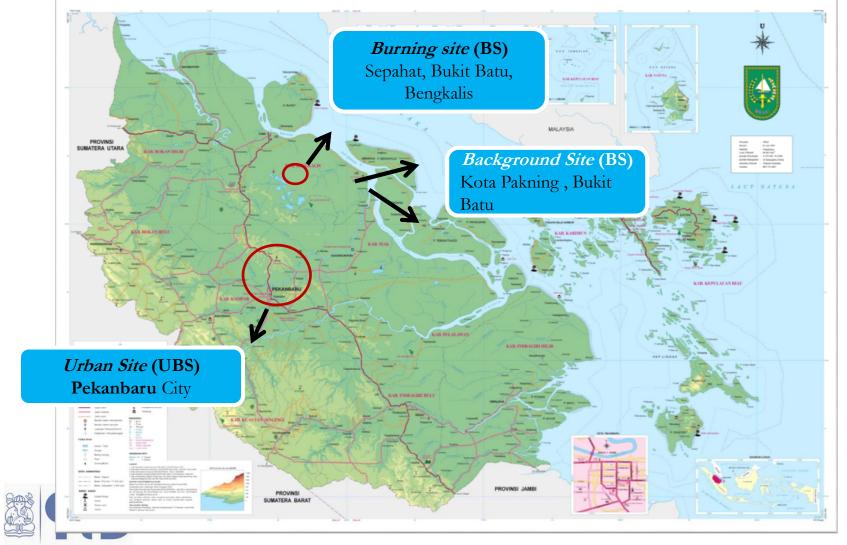
Objectives of the study

- To analyze carbonaneous particulate PM_{2.5} emission from peat land fires in Riau, Sumatera.
- To Quantitatively characterized carbon fraction of Organic Carbon (OC) and Elemental Carbon (EC) of the aerosols derived from peatland fires to determine potential source profile and indicators.
- To provide useful information to evaluate the contribution of Indonesian peatland fire aerosols to the air quality in Southeast Asia



7SEAS2010, Taiwan ,June15-18

Sampling Locations: Riau Sumatra



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Sampling Collection

Background Site



□ Minivolume sampler During burning periods □ May-June 2012





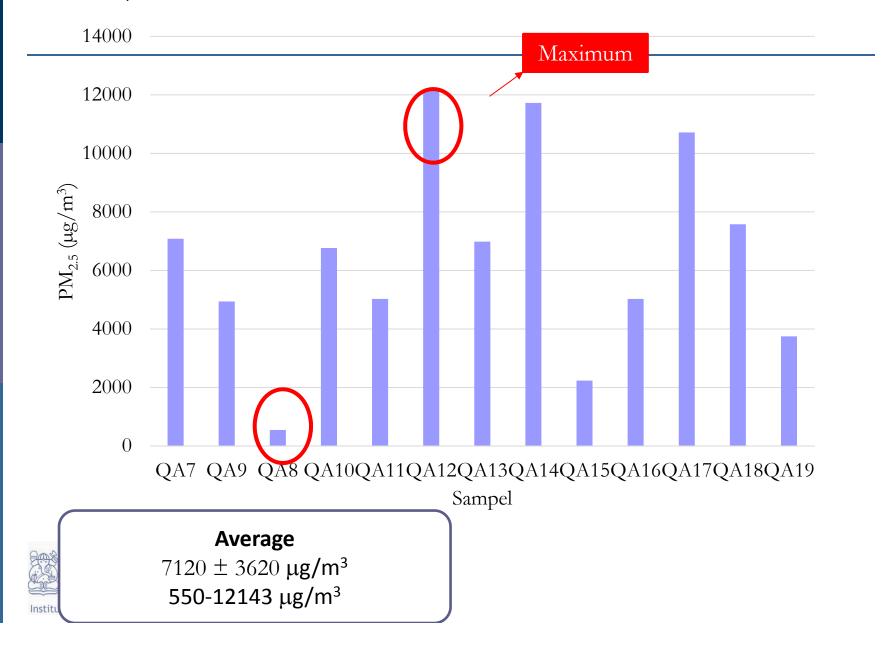


Analysis

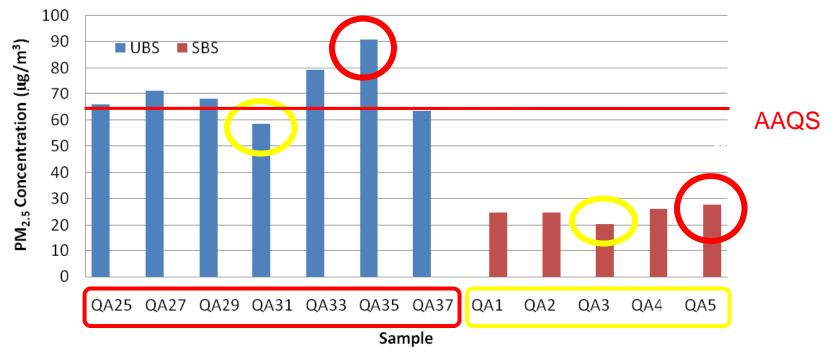
- OC and EC were analyze at Laboratory Research Institute for Advanced Science and Technology, Osaka Prefecture University, Japan (Kyoto Univ colaborator)
- The carbonaceous contents of the aerosols collected in the quartz fiber filters were quantified using a DRI Model 2001 OC/EC Carbon Analyzer, which employs thermal optical reflectance following the IMPROVE_A protocol.
- The IMPROVE_A temperature defines temperature plateaus for thermally-derived carbon fractions as follows: 140 C for OC1, 280 C for OC2, 480 C for OC3, and 580 C for OC4 in helium (He) carrier gas; 580 C for EC1, 740 C for EC2, and 840 C for EC3 in a mixture of 98% He and 2% oxygen (O2) carrier gas (Chow et al., 2007).
- □ BC were analyze using EEL type Smoke Stain Reflectometer
- OC, EC, and total carbon (TC) were calculated from the eight carbon fractions as follows:
- \Box OC= OC1 + OC2 + OC3 + OC4 + OP (1)
- $\square \quad \text{EC} = \text{EC1} + \text{EC2} + \text{EC3} \text{OP} (2)$
- $\Box TC = OC + EC$



PM_{2,5} Concentration in Burning Site



Concentrations of PM2.5 in Urban & Background sites

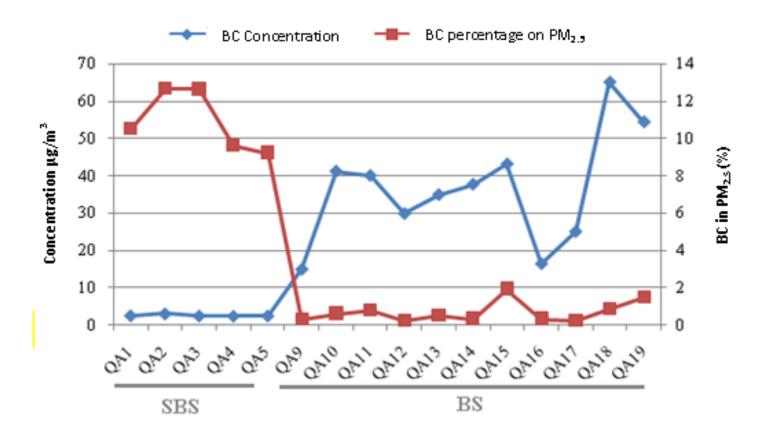


Urban site



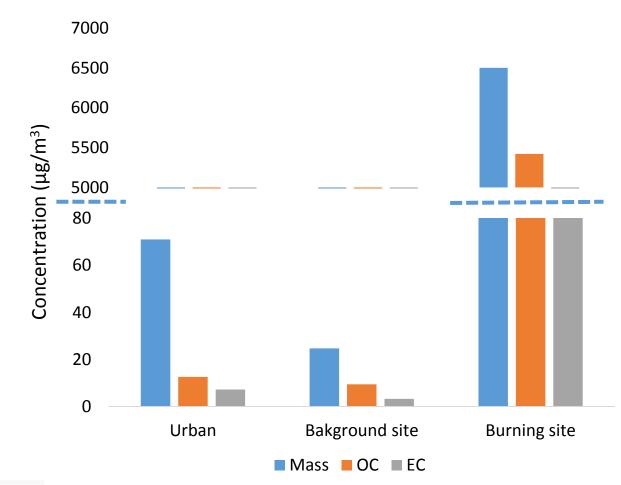
Background site

Concentration of BC at Burning Site and Background site (SBS)



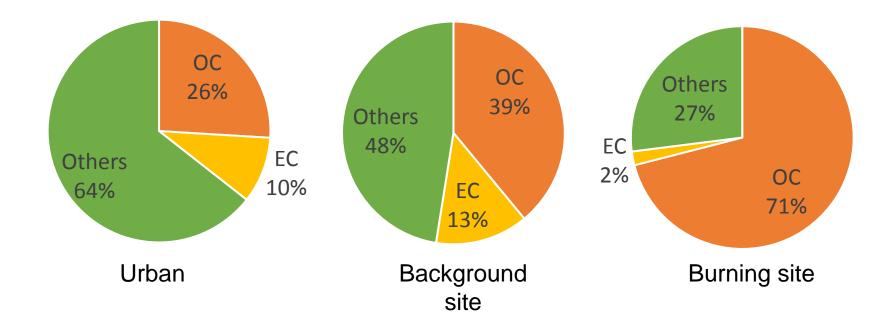


Comparison of OC/EC Concentrations in Burning, Urban and Background sites





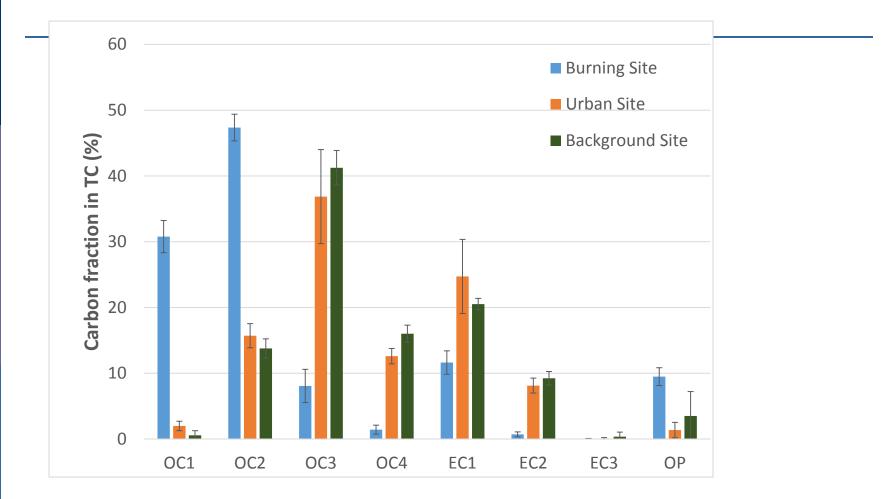
OC/EC contribution to PM2.5





Carbon Fraction in Total Carbon

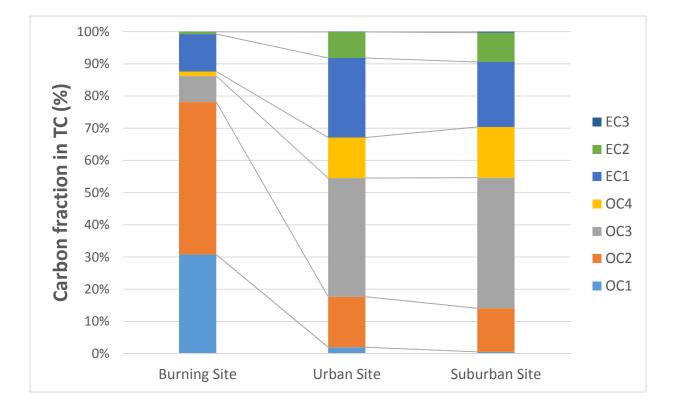
Abundances of eight thermally-derived carbon fractions differ by carbon sources



OC1 normally rich in biomass burning OC3 and OC4 relatively came from road dust OC2 was found in samples of coal combustion (decay plants)

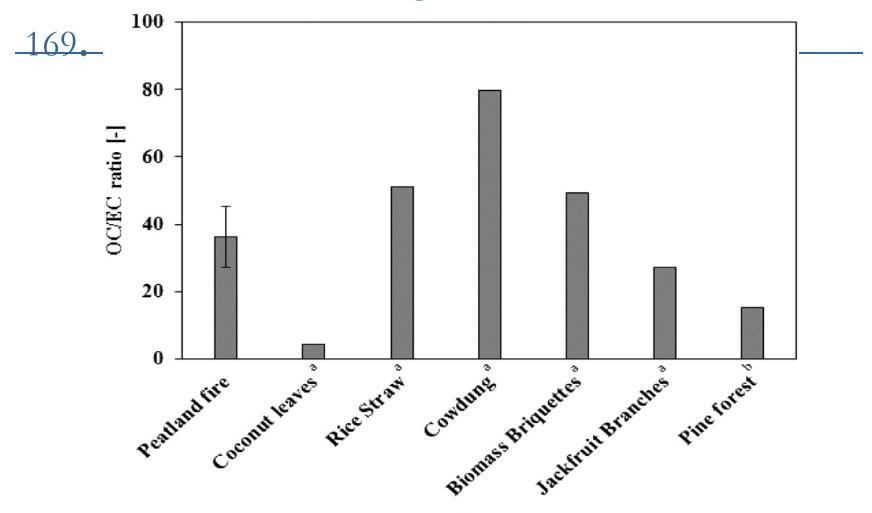
EC2 and EC3 mainly emitted by motor vehicles

Carbon Fraction in different locations





Comparison of OC/EC ratio from PM2.5 emitted from Peat land and other burning sources (AE 87(2014) 164-



^a data from Sheesley and Schauer, 2003, ^b data from Lee *et al.*, 2005.



(OC/EC) provide some indication of the origins of carbonaceous PM2.5

Conclusions:

- PM2.5 carbonaceous aerosols collected at a peatland fire hotspots had very high concentration of about 300 and 100 times of that colected at background and an urban sites respectively.
- Average PM2.5 aerosols emitted from peatland fire were observed in very high concentrations (7120 ± 3620 ug/ m3) and were primarily composed of OC (71.0 ±5.11% of PM2.5 mass)
- The OC/EC ratios (36.4 ± 9.08 for peatland fire), abundances of eight thermally-derived carbon fractions,
- PM2.5 from peat fire emissions is characterized by an abundance of OC1 and OC2 fractions

Emissions on burning site produces more volatile compounds with low molecular weight compared to the background site
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Thank you

