

Development of spin-polarized slow positron beam using a new ^{68}Ge - ^{68}Ga /GaN positron source and its application for positron annihilation spectroscopy

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Positrons emitted from radioisotopes are longitudinally spin-polarized due to the parity non-conservation in the weak interaction. Positron annihilation studies have shown that the ferromagnetic band structure can be investigated using the Doppler broadening of annihilation radiation (DBAR) technique with spin-polarized positrons [1,2]. Considering the fact that various novel spin phenomena have recently been found to occur at surfaces/interfaces and in thin films, spin-polarized positron beams is useful in spintronics studies. We developed a highly spin-polarized beam from a new ^{68}Ge - ^{68}Ga source.

The spin polarization of positrons emitted from a positron source P is given by

$$P = \frac{v}{c} \frac{(1 - \cos\theta)}{2},$$

here v is the positron speed, c is the speed of light and θ is an open angle in a specific direction [3]. This equation suggests that positron sources with a higher energy end-point are better for generating highly spin-polarized positron beams.

The ^{68}Ge - ^{68}Ga positron source has relatively long half-life and higher energy end-point (288 days and 1.9 MeV). In addition, higher cross section ($0.465 \times 10^{-24} \text{ cm}^2$ at $E_{\text{proton}} = 20 \text{ MeV}$) of nuclear reaction $^{69}\text{Ga}(p, 2n)^{68}\text{Ge}$ allows the source production using ion beam. Figure 1 shows schematic diagrams of the source capsule. High-purity metal ^{69}Ga should be used as a target material, however, metal Ga is molten during irradiation because of its low melting point (29 °C). Such molten Ga reacts with the other metal components of the source capsule and eventually destroys the capsule. To avoid this problem, we propose GaN as a target material, although the amount of $^{69}\text{Ga}(p, 2n)^{68}\text{Ge}$ is decreased. To avoid metallization during irradiation, the GaN target was placed in a carbon tray. The target was directly irradiated with 20 MeV proton beam with a maximum current of 3.5 μA using the cyclotron at the Takasaki Advanced Radiation Research Institute. The efficiency of the source production was estimated to be 0.5 MBq/ $\mu\text{A}/\text{h}$ from the gamma ray energy spectrum. This was in good agreement with the results from our simulation. The irradiation was carried out over 400 h in total.

The positrons emitted from the GaN target were moderated by a tungsten mesh moderator. The slow positrons extracted by a grid electronode were transported to the sample position by an electrostatic system in order to avoid the depolarization during the beam transportation.

Consequently, a positron beam with a diameter of 5 mm and a flux of $5 \times 10^3 \text{ e}^+/\text{s}$ was obtained with the positron energy of $E=5\text{--}15 \text{ keV}$. The spin polarization of the positron beam was determined to be $47 \pm 8\%$ from the magnetic quenching of the positronium in fused silica [4]. Thus, it is important to use positron sources with higher energy end-points to generate highly spin-polarized positron beam.

The field-reversal asymmetry of the DBAR spectra for ferromagnetic Fe was measured using the positron beam. Figure 2 shows the differential DBAR spectra in a magnetic field of 0.55 T [4]. The differential DBAR spectra [$N_{\uparrow}(p)$ and $N_{\downarrow}(p)$] were obtained by altering the field polarity. The finite

differential intensity means that there is a field-reversal asymmetry, which arises from the enhanced annihilation between the spin-up positrons and spin-down 3d unpaired electrons [5]. This also shows that our positron beam was sufficiently spin-polarized.

In summary, we produced a ^{68}Ge - ^{68}Ga /GaN source by irradiating a GaN target with proton beam and a spin-polarized positron beam was generated using this source. Measurements of the DBAR spectra of fused silica and ferromagnetic Fe in magnetic field demonstrated that the beam was sufficiently spin-polarized.

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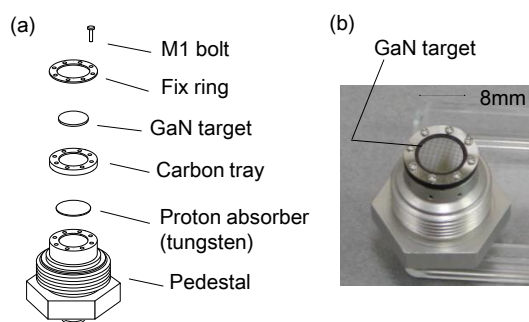


Fig.1 (a) Schematic and (b) photograph of the source capsule.

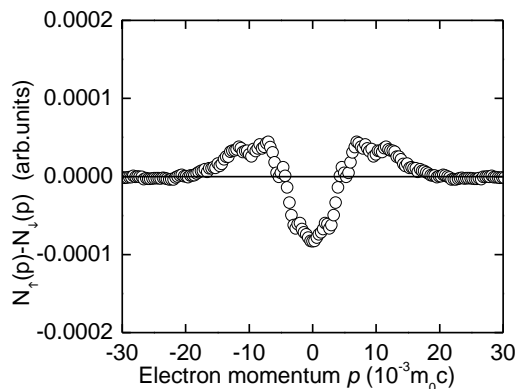


Fig.2 Differential DBAR spectra of Fe sample obtained in the external magnet field of 0.55 T.

References

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