Potentiality for Over-Estimation of Total Alkalinity observed in Arctic Ocean by Spectrophotometric Method

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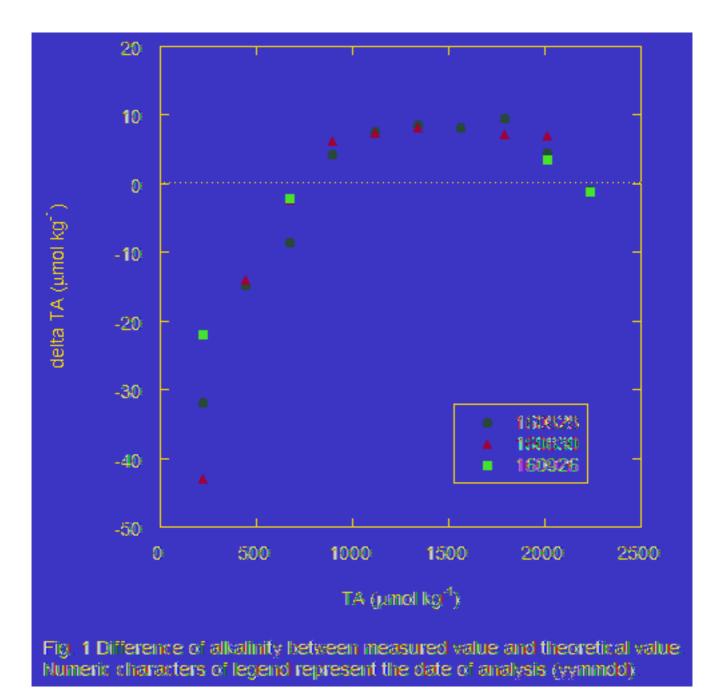
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Total alkalinity is an important component of oceanic carbonate system, which deeply affects and is infected to the oceanic chemistry, and also oceanic biology. Measurement of total alkalinity is conducted by potentiometric or spectrophotometric method. Potentionetric method is a traditional method that has been employed for a long time, while spectrophotometric method, that avoids problems attributed to glass-made pH electrodes and has higher precision than potentiometric method, was developed in 1990's and goes into use in this century. A spectrophotometric method was introduced to the laboratory of Research and Development Center for Global Change, Japan Agency for Marine-Earth Science and Technology (JAMSTEC) and the onboard measurement has been conducted for around a decade. Observations were also done in Arctic Ocean, where low alkalinity with its value under 2000 μ mol kg⁻¹ was often observed especially in sea surface. However, spectrophotometric method is suitable for the measurement of oceanic seawater with its alkalinity values ranges from about 2000 μ mol kg⁻¹ to about 2500 μ mol kg⁻¹, and its application for lower alkalinity has not been examined. In this study, we prepared some seawater samples with their alkalinity under 2000 μ mol kg⁻¹, and measured their value by spectrophotometric method.

Reliability of low alkalinity value measured by the spectrophotometric alkalinity system is evaluated by measurements of CRM and its dilutions with ultra pure water under assumption that alkalinity of ultra pure water is 0 μ mol kg⁻¹. Ten types of dilutions with its dilution ratio from 0 % (stock solution) to 90 % are prepared for measurements. Titration should be terminated at pH ~ 3.8 –4.2 in the case of using BCG as pH indicator. To examine the pH termination ranges are appropriate for analysis of seawater with low alkalinity, the CRM dilutions were analyzed at their pH termination ~ 3.1 - 4.7.

Measured alkalinity is not necessarily equal for the designated terminated pH ~ 3.8 - 4.2. The difference of measured alkalinity is in 2 - 3 μ mol kg⁻¹ for the dilution ratio under 40 % (over alkalinity of 1341.9 μ mol kg⁻¹), however, the difference of alkalinity in this pH range increases as the dilution ratio increase (lower alkalinity). The difference becomes about 10 μ mol kg⁻¹ with its dilution ratio of 50 % (1118.3 μ mol kg⁻¹), and over 20 μ mol kg⁻¹ with its ratio of 80 % (447.3 μ mol kg⁻¹). Trueness of measured alkalinity is then discussed. Figure 2 shows the difference between the measured alkalinity and theoretical alkalinity. The difference is related to their alkalinity and shows a convex distribution. Around alkalinity of 2250 μ mol kg⁻¹, the averaged alkalinity and theoretical value has good agreement. Below 2250 μ mol kg⁻¹. Lower than the alkalinity. Its difference is about 10 μ mol kg⁻¹ around the alkalinity of 1500 μ mol kg⁻¹. Lower than the alkalinity of 1500 μ mol kg⁻¹, the difference decreases as the alkalinity decrease, and the measured value and the theoretical value is nearly equal again at around alkalinity of 750 μ mol kg⁻¹. Lower than the alkalinity of 750 μ mol kg⁻¹, the measured alkalinity turns to be lower than the theoretical alkalinity and its absolute difference increases with alkalinity decrease.

In Arctic Ocean, where sea-ice melting and increase of inflow of river water has a lot of attention in these decades, alkalinity less than 2000 μ mol kg⁻¹ was often observed. On edge of sea-ice melting area, alkalinity less than 1700 μ mol kg⁻¹ is occasionally observed. Our analysis shows that measured alkalinity with its value of around 1300 - 1800 μ mol kg⁻¹ is considered to be overestimated with its difference of around 10 μ mol kg⁻¹. The degree of overestimation is less at external side of this range.



Keywords: Alkalinity, Spectrophotometric Method, Arctic Ocean

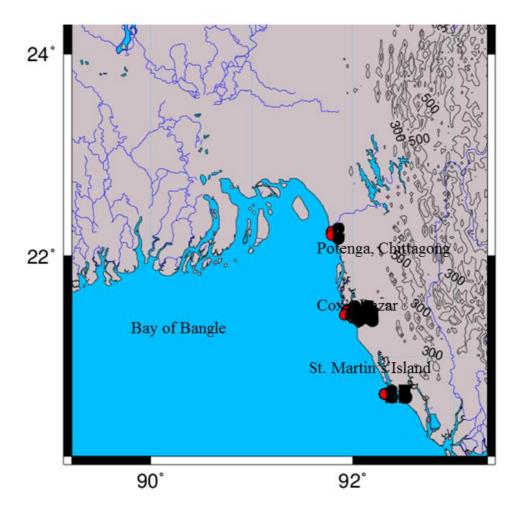
Distribution of Heavy Metals in Beach Water and Sediment of the Bay of Bengal Coast, Bangladesh

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The objectives of this research are to characterize the distribution of arsenic (As) including other heavy metals in water and sediment, types of coastal sediment in 3 coastal sites, i) located at Cox's Bazar, is the longest unbroken sea beach in the world, running 120 kilometres, ii) Patenga, a sea beach located 14 kilometres south of the port city of Chittagong, iii) St. Martin's Island, a small island (area only 8 km²) from watershed of the Bay of Bangle was investigated. In order to evaluate the occurrence of trace metals, sixty (60) sediment samples, thirty (30) seawater samples, were collected from the three coastal sites. The studied sediments show lower values (2.0-18.7 μ g/L) indicating that the sediments are unpolluted. The As concentration in water samples (average = 8.57-34.7 μ g/L) considered high. This research first investigated the water qualities and distribution pattern of rare-earth concentration in coastal sediment and water, providing a baseline in the Bay of Bengal, Bangladesh. We have, therefore, drawn new color maps for As, heavy metals and rare-earth using computerized software techniques as *Generic Mapping* Tools (GMT), Ocean Data View (*ODV*) and find the current geochemical pattern. This type of map may be used to establish general baselines against which more specific natural geochemical variations and human-induced perturbations can be appraised.

Keywords: Bay of Bangle, Arsenic (As), Rare-Earth, Sediment, Coastal Water



Stable nitrogen isotopic composition (δ^{15} N) of dissolved organic nitrogen (DON) in the South China Sea

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Dissolved organic nitrogen (DON) represents an essential N pool, and its stable isotopic composition (δ^{15} N) may provide unique insights into marine N cycle. However, few results of DON δ^{15} N have been reported to date in the global ocean. In this study, we will report DON δ^{15} N values in the upper layer waters of the South China Sea (SCS), which is the largest marginal sea west of the Pacific Ocean.

Keywords: South China Sea, δ 15N, DON