

Comments

Reply to “Comment on Precursor Morphology Controlled Formation of Rutile VO₂ Nanorods and Their Self-assembled Structures”

In the paper “Precursor Morphology Controlled Formation of Rutile VO₂ Nanorods and Their Self-Assembled Structures” (*Chem. Mater.* **2002**, *14*, 5053), we reported the rational synthesis of one-dimensional VO₂ nanomaterials through the controlled transformation from their hydrate precursors. Our use of a somewhat confusing description of the crystal structure of the VO₂ produced was the subject of a recent comment by Malavasi.¹

As a number of VO₂ polymorphs exist it is important to describe precisely the structure of the product. Malavasi’s comment pointed out that our VO₂ product has the monoclinic structure, not which is of the rutile form but which is of course tetragonal. Largely, the problem in describing the structure of VO₂ arises from the fact that this material exists in both structure types in different temperature ranges. It is well-known that VO₂ is monoclinic (which is a distorted form of the tetragonal rutile structure) at room temperature and transforms to the tetragonal rutile form at ~ 60 °C. In our Figure 4 is shown the X-ray diffraction pattern of our VO₂ product which is indexed in accord with JCPDS 9-142. Malavasi points out that this file corresponds to the monoclinic form which is correct, although the figure is labeled “Rutile VO₂”. In order to avoid further misunderstanding we agree that our product should be described as monoclinic VO₂ or distorted rutile VO₂ rather than “rutile VO₂”. In Supporting Information we show a resistance-temperature curve for our VO₂ which demonstrates clearly that the material is semiconducting at low temperatures and a

reversible phase transition occurs at ~55 °C. This is further evidence that our as-prepared VO₂ is the monoclinic form. Finally, we wish to observe that there exists an error in the Supporting Information which accompanies Malavasi’s comment. It is stated that JCPDS 70-2298 is that for the tetragonal rutile form of VO₂, whereas it is that for thallium lead chloride instead.

Experimental Information. The as-prepared VO₂ hydrate was synthesized according the paper.² The sample was heated at 400 °C for 24 h in a flowing N₂ atmosphere for dehydration, and the color of the products changes from pink to black. Then the black powder was pressed into pellets for electrical resistance measurement. Resistance measurements were performed using the ac four-probe method with an ac resistance bridge system (Linear Research, Inc., LR-700P).

Supporting Information Available: The resistance–temperature curve of the monoclinic rutile VO₂ nanorods we synthesized (PDF). This material is available free of charge via the Internet at <http://pubs.acs.org>.

Zhou Gui,^{*,†} Rong Fan,[‡] Weiqin Mo,[‡] Xianhui Chen,^{‡,§}
Ling Yang,[†] Shuyuan Zhang,[§] Yuan Hu,[†]
Zhangzhou Wang,[†] and Weicheng Fan[†]

*State Key Lab of Fire Science, Department of Physics, and
Structural Research Laboratory, University of Science and
Technology of China, Hefei 230026, P.R. China*

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[†] State Key Lab of Fire Science.

[‡] Department of Physics.

[§] Structural Research Laboratory.

(1) Malavasi, L. *Chem. Mater.* **2006**, *18*, 2774.

(2) Gui, Z.; Fan, R.; Mo, W.; Chen, X.; Yang, L.; Zhang, S.; Hu, Y.; Wang, Z.; Fan, W. *Chem. Mater.* **2002**, *14*, 5053.