

# Conditional complexation constants for $\text{Hg}^{2+}$ and $\text{CH}_3\text{Hg}^+$ in natural organic matter. (S02-bloom123738-Oral)

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## Abstract:

Mercury  $2+$  and methyl-Hg bind strongly with natural organic matter (NOM) in soils and waters. Our previous x-ray spectroscopy studies showed that in NOM reduced sulfur is involved in very strong binding of  $\text{Hg}^{2+}$  and methyl-Hg when these ions are added at low Hg to reduced S ratios. Because of the strong binding it is necessary to use competitive ligands,  $\text{Br}^-$ , EGTA, or DTPA, for the determination equilibrium binding strength. Bromide works well for methyl-Hg yielding apparent complexation constants in the range of  $10^{16}$  for soil organic matter (SOM). For  $\text{Hg}^{2+}$  the apparent  $K_d$  for adsorption in SOM is in the range of  $10^{22}$  to  $10^{23}$  but we found evidence of adsorption of Hg-bromide complexes. EGTA and DTPA do not complex as strongly as  $\text{Br}^-$  but these ligands do increase the Hg in solution and there is little possibility of multi-ligand complex formation with NOM. The apparent  $K_d$  values were lower than for  $\text{Br}^-$ . However, the two ligands gave quite different  $K_d$  results for soil HA;  $10^{20}$  for EGTA and  $10^{15}$  for EGTA. Part

of the problem may be non equilibrium conditions for the added  $\text{Hg}^{2+}$ . After heating the  $\text{HA-Hg}^{2+}$  complex to  $70^\circ\text{C}$  for a minimum of 3 days the binding strength increased such that EGTA and DTPA were not strong enough complexing agents to determine  $K_d$ .

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