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Corrigendum

Corrigendum to Mercury photoreduction and photooxidation in lakes: effects of filtration and dissolved organic carbon concentration. Journal of Environmental Sciences 68 (2018) 151-159

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The authors regret that,

Abstract:

“Filtered (0.2 μm) and unfiltered samples were analyzed for gross photoreduction, gross photooxidation, and net reduction rates of mercury using pseudo first-order curves. Unfiltered samples had higher concentrations ($p=0.04$) of photo-reducible divalent mercury ($\text{Hg(II)}_{\text{RED}}$) (mean of 754 ± 253 pg/L) than filtered samples (mean of 482 ± 206 pg/L); however, gross photoreduction and photooxidation rate constants were not significantly different in filtered or unfiltered samples in early summer. DOC was not significantly related to gross photoreduction rate constants in filtered ($R^2=0.43$; $p=0.08$) and unfiltered ($R^2=0.02$; $p=0.71$) samples; DOC was also not significantly related to gross photooxidation rate constants in filtered or unfiltered samples. However, DOC was significantly negatively related with $\text{Hg(II)}_{\text{RED}}$ in unfiltered ($R^2=0.53$; $p=0.04$), but not in filtered samples ($R^2=0.04$; $p=0.60$).”

Should be revised to

“Filtered (0.2 μm) and unfiltered samples were analyzed for gross photoreduction, gross photooxidation, and net reduction rates of mercury using pseudo first-order curves. Unfiltered samples had higher concentrations ($p=0.037$) of photo-reducible divalent mercury ($\text{Hg(II)}_{\text{RED}}$) (mean of 801 ± 260 pg/L) than filtered samples (mean of 502 ± 225 pg/L); however, gross photoreduction and photooxidation rate constants were

not significantly different in filtered or unfiltered samples in early summer. DOC was weakly related to gross photoreduction rate constants in filtered ($R^2=0.41$; $p=0.08$) and unfiltered ($R^2=0.45$; $p=0.07$) samples; DOC was also not significantly related to gross photooxidation rate constants in filtered or unfiltered samples. However, DOC was weakly and negatively related with $\text{Hg(II)}_{\text{RED}}$ in unfiltered ($R^2=0.53$; $p=0.07$), but not in filtered samples ($R^2=0.03$; $p>0.10$).”

Section 1.2:

$$\text{Hg}(0)_t = [\text{Hg(II)}_{\text{RED}}]_0 - e^{kt} [\text{Hg(II)}_{\text{RED}}]_0 \quad (2)$$

This can be rewritten as:

$$\text{Hg}(0)_t = [\text{Hg(II)}_{\text{RED}}]_0 (1 - e^{kt}) \quad (3)$$

Should be revised to

$$\text{Hg}(0)_t = [\text{Hg(II)}_{\text{RED}}]_0 - e^{-kt} [\text{Hg(II)}_{\text{RED}}]_0 \quad (2)$$

This can be rewritten as:

$$\text{Hg}(0)_t = [\text{Hg(II)}_{\text{RED}}]_0 (1 - e^{-kt}) \quad (3)$$

Section 1.3:

“Once the gross reduction and net reduction rate constants were determined, gross photooxidation rate constants were derived by subtracting net photoreduction data points from gross photoreduction data, and a pseudo first order reaction

DOI of original article: [10.1016/j.jes.2017.12.010](https://doi.org/10.1016/j.jes.2017.12.010)

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<https://doi.org/10.1016/j.jes.2021.02.005>

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equation was fitted to the resulting curve using Sigma-Plot 12.0 (Fig. 2)."

Should be revised to

"Once the gross reduction and net reduction rate constants were determined, gross photooxidation rate constants were derived by subtracting net photoreduction rate constants from gross photoreduction rate constants. Curves of gross photooxidation can then be derived using the resulting equation for a pseudo first order reaction (Fig. 2)."

Section 2.1:

"In all lakes, unfiltered samples had significantly higher concentrations of photoreducible mercury, $\text{Hg(II)}_{\text{RED}}$ (mean = 754 ± 253 pg/L) than 0.2 μm filtered samples (mean = 482 ± 206 pg/L) (t-test; $p = 0.04$; Shapiro Wilk normality $P = 0.63$)."

Should be revised to

"In all lakes, unfiltered samples had significantly higher concentrations of photoreducible mercury, $\text{Hg(II)}_{\text{RED}}$ (mean = 801 ± 260 pg/L) than 0.2 μm filtered samples (mean = 502 ± 225 pg/L) (t-test; $p = 0.04$; Shapiro Wilk normality $P = 0.61$).

Section 2.2

"In this work, gross photoreduction rate constants ranged from $1.63 \times 10^{-3} \text{ hr}^{-1}$ to $3.42 \times 10^{-3} \text{ hr}^{-1}$ in filtered samples, and $1.29 \times 10^{-3} \text{ hr}^{-1}$ to $2.93 \times 10^{-3} \text{ hr}^{-1}$ in unfiltered samples, for lakes sampled in May of 2008 and 2009 (Table 1). Similar results were found for the gross photooxidation rate constants, which ranged between $1.42 \times 10^{-3} \text{ hr}^{-1}$ and $3.04 \times 10^{-3} \text{ hr}^{-1}$ for filtered samples, and $1.21 \times 10^{-3} \text{ hr}^{-1}$ to $2.78 \times 10^{-3} \text{ hr}^{-1}$ for unfiltered samples (Table 1). It can be seen in Table 1 that the lakes sampled late in the summer season (August of 2010; Pebbleogitch, Peskowsk, and Beaverskin lakes) have significantly larger gross photoreduction and gross photooxidation rate constants than the lakes sampled in early summer (Table 1). It is interesting to note, however, that the net photoreduction rate constants are in a similar range for all lakes; this suggests that while the rate of both photooxidation and photoreduction reactions has increased, the overall effect on the net mercury reduction rate constant did not change substantially, due to the close balance of photoreduction and photooxidation reactions occurring in these lakes. The near-balance of mercury photooxidation and photoreduction observed in these lakes may suggest a common linking component between lakes, such as the atmospheric deposition of reactive mercury being the dominant mercury pool undergoing these photoreactions (Orihel et al., 2007).

Our data show that gross photoreduction (mean = $2.46 \times 10^{-3} \text{ hr}^{-1}$, SD = $6.72 \times 10^{-4} \text{ hr}^{-1}$ for filtered; mean = $2.07 \times 10^{-3} \text{ hr}^{-1}$, SD = $5.08 \times 10^{-4} \text{ hr}^{-1}$ for unfiltered) and photooxidation rate constants (mean = $2.04 \times 10^{-3} \text{ hr}^{-1}$, SD = $5.57 \times 10^{-4} \text{ hr}^{-1}$ for filtered; mean = $2.00 \times 10^{-3} \text{ hr}^{-1}$, SD = $4.76 \times 10^{-4} \text{ hr}^{-1}$ for unfiltered) are not significantly different between filtered and unfiltered lake waters, sampled in early summer (respective t-tests; $p = 0.2$; $p = 0.8$); this demonstrates that the rate constants of these mercury photoreactions are not significantly affected by the presence of particles or particle-bound mercury species in solution. This result is in contrast to the results for photoreducible mercury amounts ($\text{Hg(II)}_{\text{RED}}$) presented above, which do show significant differences between filtered and unfiltered

samples. This lack of an effect on rate constants by the presence or absence of 0.2 μm filterable material supports the conclusions of Qureshi et al. (2010) and Beucher et al. (2002), who observed no substantial influence of filtration (or biotic activity) on mercury reduction rate constants in ocean water. Relative standard error associated with the derivation of the rate constant for gross photoreduction was <1% in all cases and was higher for gross photooxidation (<10% in all cases except 34% for BDE). The higher error on the gross photooxidation results are a result of the higher error associated with net photoreduction experiments due to the low masses of mercury being quantified. There is very limited data available in the literature for mercury photooxidation rate constants. Lalonde et al. (2001) found the net mercury photooxidation rate constant for a freshwater river was 0.26 hr^{-1} which is lower but not a good point of comparison for the gross photooxidation rates measured here. Our results are more comparable to the work of Garcia et al. (2005b) who calculated gross mercury photooxidation rate constants for freshwaters ranging from 0.02 to 0.07 hr^{-1} ."

Should be revised to

"In this work, gross photoreduction rate constants ranged from $1.27 \times 10^{-1} \text{ hr}^{-1}$ to $2.66 \times 10^{-1} \text{ hr}^{-1}$ in filtered samples, and $1.01 \times 10^{-1} \text{ hr}^{-1}$ to $2.29 \times 10^{-1} \text{ hr}^{-1}$ in unfiltered samples, for lakes sampled in May of 2008 and 2009 (Table 1). Similar results were found for the gross photooxidation rate constants, which ranged between $1.08 \times 10^{-1} \text{ hr}^{-1}$ and $4.41 \times 10^0 \text{ hr}^{-1}$ for filtered samples, and $1.29 \times 10^{-1} \text{ hr}^{-1}$ to $1.76 \times 10^0 \text{ hr}^{-1}$ for unfiltered samples (Table 1). It can be seen in Table 1 that the lakes sampled late in the summer season (August of 2010; Pebbleogitch, Peskowsk, and Beaverskin lakes) have similar gross photoreduction and gross photooxidation rate constants as compared to the lakes sampled in early summer (Table 1). The near-balance of mercury photooxidation and photoreduction observed in these lakes may suggest a common linking component between lakes, such as the atmospheric deposition of reactive mercury being the dominant mercury pool undergoing these photoreactions (Orihel et al., 2007).

Our data show that gross photoreduction (mean = $1.95 \times 10^{-1} \text{ hr}^{-1}$, SD = $5.19 \times 10^{-2} \text{ hr}^{-1}$ for filtered; mean = $1.67 \times 10^{-1} \text{ hr}^{-1}$, SD = $3.65 \times 10^{-2} \text{ hr}^{-1}$ for unfiltered) and photooxidation rate constants (mean = $1.05 \times 10^0 \text{ hr}^{-1}$, SD = $1.32 \times 10^0 \text{ hr}^{-1}$ for filtered; mean = $7.2 \times 10^{-1} \text{ hr}^{-1}$, SD = $4.84 \times 10^{-1} \text{ hr}^{-1}$ for unfiltered) are not significantly different between filtered and unfiltered lake waters, sampled in early summer (respective t-tests; $p = 0.27$; $p = 0.87$); this demonstrates that the rate constants of these mercury photoreactions are not significantly affected by the presence of particles or particle-bound mercury species in solution. This result is in contrast to the results for photoreducible mercury amounts ($\text{Hg(II)}_{\text{RED}}$) presented above, which do show significant differences between filtered and unfiltered samples. This lack of an effect on rate constants by the presence or absence of 0.2 μm filterable material supports the conclusions of Qureshi et al. (2010) and Beucher et al. (2002), who observed no substantial influence of filtration (or biotic activity) on mercury reduction rate constants in ocean water. Relative standard error associated with the derivation of the rate constant for gross photoreduction was <1% in all cases and was substantially higher for net photoreduction and gross

photooxidation (ranging 34%–1747 %). The high error on the gross photooxidation results are a result of the high error on curve fitting associated with net photoreduction experiments due to the low masses of mercury being quantified. There is also some indication of declines in Hg(0) concentrations in the net photoreduction experiments after 12 hr such that a pseudo first order reaction equation was not always a good fit (Appendix A Fig. SI-5). There is very limited data available in the literature for mercury photooxidation rate constants. Lalonde et al. (2001) found the net mercury photooxidation rate constant for a freshwater river was 0.26 hr^{-1} which is lower but not a good point of comparison for the gross photooxidation rates measured here. Our results are more comparable to the work of Garcia et al. (2005b) who calculated gross mercury photooxidation rate constants for freshwaters ranging from 0.02 to 0.07 hr^{-1} .”

Section 2.3

“The mean concentration of net Hg(0) (i.e. DGM) measured in water from all lakes was approximately 8.5% of the total Hg(II)_{RED} for the gross photoreduction experiments (ranging 15 to 80 pg/L; mean = $41 \pm 17.5 \text{ pg/L}$). This result indicates that there is a much larger capacity for mercury photoreduction and volatilisation from these lakes that might be released with decreases in gross photooxidation kinetics. The net mercury reduction rate constants derived from the data for filtered samples (ranging 1.28×10^{-5} to $8.30 \times 10^1 \text{ hr}^{-1}$; mean = $1.15 \times 10^1 \text{ hr}^{-1}$; SD = $2.71 \times 10^1 \text{ hr}^{-1}$) and for unfiltered samples (ranging 3.20×10^{-1} to $1.93 \times 10^0 \text{ hr}^{-1}$; mean = $8.77 \times 10^{-1} \text{ hr}^{-1}$; SD = $4.96 \times 10^{-1} \text{ hr}^{-1}$) were more variable than those derived from the gross photoreduction experiments (Table 2). Many studies have focussed on the *in situ* net photoreduction of mercury in aquatic systems; for example, Poulain et al. (2004) determined that the DGM formation rate constant ranged from 0.76 – 1.4 h^{-1} in a wetland area, and 0.21 – 0.47 h^{-1} for a pelagic area. Another study by Amyot et al. (1994) determined the mean net mercury reduction rate constant to be 0.10 h^{-1} . A review by Vost et al. (2011) found that net mercury reduction rate constants ranged between 0.1 and 2.2 h^{-1} for freshwater samples, which are within the range of net mercury photoreduction constants measured in this study for unfiltered water samples. However, the unfiltered water samples show slower rates possibly due to slow release of reducible mercury from solid particles in unfiltered samples similar to what has been proposed between soil particles and soil solution (Pannu et al., 2014).”

Should be revised to

“In addition, the decline of Hg(0) after 12 hr in some experiments suggest a pseudo first order may not be best fit for these data (Appendix A Fig. SI-5). The mean concentration of net Hg(0) (i.e. DGM) measured at the end of each 24 h experiment in water from all lakes ranged from 1.6% to 22.7% of the total Hg(II)_{RED} for the gross photoreduction experiments (ranging 7 to 119 pg/L; mean = $30 \pm 24 \text{ pg/L}$). This result indicates that there is a much larger capacity for mercury photoreduction and volatilisation from these lakes that might be released with decreases in gross photooxidation kinetics. The net mercury reduction rate constants derived from the data for filtered samples (ranging 2.49×10^{-1} to $4.6 \times 10^0 \text{ hr}^{-1}$; mean = $1.24 \times 10^0 \text{ hr}^{-1}$; SD = $1.32 \times 10^0 \text{ hr}^{-1}$) and for unfiltered samples (ranging 3.18×10^{-1} to $1.99 \times 10^0 \text{ hr}^{-1}$;

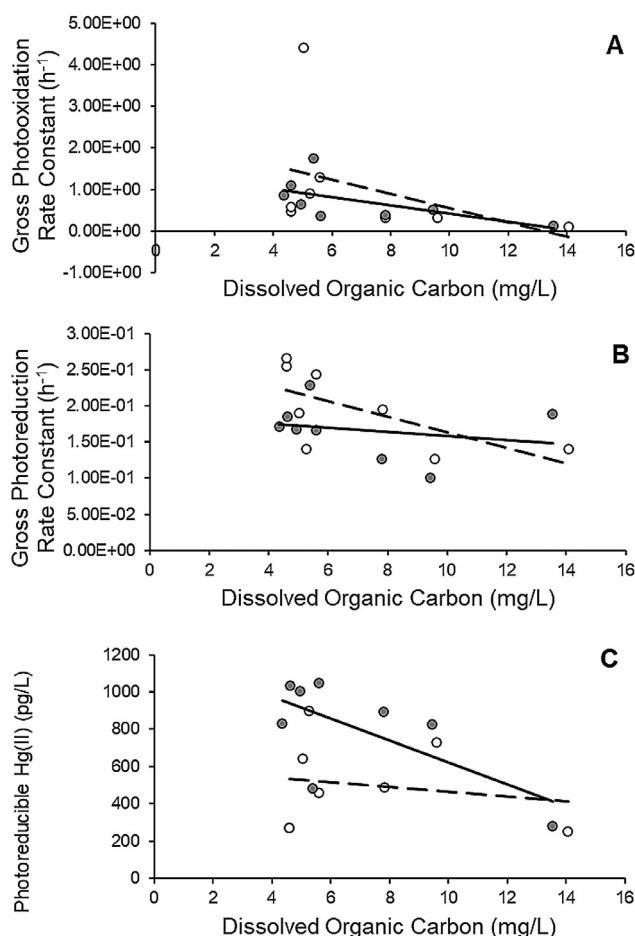


Figure 3 – DOC (mg/L^{-1}) plotted against (A) the gross photooxidation rate constant (hr^{-1} ; filtered $R^2=0.16$, $p>0.10$ and unfiltered $R^2<0.36$, $p>0.10$) (B) the gross photoreduction rate constant (hr^{-1} ; filtered $R^2=0.42$, $p=0.08$ and unfiltered $R^2=0.06$, $p>0.10$), and (C) photoreducible Hg(II) (pg/L ; filtered $R^2=0.03$, $p>0.10$ and unfiltered $R^2=0.45$, $p=0.07$) for both filtered (open circles) and unfiltered samples (shaded circles), respectively, from 7 lakes in KNP (lakes sampled in 2008 and 2009). No significant relationship was observed with photooxidation rate constants. Lakes sampled in August 2010 have substantially larger gross photoreduction rate constants (see Table 1) and are excluded from this graph.”

mean = $8.87 \times 10^{-1} \text{ hr}^{-1}$; SD = $5.07 \times 10^{-1} \text{ hr}^{-1}$) were more variable than those derived from the gross photoreduction experiments (Table 2). Many studies have focussed on the *in situ* net photoreduction of mercury in aquatic systems; for example, Poulain et al. (2004) determined that the DGM formation rate constant ranged from 0.76 – 1.4 h^{-1} in a wetland area, and 0.21 – 0.47 h^{-1} for a pelagic area. Another study by Amyot et al. (1994) determined the mean net mercury reduction rate constant to be 0.10 h^{-1} . A review by Vost et al. (2011) found that net mercury reduction rate constants ranged between 0.1 and 2.2 h^{-1} for freshwater samples, which are within the range of net mercury photoreduction constants measured in this study

Lake	0.2 um Filtered Samples						
	Gross Photoreduction K (h ⁻¹)	Standard Error	Hg(II)RED (pg L ⁻¹)	Net Photoreduction K (h ⁻¹)	Standard Error	Gross Photooxidation K (h ⁻¹)	Standard Error
Big Dam East (2009)	2.55E-01	1.30E-03	270	7.30E-01	1.28E+01	4.75E-01	1.28E+01
Big Dam West (2008)	1.95E-01	8.70E-04	490	5.09E-01	2.41E-01	3.14E-01	2.41E-01
Big Dam West (2009)	1.27E-01	4.82E-04	730	4.55E-01	3.05E-01	3.29E-01	3.05E-01
North Cranberry (2009)	1.91E-01	1.18E-03	645	4.60E+00	3.31E+01	4.41E+00	3.31E+01
Grafton (2009)	1.40E-01	1.17E-03	900	1.04E+00	3.50E-01	8.97E-01	3.50E-01
Puzzle (2009)	2.43E-01	4.22E-04	460	1.53E+00	1.85E+00	1.29E+00	1.85E+00
Big Red (2009)	1.41E-01	8.47E-04	250	2.49E-01	1.31E-01	1.08E-01	1.31E-01
Mountain (2009)	2.66E-01	1.73E-03	270	8.38E-01	4.49E-01	5.72E-01	4.49E-01
Beaverskin (2010)	8.15E-01	5.40E-03	NA	1.81E-01	1.36E+00	-6.34E-01	1.36E+00
Peskowesk (2010)	3.07E-01	1.70E-03	NA	6.43E-01	2.57E-01	3.36E-01	2.57E-01
Pebbleloggitch (2010)	8.16E-02	4.00E-04	NA	4.36E-01	2.76E-01	3.54E-01	2.76E-01
Lake	Unfiltered Samples						
	Gross Photoreduction K (h ⁻¹)	Standard Error	Hg(II)RED (pg L ⁻¹)	Net Photoreduction K (h ⁻¹)	Standard Error	Gross Photooxidation K (h ⁻¹)	Standard Error
Big Dam East (2009)	1.71E-01	8.79E-04	830	1.04E+00	8.60E-01	8.65E-01	8.60E-01
Big Dam West (2008)	1.27E-01	7.92E-04	895	5.03E-01	2.72E-01	3.76E-01	2.72E-01
Big Dam West (2009)	1.01E-01	1.17E-03	825	6.22E-01	2.85E-01	5.21E-01	2.85E-01
North Cranberry (2009)	1.68E-01	3.14E-04	1005	8.12E-01	1.34E+00	6.44E-01	1.34E+00
Grafton (2009)	1.67E-01	7.29E-04	1050	5.38E-01	5.00E-01	3.72E-01	5.00E-01
Puzzle (2009)	1.85E-01	5.95E-04	1035	1.29E+00	1.09E+00	1.10E+00	1.09E+00
Big Red (2009)	1.89E-01	1.33E-03	280	3.18E-01	1.06E-01	1.29E-01	1.06E-01
Mountain (2009)	2.29E-01	7.60E-04	485	1.99E+00	2.27E+00	1.76E+00	2.27E+00
Beaverskin (2010)	3.39E-01	2.35E-02	NA	1.05E+00	6.85E-01	7.11E-01	6.85E-01
Peskowesk (2010)	1.64E-01	8.00E-04	NA	7.62E-01	2.89E-01	5.98E-01	2.89E-01
Pebbleloggitch (2010)	1.12E-01	4.00E-04	NA	5.31E-01	5.14E-01	4.19E-01	5.14E-01

for both filtered and unfiltered water samples. The unfiltered water samples show slower rates possibly due to slow release of reducible mercury from solid particles in unfiltered samples similar to what has been proposed between soil particles and soil solution (Pannu et al., 2014)."

Section 2.4

"Gross photooxidation rate constants were not significantly related to DOC in filtered ($R^2=0.20$, $p > 0.10$) or unfiltered samples ($R^2 = 0.0007$; $p > 0.10$). There was no significant linear relationship between DOC and the gross photoreduction rate constants in either filtered ($R^2 = 0.43$; $p = 0.08$) or unfiltered ($R^2 = 0.02$; $p = 0.71$) samples (Fig. 3B). While not statistically significant, the trend suggests that in filtered water samples, as DOC increases, the rate at which photoreducible mercury is converted to Hg(0) may slow; this slowing of mercury photoreduction with increasing DOC in filtered water agrees with findings of a study done by Garcia et al. (2005a), who examined DOC fluorescence and DGM, finding that DGM was negatively correlated with DOC. However more data is required to determine if this trend is significant in larger numbers of samples. In the unfiltered samples it is possible that particulate-bound Hg(II) and Hg(0) that are variable with respect to biological transformations and photoreactivity interfere with this relationship and so a much weaker interaction is observed.

In contrast to the lack of significant relationships between DOC and photoreduction rate constants, there is a significant negative linear relationship ($R^2=0.53$; $p=0.04$) between DOC and Hg(II)_{RED} observed in unfiltered samples, and no relationship ($R^2=0.04$; $p=0.64$) observed in filtered samples (Fig. 4C)."

Should be revised to

"Gross photooxidation rate constants were not significantly related to DOC in filtered ($R^2=0.16$, $p > 0.10$) or unfiltered samples ($R^2 = 0.36$; $p > 0.10$). There was a weak linear relationship between DOC and the gross photoreduction rate constants in both filtered ($R^2 = 0.42$; $p = 0.08$) and unfiltered ($R^2 = 0.45$; $p = 0.07$) samples (Fig. 3B). While not statistically significant, the trend suggests that in filtered and unfiltered water samples, as DOC increases, the rate at which photoreducible mercury is converted to Hg(0) may slow; this slowing of mercury photoreduction with increasing DOC water agrees with findings of a study done by Garcia et al. (2005a), who examined DOC fluorescence and DGM, finding that DGM was negatively correlated with DOC. However more data is required to determine if this trend is significant in larger numbers of samples. In the unfiltered samples it is also possible that particulate-bound Hg(II) and Hg(0) that are variable with respect to biological transformations and photoreactivity interfere with this relationship.

There is a weak negative linear relationship ($R^2=0.45$; $p=0.06$) between DOC and Hg(II)_{RED} observed in unfiltered samples, and no relationship ($R^2=0.03$; $p>0.10$) observed in filtered samples (Fig. 4C)."

Table 1 and Figure 3 should be revised to the following.

Table 1: Rate constants (k; hr⁻¹) derived for gross photoreduction, net photoreduction, and gross photooxidation of mercury with standard error from curve fitting technique and total reducible (Hg(II)_{RED}; pg/L). Note that lakes sampled in August 2010 are highlighted in grey, lakes sampled in May of 2008 and 2009 are not highlighted.

The authors would like to apologise for any inconvenience caused.