Isotope constraints on dynamic nitrogen transformations in a nitrogen-replete coastal bay in southern China

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Nitrogen is essential for organisms and regulates marine productivity. Growing input of anthropogenic nitrogen from land significantly altered the original N cycle in coastal seas and consequently, the community structure and ecosystem function. However, it is a difficult task to resolve all nitrogen transformation processes at the same time due to the complexity and dynamic feature of nitrogen reaction web (five major N components, particulate/dissolved organic nitrogen, ammonium, nitrite and nitrate, interact rapidly). By adding one single ¹⁵N-labelled tracer (e.g., NH₄⁺) and measuring the and isotopic composition concentration changes simultaneously for PON, NH4+, NO2 and NO3, we created matrix equation to derive rates of well-known transformation processes in the reaction web under the assumption of mass conservation. A 24-hr incubation experiment was conducted in January 2014 in Wuyuanwan, a nutrient-replete bay in southern China. Two water depths with distinctive light intensity (2% and 80% PAR) were chosen. The solutions of matrix equation provided us rates for NH4+, NO2- and NO3uptake, ammonia oxidation, nitrite oxidation, nitrite excretion. Results showed that phytoplankton uptake dominates ammonium consumption, with rates of (691-1155) and (465-931) nmol/L/h under 80% and 2% PAR, respectively, which are orders of magnitude higher than ammonium oxidation rates at corresponding depth. Allochthonous NH4+ input passing through phytoplankton assimilation instead of nitrification means an enhancement of oxygen producing rather than consuming. The high ammonium (~25 uM) likely inhibit NO3⁻ uptake (175 and 96 nmol/L/h) though nitrate concentration (~30 uM) is also high. In addition, NO2⁻ uptake is undetectable while NO2⁻ excretion from phytoplankton is light sensitive. This is the first attempt to untangle the specific rate of individual N processes at once in costal water where characterized by serious anthropogenic influence.