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<td>Singh, Ram Sevak; Tay, Roland Yingjie; Chow, Wai Leong; Tsang, Siu Hon; Mallick, Govind; Teo, Edwin Hang Tong</td>
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Band gap effects of hexagonal boron nitride using oxygen plasma
Ram Sevak Singh, Roland Yingjie Tay, Wai Leong Chow, Siu Hon Tsang, Govind Mallick, and Edwin Hang Tong Teo

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Band gap effects of hexagonal boron nitride using oxygen plasma

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Tuning of band gap of hexagonal boron nitride (h-BN) has been a challenging problem due to its inherent chemical stability and inertness. In this work, we report the changes in band gaps in a few layers of chemical vapor deposition processed as-grown h-BN using a simple oxygen plasma treatment. Optical absorption spectra show a trend of band gap narrowing monotonically from 6 eV of pristine h-BN to 4.31 eV when exposed to oxygen plasma for 12 s. The narrowing of band gap causes the reduction in electrical resistance by ~100 fold. The x-ray photoelectron spectroscopy results of plasma treated hexagonal boron nitride surface show the predominant doping of oxygen for the nitrogen vacancy. Energy sub-band formations inside the band gap of h-BN, due to the incorporation of oxygen dopants, cause a red shift in absorption edge corresponding to the band gap narrowing. © 2014 AIP Publishing LLC. [http://dx.doi.org/10.1063/1.4872318]

Two-dimensional (2D) hexagonal boron nitride (h-BN) materials have received enormous interests due to its intriguing electronic, thermal, and mechanical properties.1 Due to the similarity in lattice constants, the hybridized structures of h-BN and graphene can be used as composite electronic devices.2,3 Since, h-BN is an insulator with a large direct band gap of almost 6 eV,4 there has been a great deal of interest from physicists, chemists, material scientists, and engineers to reduce and/or to manipulate the energy to achieve desired future electronic applications. Energy gap of h-BN can be tuned using various strategies, including atomic doping, composites, and surface functionalization.2,5,6 Recently, it was predicted that optical, electrical, and magnetic properties of boron nitride can be tuned by oxygen substitution using theoretical calculations.7–9 In this Letter, we present our observations of band gap narrowing in a few layers of h-BN with respect to doping of oxygen atoms into the lattice of h-BN via a simple, economical, and environmentally safe method of oxygen plasma treatment. Though, there have been a few attempts to alternate the h-BN band gap using carbon doping5 and through hydrogenation,10 experimental observation of oxygen induced reduction of the gap is not reported. Optical absorption measurements exhibit a monotonically narrowing of band gap in oxygen plasma treated h-BN samples, indicating the decrease in magnitude with increase of oxygen concentration. Furthermore, the decrease of band gap results in a reduction (~100 fold) of electrical resistance of oxygen doped h-BN films as substantiated by electrical measurements. X-ray photoelectron spectroscopy (XPS) suggests that h-BN under O2-plasma undergoes the changes in surface structure by creation of nitrogen (N) vacancies enabling the substitution of N-sites by oxygen atoms. This is generally symbolized as O_N-substitution doping. Tunable band gap of h-BN, realized in this study, could be a platform towards applications in nano- or opto-electronic devices.

h-BN used in this study was prepared by chemical vapor deposition (CVD) on copper substrate (25 μm thick, purchased from Alfa Aesar) using ammonia borane (Sigma Aldrich) as precursor.11,12 Prior to the deposition, copper substrate was submersed into dilute nitric acid followed by rinsing it using deionized (DI) water. Synthesis of h-BN involved annealing of copper substrate with two major steps: (1) Cu substrate was heated to 800 °C for 20 min in Ar/H2 (425:75 sccm), and (2) it was gradually heated up to 1050 °C for 40 min under the vapor of ammonia borane. 5 mg of the precursor, ammonia borane, was placed in a ceramic holder within the quartz tube and was heated at 60 °C with a flexible heating belt to vaporize it. As-grown h-BN on copper substrate was investigated using scanning electron microscopy (SEM; LEO 1550 Gemini) as shown in Fig. 1(a). In order to determine the thickness, the synthesized h-BN was transferred onto SiO2 (285 nm)/Si substrate.13 A thin layer of poly(methyl methacrylate) (PMMA) was spin-coated on top of the h-BN/Cu substrate, baked at 105 °C for 2 min, and subsequently put on Fe(NO3)3 solution for few hours to etch away the Cu. The free standing PMMA coated h-BN was then rinsed in DI water for several times and extracted onto SiO2/Si or quartz substrate. Finally, the sample was submersed in acetone to remove the PMMA. Atomic force microscopy (AFM) was used to determine the thickness of as-grown h-BN films. Fig. 1(b) shows an AFM image of a continuous h-BN film on SiO2/Si substrate with measured thickness of 1.89 nm.

Ultraviolet-visible (UV-Vis) spectroscopy is a powerful tool to investigate the optical properties such as absorption, transmittance, and optical band gap of materials. In order to

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conduct UV-vis absorption measurements, the h-BN film was transferred on optical quartz substrate from copper substrate using the methods described above. Two identical quartz substrates were used for base line corrections. The measurements were conducted by a UV-Vis (Shimadzu UV-2450) system at room temperature and in ambient conditions.

The h-BN sample was exposed to oxygen plasma cumulatively for 3 s, 6 s, and 12 s. The experiments were carried out using a commercial plasma system (AST Clen100 Descum) with RF power of 25 W. O₂-plasma was chosen for oxygen substitution because atomic oxygen can generate and directly substitute nitrogen atom at a site where a B–N bond is broken. Since, the binding energy of a hexagonal B–N bond is 7–8 eV, sufficient number of electrons or ions of at least this amount of energy is needed to break these bonds while adjusting the total energy so as to minimize the damage to the surface of h-BN. O₂-plasma treatment over ultrathin layers of h-BN is a crucial process. We managed to implement this by using low power (25 W) and varying the exposure time of the plasma. The power used in our study can be considered “mild” as compared to the power (100 W) used previously for BN nanotubes. Furthermore, this power (25 W) is sufficient to generate ions with its bombardment energy >10 eV, which is strong enough to break a B–N bond in a h-BN system. The effect of O₂-plasma modification over h-BN films was investigated using UV-Vis spectroscopy, electrical resistance measurements, and XPS.

Fig. 2 shows the UV-Vis absorption spectra acquired from O₂-plasma treated h-BN films corresponding to exposure times of 3 s, 6 s, and 12 s, respectively. The inset shows the spectrum obtained from a pristine (before exposure) h-BN sample. The spectrum of the pristine h-BN shows a prominent absorption at round 202 nm which is the result of inter-band transition from valence to conduction band of the h-BN. This absorption band relates to the optical band gap in h-BN. Under O₂-plasma, it is evident that the spectrum undergoes significant changes in absorption edges, indicating a band gap narrowing in h-BN. h-BN is a direct band gap semiconductor and the derived formula to estimate its band gap is given by its absorption coefficient

\[ \alpha = \frac{A}{d} = \left( \frac{E - E_g}{E} \right)^{1/2} / E, \]

where \( \alpha \) is the absorption constant, \( A \) is the absorption coefficient, \( d \) is the thickness of the film, \( E \) is the energy of the photon, \( E_g \) is the band gap of the material, and \( E \) is the photo energy.

The h-BN sample was exposed to oxygen plasma cumulatively for 3 s, 6 s, and 12 s, respectively. The inset shows the spectrum of pristine (before exposure) h-BN sample. (b) Tauc’s plots obtained from (a) for the determination of band gaps.

FIG. 2. (a) UV-Vis absorption spectra acquired from h-BN samples exposed to O₂-plasma for 3 s, 6 s, and 12 s, respectively. The inset shows the spectrum of pristine (before exposure) h-BN sample. (b) Tauc’s plots obtained from (a) for the determination of band gaps.
700 μm in diameter while the maximum analysis depth lay in the range of 4–8 nm. Charge correction was made based on adventitious C1s (C-C/C-H) at 285.0 eV using standard software. For chemical state analysis, a spectral deconvolution was performed by a curve-fitting procedure based on Lorentzian broadened by a Gaussian. The XPS N1s and B1s spectra of the h-BN before and after exposure to O2-plasma are shown in Fig. 4. Before exposure, B1s spectrum (Fig. 4(e)) shows a peak at 190.7 eV (Ref. 8) with a very weak additional peak, labelled “BOxNy”, at higher energy (191.8 eV). After exposure, a pronounced peak of BOxNy at 191.8 eV was observed (Fig. 4(f)). This BOxNy peak corresponds to boron atoms bound to oxygen atoms (replacement of nitrogen) as labeled in the schematic diagram in Fig. 4(b). Substitution doping mechanism was confirmed in our analysis by analysing the N1s (398.3 eV) and B1s (190.7 eV) spectra before (Figs. 4(c) and 4(e)) and after exposure (Figs. 4(d) and 4(f)). After O2-plasma exposure, it was observed that total number of B changed a little, while content of N was reduced indicating the creation of nitrogen vacancy (VN) accompanied by an increase in oxygen content. The relative compositional change (substrate related elements or chemical states are excluded) in h-BN before and after exposure is presented in Table I. As such, the observations suggest that the oxygen plasma breaks the B–N bond to create the VN where an oxygen atom is replaced. This phenomenon is supported by the theoretical works9,16,17 where O substitution for N is more favorable due to the lower formation energy.

For electrical measurements, h-BN devices were fabricated using photolithography process, metal deposition, and oxygen plasma etching. Two Cr (10 nm)/Au (50 nm) metal electrodes were deposited using electron beam evaporator. To define the channel, h-BN film was etched under oxygen plasma. The fabricated h-BN device, in this study, is shown in the inset of Fig. 5. We performed the electrical resistance measurements of pristine h-BN device before and after exposing to oxygen plasma for the duration of 3 s, 6 s, and

FIG. 3. A plot showing the change in band gap of h-BN with respect to oxygen plasma exposure time.

FIG. 4. Chemical structure of untreated (before exposure to O2-plasma) and treated (after exposure) h-BN samples. A schematic showing the structure of h-BN (a) before exposure, and (b) with oxygen doping after exposure. (c) XPS N1s spectrum before exposure. (d) XPS N1s spectrum after exposure. (e) XPS B1s before exposure. (f) XPS B1s spectrum after exposure.
12 s, respectively. All the measurements were conducted at room temperature and at ambient conditions using Keithley 4200 semiconductor characterization system. Fig. 5 shows the measured resistances of h-BN with respect to oxygen plasma exposure times. It is evident that h-BN under O₂-plasma undergoes a change in resistance from 2.4 GΩ (0 s or pristine h-BN) to 0.02 GΩ (sample exposed for 12 s) which is ~100 fold reduction in resistance compared to unexposed h-BN sample. The reduction in resistance due to oxygen doping is the indication of a band gap narrowing phenomenon in h-BN. The resistance of 0.02 GΩ observed is due to the saturated 4.31 eV energy gap at 12 s. Nonetheless, the trend of cumulative reduction in resistance supports our observations of optical band gap narrowing effects in UV-Vis spectra (Fig. 2).

In summary, the narrowing of band gap in a CVD grown h-BN using oxygen plasma treatment is realized. Optical absorption spectra show a reducing band gap trend from 6 eV of pristine h-BN to 4.31 eV of h-BN exposed for the duration of 12 s. The narrowing of band gap is also responsible for the reduction in electrical resistance (~100 fold) of exposed h-BN film. XPS measurements reveal that h-BN under O₂-plasma undergoes the breaking of B-N bond, creating nitrogen vacancy followed by oxygen doping. This chemical doping of h-BN results in the formation of a sub-band inside the band gap and causes a red shift in absorption edge corresponding to the band gap narrowing. The study could be promising to modulate the electronic, magnetic, and optical property of h-BN material towards future nano-electronic device applications.

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