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A modeling study of the influence of sea salt on inorganic aerosol concentration, size distribution, and deposition in the western Pacific Ocean

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14	Abstract. A regional air quality model system (RAQMS) was developed by coupling the
15	treatment of heterogeneous reactions between sea salt aerosols (SSAs) and trace gases and
16	applied to the investigation of aerosol properties and evolutionary features in the western
17	Pacific Ocean in the spring of 2014. Model results for meteorological variables, PM
18	concentrations, and size-resolved water soluble inorganic aerosol (WSIA) concentrations were

compared and analyzed with a variety of observations from in situ measurements and the 19 research cruise Dongfanghong II. Model validation demonstrated that the model can simulate 20 the spatial-temporal distribution and size distribution of aerosol inorganic components in the 21 marine atmosphere over East Asia, and the inclusion of heterogeneous reactions on SSAs 22 apparently improved the model simulation for WSIA concentration, especially for aerosol size 23 distribution. In the western Pacific Ocean, the non-sea salt SO_4^{2-} and NO_3^{-} formed on SSAs 24 accounted for up to 30% and 90% of surface SO_4^{2-} and NO_3^{-} concentrations on average, 25 respectively. The atmospheric depositions of total inorganic sulfur and nitrogen were estimated 26 to be 13184×10^3 kgS/d and 10728×10^3 kgN/d, respectively. Wet deposition was the dominant 27 removal pathway, which accounted for 75% and 68% of sulfur and nitrogen depositions, 28 respectively. The deposition of fine-mode SO_4^{2-} exceeded that of coarse-mode SO_4^{2-} , whereas 29 the deposition of coarse-mode NO_3^- was comparable to that of fine-mode NO_3^- . The non-sea 30 salt SO_4^{2-} and NO_3^{-} formed on SSAs contributed 16% and 9% of total sulfur and nitrogen 31

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