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Chemical Analysis of Impurity Boron Atoms in Diamond using Soft X-Ray Emission Spectroscopy

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Abstract

To analyze the local structure and/or chemical states of boron atoms in boron-doped diamond, which can be synthesized by the microwave plasma-assisted chemical vapor deposition method (CVD-B-diamond) and the temperature gradient method at high pressure and high temperature (HPT-B-diamond), we measured the soft X-ray emission spectra in the *CK* and *BK* regions of B-diamonds using synchrotron radiation at the Advanced Light Source (ALS). X-ray spectral analyses using the fingerprint method and molecular orbital calculations confirm that boron atoms in CVD-B-diamond substitute for carbon atoms in the diamond lattice to form covalent B-C bonds, while boron atoms in HPT-B-diamond react with the impurity nitrogen atoms to form hexagonal boron nitride. This suggests that the high purity diamond without nitrogen impurities is necessary to synthesize *p*-type B-diamond semiconductors.

Introduction

Impurity-doped diamonds can be wide-gap semiconductors, which can be utilized as electron emitters and high-power amplifiers in electronic devices.¹ In particular, boron-doped diamond (denoted as B-diamond) has recently attracted much attention from technological application and materials science points of view. High-quality B-diamond has been synthesized by a microwave plasma-assisted chemical vapor deposition (CVD) method where the boron concentration controls its electrical properties.^{2,3} To clarify the relationship between the electronic structure and local structure of the boron atoms in B-diamond, the valence band and conduction band have been analyzed by soft X-ray emission spectroscopy (XES) and absorption spectroscopy (XAS) using synchrotron radiation (SR).⁴⁻⁶ These spectroscopic studies have elucidated that covalent B-C bond formation in a diamond matrix is important for acceptor formation in *p*-type B-diamond semiconductors. Additionally, superconducting characteristics have been discovered in heavily boron-doped diamond,⁷ and its electronic structure has been analyzed by SR photoemission spectroscopy⁸ and the X-ray scattering method.⁹ These analytical studies on B-diamond using SR have suggested that the electronic structure strongly depends on the local structure and/or chemical states of the dopant boron atoms in the diamond lattice. On the other hand, chemical analyses of boron compounds using XES including the conventional X-ray fluorescence spectrometry have been achieved by many researchers.¹⁰⁻¹⁹ This shows that XES is a powerful tool to investigate the chemical states of boron compounds.

As previously mentioned, the electronic structure of the high-quality B-diamond synthesized by the CVD method can be successfully analyzed by XES and XAS. However, B-diamond synthesized by a temperature gradient method at high pressure and high temperature (denoted by HPT)^{20,21} has not been analyzed by XES and XAS yet. The difference between the CVD and HPT methods is the impurity nitrogen in the diamond matrix; CVD-diamond has a negligible nitrogen impurity, while HPT-diamond generally has ppm- or sub-ppm-orders of nitrogen as an impurity.²² Therefore, it should be interesting to investigate the effects of impurity nitrogen on the local structure and/or chemical states of dopant boron.

In the present study, we analyzed dopant boron in the B-diamond synthesized by the HPT method using XES and compared it to CVD-B-diamond. Moreover, the local structure and/or

chemical states of dopant boron atoms in the B-diamonds are discussed using the spectral analysis of XES spectra and molecular orbital calculations.

Experiments

Two types of B-diamond samples were prepared. One was a boron-doped diamond-single-crystal film synthesized by the CVD method with methane, hydrogen, and diborane gases on single-crystal diamond substrates (denoted as CVD-B-diamond). The concentration of boron in CVD-B-diamond was evaluated by SIMS as 920 ppm. The other B-diamond sample was synthesized by the HPT method with a high-purity carbon source and boron source for the dopant (denoted as HPT-B-diamond). The concentration of boron in the HPT-B-diamond sample was adjusted to 300 ppm. Although the concentration of impurity nitrogen in the measured HPT-B-diamond sample was not unfortunately analyzed by SIMS, it can be estimated to be not less than several tens ppm. The High-purity CVD- and HPT-diamonds without boron doping were also synthesized as references. In addition, commercially available highly oriented pyrolytic graphite (HOPG), hexagonal boron nitride (*h*-BN), cubic boron nitride (*c*-BN), boron carbide (B_4C), and amorphous boron (*a*-B) were prepared as reference compounds.

Spectroscopic measurements of the soft X-ray emission in the *BK* and *CK* regions were performed with a grating X-ray spectrometer installed at beamline BL-8.0.1²³ of the Advanced Light Source (ALS). The emission spectra were obtained with excitation energies tuned to 230 eV for the *BK* region and 320 eV for the *CK* region in order to effectively excite the 1s-electrons and prevent multiple ionizations. The estimated resolving power ($E/\Delta E$) of the spectrometer was 1300 for the *BK* measurements using a 40- μ m entrance slit and a 1500-lines/mm spherical grating with a 5-m radius, and 680 for the *CK* measurements using a 600-lines/mm grating with a 10-m radius.

Results and Discussion

Figure 1 shows the *CK* X-ray emission spectra of the B-diamonds and reference compounds. All the diamond samples exhibited similar spectral profiles; a plateau in the region of 278 – 281 eV, a low-energy satellite at 272 eV, and a high-energy peak edge at 284 eV. This spectral profile

corresponds to previously published data.^{24,25} Therefore, it is confirmed that CVD-B-diamond and HPT-B-diamond essentially take a typical diamond crystal structure. Figure 2 shows the BK X-ray emission spectra of CVD-B- and HPT-B-diamonds and reference boron compounds. The CVD-B-diamond exhibited a main peak around 184 eV with a steep edge at 185 eV and lower-energy tailing, which apparently differs from the spectral profiles of HTP-B-diamond, *h*-BN, and *c*-BN. Although B₄C and *a*-B exhibited similar spectral profiles with a main peak and lower-energy tailing of CVD-B-diamond, neither exhibited the steep 185-eV edge. This demonstrates that the chemical states of the dopant B atoms in the CVD-B-diamond differ from those of HTP-B-diamond and the reference compounds. The HPT-B-diamond exhibited a main peak at 182 eV, a high-energy shoulder at 185 eV, a low-energy shoulder at 178 eV, and a lower-energy satellite at 170 eV. This spectral profile corresponds to that of *h*-BN. Although *c*-BN also exhibited a similar profile, the main peak is broader than that of HPT-B-diamond and *h*-BN. This indicates that dopant B atoms in the HPT-B-diamond mainly form *h*-BN in the diamond matrix and the concentration of impurity N atoms in the HPT-B-diamond will be not less than 300 ppm.

To theoretically clarify the measured BK X-ray spectral profiles of the B-diamonds, we calculated the density of state (DOS) of the boron atoms with appropriate cluster models using the discrete variational (DV) $-X\alpha$ method.^{26,27} The calculated DOS was compared to the measured BK X-ray emission spectra. The upper panel of Fig. 3 shows the hydrogen-terminated cluster model, BC₁₄₆(H₁₄₈), for CVD-B-diamond where one B atom is substituted for the C atom bonded to the centered C atom in the diamond C₁₄₇H₁₄₈ cluster. The lower panel shows the occupied 2*p*- and 2*s*-DOS of the B atom and the centered C atom (denoted by asterisks in the figure) as well as compares them to the measured BK X-ray emission spectra of CVD-B-diamond. The B2*p*-DOS well reproduced the measured spectral profile. The B2*p*-DOS consisted of a main peak at -3 eV in the MO energy, a higher-energy shoulder at -1 eV, and lower-energy tailing. The high-energy shoulder in the B2*p*-DOS corresponds to the steep edge at 185 eV in the measured spectrum. The B2*p* orbitals at the higher-energy shoulder hybridize with the highest C2*p* orbitals, and the hybridized DOS can form the acceptor level in both the B and C atoms.⁶ The lower-energy tailing in the B2*p*-DOS is mainly formed by the hybridization of B2*p* with B2*s* orbitals. Thus, the BK X-ray emission spectral profile of

CVD-B-diamond can be understood from the DOS of the B atoms in the diamond lattice. Hence, it is confirmed that the dopant B atoms in the CVD-B-diamond can substitute for carbon atoms in the diamond lattice to form covalent B-C bonds.

The upper panel of Fig. 4 shows the hydrogen-terminated cluster model, $B_{48}N_{48}(H_{24})$, for *h*-BN,²⁸ which was observed in the HPT-B-diamond. The lower panel shows the calculated occupied *2p*- and *2s*-DOS of the centered B and N atoms (denoted by asterisks in the figure) as well as compares them to the measured BK X-ray spectra of HPT-B-diamond and *h*-BN. The B2*p*-DOS well reproduced the measured spectral profile. The B2*p*-DOS consisted of a main peak at -5 eV, a higher-energy shoulder around -3 eV, a lower-energy shoulder around -9 eV, and a lower-energy satellite around -16 eV. The main peak and the higher-energy shoulder are mainly formed by the hybridization of B2*p* with the N2*p* orbitals. The lower-energy shoulder can be formed by the hybridization of B2*p* with B2*s* and N2*p*, while the lower-energy satellites are mainly formed by the hybridization of B2*p* with N2*s*. This satellite is a characteristic of the sp^2 -coordinated B atoms in *h*-BN. Therefore, the BK X-ray emission spectral profile of HPT-B-diamond can be understood from the DOS in *h*-BN. Hence, it is confirmed that the dopant B atoms in the HPT-B-diamond can react with impurity N atoms in the diamond lattice to form *h*-BN clusters.

Conclusion

To analyze the local structures and/or chemical states of boron atoms in B-diamonds, which can be synthesized by the CVD method and the HPT method, the soft X-ray emission spectra in the BK and CK regions of the CVD-B-diamond and HPT-B-diamond were measured using synchrotron radiation at the ALS. Although the CK X-ray profiles of the CVD-B- and HPT-B-diamonds exhibited typical diamond profiles, CVD-B and HPT-B diamonds had different BK X-ray profiles. Spectral analysis using a fingerprint method suggested that the boron atoms in the HPT-B-diamond reacted with the impurity nitrogen atoms to form boron nitride in HPT-diamond. To clarify the local structure and/or chemical states of the boron atoms in the B-diamonds, the BK X-ray emission profiles of the CVD-B- and HPT-B-diamonds were compared to the B2*p*-DOS of the appropriate cluster models. The BK X-ray emission profile of CVD-B-diamond successfully explained the boron atoms substituting for

the carbon atoms in the diamond lattice, while *h*-BN profile explained the profile of HPT-B-diamond. Thus, it is elucidated that impurity nitrogen in the diamond interferes with the formation of covalent B-C bond formation in the diamond lattice, and that the CVD method is preferable for doping boron atoms into diamond in order to develop *p*-type B-diamond semiconductors.

Acknowledgments

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Figure Caption

Fig. 1 CK X-ray emission spectra of B-doped diamonds (CVD-B- and HPT-B-diamonds) and reference compounds of CVD-diamond, HPT-diamond, and HOPG.

Fig. 2 BK X-ray emission spectra of B-doped diamonds (CVD-B- and HPT-B-diamonds) and reference compounds of *h*-BN, *c*-BN, B₄C, and *a*-B.

Fig. 3 Upper panel shows the cluster model (BC₁₄₆H₁₄₈) for B-doped diamond where the terminating H atoms are not shown. Lower panel shows the calculated *2p*- and *2s*-DOS of the B and centered C atoms (denoted by asterisks in the cluster model) compared to the BK X-ray emission spectrum of CVD-B-diamond.

Fig. 4 Upper panel shows the cluster model (B₄₈N₄₈H₂₄) for *h*-BN where the terminating H atoms are not shown. Lower panel shows the calculated *2p*- and *2s*-DOS of the B and centered N atoms (denoted by asterisks in the cluster model) compared to the BK X-ray emission spectrum of HPT-B-diamond and *h*-BN.

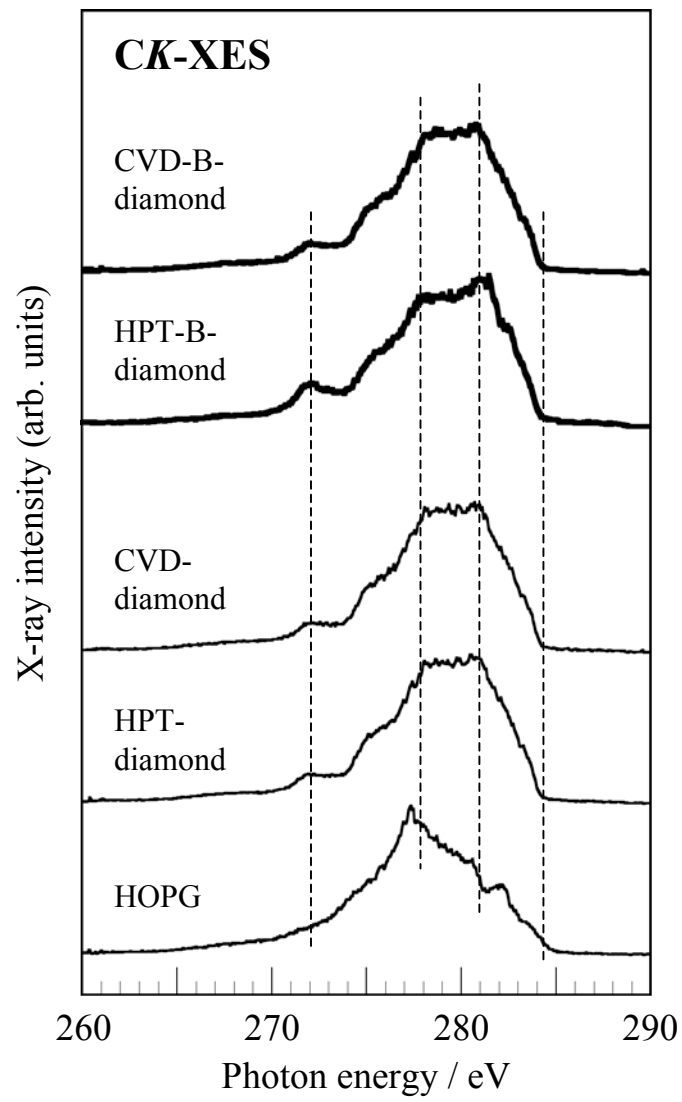


Figure 1 Y. Muramatsu et al.

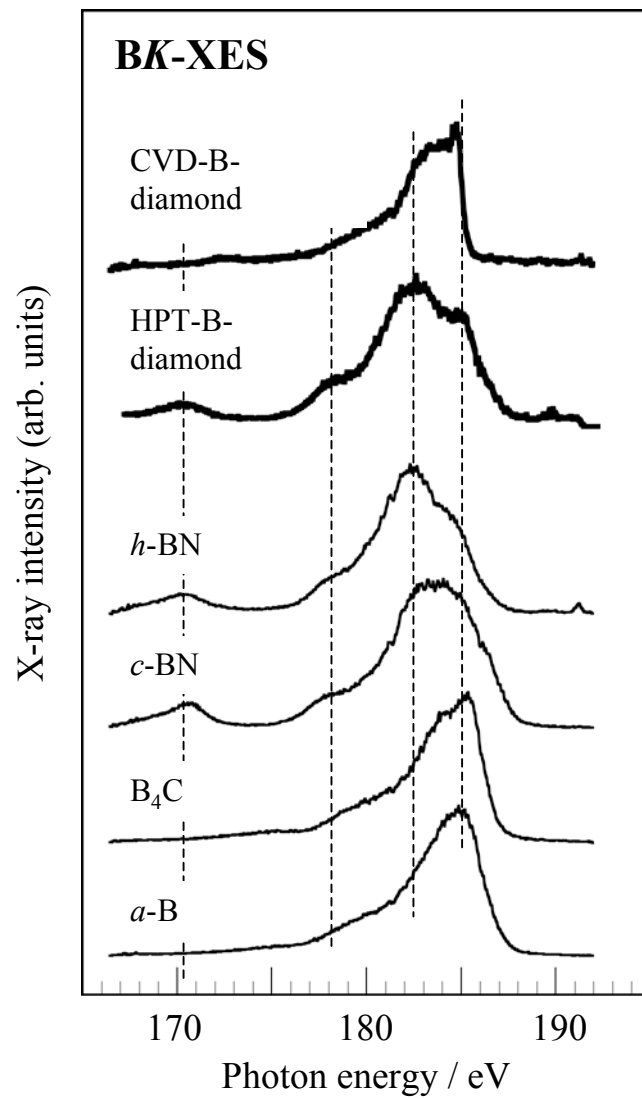


Figure 2 Y. Muramatsu et al.

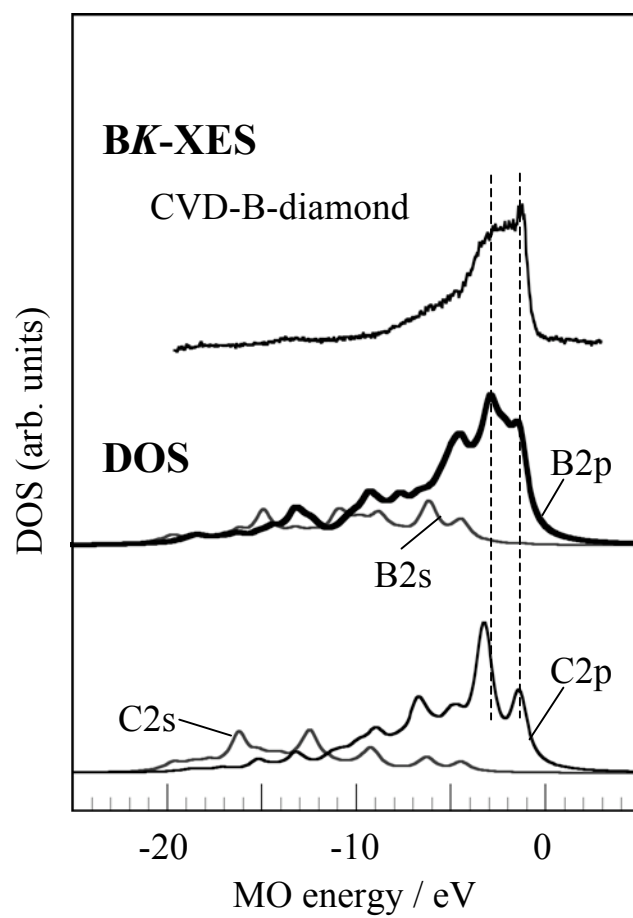
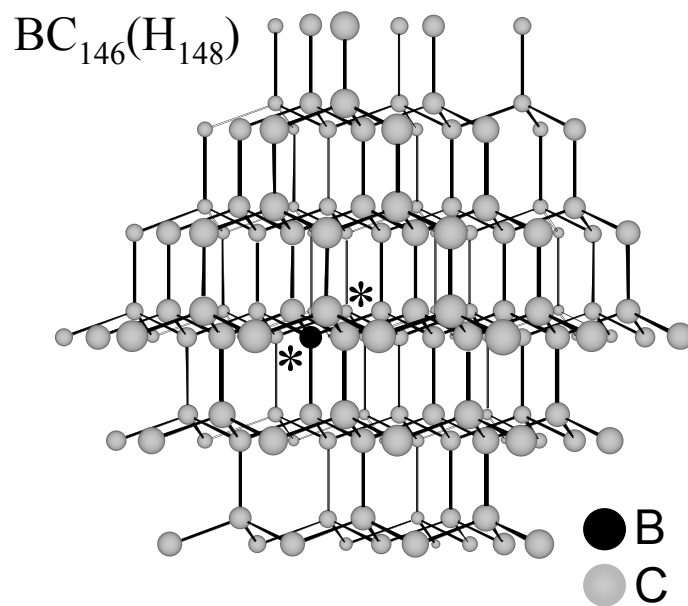


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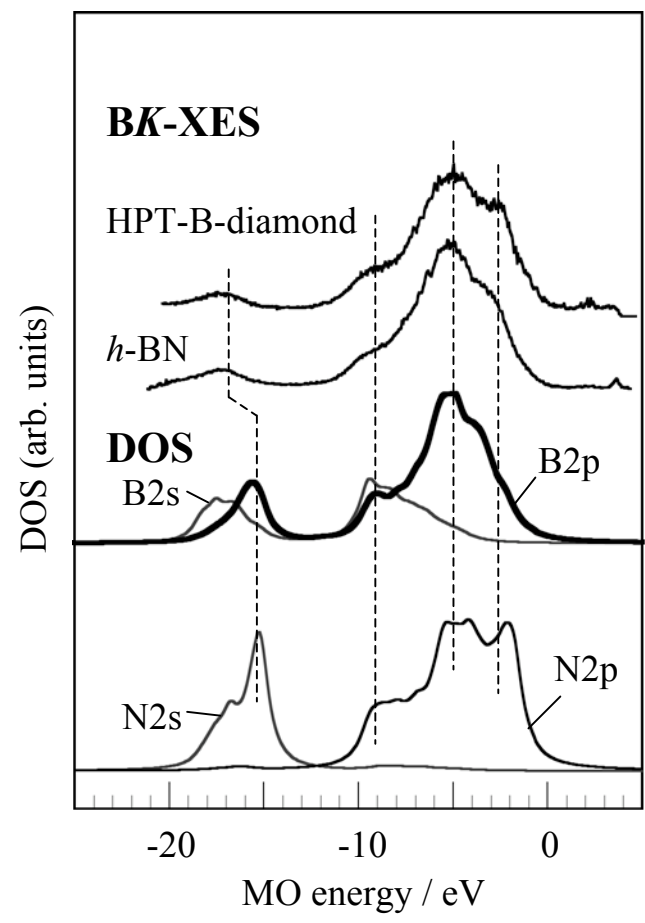
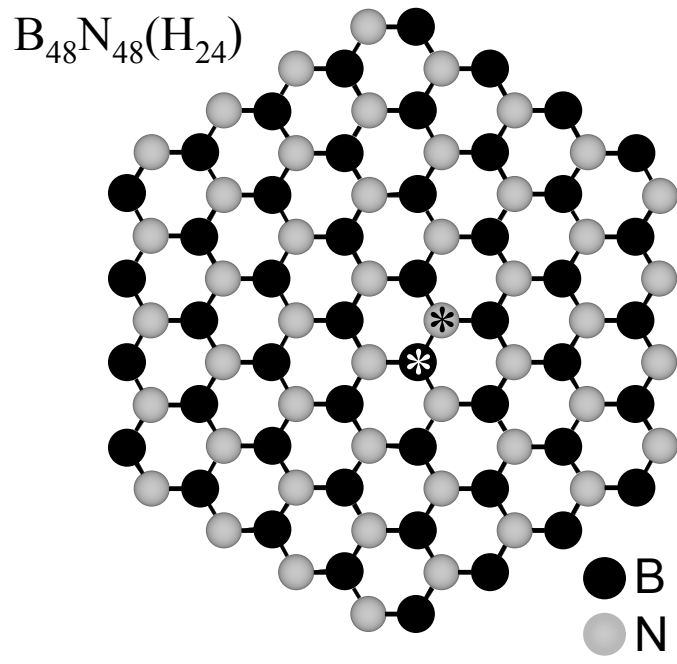


Figure 4 Y. Muramatsu et al.