Inorg. Chem. 2009, 48, 1258

Inorganic Chemistry Additions and Corrections

2008, Volume 47

Marta Viciano-Chumillas, Stefania Tanase, Ilpo Mutikainen, Urho Turpeinen, L. Jos de Jongh,* and Jan Reedijk*: Mononuclear Manganese(III) Complexes as Building Blocks for the Design of Trinuclear Manganese Clusters: Study of the Ligand Influence on the Magnetic Properties of the $[Mn_3(\mu_3-O)]^{7+}$ Core

Page 5926. The *J* and *g* values presented in the Table 4 of compounds $[Mn_3(\mu_3-O)(phpzMe)_3(MeOH)_3(MeCO_2)]$ (2) and $[Mn_3(\mu_3-O)(phpzH)_3(MeOH)_4(N_3)]$ (3) should be $J_1 = -7.2 \text{ cm}^{-1}$, $J_2 = +7 \text{ cm}^{-1}$, and g = 2.13 for compound 2 and $J_1 = -5.7 \text{ cm}^{-1}$, $J_2 = -3.9 \text{ cm}^{-1}$, and g = 2.17 for 3. These values correspond with the ones reported in the text.

Page 5929. One of the planes that defines a dihedral angle with a value of 40.11° is incorrect. The correct sentence should read as follows:

Complex **2** shows the most structurally distorted $[Mn_3^{III}(\mu_3 - O)]^{7+}$ core, with one of the Mn–O_{oxido}–Mn angles of 116.10° and a deviation of the oxido bridge from the Mn₃ plane of 0.172 Å. In view of the arguments presented for Group II, the smaller Mn–O_{oxido}–Mn angle, corresponding with Mn(2)–O(1)–Mn(3), together with a large dihedral angle ($\delta_{MnNNMN-MnOMn} = 40.11^{\circ}$) between the Mn–O–Mn plane and the Mn(2)–N_{pz}–N_{pz}–Mn(3) coordination plane, could be held responsible for the observed ferromagnetic interaction along this path.

Note: None of the conclusions are affected by these changes.

IC802339B

10.1021/ic802339b Published on Web 12/22/2008