

Development of a nano-Al/CuO based energetic material on silicon substrate

Kaili Zhang, Carole Rossi, Gustavo Adolfo Ardila Rodriguez, Christophe

Tenailleau, Pierre Alphonse

▶ To cite this version:

Kaili Zhang, Carole Rossi, Gustavo Adolfo Ardila Rodriguez, Christophe Tenailleau, Pierre Alphonse. Development of a nano-Al/CuO based energetic material on silicon substrate. Applied Physics Letters, 2007, 91 (11), pp.3117. 10.1063/1.2785132 . hal-03592695

HAL Id: hal-03592695 https://hal.science/hal-03592695

Submitted on 1 Mar 2022

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Open Archive Toulouse Archive Ouverte (OATAO)

OATAO is an open access repository that collects the work of Toulouse researchers and makes it freely available over the web where possible.

This is an author-deposited version published in: <u>http://oatao.univ-toulouse.fr/</u> Eprints ID : 2459

> To link to this article : URL : http://dx.doi.org/10.1063/1.2785132

To cite this version : Zhang, Kaili and Rossi, Carole and Ardila Rodriguez, G.A. and Tenailleau, Christophe and Alphonse, Pierre (2007) *Development of a nano-Al/CuO based energetic material on silicon substrate.* Applied Physics Letters, vol. 91 (n° 11). p. 3117. ISSN 0003-6951

Any correspondence concerning this service should be sent to the repository administrator: staff-oatao@inp-toulouse.fr

Development of a nano-Al/CuO based energetic material on silicon substrate

Kaili Zhang,^{a),b)} Carole Rossi,^{a),c)} and G. A. Ardila Rodriguez *LAAS-CNRS*, *Université de Toulouse*, *31077 Toulouse*, *France*,

Christophe Tenailleau and Pierre Alphonse CIRIMAT, 31062 Toulouse, France

Nanoenergetic materials (nEMs) have improved performances compared to their bulk counterpart or microcounterpart. The authors propose an approach to synthesize an Al/CuO based nEM that has several advantages over previous investigations such as enhanced contact, reduced impurities and Al oxidation, tailored dimensions, and easier integration into microsystem. CuO nanowires are synthesized by thermally annealing Cu film deposited onto silicon. Nano-Al is integrated with the nanowires to realize an Al/CuO based nEM. The synthesized nEM is characterized by scanning electron microscopy, high resolution transmission electron microscopy, x-ray diffraction, differential thermal analysis, and differential scanning calorimetry. © 2007 American Institute of Physics. [DOI: 10.1063/1.2785132]

Nanoenergetic materials (nEMs) have shown improved performances in terms of energy release, ignition, and mechanical properties compared to their bulk or microcounterparts. As a class of nEMs, nanothermite composites, also termed as metastable intermolecular composites (MICs), have been an extensive research subject.^{1,2} In MIC, oxidizer and fuel are mixed at nanoscale, thus leading to the enhanced reactivity caused by the reduced mass transport limitations between the reactants. Different approaches have been employed to synthesize MIC such as physical mixing of nanopowders of fuel and oxidizer,^{3–5} sol-gel,⁶ aerogel,⁷ atomic layer deposition,⁸ deposition of multilayer foils of oxidizer and fuel,⁹ integration of porous silicon with oxidizers,¹⁰ embedding Fe_2O_3 nanowires inside an Al film,¹¹ and employment of molecular self-assembly.^{12,13} In this study, Al/CuO based MICs are realized by integrating nano-Al with CuO nanowires grown from Cu thin film deposited onto silicon substrate. The fabricated MICs are characterized by scanning electron microscopy (SEM), high resolution transmission electron microscopy (HRTEM), x-ray diffraction (XRD), differential thermal analysis (DTA), and differential scanning calorimetry (DSC).

A 30 nm thick Ti film is deposited onto silicon followed by a 50 nm thick Cu film deposition by thermal evaporation. A 1 μ m thick Cu film is then deposited by electroplating. The substrate is cut into small chips that are cleaned in a HCl solution. The chips are heated in a horizontal tube furnace at 450 °C for 5 h under static air and then cooled down naturally to 25 °C. During the heating, CuO nanowires grow from the Cu film. Al is deposited by thermal evaporation onto the substrate with CuO nanowires. The deposited thickness (on average across the sample) of Al is set to be 1.12 μ m in the thermal evaporator. The Al is deposited under a vacuum level of 5×10⁻⁶ mbar and the substrate temperature is about 30 °C during the deposition. The Al/CuO based MICs on silicon are directly characterized by SEM and

^{c)}Electronic mail: rossi@laas.fr

XRD. The heat of reaction of the MIC is determined by DTA and DSC. DTA is performed from 20 to 1000 °C with a heating rate of 15 °C/min under a 99.995% Ar flow. A mass of 16.67 mg of the MIC is peeled from the substrate. DSC is carried out from 20 to 700 °C at a heating rate of 5 °C/min under a 99.999% N₂ flow with a sample mass of 2.80 mg.

Figure 1(a) shows a 30° tilted view SEM image of the annealed Cu film before Al deposition. CuO nanowires grow from the Cu film after annealing. Figure 1(b) is a SEM image of the nanowires after Al deposition. Nano-Al is uniformly integrated around the CuO nanowires, thus enhancing the interfacial contact and the reactivity. The average diameter of the nanowires is about 40–80 nm. After Al deposition, the average diameter becomes around 150–250 nm. Figures 2(a) and 2(b) are cross-section view SEM images of the CuO nanowires before and after Al deposition. Al is both integrated with the CuO nanowires to form a core-shell nanostructure and also deposited onto the film under the nanowires to form a layered structure.

The compositions of the MIC are identified by XRD, as shown in Fig. 3. Al, CuO, and Cu₂O diffraction lines can be seen from the XRD pattern. When Cu film is annealed in static air, Cu is first converted into Cu₂O. Then, CuO is formed slowly through a second step oxidation of Cu₂O. The inset picture in Fig. 3 is a HRTEM image of an individual nanowire. Each side of the nanowire is a single crystal with a distinct fringe space pattern. The interplanar spacing for one side is 2.46 Å and 1.99 Å for the other side; these inter-



FIG. 1. 30° tilted view SEM images of the annealed Cu film (a) before Al deposition and (b) after Al deposition.

^{a)}Authors to whom correspondence should be addressed.

^{b)}Electronic mail: kaili_zhang@hotmail.com



FIG. 2. Cross-section view SEM images of the annealed Cu film (a) before Al deposition and (b) after Al deposition.

planar spacings are similar to those for $\{111\}$ and $\{111\}$ planes in monoclinic CuO.¹⁴ There are no peaks for Al₂O₃ and Cu, indicating that no significant reaction between Al and CuO/Cu₂O occurs during the Al deposition due to the high vacuum level and low substrate temperature.

The exothermic reaction of the MIC is characterized with DTA, as shown in Fig. 4. There are two major exotherms. The first exotherm is observed with an onset temperature of about 500 °C, which means that the nanocomposite seems to react prior to the melting of Al. This suggests that the first exotherm is caused by the thermite reaction between the CuO nanowires and nano-Al. The second exotherm is found with an onset temperature of around 720 °C. After melting, the remaining Al reacts with the Cu₂O (and also perhaps some CuO) film under the CuO nanowires.

To quantitatively determine the heat of reaction, the MICs are characterized with DSC, as shown in Fig. 5. The first exothermic peak with an onset temperature of about 500 °C, which corresponds to the first peak in DTA curve in Fig. 4, is caused by the reaction between the CuO nanowires and nano-Al. The second endothermic peak (660 °C) is caused by the melting of the remaining Al. Integration of the first exothermic peak in DSC gives a heat of reaction equal to $1085 \pm 160 \text{ J/g}$. Assuming that the DTA signal is proportional to the heat of reactions, we use the area of the first peak, calculated from DSC analysis, to scale the DTA curve. Though it is not accurate because the relationship between temperature and heat is not exactly linear, it allows us to roughly estimate the total heat of reaction of the MIC to be about 2950 J/g. This value is between the theoretical heat of



FIG. 3. (Color online) XRD scan of the Al/CuO based MIC before reaction and HRTEM image of an individual nanowire.



FIG. 4. DTA plot of the Al/CuO based MICs.

reaction of CuO with Al (4067 J/g) and Cu₂O with Al (2434 J/g).¹⁵ The performance of the MIC can be improved by adjusting the dimensions of the CuO nanowires¹⁶ and by tuning the Al deposition to reach a stoichiometric reaction.

Figure 6 shows the XRD pattern of the MIC after DTA testing. The final reaction products are identified to be Al_2O_3 , $CuAlO_2$, and CuO. The Al peaks seen in Fig. 3 before reaction is no longer present. $CuAlO_2$ is formed from the reactions of Al_2O_3 with CuO/Cu_2O .^{17,18} However, XRD pattern does not show the Cu lines that should be there after CuO/Cu_2O reduction by Al. The Cu produced from $Al/CuO/Cu_2O$ thermite reactions is most likely oxidized by O_2 contained in Ar flow and/or generated from the reaction of Al_2O_3 with CuO.¹⁸

This approach has some advantages compared to previous investigations. First, the MICs are realized by integrating nano-Al with CuO nanowires, thus enhancing the interfacial contact and reactivity. Second, the presence of impurities and Al oxidation is smaller because the MICs are fabricated in a



FIG. 5. DSC plot of the heat flow for the Al/CuO based MICs.



FIG. 6. XRD scan of the Al/CuO based MIC after DTA testing.

high vacuum and with a low substrate temperature. Third, since the dimensions of CuO nanowires can be tailored, ¹⁶ the oxidizer/fuel dimensions can be controlled at the nanoscale. Fourth, the process uses standard microfabrication technologies and therefore it is suitable for mass production. The MICs are deposited onto silicon, a basic material for microelectronics and microsystem. Consequently, it is straightforward to integrate the MIC into microsystem to result in nanostructures, for example, high performance nanoigniters.^{2,19}

In conclusion, nano-Al/CuO based MICs have been realized by integrating nano-Al with CuO nanowires grown from Cu film deposited onto silicon. The Cu film is converted into bicrystal CuO nanowires and Cu₂O (and also perhaps some CuO) film after annealing. After Al deposition, nano-Al is integrated with CuO nanowires to form a coreshell nanostructure and also deposited onto the film under the nanowires to form a layered structure. DTA and DSC results suggest that nano-Al reacts with CuO nanowires with an onset temperature of about 500 °C prior to the melting of Al. After melting, the remaining Al reacts with the Cu_2O (and also perhaps some CuO) film beneath the CuO nanowires. The total heat of reaction is estimated to be about 2950 J/g. The MICs are realized onto silicon. Therefore, this will probably open the door to the integration of the material into silicon based microsystem, thus leading to functional nanodevices.

- ¹A. W. Miziolek, AMPTIAC 6, 43 (2002).
- ²C. Rossi, K. Zhang, D. Estève, P. Alphonse, J. Y. C. Ching, P. Tailhades, and C. Vahlas, J. Microelectromech. Syst. 16, 919 (2007).
- ³J. J. Granier and M. L. Pantoya, Combust. Flame 138, 373 (2004).
- ⁴B. S. Bockmon, M. L. Pantoya, S. F. Son, B. W. Asay, and J. T. Mang, J. Appl. Phys. **98**, 064903 (2005).
- ⁵A. Prakash, A. V. McCormick, and M. R. Zachariah, Adv. Mater. (Weinheim, Ger.) **17**, 900 (2005).
- ⁶T. M. Tillotson, A. E. Gash, R. L. Simpson, L. W. Hrubesh, J. H. Satcher, Jr., and J. F. Poco, J. Non-Cryst. Solids **285**, 338 (2001).
- ⁷A. Prakash, A. V. McCormick, and M. R. Zachariah, Nano Lett. **5**, 1357 (2005).
- ⁸J. D. Ferguson, K. J. Buechler, A. W. Weimer, and S. M. George, Powder Technol. **156**, 154 (2005).
- ⁹K. J. Blobaum, M. E. Reiss, J. M. P. Lawrence, and T. P. Weihs, J. Appl. Phys. **94**, 2915 (2003).
- ¹⁰A. Hofmann, H. Laucht, D. Kovalev, V. Y. Timoshenko, J. Diener, N. Kunzner, and E. Gross, U.S. Patent No. 6,984,274 (10 January 2006).
- ¹¹L. Menon, S. Patibandla, K. Bhargava Ram, S. I. Shkuratov, D. Aurongzeb, M. Holtz, J. Berg, J. Yun, and H. Temkin, Appl. Phys. Lett. **84**, 4737 (2004).
- ¹²S. H. Kim and M. R. Zachariah, Adv. Mater. (Weinheim, Ger.) 16, 1821 (2004).
- ¹³S. Subramaniam, S. Hasan, S. Bhattacharya, Y. Gao, S. Apperson, M. Hossain, R. V. Shende, S. Gangopadhyay, P. Redner, D. Kapoor, and S. Nicolich, *MRS Symposia Proceedings No. 896* (Materials Research Society, Pittsburgh 2006), p. 0896-H01-05.1.
- ¹⁴Joint Committee on Powder Diffraction Standards File No. 45–0937, 1991.
- ¹⁵O. B. Kubaschewski, C. B. Alcock, and P. J. Spencer, Materials Thermochemistry, 6th ed. (Pergammon, Oxford, 1993), pp. 259, 274, and 275.
- ¹⁶K. Zhang, C. Rossi, C. Tenailleau, P. Alphonse, and J. Y. C. Ching, Nanotechnology 18, 275607 (2007).
- ¹⁷J. H. Shy and B. H. Tseng, J. Phys. Chem. Solids **66**, 2123 (2005).
- ¹⁸M. N. Spallart, S. P. Pai, and R. Pinto, "PLD Growth of CuAlO₂," Thin Solid Films (in press).
- ¹⁹T. W. Barbee, R. L. Simpson, A. E. Gash, and J. H. Satcher, U.S. Patent No. WO 2005 016850 A2 (24 February 2005).