## Dynamics of the Multifragmentation of 1A GeV Gold on Carbon

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An exclusive study of the interaction of 1A GeV Au with C shows a separability into two stages: a prompt stage involving emission of most Z = 1 and some Z = 2 particles and a second stage involving the decay of an equilibrated remnant, which typically undergoes multifragmentation. The mean mass, charge, excitation energy, and the initial temperature  $T_i$  of the remnant have been determined as a function of the total charge multiplicity, m, as has the freeze-out temperature  $T_f$ . Both  $T_i$  and  $T_f$  increase linearly with m and their values at the critical point have been determined.  $T_f$  rises monotonically with excitation energy as expected for a continuous phase transition. [S0031-9007(96)00615-1]

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We have recently extracted the critical exponents  $\gamma$ ,  $\beta$ ,  $\tau$ , and  $\sigma$  from a model independent analysis of the multifragment breakup of 1*A* GeV gold ions incident on carbon [1,2]. The values of these exponents were found to be remarkably close to those for ordinary fluids [3]. These results indicate that multifragmentation (MF) may arise from a continuous phase transition, and may belong to the same universality class as ordinary fluids. They also can provide direct information about the equation of state of nuclear matter in the vicinity of its critical point [4].

The determination of the critical exponents is based on a number of assumptions, the validity of which rests in the MF dynamics. These assumptions are (1) the formation of an equilibrated remnant with charge  $Z_i$ , mass  $A_i$ , excitation energy  $E^*$ , and temperature  $T_i$ , following the emission of prompt particles in a first stage and (2) the direct proportionality between the total charged particle multiplicity, m, and temperature [5]. The target independence of MF in heavy ion interactions [6] indicates that the entrance and exit channels are decoupled, consistent with (1). Fragment energy spectra have shown that MF occurs in the breakup of a system lighter than the original nucleus with a strongly reduced Coulomb energy [7-10], suggesting that the remnant undergoes expansion and cooling prior to MF [11] with freeze-out occurring at a temperature  $T_f$ . In this Letter we explore the validity of the above assumptions in light of the MF dynamics, determine the above properties of the remnant as a function of m, and determine  $T_f(m)$ . The values of  $T_i$  and  $T_f$  at the critical multiplicity,  $m_c$ , bracket the critical temperature of the nucleus.

The experimental apparatus, described in detail in [1], consisted of a time projection chamber [12], a time-of-flight wall, and a multiple sampling ionization chamber [13]. These detectors allowed us to identify the nuclear charges and momenta of all charged projectile fragments with high efficiency. Mass identification was possible for  $Z \leq 5$ . The analysis is based on 33 000 fully reconstructed events, where the total charge for the Au system was found to be  $79 \pm 3$ .

Figure 1 shows the laboratory rapidity distributions of several light fragments. In this reverse kinematics experiment, we see that many of the fragments are moving near beam rapidity. This is the case for all Li and higher Z fragments, which show strong peaking near  $y_{\text{beam}}$ . The symmetry of these distributions is consistent with emission from an equilibrated projectile remnant. In contrast, the lighter particles, particularly *p*, *d*, and <sup>3</sup>He, show extensive tailing at low *y* from peaks near  $y_{\text{beam}}$ . These tails are due to particles which are knocked out of the Au nucleus in the first stage of the interaction.

To determine the mass-weighted average velocity of the projectile remnant we used the Li, Be, and B fragments to compute a mean velocity  $\beta_z$ . We find  $\beta_z = 0.873$ 



FIG. 1. Laboratory rapidity distributions for light fragments.  $y_{\text{beam}} = 1.35$ .

for low *m*, decreasing monotonically to 0.867 for high *m*. A linear interpolation gives  $\beta_z(m)$ . As a check on the accuracy of this velocity we have computed the masses of various heavy fragments using the experimental momenta and  $\beta_z(m)$ . The mean masses of C, N, and O fragments agree within 1% with the corresponding values obtained from a yield-weighted average of isotopically resolved fragments in high-energy *p*-Xe interactions [14]. The mean masses of fragments with Z > 10 show a similar agreement with values deduced from the EPAX parametrization of the most probable charges of products formed in high-energy reactions [15].

We can examine the behavior of the invariant cross sections as a function of  $v_{\perp}$  and  $v_{z}$ , the transverse and longitudinal velocity components in the moving system. The results for several light particles are shown in Fig. 2. The contours are nearly symmetric about  $v_z = 0$  for low  $v_{\perp}$ , indicating approximately isotropic emission in the projectile frame. However, for higher  $v_{\perp}$  the contours skew towards negative  $v_{z}$ , as expected for prompt emission. We have made a constant 30 MeV cut in the proton kinetic energies in the moving system in order to separate the two sources in an approximate way. The same energy per nucleon cuts have been made for the other Z = 1 and Z = 2 particles. The cuts determine particle yields for the two stages. The first stage multiplicity,  $m_1$ , consists of ~70% of the Z = 1particles and  $\sim 10\%$  of the Z = 2 particles. The second stage equilibrium multiplicity,  $m_2$ , is linearly proportional to  $m_1$ , and  $m_2 \simeq \frac{1}{3}m$ . By varying the position of the cuts in a manner that preserves approximate isotropy, i.e., between 25 and 35 MeV per nucleon, we estimate a 7% uncertainty in  $m_2$ .

We determine  $Z_i$  by subtracting from 79 the charges of the prompt Z = 1 and Z = 2 particles. To obtain  $A_i$  we assume the knockout of 1.7 prompt neutrons per prompt proton on the basis of a cascade simulation [16]. Figure 3(a) shows the linear variation of  $m_2$  with m. This linearity is the chief basis of our previous assertion that



FIG. 2. Contour plot of invariant cross sections for light particle emission in the indicated multiplicity bins. The linear contours correspond to equal particle number intervals. The heavy curves are cuts separating first and second stage particles.

the inclusion of prompt particles does not significantly change the values of the critical exponents [1,17].

To determine  $E^*/A_i$  we use the energy balance relation [18]

$$E^*/A_i = (1/A_i) \left[ \sum_j (E_j^{\rm kin} + Q_j) + 3nT_i/2 + Q_n \right],$$
(1)

where  $E_i^{\rm kin}$  and  $Q_i$  are the kinetic energy in the moving system and the separation energy of the *j*th charged fragment emitted in the equilibrium stage, respectively, *n* is the second stage neutron multiplicity, and  $Q_n$  is the sum of the separation energy of these neutrons. The mean neutron kinetic energy has been assumed to be  $\frac{3}{2}T$ , corresponding to volume emission [19]. The value of n,  $\sim 10$  on average, is the difference between  $A_i$  and the sum of the second stage fragment masses. In order to relate  $E^*$ and T in Eq. (1) we assume the Fermi gas model relation,  $T = (E^*/a)^{1/2}$ , where the level density parameter a is taken as A/13. This value is consistent with the observed decrease of a with increasing  $E^*$  from  $\sim A/8$  to  $\sim A/13$ , at which point a appears to level off [20]. This fixed value of a is consistent with the range of values of aobtained when  $T_i$  is extracted from  $E^*$  as a function of m; see below. Equation (1) is quadratic in  $(E^*/A)^{1/2}$  and has only one physical solution.

As a test of the procedure embodied in Eq. (1), we have applied it to simulated events generated by the ISABEL cascade code [16] coupled with a statistical MF model (SMM) [21]. The energies of SMM fragments and neutrons lead to  $E^*$  values that are in very good agreement with those given for the same events by ISABEL.



FIG. 3. Dependence on total charge multiplicity of (a) the second stage multiplicity, (b) the mean excitation energy per nucleon of the remnant, and (c) the mean mass of the remnant. Statistical errors are smaller than the size of the points where not shown.

Figure 3(b) shows the variation of the mean  $E^*/A_i$  with m. The result is subject to systematic errors associated with the source velocity determination, mass identification, first stage cut, and neutron number and energy. These effects were investigated by making reasonable changes in the velocity cut and in the assumptions about neutron emission. The combined uncertainty from these sources is  $\sim 15\%$ . The nonlinear dependence of  $E^*/A_i$  on m reflects the increase of  $E^*$  with m and the associated decrease in  $A_i$  [Fig. 3(c)]. Thus the mass of the remnant decreases from  $\sim 190$  to  $\sim 110$ .

The temperature of the unexpanded remnant with mass  $A_i$  may be obtained from  $E^*$  according to the single component Fermi gas model. It is assumed that the remnant thermalizes in the volume formerly occupied by the Au projectile. Figure 4 shows that  $T_i$  is a practically linear function of m in the region where MF occurs,  $m \sim 10-60$ , thereby confirming one of the assumptions in the determination of the critical exponents [1,2]. At the critical multiplicity,  $m_c = 26 \pm 1$  [1],  $T_i = 8.7 \pm 0.7$  MeV, where the error reflects the uncertainty of the input parameters in the Fermi gas model.

The equilibrated remnant expands and cools prior to breakup. The freeze-out temperature may be obtained from the ratio of isotope yield ratios, where each pair of isotopes differs by one neutron [22]. Pochodzalla *et al.* [23] recently used the  ${}^{3}\text{He}/{}^{4}\text{He}$  and  ${}^{6}\text{Li}/{}^{7}\text{Li}$  yield ratios to determine  $T_{f}$  for the interaction of 600A MeV Au plus



FIG. 4. Variation of  $T_i$  and  $T_f$  with m. Two different values of  $T_f$  are shown,  $T_{\text{He-DT}}$  and  $T_{\text{He-Li}}$ . The double arrow spans the values of the critical temperature.

Au. We have used this He-Li thermometer as well as a He-DT thermometer based on the  ${}^{2}\text{H}/{}^{3}\text{H}$  yield ratio to estimate  $T_{f}$ . Both  $T_{f}$  were obtained by means of equations given in Ref. [22]. Our results have not been corrected for feedin from the decay of heavier fragments, which has been estimated as a 20% correction [23]. We believe that the  $T_{\text{He-DT}}$  are more accurate than the  $T_{\text{He-Li}}$  because they do not require a correction for low-lying excited states. In addition to the statistical error, the error in  $T_{\text{He-DT}}$  is due to the uncertainty in the separation between the first and second stages, which is ~3%.

Figure 4 shows the variation of  $T_f$  with m. Both thermometers give similar results and indicate that  $T_f$  is a linear function of m. When  $m = m_c$ ,  $T_f = 5.1 \pm 0.2$  MeV. This temperature is close to the cracking temperature predicted by various statistical models [21,24] and also agrees with a determination based on a thermal liquid drop model description of the isotopic yields of fragments from p-Xe collisions [8]. Figure 4 shows that  $T_{\text{He-DT}}$  is substantially lower than  $T_i$ , where their ratio decreases from 0.77 for low m to 0.51 for high m. Evidently, the remnant cools down on its approach to freeze-out due to both expansion and evaporation, with a higher degree of cooling occurring at high  $E^*$ . The values of  $T_f(m_c)$  and  $T_i(m_c)$  bracket the critical temperature of the nucleus, which thus has a value between ~5 and 9 MeV.

Figure 5 shows the monotonic increase of  $T_{\text{He-DT}}$  with  $E^*/A_i$ . This variation is consistent with that expected for a liquid-gas phase transition in the critical region. Pochodzalla *et al.* [23] have presented a similar curve for the interaction of 600A MeV Au with Au which displays a roughly constant value of  $T_{\text{He-Li}}$  between 2 and 10 MeV/nucleon followed by a sharp rise at higher  $E^*/A_i$ . At the highest  $E^*/A_i$ , where the difference in the



FIG. 5. Dependence of  $T_{\text{He-DT}}$  on  $E^*/A_i$ . Statistical errors are smaller than the size of the points where not shown.

shape of the two curves is largest, their remnant masses are only half as large as ours. Consequently, the two systems may differ in dynamics. A detailed comparison is presented elsewhere [25].

In summary, we have separated the first stage particles from those emitted by an equilibrated remnant in the interaction of 1A GeV <sup>197</sup>Au with carbon. Most of the Z = 1particles as well as a small fraction of the Z = 2 particles are emitted prior to equilibrium. Heavier fragments are emitted by the equilibrated remnant, the charge and mass of which have been determined as a function of m. The second stage multiplicity is a linear function of m, where  $m_2 \simeq \frac{1}{3}m$ . The mean excitation energy of the remnant has been determined and the temperature  $T_i$  derived from it is a linear function of m. These results validate the assumptions made in our previous determination of the critical exponents. The freeze-out temperature  $T_f$  has been determined using two different isotope ratio thermometers and is also a linear function of m. The values of  $T_i(m_c)$ and  $T_f(m_c)$  bracket the critical temperature of the nucleus. The monotonic increase of  $T_{\text{He-DT}}$  with  $E^*/A_i$  supports the view that MF may indeed result from a continuous phase transition near the critical point.

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- M. L. Gilkes *et al.*, Phys. Rev. Lett. **73**, 1590 (1994).
- [2] J.B. Elliott et al., Phys. Lett. B (to be published).
- [3] H.E. Stanley, Introduction to Phase Transitions and Critical Phenomena (Oxford University Press, Oxford, 1971).
- [4] K. C. Chase and A. Z. Mekjian, Phys. Rev. Lett. 75, 4732 (1995).
- [5] X. Campi, J. Phys. A 19, L917 (1986); Phys. Lett. B 208, 351 (1988); in *Proceedings of the International School* of *Physics "Enrico Fermi"* (North-Holland, Amsterdam, 1990), p. 237.
- [6] J. Hubele et al., Phys. Rev. C 46, R1577 (1992).
- [7] A. M. Poskanzer et al., Phys. Rev. C 3, 882 (1971).
- [8] A.S. Hirsch et al., Phys. Rev. C 29, 508 (1984).
- [9] N.T. Porile et al., Phys. Rev. C 39, 1914 (1989).
- [10] K. Kwiatkowski *et al.*, Phys. Rev. Lett. **74**, 3756 (1995).
- [11] W. A. Friedman, Phys. Rev. C 42, 667 (1990).
- [12] G. Rai et al., IEEE Trans. Nucl. Sci. 37, 56 (1990).
- [13] W. Christie *et al.*, Nucl. Instrum. Methods Phys. Res., Sect. A 255, 46 (1987).
- [14] R. W. Minich et al., Phys. Lett. B 118, 458 (1982).
- [15] K. Sümmerer et al., Phys. Rev. C 42, 2546 (1990).
- [16] Y. Yariv and Z. Fraenkel, Phys. Rev. C 20, 2227 (1979).
- [17] M. Gilkes et al., Phys. Rev. Lett. 75, 768 (1995).
- [18] D. Cussol et al., Nucl. Phys. A561, 298 (1993).
- [19] R. Wada et al., Phys. Rev. C 39, 497 (1989).
- [20] K. Hagel et al., Nucl. Phys. A486, 429 (1988).
- [21] J. P. Bondorf et al., Phys. Rep. 257, 133 (1995).
- [22] S. Albergo *et al.*, Nuovo Cimento So. Ital. Fis. **89A**, 1 (1985).
- [23] J. Pochodzalla et al., Phys. Rev. Lett. 75, 1040 (1995).
- [24] D. H. E. Gross, Rep. Prog. Phys. 53, 605 (1990).
- [25] M. Tincknell *et al.*, Proceedings of the 12th Winter Workshops on Nuclear Dynamics (to be published).