

## Correction to “Manganese Catalysts with Bulky Bipyridine Ligands for the Electrocatalytic Reduction of Carbon Dioxide: Eliminating Dimerization and Altering Catalysis”

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Page 5466. In Table 1, the turnover frequency (TOF) values for Mn(bpy-<sup>t</sup>Bu)(CO)<sub>3</sub>Br (last column) were reported incorrectly from ref 15. The complete corrected table is shown below.

**Table 1. Comparison of Peak  $i_{\text{cat}}/i_{\text{p}}$  and TOF Values for Both [Mn(mesbpy)(CO)<sub>3</sub>(MeCN)](OTf) (2) and Mn(bpy-<sup>t</sup>Bu)(CO)<sub>3</sub>Br in MeCN (1 mM each catalyst)<sup>a</sup>**

Brønsted acid	[Mn(mesbpy)(CO) <sub>3</sub> (MeCN)](OTf) (2)			Mn(bpy- <sup>t</sup> Bu)(CO) <sub>3</sub> Br		
	[acid] <sup>b</sup> (M)	$i_{\text{cat}}/i_{\text{p}}^c$	TOF (s <sup>-1</sup> )	[acid] <sup>d</sup> (M)	$i_{\text{cat}}/i_{\text{p}}^e$	TOF (s <sup>-1</sup> )
H <sub>2</sub> O	3.5	20	700	3.1	25	120
MeOH	3.2	30	2000	5.8	26	130
TFE	1.4	50	5000	1.4	42	340

<sup>a</sup>Solutions are saturated with (approximately 0.19–0.28 M) and under an atmosphere of CO<sub>2</sub> with added weak Brønsted acids. Data are taken from voltammograms at a scan rate of 100 mV/s. [CO<sub>2</sub>] is approximately 0.28 M in dry MeCN, 0.26 M in 3.5 M H<sub>2</sub>O, 0.27 M in 3.2 M MeOH, and 0.27 M in 1.4 M TFE.<sup>22</sup> <sup>b</sup>[Acid] at highest  $i_{\text{cat}}/i_{\text{p}}$  for 2. <sup>c</sup> $i_{\text{cat}}/i_{\text{p}}$  values are calculated at equal [acid]. <sup>d</sup>[Acid] at highest  $i_{\text{cat}}/i_{\text{p}}$  for Mn(bpy-<sup>t</sup>Bu)(CO)<sub>3</sub>Br. <sup>e</sup>Values taken from ref 15 [Smieja, J. M.; Sampson, M. D.; Grice, K. A.; Benson, E. E.; Froehlich, J. D.; Kubiak, C. P. *Inorg. Chem.* **2013**, *52*, 2484–2491].

This correction to Table 1 changes some of the discussion on the comparison between complex 2 and Mn(bpy-<sup>t</sup>Bu)(CO)<sub>3</sub>Br. These changes are described below.

Page 5465. The statement, “Catalyst 2 is more active than the most active Mn bpy catalyst previously reported, Mn(bpy-<sup>t</sup>Bu)(CO)<sub>3</sub>Br, under MeOH and TFE. However, 2 is slightly less active than Mn(bpy-<sup>t</sup>Bu)(CO)<sub>3</sub>Br under H<sub>2</sub>O. Particularly, under identical conditions with 1.4 M TFE, catalyst 2 is ~1.2 times more active than Mn(bpy-<sup>t</sup>Bu)(CO)<sub>3</sub>Br.” should be changed as follows: “Catalyst 2 is more active than the most active Mn bpy catalyst previously reported, Mn(bpy-<sup>t</sup>Bu)(CO)<sub>3</sub>Br,<sup>15</sup> under all weak Brønsted acids studied. Under similar concentrations of TFE, catalyst 2 is over 10 times more active than Mn(bpy-<sup>t</sup>Bu)(CO)<sub>3</sub>Br.”