

^{17}O -rich nitrate as a tracer for constraining nitrogen transformations in coastal sediments

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Coastal sediments are often replete in organic matter and exhibit sharp gradient in redox conditions. In addition, the oxic-anoxic interfaces are often around a few centimeters if not less below the sediment surface. This make this environment poised to harbor both oxidative and reductive nitrogen transformations. As such, the use of a single or a couple of isotope-labeling tracers to determine the rates of these many processes might suffer from the systems being underconstrained. In this work, we demonstrate how naturally-occurring, ^{17}O -rich nitrate can aid in the studying these complex systems through the ability to follow the transformations of added nitrate (NO_3^-) into other different pools of N-species and closely investigate the triple isotopic compositions ($\delta^{15}\text{N}$, $\delta^{18}\text{O}$, and $\Delta^{17}\text{O}$). While we followed five different pools of N-species namely NO_3^- , NO_2^- , N_2O , NH_4^+ , and total reduced N (NH_4^+ plus DON), we chose to focus this work on the intermediate nitrite, NO_2^- . For all intact flow-through core incubations done on the sediments collected from Sylt Island, Germany, sediments acted as sources for NO_2^- in all experimental manipulations including sediment type, dissolved oxygen level, and NO_3^- loading. Unlike in the environments that are solely driven by reductive processes where the changes in $\delta^{15}\text{N}$ and $\delta^{18}\text{O}$ are coupled, the co-occurrence of both oxic and anoxic in the sediments such as ones from this study cause the $\delta^{15}\text{N}$ and $\delta^{18}\text{O}$ to decouple primarily because O is often subject to more processes than N. By using three isotope systems along with the change in concentrations, we demonstrate how we can use a natural abundance approach and rely on a series of mathematical equations to solve for different N transformation rates.

Keywords: nitrogen, isotope, ^{17}O , sediments

