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High-resolution 14-MeV neutron spectrometer
A diamond 14 MeV neutron energy spectrometer with high energy resolution

Takehiro Shimaoka,1,a) Junichi H. Kaneko,1 Kentaro Ochiai,2 Masakatsu Tsubota,1 Hiroaki Shimmyo,1 Akiyoshi Chayahara,3 Hitoshi Umezawa,3 Hideyuki Watanabe,4 Shin-ichi Shikata,3,b) Mitsutaka Isobe,5 and Masaki Osakabe5

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A self-standing single-crystal chemical vapor deposited diamond was obtained using lift-off method. It was fabricated into a radiation detector and response function measurements for 14 MeV neutrons were taken at the fusion neutronics source. 1.5% of high energy resolution was obtained by using the 12C(n,α)9Be reaction at an angle of 100° with the deuteron beam line. The intrinsic energy resolution, excluding energy spreading caused by neutron scattering, slowing in the target and circuit noises was 0.79%, which was also the best resolution of the diamond detector ever reported. © 2016 AIP Publishing LLC. [http://dx.doi.org/10.1063/1.4940929]

I. INTRODUCTION

Diamond radiation detectors have promising properties that include high temperature operation,1,2 low leakage current, high breakdown voltage, and solar blindness3 because of the wide bandgap of 5.5 eV. They also have irradiation resistance against fast neutrons,4 fast response because of high mobility, and high saturation drift velocity.5,6 Furthermore, they are useful for ion temperature measurements based on the 14 MeV neutron measurement using the 12C(n,α)9Be reaction.7

Energy resolution of 2.15% for 12C(n,α)9Be reactions using a natural diamond detector was reported by Krasilnikov et al. in 1997.8 This energy resolution satisfies deuterium-tritium (DT) plasma ion temperature measurements. The method uses direct reactions between 14 MeV neutrons and carbon. No other reaction occurs below 1 MeV of the peak attributable to the 12C(n,α)9Be reaction. Therefore, it is possible to measure ion temperatures up to 50 keV.7 An ideal diamond radiation detector can achieve detection efficiency of 1.0 × 10−5 count/(unit neutron flux) with effective volume of 2.5 × 2.5 × 0.13 mm3.9 The detector satisfies DT plasma ion temperature measurements of 100 ms in International Thermonuclear Experiment Reactor (ITER)7,10 with small size.

Other measurement methods exist for plasma ion temperature based on 14 MeV neutron such as magnetic proton recoil (MPR) spectroscopy11 and a combination of recoil proton detection and neutron ToF spectroscopy, so-called TANSY.12 However, these measurements require large instruments of several meters. Particularly, MPR requires several meters of magnetic poles and coils. TANSY requires an approximately 1 m ring scintillator array. In stark contrast, a diamond radiation detector is only several square millimeters. Consequently, it presents benefits for plasma diagnostics, which are usually restricted by spatial limitations. Small diamond detector arrays of the future are expected to be able to supply spatial distribution measurements.

With respect to semiconductor detectors, conventional silicon surface barrier detectors have been widely used in Joint European Tours (JET), but radiation damage appears by 1011–1012 neutrons/cm2.13 The detectors endure larger neutron fluence in ITER. Moreover, the detector cannot exchange frequently once the experimental period starts. Consequently, silicon surface barrier detectors are difficult to use in such harsh conditions. However, diamond radiation detectors have 100 times radiation resistance compared with silicon because they have small displacement energy per atoms as a result of low atomic number products. Pillon et al. reported that diamond radiation detectors show stable energy spectra up to 2 × 1014 neutrons/cm2 of 14 MeV neutrons.14 This fluence is greater than the fluence which the detector irradiated during one year of ITER operation. Moreover, a diamond radiation detector is solar blind and is less gamma-ray sensitive than silicon because of its wide bandgap and low atomic number.

As described above, diamond radiation detectors have ideal properties for 14 MeV neutron spectrometry. However, detector grade natural diamond is scarce. Consequently, its application has remained limited. To produce spectrometric grade diamond routinely, chemical vapor deposition (CVD)
diamond has been applied to radiation detectors.\textsuperscript{14–18} Recent CVD growth technique has enabled routines production of spectroscopic grade diamond. Cazzaniga \textit{et al.} reported 2\%–3\% energy resolution for \textsuperscript{12}C(n,\alpha)\textsuperscript{8}Be reactions.\textsuperscript{17} Nonetheless, CVD diamond has 0\%–4\% of energy resolution for 5.486 MeV alpha particles. Energy resolution for \textsuperscript{12}C(n,\alpha)\textsuperscript{8}Be was worse considering neutron energy broadening caused by scattering and slowing in a tritide titanium target. Some room remains for improvement. Sato \textit{et al.} reported that this energy resolution reduction resulted from several percent of electron charge trapping.\textsuperscript{18–20} In 14 MeV neutron measurements, charge generation was uniform in the whole sensitive area. The contributions of induced charge were almost identical between holes and electrons. The measured peak was spreading due to electron charge collection process. Thus, energy resolution for \textsuperscript{12}C(n,\alpha)\textsuperscript{8}Be became significantly worse than the energy resolution for alpha particles.

This report describes the achievement of the best energy resolution for 14 MeV neutrons ever reported. A single-crystal CVD diamond radiation detector having excellent charge carrier transportation properties for both holes and electrons was synthesized and fabricated into a radiation detector in our laboratory. A response function for 14 MeV neutrons was measured at the fusion neutron source (FNS).

\section*{II. EXPERIMENTAL}

\subsection*{A. Crystal growth and detector preparation}

A single-crystal CVD diamond was grown homoepitaxially on a HPHT type-IIa diamond substrate with an off-angle of three degrees for the (110) direction to suppress abnormal growth. Before CVD growth, the substrate was implanted with carbon ions at energy of 3 MeV and fluence of $2 \times 10^{16}$ ions/cm\textsuperscript{2} to fabricate a graphite layer for lift-off processing.\textsuperscript{21,22} Then, CVD growth was done on the (001) direction using a microwave plasma reactor (ASTEx 5250; MKS Instruments, Inc.). We prepared two Hokkaido University’s samples: HU#1 and HU#2. The gas pressure, methane concentration, and microwave power were, i.e., 110 Torr, 0.25\%, and 750 W for HU#1 and 110 Torr, 0.20\%, and 1100 W for HU#2, respectively. The substrate temperature was kept 850–900 \degree C during CVD growth. A self-standing layer was lifted-off from the substrate by electrical chemical etching. The self-standing layer thickness and the growth rate were 70 \mu m and 0.36 \mu m/h for HU#1, and 150 \mu m and 0.86 \mu m/h for HU#2, respectively.

Then it was oxygen-terminated using dichromic acid. Subsequently, the titanium contact was fabricated on the substrate side of the crystal by evaporation. It was annealed at 400 \degree C for 30 min to form titanium carbide. Then the gold contact was fabricated on the titanium carbide. The titanium carbide/gold ohmic contact was connected to a Sub Miniature version A (SMA) receptacle using silver paste for bias and read out of signals. The aluminum Schottky contact was fabricated on as-grown side. Then it was connected to the ground.

Alpha-particle-induced charge distribution measurements were conducted for charge collection efficiency evaluation. We used 5.486 MeV alpha particles from an \textsuperscript{241}Am radioactive source, a charge-sensitive preamplifier (142A; Ortec), a spectroscopy amplifier (672; Ortec), and a multi-channel analyzer (WE7562; Yokogawa Analytical Systems, Inc.). Alpha-particles were injected into aluminum contact side. A silicon surface barrier detector (CU012-050-100; Ortec) was used as the standard for charge collection efficiency. Average energies for electron and hole pair creation of silicon and CVD diamond, i.e., 3.62 eV and 13.1 eV,\textsuperscript{23} were used for charge collection efficiency calculations.

In 14 MeV neutron measurements, a charge carrier generates in the whole sensitive area of the diamond detector. Typically, charge transportation properties of electrons were worse than those of holes for scCVD diamond. The full width at half maximum (FWHM) for the \textsuperscript{12}C(n,\alpha)\textsuperscript{8}Be reactions spreads due to worse charge collection efficiency and energy resolution of electrons. The detector with nearly perfect charge collection for both holes and electrons is required to achieve excellent energy resolutions. Table I presents charge collection efficiency and energy resolution for HU#1 and HU#2. High energy resolutions of 0.38\% were obtained for both holes and electrons. Electron charge loss was improved from 3\% to 0.2\% per 100 \mu m. Details of charge collection properties were reported elsewhere.\textsuperscript{24}

\begin{table}[h]
\centering
\begin{tabular}{|c|c|c|c|c|}
\hline
Sample I.D. & Hole & Electron & Hole & Electron \\
& & & & \\
HU#1 & 100.1 & 99.9 & 0.38 & 0.38 \\
HU#2 & 100.0 & 99.7 & 0.38 & 0.38 \\
\hline
\end{tabular}
\caption{Charge transportation properties of the single-crystal CVD diamond detector.}
\end{table}

\begin{figure}[h]
\centering
\includegraphics[width=0.5\textwidth]{setup.png}
\caption{Experimental setup for response function measurements at FNS 80° beam line. The deuteron beam energy, beam current, neutron yields were 350 keV, 0.5–1 mA and \textsuperscript{10}Be \textsuperscript{3} neutrons/s, respectively. A detector was settled from 30 to 56 cm from the target at an angle of 95°–110° with respect to the deuteron beam for suppressing energy broadening.}
\end{figure}
III. EXPERIMENTAL RESULTS AND DISCUSSION

Figure 2 depicts an example of response function for 14 MeV neutron obtained by HU#1. The angle formed with the deuterium beam was 100°. The FWHM of the peak attributable to the $^{12}$C$(n, \alpha)^{9}$Be reaction was 1.5%. The earlier best energy resolution was 2.15% obtained using natural diamond for the same neutron fields that Krasilnikov et al. reported in an earlier study. Although CVD diamond growth technology improved, no detector achieved better energy resolution. We achieved the best energy resolution as a 14 MeV neutron energy spectrometer in this study. Figure 3 shows the best energy resolution as a 14 MeV neutron obtained by HU#1 at an angle of 95°. The angle with the deuteron beam was 100°. The count rate was approximately 700 cps in maximum during the measurements.

TABLE II. Energy resolution and measurement conditions of DT neutron response function.

<table>
<thead>
<tr>
<th>Sample</th>
<th>Angle with the deuterium beam line (deg)</th>
<th>$\sigma_{Ex}$ (V/$\mu$m)</th>
<th>$\sigma_{Diamond}$ (V/$\mu$m)</th>
<th>Count rate (cps)</th>
<th>Electric field (V/$\mu$m)</th>
</tr>
</thead>
<tbody>
<tr>
<td>HU#1</td>
<td>110</td>
<td>1.9</td>
<td>0.8</td>
<td>300</td>
<td>1.3</td>
</tr>
<tr>
<td></td>
<td>100</td>
<td>1.5</td>
<td>0.8</td>
<td>400</td>
<td></td>
</tr>
<tr>
<td></td>
<td>95</td>
<td>1.7</td>
<td>0.79</td>
<td>200</td>
<td></td>
</tr>
<tr>
<td>HU#2</td>
<td>110</td>
<td>2.0</td>
<td>0.5</td>
<td>700</td>
<td>1.3</td>
</tr>
<tr>
<td></td>
<td>100</td>
<td>1.5</td>
<td>0.5</td>
<td>400</td>
<td></td>
</tr>
<tr>
<td></td>
<td>95</td>
<td>1.8</td>
<td>0.99</td>
<td>200</td>
<td></td>
</tr>
</tbody>
</table>

In that equation, $\sigma_{diam}$ is the intrinsic energy resolution of diamond detector. $\sigma_{ex}$ and $\sigma_{diam}$ denote energy broadening caused by neutron scattering and slowing in the target. $\sigma_{noise}$ represents the circuit noise. Krasilnikov et al. reported $\sigma_{ex}$ = 1.48% and $\sigma_{diam}$ = 0.23% at the angle of 95°. These values were estimated using TRIM code. The circuit noise estimated using a pulser was 0.14% for HU#1 and 0.12% for HU#2 in the present measurements. The intrinsic energy resolution calculated using Equation (1) was $\sigma_{diam} = 0.79%$ for HU#1 and 0.99% for HU#2. Krasilnikov described intrinsic energy resolution $\sigma_{diam}$ = 1.95% at an angle of 95° in a report of an earlier study. Cazzaniga reported $\sigma_{diam} = 2.5%$ obtained at an angle of 90° using CVD diamond. The intrinsic energy resolution obtained in the present work was the best ever reported.

Though the best energy resolutions were obtained, there were some inconsistencies between neutron energy calculations and experimental data. Krasilnikov et al. reported that the energy spreading of neutron minimizes at 95°, on the other hand, our experimental data show the smallest energy spreading at 100°.
intrinsic energy resolution was 0.47%. Nonetheless, the intrinsic energy resolution calculated by experimental data and neutron energy spreading considering neutron scattering and slowing only in the target was 0.79%. This difference can be caused due to scattering of assembly around the target. This neutron energy spread calculated by law of propagation of errors between calculation and experimental values was 0.6%.

To estimate intrinsic energy resolution more precisely, neutron transport calculation which includes detailed assembly around the target will be required as a future work.

Since there has not been a neutron energy spectrometer which has superior to 1.0% of high energy resolution until now, the diamond detector is expected to be applied for detailed comparison of neutron transport calculation with experiment.

IV. CONCLUSION

For this study, self-standing single-crystal CVD diamond radiation detectors were fabricated using lift-off method. Response function measurements were conducted at the FNS 80° beam line. The 1.5% high energy resolution was obtained at an angle of 100°. The intrinsic energy resolution excluding neutron scattering, slowing in the target and circuit noises was 0.79%, which was the best resolution ever reported. We will strive to achieve intrinsic energy resolution to ca. 0.4% for the 12C(n, α) 4He reaction in future works by improving the electron charge collection efficiency. Similarly, we will plan to compare neutron transport calculations including detailed assembly around the target with experiment to determine intrinsic energy resolution more precisely.

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10See www.iter.org for information about International Thermonuclear Experiment Reactor.


