# **DUST DEPOSITION TO THE SARGASSO SEA:** A COMPARISON OF ESTIMATES USING SEASONALLY-RESOLVED MEASUREMENTS OF ALUMINUM IN THE **UPPER WATER-COLUMN VERSUS AEROSOLS AND RAINWATER COLLECTED ON BERMUDA** Tara Williams<sup>\*1</sup>, Peter Sedwick<sup>1</sup>, Bettina Sohst<sup>1</sup>, Joe Resing<sup>2</sup>, Kristen Buck<sup>3</sup>, Salvatore Caprara<sup>3</sup>, Rod Johnson<sup>4</sup>, Dan Ohnemus<sup>5</sup>, Laura Sofen<sup>6</sup>, Alessandro Tagliabue<sup>7</sup>, Ben Twining<sup>6</sup>

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## INTRODUCTION

- Dust deposition is a major source of bioactive trace elements to the surface ocean, yet remains difficult to quantify in space and time
- Time-averaged dust fluxes have been estimated using surface ocean dissolved aluminum (DAI) concentrations together with assumed values for the fractional solubility of aerosol aluminum (%Al<sub>S</sub>) and the replacement time of DAl in the surface mixed layer  $(\tau_{AI})^1$
- We have obtained seasonally-resolved measurements of DAL in the upper water column in the BATS region, along with ~weekly samples of aerosols and rainwater from the Tudor Hill tower on Bermuda
- Analysis of these samples allows a comparison of two independent estimates of dust deposition, based on (1) the water-column DAL inventory, and (2) total Al in aerosols and rain from Tudor Hill



**Figure 1**. Tudor Hill tower (left), map of Bermuda and BATS sampling region (center), and Research Vessel *Atlantic Explorer* (right)

## SAMPLING AND ANALYSES

- Water-column samples were collected from the BATS site and adjacent spatial stations (Fig. 1) during cruises in March (spring), May (early summer), August (late summer) and November (fall) 2019
- Bulk aerosols and rainwater were collected over the same period on the Tudor Hill tower on Bermuda, using a high-volume aerosol sampler with Whatman-41 filters and an automatic rain collector
- Water-column samples were filtered through 0.2 µm-pore Acropak Supor capsule filters, and DAI was determined in the filtered water column samples using flow-injection analysis<sup>2</sup>
- Aerosol sample portions were subjected to strong-acid microwave digestion, and to leaching with ultrapure deionized water (DIW) then 25% acetic acid (HOAc)<sup>3</sup>; rainwater samples were acidified to pH < 2
- Al in aerosol digests, leach solutions and acidified rainwater was measured using inductively-coupled plasma mass spectrometry
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# RESULTS

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## Aerosol- and rain-based dust flux estimates

- Dry dust deposition was estimated from total aerosol Al loadings and an assumed dust deposition velocity<sup>4</sup> of 1 cm/s (Fig. 2), using an Al crustal abundance of  $8.1\%^5$ , yielding an average flux of  $0.67 \text{ g/m}^2/\text{y}$
- Wet dust deposition was estimated from total-dissolvable Al in rain and measured rainwater volumes (Fig. 2), and an Al crustal abundance of 8.1%, yielding an average flux of 0.51 g/m<sup>2</sup>/y
- These estimates then yield an average total (dry+wet) dust deposition flux of 1.18 g/m<sup>2</sup>/y over our 318 day sampling period



**Figure 2**. Dry and wet deposition of total Al for each sampling period, as estimated from aerosols and rain collected at Tudor Hill tower; dust mass fluxes were derived assuming 8.1% crustal abundance of Al



**Figure 3**. Water-column profiles of DAL concentration from four seasonal BAIT cruises in the BATS region (MLD = mixed-layer depth)







### Water-column DAI-based dust flux estimates



**Figure 4**. Comparison of dust flux estimates based on water-column DAL inventories (bars) using  $%Al_{S}$  based on aerosol leaches using (A) HOAc and (B) DIW, with estimates based on total Al in aerosols and aerosols+rainwater. Also shown are fluxes estimated by other studies.

- - $\tau_{Al} \sim 49$  years (for Al<sub>s</sub> estimated from DIW leaches)  $\tau_{AI} \sim 23$  years (for Al<sub>s</sub> estimated from DIW + HOAc leaches)

These residence times are higher than an estimate based on thorium supply<sup>6</sup>, but are compatible with values extracted from a recent data assimilation modeling study<sup>7</sup>.

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 Water-column profiles of DAI (Fig. 3) from the BATS region reveal concentrations ranging from ~30 nM to ~40 nM during the year, as mixed layer depths (MLD) shoaled from ~150-200 m to ~10-30 m

• Assuming a mixed-layer  $\tau_{AI}$  value of 5 years<sup>1</sup>, average %Al<sub>S</sub> values based on our Tudor Hill aerosol leaches, and Al crustal abundance of 8.1%<sup>5</sup>, DAl inventories over the upper 200 m yield mean dust fluxes: 11.1 g/m<sup>2</sup>/y (based on average %Al<sub>S</sub> using DIW leach = 3.8%) 5.2 g/m<sup>2</sup>/y (based on average %Al<sub>S</sub> using HOAc leach = 8.1%)

• These values are ~5-10 fold higher than our estimates based on aluminum in aerosols and rain (Fig. 4), and may reflect uncertainties in aerosol deposition velocity or, more likely, the assumed  $au_{AI}$  value

### IMPLICATIONS

 One way to reconcile the aerosol- and rain-based dust flux estimates with those based on water-column DAl inventories is to require a longer replacement time for DAL in the upper 200 m

• We can estimate this  $\tau_{AI}$  value by dividing the average DAL inventory of the upper 200m by the annualized-average deposition of soluble Al in aerosols and rain based on our Tudor Hill aerosol and rain data:

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