Unusual Alkalinity in Lake Carl Blackwell, Oklahoma.

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During a study of stable isotopes in Lake Carl Blackwell, Oklahoma, total alkalinity (TA) was measured at four stations over its thalweg during 2007. Based upon historical lake data, mean TA was four-five times higher than expected in February. Between then and September mean TA declined linearly from 10.8 to 2.29 meq/l. Baseline alkalinity in 1980-1981 was 2.40 to 2.96 meq/l. It is suggested that the decline in TA was caused mainly by evasion of CO_2 to the atmosphere. High TA may have been caused by mobilization of sedimentary carbonates by CO_2 at low water levels. Alternative sources of carbonates may have been anthropogenic. © 2012 Oklahoma Academy of Science.

INTRODUCTION AND METHODS

Recently, it has been proposed that impoundments are far more important in global carbon cycling than initially believed because of their unappreciated large aerial extent and their capacity to trap carbon in their sediments (Tranvik, et al. 2009, Downing, et al. 2006 and others). Both large and small reservoirs are ubiquitous in Oklahoma, but their carbon cycles have not been well described. To close this gap the temporal and spatial distribution of the stable isotope, δ^{13} C, was described for Lake Carl Blackwell (LCB) (Toetz, et al. 2009). However, detailed analysis of total alkalinity (TA) concentrations, which were collected to support the interpretation of the stable isotope data, were not done. Usually, seasonal changes in TA in lakes are modest and often predictable. However, in the research above a much higher TA was observed when samples were first collected and thereafter a much more rapid change in TA was observed than was expected based upon historical lake data. The purpose of this paper is to describe this phenomenon and suggest its cause.

Lake Carl Blackwell is located in Payne County, Oklahoma. There is no major groundwater input. Its area is 1250 ha and has a maximum depth of 15.0 m and a mean depth 4.9 m. High turbidity is maintained by resuspension of sediments augmented by strong winds and shallow depth.

MATERIALS AND METHODS

Sampling of the lake and laboratory methods have been previously described (Toetz, et al. 2009). Only those aspects that pertain to this paper are given. LCB was sampled at four stations over the thalweg approximately monthly between February and September, 2007.

Samples were taken at 0.5 and 3 m with a Kemmerer bottle and pooled to obtain a composite water sample at all stations, except for the most upstream one where samples were taken at one meter. Filtered water (0.5μ m pore size) was used the same day or the next to determine total alkalinity (TA) which was measured by titration with $0.02 \text{ N H}_2\text{SO}_4$ to an endpoint of pH 4.8. Linear regression was performed with an Excel 2003 spreadsheet.

RESULTS AND DISCUSSION

Mean TA decreased from 10.80 to 2.29 meq/l between February and September (Table 1). The relationship of TA with time (days) was negative and significant (slope= - 0.04, P<0.00, n=24, R²=0.96). Between February Proc. Okla. Acad. Sci. 92: pp 78-80 (2012)

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Table 1. Mean total alkalinity at four stations during 2007 in Lake Carl Blackwell, Oklahoma. Standard errors are in parentheses.

Date	Mean Total Alkalinity meq/l
February 25	10.80 (0.27)
May 16	8.27 (0.18)
June 20	6.47 (0.65)
July 20	3.39 (0.10)
August 24	3.77 (0.16)
September 24	2.29 (0.03)

and September, 1981, there was a slight increase in TA from 2.42 to 2.96 meq /l (Howick, et al. 1984). Surface waters of hardwater lakes often exhibit a small reduction in TA caused by photosynthetic withdrawal of CO_2 , but the magnitude of the decline is only a few meq/l (Otsuki and Wetzel, 1974 and others).

TA in lake water is affected by exchange of carbon dioxide (CO₂) with the atmosphere, inputs from weathering in soils in the watershed, by stream flow or groundwater, photosynthetic withdrawals of CO₂, additions from respiration, and losses in effluents or sediments (Bade et al. 2004, Quay et al. 1986 and others). A carbon budget would be desirable to explain the decline in TA we observed. Nevertheless, the linear decline in TA during 2007 suggests that the major factor responsible for it operated at a constant rate for seven months and might be due to some physical factor such as evasion of CO₂ from the surface of the lake and precipitation of calcite to the sediments. In many lakes CO₂ is supersaturated with respect to the atmosphere and some lakes experience seasonal evasions of CO₂ (Wetzel, 2001, Quay, et al. 1986, Finlay, et al. 2009 and others). However, some loss of TA could also have been due to loss in effluents and dilution by influents. It is possible that

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in February the CO_2 -bicarbonate-carbonate system was not in equilibrium with the atmosphere and that there was a return to equilibrium conditions by September.

The high concentrations of TA observed in February may be due to by high concentrations of CO_2 and some source of carbonates in lake water that occurred at low water levels before 2007. At low water levels, LCB can be heterotrophic (Carroll 1975). This could lead to an ample supply of "aggressive" CO_2 , which would cause solid carbonates in the lake to dissolve. But, what would be a source of carbonates that "aggressive" CO_2 might attack?

Prior to 2007 sources of carbonate could have been natural (calcite in sediments) or anthropogenic. There was addition of rocks and broken concrete to the rip-rap of the dam and the construction of new breakwaters. There were also a large pile of broken concrete and dry-wall from building demolition that was deposited near the shore. Another source might have been deposition of aeolian dust from roads and/or concretemixing plants near the lake. Aeolian dust can be a source of calcite and as a result increase the alkalinity of acidic soils (Litaor 1986). Although the reason(s) for high TA in February will never be known, the abrupt decline in TA can be best explained by an equally abrupt cessation of input(s) of carbonate from anthropogenic sources.

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