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Dongwoon Jung, Bang-Hee Lee, Sung-Jin Kim, and Won Kang: Synthesis, Characterization, and Electronic Structure of a New Molybdenum Bronze SnMo₄O₆.

The Acknowledgment section for this article (*Chem. Mater.* **2001**, *13*, 1625–1629) should be corrected as follows:

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Kenji Okitsu, Akihiko Yue, Shuji Tanabe, and Hiroshige Matsumoto: Sonochemical Preparation and Catalytic Behavior of Highly Dispersed Palladium Nanoparticles on Alumina.

Footnote 20 in this article (*Chem. Mater.* **2000**, *12*, 3006–3011) should read as follows:

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⁽²⁰⁾ The reduction of Pd(II) with H atoms in the presence of alcohol is almost negligible because the rates of Pd(II) reduction were so much higher than that in pure water: the rate of reduction in the presence of 1-propanol was at least ca. 10-fold faster than that in pure water. Therefore, the reduction of Pd(II) in the presence of alcohol would mainly proceed by organic radicals (R_{ab}, R_{py}) generating in the sonolysis of an alcohol additive. According to the previous report, ¹⁵ it was suggested that the reduction by R_{py} would be a main reaction pathway in the presence of a hydrophobic compound. The fact that highly hydrophobic compounds can easily concentrate onto the interface of a bubble at high temperature was also reported by Grieser and co-workers^{14b} and Henglein and co-workers [Henglein, A.; Kormann, C. Int. J. Radiat. Biol. **1985**, 48, 251]. On the other hand, when methanol, a hydrophilic compound, was used as an organic additive, the reduction by ' R_{ab} would be a main reaction pathway because methanol molecules at low concentration could not sufficiently accumulate onto the interface of Pd(II) reduction in the presence of methanol (25 µM/min) almost corresponds to the combined rates of formation of H atoms and OH radicals, which were roughly estimated by the rate of formation of H₂O₂. The mechanism for the sonochemical reduction of Pd(II) was discussed in detail in ref 15.