## In situ photoelectron spectra of an electron-beam irradiated C<sub>60</sub> film

J. Onoe<sup>1</sup>\*, T. Ito<sup>2</sup>, S. Kimura<sup>3</sup>, H. Shima<sup>4</sup> and H. Yoshioka<sup>5</sup>

<sup>1</sup>Research Laboratory for Nuclear Reactors and Department of Nuclear Engineering, Tokyo Institute of Technology, Tokyo 152-8550, Japan

<sup>2</sup>Department of Materials, Physics and Energy Engineering, Nagoya University, Nagoya 464-8601, Japan

<sup>3</sup>UVSOR Facility, Institute for Molecular Science, and School of Physical Sciences, The Graduate University for Advanced Studies (SOKENDAI), Okazaki 444-8585, Japan
<sup>4</sup>Department of Applied Physics, Hokkaido University, Sapporo 060-8628, Japan
<sup>5</sup>Department of Physics, Nara Women's University, Nara 630-8506, Japan

We have found that electron-beam (EB) irradiation of a C<sub>60</sub> film gives rise to formation of a with peanut-shaped  $C_{60}$ polymer metallic electron-transport properties in air at room temperature [1]. The temperature dependence of the photo-excited carriers lifetime for the peanut-shaped polymer indicated the energy gap formation at below 50 K in a similar manner to the Peierls instability for quasi-one-dimensional (1D) metallic materials such as  $K_{0,3}MO_3$  [2], thus suggesting that the polymer is a 1D metal as illustrated in Fig. 1.

The 1D peanut-shaped polymer is fascinating from a viewpoint of topology, because it has both positive and negative Gaussian curvatures ( $\kappa$ ) lined alternatively and periodically. As shown in Table 1, this nanocarbon can be classified into a new  $\pi$ -electron conjugated carbon allotrope that is different from graphite ( $\kappa = 0$ ), fullerenes ( $\kappa > 0$ ), nanotubes ( $\kappa = 0$  at body,  $\kappa > 0$  at cap edge), and hypothetical Mackay crystal ( $\kappa < 0$ ). Accordingly, the 1D peanut-shaped polymer is expected to exhibit physical and chemical properties different from those of the conventional  $\pi$ -electron conjugated carbon materials.

We have recently examined the valence photoelectron spectra of the polymer, using *in situ* high-resolution ultraviolet photoelectron spectroscopy [3, 4], and observed the Tomonaga-Luttinger liquids (TLL) behavior as the direct evidence for 1D metal and obtained the TLL exponent ( $\alpha$ ) to be ca. 0.6 [5], which is somewhat larger than that of ca. 0.5 for 1D metallic single-walled carbon nanotubes [6]. Using the Schrodinger equation dealing with the motion of free particles on a curved surface modulated by positive and negative Gaussian curvatures periodically and alternatively, we have first demonstrated that the increase in the exponent value is caused by a curvature-induced effective potential that works for electrons conducting along the curved surface [7].

To our best knowledge, the peanut-shaped  $C_{60}$  polymer is only an existed material with a negative Gaussian curvature, whose electronic and optical properties are revealed. Thus we believe that the present system will open a new field of "quantum

science of condensed matters in Liemannian space".



Fig. 1. Schematic illustration of one-dimensional peanut-shaped  $C_{60}$  polymer.

Table 1. Classification of  $\pi$ -electron conjugated carbon materials using Gaussian curvature.

	0
Material	Gaussian curvature (K)
Graphite	0
Fullerenes	> 0
Nanotubes	0 (body), $> 0$ (capped edge)
Mackay crystal	< 0
Peanut-shaped polymer $> 0, < 0$	

\*E-mail: jonoe@nr.titech.ac.jp

[1] J. Onoe, T. Nakayama, M Aono, and T. Hara, Appl. Phys. Letters **82** (2003) 595.

[2] Y. Toda, S. Ryuzaki, and J. Onoe, Appl. Phys. Letters **92** (2008) 094102.

[3] J. Onoe et al., Phys. Rev. B 75 (2007) 233410.

[4] J. Onoe, T. Ito, and S. Kimura, J. Appl. Phys. **104** (2008) 103706.

[5] T. Ito, J. Onoe, H. Shima, H. Yoshioka, and S. Kimura, in preparation for submission.

[6] H. Ishii et al., Nature 426 (2003) 540.

[7] H. Shima, H. Yoshioka, and J. Onoe, Phys. Rev. B **79** (2009) 201401 (R).