# Fragmentation processes of C<sub>60</sub> in multiple electron loss and capture collisions of 2-MeV Si<sup>2+</sup>

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An experimental study of ionization and fragmentation of  $C_{60}$  has been performed for 2-MeV Si<sup>2+</sup> incident ions under charge-changing conditions. Time-of-flight (TOF) spectra of product ions and number distributions of secondary electrons  $(n_e)$  were measured in coincidence with charge-selected outgoing projectiles. TOF spectra from transiently formed highly charged  $C_{60}^{r+}$  ions are obtained for individual charge states r, and  $n_e$ distributions correlated with size-fixed  $C_m^+$  ions are also obtained for m < 15. It is found that mass spectra and  $n_e$  distributions are both significantly different between loss and capture collisions, whereas a remarkable similarity in the partial  $n_e$  distributions is observed almost independently of the different charge-changing conditions when compared at given  $C_m^+$  ions size. We find also that a number of secondary electrons as high as 20 are emitted in the production of small fragment ions. Furthermore, small fragment ions are produced predominantly in close collisions even at very low charge states of  $r \approx 3$  in comparison with distant collisions. This indicates that the energy partition rate between ionization and internal excitation might be considerably different in close and distant collisions.

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## I. INTRODUCTION

In the last decade, considerable effort has been devoted to study of collision-induced ionization and fragmentation of C<sub>60</sub> to achieve microscopic understanding of excitation and relaxation mechanisms involving polyatomic molecules and clusters (see 1 and references therein). The remarkable high stability against Coulomb explosion is one of the outstanding properties of C<sub>60</sub> and has been verified in several different experiments using electron impact [2], slow highly charged ion (SHCI) impact [3-6], and strong laser fields [7]. The charge state of intact  $C_{60}^{r+}$  ions observed in these investigations reaches up to  $r \sim 9$  with lifetimes of the order of microseconds. In addition, remarkably high charge states of fullerene ions as high as r=12 were observed in recent experimental work using an intense femtosecond laser [8]. On the other hand, small-sized fragment ions resulting from multifragmentation processes are, by and large, observed in fast heavy-ion collisions [9–15]. Moreover, we found recently that multifragmentation occurs at very low charge states of  $r \sim 3$  in 2-MeV Si<sup>2+</sup> collisions when measured in coincidence with single-electron loss and capture collision events [16].

The experimental findings described above indicate clearly that the stability of a highly charged  $C_{60}^{r+}$  ion is basically dependent on its charge state *r* but also affected significantly by the amount of its internal excitation energy after interactions. This is because the internal excitation energy, denoted by  $E_{int}$  hereafter, is expected to be greatly different in the different kinds of experiments described above. Actually,  $E_{int}$  of a highly charged  $C_{60}^{-12+}$  ion produced by a laser is estimated to be only 15 eV [8]. Also, in SHCI collisions,  $E_{int}$  is supposed to be small because the predominant

production mechanism of highly charged  $C_{60}^{r+}$  ions is multiple-electron capture taking place in distant collisions [3,5]. By contrast, electron loss and capture collisions of fast heavy ions are rather close-encounter processes and, consequently, a much larger amount of  $E_{int}$  can be deposited into the molecule even if its charge state r is very small as observed experimentally [16]. It is, therefore, important to know to what extent the fragmentation pattern is affected by the charge state and the internal excitation in fast ion collisions.

Campbell *et al.* [17] measured fragment ion distributions by using a femtosecond laser. They observed the fragmentation pattern to change from the ionization regime (r) to the internal excitation regime  $(E_{int})$  as a function of the laser pulse duration ranging from 25 fs to 5 ps. Unlike laser experiments of this kind, however, ionization and excitation by energetic heavy-ion impact are essentially simultaneous processes and will appear in widely different manners according to a variety of combinations among projectile species, incident energy, and charge state. For the purpose of close inspection of these dynamical behaviors of C<sub>60</sub> fragmentation, a triple-coincidence technique [5,16] serves as a powerful tool, allowing us to obtain three kinds of information simultaneously about fragment ion distributions, number distributions of secondary electrons, and outgoing charge states of projectile particles. Actually, we found evidence for the predominant importance of internal excitation over the apparent charge state r in single-electron loss and capture collisions of 2-MeV  $Si^{2+}$  ions [16]. In that paper we pointed out that the number of free electrons  $(n_i)$  emitted from a C<sub>60</sub> molecule is in practice very important to examine the degree of C<sub>60</sub> fragmentation because the value of  $n_i$  is closely related to the amount of  $E_{int}$ .

In this work, we extended measurements to multipleelectron loss and capture collisions, in which r and  $E_{int}$  are both expected to be considerably large compared with singleelectron processes. It is emphasized that the present triple-

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FIG. 1. Schematic of experimental setup of the TOF  $n_e$  coincidence measurements.

coincidence method provides information systematically about fragmentation profiles according to individual values of  $n_i$  and r, enabling us to ascertain the relative significance of r and  $E_{int}$  in C<sub>60</sub> multifragmentation.

#### **II. EXPERIMENT**

The experiment was performed at the QSEC tandem accelerator facility of Kyoto University. A schematic diagram of the experiment is shown in Fig. 1. A beam of 2-MeV Si<sup>2+</sup> extracted from the accelerator was collimated to about 0.2  $\times 0.5$  mm<sup>2</sup> with two sets of four-jaw slits. Impurity ions of undesirable charge states produced in collisions with residual gases were removed by a charge selector consisting of four magnetic deflectors located just before a crossed-beam collision chamber. The charge-selected beam then crossed an effusive molecular beam target of  $C_{60}$  in the collision chamber. A base pressure of the chamber was below  $1 \times 10^{-6}$  Pa. The C<sub>60</sub> target was produced by sublimation of high-purity (99.98%) powder at 550 °C in a quartz oven with an exit nozzle of 2 mm in diameter. The C<sub>60</sub> beam was collimated by a thin aperture of 1 mm in diameter at 10 mm below the projectile beam axis. The diameter of the C<sub>60</sub> beam in a collision region was estimated to be smaller than 4 mm which was small enough to detect nearly all the product ions and secondary electrons.

Fragment ions as well as intact ions produced in collisions were extracted by an electrostatic field of 615 V/cm perpendicular to both the incident beam and the  $C_{60}$  beam, and were detected by a two-stage multichannel-plate (MCP) detector with a front voltage of -4.6 kV. The mass-to-charge ratio of product ions was measured with a time-of-flight (TOF) spectrometer operated under Wiley-McLaren spatial-focusing conditions [18]. Secondary electrons were extracted in the opposite direction to the positive ions and collected by a focusing lens into a semiconductor solid-state detector  $(SSD_e)$  with an active area of 150 mm<sup>2</sup>. A positive voltage of 30 kV was applied on the detector. Hence, the pulse heights of signals from the  $SSD_{\rho}$  are proportional to the total energy  $(30n_e \text{ keV})$  of electrons when  $n_e$  electrons are detected simultaneously. The pulse height distribution provides, therefore, the number distribution of secondary electrons.

Outgoing projectile particles of  $Si^{q+}$  were charge separated by an electrostatic deflector and detected by a movable



FIG. 2. Comparison of TOF spectra for 2e loss collisions measured at two scattering angles: (a)  $\theta = 0 \pm 0.4$  mrad and (b)  $0.8 \pm 0.4$  mrad.

solid-state detector  $(SSD_p)$  which had a rectangular entrance slit of 0.5 mm (horizontal)  $\times$  5 mm (vertical). The corresponding acceptance angles for scattered particles were  $\pm 0.4$  mrad (horizontal) and  $\pm 4$  mrad (vertical), respectively. Measurements were made at two  $SSD_p$  positions in the horizontal direction, corresponding to scattering angles of  $\theta=0$ and 0.8 mrad, respectively. This is because charge-changing collisions involving multiple electrons are supposed to take place at small impact parameters resulting in noticeable deflection of incident particles. Comparison between different scattering angles is, therefore, important to achieve a closer inspection of the TOF spectra. An example is shown in Fig. 2 obtained for two-electron-loss collisions. Obviously, two spectra are completely different and, in particular, only small fragment ions are produced at  $\theta = 0.8$  mrad. These angulardependent TOF spectra will be discussed later.

Triple-coincidence measurements of TOF spectra and  $n_e$  distributions at fixed final charge states q were performed by using the SSD<sub>p</sub> signals as start triggers. We investigated oneand two-electron capture (q=0,1) collisions and onethrough three-electron loss (q=3-5) collisions. Data were recorded two-dimensionally in  $1024 \times 1024$  channels for each charge-changing condition. Symbols used in this paper are as follows: r is the charge state of a prefragmented ion  $C_{60}^{r+}$ ,  $n_e$  is the number of electrons detected by the SSD<sub>e</sub>,  $n_i$ is the number of free electrons emitted from  $C_{60}$ , referred to as pure-ionization,  $n_c$  is the number of electrons captured by a projectile, and  $n_l$  is the number of electrons lost from a projectile.

Also, the following relationships hold:

$$n_e = n_i$$
,  $r = n_i + n_c = n_e + n_c$  for electron capture, (1)

$$n_e = n_i + n_l, \quad r = n_i = n_e - n_l \quad \text{for electron loss.}$$
(2)



FIG. 3. (Color) Two-dimensional coincidence maps of TOF and  $n_e$  spectra obtained for various charge-changing collisions as noted; 1*e*,2*e* capture collisions (Si<sup>2+</sup> $\rightarrow$ Si<sup>+</sup>,Si<sup>0</sup>) and 2*e*,3*e* loss collisions (Si<sup>2+</sup> $\rightarrow$ Si<sup>4+</sup>,Si<sup>5+</sup>). Insets show  $n_e$  spectra correlated with intact parent ions C<sub>60</sub><sup>r+</sup>.

# **III. EXPERIMENTAL RESULTS**

Figure 3 shows four examples of two-dimensional coincidence maps obtained for various charge-changing collisions. The vertical and horizontal axes represent, respectively, the mass-to-charge ratio of product ions and the pulse height of  $SSD_e$  signals, which corresponds to the number of secondary electrons  $n_e$  emitted simultaneously. An  $n_e$  distribution correlated with a given size of product ion is obtained from the data points lined up horizontally at the position of the corresponding TOF and will be called the partial  $n_e$  spectrum hereafter. Similarly, a partial time-of-flight spectrum correlated with the emission of  $n_e$  electrons is obtained from the data points lined up vertically at a position of  $n_e$ . A series of partial TOF spectra associated with fixed charge states r of prefragmented  $C_{60}^{r+}$  ions were derived by using Eqs. (1) and

(2). Total  $n_e$  distributions and total TOF spectra are obtained by projecting all these data points onto the two axes, respectively, as demonstrated in the figures.

The insets show the partial  $n_e$  spectra correlated with intact parent ions of  $C_{60}^{r+}$  (r=1-3). As described in our previous paper [16], analysis of these partial spectra of parent ions provides the electron-detection efficiency of the SSD<sub>e</sub> detector. This is because the value of  $n_e$  is uniquely determined from Eqs. (1) and (2) if all the emitted electrons as well as intact  $C_{60}^{r+}$  ions are detected safely. As shown by solid lines, all the  $n_e$  spectra can be reproduced almost perfectly by using fitting parameters of 0.17 as the backscattering probability and 0.6 as the backscattering K factor [16,19]. The electron-detection efficiency obtained in this way was 0.94, implying that nearly all the secondary electrons including also fast electrons (~39 eV) lost from a projectile ion are detected successfully in the present experiment. It is noted that all the partial  $n_e$  distributions used hereafter were deduced precisely by this fitting calculation carried out with the same fitting parameters mentioned above.

Overall features derived from Fig. 3 are as follows. First, the total TOF spectra are significantly different between loss and capture collisions. The dominant products are smallsized fragment ions in loss collisions but intact  $C_{60}^{r+}$  ions in capture collisions. In electron loss collisions, yields of small fragment ions  $C_m^+$  ( $m \leq 3$ ) were found to increase rapidly with increasing number of  $n_l$ . For instance, in collisions of 3eloss from  $Si^{2+}(2p^63s^2)$  to  $Si^{5+}(2p^5)$ , the spectrum is completely dominated by the smallest fragments of  $m \leq 3$  and exhibits no intact  $C_{60}^{r+}$  ions, which are still visible in 1*e* and 2e loss collisions. This implies certainly that, compared to the outer-shell ionization of 3s electrons, additional 2p ionization induces a larger amount of energy deposition, resulting in nearly complete cage disintegration. Unlike for loss collisions, spectra for 1e and 2e capture collisions do not show a noticeable difference in the TOF spectra except for intact parent ions.

Second, the total  $n_e$  spectra exhibit also strong dependence on the charge-changing condition. The spectra for electron loss collisions show clearly a large number of electrons as high as 20 to be emitted, whereas only a few electrons in capture collisions. More detailed comparisons between loss and capture collisions are shown in Fig. 4, where total and partial distributions are plotted as a function of the charge state *r* of prefragmented  $C_{60}^{r+}$  ions. The distributions show roughly double-peak structures attributable to intact  $C_{60}^{r+}$  ions and small fragment  $C_m^{+}$  ions. The main reason for this capture-loss difference in distribution shape is due to the difference in relative magnitude between these two components. Obviously the former component is confined to smaller values of r compared to the latter which reveals much larger r and broader peak profiles. One can see, therefore, that electron capture collisions, where intact ions are produced preferentially, are soft or distant collisions resulting in low charge states of intact ions. On the other hand, in electron loss collisions the latter component contributes predominantly, particularly for  $n_1 \ge 2$ .

Despite considerable broadness of distribution profiles, it is helpful to examine mean charge states  $\overline{r}$  for obtaining an overall picture of ionization and fragmentation. From the data of total  $n_{e}$  distributions the values of  $\overline{r}$  were obtained as  $\overline{r} \simeq n_c + 1.7$  ( $n_c = 1, 2$ ) for capture and  $\overline{r} \simeq 3.2n_l + 3.8$  $(n_l=1-3)$  for loss collisions. Hence, in loss collisions about 3.2 electrons are released additionally from  $C_{60}$  per each electron loss event, while no such additional ionization occurs in capture events. Using Eqs. (1) and (2), the average number of pure ionization events,  $n_i$ , can also be derived as  $\bar{n}_i = \bar{r} - n_c \approx 1.7$  for capture and  $\bar{n}_i = \bar{r} = 3.2n_l + 3.8$  for loss collisions. It is pointed out that  $n_i$  is supposed to be correlated closely with the amount of electronic energy deposition  $E_d$ [11,16], and once the impact parameter is given, both  $n_i$  and  $E_d$  may be determined. In this sense, the present result of completely different  $n_e$  distributions indicates that a specific charge-changing collision takes place preferentially in a spe-



FIG. 4. (Color) Total and partial distributions of secondary electrons as a function of the charge state r of prefragmented  $C_{60}^{r+}$  ions obtained in various electron capture and loss collisions.

cific impact parameter region. Therefore, the result of  $n_i \approx 1.7$  obtained for 1e and 2e capture collisions implies that both processes occur at nearly equivalent impact parameters,



FIG. 5. (Color) Mean charges associated with  $C_m^+$  as a function of the cluster size *m* obtained for charge-changing collisions.

while in loss collisions the impact parameter becomes smaller with increasing  $n_l$ .

In contrast to the total distributions, the partial  $n_{\rho}$  distribution correlated with a specific fragment ion reveals surprisingly little dependence on the collision condition. As a typical example, the distribution profile correlated with  $C_1^{+}$  is found to peak at nearly the same position of r=8-11 for all the collision processes including even capture events. The characteristics of this independency are demonstrated more clearly in Fig. 5 by plotting  $\overline{r}$  as a function of the cluster size *m* of  $C_m^+$  (*m*=1-13). One can see  $\overline{r}$  at the same *m* to be essentially equivalent for all the collision processes except for 3e loss, for which the value is about 20% higher than others. It is interesting to note that the overall dependence on the size m can roughly be expressed as  $\overline{r} = (11 \pm 1) \times \exp(-m/10)$  except for 3e loss events. Consequently, it is plausible to state that a fragment ion of fixed m may be produced in equivalent impact parameter collisions independently of the charge-changing condition. This leads us again to the conclusion that  $E_d$  is almost equivalent in the production of size-fixed fragment ions. Otherwise it is hard to explain such a high degree of multiple ionization as observed in single-capture collisions. However, one should be reminded from Fig. 4 that violent collisions leading to multifragmentation are very minor events, accounting for at most only a few percent of the total capture events; see the relative intensities of  $C_1^{+}$  in the top two figures where contributions from intact ions are predominant.

# **IV. DISCUSSION**

We discuss here in more detail the relative importance of the charge state r of prefragmented parent ions and internal excitation. First, the intensities of product ions measured for multiple charge-changing collisions are examined in terms of the number of pure ionization events  $n_i$ . This is based on the idea that the total energy deposition  $E_d$  is shared by ioniza-



FIG. 6. (Color) Relative intensities of fullerenelike ions  $(R_p)$ , medium-sized fragments  $(R_m)$ , and small-sized fragments  $(R_s)$  obtained for electron capture collisions. Data of 1e loss are drawn for comparison.

tion  $(n_i)$  and internal excitation  $(E_{int})$  with a certain partition rate, and therefore the value of  $n_i$  is regarded as a good measure of the amount of  $E_{int}$ . For convenience, product ions are divided into three groups of parent ions including large daughter ions  $(\sum_{m=0}^{5} C_{60-2m}^{r+})$ , medium sized ions  $(\sum_{m=4}^{15} C_m^{-+})$ and the smallest three fragments  $(\sum_{m=1}^{3} C_m^{-+})$ . The relative intensities of these product ions are denoted by  $R_p$ ,  $R_m$ , and  $R_s$ ,



FIG. 7. (Color) As in Fig. 6, obtained for electron loss collisions.

respectively, with a relationship of  $R_p+R_m+R_s=1$ . Results for electron capture and loss collisions are shown in Figs. 6 and 7, respectively, where data of 1*e* loss are plotted in both figures for comparison. These figures show clearly that multifragmentation is always dominant at  $n_i > 8$  for all the collision processes (see also Fig. 4). This implies evidently that the Coulomb repulsion force and internal excitation are both high enough to induce multifragmentation in such violent collisions. Since  $R_s \approx 1$  in this region, the following discussion is focused on the region of  $n_i < 8$ .

At  $n_i < 5$  in Fig. 6, fragmentation yields  $(R_m \text{ and } R_s)$  in 2ecapture collisions are slightly enhanced. This indicates that the instability due to Coulomb repulsion becomes more important in the fragmentation of low-charged  $C_{60}^{r+}$  ions in comparison with  $E_{int}$ . At  $n_i = 4$ , for instance, the charge states r for 1e loss, 1e capture, and 2e capture collisions are 4, 5, and 6, respectively, and corresponding total Coulomb potentials  $V_c$  (eV) are known from [2,20] to be 15.10 (r=4), 26.64 (r=5), and 41.08 (r=6). On the other hand,  $E_{int}$  is estimated from a statistical partitioning model to be 53 eV at  $n_i$ =4 as described in [16]. If we consider the sum of  $V_c$  and  $E_{int}$  as the total "instability energy," we obtain 68.1, 79.6, and 94.1 eV for r=4, 5, and 6, respectively. Here, it is worth noting the theoretical results of maximum entropy calculations made for a neutral  $C_{60}$  by Campbell *et al.* [21]. They showed that the multifragmentation starts at  $E_{int} \approx 85 \text{ eV}$  and entire destruction into small fragments occurs beyond 225 eV, whereas the molecule keeps its fullerene structure below 85 eV. Hence, in a qualitative sense, our instability energy seems to explain reasonably the enhancement of multifragmentation at  $n_i < 5$  in 2e capture collisions. It should be noted that the instability due to Coulomb explosion is the predominant fragmentation mechanism in SHCI collisions, where the main process of  $C_{60}^{r+}$  production is distant electron capture collisions accompanying small internal excitation energies [5]

In electron loss collisions (Fig. 7), there are some discrepancies in fragmentation yields at  $n_i < 8$ . For instance, at  $n_i = 5$ , the values of  $R_s$  are 0.50, 0.76, and 0.98 for  $n_l = 1, 2$ , and 3, respectively. These results are in contrast to our energy partitioning model, because both the internal energy and charge states are supposed to be constant at a fixed value of  $n_i$ , irrespective of  $n_\ell$ , in electron loss collisions. However, our experimental results indicate that the internal energy at the same  $n_i$  becomes higher with increasing  $n_\ell$ .

Similar behaviors are observed more directly in the scattering angle dependence of loss collisions. As we saw already in Fig. 2, TOF spectra are completely different for zero-degree and off-center angles. By taking account of the geometry of our detection system, zero-degree and scattering components were deduced separately as plotted in Fig. 8 with blank bars ( $\theta$ =0) and solid bars ( $\theta$ =0.8 mrad), respectively. One can see that, even at low charge states r (= $n_i$ ), small fragment ions are predominantly produced in offcenter scatterings. Consequently, these components lead to enhancement of multifragmentation shown in Fig. 7. Actually, experimental values of  $R_s$  for 2e loss collisions become identical with those for 1e loss collisions after subtracting the scattering components as depicted by arrows in Fig. 7.



FIG. 8. Comparison of intensity distributions of product ions between zero-degree components (blank bars) and scattering components (solid bars) deduced by taking account of our detector geometry.

We note here that scattering components were negligibly small for 1*e* loss and capture collisions ( $n_c=1,2$ ) but predominantly large for 3*e* loss collisions. These results indicate that the impact parameter becomes smaller as  $n_\ell$  increases in loss collisions.

Enhanced multifragmentation observed at nonzero scattering angles implies evidently that the internal excitation energy is higher than small-angle scattering when compared at the same  $n_i$  (=r). Note that the recoiling effect of a target carbon atom is also ruled out from consideration because the recoil energy is estimated to be only 6.7 eV at most ( $\theta$ =1.2 mrad) in a collision of Si+C. The present results indicate that the partition rate in close collisions may be considerably different from that in distant collisions; namely, close collisions are supposed to induce internal excitation greatly in comparison with distant collisions, whereas the degree of pure ionization is the same in both collisions.

As another explanation, we speculate that the energy partitioning rate itself has a certain statistical width around the mean value. It might, therefore, be possible to result in relatively larger  $E_{int}$  but smaller  $n_i$  than expected from the mean partition rate, leading to a higher degree of multifragmentation at small  $n_i$ 's. The present results of enhanced fragmentation in 2e and 3e loss collisions at  $n_i < 8$  seem to reflect this statistical nature of the energy partitioning process.

#### V. SUMMARY

The correlated processes of fragmentation and ionization of  $C_{60}$  induced in charge-changing collisions of 2-MeV Si<sup>2+</sup> have been investigated in detail by specifying the charge state of prefragmented  $C_{60}^{r+}$  ions and the number of free electrons ( $n_e$ ) emitted simultaneously from both target and incident particles. Some important findings are listed below. First, a huge number of secondary electrons as high as 20 are emitted in the production of small fragment ions like  $C_1^+$  and these  $n_e$  distributions exhibit nearly the same shape in all the charge-changing conditions including also 1*e* capture events. Second, multiple-electron loss collisions  $(n_l \ge 2)$  take place predominantly at smaller impact parameters, resulting in the deflection of incident ion trajectories and the corresponding TOF spectra are dominated by only the few smallest fragment ions. By contrast, electron capture of both 1*e* and 2*e* is a distant-collision process without accompanying noticeable deflection of incident ions.

The present results are examined with an energy partition model stating that the internal excitation energy  $E_{int}$  may possibly be estimated from a certain partition rate once r and  $n_e$  are known simultaneously. With this model, various fragmentation phenomena, which are significantly different in different charge-changing conditions, are successfully analyzed in terms of the degree of pure ionization  $n_i$  of a target molecule. Although the C<sub>60</sub> fragmentation is induced mainly via internal excitation, the instability due to Coulomb repulsive force is found to become also important at  $n_i \leq 4$  for electron capture collisions. As for the region of large values of  $n_i$  (>8), it is concluded that the C<sub>60</sub> fragmentation is induced via both internal excitation and the Coulomb repulsion force, irrespective of electron capture and loss collisions.

Finally, the predominant multifragmentation observed at nonzero scattering angles implies that the partition rate may be considerably different in close and distant collisions. It is, therefore, important to know the impact-parameterdependent partition rate to achieve further detailed understanding of the fragmentation mechanisms.

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