

## Correction to "Simple, Chemoselective, Catalytic Olefin Isomerization"

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Page 16788 and Supporting Information page SI-4. Entry 13 was inadvertently omitted from Table 1 and S.I. Table 1. This entry suggests mechanistic overlap between our alkene

isomerization and hydrogen atom transfer radical polymerization, more so than entry 12 alone. The complete S.I. Table 1 is shown below.

S.I. Table 1. Distribution of products from common Mukaiyama hydrofunctionalization catalysts

entry	conditions <sup>a</sup>	%1 <sup>b</sup>	%2	%3
1	1 mol% Co(Sal <sup>(Bu,fBu)</sup> Cl, 2 mol% PhSiH <sub>3</sub> , PhH, 22 °C, 3 h	4	96	0
2	10 mol% Mn(dpm) <sub>3</sub> , 2 equiv. PhSiH <sub>3</sub> , i-PrOH, 22 °C, 3 h	39 <sup>c</sup>	0	16
3	2 equiv. Fe <sub>2</sub> (oxalate) <sub>3</sub> , 6.4 equiv. NaBH <sub>4</sub> , ACN/H <sub>2</sub> O/i-PrOH (1:1:0.2), 0 °C, 1 h, air atm.	34	1	1
4	50 mol% Fe(acac) <sub>3</sub> , 1 equiv. PhSiH <sub>3</sub> , EtOH, 60 °C, 1 h	28 <sup>c</sup>	0	57
5	50 mol% Fe(acac) <sub>3</sub> , 1 equiv. PhSiH <sub>3</sub> , PhH, 60 °C, 1 h	65 <sup>c</sup>	0	0
6	50 mol% Co(acac) <sub>3</sub> , 1 equiv. PhSiH <sub>3</sub> , PhH, 60°C, 1 h	99	0	0
7	5 mol% Salcomine-Cl, 50 mol% PhSiH <sub>3</sub> , PhH, 60 °C, 3 h	65	20	6
8	1 mol%, Co(Sal <sup>rBu,rBu</sup> )Cl, 10 mol% PhSiH <sub>3</sub> , Me <sub>2</sub> CO, 22 °C, 3 h.	5	94	<1
9	1 mol% Co(Sal <sup>fBu,fBu</sup> )Cl, 10 mol% PhSiH <sub>3</sub> , CH <sub>2</sub> Cl <sub>2</sub> , 22 °C, 3 h	17 <i>c</i>	63	2
10	5 mol% Co(Sal <sup>fBu,fBu</sup> )Cl, 2 equiv. PMHS, PhH, 22 °C, 24 h	<b>4</b> <sup>c</sup>	78	0
11	2 mol% Co(Sal <sup>fBu,fBu</sup> )Cl, 40 mol% TESH, PhH, 22 °C, 24 h	98	0	0
12	5 mol% Co(Sal <sup>fBu,fBu</sup> )Cl, 50 mol% AIBN, PhH, 80 °C, 2 h	86	11	0
13	5 mol% Co(Sal <sup>fBu,fBu</sup> ), 50 mol% AIBN, PhH, 80 °C, 2 h	70	30	0

 $^a$ under Ar unless otherwise noted;  $^b$ according to GC-FID;  $^c$ other unidentified product(s) were observed