Evaluate the spatial and temporal variation of  $N_2O$  and associated flux into the air

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In order to evaluate the spatial and temporal variation of N<sub>2</sub>O and associated flux into the air in a granite unconfined aquifer of Ikuchi Island, water samples were collected from 9 observation wells with different depths and 6 observation wells in the groundwater discharge area from 2013 to 2015 and analyzed for N<sub>2</sub>O, NO<sub>3</sub><sup>-</sup>-N and Cl<sup>-</sup>. The results showed that the concentrations of dissolved N  $_2$ O changed with water depth, which can be attributed to the C/N ratio. When the C/N ratio  $\leq$ 5, high concentrations of dissolved N<sub>2</sub>O occur. In addition, the N<sub>2</sub>O concentrations increased with the redox condition of water changes from oxidation to reductive. However, when water was in strong reductive conditions such as ORP  $\leq$ -200mV, the relative low concentrations of N<sub>2</sub>O took place, since N<sub>2</sub>O can change into N<sub>2</sub> due to the complete denitrification. Dissolved N<sub>2</sub>O concentrations also increased in dry seasons, when most of observation wells being in a reductive state. In the groundwater discharge area, dissolved N<sub>2</sub>O and NO<sub>3</sub><sup>-</sup>-N concentrations decreased along the groundwater flow pathway, which results from the dilution of seawater and denitrification. The flux of N<sub>2</sub>O into air was estimated to be 49gha<sup>-1</sup>year<sup>-1</sup>, the same level as seawater emission rate. Whereas, in a high precipitation event(precipitation  $\leq$ 30mm/day), 5gha<sup>-1</sup>day<sup>-1</sup> (about 10 percent of annual N<sub>2</sub>O emission) would emission into air.