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TEM study of boron phosphide: Discovery of rhombohedral BP

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Microstructure of sphalerite (3C) boron phosphide, BP, produced by self-propagated high-temperature synthesis has been studied by high-resolution transmission electron microscopy. Along with numerous twins on the $\{111\}_{3C}$ plane, layers of wurtzite (2H) polymorphic modification and previously unknown for BP rhombohedral (9R) structure were found which indicates trimorphism of BP.

Keywords: boron phosphide, transmission electron microscopy, polymorphism, twinning.

Introduction

Boron phosphide BP is a hard (Vickers hardness $H_V \sim 30$ GPa [1]) refractory (melting temperature at ambient pressure is 2840 K [2]) and low-compressible (300-K bulk modulus is 174 GPa [3]) wide bandgap ($E_g = 2.1$ eV [4]) semiconductor with outstanding chemical and high-temperature stability that makes it a promising material for a wide range of applications [5]. Under ambient conditions, BP crystallizes in cubic sphalerite (*F*-43*m*) structure with boron and phosphorus tetrahedrally coordinated to each other. Another tetrahedral structure of boron phosphide, hexagonal wurtzite (*P*6₃*mc*) polymorph, has also been reported [6], however, in the literature there is no data on the structure and properties of this phase.

The main limitation for the use of boron phosphide is the lack of relatively simple and economical methods of its production, especially, of BP single crystals. The disadvantages of the existing methods are the use of toxic and aggressive reagents, complicated technical implementation and high time consumption. Recently, two new methods of boron phosphide synthesis have been

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developed i.e. self-propagated high-temperature synthesis [7] and mechanochemical synthesis [8] that are characterized by simplicity, high efficiency, low cost and good perspectives for large-scale production. Here we report the results of transmission electron microscopy studies of BP produced by self-propagated high-temperature synthesis.

Experimental

Microcrystalline powder of boron phosphide has been synthesized by self-propagating hightemperature reaction of boron phosphate and metallic magnesium: BPO₄ + 4Mg = BP + 4MgO using the method described earlier [7]. According to X-ray diffraction study (TEXT 3000 Inel, CuKa1 radiation) the sample is well-crystallized sphalerite BP with lattice parameter a = 0.45356(9) nm, which was close to the literature value 0.4538(2) nm [9]. The amount of B₁₂P₂ boron subphosphide impurity caused by a partial decomposition of as-forming BP due to a high local temperatures in the reaction mixture did not exceed 1 vol.%.

Microstructure of boron phosphide has been studied by high-resolution transmission electron microscopy (HRTEM) using JEM-2010 microscope with energy dispersive x-ray spectroscopy (EDS) attachment.

Results and discussion

According to TEM data (Fig. 1), the boron phosphide powder consists of flat polyhedral particles having dimensions from 50 to several hundred nanometers (Fig.1a); significant part of them contains twins. Fig.1b shows two particles containing twins. Traces of twinning planes are shown by arrows. Such particles with a sphalerite structure predominate in the sample. At the same time, other phases were detected in small quantities.

Fig. 2 presents TEM and HRTEM images of sphalerite BP particle containing two systems of twins. White rectangles depict stacking faults which change the layer stacking sequence from *ABCABC* for cubic (3C) sphalerite structure to *ABA* for hexagonal (2H) wurtzite structure. The corresponding fast Fourier transform (FFT) image is shown in the inset. Strong streakings parallel to (111) and (1-1-1) are caused by diffuse scattering from {111} type stacking faults. Similar structural transformations were previously observed in carbon (diamond \rightarrow lonsdaleite) [10] and in silicon (SiI \rightarrow SiIV) [11]. According to [12], the lattice parameters of wurtzite BP are *a* = 0.320 nm and *c* = 0.531 nm, which are noticeably different from the lattice parameters of its closest structural analog, wurtzite BN (*a* = 0.25505 nm; *c* = 0.4210 nm [13]).

Fig. 3 shows a BP particle with a complex structure. In the upper part of HRTEM image twins in sphalerite 3C structure along the (111) plane are observed. The trace of the twinning plane is indicated by a vertical line at the top of the figure. In the lower part on the right, twins along the

(11-1) plane are visible. Both twinning planes are marked in Figure. The red circles (Fig. 3b) indicate the reflexes related to the 3C structure located in the upper right part, while the yellow circles indicate the reflexes of the 3C structure located in the upper left part of the figure. The yellow color line indicates the line separating the different phases in the particle. The rhombohedral (9R) phase was formed on both sides of the (111) plane in the lower part of the image. On the right side the tripling occurred along the [11-1]_{3C} direction. On the left part – along the [111]_{3C} direction. The crystal lattice in the upper part corresponds to the sphalerite (3C) structure with lattice parameter a = 0.4538 nm, whereas the structure in the lower part is a rhombohedral (9R) lattice with trippled value of interplanar distance for (111) sphalerite planes i.e. $0.262 \times 3 = 0.786$ nm. The tripling is visible both on HRTEM and FFT imags. As follows from the comparison of FFT images of two neighboring fragments (Fig. 3b) the (11-1)_{3C} || (001)_{9R} and [1-10]_{3C} || [11-20]_{9R} orientation relation is observed. Red circles indicate reflexes from 3C crystalline lattice. According to our assessment, the lattice parameters of the rhombohedral phase are: a = 0.321 nm, c = 2.358 nm. Thus, Fig. 3 shows a twinned BP particle which also contains two fragments of rhombohedral phase. The rhombohedral phase in the right part is formed as a result of twinning.

Fig. 4 shows the schematic representation of different structures of the BP composition: a) *ABAB* packing (2H); b) *ABCABC* packing (3C) and c) *ABABCBCAC* packing (9R). Drawings containing models of crystal structures are made in a software package for visualization and analysis Ovito [14]. We could not find any information on the existence of rhombohedral BP in the literature, however, there is the experimental evidence for formation of rhombohedral polymorphic modification of zinc sulfide (ZnS) [15], the structural analogue of BP. This rhombohedral structure of ZnS was found to occur in the 600-1020°C temperature range as an intermediate phase between sphalerite (3C) and wurtzite (2H) structures. In our case, the appearance of rhombohedral BP can be associated with deformation of as-formed sphalerite lattice in the course of ultrafast self-propagated high-temperature reaction, which leads to a change of interplanar distances for {111}_{3C} planes and, consequently, results in the change of the lattice symmetry.

Fig. 5a shows a two-phase particle. The left part of the particle corresponds to $B_{12}P_2$ composition and contains polytypes, as evidenced by the elongated streaks passing through the spots in the corresponding FFT-image (Fig. 4b). The right part contains a twice-twinned fragment of BP composition (the corresponding FFT-image is shown in Fig. 4c). The mutual orientation of these two phases corresponds to the $(003)_{B_{12}P_2} \parallel (111)_{BP}$ orientation relation. The arrows denote the $\{111\}_{BP}$ planes. One of the planes separates $B_{12}P_2$ and BP crystals, while the other two are twinning planes in BP crystal. Such a junction of twinning planes is generally denoted as $\Sigma = 9$ [16]. In other words, $\Sigma = 9$ (second order twin) boundary is created when two $\Sigma = 3$ (first order twin) boundaries intersect. EDS spectra of $B_{12}P_2$ is shown in Fig. 4d.

Previously, in boron subphosphide $B_{12}P_2$ synthesized by the same self-propagating hightemperature method we have discovered two systems of twins i.e. conventional twins on the (0003)_h plane and nanotwins resulting from duplication of the rhombohedral unit cell of $B_{12}P_2$ along one of the basic vectors [17]. In the present study, neither these nor other twins were observed in the only few $B_{12}P_2$ particles found. The latter can be explained by the fact that $B_{12}P_2$ is a secondary phase formed due to the thermal decomposition of as-synthesized BP at points of the local overheating in the reaction front.

Conclusions

In boron phosphide produced by self-propagated high-temperature synthesis, along with traditional sphalerite (3C) polymorphic modification, fragments of wurtzite (2H) polymorph and previously unknown for BP rhombohedral (9R) structure were found which shed new light on polymorphism of boron phosphide.

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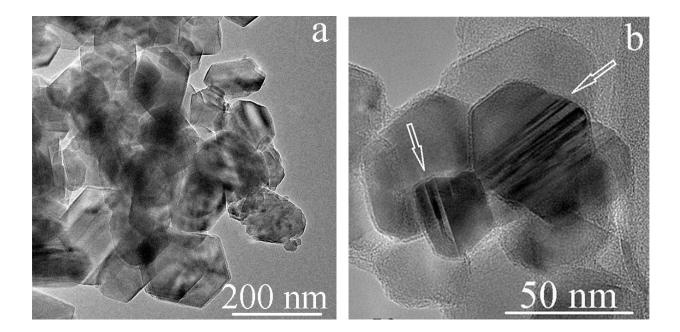


Fig. 1. A characteristic TEM image of boron phosphide particles(a); two particles contain twins (b). Traces of twinning planes are shown by arrows.

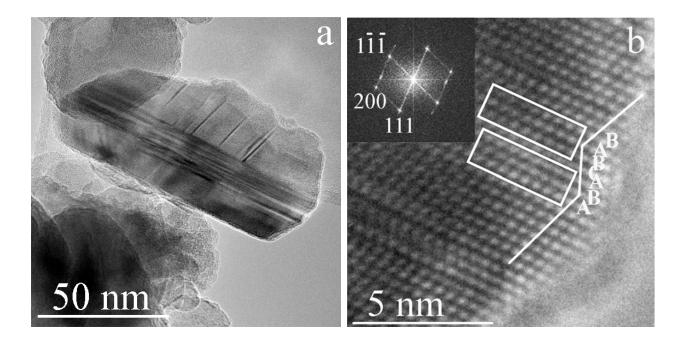


Fig. 2. Two systems of twins in BP structure: (a) overview of the particle and (b) high resolution image of a portion of the particle where white rectangles depict the fragments of wurtzite (2H) structure with BAB and ABA layer stacking sequence; the corresponding FFT image is shown in the inset.

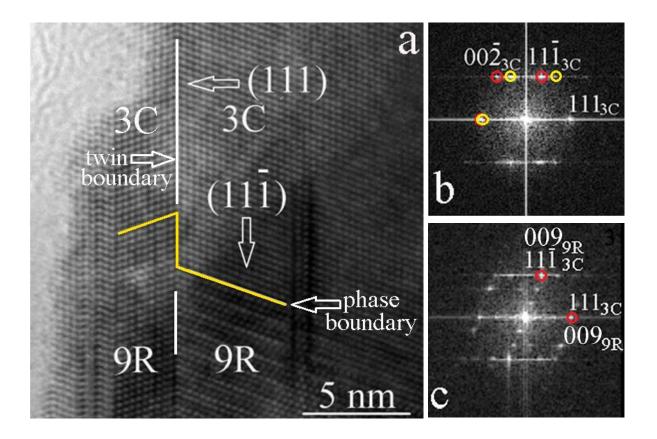


Fig. 3. (a) HRTEM image of BP particle which contains twinned sphalerite phase, as well as two fragments of rhombohedral (9R) polymorph: the upper part shows BP twins on the (111)_{3C} plane; the trace of the twinning plane is indicated by a vertical line at the top of the figure; b) the corresponding FFT-image. The red circles indicate the reflexes related to the 3C structure located in the upper right part, while the yellow circles indicate the reflexes of the 3C structure located in the upper left part of the figure. The yellow color indicates the line separating the different phases in the particle. At the lower part on both sides of the (111)_{3C} plane (vertical white dash at the bottom of the figure), two fragments of rhombohedral phase were formed. On the right side the tripling occurred relative to the (111-1)_{3C} plane; while on the left side the tripling occurred relative to the (111)_{3C} plane. (*c*) FFT-image of the right fragment of a particle containing 9R and 3C structures. Red circles indicate reflexes from 3C crystalline lattice.

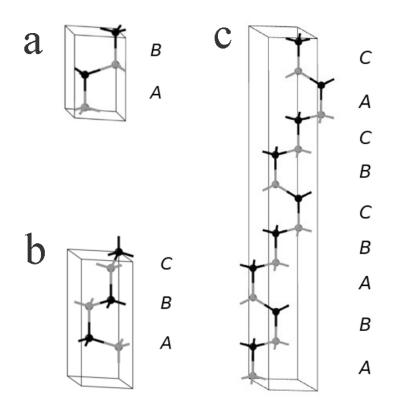


Fig. 4. Schematic representation of different structures of the BP composition: a) packing *ABAB* (2H); b) packing *ABCABC* (3C) and c) packing *ABABCBCAC* (9R).

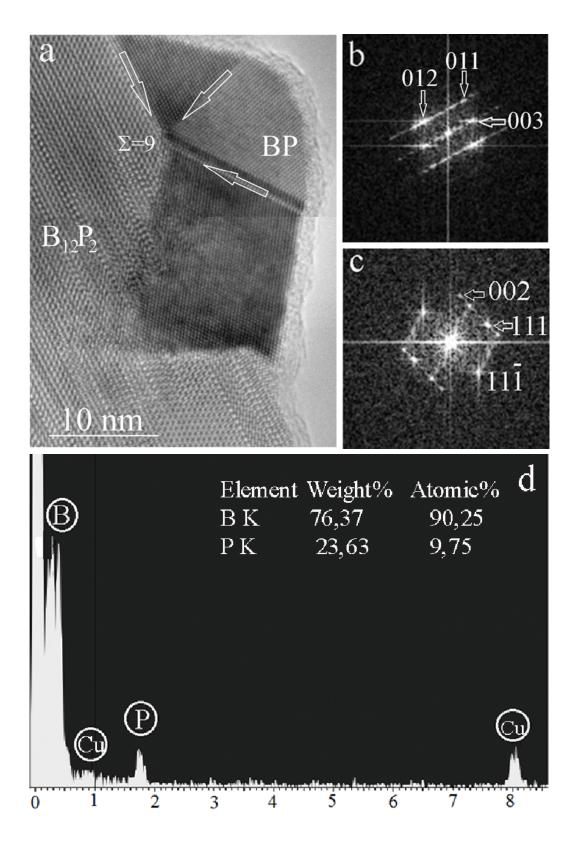


Fig. 5. (a) HRTEM image of two-phase particle: the left part is of B₁₂P₂ composition and contains polytypes; the right part is twice-twinned fragment of BP composition. The arrows indicate {111}_{BP} planes. (b) FFT image of the left part of (a); (c) FFT image of the right part of (a); d) EDS-spectra from B₁₂P₂.