Integrated emission inventory and modeling to assess distribution of particulate matter mass and black carbon composition in Southeast Asia

Didin Agustian Permadi[1], Nguyen Thi Kim Oanh[1*] and Robert Vautard[2]

[1] Environmental Engineering and Management; School of Environment, Resources and Development; Asian Institute of Technology; Klong Luang, Pathumthani 12120, Thailand.
[2] Laboratoire des Sciences du Climate de l'Environment (LSCE), Institut Pierre Simon Laplace (IPSL), Gif Sur Yvette, France.

Correspondence to: Nguyen Thi Kim Oanh (kimoanh@ait.ac.th)

Abstract. This is a part of a research addressing the potential co-benefits associated with selected black carbon (BC) emission reduction measures on mitigation of air pollution and climate forcing in Southeast Asia (SEA). This paper presents details on the emission inventory (EI) results and the WRF/CHIMERE model performance evaluation. The SEA regional emissions for 2007 were updated with our EI results for Indonesia, Thailand and Cambodia and used for the model input. WRF/CHIMERE simulated PM_{10}, $PM_{2.5}$ and BC over the SEA domain (0.25º x 0.25º) of the year 2007 and the results were evaluated against the available monitoring data in the domain. WRF hourly simulation results were evaluated using the observed data at 8 international airport stations in 5 SEA countries and showed a satisfactory performance. WRF/CHIMERE results for PM_{10} and $PM_{2.5}$ showed strong seasonal influence of biomass open burning while BC distribution showed the influence of urban activities in big SEA cities. Daily average PM_{10} constructed from the hourly concentrations were obtained from the automatic monitoring stations in three SEA large cities, i.e. Bangkok, Kuala Lumpur and Surabaya for model evaluation. The daily observed $PM_{2.5}$ and BC concentrations obtained from the Improving Air Quality in the Asian Developing Countries (AIRPET) project for 4 cities (i.e. Bangkok, Hanoi, Bandung, and Manila) were also used for model evaluation. In addition, hourly BC concentrations were taken from the measurement results of the Asian Pacific Network (APN) project at a sub-urban site in Bangkok. The modeled PM_{10} and BC satisfactorily met all suggested statistical criteria for PM evaluation. The modeled $PM_{2.5}/PM_{10}$ ratios estimated for four AIRPET sites ranged between 0.47 - 0.59, lower than observed values of 0.6 - 0.83. Better agreement was found for $BC/PM_{2.5}$ ratios with the modeled values of 0.05-0.33 as compared to the observation values of 0.05 – 0.28. AODEM (extended aerosol optical depth module) was used to calculate the total columnar aerosol optical depth (AOD) and BC AOD up to the top of the domain at 500 hPa (~5,500 m) which did not include the free tropospheric long-range transport of the pollution. The model AOD results calculated using the internal mixing assumption, were evaluated against the observed AOD by both AERONET and MODIS satellite in 10 countries in the domain. Our model results showed that the BC AOD contributed 7.5 – 12% of the total AOD, which was in the same ranges reported by other studies for places with intensive emissions. The results of this paper are used to calculate the regional aerosol direct radiative forcing under different emission reduction scenarios to explore potential co-benefits for air quality improvement, reduction in number of premature deaths and climate forcing mitigation in SEA in 2030 (Permadi et al., 2017a).

1 Introduction

The Southeast Asia (SEA), with a large population and fast-growing economy, is an important contributor to the emissions of air pollution and greenhouse gases in Asia (Streets et al., 2003; Zhang et al., 2009). The emissions of anthropogenic aerosol from Asia and specifically from SEA, are expected to rise in the near future due to the increase in the energy demand and rapid industrialization (Lawrence and Lelieveld, 2010; Ohara et al., 2007). High levels of fine particulate matter (PM with diameter less than 2.5 micron or $PM_{2.5}$), the most detrimental air pollutant from the health effect point (Janssen et al., 2011; WHO, 2012), are observed in many developing Asian cities with the annual average often exceeding the WHO guideline of 10 μg m^{-3} by many times (Kim Oanh et al., 2006; Hopke et al., 2008). Being components of PM, e.g. $PM_{2.5}$ and PM_{10} (PM with diameter less than 10 micron), black carbon (BC) and organic carbon (OC), have been monitored in some Asian cities and the results, although fragmented, showed considerably high levels (Kondo et al., 2009; Kim Oanh et al., 2006; Hopke et al., 2008). The fine particles and their precursors are also involved in the long-range transport (LRT) hence causing the regional phenomenon such as Atmospheric Brown Clouds (ABC) (UNEP and C^4, 2002; Ramanathan et. al., 2001) and can affect the climate (UNEP-WMO, 2011). Globally, measures aim to reduce emissions of BC (and co-emitting pollutants) were shown to reduce the number of premature deaths and slow down the near future temperature increase and more benefits to be gained in Asia, where current emissions are high (UNEP-WMO, 2011; Shindell et al., 2012).

To comprehensively assess the co-benefits of emission reduction measures at regional scales, finer temporal and spatial resolutions of the modeling results are required. Several studies have been conducted for various Asian domains using the regional climate model (RCM) with chemistry (Nair et al., 2012) or chemical transport models (CTMs) with an additional aerosol optical module. Most of the Asian regional modeling studies mainly focused on the domains of East (Han et al., 2011; Park et al., 2011; Chen et al., 2013; Zhang et al., 2016)), South (Goto et al., 2011) and Continental East-Southeast Asia (Lin et al., 2014). These studies also highlighted several challenges for models to reproduce the ground-observed PM due to inaccurate emission inventory (EI), simulated meteorological fields and the extent of model representations (e.g., secondary organic aerosol formation, gas/particle partitioning, dry and wet deposition).

There is not yet detail modeling studies conducted for the SEA domain, especially the maritime SEA which includes Indonesia with its large biomass open burning (OB) emissions. For such a modeling effort, first reasonably accurate regional EI database should be prepared to generate input data. Several global and regional EI databases are available which also cover the SEA domain. These datasets have been developed using the activity data taken from several international data sources (Zhang et al., 2009; EC-JRC/PBL, 2010) or based on the results of large scale energy model (Streets et al., 2003). Efforts therefore should be put forward to update the SEA EI databases to generate the emission input data for the SEA regional modeling studies.

Our research used integrated EI and modeling tool to provide the spatial and seasonal distributions of aerosol species (PM_{10}, $PM_{2.5}$, and BC) in SEA for 2007 and the co-benefits (on air quality, health and climate forcing) of selected emission reduction measures for 2030. This paper presents the SEA emissions for the base year of 2007 and the WRF-

CHIMERE performance evaluation. CHIMERE (Menut et al., 2013 and references therein) was used to simulate three-dimensional (3D) aerosol concentrations in the domain using the meteorological fields generated by the Weather Research and Forecasting (WRF) model (Michalakes et al., 2004). The model results were evaluated using available ground-based measurements of PM_{10}, $PM_{2.5}$ and BC in several SEA cities. The extended aerosol optical depth (AOD) module (AODEM), detailed in Landi and Curci (2011), was applied to calculate the total columnar AOD and BC AOD. The modeled total AOD was evaluated using the observed AOD from both ground based Aerosol Robotic Network (AERONET) and the Moderate Resolution Imaging Spectroradiometer (MODIS) satellite product. The results are used in the follow up study which investigated the potential co-benefits of various emission reduction measures implemented in Indonesia and Thailand on air quality improvement, number of premature death reduction and climate forcing mitigation in 2030 (Permadi et al., 2017a).

2 Methodology

2.1 Emission inventory and emission input data

The emissions from major anthropogenic sources in Indonesia, Thailand and Cambodia were developed following the EI framework given in the "Atmospheric Brown Cloud Emission Inventory Manual (ABC EIM)" (Shrestha et al., 2013) using the activity data summarized in Table 1. Detail EI methodology for Indonesia was presented in Permadi et al. (2017b).

The biomass OB categories considered in this study included crop residue open burning (CROB) and forest fires (above-ground forest fires and peatland fires). The CROB emission (aerosol and trace gases) for Thailand for 2007 was taken from Kanabkaew and Kim Oanh (2011), and both CROB and above-ground forest fire emissions for Indonesia were from Permadi and Kim Oanh (2013) also for 2007. For Cambodia, CROB emission for 2007 was also included in the inventory (Permadi, 2013) but forest fire emissions of the country were taken from the Global Fire Emission Database v3 (GFEDv3) database (van der Werf et al., 2010). CROB emissions for Thailand, Indonesia and Cambodia were estimated from crop production statistics, residue to crop ratio, dry matter to crop residue ratio, fraction of biomass burned in the field, combustion efficiency, and EFs. The emission results were higher than other databases that used the MODIS product (e.g. GFEDv3) because the satellite may not efficiently capture short-lived, small size and sporadic CROB fires (Permadi and Kim Oanh, 2013). For other countries in the domain, the emissions from the above-ground forest fires were from Song et al. (2009) while those from CROB were from GFEDv3 for the base year of 2007. The 2007 emissions from peatland fires of all countries in the SEA domain were also taken from GFEDv3 (van der Werf et al., 2010). The GFEDv3 database was developed using a combination of MODIS burned area and active fires which is believed to better detect the peatland fires than those used the MODIS burn scars area product of MCD45A1 for forest fires detection as quoted by Shi et al. (2014).

Biogenic emissions were calculated on-line in the CHIMERE model using the methodology described in Simpson et al. (1999) that considers seasons and vegetation cover types taken from the Global Land Cover Facility (GLCF) (http://glcf.umd.edu) with a resolution of 1 x 1 km^2. CHIMERE incorporates the Model of Emissions of Gases and Aerosol from Nature (MEGAN) module (Guenther et al., 1995) for estimation of VOC from natural vegetation and

NO emission from soil. We did not include the emission from the international shipping in this simulation however the "inland waterway" sources were included in the inventories for the three countries. Other sources of PM such as un-paved road and wind-blown dust emissions were also not included in this study.

Spatial distribution was made based on source activity data or relevant proxies collected for the administrative boundaries. For example, for Indonesia, the emissions were disaggregated into district level (300 districts) while for Thailand and Cambodia, emissions were presented for provincial level (76 and 24 provinces, respectively). Further the emissions were gridded into 0.25° x 0.25° using the Geographic Information System (GIS) tool. For other countries in SEA, the emissions of SO_2, NOx, CO, VOC, PM_{10}, $PM_{2.5}$, BC and OC were taken from the available online gridded EI databases (grid size of 0.5° ~ 50 km) compiled by the Center for Global and Regional Environmental Research (CGRER) (Zhang et al., 2009). The gridded CH_4 and NH_3 emissions that were not included in CGRER, were taken from the global Emission Database for Global Atmospheric Research (EDGAR) (EC-JRC/PBL, 2010), with a grid resolution of 10 x 10 km^2. Emissions from all considered sources were compiled for the base year of 2007 and were gridded to 0.25° x 0.25° (~ 30 x 30 km^2) for the modeling input.

Monthly based activity data were obtained whenever available to construct monthly emission, however when the data were not available relevant proxies were used. Specific country hourly variations of emissions were extracted from the available studies in SEA. For Indonesia, detail methodology and data sources to construct monthly and hourly emissions of major anthropogenic sources were detailed in Permadi et al. (2017b). For above-ground forest vegetation, crop residue and municipal solid waste OB emissions, the monthly and hourly profiles were adopted from Permadi and Kim Oanh (2013). For Thailand, the hourly profiles for power plant and industry were obtained from Pham et al. (2008) while that for other major anthropogenic sources were taken from Vongmahadlek et al. (2009). The survey-based information on hourly profiles reported in Kanabkaew and Kim Oanh (2011) was used for CROB emission. For other countries, relevant data from regional EI from Streets et al. (2003) were utilized. The hourly emissions for the domain were prepared using a Fortran program developed at the Asian Institute of Technology (AIT) for this purpose.

The lumping of Non-Methane Volatile Organic Compounds (NMVOCs) emissions to the model species was done according to the MELCHIOR mechanism (Middleton et al., 1990). The aggregation produced the emissions of 33 species including trace gases and aerosol in unit of mole cm^{-2} s^{-1}. Aerosol fluxes were also converted to the "molecule-like" units in the emission input data using a fictive molar mass equal to 100 g $mole^{-1}$ (Bessagnet et al., 2004).

2.2 Modeling domain

The choice of domain size and resolution affects the balance between the boundary and internal modeling forcing in the simulated concentrations (Seth and Giorgi, 1998). For this study, it is important that the defined domain allows the transport of air pollutants by the monsoon circulation across SEA. Therefore, we set the domain to cover as much as possible the major upwind emission sources and to capture meteorological processes in the region of interest.

The SEA domain horizontally covered 9 (nine) countries of the Association of Southeast Asian Nations (ASEAN) and 3 provinces of southern China (Fig. S1, supplementary information, SI). The WRF domain extended from middle part

of Myanmar to the northern part of Australia covering 230 x 200 grids. The CHIMERE domain extended from southern China (24ºN, 95ºE) to eastern parts of Indonesia (9ºS, 137ºE) consisting of 169 x 133 grids. The grid resolution of the WRF and CHIMERE was set to be the same, 0.25° x 0.25° (~30 x 30 km^2).

2.3 WRF and CHIMERE model configuration

5　WRF version 3.3 was used with lateral boundaries and initial meteorological conditions taken from the National Centers for Environmental Prediction (NCEP) final (FNL) global analyses that are available at 1º x 1º grid resolution for every six hours (http://rda.ucar.edu/datasets/ds083.2/). WRF Pre-processing System (WPS) of geographical input data (i.e. land use, vegetation index, soil type, and albedo) was also obtained from the NCEP database. Totally, 28 vertical levels were simulated with the lowest level having the physical height of about 38 m. Analysis nudging was

10　performed in the Planetary Boundary Layer (PBL) and other layers for wind components (u and v), temperature (T) and relative humidity (RH). Nudging coefficients were set for all parameters at 0.00005 s^{-1}. The time interval between analyses was set at 360 minutes, which is equivalent to 6-hourly boundary input data used in our study. This analysis nudging was performed as it is suitable for a coarse resolution simulations (30 x 30km^2) to drive regional air quality models as it can improve the accuracy for the downscaled/nested fields (Dudhia, 2012; Bowden et al., 2012). Note

15　that, due to the insufficiency of spatially distributed meteorological observations in the domain, the observation nudging was not performed.

In the WRF simulation, the following physics options were used: simple ice microphysics, Unified Noah land-surface model for land-surface scheme, Rapid Radiative Transfer Model (RRTM) and Dudhia schemes for long and short wave radiation, PBL parameterization scheme of Yonsei University (YU), and Kain-Fritsch (KF) scheme with deep

20　and shallow convection option for cumulus parameterization. These schemes were selected as they are suitable for mesoscale grid size and have been used in the previous studies in the world (Jankov et al., 2005; Osuri et al., 2012).

This study used CHIMERE version 2008c with the MELCHIOR 2 chemical mechanism that was adapted from the original European Monitoring and Evaluation Programme (EMEP) and consisted of around 120 reactions and 40 chemical species. The vertical profiles of updated reaction rates in MELCHIOR 2 have been developed using tabulated

25　clear-sky photolysis rates taken from the Tropospheric Ultraviolet and Visible (TUV) model (EC4MACS, 2012). This version of CHIMERE has an aerosol module which consists of the total primary PM emission (BC, OC and other primary particles) and secondary inorganic PM species, such as nitrate, sulfate and ammonium, and secondary organic aerosol (SOA) (Bessagnet et al., 2004). CHIMERE applies the sectional approach to discretize particle size distribution into a finite number of bins. The considered particles size range was from 40 nm to 10 μm that are

30　distributed into 8 bins (0.039, 0.078, 0.156, 0.312, 0.625, 1.25, 2.5, 5, 10 μm) (Pere et al., 2011). Most of aerosol-related dynamic processes, such as condensation, coagulation, wet and dry deposition, adsorption and scavenging, are incorporated in the model (http://www.lmd.polytechnique.fr/chimere/). This version of CHIMERE only allows tropospheric simulations below 200 hPa (~12 km).

We used 8 vertical layers in this study, from sigma level 0.999 (~20 m) to the sigma level ~0.5 (~5,500 m), equivalent to the 500 hPa pressure level. This upper limit was selected based on a suggestion that in the modeling of anthropogenic pollution, extending the vertical dimension beyond 500 hPa would not substantially change the modeled aerosol concentrations for the ground level (Menut et al., 2013). However, it is recognized that the top of the domain at 500 hPa may not be able to include the free tropospheric LRT of the pollution and it brings in uncertainty to the total column AOD results. Note that in the CHIMERE version used in this study, the photolysis rates are calculated under clear sky conditions as a function of height using the TUV model (Madronich et al., 1998) and photolysis rates are estimated only up to 9,000 m. However, with the present formulation for cloud-radiation-photolysis, assuming that model domain is below the cloud, hence no cloud albedo was taken into account and this puts constraints on the model top of 500 hPa (IPSL/INERIS, 2008). Monthly mean boundary conditions of gases and aerosol were taken the simulation results for a period of 1998-2002 by the Laboratoire de Météorologie Dynamique (LMDZ) - Interaction avec la Chimie et les Aérosols (INCA) (Schulz et al. 2006) which are available at the CHIMERE website. To assess the effects of the somewhat aged boundary conditions compared to the model year, a comparison between the monthly average concentrations at the boundaries between 2007 and 1998-2002 was made which showed that the difference among the datasets was differing by 0.98 – 1.23 considering the ratios for aerosol and PM precursor gases (i.e. BC, OC, NO_2, CO, SO_2, C_2H_4, CH_3CHO, and NH_3) between the two datasets. This implies that basically the two datasets were almost similar. The impacts of the aged boundary conditions on the simulation are expected but with a small magnitude. Initial conditions of gases and aerosol concentrations in every grid were interpolated from the outputs of the global CTM of LMDZ-INCA simulation. One year simulation (January 1–December 31 2007) was performed by both WRF and CHIMERE with a spin up period of one week prior to the main simulation period.

2.4 Aerosol optical depth calculation

A stand-alone post processing tool, namely AODEM, developed by Istituto di Scienze dell'Atmosfera e del Clima - Consiglio Nazionale delle Ricerche (ISAC-CNR) of Italy (Landi and Curci, 2011) was used to calculate optical parameters of AOD (extinction coefficients and single scattering albedo) using the 3D aerosol species mass concentration fields output of WRF/CHIMERE for different size bins. AODEM calculates 3D particle number concentrations from these mass concentrations and provides the extinction coefficients for each grid-cell assuming the spherical shape of particles (http://people.isac.cnr.it/~landi/PhD.html). Three options of the aerosol mixing state were provided in AODEM: external, internal homogeneous and internal coated spheres. Aerosol optical properties are simulated by AODEM following the Mie theory (Bohren and Huffman, 1983) for the wavelength range from 340 nm to 1,640 nm. We selected the "aerosol internal mixing" option in the calculation because existing field measurements confirmed that aerosol is typically found in the internally mixed state (Lesins et al., 2002) largely due to coagulation and growth of aerosol particles (Jacobson, 2000). Note that, AOD was calculated from the surface to the model top layer of 500 hPa hence it could not represent the transport through the convective processes taking place above the model top layer as well as the LRT in the free troposphere mentioned above.

For calculation of optical aerosol properties, AODEM provides the particle number concentrations separately for five components: BC, OC, sea salt, dust, and secondary inorganics (nitrate, sulfate and ammonium). The AOD scattering

was simulated using "brute force" by excluding BC in the simulation (Landi and Curci, 2011). BC AOD was calculated by substracting the AOD scattering from the total AOD.

2.5 Model evaluation

The evaluation of WRF outputs was done using observed data from eight (8) airport meteorological stations in 5 SEA countries that captured major sub-climate zones (upper, near-equator and lower latitude) in the domain. Hourly observations from all these airport stations in 2007 were obtained from http://weather.uwyo.edu/surface/meteorogram/. The statistical evaluation of WRF outputs was done using the criteria provided by Emery et al. (2001) which includes the mean bias error (MBE), mean absolute gross error (MAGE) and root mean squared error (RMSE). The modeled surface pressure and upper wind fields (at 850 hPa) were compared with the European Centre for Medium-Range Weather Forecasts (ECMWF) Reanalysis (ERA) Interim data extracted from http://apps.ecmwf.int/datasets/ for the same vertical levels. Simulated monthly accumulated precipitation fields were compared with the satellite-based observations provided by the Tropical Rainfall Measuring Mission (TRMM-3B43) (https://disc.sci.gsfc.nasa.gov/datasets/TRMM_3B43_V7/summary?keywords=TRMM).

Only limited air pollution data was available in SEA for the model performance evaluation. This study collected the observed concentrations of aerosol (BC, OC, $PM_{2.5}$ and PM_{10}) and related gases from various sources. For examples, daily (24h) concentrations of PM_{10}, $PM_{2.5}$, BC, and OC in four SEA cities (i.e. Manila, Hanoi, Bandung, and Bangkok) in 2007 were taken from the measurement data generated by the AIRPET project "Improving Air Quality in Asian Developing Countries" (Kim Oanh et al., 2006; Kim Oanh et al., 2014). Hourly BC and OC concentrations were taken from the measurement results of the Asian Pacific Network (APN) project at the AIT located in Pathumthani province of the Bangkok Metropolitan Region, Thailand (Kondo et al., 2009). Hourly PM_{10} in Bangkok (Thailand), Kuala Lumpur (Malaysia) and Surabaya (Indonesia) in 2007 were collected from the respective national monitoring networks. The statistical evaluation of simulated aerosol levels was done using mean fractional bias (MFB) and mean fractional error (MFE) (Boylan and Russel, 2006). Definitions of the statistical measures used in the model performance evaluation are given in Table S1, SI.

The monthly AERONET data for 2007 was downloaded from the National Aeronautics and Space Administration (NASA) website (http://aeronet.gsfc.nasa.gov/) for the evaluation of the modeled AOD The AERONET data was of level 2 quality controlled, recorded at 10 AERONET stations (using Sun Photometer) listed in Table S2, SI. This provided AERONET dataset that has already been pre- and post-field calibrated with cloud screening and quality assurance. The selected 10 AERONET stations had more complete datasets in 2007 and they represent all sub-climate zones in the domain. Sun Photometer measures AERONET AOD at six different wavelengths (1020, 870, 675, 500, 440 and 380 nm). Therefore, to compare with the modeled AOD at 550 nm, the AERONET AOD at 500 nm were converted to that 550 nm using a logarithmic interpolation (Chung et al., 2012).

For a qualitative evaluation of the spatial distributions of the modeled AOD, the satellite MODIS Terra and MODIS Aqua AOD were used. The consistency between the modeled AOD spatial distribution over the SEA domain for

7

monthly MODIS AOD (level-3 data measured at 550 nm wavelength downloaded from https://giovanni.sci.gsfc.nasa.gov/giovanni/#service=TmAvMp&starttime=&endtime=&variableFacets=dataFieldM easurement%3AAerosol%20Optical%20Depth%3B).

3   Results and Discussion

3.1 Base year emissions

The obtained total national emission estimates of Indonesia, Thailand and Cambodia for 2007 were compared with the existing regional EI databases of EDGAR for 2007 and CGRER for 2006. Table 2 shows a reasonable agreement in the ranges of the estimates between the emission databases for three countries, Cambodia, Indonesia and Thailand. Detail EI results for Indonesia was presented in Permadi et al. (2017b). There are certain discrepancies between the databases that may be explained by several factors, including the uncertainty in activity data levels and EFs used, as well as the different coverage of the emission sources by different EI works. Specifically, for the emission sources of N_2O, our EI for the three (3) countries did not cover the direct emissions from cultivated soil (fertilized land) and the indirect N_2O emissions from agriculture-related activities (microbial nitrification and denitrification) hence resulting in lower N_2O emission estimates. Similar reason may be used to explain our lower estimate for CH_4 as compared to EDGAR for all 3 countries.

For Indonesia, our emission estimates were in between those of CGRER and EDGAR for a number of species. The estimates for PM_{10}, $PM_{2.5}$ and BC actually agreed well between these databases while OC of CGRER appeared to be higher. However, the SO_2 estimates differed a lot between the databases and our value was lower than others (mainly for on-road transportation and industry) which may be attributed to a more bottom up approach used in our EI that relied on actual S content used in the country and implementation of air pollution control devices (Permadi et al., 2017b). For example, our SO_2 estimate for the power plants of Indonesia was 300 Gg y^{-1} which was more comparable with the CGRER estimate (409 Gg y^{-1}) but much lower than the EDGAR estimate (1,000 Gg y^{-1}). The most striking difference was for the CO_2 emission that showed a much higher value by EDGAR, and this could be clearly explained by the inclusion of two major sources in the EDGAR dataset: i) the forest fire post burn decay (698,000 Gg y^{-1}) and ii) decay of drained peatland (504,000 Gg y^{-1}). If these two emission sources are excluded from the EDGAR results, the CO_2 estimates of all 3 databases are similar for Indonesia. The EI results for Thailand and Cambodia also showed reasonable agreements between the available datasets, except for CO_2 which was estimated higher by EDGAR for Cambodia. The BC and OC emissions for these two countries were mostly comparable with the results differed by a factor less than 2.0 among the datasets.

The emission shares by source category for the three countries are presented in Fig. S2, SI. The emissions of aerosol species (PM_{10}, $PM_{2.5}$, BC and OC) were mainly from the residential and commercial combustion in Indonesia (43-80%) and Cambodia (55-78%) while for Thailand the biomass OB (forest fire and crop residue) emissions were dominant, i.e. 31-74%. For SO_2, the emission in Indonesia was mainly contributed by the transport sector (36%) and thermal power plants (33%) but the industry was the main contributor in both Thailand (66%) and Cambodia (33%).

8

For NOx, the total emission in Indonesia was dominated by the fugitive emissions from oil and gas operation (44%), in Thailand by power plant (34%) and in Cambodia by forest fires (60%). The total emission of NH_3, an important precursor for $PM_{2.5}$, in all three countries was mainly from the manure management and fertilizer application (others), i.e. 63% for Indonesia, 75% for Thailand and 78% for Cambodia.

The emissions from other SEA countries and from the non-SEA part (southern part of China) of the domain used in our modeling study are also included in Table 2. The emissions from the southern China had high shares in the total emissions from the modeling domain. It is seen that Indonesia and Thailand were collectively the largest emitters of all pollutants, sharing of 25-66% of 2007 SEA emissions and 17-44% of the modeling domain emissions. Thus, emission reduction measures implemented for these 2 countries are expected to contribute remarkably to the air quality improvement in the region which will be analyzed in the companying paper (Permadi et al., 2017a). The spatial distributions of the annual average emissions of BC and CO at 0.25° x 0.25° (~30 x 30 km^2) resolution over are presented in Fig. 1 that showed higher emission intensity over large urban areas in the domain.

3.2 WRF model results and evaluation

3.2.1 Model statistical performance evaluation

The WRF hourly outputs, including surface temperature (T), relative humidity (RH) and wind speed (WS) for 2007 were compared with the observed data at 8 international meteorological stations in 5 SEA countries (Table 3). The comparison was done for two seasons, 3 months, 1 January – 31 March, to represent the dry season in the continental SEA (but the wet season in Indonesia) and 3 months, 1 August – 31 October, to represent the wet season in the continental SEA (but the dry season in Indonesia). The time series of daily average modeled vs. observed meteorological parameters, as shown in Figs. S3a-S3b, SI, showed that the model appeared to reasonably reproduce all parameters for the considered stations. In general, the model performance for temperature and WS simulations at all the stations was better than for RH during both periods.

The statistical performance evaluation for the hourly simulated values, against the MBE, MAGE and RMSE criteria is given in Table 3. MBE for the January-March period range was -1.9 – +0.7 °C for T, -0.3 – +2.7 m s^{-1} for WS, and -7.1 – +7.7% for RH. The corresponding range obtained for the August-October period was -0.1 – +2.3 °C, -0.6 - +2.1 m s^{-1} and -5.4 – +2.6%. Other statistical measures of MAGE and RMSE varied between the stations and the deviations from the suggested criteria were generally small. This suggested a relatively good model performance of WRF for both dry and rainy seasons. Overall, for the stations located in the north latitudes (above the equator line), the model performed better in the wet season (August – October) while for those located near and lower than the equator line the model performance was equally good for both dry and wet seasons. The discrepancy between model results and observations was perhaps partly due to the fact that the domain covers some regions, such as the Indonesian maritime – continent, that are principally characterized by active convection with a frequent presence of deep convection. These local processes, e.g. deep convection, are difficult to simulate using the mesoscale meteorological model of WRF with a rather coarse resolution (0.25º ~ 30 x 30 km^2) used in this SEA modeling study. Therefore, finer resolutions are

required to capture the dynamical processes undergoing at smaller scales. Different physics options may be required for sub-region domains to capture the processes and this should be done in future studies. In addition, a certain discrepancy is always expected because the model provided a grid average value, i.e. one for a 30 x 30 km^2 cell, while the observation is point based at individual stations.

3.2.2 Synoptic scale model evaluation

Spatial distribution of surface pressure over the WRF domain is presented together with the ERA interim dataset in Fig. S4 for three selected days (Jan 1, 2013; Oct 8, 2007; and Nov 7, 2007; 00:00 UTC). Both modeled and ERA data showed similar spatial distribution patterns of pressure but WRF appeared to produce slightly lower surface pressure over central Papua of Indonesia for all 3 cases presented. In fact, both datasets showed lower pressure zones over the high mountain areas of Himalaya, eastern parts of China and central Papua of Indonesia that indicated the effects of the topography.

The simulated wind fields at 850 hPa (~ 1,500 m) are compared with the ERA interim upper wind fields in Fig. S5 that also showed a consistency of the two datasets and more in the center of the domain both for wind speeds and wind directions. A large discrepancy was seen at the NW corner of the modeling domain and this may be attributed to the boundary conditions (taken from NCEP FNL in this study). The modeled monthly precipitations for two selected months (August and October, 2007) were compared with the TRMM-3B43 dataset in Fig. 2 which showed good agreement in the distribution patterns but the model somehow underestimated the domain maximum monthly precipitation column, for example, that occurred over Myanmar in August 2007 or over the central part of Vietnam in October.

The domain maximum hourly values of simulated PBL in different months of 2007 (Fig. S6) showed the PBL of 1,800 – 3,900 m. The maximum value of 3,900 m occurred in March, which was lower than the model top level of 500 hPa (~5,500 m) mentioned above, while the lowest PBL was in August.

3.3   CHIMERE model results and evaluation

Aerosol simulation always presents a big challenge due to the complex multiphase chemistry and transport processes. Lack of ground monitoring data of aerosol in the SEA region is an obstacle to a comprehensive model performance evaluation. For model performance evaluation, the CHIMERE results of PM_{10}, $PM_{2.5,}$ BC and the ratios of $PM_{2.5}/PM_{10}$ and BC/PM are discussed when comparing with available observed data in the domain in 2007.

3.3.1 PM_{10}

The daily (24h) modeled PM_{10} concentrations were estimated using the hourly data and the results were compared with the data gathered from the governmental monitoring networks that are available in three big cities of SEA, (i.e. 1 station in Kuala Lumpur, 2 stations in Bangkok, and 1 station in Surabaya). Note that the same two periods, as for WRF evaluation above, were used to represent dry and rainy season for both northern and southern parts of the equator.

Overall, model results ranged from near zero to 85 µg m^{-3} while the observations ranged from 5 to 90 µg m^{-3} at the three cities. The period average of modeled PM$_{10}$ in the three cities ranged from 21.7 – 29.2 µg m^{-3} while the corresponding observations ranged from 25.9 to 45.2 µg m^{-3} (Fig. 3).

Scatter plots of daily average observed and modeled values are presented in Fig. 3 showed that the model appeared to reasonably capture the range of 24h PM$_{10}$ in the cities but it showed non-linearity correlation. The model underestimated the low observed values at the Kuala Lumpur station (one station), i.e. the observed levels were 30-60 µg m^{-3} while the modeled were fluctuating from near zero to about 60 µg m^{-3}. A better agreement in the range of 24h PM$_{10}$ was shown for Surabaya, i.e. both were from 5 µg m^{-3} to 85 µg m^{-3}, but the linear correlation was still quite low. For Bangkok, the modeled 24h PM$_{10}$ ranged from 10 - 60 µg m^{-3} while the upper limit of the observed values was 90 µg m^{-3}. It is noted that although the ranges of the modeled 24h PM$_{10}$ were comparable with the observed ranges but the correlations were not clear for all three cities.

The reason for the discrepancy in the day to day variations between the modeled and observed 24h PM$_{10}$ values could be due attributed to the less accuracy of the temporal variations of the emission input data, as well as due to the coarse resolution of the model which may not be able to represent the weather variables in a convection-dominated climate. It is always challenging to compare the regional scale modeling results obtained for a coarse resolution (i.e. 30 x 30 km^2) with the point-based observations especially in complex mixed urban areas. Lack of systematic monitoring data for PM$_{10}$ in rural sites of the domain during the modeling periods prevented from making a more comprehensive model performance evaluation. The statistical evaluation showed that in all 3 cities, the MFB and MFE values for 24h PM$_{10}$ (totally 179 data points for each city) were within the suggested criteria (Table 4). The MFB values in Bangkok, Kuala Lumpur and Surabaya were -53%, -56% and -9%, respectively, i.e. lower than the criteria of ≤±60%. The MFE values in Bangkok, Kuala Lumpur and Surabaya were 55%, 56% and 18%, respectively, which were also well within the criteria of ≤+75%. The simulated monthly average of PM$_{10}$ in Kuala Lumpur and Bangkok, were consistently lower than the observed values in all months (Fig. 3) which should be expected in principle due to the grid averaging of the model results. For Surabaya, however, the model simulated monthly PM$_{10}$ values were higher than the observed during the period of January – March 2007 but lower than the observed for the period of August – October.

Overall, the discrepancy between the modeled and observed PM$_{10}$ and other parameters may be caused by several factors including the input fields of meteorology and emission data for the simulation. The WRF model evaluation presented above showed an acceptable performance (see section 3.2) but still with discrepancies with the observed data. More uncertainty however was expected from the EI input data. In addition, the uncertainty may arise from the monitoring data, especially when a large number of missing data points such as at the Surabaya, Indonesia. Overall, the simulation of urban areas would require more refine emission input data to capture the local emission sources, such as roads or industries, and this should be addressed in future studies.

11

3.3.2 PM$_{2.5}$

Only some fragmented measurement data of PM$_{2.5}$ was available in the domain in 2007 for the model evaluation (Fig. 4). This study used the 24h PM$_{2.5}$ data monitored in the SEA cities of Bandung, Bangkok, Hanoi and Manila, under the AIRPET project (Kim Oanh et al., 2006; Kim Oanh et al., 2014). The observation data were only available for some specific periods in 2007 at different sites hence the modeled results were extracted for the corresponding periods for comparison. The observed sites were the mixed sites which were influenced by typical emission sources in the respective cities. The AIT site, located of about 650 m away from a heavily travelled road, represented a sub-urban site with the influences of emissions from traffic and OB of rice straw (Kim Oanh et al. 2009). Thuong Dinh (TD) of Hanoi was a mixed urban site influenced by traffic and residential combustion among other sources (Hai and Kim Oanh, 2013). Both Tegalega (TG) located in Bandung, Indonesia and Manila observatory (MO) in Manila, Philippines were urban mixed sites with strong influence of traffic and other urban typical sources. The data therefore represents different periods of the year and different urban characteristic sites hence was not meant to compare the levels between the cities but only for model performance evaluation.

Overall, the available observed 24 PM$_{2.5}$ data in four AIRPET cities ranged from 4 to 120 µg m^{-3} while the modeled values for the same data periods ranged from 5 to 64 µg m^{-3}. The average levels of the observed PM$_{2.5}$ over all the data periods ranged from 35 to 43 µg m^{-3} as compared to the modeled, i.e. from 9.7 to 21 µgm^{-3}. Scatter plots of observed and modeled 24h PM$_{2.5}$ at four AIRPET stations (Fig. 4) clearly showed that the model underestimated 24h PM$_{2.5}$ in all stations. In the mixed polluted urban site in Bandung (TG), modeled 24h PM$_{2.5}$ was within a range of 11-33 µg m^{-3} while the observed were 27-69 µg m^{-3}. In the TD urban site in Hanoi (close to a busy road), the simulated 24h PM$_{2.5}$ were 5-64 µg m^{-3} as compared to the observed of 20-120 µg m^{-3}. In the mixed urban site of MO in Manila the simulated 24h PM$_{2.5}$ were 6-37 µg m^{-3} as compared to the observed range of 4-55 µg m^{-3}. As discussed above, the four selected AIRPET sites were located quite close to heavily travelled roads (although were not directly at roadside) hence the local traffic emissions could directly affect the monitored pollution levels. This may be an important reason for the discrepancy between the monitored levels and the simulated grid average values. In addition, the observed data points were quite limited for 2007 ($\leq$30 at each site) hence were not sufficient for the statistical model performance evaluation. The PM$_{2.5}$ monitoring efforts should be enhanced to characterize the pollution in SEA and also provide sufficient data points for the model evaluation.

3.3.3 Black carbon

For the model evaluation purpose, we used available measurements in the previous projects for SEA. The 24h BC measured by the optical method available at several SEA sites under the AIRPET project (Kim Oanh et al., 2014). The hourly-based EC (elemental carbon by a Sunset analyzer) measurements, available from the APN project (Kondo et al., 2009) for the AIT site (sub-urban) which were used to calculate 24h BC levels. EC was measured using thermal optical method while BC was measured using light absorption method (continuous soot monitoring system or COSMOS). The model performance evaluation was done using 24h BC data of both APN and AIRPET projects.

The APN hourly EC dataset for the AIT site was available for both dry and wet seasons, from March to December 2007. The hourly EC and hourly BC the measured simultaneously by the APN project at AIT were found to have a strong linear correlation (Kim Oanh et al., 2009). Therefore, we used the observed Sunset EC to compare with the modeled output of BC. This is because for the PM mass closure, EC seems to be better while BC is suitable for radiative transfer budget analysis (Gelencsér, 2004). Fig. 5 presents the time series of the modeled and observed 24h BC for the AIT site. The modeled 24h BC was from 1.0 to 10 $\mu g\ m^{-3}$ that is comparable with the observed range from 0.8 – 10 $\mu g\ m^{-3}$. However, correlation between the modeled and observed BC shown in the scatter plot was not clear. The discrepancy between the modeled and observed BC seen in the time series may principally be due to the gridded average of the model output as compared to the point-based measurement. Higher BC levels measured at the AIT site were contributed by multiple local sources, such as nearby highway traffic activity and biomass OB (of rice straw) that occurred more intensively during the dry season (December). However, these sources especially small-scale rice straw field burning activity may not be well represented spatially by the EI input data made for a large resolution (30 x 30 km^2). Three (3) statistical measures of MBE, MFB and MFE were considered for the model performance evaluation in the BC simulation at the AIT site (Table 4). The MFB and MFE values were -24% and 49%, respectively, which all meet the suggested criteria (for PM). The MBE value was -0.12 $\mu g\ m^{-3}$ for AIT site which showed that the model somewhat underestimated the observed BC values but there is no criteria MBE available for PM for comparison.

The 24h BC (optically) measured on the 24h $PM_{2.5}$ sampled filters collected in the same locations of $PM_{2.5}$ measurements in SEA under the AIRPET project (Kim Oanh et al., 2006; Kim Oanh et al., 2014) were compared with the 24h modeled BC extracted for the sites and dates of 2007. Fig. 6 shows that the modeled 24h BC were lower than the observed at all the sites. The ranges of observed values and the modeled values were in somewhat better agreement for the AIT site and MO Manila site than the other 2 sites. At AIT, the observed BC values was 1.3 – 3.4 $\mu g\ m^{-3}$ (January, February and May) were higher but quite comparable to the modeled range of 0.5 – 1.8 $\mu g\ m^{-3}$. At MO, the observed 24h BC was 7 – 13 $\mu g\ m^{-3}$ (January and February) was quite close to the modeled 24h BC of 4.2 - 13 $\mu g\ m^{-3}$. More discrepancies were found for the Bandung site with the observed 24h BC values ranged from 4.2 to 9.8 $\mu g\ m^{-3}$ (July 2007) as compared to the modeled values of 1.3 – 3.2 $\mu g\ m^{-3}$. Similarly, the observed BC values at the mixed site of TD, Hanoi ranged from 12 to 23 $\mu g\ m^{-3}$ (January 2007), much higher than the modeled values of 1-7 $\mu g\ m^{-3}$. The effects of local sources, especially traffic emissions, at the quoted sites should be a main cause of the discrepancies when compared to the grid average modeled BC with the observed values. The limited measurement data available prevented from a more comprehensive model performance evaluation. Note that due to the limited measurement data points, a statistical performance evaluation was not conducted for the BC simulation.

3.3.4 Ratios between fine and coarse PM, and between BC and PM

In fact, $PM_{2.5}$ mass is principally contributed by both local combustion sources and secondary particles formation by chemical reactions in the atmosphere. The gaseous precursors of NOx, SOx and VOCs for the $PM_{2.5}$ formation may be of both local and LRT origins. The coarse fraction ($PM_{10-2.5}$) would mainly consist of primary particles of the

13

geological origin (Chow et al., 1998), and these are mainly contributed by local sources of soil, road dust and also construction activities (Hai and Kim Oanh, 2013). Due to its formation process as well as the ability to participate in the regional transportation, the fine particles (PM$_{2.5}$) are more uniformly distributed in an urban area than the coarser particles. The PM$_{2.5}$/PM$_{10}$ ratios could provide some information of the dominance of local sources of PM$_{2.5}$. We

5   compare the PM$_{2.5}$/PM$_{10}$ ratios based on the modeled 24h PM$_{2.5}$ and 24h PM$_{10}$ (PM$_{10}$ = PM$_{2.5}$ + PM$_{10-2.5}$) with those computed from the observed PM data available at the four (4) AIRPET monitoring sites discussed above. Overall, the modeled PM$_{2.5}$/PM$_{10}$ ratios ranged from 0.47 to 0.59 while the observed values were higher, 0.6 - 0.83. More pronounced difference was for TD of Hanoi, i.e. 0.74 observed vs. 0.47 modeled, and for TG of Bandung, 0.83 observed vs. 0.55 modeled. Better agreements were obtained for MO of Manila, 0.61 observed vs. 0.47 modeled, and

10 the AIT site, 0.60 observed vs. 0.59 modeled. The urban mixed sites of TD in Hanoi and TG in Bandung located in the traffic areas hence higher contributions of the primary PM$_{2.5}$ emitted from traffic may be seen better for TD and TG sites to the total measured PM$_{10}$, as compared of MO in Manila and AIT sites. However, to evaluate the variations in the PM$_{2.5}$/PM$_{10}$ ratios, contributions of various sources of the coarse particles, such as road dust and construction dust, should be further analyzed. It is noted that the ratios used to compare with the model simulated values were all

15 derived from the observations made in large cities in SEA. Lack of the observation data in rural areas and remote sites presents an obstacle for more in-depth analysis of the model performance. The remote sites, with less influence of the local sources, could be valuable for the model performance evaluation, both for the PM mass concentrations and their ratios.

20 BC is emitted directly from the combustion sources with higher fractions in PM emitted from the diesel exhaust (Kim Oanh et al., 2010) and lower fractions from biomass OB (Kim Oanh et al., 2011). Hence the ratio of BC/PM$_{2.5}$, for example, can infer the contribution of the primary particles from these combustion activities. BC/PM$_{2.5}$ and BC/PM$_{10}$ ratios were calculated using the observed 24h data at four AIRPET sites. Modeled BC/PM$_{2.5}$ ratios ranged from 0.05 to 0.33 as compared to the observed ratios of 0.05 – 0.28. For BC/PM$_{10}$, the modeled values ranged from 0.03 to 0.16

25 while the observed values ranged from 0.034 to 0.17. Observed BC/PM$_{2.5}$ ratios were higher than the modeled values at TG of Bandung (0.16 vs. 0.1) and AIT (0.055 vs. 0.05) sites. In TD of Hanoi and MO of Manila, the observed ratios (0.22 and 0.23) were lower than the modeled (0.28 and 0.33). As for BC/PM$_{10}$, the observed ratios at three (3) AIRPET sites of TG, TD and AIT (0.13, 0.17, and 0.034) were higher than the modeled values (0.06, 0.13 and 0.03) while for MO of Manila the opposite was shown with a lower observed (0.14) as compared to the modeled (0.16) value. The

30 simulated BC/PM ratio was the highest in TD of Hanoi, 22% of PM$_{2.5}$ and 17% of PM$_{10}$, during the dry period of January – February 2007 which confirmed the strong influence of traffic emission at this site.

The lack of data for the areas outside the cities is an issue remained. Generally, we expect that PM$_{2.5}$ mass may be more uniform in an urban area, for example measurements conducted in several mountain areas in Asia showed high

35 PM$_{2.5}$ concentrations which were mainly due to the regional transport (Hang and Kim Oanh, 2014; Co et al., 2014) or local combustion sources (e.g. residential cooking, biomass OB) such as found in China (Liu et al., 2017). However,

the BC fraction of PM may vary a lot with much lower values in remote sites but the lack of the data would prevent from a more in-depth analysis.

As seen in the statistical model evaluation, a negative MB was obtained for PM_{10}, -3 to -17, and BC, -0.12 (BC) at all sites (not enough data for statistical evaluation of $PM_{2.5}$) which showed an underestimation of PM_{10} and BC concentrations by the model at all sites. This may be explained by the coarse resolution (30 x 30 km^2) of emission input data which could adequately represent the spatial distributions of local sources of a smaller scale such as road traffic. These local sources, for example road traffic and residential cooking, affect PM measured at all sites hence affecting the $PM_{2.5}/PM_{10}$ and BC/PM ratios. The road/soil dust emission contribute more to $PM_{10-2.5}$ hence lowering $PM_{2.5}/PM_{10}$ ratios in urban areas but this coarse fraction of PM emission was not included in our emission input file. In addition, the LRT pollution above the model top layer (> 500 hPa) may contribute to the pollution in the domain, more to $PM_{2.5}$ and BC than the coarse PM. The free tropospheric LRT of aerosol and high convective processes should be also considered by extending the vertical model set-up in future studies.

3.4 Spatial distribution of modeled monthly PM_{10}, $PM_{2.5}$ and BC

Spatial distributions of the modeled monthly average PM_{10}, $PM_{2.5}$ and BC are presented in Fig. 7 for January, August and November while those of the respective annual averages are presented in Fig. S7, SI. The highest monthly average concentrations of PM_{10} in January, August and November 2007 simulated in the domain (one value for the whole domain) were 69, 58, and 44 µg m^{-3} while corresponding values of $PM_{2.5}$ were 40, 37 and 27 µg m^{-3}, respectively. The simulated maximum monthly average BC concentration in the domain was higher in January (8.2 µg m^{-3}) as compared to August (7.8 µg m^{-3}) and November (5.9 µg m^{-3}).

The simulated highest hourly PM_{10} in the considered months of January, August and November 2007 were 325, 245 and 164 µg m^{-3}, respectively, while the $PM_{2.5}$ corresponding values were and 188, 150 and 99 µg m^{-3}. The highest values of simulated annual average in the domain for PM_{10} and $PM_{2.5}$ were 51 and 32 µg m^{-3}, respectively. The maximum simulated annual average in the domain for BC was 6 µg m^{-3}. A summary of the simulated pollutant levels in the domain is presented in Table S3, SI.

For all considered pollutants over the domain, higher concentrations were observed over East Java, Indonesia, particularly over Surabaya city, which show the effects of emission from residential and traffic in the city and surrounding satellite cities as well as the crop residue OB (Permadi and Kim Oanh, 2013; Permadi et al., 2017b). High concentrations were consistently observed in several places in Indonesia including Java Island, West Sumatera (Padang), and West Kalimantan (Pontianak), and over Bangkok, Thailand. Large hotspots but with lower concentrations were also observed over the Southern China, over Hanoi and Ho Chi Minh of Vietnam which can be largely explained by the influence of the local sources (Fig. 7).

The monsoon circulation plays an important role in transporting PM from the emission source regions to other parts of the domain. In the dry months, higher emissions of biomass OB are expected hence higher concentrations of PM should be seen in the region near and downwind sources. Accordingly, at the northern part of the domain, higher PM levels were found in January-April while at the southern part of the domain higher concentrations were found during the period of April-August. In January in the Northern hemisphere, NE monsoon transports pollutants from the source regions to the southwest direction, while in the Southern hemisphere part (Indonesia) the plume moved to the northeast/east direction. The opposite is seen in August and November months (Fig. 7). In August, at the Southern hemisphere the plumes of PM moved northwesterly and turned into northeasterly after reaching the equator line. The plumes of PM_{10} and $PM_{2.5}$ were converged in the South China Sea in January and November when the NE monsoon prevalent that brought PM pollution from the Southern part of China mainland to the South China Sea (Figs. S8a-d). WRF result showed no rainfall over the South China Sea during the particular period that may also contribute to the converged PM levels (Figs. S8e-f).

In August and November, the dry months in the Southern domain, the PM_{10} and $PM_{2.5}$ plumes showing the effects of biomass OB (crop residue and forest fire) emissions in Indonesia originated in Riau province (Sumatera Island) and western and southern parts of Borneo Island, were seen clearly moving northeast-ward. In the dry season month in northern domain of January, the plumes of PM_{10} and $PM_{2.5}$ intensified by biomass OB in the central and northern parts of Thailand were shown moving southwest-ward. BC plumes were generally seen originated from big cities in the domain showing a significant influence of the fossil fuel combustion emission, specifically traffic, and other urban activities for all months of the year. During the dry period, BC plumes from the areas that have intensive biomass OB emissions were not as clearly seen as the PM plumes and this maybe because biomass OB contributed more to OC rather than BC emissions.

Effects of precipitation on the PM levels were also seen, e.g. higher PM levels (Fig. 7) were simulated over Indochina in January, October and November as compared to August because the latter was a rainy month in this part of the domain, i.e. less biomass open burning and more wet removal in principle. The opposite was actually seen in the Southern part of modeling domain, e.g. above Indonesia, where lower PM levels were simulated in October (more rainy month in this part) than other months.

3.5   Aerosol optical depth

Both total AOD and BC AOD were considered for the model evaluation. The monthly average of the total columnar AOD (scattering and absorbing), at the wave length of 550 nm, was produced from the AODEM simulation for 2007. The simulated monthly AOD data was compared with the monthly Terra-MODIS AOD, also at 550 nm, retrieved from the NASA website. Fig. 8 showed that the modeled AOD was lower than the MODIS observed for example in January, the maximum AOD simulated for the Southern China part of the domain was about 0.36 as compared to the MODIS AOD of 0.42-0.58. In the same month, the modeled AOD values over the Java Island of Indonesia were 0.072-0.28 while the MODIS AOD was 0.26-0.42. In April, the model results over the Southern China was 0.25-0.75

16

while the observed MODIS AOD was 0.42-0.90. Near the border between Myanmar and Bangladesh (northwest corner of the domain), the modeled AOD and the observed MODIS AOD was similar 0.74-0.75. However, the modeled AOD values over the Java Island in April were higher, i.e. 0.02-1.0, than the observed MODIS AOD of 0.26-0.42. The simulated hourly maximum and monthly average PM_{10} and $PM_{2.5}$ concentrations over Java Island were the highest throughout the year in April which may be due to higher emissions from residential combustion and traffic during the big Moslem Festival in the country (Permadi et al., 2017b).

In October, a hotspot with the maximum AOD of 0.8 was observed by MODIS over Riau, Sumatra Island and Singapore that was well above the model result for the grid of 0.4. The model was also not able to capture AOD hotspots over the Southern China mainland in this month. The results for August and November both showed some significant underestimation of AOD as compared to the MODIS observed values. There are several reasons for these discrepancies, including the temporal and spatial inconsistency in the observed and modeled values used for comparison. For example, the Terra MODIS satellite daily passed a region for a particular time (i.e. 13:30) hence giving only a snapshot of the value while the model provided the hourly average for 13:00 – 14:00 hence there is certain inconsistency in the monthly averages derived from these two datasets. The discrepancy may come from the fact that in the simulation AOD was covering up to 500 hPa hence could not include aerosol in the upper layers as mentioned above. Different spatial resolutions of modeled AOD (30 x 30 km^2) and MODIS AOD (10 x 10 km^2) can be another reason. In addition, shipping emission and the natural sources of aerosol, such as wind-blown dust, were not included in our emission input data hence the model would produce lower AOD (as well as PM_{10}) values. Consistently with the PM results, the effects of precipitation on AOD were captured, i.e. higher in the dry months and lower in the wet months in the respective parts of the domain. Overall, this qualitative analysis of the modeled vs. MODIS AOD provided only some insight into the regional distributions. Further efforts to conduct more comprehensive model evaluation are still required.

Simulated monthly AOD values were also compared with the observed data retrieved from 10 AERONET stations located in the domain, i.e. in Vietnam, Singapore, Hong Kong, Taipei, Thailand and Indonesia, which also showed lower simulated AOD values than the AERONET observed (Fig. 9). The model appeared to capture better seasonal variability in Bac Lieu (Vietnam), Silpakorn University and Songkhla (Thailand) stations while in other stations model underestimated the AOD. Model seemed to be able to simulate well the monthly average AOD in Hok Sui station (Hong Kong) but only for the months of October, November and December 2007. The strong seasonal variation of aerosol in SEA, largely caused by the biomass OB and meteorological conditions, creates a huge challenge for models to reproduce. At Puspitek Serpong (Indonesia) where emissions of urban activities from the capital city of Jakarta would dominate, the high AOD in October was reasonably captured by the model. The seasonal variation in the emission input file for anthropogenic sources would need to be further refined to improve the situation. Better proxies should be used for transport, industry, residential combustion and thermal power generation which should reflect the actual variation in the monthly activity data of each sector.

17

The BC AOD (absorbing) was calculated as the difference between the total AOD (scattering + absorbing) and the scattering AOD following the same method presented in Landi and Curci (2011). The spatial distribution of monthly average BC AOD is presented in Fig. S9, SI. In January, the dispersion plumes of high BC AOD spreading over in the Southern China (maximum AOD of 0.027) and eastern part of Indonesia (maximum 0.018) which shared 7.5-10% of the total AOD of 0.36 and 0.18, respectively, in these areas. In April, the highest value of the modeled BC AOD was seen over Surabaya (East Java province, Indonesia) with the range of 0.051-0.078, followed by relatively high values over Hong Kong and Shenzhen of 0.06-0.069. The contributions of the BC AOD to the total AOD in Surabaya, Hong Kong and Bangladesh were 9% (of 0.89), 11% (of 0.6) and 12% (of 0.54), respectively.

In other months, the highest monthly average BC AOD was shown in different parts of the domain ranging between 0.015-0.027 while the total AOD were 0.18-0.36, hence the shares of BC AOD in the total AOD was 7.5-8.6%. Our BC AOD contributions to the total AOD were higher than the reported global average value of 3% (Reddy et al., 2005), but in the same range of those reported for different regions with intensive emission sources. The relative contribution of BC to total AOD has been reported to depend on wavelength, i.e. increasing with decreasing wavelength, and on the dominant emission sources. For example, measurements showed typical contributions of around 12% under the influence of natural dust (Chiapello et al., 1999) and around 5-12% if biomass OB is dominant (Eck et al., 1999; Dubovik et al., 2002). The modeled BC AOD serves as input to estimate BC direct radiative forcing of anthropogenic emissions for the SEA domain which will be analyzed in our companying paper (Permadi et al., 2017a).

4 Summary and conclusions

This study developed and evaluated the EI databases for Indonesia, Thailand and Cambodia for 2007. The results were compiled with the existing CGRER and EDGAR emission datasets to generate the emission input data of the entire SEA domain for regional WRF/CHIMERE modeling. Our EI results for the three countries were comparable to other existing databases and the differences are explained mainly by the differences in the sources covered by different EI works. The BC emissions were mainly from residential and commercial combustion in Indonesia (71%) and Cambodia (70%) but was dominated by biomass OB emissions in Thailand (31%).

The model performance for 2007 was evaluated using the hourly and daily observed data in the SEA domain. The WRF model outputs were in good agreement with the observed data at 8 international airport stations in Indonesia, Thailand, Vietnam, Cambodia and Philippines. The WRF/CHIMERE satisfactorily reproduced the aerosol species of PM_{10}, $PM_{2.5}$ and BC in terms of the spatial distributions and seasonal variations. The statistical evaluation was conducted for 24h PM_{10} and 24h BC which had sufficient observed data points for the analyses. The modeled 24h PM_{10} in three cities (Thailand, Malaysia, and Indonesia) had the MFB and MFE values met the suggested criteria. Similarly, the modeled 24h BC values met the MFB and MFE criteria for PM when compared the observed data at a sub-urban site in Thailand (AIT).

The PM$_{2.5}$/PM$_{10}$ ratios calculated from the modeled outputs were lower than those estimated from the observed data at four AIRPET sites and this would imply a necessity of further improvement of the PM speciation of the emission input data. The modeled BC/PM$_{2.5}$ ratios were in compatible range (0.05 - 0.33) with the observed values (0.05 – 0.28) and were lower in two sites (AIT and Bandung) but higher in the others (Hanoi and Manila). The modeled BC/PM$_{10}$ ratios ranged between 0.03 - 0.16 which were comparable to while the observed values range (0.034 - 0.17). Lack of systematic observed BC data prevented from a more comprehensive model performance evaluation. Nevertheless, further improvement of the EI for primary aerosol, especially the PM speciation of major sources, as well as inclusion of un-paved road and wind-blown dust emissions are highly required. Vertical model set-up should be extended beyond of 500 hPa (~5,500 m) in the future studies to better incorporate the free tropospheric LRT of aerosol.

The spatial distributions of the columnar total AOD estimated for the WRF/CHIMERE output PM concentrations using AODEM were comparable with the observed (MODIS and AERONET) in 2007. In particular, exclusion of the un-paved road and wind-blown dust emissions (coarse particles) from the emission input in this study was a reason for the discrepancy in the modeled and observed total AOD which may underestimate the coarse PM concentrations. The lower values of aerosol species simulated by the model were explained by the grid averaging effects: WRF/CHIMERE had a larger grid of 30 km, as compared to MODIS AOD of 10 km, while AERONET is actually point based. Therefore, the spatial distribution of local sources of a smaller size can not be captured well by WRF/CHIMERE.

The spatial distribution patterns of the modeled aerosol species in the domain may be explained by the intensive biomass OB emissions. The plumes of PM$_{10}$ and PM$_{2.5}$ originated from Sumatera and Borneo Island of Indonesia in August-November, and from Central and Northern Thailand during January-April, which coincided with the dry months in the respective areas with subsequent more biomass OB. Spatial distributions of BC showed the influence of the traffic emission and residential combustion in big SEA cities. Based on the model results, the contribution of BC AOD to total AOD in the domain was around 7.5 – 12% which is consistent with the literature reported values for intensive emission areas. Effects of precipitation were captured by the model that produced lower PM and AOD levels in the months with higher precipitation simulated.

The EI data and WRF/CHIMERE performance for 2007 were satisfactory in terms of reproduction of the key aerosol species in the domain. In the accompanying paper (Permadi et al., 2017a), we present the WRF/CHIMERE simulation results for PM and BC for the SEA domain in the business as usual emission scenario (BAU2030) and in the emission reduction scenario (RED2030) to quantify potential co-benefits on the air quality improvement, reducing number of premature deaths, and radiative forcing mitigation in Southeast Asia.

Acknowledgments

References

Bessagnet, B., Hodzic, A., Vautard, R., Beekmann, S., et al.: Aerosol modeling with CHIMERE—Preliminary evaluation at the continental scale, Atmospheric Environment, 38, 2803–2817. 2004.

Bohren, C.F. and Huffman, D.R.: Absorption and Scattering of Light by Small Particles, New York: Wiley, 1983.

Bowden, H.J., Otte, L., Nolte, C.G., Otte, M.J.: Examining interior grid nudging techniques using two-way nesting in the WRF model for regional climate modeling, Journal of Climate, 25, 2805-2823. 2012.

Boylan, J.W. and Russell, A.G.: PM and light extinction model performance metrics, goals, and criteria for three-dimensional air quality models, Atmospheric Environment, 40, 4946–4959, 2006.

Chen, T.F., Tsai, C.Y., Chang, K.H.: Performance evaluation of atmospheric particulate matter modeling for East Asia, Atmospheric Environment, 77, 365-375, 2013.

Chiapello, I., Bergametti, G., Chatenet, B., Dulac, F., Jankowiak, I., Liousse, C., Soares, E.S.: Contribution of the different aerosol species to the aerosol mass load and optical depth over the northeastern tropical Atlantic, Journal of Geophysical Research, 104 (D4), 4025–4035, 1999.

Chung, C.E., Lee, K., Muller, D.: Effect of internal mixture on black carbon radiative forcing, Tellus B, 64, 10925, 2012.

Chow, J.C., Watson, J.G., Lowenthal, D.H., Egami, R.T., Solomon, P.A., Thuillier, R.H., Magliano, K.L., Ranzieri, A.J.: Spatial and temporal variations of particulate precursor gases and photochemical reaction products during SJVAQS/AUSPEX ozone episodes. Atmos. Environ., 32(16), 2835-2844, 1998.

Co, H.X., Dung, N.T., Kim Oanh, N.T., Hang, N.T., Phuc, N.H., Le, H.A.: Levels and Composition of Ambient Particulate Matter at a Mountainous Rural Site in Northern Vietnam. Aerosol and Air Quality Research 14(7), 1917-1928, 2014.

Dubovik, O., Holben, B., Eck, T.F., Smirnov, A., Kaufman, Y.J., King, M.D., Tanre, D., Slutsker, I.: Variability of absorption and optical properties of key aerosol types observed in worldwide locations, J. Atmos. Sci, 59 (3), 590–608, 2002.

Dudhia, J.: WRF Four-dimensional data assimilation, Papers presented at the 13th Annual WRF Users' Workshop, Colorado, 24-26 June 2012, USA, 2012.

Eck, T.F., Holben, B.N., Reid, J.S., Dubovik, O., Smirnov, A., O'Neill, N.T., Slutsker, I., Kinne, S.: Wavelength dependence of the optical depth of biomass burning, urban, and desert dust aerosol, J. Geophys. Res, 104 (D24), 31333–31349, 1999.

EC4MACS.: Modeling methodology: The CHIMERE Atmospheric Model, European Consortium for Modelling of Air Pollution and Climate Strategies, INERIS, France, 2012.

EC-JRC/PBL.: European Commission, Joint Research Centre /Netherlands Environmental Assessment Agency. Emission Database for Global Atmospheric Research (EDGAR), release version 4.1. http://edgar.jrc.ec.europa.eu, 2010.

Emery, C.A., Tai, E., Yarwood, G.: Enhanced Meteorological Modeling and Performance Evaluation for Two Texas Ozone Episodes, Retrieved from http://www.tceq.state.tx.us/assets/public/implementation/air/am/contracts/reports/mm/EnhancedMetModelingAndPerformanceEvaluation.pdf, 2001.

Gelencsér, A.: Carbonaceous aerosol. Atmospheric and Oceanographic Sciences Library, Springer, 2004.

Guenther, A., Hewitt, C. N., Erickson, D., Fall, R., Geron, C., Graedel, T., Harley, P., Klinger, L., Lerdau, M., McKay, W. A., Pierce, T., Scholes, B., Steinbrecher, R., Tallamraju, R., Taylor, J., Zimmerman, P.: A global model of natural volatile organic compound emissions, J. Geophys. Res., 100(D5), 8873 – 8892, 1995.

Goto, D., Badarinath, K.V.S., Takemura, T., Nakajima, T.: Simulation of aerosol optical properties over a tropical urban site in India using a global model and its comparison with ground measurements, Annales Geophysicae, 29, 955-963, 2011.

Han, X., Zhang, M., Han, Z., Xin, J., Liu, X.: Simulation of aerosol direct radiative forcing with RAMS-CMAQ in East Asia, Atmospheric Environment, 45, 6576-6592, 2011.

Hai, C.D. and Kim Oanh, N.T.: Effects of local, regional meteorology and emission sources on mass and compositions of particulate matter in Hanoi, Atmospheric Environment, 78, 105-112, 2013.

Hang, N.T. and Kim Oanh, N.T.: Chemical characterization and sources apportionment of fine particulate pollution in a mining town of Vietnam. Atmospheric Research,145–146, 214-225, 2014.

Hopke, P.K., Cohen, D.D., Begum, B.A., Biswas, S.K., Ni, B., Pandit, G.G., Santoso, M., Chung, Y.S., Davy, P., et al.: Urban air quality in the Asian region, Science of the Total Environment, 404, 103–112, 2008.

Jacobson, M.Z.: A physically-based treatment of elemental carbon optics: Implications for global direct forcing of aerosol, Geophysical Research Letters, 27 (2), 217-220, 2000.

Jankov, I., Gallus, W., Segal, M., Shaw, B., Koch, S.E.: The impact of different WRF model physical parameterizations and their interactions on warm season MCS rainfall, Weather and Forecasting, 20, 2005.

Janssen, N.A.H., Hoek, G., Simic-Lawson, M., Fischer, P., van Bree, L., et al.: Black carbon as an additional indicator of the adverse health effects of airborne particles compared with PM_{10} and $PM_{2.5}$, Environmental Health Perspectives, 119 (12), 1691-1699, 2011.

Kanabkaew, T. and Kim Oanh, N.T.: Development of spatial and temporal emission inventory for crop residue field burning, Environmental Modeling and Assessment, 16(5), 453-464, 2011.

Kim Oanh, N.T., Upadhyay, N., Zhuang, Y.H, Hao, Z.P., Murthy, D.V.S, Lestari, P., Villarine, J.T., Chengchua, K., Co, H.X., Dung, N.T., Lindgren, E.S.: Particulate air pollution in six Asian cities: Spatial and temporal distributions, and associated sources, Atmospheric Environment, 40, 3367–3380, 2006.

Kim Oanh, N.T., Kondo, Y., Co, H.X.: Investigation on the Impacts of Urban-Rural Air Pollution on Air Quality and Climate in Southeast Asia, Final report for APN project: ARCP2007-07CMY-Oanh, Asia Pacific Network for Global Change Research, 2009.

Kim Oanh, N.T., Thiansathit, W., Bond, T.C., Subramanian, R., Winijkul, E., Paw-armart, I.: Compositional characterization of PM2.5 emitted from in-use diesel vehicles, Atmospheric Environment, 44(1), 15-22, 2010.

Kim Oanh, N.T., Ly, B.T., Tipayarom, D., Manandhar, B.R., Prapat, P., Simpson, C.D., Liu L.J.S.: Characterization of particulate matter emission from open burning of rice straw, Atmospheric Environment, 45(2), 493-502, 2011.

Kim Oanh, N.T., Pongkiatkul, P., Templonuevo Cruz, M., Nghiem, T.D., Philip, L., Zhuang, G., Lestari, P.: Monitoring and source apportionment for particulate matter pollution in six Asian Cities, Chapter 3 in Integrated Air Quality Management: Asian Case Studies, CRC Press, Boca Raton, Florida, USA., 2014.

Kondo, Y., Sahu, L., Kuwata, M., Miyazaki, Y., Takegawa, N., Moteki, N., Imaru, J., Han, S., Nakayama, T., Kim Oanh, N.T., Hu, M., Kim, Y.J., Kita, K.: Stabilization of the Mass Absorption Cross Section of Black Carbon for Filter-Based Absorption Photometry by the use of a Heated Inlet, Aerosol Science and Technology, 43, 741–756, 2009.

Landi, T.C. and Curci, G.: Modeling aerosol optical properties with AODEM: accounting for non-sphericity of dust particles, Geophysical Research Abstracts, 13, EGU2011-3958, 2011.

Lawrence, M.G. and Lelieveld, J.: Atmospheric pollutant outflow from Southern Asia: a review. Atmospheric Chemistry and Physics, 10, 11017 – 11096, 2010.

Lesins, G., Chylek, P., Lohmann, U.: A study of internal and external mixing scenarios and its effect on aerosol optical properties and direct radiative forcing, Journal of Geophysical Research, 107, 0148-0227, 2002.

Lin, C.Y., Zhao, C., Liu, X., Lin, N.H., Chen, W, N.: Modelling of long-range transport of Southeast Asia biomass-burning aerosol to Taiwan and their radiative forcings over East Asia, Tellus B, 66, 23733, 2014.

Liu, J., Wang, Y., Li, P.H., Shou, Y.P., Li T., Yang, M.M., Wang, L., et al.: Polycyclic Aromatic Hydrocarbons (PAHs) at High Mountain Site in North China: Concentration, Source and Health Risk Assessment. In press. DOI: 10.4209/aaqr.2017.01.0011, 2017.

Menut L., Bessagnet B., Khvorostyanov D., Beekmann M., Blond N., Colette A., Coll I., Curci G., Foret G., Hodzic A., Mailler S., Meleux F., Monge J.L., Pison I., Siour G., Turquety S., Valari M., Vautard R., Vivanco M.G.: CHIMERE 2013: a model for regional atmospheric composition modelling, Geosci. Model Dev., 6, 981-1028, 2013.

Michalakes, J., Dudhia, J., Gill, D., Henderson, T., Klemp, J., Skamarock, W., Wang, W.: The Weather Research and Forecast Model: Software Architecture and Performance. Proceedings of the 11th ECMWF Workshop on the Use of High Performance Computing in Meteorology, 25-29 October 2004, Reading, U.K. Ed. George Mozdzynski.

Middleton, P., Stockwell, W. R., Carter, W. P.: Agregation and analysis of volatile organic compound emissions for regional modelling, Atmospheric Environment, 24, 1107–1133, 1990.

Nair, V.S., Solmon, F., Giorgi, F., Mariotti, L., Babu, S., Moorthy, K.K.: .Journal of Geophysical Research, 117, D04209, 2012.

Ohara, T., Akimoto, H., Kurokawa, J., Horii, N., Yamaji, K., Yan, X., Hayasaka, T.: An Asian emission inventory of anthropogenic emission sources for the period 1980–2020, Atmospheric Chemistry and Physics, 7, 4419–4444, 2007.

Osuri, K,K., Mohanty, U.C., Routray, A., Kulkarni, M.A., Mohapatra, M.: Customization of WRF-ARW model with physical parameterization schemes for the simulation of tropical cyclones over North Indian Ocean, Nat Hazards, 63, 1337–1359, 2012.

Park, R.S., Song, C.H., Han, K.M., Park, M.E., Lee, S.S., Kim, S.B., Shimizu, A.: A study on the aerosol optical properties over East Asia using a combination of CMAQ-simulated aerosol optical properties and remote-sensing data via data assimilation technique, Atmospheric Chemistry and Physics, 11, 12275-12296, 2011.

Permadi, D.A. and Kim Oanh, N.T.: Assessment of biomass open burning emissions in Indonesia and potential climate forcing impact. Atmospheric Environment, 78, 250-258, 2013.

Permadi, D.A., Kim Oanh, N.T., Vautard, R.: Assessment of emission scenarios for 2030 and co-benefits of black carbon emission reduction measures on air quality and climate forcing in Southeast Asia, Submitted to Atmospheric Chemistry and Physics, 2017a.

Permadi, D.A., Sofyan, A., Kim Oanh, N.T.: Assessment of emissions of greenhouse gases and air pollutants in Indonesia and impacts of national policy for elimination of kerosene use in cooking. Atmospheric Environment, 154, 82-94, 2017b.

Pere, J.C., Mallet, M., Pont, V., Bessagnet, B.: Impact of aerosol direct radiative forcing on the radiative budget, surface heat fluxes, and atmospheric dynamics during the heat wave of summer 2003 over western Europe: A modeling study, J. Geophys. Res., 116, D23119, 2011.

Ramanathan, V., Crutzen, P.J., Lelieveld, J., Mitra, A.P., Althausen, D., Anderson, J., Andreae, M.O., et al.: Indian Ocean Experiment: An integrated analysis of the climate forcing and effects of the great Indo-Asian haze, Journal of Geophysical Research, 106, 28,371– 28,398, 2001.

Reddy, M.S., Boucher, O., Bellouin, N., Schulz, M., Balkanski, Y., Dufresne, J.L., Pham, M.: Estimates of global multicomponent aerosol optical depth and direct radiative perturbation in the Laboratoire de Meteorologie Dynamique general circulation model, Journal of Geophysical Research, 110, D10S16, 2005.

Schulz, M., Textor, C., Kinne, S., Balkanski, Y., Bauer, S., Berntsen, T., Berglen, T., Boucher, O., Dentener, F., et al.: Radiative forcing by aerosol as derived from the AeroCom present-day and pre-industrial simulations, Atmospheric Chemistry and Physics, 6, 5225–5246, 2006.

Seth, A. and Giorgi, F. The effects of domain choice on summer precipitation simulation and sensitivity in a regional climate model, Journal of Climate, 11, 2698 – 2712, 1998.

Shrestha, R. M., Kim Oanh, N.T., Shrestha, R., Rupakheti, M., Permadi, D.A., Kanabkaew, T., Salony, R.: Atmospheric Brown Cloud (ABC) Emission Inventory Manual (EIM), United Nation Environmental Programme (UNEP), Nairobi, Kenya, 2013.

Shindell, D., Kuylenstierna, J., Vignatti, E., Van Dingenen, R., Amann, M. et al.: Simultaneously mitigating near-term climate change and improving human health and food security, Science, 335, 183, 2012.

Shi, Y., Sasai, T., and Yamaguchi.: Spatio-temporal evaluation of carbon emissions from biomass burning in Southeast Asia during the period 2001–2010, Ecological Modelling, 272, 98-115.

Simpson, D., Winiwarter, W., Borjesson, G., Cinderby, S., Ferreiro, A., Guenther, A., et al.: Inventorying emissions from nature in Europe, Journal of Geophysical Research, 104, 8113–8152, 1999.

Song, Y., Chang, D., Liu, B., Miao, W., Zhu, L & Zhang, Y.: A new emission inventory for nonagricultural open fires in Asia from 2000 to 2009, Environ. Res. Lett., 5, 014014. Doi:10.1088/1748-9326/5/1/014014, 2009.

Streets, D.G., Bond, T.C., Carmichael, G.R., et al.: An inventory of gaseous and primary aerosol emissions in Asia in the year 2000, Journal of Geophysical Research, 108, NO. D21, 8809, 2003.

UNEP-C[4].: The Asian Brown Cloud: Climate and Other Environmental Impacts, UNEP, Nairobi, 2002.

UNEP-WMO.: Integrated assessment of black carbon and tropospheric ozone, United Nations Environment Programme and World Meteorological Organization, Nairobi, Kenya, 2011.

van der Werf, G.R., Randerson, J.T., Giglio, L., Collatz, G.J., et al.: Global fire emissions and the contribution of deforestation, savanna, forest, agricultural, and peat fires (1997–2009), Atmos. Chem. Phys., 10, 11707-11735, 2010.

WHO.: Health effects of black carbon. Copenhagen, World Health Organization Regional Office for Europe, 2012.

Zhang, Q., Streets, D.G., Carmichael, G.R., He, K.B., Huo,H., Kannari, A., Klimont, Z., Park, S., Reddy, S., Fu, JS., Chen, D., Duan, L., et al.: Asian emissions in 2006 for the NASA INTEX-B mission, Atmos. Chem. Phys., 9, 5131–5153, 2009.

Zhang, Y., Zhang, X., Wang, L., Zhang, Q., Duan, F., He, K.: Application of WRF/Chem over East Asia: Part I. Model evaluation and intercomparison with MM5/CMAQ, Atmospheric Environment, 124 285-300, 2016.

Table caption

5

Table 1. Summary of activity data level from different emission sources in 3 countries.

Sectors	Types of activity data	Activity data		
		Indonesia	Thailand	Cambodia
Power generation	Fuel consumption (Mt y^{-1}):			
	- Coal	23.4	20.5	-
	- Natural gas	3.2	29.8	-
	- Fuel oil	9.4	0.75	0.62
	- Biomass	6.3	-	-
Manufacturing industry	Fuel consumption (Mt y^{-1})			
	- Coal	5.4	12.3	-
	- Gasoline	0.34	0.013	-
	- Fuel oil	1.8	2.4	0.52
	- Biomass	-	20.7	-
On road transport	Number of registered vehicle (Million y^{-1})	48	26	1.9
Air traffic	LTO (*1000/year)	344	555	39
Residential & commercial	Fuel consumption (Mt y^{-1})			
	- Coal	0.028	-	-
	- Wood	100.5	7.6	0.4
	- Kerosene	7.3	0.13	0.003
	- LPG	1	1.15	0.005
	- Charcoal	20.4	3.9	0.042
	- Other biomass	-	0.14	-
Fugitive emission from fuel	- Coke production (Kt y^{-1})	182	-	-
	- Gas production (Tg y^{-1})	8,654	31.2	-
	- Oil production (Tg y^{-1})	29	6.2	-
	- Gasoline distributed (Mt y^{-1})	13.7	5.4	-
Agro residue open burning	Total dry crop residue openly burned (Mt y^{-1})	43.5	18.2	4.3
Forest fire	Total forest area burned, including peatland fire (ha y^{-1})	545,881	1,851,850	98,761
Solid waste open burning	Total dry solid waste burned (Mt y^{-1})	1.26	0.28	0.175
Agriculture related activities	- Total number of livestock population (head, *10^6)	1,359	328	22.3
	- Fertilizer consumption (Mt y^{-1})	6.8	3.6	-
Solvent and product use	- Paint (Kt y^{-1} of paint)	606	ne	ne
	- Degrease (t y^{-1} of solvent consumed)	103		
	- Chemicals (Kt y^{-1} of products)	1,269		
	- Other products use (i.e. ink, domestic solvent, glue and adhesives) (Kt of products)	161		

5

Table 2. EI results for base year in comparison with the existing regional EI datasets (Gg y^{-1}).

Species	Indonesia			Thailand			Cambodia			Other SEA countries[d]	Southern part of China	Total domain
	This study[a]	EDGAR[b]	CGRER[c]	This study[a]	EDGAR[b]	CGRER[c]	This study[a]	EDGAR[b]	CGRER[c]			
SO_2	997	2,433	1,499	827	721	961	41	42	34	2,695	6,204	10,781
NO_x	3,282	2,162	1,896	701	882	1,086	97	92	27	2,623	4,166	10,910
CO	24,169	32,246	26,703	9,095	12,553	10,815	2,877	2,453	570	19,054	33,377	89,252
NMVOC	3,840	4,528	8,225	1,120	525	3,052	331	18	113	5,644	4,441	15,613
NH_3	1,258	1,617	1,390	469	675	388	110	95	86	1,543	2,247	5,645
CH_4	3,950	10,300	6,443	1,053	4,541	3,567	713	1,969	708	13,833	14,640	34,218
PM_{10}	2,046	3,454	1,838	782	1,196	474.9	115	268	68	1,763	3,644	8,458
$PM_{2.5}$	1,644	2,023	1,609	607	781	388.5	65	201	61	1,466	2,653	6,519
BC	226	173	229	47	73	72	7	13	7	159	362	821
OC	674	711	1,246	240	234	364	40	73	32	604	643	2,245
CO_2	508,022	1,700,450 (514,882)	587,000	260,988	235,644 (229,500)	351,000	28,296	185,211 (174,300)	36,000	856,225	1,406,860	3,092,654
N_2O	180	329	ne	84	71	ne	60	73	ne	271	346	941

Note: ne – not estimated

[a] EI conducted for base year of 2007 using the ABC EIM framework (Shrestha et al., 2013). Detail methodology and results were presented in Permadi et al. (2017).
[b] EDGAR for base year of 2007 (Oliver et al., 2001). Retrieved from http://edgar.jrc.ec.europa.eu/datasets_list.php?v=42. The CO_2 emission, excluding forest fire post burn decay and decay of drained peatland, are given in brackets for comparison with our estimates
[c] CGRER for base year of 2006 (Zhang et al., 2009). Retrieved from http://www.cgrer.uiowa.edu/EMISSION_DATA_new/index_16.html. For NH_3, CH_4, and CO_2, emissions were taken from CGRER inventory in 2000 (Streets et al., 2003). Peatland fire for SEA for 2007 was taken from GFED v3. Retrieved from https://daac.ornl.gov/cgi-bin/dsviewer.pl?ds_id=1191.
[d] Other SEA countries include Brunei, Lao PDR, Malaysia, Myanmar, Philippines, Singapore and Vietnam.

Table 3. Statistical parameters for WRF Model Performance evaluation for two periods

Station	MBE			MAGE			RMSE			N
	RH (%)	T (ºC)	WS (m s⁻¹)	RH (%)	T (ºC)	WS (m s⁻¹)	RH (%)	T (ºC)	WS (m s⁻¹)	
January –March 2007										
Olongapo-Philippines	5.4	-0.9	1.3	15.8	3.1	0.18	19.3	3.9	2.7	1,861
Davao-Philippines	-4	-1.6	2.7	10.4	2.8	2.9	12.6	3.6	3.4	1,250
Don Muang-Thailand	-13	**-0.4**	1	16	2.5	1.6	18.5	3.8	**2**	2,148
Trat-Thailand	5.7	-1.9	1.6	12.7	3.1	3.1	15.2	3.7	3.6	996
Pnom Penh-Cambodia	**7.7**	0.7	**0.5**	15.5	3.6	2.1	18.1	4.5	2.6	1,513
Jakarta-Indonesia	**-7.1**	-0.8	0.7	14.5	3.2	2.1	19.7	4.2	2.6	2,036
Kuala Lumpur-Malaysia	**-2.5**	**-0.14**	**0.14**	6.8	**1.2**	1.1	10.3	2.2	**1.4**	2,143
Sarawak-Malaysia	**-1.8**	**-0.13**	**-0.3**	5.6	**1.2**	0.9	9.2	2.1	**1.2**	2,148
August – October 2007										
Olongapo-Philippines	**1.5**	2.3	**0.5**	8.3	2.5	1.3	17.6	3.5	3.1	1,958
Davao-Philippines	**-0.81**	**-0.13**	**0.2**	6.4	2.2	0.7	12.8	4.8	**1.3**	1,262
Don Muang-Thailand	**2.6**	**-0.4**	**-0.1**	11.1	2.1	1.4	15.4	3.5	2.2	2,139
Trat-Thailand	**0.86**	**-0.1**	2.1	5.4	**1.3**	2.7	11.6	3.8	3.3	1,017
Pnom Penh-Cambodia	**-6.7**	0.7	**0.1**	10.1	**1.7**	1.1	14.6	3.4	**1.7**	1,602
Jakarta-Indonesia	**0.8**	2.3	**0.47**	5.6	2.4	2.3	11.2	3.3	3.1	1,958
Kuala Lumpur-Malaysia	**-5**	**0.3**	**-0.1**	10.4	**1.5**	0.98	13.2	1.9	**1.23**	2,159
Sarawak-Malaysia	**-5.4**	**-0.1**	-0.6	8.9	3.5	1.1	11.56	4.2	**1.4**	2,159

Note: bolded values represent satisfactorily model output; Criteria for MBE: WS≤±0.5 m s⁻¹, T≤±0.5ºC, RH≤±10%; Criteria for MAGE: T≤ 2ºC, RH≤ 2 %; criteria for RMSE: WS≤ 2 m s⁻¹, N – number of data points. Description of the statistics measures are presented in Table S1, SI.

5

10

15

20

25

Table 4. Statistical parameters for CHIMERE model performance (PM$_{10}$ and BC) evaluation.

Parameters and station name	Statistical measures		
	MBE (μg m^{-3})	MFB (%)	MFE (%)
PM$_{10}$[a]			
1. BMR (average of 10T and 11T)[b]	-17.5	**-53.3**	**55.7**
2. SUF1 (Surabaya)[c]	-2.6	**-8.9**	**18**
3. Jerantut, Kuala Lumpur[d]	-13.6	**-56.3**	**66.5**
4. Petaling Jaya, Kuala Lumpur[e]	-10.3	**-41.1**	**56.1**
BC[f]			
AIT site	-0.12	**-3.3**	**20.8**

Note: Criteria from Boylan and Russel (2006). MFB: PM ≤ ±60%, and MFE: PM ≤+75%, bold showed the parameters that satisfy the criteria. No criteria is available for MBE.

[a] Period taken was from January – March and August – October 2007 for all stations (daily average concentrations)

[b] Urban mixed site

[c] Urban mixed site

[d] Background concentration

[e] Urban mixed site

[f] Period taken was from March to December 2007 (daily average concentrations).

Figure captions

Figure 1: Gridded (0.25º x 0.25º) annual emissions for the selected pollutants over the SEA domain
Figure 2: Comparison of modeled monthly precipitation and the TRMM-3B43 dataset in August and October 2017
Figure 3: Comparison of modeled and observed 24h PM_{10} in Kuala Lumpur, Malaysia (1 station), Surabaya, Indonesia (1 station) and Bangkok, Thailand (3 stations). Note that the stations included in the comparison are those located within the cell
Figure 4: Scatter plots of modeled vs. observed 24h $PM_{2.5}$ at four AIRPET sites, 2007
Figure 5: Time series comparison and scatter plot of modeled vs. observed 24h EC in AIT site, 2007
Figure 6: Comparison of 24h simulated and observed BC at four AIRPET sites in SEA domain, 2007
Figure 7: Spatial distribution of monthly average PM_{10}, $PM_{2.5}$ and BC in the selected months, 2007
Figure 8: Spatial distribution of monthly modeled AOD as compared to the MODIS Terra AOD for the selected months, 2007
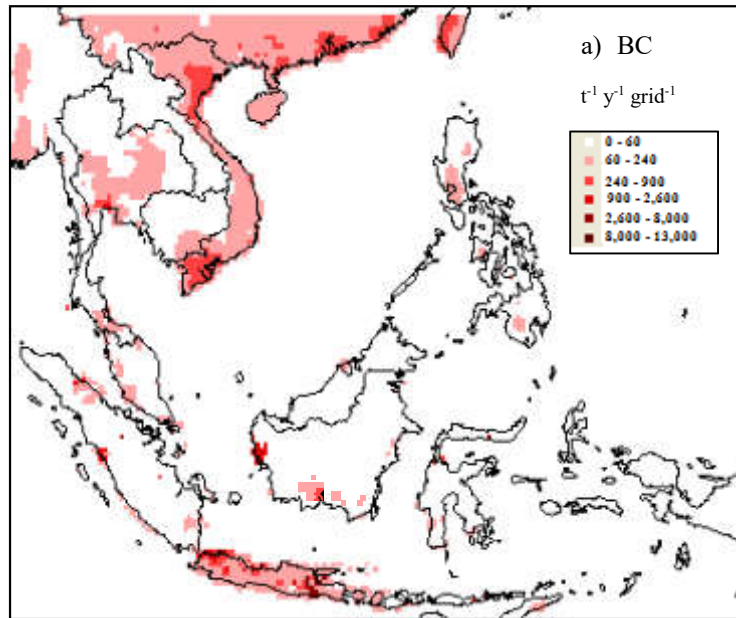**Figure 9: Monthly average of simulated vs. observed AOD at 10 AERONET stations, 2007**
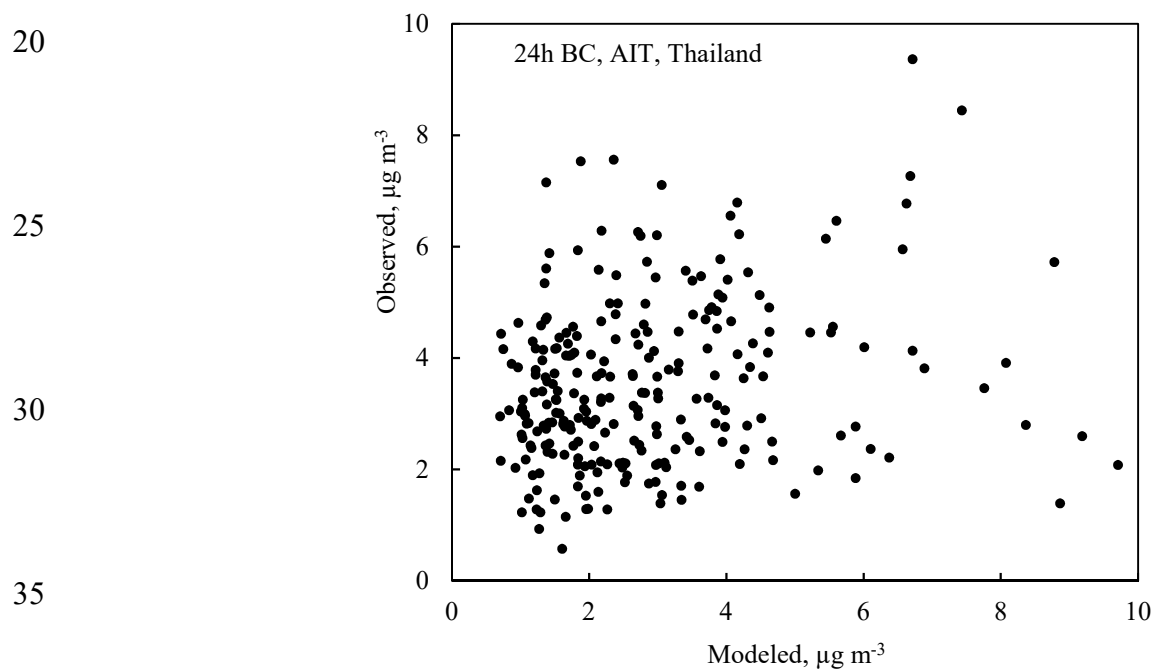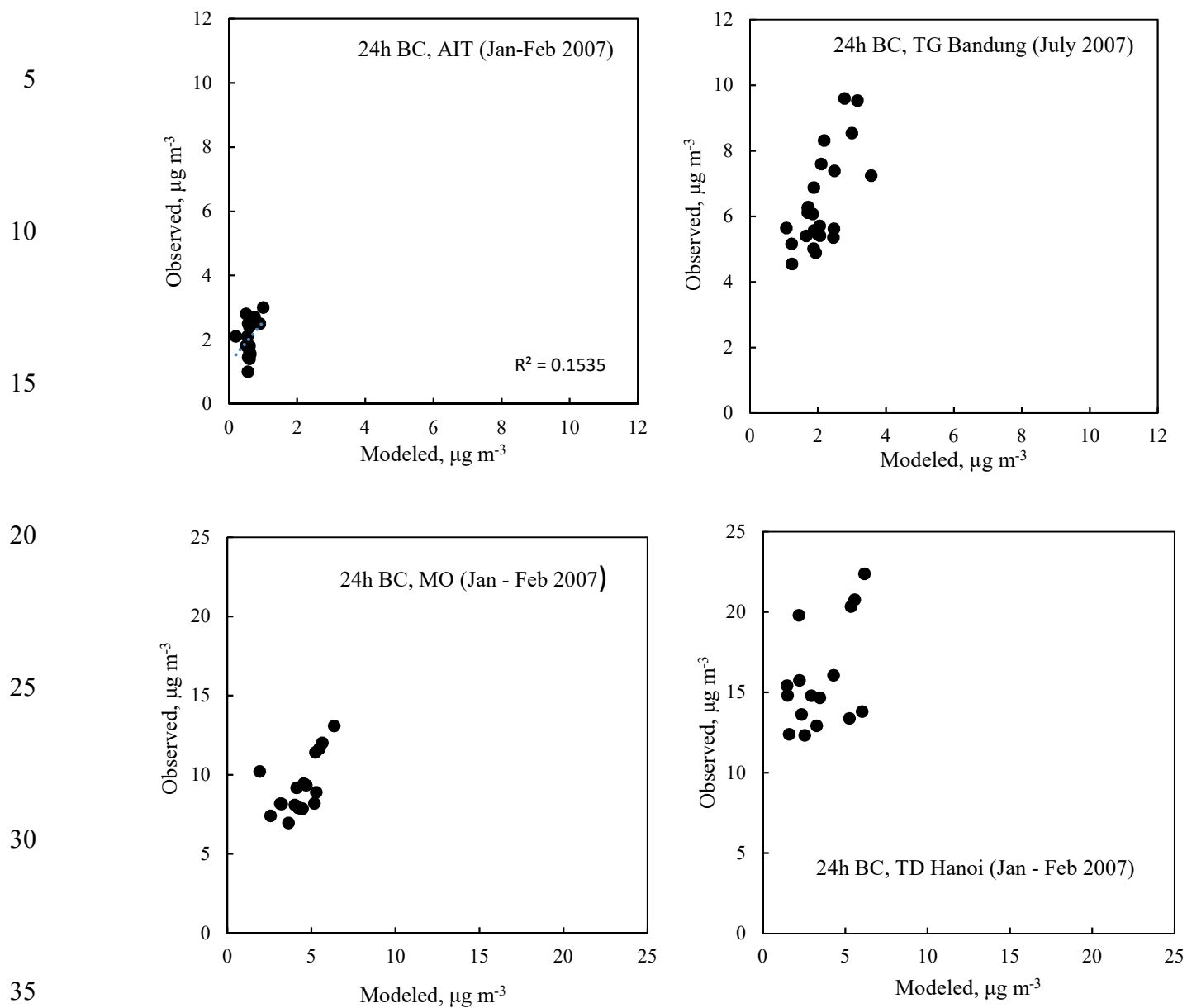
30

a) BC

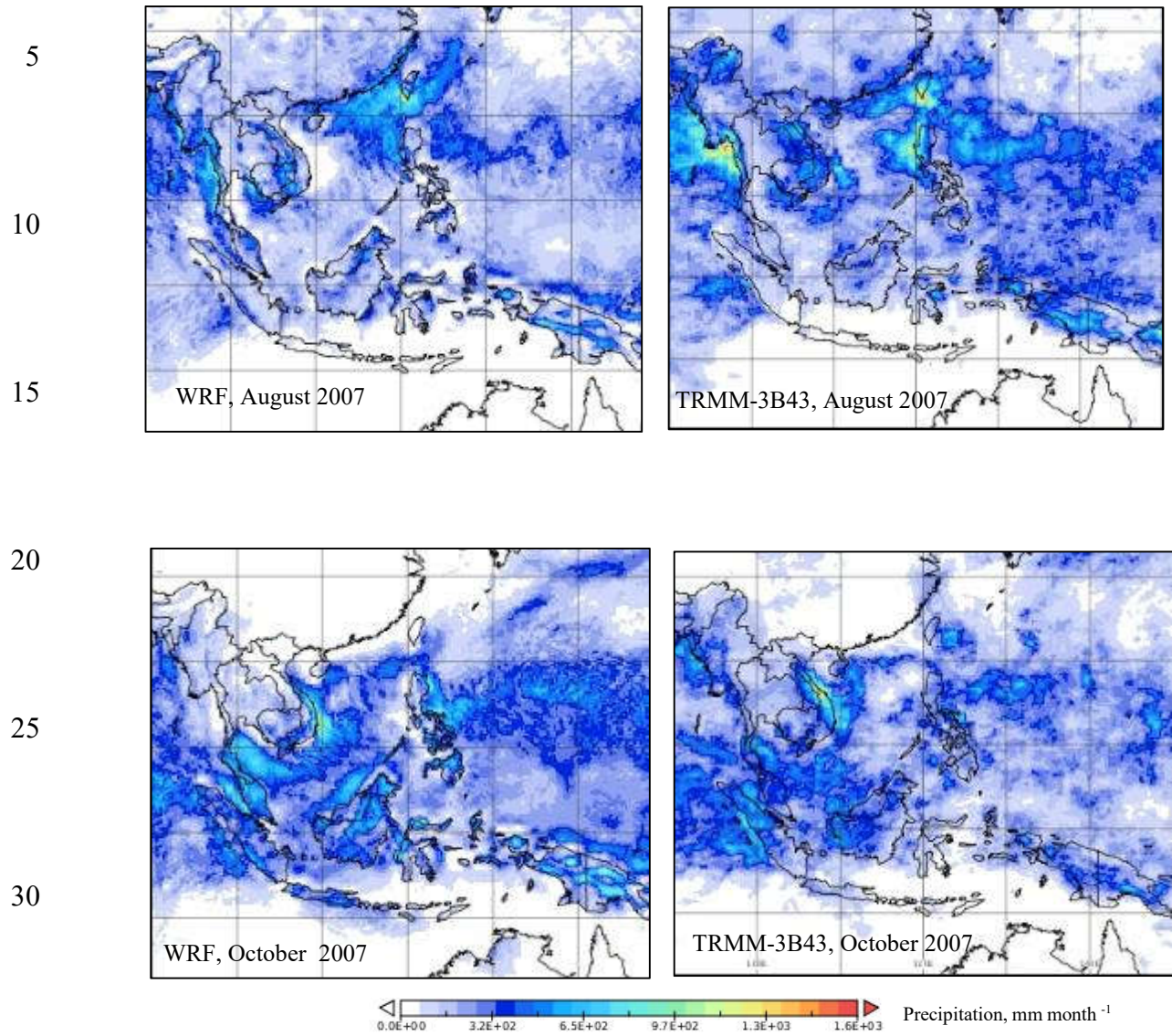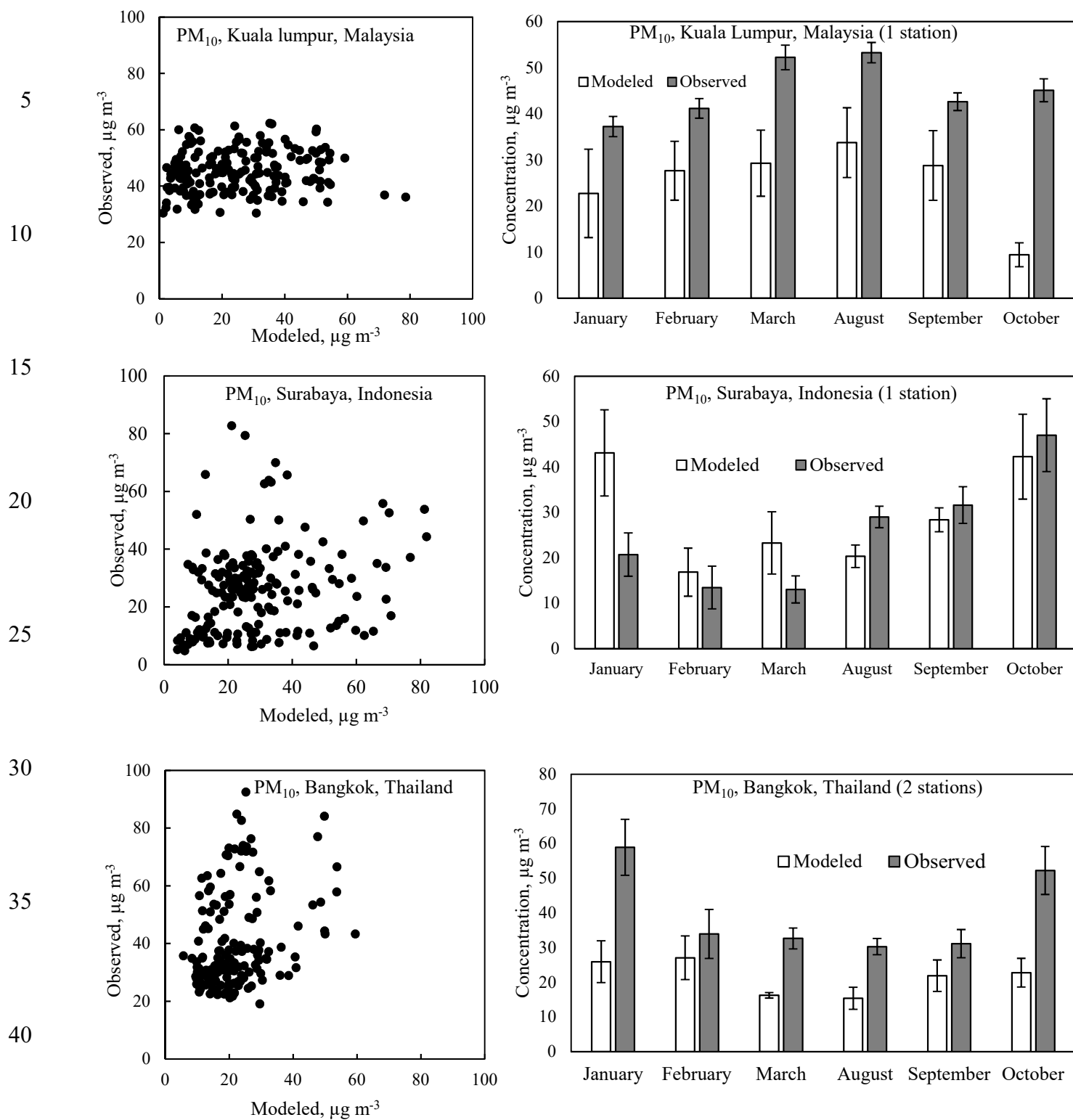$t^{-1} y^{-1} grid^{-1}$

0 - 60
60 - 240
240 - 900
900 - 2,600
2,600 - 8,000
8,000 - 13,000

b) CO

$t^{-1} y^{-1} grid^{-1}$

0 - 10,000
10,000 - 40,000
40,000 - 180,000
180,000 - 400,000
400,000 - 900,000

Figure 1. Gridded (0.25º x 0.25º) annual emissions for the selected pollutants over the SEA domain.

5

10

15

WRF, August 2007

TRMM-3B43, August 2007

20

25

30

WRF, October 2007

TRMM-3B43, October 2007

Precipitation, mm month $^{-1}$

0.0E+00   3.2E+02   6.5E+02   9.7E+02   1.3E+03   1.6E+03

35

Figure 2. Comparison of simulated monthly precipitation and the TRMM-3B43 dataset in August and October
40   **2017.**

45

Figure 3. Comparison of modeled and observed 24h PM₁₀ in Kuala Lumpur, Malaysia (1 station), Surabaya, Indonesia (1 station) and Bangkok, Thailand (2 stations).
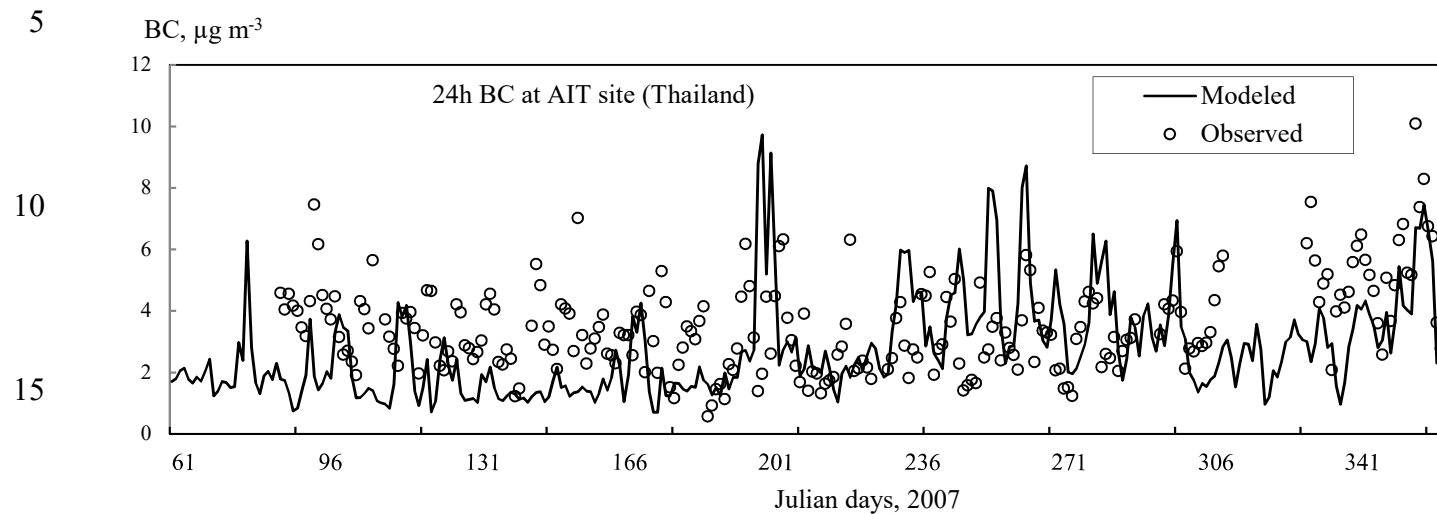**Note: the stations included in the comparison are those located within the cell.**

PM$_{2.5}$, AIT Thailand (Jan-Feb 2007)

Observed, μg m^{-3}

Modeled, μg m^{-3}

PM$_{2.5}$ Tegal Lega (TG), Bandung
(July 2007)

Observed, μg m^{-3}

Modeled, μg m^{-3}

PM$_{2.5}$, Manila Observatory (MO)
(Jan-Feb 2007)

Observed, μg m^{-3}

Modeled, μg m^{-3}

PM$_{2.5}$ Thuong Dinh (TD), Hanoi
(Jan-Feb 2007)

Observed, μg m^{-3}

Modeled, μg m^{-3}

Figure 4. Scatter plots of modeled vs. observed 24h PM$_{2.5}$ at four AIRPET sites, 2007.

34

Figure 5. Time series comparison and scatter plot of modeled vs. observed 24h EC in AIT site, 2007.

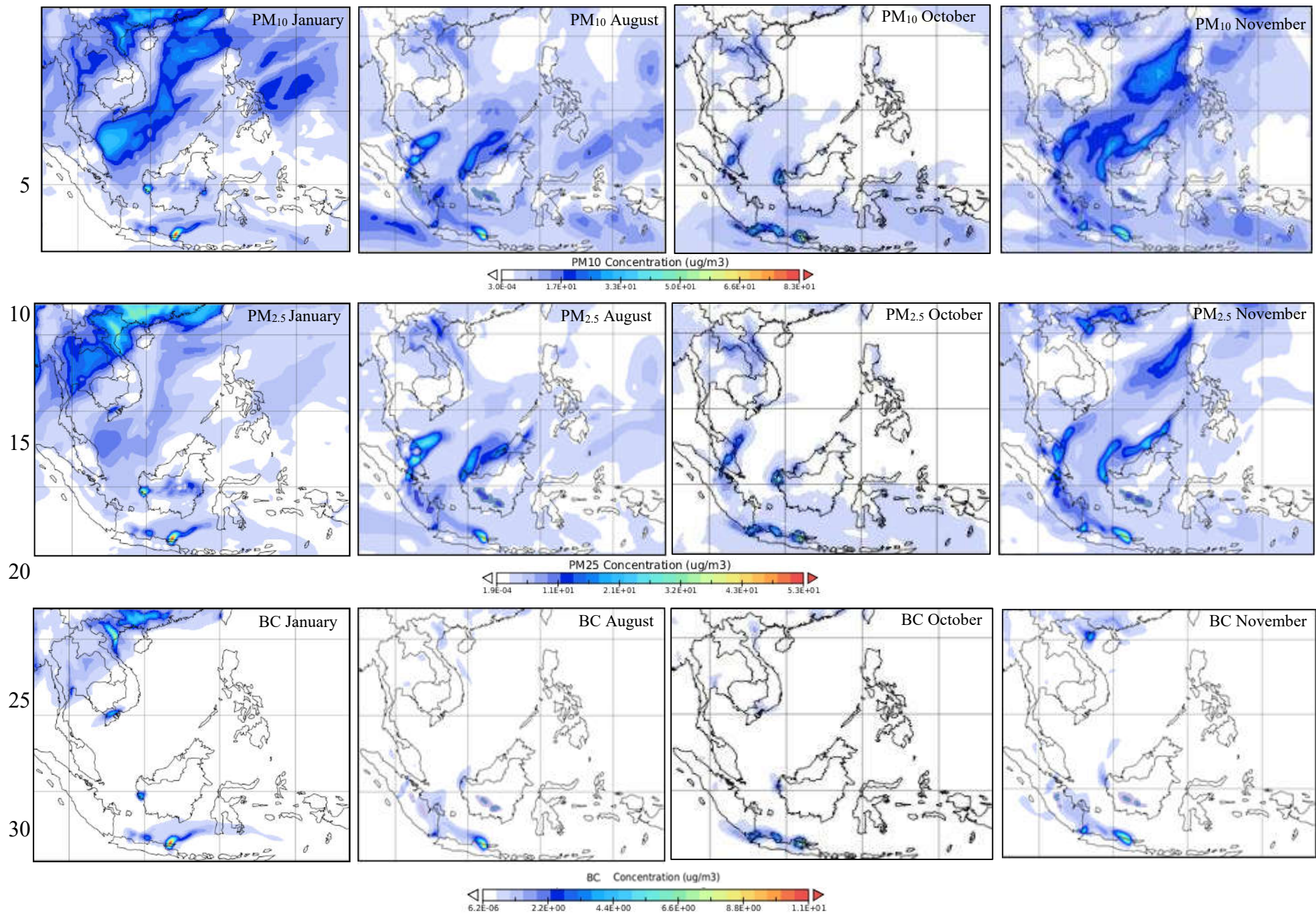**Figure 6. Comparison of 24h simulated and observed BC at four AIRPET sites in SEA domain, 2007.**

Figure 7. Spatial distribution of monthly average PM₁₀, PM₂.₅ and BC in the selected months, 2007.

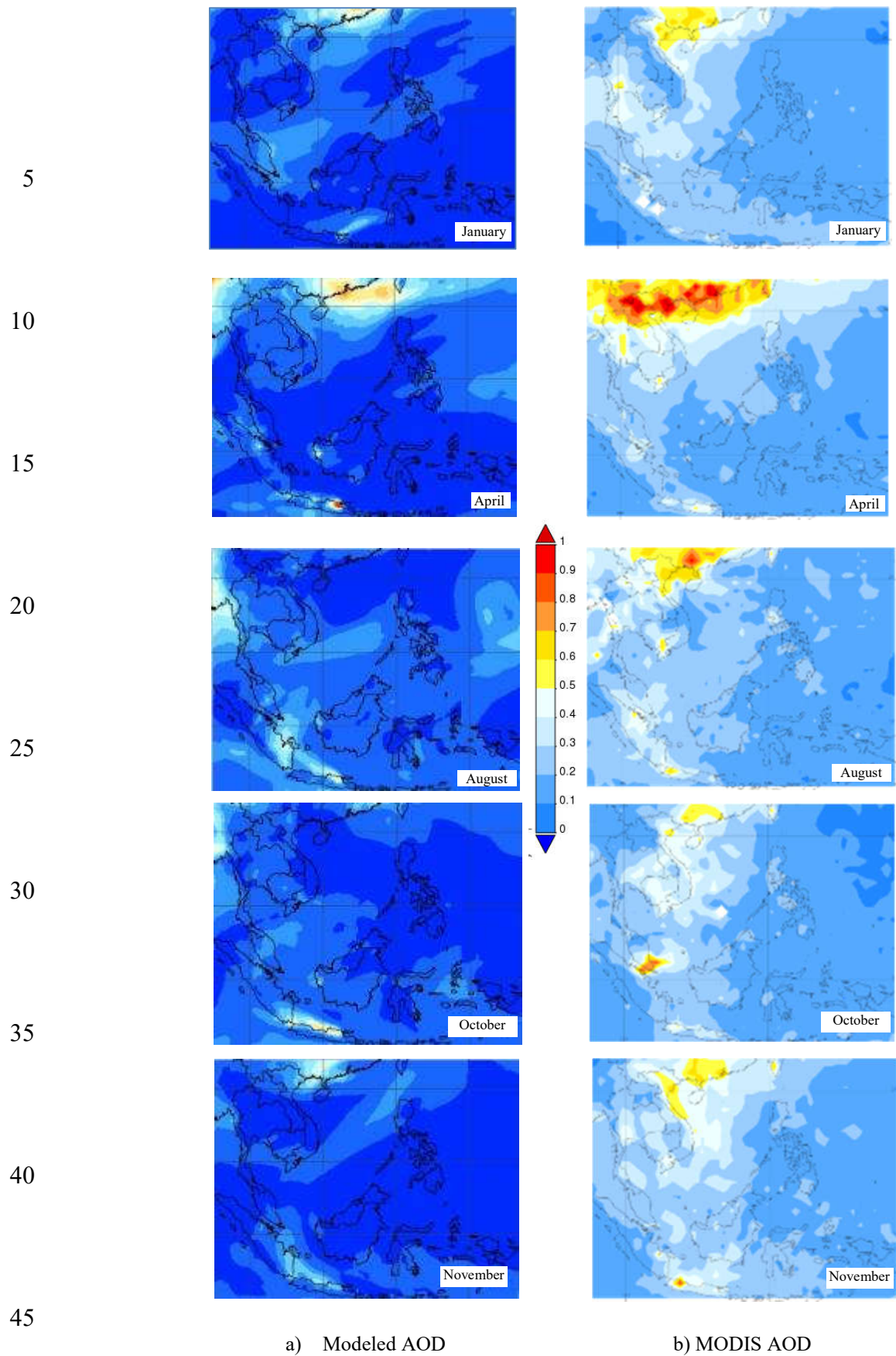a) Modeled AOD   b) MODIS AOD

Figure 8. Spatial distribution of monthly modeled AOD as compared to the MODIS Terra AOD for the selected months, 2007.
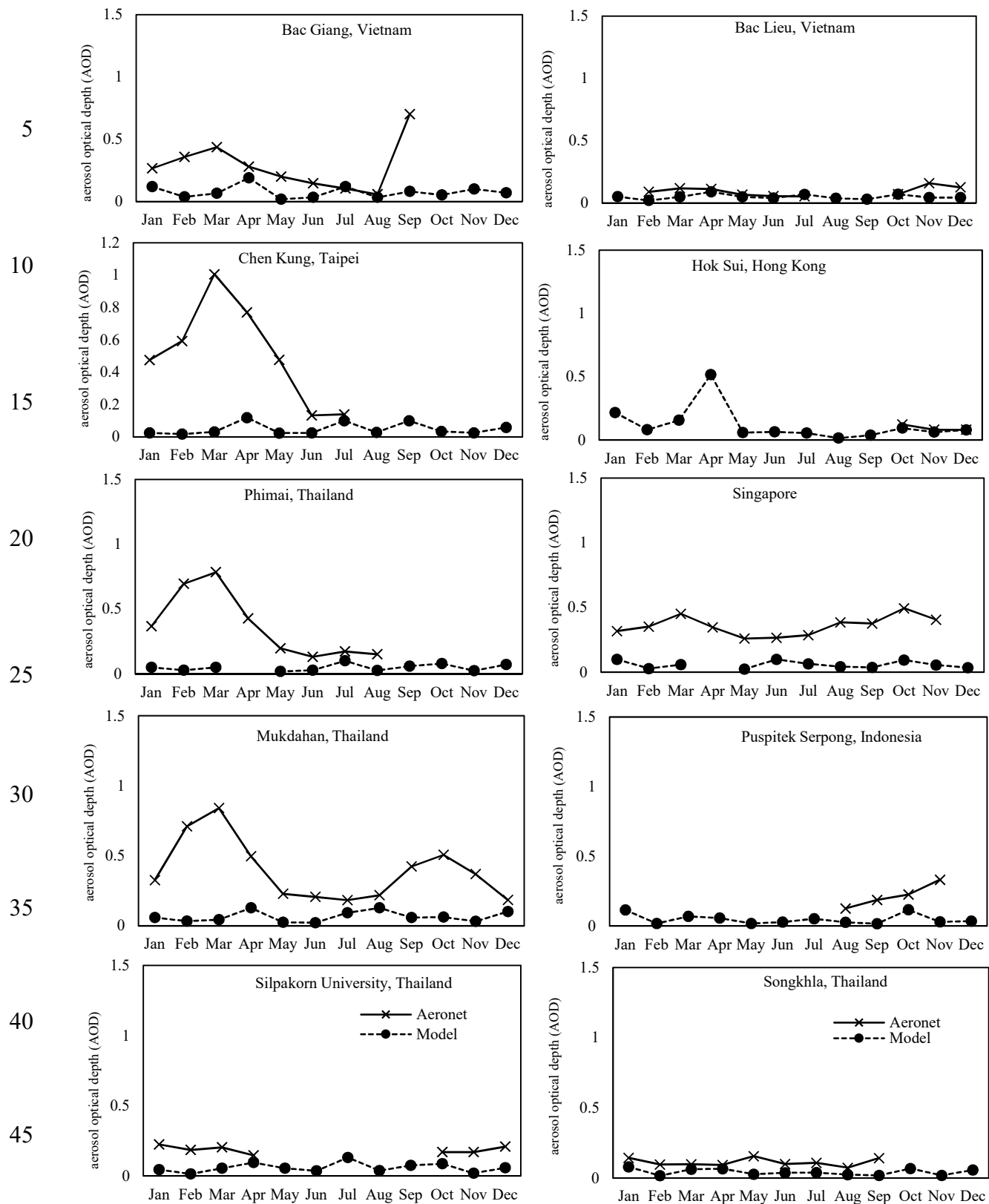
Figure 9. Monthly average of simulated vs. observed AOD at 10 AERONET stations, 2007.