

# Global relationships of total alkalinity with salinity and temperature in surface waters of the world's oceans

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Received 12 June 2006; revised 29 August 2006; accepted 5 September 2006; published 5 October 2006.

[1] A simple function of sea surface salinity (SSS) and temperature (SST) in the form  $A_T = a + b(SSS - 35) + c(SSS - 35)^2 + d(SST - 20) + e(SST - 20)^2$  fits surface total alkalinity ( $A_T$ ) data for each of five oceanographic regimes within an area-weighted uncertainty of  $\pm 8.1 \mu\text{mol kg}^{-1}$  ( $1\sigma$ ). Globally coherent surface  $A_T$  data ( $n = 5,692$ ) used to derive regional correlations of  $A_T$  with SSS and SST were collected during the global carbon survey in the 1990s. Such region-specific  $A_T$  algorithms presented herein enable the estimation of the global distribution of surface  $A_T$  when observations of SSS and SST are available. **Citation:** Lee, K., L. T. Tong, F. J. Millero, C. L. Sabine, A. G. Dickson, C. Goyet, G.-H. Park, R. Wanninkhof, R. A. Feely, and R. M. Key (2006), Global relationships of total alkalinity with salinity and temperature in surface waters of the world's oceans, *Geophys. Res. Lett.*, 33, L19605, doi:10.1029/2006GL027207.

## 1. Introduction

[2] Total alkalinity ( $A_T$ ) variability in the surface ocean is controlled mainly by freshwater addition (precipitation and sea-ice melting) or removal (evaporation and sea-ice formation) which also acts to change salinity [Brewer *et al.*, 1986; Millero *et al.*, 1998a]. In the (Sub)tropical oceans (i.e.,  $30^\circ\text{N}$ – $30^\circ\text{S}$ ), surface  $A_T$  variations associated with water balance-induced changes in salinity account for more than  $\sim 80\%$  of total variability in  $A_T$  [Millero *et al.*, 1998a]. At higher latitudes (i.e., north of  $\sim 30^\circ\text{N}$  or south of  $\sim 30^\circ\text{S}$ ), a progressive increase in the convective mixing of deep waters rich in  $A_T$  during seasonal cooling is an important additional factor that acts to increase surface  $A_T$  concentrations. The greater the intensity of seasonal convective mixing; the higher the  $A_T$  associated with the dissolution of  $\text{CaCO}_3$  and the lower the water temperature. In this case,  $A_T$  is negatively correlated with sea surface temperature. Coccolithophorid

blooms during seasonal warming are key biological factors that decrease surface  $A_T$  in a measurable way [Balch *et al.*, 2005]. The formation and destruction of organic matter also impact surface  $A_T$ , but only in a minor way, except in biologically productive waters [Kim *et al.*, 2006]. Such decrease in  $A_T$  due to marine biota generally occurs in tandem with sea surface temperature increase. Overall, variations in both sea surface salinity (SSS) and temperature (SST) can be good proxies for surface  $A_T$  variations.

[3] Identifying the controls on surface  $A_T$  is becoming increasingly important for understanding the effects of ocean acidification resulting from the addition of anthropogenic  $\text{CO}_2$  into surface waters [Feely *et al.*, 2004]. However, determining the global distribution of surface  $A_T$  is problematic because in many parts of the ocean  $A_T$  data are severely limited compared to the SSS and SST data sets, which are several orders of magnitude larger. In this case, empirical algorithms that relate surface  $A_T$  to SSS and SST are particularly useful in constructing the global distribution of  $A_T$  when combined with the global fields of SSS and SST. Such empirical  $A_T$  algorithms can aid in predicting the  $\text{CO}_2$  flux across the air-sea interface in a numerical ocean model because surface  $\text{pCO}_2$  could be calculated from the values of  $A_T$  and total inorganic carbon ( $C_T$ ) for the model's surface box. In such a model,  $A_T$  and  $C_T$  are transported between the ocean boxes. Alternatively, surface  $\text{pCO}_2$  and  $A_T$  data are used to calculate the surface  $C_T$ . The surface  $A_T$  algorithms can also be used to study other aspects of the marine carbonate system. For example, the annual cycle of  $A_T$  in the global ocean derived from the surface  $A_T$  algorithms along with hydrographic parameters can be extended to estimate the annual export production of biogenic  $\text{CaCO}_3$  by integrating the seasonal decrease in the mixed-layer inventory of  $A_T$ , once it is corrected for salinity effects [Lee, 2001].

[4] The first set of global relationships of salinity-normalized  $A_T$  with SST [Millero *et al.*, 1998a] was derived using subsets ( $n = 1,740$ ) of historical  $A_T$  data as well as those collected during the global carbon survey of the 1990s. Since the global  $A_T$  relationships first became available, a significant number of new  $A_T$  measurements have been added to the global data set. Here, we use surface  $A_T$  measurements ( $n = 5,692$ ) from the Global Ocean Data Analysis Project v1.1 data set [Key *et al.*, 2004] which have been carefully quality controlled to determine the relationships of  $A_T$  with SSS and SST for different ocean regimes. The functional form of the model used in our analysis of surface  $A_T$  results is also distinct from that used by Millero *et al.* [1998a]. In the previous analysis, the equations that relate salinity-normalized  $A_T$  to SST probably include erroneous trends

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**Table 1.** Regional Representations of Surface Total Alkalinity ( $A_T$ ) in the World's Oceans<sup>a</sup>

Zone	Ocean	Approximate Region <sup>b</sup>	$A_T$ , $\mu\text{mol kg}^{-1}$	$\sigma^c$	$N^d$
1	(Sub)tropics <sup>e</sup>	30°S–30°N (Atlantic, Indian, and Pacific Oceans excluding Zone 2), SST > 20°C, 31 < SSS < 38	$2305 + 58.66 (\text{SSS} - 35) + 2.32 (\text{SSS} - 35)^2 - 1.41 (\text{SST} - 20) + 0.040 (\text{SST} - 20)^2$	8.6	2756
2	Equatorial upwelling Pacific	75°W–110°W → 20°N–20°S and 110°W–140°W → 10°N–10°S, SST > 18°C, 31 < SSS < 36.5	$2294 + 64.88 (\text{SSS} - 35) + 0.39 (\text{SSS} - 35)^2 - 4.52 (\text{SST} - 29) - 0.232 (\text{SST} - 29)^2$	7.5	644
3	North Atlantic	30°N–80°N, 0°C < SST < 20°C, 31 < SSS < 37	$2305 + 53.97 (\text{SSS} - 35) + 2.74 (\text{SSS} - 35)^2 - 1.16 (\text{SST} - 20) - 0.040 (\text{SST} - 20)^2$	6.4	326
4	North Pacific	north of 30°N, SST < 20°C, 31 < SSS < 35	$2305 + 53.23 (\text{SSS} - 35) + 1.85 (\text{SSS} - 35)^2 - 14.72 (\text{SST} - 20) - 0.158 (\text{SST} - 20)^2 + 0.062 (\text{SST} - 20) (\text{LONG})$	8.7	258
5	Southern Ocean	30°S–70°S, SST < 20°C, 33 < SSS < 36	$2305 + 52.48 (\text{SSS} - 35) + 2.85 (\text{SSS} - 35)^2 - 0.49 (\text{SST} - 20) + 0.086 (\text{SST} - 20)^2$	8.4	1708

<sup>a</sup>For predicting  $A_T$  from SSS and SST values in a given zone, the primary criterion for choosing an appropriate  $A_T$  algorithms is the SST. That is, when the temperature of a water sample in the zone of interest is out of the range of the applicable equation, the equation for an adjacent zone sharing a common boundary should be used instead.

<sup>b</sup>Geographic boundaries between the South Pacific–South Atlantic, between the South Atlantic–South Indian, and between the South Indian–South Pacific, are 70°W, 20°E, and 120°E, respectively.

<sup>c</sup>Root mean square deviation =  $\{(\Delta^2/(N - 1))^{0.5}\}$ , where  $\Delta$  is the difference between measured  $A_T$  values and those calculated from the equations.

<sup>d</sup>The number of data points used for fitting.

<sup>e</sup>Arabian Sea and Bay of Bengal are included.

[Friis *et al.*, 2003]. Therefore, for each ocean regime we relate  $A_T$  results to both SSS and SST.

## 2. Methods

### 2.1. Optimal Polynomial Model for Fitting $A_T$ Data

[5] An optimal functional form for fitting  $A_T$  data for each ocean regime was chosen on the basis of results obtained from analysis of  $A_T$  data using the 10-fold cross validation method, which tests which polynomial form (first, second, or third order) gives a better fit [Stone, 1974; Breiman, 1996]. For each ocean regime, we randomly divided the  $A_T$  data set into 10 subsets. The first step of the 10-fold cross validation analysis was to remove 1 subset from the 10 data sets. Then we derived a regression model without this subset, predicted the output values for this subset using the derived regression model, and computed the residuals. This calculation routine was repeated for each subset and the squares of the resulting residuals were summed. This cross-validation analysis applied for polynomial models of first, second, and third order. We found that a second-order polynomial model yielded the lowest value of the sum of the squares of the residuals; hence we chose a second-order model as the optimal regression model for fitting  $A_T$  data for each ocean regime.

[6] We also tested whether the cross-correlation term between the two predictor variables (SSS and SST) should be included in the selected functional form. In all regimes, lower tolerance values (i.e., <0.2), calculated from  $1 - r^2$  for the regression of the interaction term on SSS and SST, indicate a high correlation between the two predictor variables and their interactions. Therefore, the interaction term was not included in the polynomial model that was used for generating  $A_T$  algorithms. However, for the North Pacific, the model including the interaction term between SST and longitude significantly reduced systematic errors.

### 2.2. Global Surface $A_T$ Data Used to Derive $A_T$ Algorithms

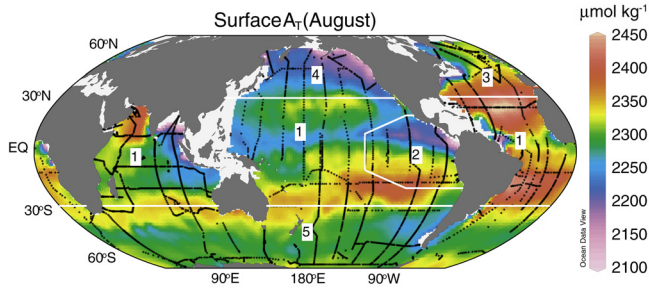
[7] Global surveys of marine inorganic carbon parameter distributions were conducted from 1990 to 1998 as part of the Joint Global Ocean Flux Study, the Ocean Atmosphere Carbon Exchange Study, and the World Ocean Circulation Experiment. The final  $A_T$  data set is internally consistent to  $\pm 5 \mu\text{mol kg}^{-1}$  [Key *et al.*, 2004]. Detailed descriptions of measurement techniques and calibration procedures for  $A_T$  data are provided elsewhere [Millero *et al.*, 1998b; Key *et al.*, 2004]. Subsets ( $n = 5,692$ ) of the final  $A_T$  data collected from waters of less than 20 m depth in the (Sub)tropics (i.e., 30°N–30°S) and 30 m depth at latitudes poleward of  $\sim 30^\circ$  were used in deriving region-specific correlations of surface  $A_T$  with SSS and SST.

[8] Monthly mean global  $A_T$  fields on  $1^\circ$  latitude  $\times$   $1^\circ$  longitude grid cells were estimated from five regional  $A_T$  relationships presented in Table 1, along with monthly mean SSS and SST fields from the *World Ocean Atlas 2001* [Conkright *et al.*, 2002].

## 3. Results and Discussion

### 3.1. Parameterization of Alkalinity in Surface Oceans

[9] Our analysis indicates that the distribution of surface  $A_T$  can be derived by dividing the global ocean into five regimes with corresponding equations relating  $A_T$  to SSS and



**Figure 1.** Monthly mean values of surface  $A_T$  ( $\mu\text{mol kg}^{-1}$ ) for August for the world's ocean, as estimated from the regional  $A_T$  algorithms given in Table 1 applied to monthly mean sea surface salinity (SSS) and temperature (SST) fields ( $1^\circ$  latitude  $\times$   $1^\circ$  longitude) from the *World Ocean Atlas 2001* [Conkright *et al.*, 2002]. White lines represent the approximate boundaries for the five different regions with unique relationships of  $A_T$  with SSS and SST and the numerical numbers represent the five regions listed in Table 1. Crosses denote locations of sampling stations.

SST (Table 1 and Figure 1). Since the derived equations given in Table 1 are expressed as functions of SSS and SST, the geographic boundaries of the regimes where the equations apply are dynamic, depending on seasonal variations in SST. For each ocean regime, surface  $A_T$  results are fitted to a second-order polynomial in SSS and SST and are also forced to satisfy equations for adjacent regime(s) which share boundaries with the regime of interest. The use of this functional form of equation is to ensure a smooth transition in  $A_T$  between adjacent regimes, and thus to minimize errors in  $A_T$  predictions at their mutual boundaries.

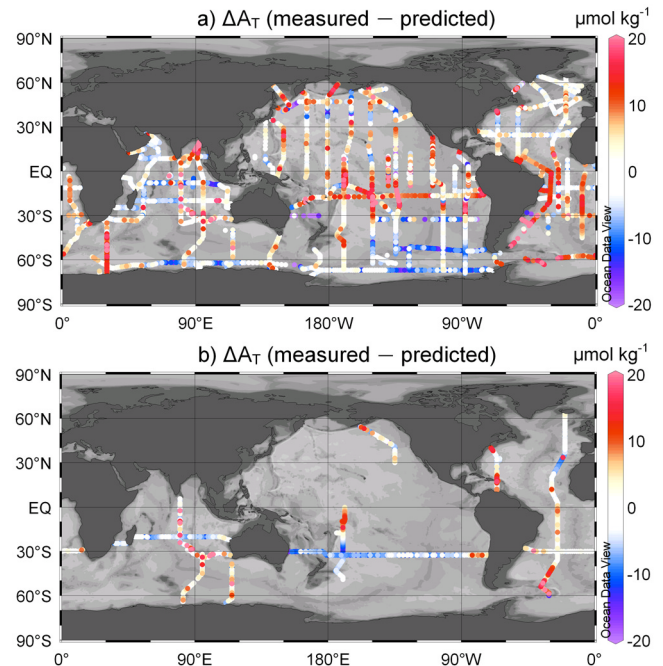
[10] In the (Sub)tropical oceans (Zone 1), except for the equatorial upwelling Pacific, waters with a temperature of  $20^\circ\text{C}$  and salinity of 35 have  $A_T$  values of about  $2305 \mu\text{mol kg}^{-1}$  for the entire (Sub)tropics. Thus, at  $\text{SST} = 20^\circ\text{C}$  and  $\text{SSS} = 35$ ,  $A_T$  values in these warm waters are forced to an intercept of  $2305 \mu\text{mol kg}^{-1}$  for the (Sub)tropics (see the first equation in Table 1). In the equatorial upwelling Pacific (Zone 2), waters have generally higher  $A_T$  values than those observed at the same salinity in similar latitude bands of the Atlantic and Indian Oceans. Thus, a separate equation is derived for this region to account for the effect of upwelling waters on the  $A_T$  trend. To make a smooth transition to values of  $A_T$  in the (Sub)tropics, the equation for this region is forced to an intercept of  $2294 \mu\text{mol kg}^{-1}$  at  $\text{SST} = 29^\circ\text{C}$  and  $\text{SSS} = 35$ . This intercept is similar to a mean  $A_T$  value for waters generally found at the boundary between the equatorial upwelling Pacific and the (Sub)tropics.

[11] In higher latitude oceans (poleward of  $30^\circ$ ), regionally contrasting sets of coefficients are derived for equations of the same functional form for the North Atlantic (Zone 3), North Pacific (Zone 4), and Southern Ocean (Zone 5). In the North Pacific, the  $A_T$  values in the northwestern Pacific are significantly higher than those observed at the same salinity in the northeastern Pacific. Such contrasting trends of  $A_T$  in the North Pacific can be attributed to the east-west differences in the intensity of convective mixing. The greater the intensity of seasonal convective mixing; the higher the  $A_T$ . Therefore, the variable “longitude” is added to the equation to account for east-west  $A_T$  differences in the North Pacific

(see the fourth equation in Table 1). The equations for the three zones at higher latitudes are all forced to the same intercept of  $2305 \mu\text{mol kg}^{-1}$  at  $\text{SST} = 20^\circ\text{C}$  and  $\text{SSS} = 35$  to ensure a smooth transition to values of  $A_T$  in the (Sub)tropics.

[12] The effect of seasonal differences in  $A_T$  data on the derived  $A_T$  relationships was also addressed by comparing  $A_T$  values predicted from the relationships in Table 1 with  $A_T$  measurements which are not included in the data set. The mean difference of  $-1.2 \pm 7.8 \mu\text{mol kg}^{-1}$  ( $1\sigma$ ) indicates that the derived  $A_T$  relationships can be used to predict seasonal extrapolation of surface  $A_T$ . Some differences are found locally (Figure 2b), but these are still close to uncertainties of our  $A_T$  relationships (see the fifth column in Table 1). The effect of seasonality on the  $A_T$  relationships can be more thoroughly evaluated when time-series measurements become available in high latitude regions in which seasonal variations in both salinity and convective mixing lead to contrasting changes in surface  $A_T$  values, either in terms of magnitude or sign.

[13] For predicting  $A_T$  from SSS and SST values in a given zone, the primary criterion for choosing an appropriate  $A_T$  algorithm is the SST. That is, when the temperature of a water sample in the zone of interest is out of the range of the applicable equation, the equation for an adjacent zone sharing a common boundary should be used instead. For example, for



**Figure 2.** Comparison of the estimated  $A_T$  values from the regional  $A_T$  algorithms given in Table 1 with measurements which are (a) used or (b) not used when deriving the  $A_T$  algorithms. The measurements which are not used when deriving the  $A_T$  algorithms given Table 1 include data from the three major ocean basins: I8N/I8S (January 1995, along  $\sim 80^\circ\text{E}$ ), I03I04 (2003, along  $\sim 20^\circ\text{S}$ ), P06 (2003, along  $\sim 30^\circ\text{S}$ ), P15S/SR05 (2001, along  $\sim 170^\circ\text{E}$ ), P17N (2001,  $135^\circ\text{W}$ – $160^\circ\text{W}$  and  $30^\circ\text{N}$ – $55^\circ\text{N}$ ), A10 (2003, along  $\sim 30^\circ\text{S}$ ), A16N/A16S (2003/2005, along  $\sim 20^\circ\text{W}$ ), and A22 (2003, along  $\sim 65^\circ\text{W}$ ).



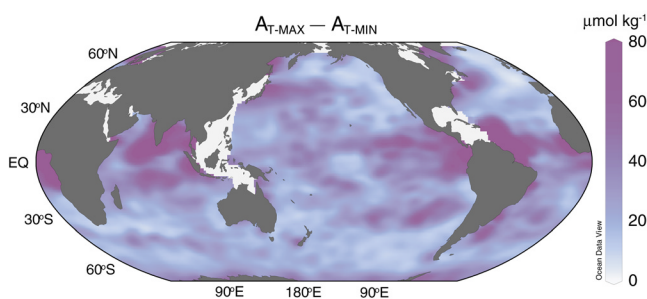
waters with SST < 20°C, which are commonly found in the northern (Sub)tropical Atlantic (Zone 1) during the winter season, the equation that applies to the North Atlantic (Zone 3) should be used to predict  $A_T$ .

### 3.2. Global Distribution of Alkalinity in the Surface Ocean

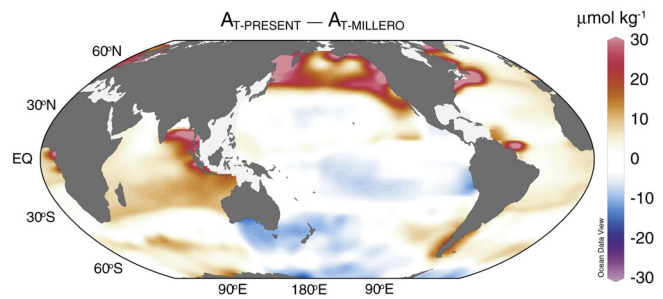
[14] The global distribution of surface  $A_T$  for August calculated using the derived  $A_T$  algorithms of Table 1 applied to climatological SSS and SST fields [Conkright *et al.*, 2002] is shown in Figure 1. The surface  $A_T$  field shows broad maxima in the central gyres centered at about 25° north and south of the Equator, similar to the pattern for salinity. Such strong salinity-dependent trends in surface  $A_T$  result from the fact that the concentrations of the key chemical species (i.e.,  $\text{HCO}_3^-$ ,  $\text{CO}_3^{2-}$ , and  $\text{B(OH)}_4^-$ ) that contribute to  $A_T$  proportionally increase with increasing salinity, and thus the ratio of  $A_T$  to salinity is approximately constant throughout the (Sub)tropics.

[15] Poleward of the (Sub)tropics, the annual excess precipitation over evaporation increases, and thus salinity decreases, with latitude. As a result, the values of  $A_T$  generally decrease with increasing latitude (Figure 1). Seasonal changes in the intensity of the convective mixing here are additional key factors that act to change surface  $A_T$  in a measurable way. During seasonal cooling, the intensive vertical mixing brings deep waters rich in  $\text{CaCO}_3$ -driven  $A_T$  to the surface, and thus increases surface  $A_T$ . During seasonal warming, however, the shoaling of the mixed layer makes the contribution of  $A_T$ -rich deep waters to the change in surface  $A_T$  minimal.

[16] The magnitude of seasonal  $A_T$  variability (Figure 3), computed by differencing the maximum and minimum monthly mean  $A_T$  values in each 1° latitude × 1° longitude grid cell, is directly proportional to the seasonal variations in salinity. Seasonal  $A_T$  variations are generally larger in the (Sub)tropics than in the higher-latitude oceans. In particular, larger amplitudes of seasonal variability are observed in areas where freshwater inputs through rivers (e.g., near the Amazon River and the Bay of Bengal) and the melting of ice (e.g., near sea-ice edges) are significant, or where tropical upwelling occurs.



**Figure 3.** Distribution of the seasonal amplitude (maximum  $A_{T\text{-MAX}}$  – minimum  $A_{T\text{-MIN}}$ ) of surface  $A_T$  predicted from the regional  $A_T$  relationships given in Table 1, applied to monthly mean sea surface salinity and temperature fields (1° latitude × 1° longitude) from the *World Ocean Atlas 2001* [Conkright *et al.*, 2002].



**Figure 4.** Differences ( $\mu\text{mol kg}^{-1}$ ) in values of surface  $A_T$  for August determined from the regional  $A_T$  algorithms listed in Table 1, versus those calculated from previously reported  $A_T$  algorithms [Millero *et al.*, 1998a].

[17] Comparison of the surface  $A_T$  values for August calculated using the five equations (Table 1) applied to climatological SSS and SST fields with the corresponding values estimated using the algorithms previously reported by Millero *et al.* [1998a] reveals that the degree of discordance between the two sets of estimates varies regionally (Figure 4). The greatest differences are found in the North Pacific and the North Atlantic and, to a lesser extent, in the Southern Ocean. Non-negligible differences also are observed in the subtropical gyres of the South Pacific and South Indian Oceans. In the (Sub)tropics, Millero *et al.* [1998a] forced the single equation to two different intercepts for SST at 20°C; one for the northern and southern (Sub)tropical Pacific and another for the rest of (Sub)tropics. However, our analysis indicates that, regardless of basins, only a single intercept of the  $A_T$  equation is needed at SSS = 35 and SST = 20°C for the entire (Sub)tropics.

[18] Finally, since the five regional  $A_T$  algorithms given in Table 1 were derived from the most comprehensive surface  $A_T$  data available using the laboratory-calibrated measurement protocol, we propose that this new set of  $A_T$  equations should be used for the prediction of surface  $A_T$  over large oceanic areas where SSS and SST values are known.

[19] **Acknowledgments.** We thank all individuals who contributed to the global alkalinity data set, including the chief scientists. This work was financially supported by the National Research Laboratory Program of the Korea Science and Engineering Foundation. Partial support was provided by the AEBRC at the POSTECH, the Korea Aerospace Research Institute, the MOEHRD (KRF-2005-070-C00143), the Korea Science and Engineering Foundation (R01-2002-000-00549-0, 2004), and NOAA's Office of Climate Observation; PMEL contribution 2938.

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