An Oxidation-Resistant Semiconductor

Stabilization of $\beta$-SiB$_3$ from Liquid Ga: A Boron-Rich Binary Semiconductor Resistant to High-Temperature Air Oxidation**

James R. Salvador, Daniel Bilc, S. D. Mahanti, and Mercouri G. Kanatzidis*

Liquid metals can be powerful solvents for the exploration of novel refractory solid-state materials ranging from intermetallic compounds to silicides, carbides and borides. Recently we have discovered, from liquid Al and Ga, a large variety of ternary and quaternary phases of the type RE/TM/Si or Ge, RE/TM/Al or Ga, RE/TM/Al/Si or Ge, and RE/TM/Ga/Ge or Si (RE = rare earth La–Lu, and TM = 1st, 2nd, or 3rd row transition metal).[1–2] Many of these phases cannot be formed by conventional high temperature solid-state synthetic techniques such as direct combination reactions with arc melting or radio-frequency (rf) induction furnace heating.[3] We have extended the flux methodology to include RE/TM/B/Si (borosilicides) in an attempt to produce lighter analogues of the RE/TM/Al/Si compounds, and we chose liquid gallium as the reaction medium because both Si and B are soluble in it and do not form binaries.[4] Also from previous work we learned that RE/TM/Ga/Si phases do not form readily, and so competing phases of this type are not anticipated.[1] Boron-containing solids have acquired renewed interest because quaternary borocarbides are superconducting,[5] and MgB$_2$ is a high temperature nonoxidic superconductor.[6]

Borosilicides are an unfamiliar class of compounds. Specifically boron-rich borosilicides are rare with only a handful of examples are known: CrSi$_3$(B$_8$)Se$_{12}$ (B$_8$Se$_3$)$_{1.33}$, REB$_2$Si$_3$ (RE = Y, Tb), Sc$_{0.83}$–B$_{1.00}$–C$_{0.17}$–Si$_{0.83}$–C$_{0.17}$ (SB$_3$, SiB$_3$, and SiB) to name a few. We have recently shown that liquid gallium can provide an excellent route to complex quaternary silicon borides such as Tb$_2$Si$_3$C$_2$(B$_{12}$)$_3$ that cannot be formed by using high-temperature techniques such as arc melting.[7] Herein, we present an illustration of the ability of liquid Ga to stabilize boride phases inaccessible by conventional synthetic routes. Namely, we describe a novel B-rich, refractory, semiconducting binary phase $\beta$-SiB$_3$.

$\beta$-SiB$_3$ is a surprising discovery because a phase with a similar formula, Si$_{1.4}$B$_{12}$ with a compositional spread of $x = 0–0.2$,[8] that has a rhombohedral structure has been known for decades.[11] We will henceforth refer to this phase as $\alpha$-SiB$_3$. Instead $\beta$-SiB$_3$ is orthorhombic $Imma$, representing a new structure type.[12] Despite the similarity in stoichiometry and the presence of B$_{12}$ icosahedra the structural features and electronic properties of the $\alpha$ and $\beta$ phases are very different. Rhombohedral $\alpha$-SiB$_3$ forms from direct combination of the elements with prolonged isothermal heating at 1225 °C or by arc melting, in contrast $\beta$-SiB$_3$ was produced from a lower temperature reaction (1000–850 °C) in Ga flux. Additionally, in $\alpha$-SiB$_3$, some Si atoms are randomly distributed in the B$_{12}$ cages resulting in very distorted icosahedra.[9] In contrast, Si atoms in $\beta$-SiB$_3$ are rigorously excluded from the cages, thus resulting in more regular icosahedra. To the best of our knowledge this is the first Si–B binary phase that is fully crystallographically ordered without any Si/B mixed sites or disorder. This exclusion of Si from the cages has a profound influence on the thermal stability of the orthorhombic form.

$\beta$-SiB$_3$ was originally found as a minor product in reactions aimed to produce Er$_{1.8}$Si$_8$C$_2$(B$_{12}$)$_3$ from the Ga flux. The new phase forms as distorted rectangular plates that are translucent and amber in color. Energy dispersive spectroscopy (EDS) on several crystals of the title compound detected only Si and B.[13] The presence of C in $\beta$-SiB$_3$ was detected by combustion elemental analysis which showed the carbon concentration to be $\sim 0.4\%$ by weight.[14] Similar combustion analysis on $\alpha$-SiB$_3$ (a phase known to contain no C) showed a C concentration of 0.16% by weight. We originally thought that there was a small amount of C present...
in β-SiB\textsubscript{3}, and that it was statistically distributed over the B\textsubscript{12} cages resulting in the stoichiometry Si\textsubscript{x}B\textsubscript{12}\textsubscript{1−x}C\textsubscript{y} (x \approx 0.4). The yield of β-SiB\textsubscript{3} was observed to increase when carbon was included in the preparation (from 40% to 60%). Although it is possible that a small amount of carbon is accommodated in the structure, we believe that it is not needed to prepare the compound. The issue of adventitious carbon inclusion is discussed below.

Importantly, no traces of the known black rhombohedral α-SiB\textsubscript{3} phase were found by powder X-ray diffraction, nor visual inspection in the flux reactions. As the crystal habit and color of the two species are quite different, they are easily distinguished by looking. An inspection of other reactions of the type TM/B/C/Si (TM = Fe, Co, Ni, Cu, Zn) in a Ga flux showed large amounts of β-SiB\textsubscript{3}, particularly when TM = Cu.

The formation of β-SiB\textsubscript{3} is exclusive to flux synthesis. Several attempts were made to produce the phase by direct combination of the elements with arc-melt alloying, rf induction heating and prolonged heating at 1000°C for two weeks. The induction furnace and the two-week 1000°C isothermal synthesis both yielded starting material, presumably because the temperatures reached by these methods were insufficient to bring about a reaction. Arc-melt alloying Si and B in a 1:3 ratio yielded the α-SiB\textsubscript{3} structure. EDS analysis was also performed on crystals produced from this reaction and found on Si and B present in similar ratios to the title compound. These results may explain why β-SiB\textsubscript{3} was not discovered previously, as flux routes to borosilicides are a new synthetic strategy.

The structure of β-SiB\textsubscript{3} has two building blocks: infinite Si\textsubscript{4} chains and B\textsubscript{12} cages (Figure 1a and 1b respectively). The B\textsubscript{12} cages are anionic and are well rationalized by the Zintl concept. [16] In contrast, the Si\textsubscript{4} chains of BaGe\textsubscript{2} are not anionic and are not regarded as Zintl species.

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The four crystallographically distinct boron atoms are arranged in the B\textsubscript{12} cage that form a sheet structure extending over the ac plane, which results in a regular four-sided ring of cages Figure 1c. The cages are bound through symmetry-generated B(4)–B(4) bonds at 1.717(2) Å. Intracage distances range from 1.934(2) Å for B(4)–B(4) to 1.756(2) Å for B(1)–B(3) with an average of 1.792 Å. The Si zig-zag chains insert between the B\textsubscript{12} icosahedral sheets forming B–Si bonds and a three-dimensional framework along the b axis (Figure 2). The B–Si bond distances are 2.027(1) Å for Si(1)–B(1), 1.973(2) Å between Si(2) and B(3), and 2.009(2) for Si(2)–B(2).

Figure 1. a) A segment of the infinite Si\textsubscript{4} ribbon as it extends down the a axis, a motif reminiscent of red phosphorous, thermal elipsoids are set at the 50% probability level. Selected bond angles Si(1)–Si(1)–Si(2) = 91.85(2)°, Si(1)–Si(2)–Si(1) = 121.53(2)°, Si(2)–Si(1)–Si(2) = 54.97(2)°, Si(2)–Si(2)–Si(1) = 62.51(3)°. b) B\textsubscript{12} icosahedral unit, thermal elipsoids are set at the 50% probability level. c) The arrangement of B\textsubscript{12} cages in the ac plane.

Figure 2. A view of the overall structure of β-SiB\textsubscript{3}, showing the way two structural moieties stack along the b axis. Si atoms: blue; B atoms: red.

β-SiB\textsubscript{3} is a semiconductor with an energy band gap of \approx 2.0 eV, (Figure 3a). [17] This wide gap is in sharp contrast to that of the α-SiB\textsubscript{3}, which is only \approx 0.2 eV, (Figure 3b), and points to the dramatically different electronic properties of...
When the experiment was repeated in air instead of oxygen, there was no evident ignition point and only a nominal relative weight gain of \( \approx 2\% \) was observed up to 1000°C. In contrast, \( \alpha \)-SiB\(_3\) shows an ignition point of 600°C in air, Figure 3c. We attribute the lack of stability in \( \alpha \)-SiB\(_3\) to the nature of the Si–B disorder within the B\(_4\) icosahedra, which causes the weakening of bonds. To test more realistically the thermal stability of the title compound, several single crystals were placed in open fused silica tubes and heated to 700°C and held there for 12 h. \( \beta \)-SiB\(_3\) remained intact at this temperature. Scanning electron micrographs of a \( \beta \)-SiB\(_3\) single crystal before and after heating can be seen in Figure 3d and 3e respectively. Only when the experiment was repeated at 1000°C was there evidence of thermal degradation.

The resistance to air oxidation at high temperature is impressive for a nonoxidic compound. As it also has a wide band gap of 2.0 eV, this new material might be useful as a refractory p-type semiconductor capable of operating in extreme environments. Currently the leading material for such applications is SiC, however, it suffers from the major drawback of having nearly 200 polytype structures, all of which have different electronic properties.\(^{[21]}\) The compound presented here has no other polytypes, and \( \beta \)-SiB\(_3\) can be made independently from \( \alpha \)-SiB\(_3\).

It is significant that the synthesis of \( \beta \)-SiB\(_3\) requires metallic flux conditions, which permit the total bypass of the rhombohedral compound \( \alpha \)-SiB\(_3\). The results reported herein warrant future investigations of the electrical properties of \( \beta \)-SiB\(_3\).

**Experimental Section**

Optimized flux synthesis of \( \beta \)-SiB\(_3\) was carried out by combining B (43 mg, 4 mmol; 99.995% submicron amorphous, Aldrich), Si (28 mg, 1 mmol; 99.999% – 325 mesh, Cerac), Cu (64 mg, 1 mmol; 99.9% 325 mesh, Cerac), and Ga (1.394 g, 20 mmol; 99.9% 2–5 mm pieces, Plasmaterials) into an Al\(_2\)O\(_3\) ceramic tube. The ceramic tube was placed in a fused silica tube that was flame-sealed under a vacuum of \(-10^{-2}\) mbar. The reactants were heated up to 1000°C over 12 h, then held at 1000°C for 96 h. The reaction system was then cooled to 150°C over 48 h. Isolation of the product was accomplished by a warm centrifugation, which removed the excess Ga. Further purification was carried out by adding the product to a solution of I\(_2\) (5m) in dimethylformamide, and subjecting the mixture to ultrasound for 30 min to remove remaining Ga. Small shiny amber colored plates were obtained approximately 0.5–1 mm at the widest point. Typical yields were 40%.

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Communications


[10] The phases SiB4 and SiB6 are listed separately in “Pearson’s Handbook of Crystallographic Data for Intermetallic Phases”, and are even listed as different structure types the former having the B6P structure type and the later the B 13C2. In fact SiB3 and SiB4 have the same structure, that of B6P, and the variation in stoichiometry is the result of a compositional spread ranging from SiB2.89 up to SiB4. This stoichiometric flexibility arises from the substitution of varying amounts of Si into the polar sites of the B6P cages.


[12] Crystal data for β-SiB3, M = 61.02 g mol⁻¹; orthorhombic, space group Inmma, a = 8.3915(5), b = 12.5680(8), c = 6.2134(4) Å, V = 655.29(2) Å³, Z = 16, pDcalc = 2.454 g cm⁻³, µ = 0.805 mm⁻¹, R1/ wR = 1.91%/5.06% for all the 558 unique reflections, 3685 total reflections (Rwp = 1.78%) and 43 parameters. X-ray diffraction data were collected at room temperature on a Bruker SMART CCD diffractometer, graphite-monochromatized MoKα radiation (0.71073 Å). The structure was solved by direct methods and refined with the SHELXL suite of programs. Further details on the crystal structure investigation may be obtained from the Fachinformationszentrum Karlsruhe, 76344 Eggenstein-Leopoldshafen, Germany (fax: (+49) 7247-808-666; e-mail: crys-data@fiz-karlsruhe.de), on quoting the depository number CSD-412621.


[14] EDS was performed on several crystals of the compound by using 8 kV accelerating voltage and 120 s acquisition time and indicated that boron and silicon were present in the amber plate crystals; the ratio of boron to silicon was approximately 2.4:1. Samples were mounted on silver paste to obtain a clear carbon content. Measurements were performed with a JEOL JSM-35C scanning electron microscope equipped with a Noran Vantage Energy Dispersive Spectroscopy (Si:Li) detector and Norvar window suitable for the standardless quantitation of light elements (Z ≥ 4).

[15] Combustion analysis was carried out by selecting single crystals of the title compound (sample size 2.272 mg) and α-SiB3 (sample size 3.730 mg) and rinsing them in aqua regia and “piranaha” solution (75 vol% conc. H2SO4/25 vol% 30% H2O2) to minimize surface carbon contamination. Elemental analysis was performed with a Perkin Elmer series II CHNS/ O analyzer 2400.


[17] Band gap measurements were made on a Shimadzu UV-3101PC UV/Vis–NIR solid-state spectrophotometer. Measurements were made in reflectance mode from a BaSO4 standard reference. The spectrum was converted to absorbance by using the Kubelka–Munk equation.

[18] Band-gap measurements for rhombohedral SiB4 were made with a Nicolet Magna IR 750 FTIR spectrophotometer in reflectance mode. The data was processed with the Kubelka-Munk equation.


[20] Band-structure calculations were performed by using the self-consistent full-potential linearized augmented plane-wave method (LAPW) within density functional theory (DFT) by using the generalized gradient approximation (GGA) for the exchange and correlation potential. Scalar relativistic corrections were added and spin-orbit interaction (SOI) was incorporated by using a second variational procedure. The calculations were performed with the WIEN97 program.