### Dry Deposition of Atmospheric Mercury to the Great Salt Lake

#### Kevin D. Perry<sup>1</sup>, Joel R. Lisonbee<sup>2</sup>, and Eric R. Pardyjak<sup>3</sup>

*<sup>1</sup>Department of Atmospheric Sciences, University of Utah <sup>2</sup>Darwin Climate Service Center, Australian Bureau of Meteorology <sup>3</sup>Department of Mechanical Engineering, University of Utah*



October 10, 2013

## Outline

- Goals and Objectives
- AMNet Site Description
- Annual, Seasonal, and Diurnal Mercury Variations
- Dry Deposition Model Description
- Dry Deposition Estimates
- Conclusions

## Objectives (2009-2012)

1) Characterize the annual, seasonal, and daily variations of speciated atmospheric mercury concentrations near the Great Salt Lake

2) Determine whether the UT96 site is representative of urban, rural or mixed conditions for atmospheric Hg

3) Use data from the UT96 site to estimate the dry deposition of Hg to the Great Salt Lake

### AMNet Sites



### UT96 Site Location



## Meteorological Wind Rose



#### Instrumentation

- Tekran Mercury Monitoring System
- Micrometeorological Measurement System
- 8-Stage Rotating Drum Impactor

#### Speciated Mercury and PM Measurements



8-Stage Drum

## Campbell Scientific Inc. CSAT 3D Sonic Anemometer

- 7.44 m height
- 10 Hz measurements
	- 3D wind
	- Temperature
- CR 1000 datalogger





#### GEM Seasonal Variations (July 1, 2009- June 30, 2012)



#### GOM Seasonal Variations (July 1, 2009- June 30, 2012)



#### PBM Seasonal Variations (July 1, 2009- June 30, 2012)



#### GEM Seasonal Variations (July 1, 2009- June 30, 2012)



# Diurnal GEM Cycles



Afternoon minimum could result from:

- 1) Higher deposition velocities during afternoon
- 2) Chemical transformation of GEM to GOM and/or PBM
- 3) Dilution of GEM through entrainment of free-tropospheric air

# Diurnal GOM Cycles



Afternoon maximum could result from:

- 1) Chemical transformation of GEM to GOM
- 2) Entrainment of air from the freetroposphere

# Diurnal PBM Cycles



Afternoon minimum could result from:

- 1) Higher deposition velocities during afternoon
- 2) Conversion of PBM to GOM and/or GEM
- 3) Dilution of PBM through entrainment of free-tropospheric air

#### Mercury Deposition Dry Wet

Speciated Mercury (GEM, GOM, PBM)

> Micro-Met Tower

**Estimated Particulate** Mercury Size **Distribution** 

Inferential Model of Dry Deposition for GEM and GOM

Inferential Model of Dry Deposition for PBM



Hg Wet **Deposition** 

(UTAH DEQ)

# Hg Dry Deposition Flux (F<sub>Ha</sub>)

$$
F_{Hg} = -V_d C_{Hg}
$$

 $V_{d}$  = deposition velocity *CHg* = speciated mercury concentration

## Resistance Model for V<sub>d</sub>

$$
V_d = \frac{1}{R_a + R_b + R_c + R_a R_b V_s} + V_s
$$

*R<sup>a</sup>* = aerodynamic resistance *R<sup>b</sup>* = boundary layer resistance *R<sup>c</sup>* = surface layer resistance *V<sup>s</sup>* = settling velocity (for particles only)



# Aerodynamic Resistance (R<sub>a</sub>)

$$
\zeta_0 = \frac{z_0}{L}
$$
  
\n
$$
R_a = \begin{cases} \frac{\text{Pr}}{\kappa u_*} \left[ \ln \left( \frac{z_1}{z_0} \right) + 4.7(\zeta - \zeta_0) \right] & \text{(stable)} \end{cases}
$$
  
\n
$$
R_a = \begin{cases} \frac{\text{Pr}}{\kappa u_*} \ln \left( \frac{z_1}{z_0} \right) & \eta_0 = \sqrt{(1 - 9\zeta_0)} \end{cases}
$$
  
\n
$$
\frac{\text{Pr}}{\kappa u_*} \left[ \ln \left( \frac{z_1}{z_0} \right) + 2 \left( \frac{(1 + \eta_0)}{(1 + \eta)} \right) \right] & \text{(unstable)} \end{cases}
$$
  
\n
$$
\eta = \sqrt{(1 - 9\zeta)}
$$

# Boundary Layer Resistance (Rb)

Gases



# Boundary Layer Resistance (Rb) Particles



# Surface Layer Resistance (R<sub>c</sub>)

$$
R_C = \frac{1}{K_L H_A} + \frac{1}{K_G}
$$

 $K_i$  = liquid-phase mass transport coefficient *K<sub>G</sub>* = gas-phase mass transport coefficient *H<sup>A</sup>* = Dimensionless Henry's Law coefficient

### Monthly-Averaged Dry Deposition Velocities







## Flux Comparison





### Dry Deposition Totals  $(\mu g m^{-2} yr^{-1})$



# Mercury Influx Pathways

- Dry Deposition
	- $-10.5$  µg m<sup>-2</sup> yr<sup>-1</sup>
- Wet Deposition (MDN UT DEQ)  $-8.1 \mu g m^{-2} yr^{-1}$
- Riverine influx (Naftz et al. 2009)  $-1.9 \mu g m^{-2} yr^{-1}$
- Coarse PBM (Carling et al. 2012)  $-$  3  $\mu$ g m<sup>-2</sup> yr<sup>-1</sup>
- Total

 $-23.5$  µg m<sup>-2</sup> yr<sup>-1</sup>

## Conclusions

- The UT96 site is a mixed rural/urban receptor site
- The UT06 site is periodically impacted by SIGNIFICANT local sources
- Dry deposition is the dominant influx pathway for mercury to the Great Salt Lake accounting for 45% of the total Hg input
- Wet deposition is responsible for 34% of the total Hg input
- Riverine input is responsible for 8% of the total Hg input
- Coarse PBM could contribute more than 10% of the total Hg input