

海水溶解磷酸盐氧同位素组成的测定

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中文摘要:生物磷酸盐和水分子间的氧同位素分馏主要受温度和生物活动控制,因此磷酸盐氧同位素组成既可以测量古温度又可以示踪磷循环。近年来磷酸盐氧同位素研究受到较多除了传统的生物体磷灰石古温度测量外,这些研究大多是关于磷循环的。磷酸盐的氧同位素组成可以示踪海洋中磷的源区和生物对磷的利用效率。由于海水的组成十分复杂,测量前对样品进行富集、分离和纯化处理。目前,加州大学(Santa Cruz)Paytan教授和耶鲁大学Blake教授的实验室已建立了海水溶解磷酸盐氧同位素的测量方法,二者各有优缺点。我们结合这两种方法的优点,并对一些步骤进行了改进,建立了海水溶解磷酸盐氧同位素组成的测量方法。通过向海水样品中加入NaOH,形成Mg(OH)₂来富集海水中的PO₄³⁻,也可同时除去杂质离子和溶解有机质;通过将PO₄³⁻转化为CePO₄沉淀来进一步除去杂质离子,然后用阳离子交换树脂去除Ce³⁺,再通过阴离子交换树脂柱来除溶解有机质。最后将磷酸盐转换为Ag₃PO₄沉淀,在1350℃裂解Ag₃PO₄,产生的O₂和石墨反应形成CO用IRMS测定。结果显示富集、分离和纯化过程可以获得纯的Ag₃PO₄颗粒,不会产生PO₄³⁻的氧同位素分馏。测量Ag₃PO₄用量仅为0.3 mg,标准偏差在±0.2%~±0.3%之间。

中文关键词:海水 溶解磷酸盐 氧同位素 方法

The Measurement of Oxygen Isotope Composition of Dissolved Inorganic Phosphate in Seawater

Abstract:The oxygen isotope fractionation between biogenic phosphate and water is primarily controlled by temperature and biological activity, so the oxygen isotope composition of phosphate ($\delta^{18}\text{O}_\text{P}$) has been used to measure the paleotemperature and trace phosphorus biogeochemical cycles. Recently the study of $\delta^{18}\text{O}_\text{P}$ has attract much attention. In addition to traditional biogenic apatite paleotemperature measurements, the studies are mostly concentrated on the phosphorus cycle. $\delta^{18}\text{O}_\text{P}$ of dissolve inorganic phosphate (DIP) is an effective proxy for tracing sources and biogeochemical cycle of phosphorus. Seawater composition is very complex, so the seawater sample must be separated and purified before the $\delta^{18}\text{O}_\text{P}$ measurement. Professor Paytan of University of California (Santa Cruz), and Professor Blake of Yale University have established their respective $\delta^{18}\text{O}_\text{P}$ measurement methods of seawater DIP, but the two method have their respective advantages and disadvantages. The authors combine the advantages of the two methods and modified some procedures to establish an improved $\delta^{18}\text{O}_\text{P}$ measurement method of seawater DIP. In this method, DIP in water samples is concentrated through Mg(OH)₂-PO₄ co-precipitation, which can exclude a portion of the dissolved organic material (DOM) and interfering ions; then the resulting solution is converted to CePO₄ precipitate to further separate Pi from dissolved salts, especially Cl⁻, and subsequently the solution is purified through a batch mode cation r to remove Ce³⁺ from the solution and through anion resin column treatment to remove the dissolved organic material. Phosphate is converted to silver phosphate ultimately which is in turn pyrolytically decomposed to CO at 1350 °C and $\delta^{18}\text{O}$ is analyzed with continuous-flow isotope ration mass spectrometry (IRMS). The results show that this