LA-UR-12-01364

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Title:	CEM03.03 User Manual
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Intended for:	RSICC and the MCNP6 Code Package



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CEM03.03 User Manual

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Abstract

The Fortran 77 code CEM03.03 is an extended and improved version of the earlier codes CEM03.01 and CEM2k+GEM2, which are based in turn on their predecessor codes CEM2k, CEM97, CEM95, CEM92M, CEM92, and MARIAG, which implement versions of the Cascade-Exciton Model (CEM) of nuclear reactions. CEM03.03 calculates total reaction and fission cross-sections, nuclear fissilities, excitation functions, nuclide distributions (yields) of all produced isotopes separately as well as their A- and Z-distributions, energy and angular spectra, double-differential cross-sections, mean multiplicities, i.e. the number of ejectiles per inelastic interaction of the projectile with the target, ejectile yields and their mean energies for $n, p, d, t, {}^{3}He, {}^{4}He, \pi^{+}, \pi^{-}, and \pi^{0}$. In addition, CEM03.03 provides in its output separately the yields of Forward (F) and Backward (B) produced isotopes, their mean kinetic energies, A- and Z-distributions of the mean emission angle, their parallel velocities, and the F/B ratio of all products in the laboratory system, distributions of the mean angle between two fission fragments, of neutron multiplicity, of the excitation energy, of momentum and angular momentum, and of mass and charge numbers of residual nuclei after the INC and preequilibrium stages of reactions, as well as for fissioning nuclei before and after fission.

CEM03.03 calculates reactions induced by nucleons, pions, bremsstrahlung and monochromatic photons on not too light targets at incident energies from ~10 MeV (~ 30 MeV, in the case of $\gamma + A$) up to several GeV. This Manual describes the basic assumptions of the improved CEM as realized in the code CEM03.03, essential technical details of the code such as the description of the input and output files, and provides the user with necessary information for practical use of and for possible modification of the CEM03.03 output, if required.

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1. Introduction

The Cascade-Exciton Model (CEM) of nuclear reactions was proposed 32 years ago at the Laboratory of Theoretical Physics, Joint Institute for Nuclear Research, Dubna, USSR by Gudima, Mashnik, and Toneev [1, 2]. Is is based on the Dubna IntraNuclear Cascade (INC) [3, 4] and the Modified Exciton Model (MEM) [5, 6]. It was extended to consider photonuclear reactions [7] and to describe fission cross sections using different options for nuclear masses, fission barriers, and level densities [8] and its 1995 version, CEM95, was released to the public via NEA/OECD, Paris as the code IAEA1247, and via the Radiation Safety Information Computational Center (RSICC) at Oak Ridge, USA, as the RSICC code package PSR-357 [9].

The International Code Comparison for Intermediate Energy Nuclear Data [10, 11] organized during 1993–1994 at NEA/OECD in Paris to address the subject of codes and models used to calculate nuclear reactions from 20 to 1600 MeV showed that CEM95 had one of the best predictive powers to describe nucleon-induced reactions at energies above about 150 MeV when compared to other models and codes available at that time.

CEM95 and/or its predecessors and its successors CEM97 [12, 13], CEM2k [14],

CEM2k+GEM2 [15]–[17], CEM03 [18, 19], CEM03.01 [20, 21], CEM03.02 [22, 23] and the latest version, CEM03.03 [23, 24] are used as stand-alone codes to study different nuclear reactions for applications and fundamental nuclear physics (see, e.g., [25]–[36] and references therein). Parts of different versions of the CEM code are used in many other stand-alone codes, like **PICA95** [37], **PICA3** [38], **CASCADO** [39], **CAMO** [40], **MCFX** [41], **ECM** [42], and **NUCLEUS** [43]. CEM95 and some of its predecessor or successor versions are incorporated wholly, or in part in different transport codes used in many applications, like **CAS-CADE** [44], **GEANT4** [45, 46], **SHIELD** [47], **RTS&T** [48], **SONET** [49], **CALOR** [50], **HETC-3STEP** [51], **CASCADE/INPE** [52], **HADRON** [53], and others. The latest version, CEM03.03 [23, 24], was recently incorporated as event generator in **MCNP6** [54], **MC-NPX2.7.0** [55], **MARS15** [56], and **MRED** [57].

All CEM code versions still have some problems to be solved, just as all similar models do. Following an increased interest in intermediate-energy nuclear data in relation to such projects as the Accelerator Transmutation of nuclear Wastes (ATW), the Accelerator Production of Tritium (APT), the Spallation Neutron Source (SNS), the Rare Isotope Accelerator (RIA), Proton Radiography (PRAD) as a radiographic probe for the Advanced Hydro-test Facility, and others, for several years the US Department of Energy has supported our work on the development of an improved version of the CEM which has led to the code CEM03.03 described here.

2. A Brief Survey of CEM03.03 Physics

The basic version of the modern CEM code is the so-called "03.01" version, namely CEM03.01 [20, 21]. The CEM03.01 code calculates nuclear reactions induced by nucleons, pions, and photons. It assumes that the reactions occur generally in three stages. The first stage is the IntraNuclear Cascade (INC), in which primary particles can be re-scattered and produce secondary particles several times prior to absorption by, or escape from the nucleus. When the cascade stage of a reaction is completed, CEM03.01 uses the coalescence model to "create" high-energy d, t, ³He, and ⁴He by final-state interactions among emitted cascade nucleons, already outside of the target. The emission of the cascade particles determines the particle-hole

configuration, Z, A, and the excitation energy that is the starting point for the second, preequilibrium stage of the reaction. The subsequent relaxation of the nuclear excitation is treated in terms of an improved version of the modified exciton model of preequilibrium decay followed by the equilibrium evaporation/fission stage of the reaction. Generally, all four components may contribute to experimentally measured particle spectra and other distributions. But if the residual nuclei after the INC have atomic numbers with $A \leq 12$, CEM03.01 uses the Fermi breakup model to calculate their further disintegration instead of using the preequilibrium and evaporation models. Fermi breakup is much faster to calculate and gives results very similar to the continuation of the more detailed models for much lighter nuclei.

The main difference of the following, so-called "03.02" version of CEM from the basic "03.01" version is that the earlier code only uses the Fermi breakup model to calculate the disintegration of light nuclei, in lieu of the preequilibrium and evaporation models, when the excited nuclei after the INC have a mass number $A \leq 12$, but not when such nuclei are produced after the preequilibrium, evaporation, or fission stages. This problem was solved in the 03.02 versions of CEM and LAQGSM, where the Fermi breakup model is used at all stages of a reaction, when producing an excited nucleus with $A \leq 12$. A schematic outline of a nuclear reaction calculation by CEM03.02 (and by CEM03.03) is shown in Fig. 1.



Figure 1: Flow chart of nuclear-reaction calculations by CEM03.03.

In addition, the routines that describe the Fermi breakup model in the basic 03.01 version of our codes were written more than twenty years ago in the group of Prof. Barashenkov at JINR, Dubna, Russia, and are not quite perfect, though they are quite reliable and are still used without any changes in some current transport codes. First, these routines allow in rare cases production of some light unstable fragments like ⁵He, ⁵Li, ⁸Be, ⁹B, etc., as a result of a

breakup of some light excited nuclei. Second, these routines allowed in some very rare cases the production of "neutron stars" (or "proton stars"), i.e., residual "nuclei" produced via Fermi breakup that consist of only neutrons (or only protons). Lastly, in some very rare cases, these routines could even crash the code, due to cases of 0/0. All these problems of the Fermi breakup model routines are addressed and solved in the 03.02 version of our codes [22, 23]. Several bugs are also fixed in 03.02 in comparison with its predecessor. On the whole, the 03.02 versions describe nuclear reactions on intermediate and light nuclei, and production of fragments heavier than ⁴He from heavy targets better than their predecessors (see Fig. 2 and Ref. [22]), rarely produce any unstable unphysical final products, and are free of the fixed bugs.



Figure 2: Mass distribution of the product yields from the reaction 730 MeV p + 27 Al calculated with CEM03.01 without considering the Fermi-break-up mode during the preequilibrium and evaporation stages of reactions (solid circles connected with a solid line) and with the extended version of the code referred to here and below as CEM03.02, that considers the Fermi break-up mode during the preequilibrium and evaporation stages of reactions (open circles connected with a dashed line) compared with experimental data available at a nearby energy of 800 MeV from the T-16 Lib compilation [58] (open red squares).

However, even after solving these problems and after implementing the improved Fermi breakup model into CEM03.02 [22], in some very rare cases, our code still could produce some unstable products via very asymmetric fission, when the excitation energy of such fragments was below 3 MeV and they were not checked and not disintegrated with the Fermi breakup model (see more details in [59]). This problem was addressed in the "03.03" versions of our

CEM (and LAQGSM) codes, where we force such unstable products to disintegrate via Fermi breakup independently of their excitation energy. Several more bugs were fixed on the "03.03" version as well. We emphasize that the occurrence of these problems even in the "03.01" version is quite rare, allowing stand-alone calculations of many nuclear reactions to proceed without problems, but are unacceptable when CEM (and LAQGSM) are used as event generators inside transport codes doing large-scale simulations.

To maintain historical clarity, we note here that the "standard 03.03" version of CEM produced as described above (see more details in [59]) is used at present only in MARS15 [56]. In MCNP6 [54] and in the latest versions of MCNPX, 2.7.A, 2.7.B, 2.7.C, 2.7.D, 2.7.E, and 2.7.0 (see [55] and references therein), as well as in the Monte Carlo Radiative Energy Deposition (MRED) code developed at Vanderbilt University for single event effect studies [57], we use now a slight modification of CEM03.02 which eliminates the rare fission-fragment problem just discussed by disallowing all fission into fragments with A < 13, a slightly simpler remedy of the difficulty. Therefore, there is no need to use the "standard 03.03" version to address this problem.

Until very recently, we have called this latest version of CEM included in MCNP6/X and in MRED as "CEM03.02" (to distinguish it from the "standard 03.03" version used in MARS15) though its physics is essentially identical. We used it to participate in the recent Benchmark of Spallation Models organized at the International Atomic Energy Agency during 2008-2009 [60], and it is referred to there as "CEM03.02". As one can see from the numerous and various results presented at the Web site of that Benchmark [60], the results by "CEM03.02" are practically the same as those by "CEM03.03", just as expected, differing only at the level of statistical fluctuations. The situation of having different names of the latest version of CEM in MCNP6/X and MRED as "CEM03.02" and as "CEM03.03" in MARS15 was confusing for people outside our Group, as kindly pointed out to us by one of the referees of our recent paper on Validation and Verification of MCNP6 [36]. To address this, we decided to call in Ref. [36] and in all our following publications the latest version of CEM we use at LANL (and in MRED at Vanderbilt University) "CEM03.03".

CEM03.03 contains one more important fix relative to its 03.02 and 03.01 precursors. Recently we discovered an error in the calculation of the fission level-density parameter of a few fissioning preactified nuclei a_f (or more exactly, the ratio of level density parameters for the fission and neutron-evaporation channels, a_f/a_n ; see details in Section 2.5.5 below and in Ref. [61]) for reactions on ¹⁸¹Ta and nearby Z = 72 or 73 nuclei in CEM03.01 [21], which we produced at the beginning of 2005. That error was not present in versions of CEM before 2005. but was introduced by accident in 2005 in CEM03.01. It migrated later also to the "03.02" and "03.03" versions of CEM, as well as to MCNP6/X, MRED, and MARS transport codes using those versions of CEM as event generators. Unfortunately, that error seriously affected the calculated fission cross section of ¹⁸¹Ta and of nearby nuclei, as well as the yield of fission fragments, and, to a lesser degree, also the spectra of secondary particles from such reactions. We discovered that error in the middle of 2011 and have fixed it in CEM03.03 (see details in Ref. [61] and Fig. 3 below). It was fixed also in the most recent versions of the transport codes MCNP6, MRED, and MARS15. However, users of versions of CEM03.01/.02/.03 and of transport codes MCNP6/X, MRED, and MARS15 produced after 2005 and before the second half of 2011 will still have it and need to update their versions of the codes to eliminate that error. For brevity's sake, when we need to refer to all 03.03, 03.02, and 03.01 versions of CEM or of LAQGSM, we use a generalized notation "03.xx".

Having discovered the 2005 error and knowing how it affects the CEM results, we can now

understand why in the recent works by Titarenko et al. [34, 35] it was found that CEM03.02 (which has practically the same physics as the version CEM03.03 described here) provided a poor agreement with the measured yields of the nuclides produced in proton interactions with ¹⁸¹Ta and the nearby target nuclei for energies above 250 MeV.



Figure 3: Comparison of Prokofiev systematics (open circles) and of several measured protoninduced fission cross section data for ¹⁸¹Ta (symbols; all references may be found in Ref. [61]) with our old CEM03.xx calculations (black dashed lines) before we found the error in the middle of 2011, and with the updated CEM03.03 (red solid line), including the fix [61].

We have collaborated with the ITEP Group of Prof. Titarenko for more than a decade and have analyzed with different versions of our CEM and LAQGSM codes practically all of the proton-induced activation data measured by this group: some 14,621 product yields, from proton reactions on 24 targets, from ^{nat}Cr to ^{nat}U, at incident energies from 40 MeV to 2.6 GeV. Generally, both the CEM and LAQGSM codes describe quite well the data measured by Titarenko et al. This group defines a mean deviation factor $\langle F \rangle$, which involves an average of the ratio of the experimental to the theoretical cross sections over all measured nuclide products for a particular reaction energy and target. For most of these reactions, our codes have a value of $\langle F \rangle$ near or less than 2, nearly the best performance in comparison with about a dozen of other popular codes compared to the ITEP data (see, e.g., Figs. 4–8 and Figs. 9–11 in [35] and, especially, Fig. 9 in Ref. [32]). However, this was only "usually," because in the case of ¹⁸¹Ta, this factor $\langle F \rangle$ for CEM03.02 presented in Tab. 4 of Ref. [34] was of 1.61, 1.85, 2.21, 1.59, 1.42, 2.86, 4.17, 4.19, 4.30, 3.43, and 3.33 at energies of the bombarding protons of 40, 70, 100, 150, 250, 400, 600, 800, 1200, 1600, and 2600 MeV, respectively. The values of $\langle F \rangle$ at proton energies above 250 MeV, are significantly higher than 2, and CEM03.02 does not provide, for these cases, the best agreement with the data in comparison with the other models tested in Ref. [34], a result that was unexpected and not understood when this paper was published. Similarly, from Fig. 14 of Ref. [35], we see that the mean deviation factor between results by CEM03.02 and measured data is usually within a factor of two, except for the Ta and nearby target nuclei at energies above ~ 400 MeV, where the agreement was found to be worse, as shown by that "red finger" in Fig. 14 of this paper. Now, we understand that this behavior was due to the 2005 error in the values of a_f in CEM; as we see from Fig. 3; all nuclides arising from fission reactions will suffer from an under-prediction of the same order as the fission cross section. After fixing that problem, CEM03.03 calculates fission cross sections (and fission fragment yields) in a good agreement with available experimental data for reactions induced by nucleons, pions, and photons on both subactinide and actinide nuclei (from ¹⁶⁵Ho to ²³⁹Pu; see details in Ref. [61]).

Finally, we replaced the random-number generator used in our model: In CEM03.01, CEM03.02, and CEM03.03 through most of 2011, we used an algorithm for the uniform randomnumber generator RNDM adopted from MARS15 [56], originally published in "Toward a Universal Random Number Generator" by George Marsaglia and Arif Zaman, Florida State University Report FSU-SCRI-87-50 (1987). It was later modified by F. James and published in "A Review of Pseudo-random Number Generators". It is considered as one of the better random number generators available in the literature and is used by various modern Monte-Carlo codes. However, recently we have found that the newer MCNP5/6 random-number generator by Forrest Brown and Yasunobu Nagaya [62] is, because of a much longer period, extensive testing, and a faster execution speed in many implementations, better for our purposes. The MCNP6 version is entirely conformant to the FORTRAN-90 standard, and optionally preserves the exact random sequence of previous MCNP versions and is completely portable. In addition, new skip-ahead algorithms have been implemented to efficiently initialize the generator for new histories, a capability that greatly simplifies parallel algorithms. Finally, it has been subjected to sets of rigorous and extensive tests to verify that it produces a sufficiently random sequence (see, e.g. [63] and references therein). Because CEM03.03 remains in Fortran 77 format, we have adapted the MCNP6 generator to fixed source format, while preserving its basic functionality, but not the parallelization features. We find that with this new generator, CEM03.03 runs about 20% faster than with the older RNDM.

In the following we highlight the main assumptions of the models contained in CEM03.xx.

2.1. The INC

The intranuclear cascade model in CEM03.xx is based on the standard (non-time-dependent) version of the Dubna cascade model [3, 4]. All the cascade calculations are carried out in a three-dimensional geometry. The nuclear matter density $\rho(r)$ is described by a Fermi distribution with two parameters taken from the analysis of electron-nucleus scattering, namely

$$\rho(r) = \rho_p(r) + \rho_n(r) = \rho_0 \{1 + exp[(r-c)/a]\}, \qquad (1)$$

where $c = 1.07A^{1/3}$ fm, A is the mass number of the target, and a = 0.545 fm. For simplicity, the target nucleus is divided by concentric spheres into seven zones in which the nuclear density is considered to be constant. The energy spectrum of the target nucleons is estimated in the perfect Fermi-gas approximation with the local Fermi energy $T_F(r) = \hbar^2 [3\pi^2 \rho(r)]^{2/3}/(2m_N)$, where m_N is the nucleon mass. The influence of intranuclear nucleons on the incoming projectile

is taken into account by adding to its laboratory kinetic energy an effective real potential V, as well as by considering the Pauli principle which forbids a number of intranuclear collisions and effectively increases the mean free path of cascade particles inside the target. For incident nucleons $V \equiv V_N(r) = T_F(r) + \epsilon$, where $T_F(r)$ is the corresponding Fermi energy and ϵ is the binding energy of the nucleons. For pions, CEM03.xx uses a square-well nuclear potential with the depth $V_{\pi} \simeq 25$ MeV, independently of the nucleus and pion energy, as was done in the initial Dubna INC [3, 4].

The interaction of the incident particle with the nucleus is approximated as a series of successive quasifree collisions of the fast cascade particles $(N, \pi, \text{ or } \gamma)$ with intranuclear nucleons:

$$NN \to NN, \qquad NN \to \pi NN, \qquad NN \to \pi_1, \cdots, \pi_i NN$$
, (2)

$$\pi N \to \pi N, \qquad \pi N \to \pi_1, \cdots, \pi_i N \qquad (i \ge 2) .$$
 (3)

In the case of pions, besides the elementary processes (3), CEM03.03 also takes into account pion absorption on nucleon pairs

$$\pi NN \to NN.$$
 (4)

The momenta of the two nucleons participating in the absorption are chosen randomly from the Fermi distribution, and the pion energy is distributed equally between these nucleons in the center-of-mass system of the three particles participating in the absorption. The direction of motion of the resultant nucleons in this system is taken as isotropically distributed in space. The effective cross section for absorption is related (but not equal) to the experimental cross sections for pion absorption by deuterons.

In the case of photonuclear reactions, CEM03.xx follows [19] the ideas of the photonuclear version of the Dubna INC proposed initially 43 years ago by Gudima, Iljinov, and Toneev [64] to describe photonuclear reactions at energies above the Giant Dipole Resonance (GDR) region [65]. [At photon energies $T_{\gamma} = 10-40$ MeV, the de Broglie wavelength $\lambda/2\pi$ is of the order of 20–5 fm, greater than the average inter-nucleonic distance in the nucleus; the photons interact with the nuclear dipole resonance as a whole, thus the INC is not applicable.] Below the pion-production threshold, the Dubna INC considers absorption of photons on only "quasi-deuteron" pairs according to the Levinger model [66]:

$$\sigma_{\gamma A} = L \frac{Z(A-Z)}{A} \sigma_{\gamma d} , \qquad (5)$$

where A and Z are the mass and charge numbers of the nucleus, $L \approx 10$, and $\sigma_{\gamma d}$ is the total photoabsorption cross section on deuterons as defined from experimental data.

At photon energies above the pion-production threshold, the Dubna INC considers production of one or two pions; the specific mode of the reaction is chosen by the Monte-Carlo method according to the partial cross sections (defined from available experimental data):

$$\gamma + p \rightarrow p + \pi^0$$
, (6)

$$\rightarrow n + \pi^+$$
, (7)

$$\rightarrow p + \pi^+ + \pi^- , \qquad (8)$$

$$\rightarrow p + \pi^0 + \pi^0 , \qquad (9)$$

$$\rightarrow n + \pi^+ + \pi^0 . \tag{10}$$

The cross sections of $\gamma + n$ interactions are derived from consideration of isotopic invariance, i.e. it is assumed that $\sigma(\gamma + n) = \sigma(\gamma + p)$. The Compton effect on intranuclear nucleons is

neglected, as its cross section is less than $\approx 2\%$ of other reaction modes (see, e.g. Fig. 6.13 in Ref. [67]). The Dubna INC does not consider processes involving production of three and more pions; this limits the model's applicability to photon energies $T_{\gamma} \leq 1.5$ GeV [for T_{γ} higher than the threshold for three-pion production, the sum of the cross sections (8)–(10) is assumed to be equal to the difference between the total inelastic $\gamma + p$ cross section and the sum of the cross sections of the two-body reactions (6)–(7)].

The integral cross sections for the free NN, πN , and γN interactions (2)–(10) are approximated in the Dubna INC model [3] used in CEM95 and its predecessors using a special algorithm of interpolation/extrapolation through a number of picked points, mapping as well as possible the experimental data. This was done very accurately by the group of Prof. Barashenkov using all experimental data available at that time, about 43 years ago. Currently the experimental data on cross sections is much more complete than at that time; therefore we have revised the approximations of all the integral elementary cross sections used in CEM95 and its predecessors. We started by collecting all published experimental data from all available sources. Then we developed an improved, as compared with the standard Dubna INC [3], algorithm for approximation of cross sections and developed simple and fast approximations for elementary cross sections which fit very well presently available experimental data not only to 5 GeV, the upper recommended energy for the present version of the CEM, but up to 50-100 GeV and higher, depending on availability of data (see details in [12, 19]). So far, we have in CEM03.xx new approximations for 34 different types of elementary cross sections induced by nucleons, pions, and gammas. Integral cross sections for other types of interactions taken into account in CEM03.xx are calculated from isospin considerations using the former as input.

The kinematics of two-body elementary interactions and absorption of photons and pions by a pair of nucleons is completely defined by a given direction of emission of one of the secondary particles. The cosine of the angle of emission of secondary particles in the c.m. system is calculated by the Dubna INC [3] as a function of a random number ξ , distributed uniformly in the interval [0,1] as

$$\cos\theta = 2\xi^{1/2} \left[\sum_{n=0}^{N} a_n \xi^n + (1 - \sum_{n=0}^{N} a_n) \xi^{N+1} \right] - 1 , \qquad (11)$$

where N = M = 3,

$$a_n = \sum_{k=0}^{M} a_{nk} T_i^k . (12)$$

The coefficients a_{nk} were fitted to the then available experimental data at a number of incident kinetic energies T_i , then interpolated and extrapolated to other energies (see details in [3, 64, 65] and references therein). The distribution of secondary particles over the azimuthal angle φ is assumed isotropic. For elementary interactions with more than two particles in the final state, the Dubna INC uses the statistical model to simulate the angles and energies of products (see details in [3]).

For the improved version of the INC in CEM03.xx, we use currently available experimental data and recently published systematics proposed by other authors and have developed new approximations for angular and energy distributions of particles produced in nucleon-nucleon and photon-proton interactions. So, for pp, np, and nn interactions at energies up to 2 GeV, we did not have to develop our own approximations analogous to the ones described by Eqs. (11) and (12), since reliable systematics have been developed recently by Cugnon et al. for

the Liege INC [68], then improved still further by Duarte for the BRIC code [69]; we simply incorporate into CEM03.xx the systematics by Duarte [69]. Similarly, for γN interactions, we take advantage of the event generators for γp and γn reactions from the Moscow INC [70] kindly sent us by Dr. Igor Pshenichnov. In CEM03.xx, we use part of a data file with smooth approximations through presently available experimental data, developed for the Moscow INC [70] and have ourselves developed a simple and fast algorithm to simulate unambiguously $d\sigma/d\Omega$ and to choose the corresponding value of Θ for any E_{γ} , using a single random number ξ uniformly distributed in the interval [0,1] (see details in [19]).

The analysis of experimental data has shown that the channel (8) of two-pion photoproduction proceeds mainly through the decay of the Δ^{++} isobar listed in the last Review of Particle Physics by the Particle Data Group as having the mass M = 1232 MeV

$$\begin{array}{l} \gamma + p \quad \to \quad \Delta^{++} + \pi^{-} \\ \Delta^{++} \quad \to \quad p + \pi^{+} \\ \end{array}, \tag{13}$$

whereas the production cross section of other isobar components $(\frac{3}{2}, \frac{3}{2})$ are small and can be neglected. The Dubna INC uses the Lindenbaum-Sternheimer resonance model [71] to simulate the reaction (13). In this model, the mass of the isobar M is determined from the distribution

$$\frac{\mathrm{d}W}{\mathrm{d}M} \sim F(E, M)\sigma(M) , \qquad (14)$$

where E is the total energy of the system, F is the two-body phase space of the isobar and π^- meson, and σ is the isobar production cross section which is assumed to be equal to the cross section for elastic $\pi^+ p$ scattering.

The c.m. emission angle of the isobar is approximated using Eqs. (11) and (12) with the coefficients a_{nk} listed in Tab. 3 of Ref. [65]; isotropy of the decay of the isobar in its c.m. system is assumed.

In order to calculate the kinematics of the non-resonant part of the reaction (8) and the two remaining three-body channels (9) and (10), the Dubna INC uses the statistical model. The total energies of the two particles (pions) in the c.m. system are determined from the distribution

$$\frac{\mathrm{d}W}{\mathrm{d}E_{\pi_1}\mathrm{d}E_{\pi_2}} \sim (E - E_{\pi_1} - E_{\pi_2})E_{\pi_1}E_{\pi_2}/E , \qquad (15)$$

and that of the third particle (nucleon, N) from conservation of energy. The actual simulation of such reactions is done as follows: Using a random number ξ , we simulate in the beginning the energy of the first pion using

$$E_{\pi_1} = m_{\pi_1} + \xi (E_{\pi_1}^{max} - m_{\pi_1}),$$

where

$$E_{\pi_1}^{max} = [E^2 + m_{\pi_1}^2 - (m_{\pi_2} + m_N)^2]/2E.$$

Then, we simulate the energy of the second pion E_{π_2} according to Eq. (15) using the Monte-Carlo rejection method. The energy of the nucleon is calculated as $E_N = E - E_{\pi_1} - E_{\pi_2}$, following which we check that the "triangle law" for momenta

$$|p_{\pi_1} - p_{\pi_2}| \le p_N \le |p_{\pi_1} + p_{\pi_2}|$$

is fulfilled, otherwise this sampling is rejected and the procedure is repeated. The angles Θ and φ of the pions are sampled assuming an isotropic distribution of particles in the c.m. system,

$$\cos \Theta_{\pi_1} = 2\xi_1 - 1, \qquad \cos \Theta_{\pi_2} = 2\xi_2 - 1, \qquad \varphi_{\pi_1} = 2\pi\xi_3, \qquad \varphi_{\pi_2} = 2\pi\xi_4,$$

and the angles of the nucleon are defined from momentum conservation, $\vec{p}_N = -(\vec{p}_{\pi_1} + \vec{p}_{\pi_2})$. More details on our new approximations for differential elementary cross sections may be found in [18, 19].

The Pauli exclusion principle at the cascade stage of the reaction is handled by assuming that nucleons of the target occupy all the energy levels up to the Fermi energy. Each simulated elastic or inelastic interaction of the projectile (or of a cascade particle) with a nucleon of the target is considered forbidden if the "secondary" nucleons have energies smaller than the Fermi energy. If they do, the trajectory of the particle is traced further from the forbidden point and a new interaction point, a new partner and a new interaction mode are simulated for the traced particle, etc., until the Pauli principle is satisfied or the particle leaves the nucleus.

In this version of the INC, the kinetic energy of the cascade particles is increased or decreased as they move from one of the seven potential regions (zones) to another, but their directions remain unchanged. That is, in our calculations, refraction or reflection of cascade nucleons at potential boundaries is neglected. CEM03.xx allows us to take into account refractions and reflections of cascade nucleons at potential boundaries; for this, one needs to change the value of the parameter **irefrac** from 0 to 1 in the subroutine **initial**. But this option provides somewhat worse overall agreement of calculations with some experimental data, therefore the option of no refractions/reflections was chosen as the default in CEM03.xx.

This INC does not take into account the so-called "trawling" effect [3]. That is, in the beginning of the simulation of each event, the nuclear density distributions for the protons and neutrons of the target are calculated according to Eq. (1) and a subsequent decrease of the nuclear density with the emission of cascade particles is not taken into account. Our detailed analysis of different characteristics of nucleon- and pion-induced reactions for targets from C to Am has shown that this effect may be neglected at incident energies below about 5 GeV in the case of heavy targets like actinides and below about 1 GeV for light targets like carbon. At higher incident energies the progressive decrease of nuclear density with the development of the intranuclear cascade has a strong influence on the calculated characteristics and this effect has to be taken into account [3]. Therefore, in transport codes that use as event generators both CEM03.xx and our high-energy code LAQGSM03.xx [23], we recommend simulating nuclear reactions with CEM03.xx at incident energies up to about 1 GeV for light nuclei like C and up to about 5 GeV for actinide nuclei, and to switch to simulations using LAQGSM03.03, which considers the "trawling" effect (target nucleon depletion during the cascade), at higher energies of transported particles.

An important ingredient of the CEM is the criterion for transition from the intranuclear cascade to the preequilibrium model. In conventional cascade-evaporation models (like ISABEL and Bertini's INC used in MCNPX [55], fast particles are traced down to some minimal energy, the cutoff energy T_{cut} (or one compares the duration of the cascade stage of a reaction with a cutoff time, in "time-like" INC models, such as the Liege INC [68]). This cutoff is usually less than $\simeq 10$ MeV above the Fermi energy, below which particles are considered to be absorbed by the nucleus. The CEM uses a different criterion to decide when a primary particle is considered to have left the cascade.

An effective local optical absorptive potential $W_{opt. mod.}(r)$ is defined from the local interaction cross section of the particle, including Pauli-blocking effects. This imaginary potential is compared to one defined by a phenomenological global optical model $W_{opt. exp.}(r)$. We characterize the degree of similarity or difference of these imaginary potentials by the parameter

$$\mathcal{P} = |(W_{opt.\ mod.} - W_{opt.\ exp.})/W_{opt.\ exp.}|.$$
(16)

When \mathcal{P} increases above an empirically chosen value, the particle leaves the cascade, and is then considered to be an exciton. From a physical point of view, such a smooth transition from the cascade stage of the reaction seems to be more attractive than the "sharp cut-off" method. In addition, as was shown in Ref. [2], this improves the agreement between the calculated and experimental spectra of secondary nucleons, especially at low incident energies and backward angles of the detected nucleons (see e.g., Figs. 3 and 11 of Ref. [2]). More details about this can be found in [2, 14, 72].

CEM03.xx uses a fixed value $\mathcal{P} = 0.3$ (at incident energies below 100 MeV), just as all its predecessors did. With this value, we find that the cascade stage of the CEM is generally shorter than that in other cascade models. This fact leads to an overestimation of preequilibrium particle emission at incident energies above about 150 MeV, and correspondingly to an underestimation of neutron production from such reactions, as was established in Ref. [14]. In Ref. [14], this problem was solved temporarily in a very rough way by using the transition from the INC to the preequilibrium stage according to Eq. (16) when the incident energy of the projectile is below 150 MeV, and by using the "sharp cut-off" method with a cutoff energy $T_{cut} = 1$ MeV for higher incident energies. This "ad hoc" rough criterion solved the problem of underestimating neutron production at high energies, providing meanwhile a reasonably good description of reactions below 150 MeV. But it provides an unphysical discontinuity in some observables calculated by MCNPX using CEM2k [14] as an event generator, observed but not understood by Broeders and Konobeev [73]. In CEM03.xx, this problem is solved by using a smooth transition from the first criterion to the second one in the energy interval from 75 to 225 MeV, so that no discontinuities are produced in results from CEM03.xx.

Beside the changes to the Dubna INC mentioned above, we also made in the INC a number of other improvements and refinements, such as imposing momentum-energy conservation for each simulated event (the Monte-Carlo algorithm previously used in the CEM provided momentum-energy conservation only statistically, on the average, but not exactly for each simulated event) and using real binding energies for nucleons in the cascade instead of the approximation of a constant separation energy of 7 MeV used in previous versions of the CEM. We have also improved many algorithms used in the Monte-Carlo simulations in many subroutines, decreasing the computing time by up to a factor of 6 for heavy targets, which is very important when performing practical simulations with transport codes like MCNPX or MARS.

Let us mention that in the CEM the initial configuration for the preequilibrium decay (number of excited particles and holes, i.e. excitons $n_0 = p_0 + h_0$, excitation energy E_0^* , linear momentum \mathbf{P}_0 , and angular momentum \mathbf{L}_0 of the nucleus) differs significantly from that usually postulated in exciton models. Our calculations [2, 74, 75] have shown that the distributions of residual nuclei remaining after the cascade stage of the reaction, i.e. before the preequilibrium emission, with respect to n_0 , p_0 , h_0 , E_0^* , \mathbf{P}_0 , and \mathbf{L}_0 are rather broad.¹

¹Unfortunately, this fact was misunderstood by the authors of the code HETC-3STEP [51]. In spite of the fact that it has been stressed explicitly, and figures with distributions of excited nuclei after the cascade stage of a reaction with respect to the number of excitons and other characteristics were shown in a number of publications (see, e.g., Fig. 5 in Ref. [2], Fig. 1 in Ref. [75], p. 109 in Ref. [74], and p. 706 in Ref. [26]), the authors of Ref. [51] misstated this fact as "Gudima et al. assumed the state of two particles and one hole at

2.2. The Coalescence Model

When the cascade stage of a reaction is completed, CEM03.xx uses the coalescence model described in Refs. [76, 77] to "create" high-energy d, t, ³He, and ⁴He by final-state interactions among emitted cascade nucleons, already outside of the target nucleus. In contrast to most other coalescence models for heavy-ion induced reactions, where complex particle spectra are estimated simply by convolving the measured or calculated inclusive spectra of nucleons with corresponding fitted coefficients (see, e.g., [78] and references therein), CEM03.xx uses in its simulation of particle coalescence real information about all emitted cascade nucleons and does not use integrated spectra. CEM03.xx assumes that all the cascade nucleons having differences in their momenta smaller than p_c and the correct isotopic content form an appropriate composite particle. This means that the formation probability for, e.g. a deuteron is

$$W_d(\vec{p}, b) = \int \int d\vec{p}_p d\vec{p}_n \rho^C(\vec{p}_p, b) \rho^C(\vec{p}_n, b) \delta(\vec{p}_p + \vec{p}_n - \vec{p}) \Theta(p_c - |\vec{p}_p - \vec{p}_n|),$$
(17)

where the particle density in momentum space is related to the one-particle distribution function f by

$$\rho^C(\vec{p}, b) = \int d\vec{r} f^C(\vec{r}, \vec{p}, b).$$
(18)

Here, b is the impact parameter for the projectile interacting with the target nucleus and the superscript index C shows that only cascade nucleons are taken into account for the coalescence process. The coalescence radii p_c were fitted for each composite particle in Ref. [76] to describe available data for the reaction Ne+U at 1.04 GeV/nucleon, but the fitted values turned out to be quite universal and were subsequently found to satisfactorily describe high-energy complex-particle production for a variety of reactions induced both by particles and nuclei at incident energies up to about 400 GeV/nucleon, when describing nuclear reactions with the Los Alamos version of the Quark-Gluon String Model (LAQGSM) [23, 36, 79] or with its predecessor, the Quark-Gluon String Model (QGSM) [80]. These parameters are:

$$p_c(d) = 90 \text{ MeV/c}; \quad p_c(t) = p_c({}^{3}\text{H}e) = 108 \text{ MeV/c}; \quad p_c({}^{4}\text{H}e) = 115 \text{ MeV/c}.$$
 (19)

As the INC of CEM03.xx is different from those of LAQGSM or QGSM, it is natural to expect different best values for p_c as well. Our recent studies show that the values of parameters p_c defined by Eq. (19) are also good for CEM03.xx for projectile particles with kinetic energies T_0 lower than 300 MeV and equal to or above 1 GeV. For incident energies in the interval 300 MeV $< T_0 \leq 1$ GeV, a better overall agreement with the available experimental data is obtained by using values of p_c equal to 150, 175, and 175 MeV/c for d, $t({}^{3}\text{He})$, and ${}^{4}\text{He}$, respectively. These values of p_c are fixed as defaults in CEM03.xx. If several cascade nucleons are chosen to coalesce into composite particles, they are removed from the distributions of nucleons and do not contribute further to such nucleon characteristics as spectra, multiplicities, etc.

2.3. Preequilibrium Reactions

The subsequent preequilibrium interaction stage of nuclear reactions is considered by the CEM in the framework of an extension of the Modified Exciton Model (MEM) [5, 6]. At

the beginning \cdots Hence, their assumption is not valid for the wide range of incident energy", claiming this as a weakness of the CEM and a priority of the code HETC-3STEP, where smooth distributions of excited nuclei after the cascade stage of reactions with respect to n_0 are used. This had already been done in the CEM [1, 2].

the preequilibrium stage of a reaction we take into account all possible nuclear transitions changing the number of excitons n with $\Delta n = +2, -2$, and 0, as well as all possible multiple subsequent emissions of $n, p, d, t, {}^{3}$ He, and 4 He. The corresponding system of master equations describing the behavior of a nucleus at the preequilibrium stage is solved by the Monte-Carlo technique [1, 2].

For a preequilibrium nucleus with excitation energy E and number of excitons n = p + h, the partial transition probabilities changing the exciton number by Δn are

$$\lambda_{\Delta n}(p,h,E) = \frac{2\pi}{\hbar} |M_{\Delta n}|^2 \omega_{\Delta n}(p,h,E) .$$
⁽²⁰⁾

The emission rate of a nucleon of the type j into the continuum is estimated according to the detailed balance principle

$$\Gamma_{j}(p,h,E) = \int_{V_{j}^{c}}^{E-B_{j}} \lambda_{c}^{j}(p,h,E,T) dT ,$$

$$\lambda_{c}^{j}(p,h,E,T) = \frac{2s_{j}+1}{\pi^{2}\hbar^{3}} \mu_{j} \Re_{j}(p,h) \frac{\omega(p-1,h,E-B_{j}-T)}{\omega(p,h,E)} T\sigma_{inv}(T) , \qquad (21)$$

where s_j , B_j , V_j^c , and μ_j are the spin, binding energy, Coulomb barrier, and reduced mass of the emitted particle, respectively. The factor $\Re_j(p, h)$ ensures the condition for the exciton chosen to be the particle of type j and can easily be calculated by the Monte-Carlo technique.

Assuming an equidistant level scheme with the single-particle density g, we have the level density of the *n*-exciton state as [81]

$$\omega(p,h,E) = \frac{g(gE)^{p+h-1}}{p!h!(p+h-1)!} .$$
(22)

This expression should be substituted into Eq. (21). For the transition rates (20), one needs the number of states taking into account the selection rules for intranuclear exciton-exciton scattering. The appropriate formulae have been derived by Williams [82] and later corrected for the exclusion principle and indistinguishability of identical excitons in Refs. [83, 84]:

$$\omega_{+}(p,h,E) = \frac{1}{2}g \frac{[gE - \mathcal{A}(p+1,h+1)]^{2}}{n+1} \left[\frac{gE - \mathcal{A}(p+1,h+1)}{gE - \mathcal{A}(p,h)}\right]^{n-1},$$

$$\omega_{0}(p,h,E) = \frac{1}{2}g \frac{[gE - \mathcal{A}(p,h)]}{n} [p(p-1) + 4ph + h(h-1)],$$

$$\omega_{-}(p,h,E) = \frac{1}{2}gph(n-2),$$
(23)

where $\mathcal{A}(p,h) = (p^2 + h^2 + p - h)/4 - h/2$. By neglecting the difference of matrix elements with different Δn , $M_+ = M_- = M_0 = M$, we estimate the value of M for a given nuclear state by associating the $\lambda_+(p,h,E)$ transition with the probability for quasi-free scattering of a nucleon above the Fermi level on a nucleon of the target nucleus. Therefore, we have

$$\frac{\langle \sigma(v_{rel})v_{rel} \rangle}{V_{int}} = \frac{\pi}{\hbar} |M|^2 \frac{g[gE - \mathcal{A}(p+1,h+1)]}{n+1} \left[\frac{gE - \mathcal{A}(p+1,h+1)}{gE - \mathcal{A}(p,h)}\right]^{n-1} .$$
(24)

Here, V_{int} is the interaction volume estimated as $V_{int} = \frac{4}{3}\pi(2r_c + \lambda/2\pi)^3$, with the de Broglie wave length $\lambda/2\pi$ corresponding to the relative velocity $v_{rel} = \sqrt{2T_{rel}/m_N}$. A value of the order of the nucleon radius is used for r_c in the CEM: $r_c = 0.6$ fm.

The averaging in the left-hand side of Eq. (24) is carried out over all excited states taking into account the Pauli principle in the approximation

$$\langle \sigma(v_{rel})v_{rel} \rangle \simeq \langle \sigma(v_{rel}) \rangle \langle v_{rel} \rangle$$
 (25)

The averaged cross section $\langle \sigma(v_{rel}) \rangle$ is calculated by the Monte-Carlo simulation method and by introducing a factor η effectively taking into account the Pauli principle exactly as is done in the Fermi-gas model (see, e.g., [85])²

$$\sigma(v_{rel}) = \frac{1}{2} [\sigma_{pp}(v_{rel}) + \sigma_{pn}(v_{rel})] \eta(T_F/T) , \text{ where}$$
(26)

$$\eta(x) = \begin{cases} 1 - \frac{7}{5}x, & \text{if } x \le 0.5 ,\\ 1 - \frac{7}{5}x + \frac{2}{5}x(2 - \frac{1}{x})^{5/2}, & \text{if } x > 0.5 . \end{cases}$$
(27)

Here, v_{rel} is the relative velocity of the excited nucleon (exciton) and the target nucleon in units of the speed of light and T is the kinetic energy of the exciton. The free-particle interaction cross sections $\sigma_{pp}(v_{rel})$ and $\sigma_{pn}(v_{rel})$ in Eq. (26) are estimated using the relations suggested by Metropolis et al. [86]

$$\sigma_{pp}(v_{rel}) = \frac{10.63}{v_{rel}^2} - \frac{29.92}{v_{rel}} + 42.9 ,$$

$$\sigma_{pn}(v_{rel}) = \frac{34.10}{v_{rel}^2} - \frac{82.2}{v_{rel}} + 82.2 ,$$
(28)

where the cross sections are given in mb.

The relative kinetic energy of colliding particles necessary to calculate $\langle v_{rel} \rangle$ and the factor η in Eqs. (26,27) are estimated in the so-called "right-angle collision" approximation [5], i.e. as a sum of the mean kinetic energy of an excited particle (exciton) measured from the bottom of the potential well $T_p = T_F + E/n$ plus the mean kinetic energy of an intranuclear nucleon partner $T_N = 3T_F/5$, that is $T_{rel} = T_p + T_N = 8T_F/5 + E/n$.

Combining (20), (22) and (24), we get finally for the transition rates:

$$\lambda_{+}(p,h,E) = \frac{\langle \sigma(v_{rel})v_{rel} \rangle}{V_{int}},$$

$$\lambda_{0}(p,h,E) = \frac{\langle \sigma(v_{rel})v_{rel} \rangle}{V_{int}} \frac{n+1}{n} \Big[\frac{gE - \mathcal{A}(p,h)}{gE - \mathcal{A}(p+1,h+1)} \Big]^{n+1} \frac{p(p-1) + 4ph + h(h-1)}{gE - \mathcal{A}(p,h)},$$

$$\lambda_{-}(p,h,E) = \frac{\langle \sigma(v_{rel})v_{rel} \rangle}{V_{int}} \Big[\frac{gE - \mathcal{A}(p,h)}{gE - \mathcal{A}(p+1,h+1)} \Big]^{n+1} \frac{ph(n+1)(n-2)}{[gE - \mathcal{A}(p,h)]^{2}}.$$
(29)

CEM considers the possibility of fast d, t, ³He, and ⁴He emission at the preequilibrium stage of a reaction in addition to the emission of nucleons. We assume that in the course of a reaction p_j excited nucleons (excitons) are able to condense with probability γ_j forming a complex particle which can be emitted during the preequilibrium state. A modification of

 $^{^{2}}$ Unfortunately, formula (27) as presented in Ref. [2] had some misprints; in the prior publication [1], it was correct.

Eq. (21) for the complex-particle emission rates is described in detail in Refs. [1, 2]. The "condensation" probability γ_j is estimated in those references as the overlap integral of the wave function of independent nucleons with that of the complex particle (cluster)

$$\gamma_j \simeq p_j^3 (V_j/V)^{p_j-1} = p_j^3 (p_j/A)^{p_j-1} .$$
(30)

This is a rather crude estimate. In the usual way the values γ_j are taken from fitting the theoretical preequilibrium spectra to the experimental ones, which gives rise to an additional, as compared to (30), dependence of the factor γ_j on p_j and excitation energy (see, e.g., Refs. [87, 88]), for each considered reaction.

The single-particle density g_j for complex particle states is found in the CEM by assuming the complex particles move freely in a uniform potential well whose depth is equal to the binding energy of this particle in a nucleus [2]

$$g_j(T) = \frac{V(2s_j + 1)(2\mu_j)^{3/2}}{4\pi^2\hbar^3} (T + B_j)^{1/2} .$$
(31)

As we stated previously, this is a crude approximation and it does not provide a good prediction of emission of preequilibrium α particles (see, e.g., [72] and references therein). In CEM03.xx, to improve the description of preequilibrium complex-particle emission, we estimate γ_j by multiplying the estimate provided by Eq. (30) by an empirical coefficient $M_j(A, Z, T_0)$ whose values are fitted to available nucleon-induced experimental complex-particle spectra. We fix the fitted values of $M_j(A, Z, T_0)$ in CEM03.xx and complement them with routines **gambetn** and **gambetp** for their interpolation outside the region covered by our fitting. As shown in one example in Fig. 9 of Appendix 3, after fitting $M_j(A, Z, T_0)$, CEM03.03 describes quite well the measured spectra of all complex particles, providing a much better agreement with experimental data than all its predecessors did.

The CEM predicts forward peaked (in the laboratory system) angular distributions for preequilibrium particles. For instance, CEM03.xx assumes that a nuclear state with a given excitation energy E^* should be specified not only by the exciton number *n* but also by the momentum direction Ω . Following Ref. [89], the master equation (11) from Ref. [2] can be generalized for this case provided that the angular dependence for the transition rates λ_+ , λ_0 , and λ_- (Eq. (29)) is factorized. In accordance with Eqs. (24) and (25), in the CEM it is assumed that

$$\langle \sigma \rangle \rightarrow \langle \sigma \rangle F(\Omega) ,$$
 (32)

where

$$F(\Omega) = \frac{d\sigma^{free}/d\Omega}{\int d\Omega' d\sigma^{free}/d\Omega'} .$$
(33)

The scattering cross section $d\sigma^{free}/d\Omega$ is assumed to be isotropic in the reference frame of the interacting excitons, thus resulting in an asymmetry in both the nucleus center-of-mass and laboratory frames. The angular distributions of preequilibrium complex particles are assumed [2] to be similar to those for the nucleons in each nuclear state.

This calculational scheme is easily realized by the Monte-Carlo technique. It provides a good description of double differential spectra of preequilibrium nucleons and a not-so-good but still satisfactory description of complex-particle spectra from different types of nuclear reactions at incident energies from tens of MeV to several GeV. For incident energies below about 200 MeV, Kalbach [90] has developed a phenomenological systematics for preequilibrium-particle angular distributions by fitting available measured spectra of nucleons and complex

particles. As the Kalbach systematics are based on measured spectra, they describe very well the double-differential spectra of preequilibrium particles and generally provide a better agreement of calculated preequilibrium complex particle spectra with data than does the CEM approach based on Eqs. (32,33). This is why we have incorporated into CEM03.xx the Kalbach systematics [90] to describe angular distributions of both preequilibrium nucleons and complex particles at incident energies up to 210 MeV. At higher energies, we use in CEM03.xx the CEM approach based on Eqs. (32,33).

By "preequilibrium particles" we mean particles which are emitted after the cascade stage of a reaction but before achieving statistical equilibrium at a time t_{eq} , which is fixed by the condition $\lambda_+(n_{eq}, E) = \lambda_-(n_{eq}, E)$ from which we get

$$n_{eq} \simeq \sqrt{2gE}$$
 . (34)

At $t \ge t_{eq}$ (or $n \ge n_{eq}$), the behavior of the remaining excited compound nucleus is described in the framework of both the Weisskopf-Ewing statistical theory of particle evaporation [91] and fission competition according to Bohr-Wheeler theory [92].

The parameter g entering into Eqs. (29) and (34) is related to the level-density parameter of single-particle states $a = \pi^2 g/6$. At the preequilibrium stage, we calculate the level-density parameter a with our own approximation [13] in the form proposed initially by Ignatyuk et al. [93], following the method by Iljinov et al. [94]:

$$a(Z, N, E^*) = \tilde{a}(A) \left\{ 1 + \delta W_{gs}(Z, N) \frac{f(E^* - \Delta)}{E^* - \Delta} \right\},\tag{35}$$

where

$$\tilde{a}(A) = \alpha A + \beta A^{2/3} B_s \tag{36}$$

is the asymptotic Fermi-gas value of the level density parameter at high excitation energies. Here, B_s is the ratio of the surface area of the nucleus to the surface area of a sphere of the same volume (for the ground state of a nucleus, $B_s \approx 1$), and

$$f(E) = 1 - exp(-\gamma E) . \tag{37}$$

 E^* is the total excitation energy of the nucleus, related to the "thermal" energy U by: $U = E^* - E_R - \Delta$, where E_R and Δ are the rotational and pairing energies, respectively.

We use the shell correction $\delta W_{gs}(Z, N)$ by Möller et al. [95] and the pairing energy shifts from Möller, Nix, and Kratz [96]. The values of the parameters α , β , and γ were derived in Ref. [13] by fitting the the same data analyzed by Iljinov et al. [94] (we discovered that Iljinov et al. used $11/\sqrt{A}$ for the pairing energies Δ in deriving their level-density systematics instead of the value of $12/\sqrt{A}$ stated in Ref. [94] and we also found several misprints in the nuclear level-density data shown in their Tables. 1 and 2 used in the fit). We find:

$$\alpha = 0.1463, \beta = -0.0716, \text{ and } \gamma = 0.0542$$

As mentioned in Section 2.1, the standard version of the CEM [2] provides an overestimation of preequilibrium particle emission from different p+A and A+A reactions we have analyzed (see more details in [14, 15]). One way to solve this problem suggested in Ref. [14] is to change the criterion for the transition from the cascade stage to the preequilibrium one, as described in Section 2.1. Another easy way suggested in Ref. [14] to shorten the preequilibrium stage of a reaction is to arbitrarily allow only transitions that increase the number of excitons, $\Delta n = +2$, i.e., only allow the evolution of a nucleus toward the compound nucleus. In this case, the time of the equilibration will be shorter and fewer preequilibrium particles will be emitted, leaving more excitation energy for the evaporation. Such a "never-come-back" approach is used by some other exciton models, for instance, by the Multistage Preequilibrium Model (MPM) used in LAHET [97] and by FLUKA [98]. This approach was used in the CEM2k [14] version of the CEM and it allowed us to describe much better the p+A reactions measured at GSI in inverse kinematics at energies around 1 GeV/nucleon. Nevertheless, the "never-come-back" approach seems unphysical, therefore we no longer use it. We now address the problem of emitting fewer preequilibrium particles in the CEM by following Veselský [99]. We assume that the ratio of the number of quasiparticles (excitons) n at each preequilibrium reaction stage to the number of excitation energy, to be a crucial parameter for determining the probability of preequilibrium emission P_{pre} . This probability for a given preequilibrium reaction stage is evaluated using the formula

$$P_{pre}(n/n_{eq}) = 1 - \exp\left(-\frac{(n/n_{eq} - 1)}{2\sigma_{pre}^2}\right)$$
(38)

for $n \leq n_{eq}$ and equal to zero for $n > n_{eq}$. The basic assumption leading to Eq. (38) is that P_{pre} depends exclusively on the ratio n/n_{eq} as can be deduced from the results of Böhning [100] where the density of particle-hole states is approximately described using a Gaussian centered at n_{eq} . The parameter σ_{pre} is a free parameter and we assume no dependence on excitation energy [99]. Our calculations of several reactions using different values of σ_{pre} show that an overall reasonable agreement with available data can be obtained using $\sigma_{pre} = 0.4$ -0.5 (see Fig. 11 in Ref. [15]). In CEM03.xx, we choose the fixed value $\sigma_{pre} = 0.4$ and use Eqs. (34,38) as criteria for the transition from the preequilibrium stage of reactions to evaporation, instead of using the "never-come-back" approach along with Eq. (34), as was done in CEM2k.

2.4. Evaporation

CEM03.xx uses an extension of the Generalized Evaporation Model (GEM) code GEM2 by Furihata [101]–[103] after the preequilibrium stage of reactions to describe evaporation of nucleons, complex particles, and light fragments heavier than ⁴He (up to ²⁸Mg) from excited compound nuclei and to describe their fission, if the compound nuclei are heavy enough to fission ($Z \ge 65$). The GEM is an extension by Furihata of the Dostrovsky evaporation model [104] as implemented in LAHET [97] to include up to 66 types of particles and fragments that can be evaporated from an excited compound nucleus plus a modification of the version of Atchison's fission model [105, 106] used in LAHET. Many of the parameters were adjusted by Furihata for a better description of fission reactions when using it in conjunction with the extended evaporation model.

A very detailed description of the GEM, together with a large amount of results obtained for many reactions using the GEM coupled either with the Bertini or ISABEL INC models in LAHET may be found in [101, 102]. Therefore, we present here only the main features of the GEM, following mainly [102] and using as well information obtained in private communications with Dr. Furihata.

Furihata did not change in the GEM the general algorithms used in LAHET to simulate evaporation and fission. The decay widths of evaporated particles and fragments are estimated using the classical Weisskopf-Ewing statistical model [91]. In this approach, the decay probability P_j for the emission of a particle j from a parent compound nucleus i with the total kinetic energy in the center-of-mass system between ϵ and $\epsilon + d\epsilon$ is

$$P_j(\epsilon)d\epsilon = g_j\sigma_{inv}(\epsilon)\frac{\rho_d(E-Q-\epsilon)}{\rho_i(E)}\epsilon d\epsilon,$$
(39)

where E [MeV] is the excitation energy of the parent nucleus i with mass A_i and charge Z_i , and d denotes a daughter nucleus with mass A_d and charge Z_d produced after the emission of ejectile j with mass A_j and charge Z_j in its ground state. σ_{inv} is the cross section for the inverse reaction, ρ_i and ρ_d are the level densities [MeV]⁻¹ of the parent and the daughter nucleus, respectively. $g_j = (2S_j + 1)m_j/\pi^2\hbar^2$, where S_j is the spin and m_j is the reduced mass of the emitted particle j. The Q-value is calculated using the excess mass M(A, Z) as $Q = M(A_j, Z_j) + M(A_d, Z_d) - M(A_i, Z_i)$. In GEM2, four mass tables are used to calculate Q-values, according to the following priorities, where a lower priority table is only used outside the range of validity of the higher priority one: (1) the Audi-Wapstra mass table [107], (2) theoretical masses calculated by Möller et al. [95], (3) theoretical masses calculated by Comay et al. [108], (4) the mass excess calculated using the old Cameron formula [109]. As does LAHET, GEM2 uses Dostrovsky's formula [104] to calculate the inverse cross section σ_{inv} for all emitted particles and fragments

$$\sigma_{inv}(\epsilon) = \sigma_g \alpha \left(1 + \frac{\beta}{\epsilon} \right) , \qquad (40)$$

which is often written as

$$\sigma_{inv}(\epsilon) = \begin{cases} \sigma_g c_n (1 + b/\epsilon) & \text{for neutrons} \\ \sigma_g c_j (1 - V/\epsilon) & \text{for charged particles} \end{cases},$$

where $\sigma_g = \pi R_b^2$ [fm²] is the geometrical cross section, and

$$V = k_j Z_j Z_d e^2 / R_c \tag{41}$$

is the Coulomb barrier in MeV.

One new ingredient in GEM2 in comparison with LAHET, which considers evaporation of only 6 particles (n, p, d, t, ³He, and ⁴He), is that Furihata includes the possibility of evaporation of up to 66 types of particles and fragments and incorporates into GEM2 several alternative sets of parameters b, c_j , k_j , R_b , and R_c for each particle type.

The 66 ejectiles considered by GEM2 for evaporation are selected to satisfy the following criteria: (1) isotopes with $Z_j \leq 12$; (2) naturally existing isotopes or isotopes near the stability line; (3) isotopes with half-lives longer than 1 ms. All the 66 ejectiles considered by GEM2 are shown in Table 1.

$\overline{Z_i}$	Eiectil	es					
$\frac{J}{0}$	n						
1	р	d	t				
2	³ He	$^{4}\mathrm{He}$	⁶ He	⁸ He			
3	⁶ Li	$^{7}\mathrm{Li}$	⁸ Li	⁹ Li			
4	$^{7}\mathrm{Be}$	⁹ Be	$^{10}\mathrm{Be}$	$^{11}\mathrm{Be}$	$^{12}\mathrm{Be}$		
5	$^{8}\mathrm{B}$	$^{10}\mathrm{B}$	$^{11}\mathrm{B}$	$^{12}\mathrm{B}$	$^{13}\mathrm{B}$		
6	$^{10}\mathrm{C}$	$^{11}\mathrm{C}$	$^{12}\mathrm{C}$	$^{13}\mathrm{C}$	$^{14}\mathrm{C}$	$^{15}\mathrm{C}$	$^{16}\mathrm{C}$
$\overline{7}$	^{12}N	$^{13}\mathrm{N}$	$^{14}\mathrm{N}$	$^{15}\mathrm{N}$	^{16}N	$^{17}\mathrm{N}$	
8	$^{14}\mathrm{O}$	$^{15}\mathrm{O}$	$^{16}\mathrm{O}$	$^{17}\mathrm{O}$	$^{18}\mathrm{O}$	$^{19}\mathrm{O}$	^{20}O
9	$^{17}\mathrm{F}$	$^{18}\mathrm{F}$	$^{19}\mathrm{F}$	20 F	$^{21}\mathrm{F}$		
10	$^{18}\mathrm{Ne}$	$^{19}\mathrm{Ne}$	20 Ne	$^{21}\mathrm{Ne}$	22 Ne	$^{23}\mathrm{Ne}$	$^{24}\mathrm{Ne}$
11	21 Na	22 Na	23 Na	24 Na	25 Na		
12	^{22}Mg	$^{23}\mathrm{Mg}$	$^{24}\mathrm{Mg}$	$^{25}\mathrm{Mg}$	$^{26}\mathrm{Mg}$	$^{27}\mathrm{Mg}$	$^{28}\mathrm{Mg}$

Table 1. The evaporated particles considered by GEM2

GEM2 includes several options for the parameter set in expressions (40,41):

1) The "simple" parameter set is given as $c_n = c_j = k_j = 1$, b = 0, and $R_b = R_c = r_0(A_i^{1/3} + A_d^{1/3})$ [fm]; users need to input r_0 .

2) The "precise" parameter set is used in GEM2 as the default, and we use this set in our present work.

A) For all light ejectiles up to α $(A_j \leq 4)$, the parameters determined by Dostrovsky et al. [104] are used in GEM2, namely: $c_n = 0.76 + c_a A_d^{-1/3}$, $b = (b_a A_d^{-2/3} - 0.050)/(0.76 + c_a A_d^{-1/3})$ (and b = 0 for $A_d \geq 192$), where $c_a = 1.93$ and $b_a = 1.66$, $c_p = 1 + c$, $c_d = 1 + c/2$, $c_t = 1 + c/3$, $c_{^3He} = c_{\alpha} = 0$, $k_p = k$, $k_d = k + 0.06$, $k_t = k + 0.12$, $k_{^3He} = k_{\alpha} - 0.06$, where c, k, and k_{α} are listed in Table 2 for a set of Z_d . Between the Z_d values listed in Table 2, c, k, and k_{α} are interpolated linearly. The nuclear distances are given by $R_b = 1.5A^{1/3}$ for neutrons and protons, and $1.5(A_d^{1/3} + A_j^{1/3})$ for d, t, ³He, and α .

Z_d	k	k_{lpha}	С
≤ 20	0.51	0.81	0.0
30	0.60	0.85	-0.06
40	0.66	0.89	-0.10
≥ 50	0.68	0.93	-0.10

Table 2. k, k_{α} , and c parameters used in GEM2

The nuclear distance for the Coulomb barrier is expressed as $R_c = R_d + R_j$, where $R_d = r_0^c A^{1/3}$, $r_0^c = 1.7$, and $R_j = 0$ for neutrons and protons, and $R_j = 1.2$ for d, t, ³He, and ⁴He. We note that several of these parameters are similar to the original values published by Dostrovsky et al. [104] but not exactly the same. Dostrovsky et al. [104] had $c_a = 2.2$, $b_a = 2.12$, and $r_0^c = 1.5$. Also, for the k, k_{α} , and c parameters shown in Table 2, they had slightly different values, shown in Table 3.

Z_d	k_p	c_p	k_{α}	c_{α}
10	0.42	0.50	0.68	0.10
20	0.58	0.28	0.82	0.10
30	0.68	0.20	0.91	0.10
50	0.77	0.15	0.97	0.08
≥ 70	0.80	0.10	0.98	0.06

Table 3. k_p , c_p , k_{α} , and c_{α} parameters from Ref. [104]

B) For fragments heavier than α $(A_j \ge 4)$, the "precise" parameters of GEM2 use values by Matsuse et al. [110], namely: $c_j = k = 1$, $R_b = R_0(A_j) + R_0(A_d) + 2.85$ [fm], $R_c = R_0(A_j) + R_0(A_d) + 3.75$ [fm], where $R_0(A) = 1.12A^{1/3} - 0.86A^{-1/3}$.

3) The code GEM2 contains two other options for the parameters of the inverse cross sections.

A) A set of parameters due to Furihata for light ejectiles in combination with Matsuse's parameters for fragments heavier than α . Furihata and Nakamura determined k_j for p, d, t, ³He, and α as follows [103]:

$$k_j = c_1 \log(Z_d) + c_2 \log(A_d) + c_3.$$

The coefficients c_1 , c_2 , and c_3 for each ejectile are shown in Table 4.

Table 4. c_1 , c_2 , and c_3 for p, d, t, ³He, and α from [103]

Ejectile	c_1	c_2	c_3
р	0.0615	0.0167	0.3227
d	0.0556	0.0135	0.4067
\mathbf{t}	0.0530	0.0134	0.4374
$^{3}\mathrm{He}$	0.0484	0.0122	0.4938
α	0.0468	0.0122	0.5120

When these parameters are chosen in GEM2, the following nuclear radius R is used in the calculation of V and σ_g :

	0	for $A = 1$,
	1.2	for $2 \le A \le 4$,
	2.02	for $5 \le A \le 6$,
$R = \langle$	2.42	for $A = 7$,
	2.83	for $A = 8$,
	3.25	for $A = 9$,
	$1.414A_d^{1/3} + 1$	for $A \ge 10$.

B) The second new option in GEM2 is to use Furihata's parameters for light ejectiles up to α and the Botvina *et al.* [111] parameterization for inverse cross sections for heavier ejectiles. Botvina et al. [111] found that σ_{inv} can be expressed as

$$\sigma_{inv} = \sigma_g \begin{cases} (1 - V/\epsilon) & \text{for } \epsilon \ge V + 1 \text{ [MeV]},\\ \exp[\alpha(\epsilon - V - 1)]/(V + 1) & \text{for } \epsilon < V + 1 \text{ [MeV]}, \end{cases}$$
(42)

where

$$\alpha = 0.869 + 9.91/Z_i,$$

$$V = \frac{Z_j Z_d}{r_0^b (A_j^{1/3} + A_d^{1/3})},$$

$$r_0^b = 2.173 \frac{1 + 6.103 \times 10^{-3} Z_j Z_d}{1 + 9.443 \times 10^{-3} Z_j Z_d} \text{ [fm]}$$

The expression of σ_{inv} for $\epsilon < V + 1$ shows the fusion reaction in the sub-barrier region. When using Eq. (42) instead of Eq. (40), the total decay width for a fragment emission can not be calculated analytically. Therefore, the total decay width must be calculated numerically and takes much CPU time.

The total decay width Γ_j is calculated by integrating Eq. (39) with respect to the total kinetic energy ϵ from the Coulomb barrier V up to the maximum possible value, (E-Q). The good feature of Dostrovsky's approximation for the inverse cross sections, Eq. (40), is its simple energy dependence that allows the analytic integration of Eq. (39). By using Eq. (40) for σ_{inv} , the total decay width for the particle emission is

$$\Gamma_j = \frac{g_j \sigma_g \alpha}{\rho_i(E)} \int_V^{E-Q} \epsilon \left(1 + \frac{\beta}{\epsilon}\right) \rho_d(E - Q - \epsilon) d\epsilon.$$
(43)

The level density $\rho(E)$ is calculated in GEM2 according to the Fermi-gas model using the expression [112]

$$\rho(E) = \frac{\pi}{12} \frac{\exp(2\sqrt{a(E-\delta)})}{a^{1/4}(E-\delta)^{5/4}},\tag{44}$$

where a is the level density parameter and δ is the pairing energy in MeV. As does LAHET, GEM2 uses the δ values evaluated by Cook et al. [113]. For those values not evaluated by Cook et al., δ 's from Gilbert and Cameron [112] are used instead. The simplest option for the level-density parameter in GEM2 is $a = A_d/8$ [MeV⁻¹], but the default is the Gilbert-Cameron-Cook-Ignatyuk (GCCI) parameterization from LAHET [97]:

$$a = \tilde{a} \frac{1 - e^{-u}}{u} + a_I \left(1 - \frac{1 - e^{-u}}{u} \right), \tag{45}$$

where $u = 0.05(E - \delta)$, and

$$a_{I} = (0.1375 - 8.36 \times 10^{-5} A_{d}) \times A_{d},$$

$$\tilde{a} = \begin{cases} A_{d}/8 & \text{for } Z_{d} < 9 \text{ or } N_{d} < 9, \\ A_{d}(a' + 0.00917S) & \text{for others.} \end{cases}$$

For deformed nuclei with $54 \leq Z_d \leq 78$, $86 \leq Z_d \leq 98$, $86 \leq N_d \leq 122$, or $130 \leq N_d \leq 150$, a' = 0.12 while a' = 0.142 for other nuclei. The shell corrections S is expressed as a sum of separate contributions from neutrons and protons, i.e. $S = S(Z_d) + S(N_d)$ from [112, 113] and are tabulated in [101].

The level density is calculated using Eq. (44) only for high excitation energies, $E \ge E_x$, where $E_x = U_x + \delta$ and $U_x = 2.5 + 150/A_d$ (all energies are in MeV). At lower excitation energies, the following [112] is used for the level density:

$$\rho(E) = \frac{\pi}{12} \frac{1}{T} \exp((E - E_0)/T), \qquad (46)$$

where T is the nuclear temperature defined as $1/T = \sqrt{a/U_x} - 1.5/U_x$. To provide a smooth connection of Eqs. (44) and (46) at $E = E_x$, E_0 is defined as $E_0 = E_x - T(\log T - 0.25 \log a - 1.25 \log U_x + 2\sqrt{aU_x})$.

For $E - Q - V < E_x$, substituting Eq. (46) into Eq. (44) we can calculate the integral analytically, if we neglect the dependence of the level density parameter a on E:

$$\Gamma_j = \frac{\pi g_j \sigma_g \alpha}{12\rho_i(E)} \{ I_1(t,t) + (\beta + V) I_0(t) \},$$
(47)

where $I_0(t)$ and $I_1(t, t_x)$ are expressed as

$$I_0(t) = e^{-E_0/T}(e^t - 1),$$

$$I_1(t, t_x) = e^{-E_0/T}T\{(t - t_x + 1)e^{t_x} - t - 1\},$$

where t = (E - Q - V)/T and $t_x = E_x/T$. For $E - Q - V \ge E_x$, the integral of Eq. (43) cannot be solved analytically because of the denominator in Eq. (44). However, it is approximated as

$$\Gamma_j = \frac{\pi g_j \sigma_g \alpha}{12\rho_i(E)} [I_1(t, t_x) + I_3(s, s_x)e^s + (\beta + V)\{I_0(t_x) - I_2(s, s_x)e^s\}],$$
(48)

where $I_2(s, s_x)$ and $I_3(s, s_x)$ are given by

$$I_2(s, s_x) = 2\sqrt{2} \{ s^{-3/2} + 1.5s^{-5/2} + 3.75s^{-7/2} - (s_x^{-3/2} + 1.5s_x^{-5/2} + 3.75s_x^{-7/2})e^{s_x - s} \}$$

$$\begin{split} I_3(s,s_x) &= (\sqrt{2}a)^{-1} [2s^{-1/2} + 4s^{-3/2} + 13.5s^{-5/2} + 60.0s^{-7/2} + 325.125s^{-9/2} \\ &- \{(s^2 - s_x^2)s_x^{-3/2} + (1.5s^2 + 0.5s_x^2)s_x^{-5/2} + (3.75s^2 + 0.25s_x^2)s_x^{-7/2} + (12.875s^2 + 0.625s_x^2)s_x^{-9/2} + (59.0625s^2 + 0.9375s_x^2)s_x^{-11/2} + (324.8s_x^2 + 3.28s_x^2)s_x^{-13/2}\}e^{s_x - s}], \end{split}$$

with $s = 2\sqrt{a(E - Q - V - \delta)}$ and $s_x = 2\sqrt{a(E_x - \delta)}$.

The particle type j to be evaporated is selected in GEM2 by the Monte-Carlo method according to the probability distribution calculated as $P_j = \Gamma_j / \sum_j \Gamma_j$, where Γ_j is given by Eqs. (47) or (48). The total kinetic energy ϵ of the emitted particle j and the recoil energy of the daughter nucleus is chosen according to the probability distribution given by Eq. (39). The angular distribution of ejectiles is simulated to be isotropic in the center-of-mass system.

According to Friedman and Lynch [114], it is important to include excited states in the particle emitted via the evaporation process along with evaporation of particles in their ground states, because it greatly enhances the yield of heavy particles. Taking this into consideration, GEM2 includes evaporation of complex particles and light fragments both in the ground states and excited states. An excited state of a fragment is included in calculations if its half-lifetime $T_{1/2}(s)$ satisfies the following condition:

$$\frac{T_{1/2}}{\ln 2} > \frac{\hbar}{\Gamma_j^*},\tag{49}$$

where Γ_j^* is the decay width of the excited particle (resonance). GEM2 calculates Γ_j^* in the same manner as for a ground-state particle emission. The *Q*-value for the resonance emission is expressed as $Q^* = Q + E_j^*$, where E_j^* is the excitation energy of the resonance. The spin state of the resonance S_j^* is used in the calculation of g_j , instead of the spin of the ground state S_j .

GEM2 uses the ground state masses m_j for excited states because the difference between the masses is negligible.

Instead of treating a resonance as an independent particle, GEM2 simply enhances the decay width Γ_j of the ground state particle emission as follows:

$$\Gamma_j = \Gamma_j^0 + \sum_n \Gamma_j^n,\tag{50}$$

where Γ_j^0 is the decay width of the ground state particle emission, and Γ_j^n is that of the *n*th excited state of the particle *j* emission which satisfies Eq. (49).

The total-kinetic-energy distribution of the excited particles is assumed to be the same as that of the ground-state particle. S_j^* , E_j^* , and $T_{1/2}$ used in GEM2 are extracted from the Evaluated Nuclear Structure Data File (ENSDF) database maintained by the National Nuclear Data Center at Brookhaven National Laboratory [115].

Note that when including evaporation of up to 66 particles in GEM2, its running time increases significantly compared to the case when evaporating only 6 particles, up to ⁴He. The major particles emitted from an excited nucleus are n, p, d, t, ³He, and ⁴He. For most cases, the total emission probability of particles heavier than α is negligible compared to those for the emission of light ejectiles. Our detailed study of different reactions (see, e.g., [116] and references therein) shows that if we study only nucleon and complex-particle spectra or only spallation and fission products and are not interested in light fragments, we can consider evaporation of only 6 types of particles in GEM2 and save much time, getting results very close to the ones calculated with the more time consuming "66" option. In CEM03.xx, we have introduced an input parameter called **nevtype** that defines the number of types of particles to be considered at the evaporation stage. The index of each type of particle that can be evaporated corresponds to the particle arrangement in Table 1, with values, e.g., of 1, 2, 3, 4, 5, and 6 for n, p, d, t, ³He, and ⁴He, with succeeding values up to 66 for ²⁸Mg. All 66 particles that can possibly evaporate are listed in CEM03.xx together with their mass number, charge, and spin values in the **block data bdejc**. For all ten examples of inputs and outputs of CEM03.03 included in Appendices 1 and 2, whose results (when run with much larger numbers of events to improve statistics) are plotted in the figures in Appendix 3, we have performed calculations taking into account only 6 types of evaporated particles (nevtype = 6) as well as with the "66" option (nevtype = 66) and we provide the corresponding computing time for these examples in the captions to the appropriate figures shown in Appendix 3. The "6" option can be up to several times faster than the "66" option, providing meanwhile almost the same results. Therefore we recommend that users of CEM03.03 use 66 for the value of the input parameter **nevtype** only when they are interested in all fragments heavier than ⁴He; otherwise, we recommend the default value of 6 for **nevtype**, saving computing time. Alternatively, users may choose intermediate values of **nevtype**, for example 9 if one wants to calculate the production of ⁶Li, or 14 for modeling the production of ⁹Be and lighter fragments and nucleons only, while still saving computing time compared to running the code with the maximum value of 66.

2.5. Fission

The fission model used in GEM2 is based on Atchison's model [105, 106] as implemented in LAHET [97], often referred in the literature as the Rutherford Appleton Laboratory (RAL) fission model, which is where Atchison developed it. In GEM2 there are two choices of parameters for the fission model: one of them is the original parameter set by Atchison [105, 106] as implemented in LAHET [97], and the other is a parameter set developed by Furihata [101, 102].

2.5.1. Fission Probability. The Atchison fission model is designed to describe only fission of nuclei with $Z \ge 70$ (we extended it in our CEM03.xx and LAQGSM03.xx codes down to $Z \ge 65$). It assumes that fission competes only with neutron emission, i.e., from the widths Γ_j of n, p, d, t, ³He, and ⁴He, the RAL code calculates the probability of evaporation of any particle. When a charged particle is selected to be evaporated, no fission competition is taken into account. When a neutron is selected to be evaporated, the code does not actually simulate its evaporation, instead it considers that fission probability P_f . This quantity is treated by the RAL code differently for the elements above and below Z = 89. The reasons Atchison split the calculation of the fission probability P_f are: (1) there is very little experimental information on fission in the region Z = 85 to 88, (2) the marked rise in the fission barrier for nuclei with Z^2/A below about 34 (see Fig. 2 in [106]) together with the disappearance of asymmetric mass splitting, indicates that a change in the character of the fission process occurs. If experimental information were available, a split between regions around $Z^2/A \approx 34$ would be more sensible [106].

1) 65 $\leq Z_j \leq 88$. For fissioning nuclei with 65 $\leq Z_j \leq 88$, GEM2 uses the original Atchison calculation of the neutron emission width Γ_n and fission width Γ_f to estimate the fission probability as

$$P_f = \frac{\Gamma_f}{\Gamma_f + \Gamma_n} = \frac{1}{1 + \Gamma_n / \Gamma_f}.$$
(51)

Atchison uses [105, 106] the Weisskopf and Ewing statistical model [91] with an energyindependent pre-exponential factor for the level density (see Eq. (44)) and Dostrovsky's [104] inverse cross section for neutrons and estimates the neutron width Γ_n as

$$\Gamma_n = 0.352(1.68J_0 + 1.93A_i^{1/3}J_1 + A_i^{2/3}(0.76J_1 - 0.05J_0)),$$
(52)

where J_0 and J_1 are functions of the level density parameter a_n and $s_n (= 2\sqrt{a_n(E - Q_n - \delta)})$,

$$J_0 = \frac{(s_n - 1)e^{s_n} + 1}{2a_n},$$
$$J_1 = \frac{(2s_n^2 - 6s_n + 6)e^{s_n} + s_n^2 - 6}{8a_n^2}$$

Note that the RAL model uses a fixed value for the level density parameter a_n , namely

$$a_n = (A_i - 1)/8, (53)$$

and this approximation is kept in GEM2 when calculating the fission probability according to Eq. (51), although it differs from the GCCI parameterization (45) used in GEM2 to calculate particle evaporation widths. The fission width for nuclei with $65 \le Z_j \le 88$ is calculated in the RAL model and in the GEM as

$$\Gamma_f = \frac{(s_f - 1)e^{s_f} + 1}{a_f},$$
(54)

where $s_f = 2\sqrt{a_f(E - B_f - \delta)}$ and the level density parameter in the fission mode a_f is fitted by Atchison to describe the measured Γ_f/Γ_n to be [106]:

$$a_f = a_n \Big(1.08926 + 0.01098 (\chi - 31.08551)^2 \Big), \tag{55}$$

and $\chi = Z^2/A$. The fission barriers B_f [MeV] are approximated by

$$B_f = Q_n + 321.2 - 16.7 \frac{Z_i^2}{A} + 0.218 \left(\frac{Z_i^2}{A_i}\right)^2.$$
 (56)

Note that neither the angular momentum nor the excitation energy of the nucleus are taken into account in finding the fission barriers.

2) $Z_j \geq 89$. For heavy fissioning nuclei with $Z_j \geq 89$ GEM2 follows the RAL model [105, 106] and does not calculate at all the fission width Γ_f and does not use Eq. (51) to estimate the fission probability P_f . Instead, the following semi-empirical expression obtained by Atchison [105, 106] by approximating the experimental values of Γ_n/Γ_f published by Vandenbosch and Huizenga [117] is used to calculate the fission probability:

$$\log(\Gamma_n/\Gamma_f) = C(Z_i)(A_i - A_0(Z_i)), \tag{57}$$

where C(Z) and $A_0(Z)$ are constants depending on the nuclear charge Z only. The values of these constants are those used in the current version of LAHET [97] and are tabulated in Table 5 (note that some adjustments of these values have been done since Atchison's papers [105, 106] were published).

Z	C(Z)	$A_0(Z)$
89	0.23000	219.40
90	0.23300	226.90
91	0.12225	229.75
92	0.14727	234.04
93	0.13559	238.88
94	0.15735	241.34
95	0.16597	243.04
96	0.17589	245.52
97	0.18018	246.84
98	0.19568	250.18
99	0.16313	254.00
100	0.17123	257.80
101	0.17123	261.30
102	0.17123	264.80
103	0.17123	268.30
104	0.17123	271.80
105	0.17123	275.30
106	0.17123	278.80

Table 5. C(Z) and $A_0(Z)$ values used in GEM2

In this approach the fission probability P_f is independent of the excitation energy of the fissioning nucleus and its angular momentum.

2.5.2. Mass Distribution. The selection of the mass of the fission fragments depends on whether the fission is symmetric or asymmetric. For a pre-fission nucleus with $Z_i^2/A_i \leq 35$, only symmetric fission is allowed. For $Z_i^2/A_i > 35$, both symmetric and asymmetric fission are

allowed, depending on the excitation energy of the fissioning nucleus. No new parameters were determined for asymmetric fission in GEM2.

For nuclei with $Z_i^2/A_i > 35$, whether the fission is symmetric or not is determined by the asymmetric fission probability P_{asy}

$$P_{asy} = \frac{4870e^{-0.36E}}{1 + 4870e^{-0.36E}}.$$
(58)

2.5.2.a. Asymmetric fission. For asymmetric fission, the mass of one of the post-fission fragments A_1 is selected from a Gaussian distribution of mean $A_f = 140$ and width $\sigma_M = 6.5$. The mass of the second fragment is $A_2 = A_i - A_1$.

2.5.2.b. Symmetric fission. For symmetric fission, A_1 is selected from the Gaussian distribution of mean $A_f = A_i/2$ and two options for the width σ_M as described below.

The first option for choosing σ_M is the original Atchison approximation:

$$\sigma_M = \begin{cases} 3.97 + 0.425(E - B_f) - 0.00212(E - B_f)^2, \\ 25.27, \end{cases}$$
(59)

for $(E - B_f)$ below or above 100 MeV, respectively. In this expression all values are in MeV and the fission barriers B_f are calculated according to Eq. (56) for nuclei with $Z_i \leq 88$. For nuclei with $Z_i > 88$, the expression by Neuzil and Fairhall [118] is used:

$$B_f = C - 0.36(Z_i^2/A_i), (60)$$

where C = 18.8, 18.1, 18.1, and 18.5 [MeV] for odd-odd, even-odd, odd-even, and even-even nuclei, respectively.

The second option in GEM2 for σ_M (used here) was found by Furihata as:

$$\sigma_M = C_3 (Z_i^2 / A_i)^2 + C_4 (Z_i^2 / A_i) + C_5 (E - B_f) + C_6.$$
(61)

The constants $C_3 = 0.122$, $C_4 = -7.77$, $C_5 = 3.32 \times 10^{-2}$, and $C_6 = 134.0$ were obtained by fitting with GEM2 the recent Russian collection of experimental fission-fragment mass distributions [119]. In this expression, the fission barriers B_f by Myers and Swiatecki [120] are used. More details may be found in Ref. [102].

2.5.3. Charge Distribution. The charge distribution of fission fragments is assumed to be a Gaussian distribution of mean Z_f and width σ_Z . Z_f is expressed as

$$Z_f = \frac{Z_i + Z_1' - Z_2'}{2},\tag{62}$$

where

$$Z'_{l} = \frac{65.5A_{l}}{131 + A_{l}^{2/3}}, l = 1 \text{ or } 2.$$
(63)

The original Atchison model uses $\sigma_Z = 2.0$. An investigation by Furihata [102] suggests that $\sigma_Z = 0.75$ provides a better agreement with data; therefore $\sigma_Z = 0.75$ is used in GEM2 and in our code.

2.5.4. Kinetic Energy Distribution. The kinetic energy of fission fragments [MeV] is determined by a Gaussian distribution with mean ϵ_f and width σ_{ϵ_f} .

The original parameters in the Atchison model are:

$$\epsilon_f = 0.133 Z_i^2 / A_i^{1/3} - 11.4,$$

 $\sigma_{\epsilon_f} = 0.084 \epsilon_f.$

Furihata's parameters in the GEM, which we also use, are:

$$\epsilon_f = \begin{cases} 0.131 Z_i^2 / A_i^{1/3}, \\ 0.104 Z_i^2 / A_i^{1/3} + 24.3, \end{cases}$$
(64)

for $Z_i^2/A_i^{1/3} \leq 900$ and $900 < Z_i^2/A_i^{1/3} \leq 1800$, respectively, according to Rusanov et al. [119]. By fitting the experimental data by Itkis et al. [121], Furihata found the following expression for σ_{ϵ_f}

$$\sigma_{\epsilon_f} = \begin{cases} C_1(Z_i^2/A_i^{1/3} - 1000) + C_2, \\ C_2, \end{cases}$$
(65)

for $Z_i^2/A_i^{1/3}$ above and below 1000, respectively, and the values of the fitted constants are $C_1 = 5.70 \times 10^{-4}$ and $C_2 = 86.5$. The experimental data used by Furihata for fitting are the values extrapolated to the nuclear temperature 1.5 MeV by Itkis et al. [121]. More details may be found in [102].

We note that Atchison has also modified his original version using recent data and published [122] improved (and more complicated) parameterizations for many quantities and distributions in his model, but these modifications [122] have not been included either in LAHET or in GEM2.

2.5.5. Modifications to GEM2 in CEM03.xx. First, we fixed several observed uncertainties and small errors in the 2002 version of GEM2 Dr. Furihata kindly sent us. Then, we extended GEM2 to describe fission of lighter nuclei, down to $Z \ge 65$, and modified it [17] so that it provides a good description of fission cross sections when it is used after our INC and preequilibrium models.

If we had merged GEM2 with the INC and preequilibrium-decay modules of CEM03.xx without any modifications, the new code would not describe correctly fission cross sections (and the yields of fission fragments). This is because Atchison fitted the parameters of his RAL fission model when it followed the Bertini INC [123] which differs from ours. In addition, Atchison did not model preequilibrium emission. Therefore, the distributions of fissioning nuclei in A, Z, and excitation energy E^* simulated by Atchison differ significantly from the distributions we get; as a consequence, all the fission characteristics are also different. Furihata used GEM2 coupled either with the Bertini INC [123] or with the ISABEL [124] INC code, which also differs from our INC, and did not include preequilibrium particle emission. Therefore the distributions of fissioning nuclei simulated by Furihata differ from those in our simulations, so the parameters adjusted by Furihata to work well with her INC are not appropriate for us. To get a good description of fission cross sections (and fission-fragment yields) we have modified at least two parameters in GEM2 as used in CEM03.xx (see more details in [15, 16]).

The main parameters that determine the fission cross sections calculated by GEM2 are the level density parameter in the fission channel, a_f (or more exactly, the ratio a_f/a_n as calculated by Eq. (55)) for preactinides, and parameter C(Z) in Eq. (57) for actinides. The sensitivity of results to these parameters is much higher than to either the fission barrier heights used in a calculation or other parameters of the model. Therefore we choose [17] to adjust only these two parameters in our merged code. We do not change the form of systematics (55) and (57) derived by Atchison. We only introduce additional coefficients both to a_f and C(Z), replacing $a_f \to C_a \times a_f$ in Eq. (55) and $C(Z_i) \to C_c \times C(Z_i)$ in Eq. (57) and fit C_a and C_c to experimental proton-induced fission cross sections covered by Prokofiev's systematics [125]. No other parameters in GEM2 have been changed. For preactinides, we fit only C_a . The values of C_a found in our fit to Prokofiev's systematics are close to one and vary smoothly with the proton energy and the charge or mass number of the target. This result gives us some confidence in our procedure, and allows us to interpolate the values of C_a for nuclei and incident proton energies not analyzed by Prokofiev. For actinides, as described in [15, 16], we have to fit both C_a and C_c . The values of C_a we find are also very close to one, while the values of C_c are more varied, but both of them change smoothly with the proton energy and Z or A of the target, which again allows us to interpolate them for nuclei and energies outside Prokofiev's systematics.

We fix the fitted values of C_a and C_c in data blocks in our code and use the routines **fitafpa** and **fitafac** to interpolate to nuclei not covered by Prokofiev's systematics. We believe that such a procedure provides a reasonably accurate fission cross section calculation, at least for proton energies and target nuclei not too far from the ones covered by the systematics.

2.6. The Fermi Break-Up Model

After calculating the coalescence stage of a reaction, CEM03.xx moves to the description of the last slow stages of the interaction, namely to preequilibrium decay and evaporation, with a possible competition of fission. But as mentioned above, if the residual nuclei have atomic numbers with A < 13, CEM03.xx uses the Fermi break-up model [126] to calculate their further disintegration instead of using the preequilibrium and evaporation models.

All formulas and details of the algorithms used in the version of the Fermi break-up model developed in the former group of Prof. Barashenkov at Joint Institute for Nuclear Research (JINR), Dubna, Russia and released in CEM03.xx may be found in [45]. All the information needed to calculate the break-up of an excited nucleus is its excitation energy U and the mass and charge numbers A and Z. The total energy of the nucleus in the rest frame will be E = U + M(A, Z), where M is the mass of the nucleus. The total probability per unit time for a nucleus to break up into n components in the final state (e.g., a possible residual nucleus, nucleons, deuterons, tritons, alphas, etc.) is given by

$$W(E,n) = (V/\Omega)^{n-1}\rho_n(E), \tag{66}$$

where ρ_n is the density of final states, V is the volume of the decaying system and $\Omega = (2\pi\hbar)^3$ is the normalization volume. The density $\rho_n(E)$ can be defined as a product of three factors:

$$\rho_n(E) = M_n(E)S_nG_n. \tag{67}$$

The first one is the phase space factor defined as

$$M_n(E) = \int_{-\infty}^{+\infty} \cdots \int_{-\infty}^{+\infty} \delta\left(\sum_{b=1}^n \vec{p}_b\right) \delta\left(E - \sum_{b=1}^n \sqrt{p^2 + m_b^2}\right) \prod_{b=1}^n d^3 p_b,\tag{68}$$

where \vec{p}_b are fragment momenta. The second one is the spin factor

$$S_n = \prod_{b=1}^n (2s_b + 1), \tag{69}$$

which gives the number of states with different spin orientations. The last one is the permutation factor

$$G_n = \prod_{j=1}^k \frac{1}{n_j!},$$
(70)

which takes into account identical particles in the final state $(n_j \text{ is the number of components})$ of *j*-type particles and *k* is defined by $n = \sum_{j=1}^{k} n_j$. For example, if we have in the final state six particles (n = 6) and two of them are alphas, three are nucleons, and one is a deuteron, then $G_6 = 1/(2!3!1!) = 1/12$. For the non-relativistic case, the integration in Eq. (68) can be evaluated analytically (see, e.g., [45]) and the probability for a nucleus to disintegrate into *n* fragments with masses m_b , where $b = 1, 2, 3, \ldots, n$ is

$$W(E,n) = S_n G_n \left(\frac{V}{\Omega}\right)^{n-1} \left(\frac{1}{\sum_{b=1}^n m_b} \prod_{b=1}^n m_b\right)^{3/2} \frac{(2\pi)^{3(n-1)/2}}{\Gamma(3(n-1)/2)} E^{(3n-5)/2},\tag{71}$$

where $\Gamma(x)$ is the gamma function.

The angular distribution of n emitted fragments is assumed to be isotropic in the c.m. system of the disintegrating nucleus and their kinetic energies are calculated from momentumenergy conservation. The Monte-Carlo method is used to randomly select the decay channel according to probabilities defined by Eq. (71). Then, for a given channel, CEM03.xx calculates kinematical quantities for each fragment according to the *n*-body phase space distribution using Kopylov's method [127]. Generally, CEM03.xx considers formation of fragments only in their ground and those low-lying states which are stable for nucleon emission. However, several unstable fragments with large lifetimes: ⁵He, ⁵Li, ⁸Be, ⁹B, etc. were considered as well in the version of the Fermi break-up model as incorporated at JINR in a FORTRAN routine used by several transport codes, as described in [45]. (Let us recall here that, as was mentioned above in Section 2, we have addressed and fixed at LANL the problem of production of unstable or/and unphysical fragments in the "03.02" versions of our CEM and LAQGSM codes; CEM03.03 uses the fixed version of the Fermi break-up model, which does not provide unstable or/and unphysical fragments). The randomly chosen channel will be allowed to decay only if the total kinetic energy E_{kin} of all fragments at the moment of break-up is positive, otherwise a new simulation will be performed and a new channel will be selected. The total kinetic energy E_{kin} can be calculated according to the equation:

$$E_{kin} = U + M(A, Z) - E_{Coulomb} - \sum_{b=1}^{n} (m_b + \epsilon_b),$$
 (72)

where m_b and ϵ_b are masses and excitation energies of fragments, respectively, and $E_{Coulomb}$ is the Coulomb barrier for the given channel. It is approximated by

$$E_{Coulomb} = \frac{3}{5} \frac{e^2}{r_0} \left(1 + \frac{V}{V_0} \right)^{-1/3} \left(\frac{Z^2}{A^{1/3}} - \sum_{b=1}^n \frac{Z_b^2}{A_b^{1/3}} \right),$$
(73)

where A_b and Z_b are the mass number and the charge of the *b*-th particle of a given channel, respectively. V_0 is the volume of the system corresponding to normal nuclear density and $V = kV_0$ is the decaying system volume (we assume k = 1 in CEM03.xx). Thus, the Fermi break-up model used here has only one free parameter, V or V_0 , the volume of decaying system, which is estimated as follows:

$$V = 4\pi R^3/3 = 4\pi r_0^3 A/3,\tag{74}$$

where we use $r_0 = 1.4$ fm. This parameter is used to calculate the quantity **bl** in the routine **gitab**.

There is no limitation on the number n of fragments a nucleus may break up into in our version of the break-up model, in contrast to implementations in other codes, such as $n \leq 3$ in MCNPX, or $n \leq 7$ in LAHET.

2.7. Total Reaction Cross Sections (Normalization)

CEM03.xx (just like many other INC-based models) calculates the total reaction cross section, σ_{in} , by the Monte-Carlo method using the geometrical cross section, σ_{geom} , and the number of inelastic, N_{in} , and elastic, N_{el} , simulated events, namely: $\sigma_{in} = \sigma_{geom} N_{in}/(N_{in}+N_{el})$. The value of the total reaction cross section calculated this way is printed in the beginning of the CEM03.xx output labeled as *Monte Carlo inelastic cross section*. This approach provides a good agreement with available data for reactions induced by nucleons, pions, and photons at incident energies above about 100 MeV, but is not reliable enough at energies below 100 MeV (see, e.g., Fig. 4 and Ref. [16] and Figs. 4 and 5 in Ref. [19]).

To address this problem, we have incorporated [16] into CEM03.xx the NASA systematics by Tripathi et al. [128] for all incident protons and for neutrons with energies above the maximum in the NASA reaction cross sections, and the Kalbach systematics [129] for neutrons of lower energy. For reactions induced by monochromatic and bremsstrahlung photons, we incorporate into CEM03.xx [19] the recent systematics by Kossov [130]. Details on these systematics together with examples of several total inelastic cross sections calculated with them compared with available experimental data may be found in [16, 19]. Our analysis of many different reactions has shown that at incident energies below about 100 MeV these systematics generally describe the total inelastic cross sections better that the Monte-Carlo method does, and no worse than the Monte-Carlo method at higher energies. Therefore we choose these systematics as the default for normalization of all CEM03.xx results. The total reaction cross sections calculated by these systematics are printed in the CEM03.xx output labeled as *Inelasic cross section used here.* (Of course, users may renormalize all the CEM03.xx results to the Monte-Carlo total reaction cross sections by making a small change to the code in the subroutine **typeout**).

3. Storage of Simulation Results

Although we have extended significantly the variety of characteristics printed in the CEM03.xx output as compared to its predecessors, no predetermined outputs can satisfy the needs of all users. Therefore we provide below the necessary information to help users to modify the output according to their specific needs.

Almost all information about all particles, light fragments, and residual nuclei (there may be two residual nuclei in the case of fission) from every inelastic simulated event is stored in two arrays, spt(5,150) and parz(6,150). The second index of both these arrays shows the serial number k of a particular particle or nucleus stored in these arrays. These arrays contain physical information only for $1 < k \leq k_{max}$, where k_{max} is the number of all products from a particular inelastic event; all their elements for $k > k_{max}$ are equal to zero. For historical reasons, there is some redundancy in the arrays; for example Θ , T_k , and Z are available from more than one array element. The contents of the arrays spt(i,k) and parz(j,k) are as follows (all values are in the laboratory system; all energies and masses are in MeV; all angles are in degrees):

1) $\operatorname{spt}(1,k) = \sin \Theta$ of particle k 2) $\operatorname{spt}(2,k) = \cos \Theta$ of particle k 3) $\operatorname{spt}(3,k) = T_k$, kinetic energy of particle k 4) $\operatorname{spt}(4,k) = \operatorname{electric} \operatorname{charge}(Z)$ of particle k 5) $\operatorname{spt}(5,k) = \operatorname{rest} \max$ of particle k

1) parz(1,k) = particle type (index), defined as:

$$1 = n$$

$$2 = p$$

$$3 = d$$

$$4 = t$$

$$5 = {}^{3}\text{He}$$

$$6 = {}^{4}\text{He}$$

$$7 = \pi^{-}$$

$$8 = \pi^{0}$$

$$9 = \pi^{+}$$

$$1000\text{Z} + \text{N} = \text{A} + 999\text{Z}, \text{ for products heavier than } {}^{4}\text{He}$$

$$2) \text{ parz}(2, \text{k}) = T_{k}, \text{ kinetic energy of particle k}$$

3) $parz(3,k) = \Theta$ of particle k

4) $parz(4,k) = \varphi$ of particle k

5) parz(5,k) = reaction mechanism type (index), as following:

- if < 100, the "k" particle was emitted at the INC stage of reaction; the value stored here is equal to the number of successive interaction acts n_c before emission of particle k (see more details in [26])
- = 100 for preequilibrium emission
- = 200 for particles produced via coalescence
- = 1000 for evaporation from spallation residue (and for residue itself)
- = 1500 for Fermi breakup

= 2000 for evaporation from fission product (and for fragments themselves) 6) parz(6,k) = electric charge of particle k.

The code does not store the value of k_{max} for each simulated event. To retrieve information about the products from an inelastic event, users should read (in **vlobd**) either array in a loop over k from 1 to 150 looking, e.g., at the mass of products stored in **spt(5,k)** until they get for $k = (k_{max} + 1)$ a zero value, indicating that there are no more products from this event, thereby determining k_{max} .

CEM03.xx does not describe emission of γ 's from residual nuclei with an excitation energy below the threshold of particle evaporation, i.e. a few MeV. (It neglects also emission of γ 's with higher energy, as a competitor to evaporation and preequilibrium-particle emission, since the cross sections of such processes are insignificant compared to those of particle emission.) When using CEM03.xx as an event generator in a transport code, it should be supplemented by a module with the same function as the **PHT** code from LAHET [97], which can describe the cooling of such excited nuclei via γ emission. For this, one needs to know the excitation energy of all residual nuclei provided by CEM03.xx. (CEM03.03 was incorporated in just this way into the LANL transport codes MCNP6 [54] and MCNPX [55].)

Note that in the case of Fermi break-up, we have no excited residual nuclei; it is assumed that all fragments are already in their ground states (unstable fragments are allowed to decay before filling the arrays **spt** and **parz**. For such events, we do not have to look for a residual nucleus to deposit its excitation energy via γ -emission. To know if this is the case, we have to look at the value of the parameter **fusion** stored in the common block: **common /dele/ sfu**, **wf, fusion, sigfw**. If **fusion = 0**, this is an event that ended with Fermi break-up and we have no excited residual nuclei.

If fusion = -1, this is an event without Fermi break-up and without fission and we have only one residual nucleus. Its excitation energy, ut (in MeV), is stored in the common block: common /bl1003/ ut, at, zt.

If fusion = +1, this is an event that ended with fission, so that we have two excited residual nuclei. Their excitation energy (in MeV) are equal to ex12(1) and ex12(2), their mass and charge numbers are equal to af12(1) and af12(2), and zf12(1) and zf12(2), respectively, and their kinetic energy (in MeV) are equal to tf12(1) and tf12(2), correspondingly. All this information is stored in the common block: common /ifiss/ af12(2), zf12(2), tf12(2), ex12(2), bf12(2,3), ifiss. In the case of fission, the parameter ifiss has a values of 1 (ifiss = 0 for events without fission); its value can be also used to determine if fission occurred in a particular event, in addition to checking the parameter fusion.

Unfortunately, GEM2 as used in CEM03.xx does not consider at all the angular momenta of evaporated particles, the residual nucleus, and fission fragments. This is why the code does not provide values of angular momenta for the final reaction products. This is one problem we plan to address in the next version of CEM. We do calculate angular momenta of excited nuclei after the cascade and preequilibrium stages of reactions. After the cascade stage, their values (in units of \hbar), $L_x = amnucl(1)$, $L_y = amnucl(2)$, and $L_z = amnucl(3)$ are provided as output of the **subroutine cascad (enext, atwght, charge, pnucl, amnucl, kstart, obr, nel**). After the preequilibrium stage, their values $L_x = angmon(1)$, $L_y = angmon(2)$, and $L_z = angmom(3)$ are stored in the common block: **common /resid/ angmom(3), v(3), remn**. These values are used to build several distributions printed in the CEM03.xx output. They are presently of only "academic" interest to study nuclear reactions but are not ready for applications, as GEM2 does not consider angular momenta and CEM03.xx does not provide angular momenta for the final reaction products.

With this information and our routines **vlobd**, **resdist**, **opandis**, and **disnmul** as examples of how to build the histograms of needed characteristics and our routines **prinp** and **typeout** that write them into our output file, users should be able to write their own customized output tables.
4. Input File

CEM03.xx has four data files: mass.tbl, level.tbl, gamman.tbl, and level.tbl, which should not be changed by users. It uses one user-specified input file called cem03.inp that must be prepared to define a calculation. It has 25 obligatory lines that describe the reaction to be calculated and the desired format of the output, and can contain also up to 10 lines of text with comments to be printed near the beginning of the output file.

4.1. 1st Input Line

This line defines the name of an auxiliary output file where some diagnostic information is printed (no results of calculations are stored in this file; the information printed in it is very useful if we encounter an unexpected problem in calculation of a specific reaction, like a "bug"). This name may contain up to 30 characters.

4.2. 2nd Input Line

This line defines the name for the CEM03.03 output file, again with up to 30 characters.

4.3. 3rd to 5th Input Lines

The 3rd line defines the projectile. Use prot for protons, neut for neutrons, pipl for π^+ , pimi for π^- , pize for π^0 , gamm for monochromatic γ 's, and gamb for bremsstrahlung γ 's. The 4th and 5th lines continue the description of the line 3 input, which is too long to fit entirely on the 3rd line.

4.4. 6th Input Line

This line defines the projectile kinetic energy in MeV, t0mev, for the case where we need to calculate only one energy and the reaction is not induced by bremsstrahlung γ 's. When we need to calculate a reaction at several energies (with a step defined by the 10th line and the final energy by the 11th line), t0mev is the initial energy. For reactions induced by bremsstrahlung (only), we recommend using 30.0 on this line; it is the value of E_{min} , the minimum energy of the bremsstrahlung γ spectrum to be considered in calculations (above the GDR region), as described in [19].

4.5. 7th Input Line

This line defines the mass number A of the target nucleus.

4.6. 8th Input Line

This line defines the atomic number Z of the target nucleus.

4.7. 9th Input Line

This line defines the number of inelastic events to be simulated for this particular case. The appropriate value for this number depends on the characteristics of a reaction in which we are interested, as well as on the target and projectile: To calculate only fission cross sections of actinides or the mean multiplicities of nucleons from any reaction, 5000 inelastic events would be more than enough. To get double differential spectra of particles with small energy and angle bins, we may need to simulate 1,000,000 inelastic events. To have satisfactory statistics for the cross sections for production of isotopes in regions between spallation and fission and between fission and fragmentation (whose yields are very small) and for their mean energies and angles of emission, we may need to simulate 10,000,000 or even more inelastic events. The 10 examples shown in Appendix 3 may give guidance on choosing this number for different reactions.

4.8. 10th Input Line

This line defines the step of the projectile kinetic energy in MeV for calculating several incident energies in a single run. For only one incident energy, use a negative number, for example -5.0 for this parameter.

4.9. 11th Input Line

This line defines the maximum (final) kinetic energy of the projectile in MeV when we calculate a reaction at several energies, with a step defined on line 10 and the initial energy on line 6. The number must be greater than t0mev in order to calculate a single energy. Alternatively, for reactions induced by bremsstrahlung photons only, this parameter defines the end-point (maximum) energy in MeV of the bremsstrahlung γ spectrum, usually denoted in the literature as E_0 ; see details on bremsstrahlung reactions in [19] and in our Example 10 below.

Useful Hints:

When we need to calculate only one incident energy, e.g., 100 MeV, we suggest using for the t0mev parameter on the 6th line the value 100.0, for the parameter dt0 on the 10th line, any negative number, like -5.0, and for the parameter t0max on this line, any number bigger than 100.0, e.g., 5000.0.

If we need to calculate a reaction at three incident energies, e.g., 100, 200, and 300 MeV, use for t0mev on the 6th line the value 100.0, for the parameter dt0 on the 10th line, use the value 100.0, and for t0max on this line, use the values 300.5: The code will run in a loop at 100, 200, and 300 MeV, and will stop after 300 MeV, as with the incident energy step of 100 MeV the next incident energy would be 400 MeV, which is higher than 300.5 MeV given by the t0max on this line, so the code will stop.

4.10. 12th Input Line

This line defines the step-size $\Delta\Theta$ in degrees in the ejectile angular spectra $d\sigma/d\Omega$ [mb/sr]. It is used only when we calculate $d\sigma/d\Omega$, which is selected by the input parameter mang defined on the 18th line having the value 1 or 2.

4.11. 13th Input Line

This line defines whether or not we wish to calculate the angle-integrated energy spectra of ejectiles $d\sigma/dT$ in mb/MeV. Use 0 for the parameter mspec on this line to not calculate the

energy spectra, and 1 to calculate them. When we use mspec = 1, the code provides $d\sigma/dT$ for particles produced by all nuclear-reaction mechanisms considered in CEM03.03 labeled as Total, as well as their components from particles emitted at the intranuclear cascade (or produced via coalescence, in the case of complex particles), preequilibrium, and evaporation stages of reactions labeled as Cascade, Precompound, and Total Evaporation, respectively. Particles produced via Fermi break-up are included into the Total Evaporation component; in case of evaporation from heavy nuclei that fission, this component includes particles evaporated both before fission (often called in the literature "pre-fission") and evaporated from fission fragments after fission (called in the literature "post-fission"). At the end of each spectrum, its integral over the entire energy range (i.e., particle "yield" in mb) is provided, labeled as Integrated.

There is an additional option for this parameter, namely having it equal to 2 instead of 1, to study in more detail fission reactions: If we choose the value 2 for mspec, the code will provide energy spectra of ejectiles only from events that do fission (particles from events of this reaction that do not fission will be not included into $d\sigma/dT$ calculated with the option mspec = 2) and the normalization of spectra is made to the fissioning events only and not to the total number of simulated events as is done for mspec = 1. These spectra do not represent the spectra of all emitted particles and they can not be compared directly with the measured spectra that include all particles produced from events both with and without fission. This option is useful for comparing to coincidence experiments, where particles are measured in coincidence with fission fragments.

In addition to $d\sigma/dT$ in mb/MeV, the option mspec = 2 provides also spectra of particles normalized to one (i.e. probability spectra), often used in the literature when studying fission induced by low energy projectiles. The option mspec = 2 provides separately "pre-fission" and "post-fission" components of particle spectra labeled as Prefission and Fission Fragments, in addition to the Total, Cascade, Precompound, and Total Evaporation components of the spectra.

4.12. 14th Input Line

This line defines whether or not we wish to calculate secondary-particle multiplicities, yields (in mb), and mean kinetic energies (in MeV). If we use 0 for the parameter mpyld on this line, these characteristics will not be calculated. When mpyld = 1, these characteristics will be calculated and printed in a table near the beginning of the output. This table contains the total (labeled as T) mean multiplicity, yield, and kinetic energy of n, p, d, t, ³He, ⁴He, π^{-} , π^{0} , and π^+ , when these values are non-zero, as well as their components from cascade, preequilibrium, evaporation from events without fission (spallation), evaporation events that will fission just before fission, evaporation from fission fragments, the sum of all evaporated particles, and from coalescence, labeled as C, P, Sp, Pf, F, E, and Co, respectively. Particles produced via Fermi Break-up are included in "evaporation" (E). As in the case of the parameter mspec defined on the 13th line, there is an additional option for this parameter, mpyld = 2, to study in more detail fission. In this case, only events with fission will be considered and the mean multiplicities, yields, and kinetic energies of particles produced only in fission events will be included in this table; their normalization is done to the fission events only. As mentioned above about particle spectra, these characteristics obtained with the option mpyld = 2 can be compared directly only with models or data from coincidence measurements which select fission reactions.

4.13. 15th Input Line

This line defines whether or not we wish to calculate cross sections of 192 possible specific "channels" of reactions that contribute to the production of final isotopes. Knowledge of excitation functions for such "channels" are of mainly "academic" interest rather than for applications, as it splits the contributions to the production of a specific final isotope into different reaction channels. The γ -spectrometry method frequently used to measure nuclide production cross sections from, let us say, a photonuclear reaction on a target [Z,A], provides us only the cross section (yield) of a final isotope, e.g., [(Z-y),(A-x-y)] considering it to be produced via a $(\gamma, xnyp)$ reaction, without any information about the emitted x neutrons and y protons; have they been emitted as independent nucleons or contained in emitted complex particles? The option mchy = 1 of CEM03.03 defined on this line helps us to address this question, as it provides cross sections for different possible "channels" of a reaction that lead to the same final product nuclide. The option mchy = 1 requires significant additional computing time, so we recommend using it only when the contributions to the production of final isotopes from different processes need to be studied in detail. When not needed, use the faster option mchy = 0, in order to ignore such "channel" cross sections.

4.14. 16th Input Line

This line defines whether or not we wish to calculate cross sections ("yields") in mb of all nuclide products. The option misy = 0 does not provide such yields.

With the option misy = 1, CEM03.03 calculates yields of all isotopes produced in a reaction, as well as the integrated mass and charge yields in mb and the mass and charge distributions of their mean kinetic energies and their variances in MeV. In all cases, lines with all zero values are not written to the output file.

The option misy = 2 provides the same as misy = 1, plus:

it provides also yields of all nuclei, mass and charge distributions of cross sections and of mean kinetic energy of products emitted separately in the forward and backward directions in the laboratory system;

the average kinetic energy in MeV of all products;

mass yield and mean and variance of the laboratory emission angle in degrees as functions of the product mass number;

mean and variance of the z-velocity (parallel to the projectile beam) of all products in units of v/c and the ratio of the mass yields of products emitted in the forward direction in the laboratory system to those for the backward direction (the F/B ratio) and its variance as functions of product mass numbers;

similar distributions as functions of the atomic number of the products.

Finally, the most detailed option misy = 3 provides the same as misy = 2, plus:

mass distributions of excited nuclei after the cascade and preequilibrium stages of a reaction and distributions at the beginning of the evaporation stage of nuclei that will not fission, of ones that will fission, and of all nuclei just prior to fission, after any possible prefission evaporation;

similar distributions of nuclei as functions of their atomic number;

excitation energy distributions in 1/MeV of nuclei after the cascade and preequilibrium stages and distributions at the beginning of the evaporation stage of nuclei that will not fission, of ones that will fission, and of all nuclei just prior to fission;

similar distributions for linear momentum of nuclei in 1/MeV/c;

similar distributions for angular momentum of nuclei in $1/\hbar$;

distributions of fission-fragment opening angles in the laboratory system in 1/degrees in

different bins of neutron multiplicity < n >, namely, for < n >= 0 - 5, < n >= 6 - 8, < n >= 9 - 12, < n >= 13 - 15, < n >= 16 - 19, and $< n >\geq 20$, respectively;

neutron-multiplicity probabilities for all events, as well as for the emission of neutrons only during the cascade and preequilibrium stages of reaction, and for evaporation from events without fission (labeled in the output as Evap. res., from evaporation before fission for events that will fission (labeled in the output as Pre-fiss, and from evaporation from fission fragments after fission (labeled in the output as Post-fiss).

At the end of tables with these distributions, the mean values (labeled as < ... >), their standard deviations (labeled as St dv), and the corresponding normalization factor (labeled as norm.) are printed in the output, respectively.

4.15. 17th Input Line

The parameter mdubl on this line defines whether or not to calculate double-differential spectra of ejectiles $d^2\sigma/(dTd\Omega)$ in mb/(MeV sr). Use for the parameter mdubl values of 0, 1, or 2 to not calculate, to calculate, or to calculate only for fission events $d^2\sigma/(dTd\Omega)$ of ejectiles, in the same manner as described above for the parameter mspec defined on the 13th line.

4.16. 18th Input Line

The parameter mang on this line defines whether or not to calculate energy-integrated angular spectra of ejectiles $d\sigma/d\Omega$ in mb/sr. Use for the parameter mang values of 0, 1, or 2 to not calculate, to calculate, or to calculate only for fission events $d\sigma/d\Omega$ of ejectiles, in the same manner as described above for the parameter mspec defined on the 13th line.

4.17. 19th Input Line

The parameters ipar1 and ipar2 on this line define the range of the ejectile types for calculations of spectra $d\sigma/dT$, $d\sigma/d\Omega$, and $d^2\sigma/(dTd\Omega)$, with the notation of the particle "type" (ID) as described above: 1, 2, 3, 4, 5, 6, 7, 8, and 9 for n, p, d, t, ³He, ⁴He, π^- , π^0 , and π^+ , respectively. Note that the code does not allow us to calculate electively spectra of only several particles when their ID are not ordered: e.g., if we need spectra of only n, d, and π^+ , we will have to calculate in a loop spectra of all 9 types of particles, using ipar1 = 1 and ipar1 = 9. To calculate spectra (and multiplicities) of fragments heavier than ⁴He, users will have to modify the writing to the output file for themselves.

4.18. 20th Input Line

This line defines up to 10 angle bins $\Theta(j)$ from $\Theta_1(j)$ to $\Theta_2(j)$ in degrees for doubledifferential spectra calculations (when mdubl = 1). The code calculates in a loop either 10 spectra starting with j = 1 to j = 10, or until a value of $\Theta_1(j)$ read from this line is negative. If we need, e.g., $d^2\sigma/(dTd\Omega)$ for only two angles ($\Theta(1) < \Theta(2)$), define on this line the corresponding values of $\Theta_1(1)$, $\Theta_2(1)$, and $\Theta_1(2)$, $\Theta_2(2)$, and use any negative number for $\Theta_1(3)$, like -5.0, then include values for the rest of $\Theta_1(j)$, $\Theta_2(j)$, up to j = 10 as well, since the code expects to encounter all 20 values.

4.19. 21st Input Line

This line defines energy bins $\Delta T(j)$ for four energy regions j, from $T_1(j)$ to $T_2(j)$ (j = 1, 4)in MeV for angle-integrated energy spectra $d\sigma/dT$ and double-differential spectra $d^2\sigma/(dTd\Omega)$ calculations (when mspec ≥ 1 or/and mdubl ≥ 1). Take care that the whole possible energy region is covered by the four energy regions chosen on this line, so that $T_2(1) = T_1(2)$, $T_2(2) = T_1(3)$, and $T_2(3) = T_1(4)$, as is done in our examples of inputs shown in Appendix 1. It is also important that the value of $T_2(4)$ be as large as the maximum energy particle to be encountered.

4.20. 22nd Input Line

The parameter **nevtype** on this line defines the number of up to 66 different types of particles to be possibly evaporated, as described above in Section 2.4. We recommend using for the parameter **nevtype** on this line values in the range **7–66** only when fragments heavier than ⁴He, need to be considered; otherwise, we suggest using a default value of **6**, saving much computing time. See more details at the end of Section 2.4.

4.21. 23rd Input Line

The parameter ityp on this line defines the version of the random-number generator to be used in simulations, as defined by the first seven "indices" in Table 1 of Ref. [62]. See also the comments in the randmc.f source file. Note that the final CEM03.03 results calculated with good statistics should not depend on this parameter, so a user could always use the default value of **1** for it. Rerunning a particular simulation with a different value of ityp can provide an independent estimate of statistical uncertainties.

4.22. 24rd Input Line

The parameter nh on this line defines the number of up to 10 lines of commentary text to be printed in the beginning of the output as a header describing the given calculation. It could be zero, if users do not need to have any comments in their output.

4.23. Input Lines from 24 to 34

Here, users put up to 10 lines of commentary text to be printed in the beginning of the output as a header, as described above (up to 72 characters per line).

4.24. Input lines from 24 + nh + 1 to ...

All lines from 3 to 24 + nh may be repeated (with appropriately changed input values) as many times as desired in order to study other energy ranges, target isotopes, projectiles, etc.

4.25. The Last Input Line

On the last line of the input should be: **stop**.

5. Output File

The CEM03.03 output has plenty of captions and descriptions of all quantities printed, therefore we hope that users will have few problems in understanding it, given the information in the previous Section. We mention here only a few points about the output.

In the beginning of the output, all CEM03.03 input parameters and approximations for the level-density parameters used at the preequilibrium stage of the reaction are listed.

Then the total reaction cross section is listed, as described in Section 2.7.

The total number of inelastic and elastic simulated events are printed. Note that the elastic cross section printed after that is only to give users an idea about its order of magnitude: CEM03.03, like many other INC-based models does not pretend to describe reliably elastic cross sections.

Following this are several tables with statistical information about the mean values of the excitation energy, charge, mass, and angular momentum of nuclei after the cascade and preequilibrium stages of reactions, as well as similar distributions (plus a little more) for fissioning nuclei. Possible negative values for the minimum excitation energy of nuclei after the INC labeled as E*min may occur for some reactions. This does not indicate that CEM03.03 met a problem in calculating that specific reaction or something was wrong. A negative "excitation energy" may appear in occasional cases at the beginning of the preequilibrium stage of a reaction, when the real excitation energy of a nucleus after the INC is positive but very small. We subtract from it at the preequilibrium stage the pairing energy and the rotational energy (which are ignored by the classical INC). These rare cases are handled internally by counting them the same as any other case exiting from the cascade with an excitation energy less than a particle binding energy (listed as a residual nucleus with only cascade particles in the particle arrays **spt** and **parz**).

Following these tables, the fission probability (labeled as Fissility) and the fission cross section in mb, if the target was heavy enough to fission, are presented. CEM03.03 calculates the fission probability and cross section in two different ways: by the direct Monte-Carlo method and using the statistical weight-function method (see details in [9]), and results from both methods are printed in the output. All the yields of all products are calculated in CEM03.03 using the direct Monte-Carlo method, using the fission cross section calculated by this method. Therefore, we suggest that users use the fission probability and cross section calculated by the direct Monte-Carlo method, if the statistics of the calculation are high enough so that the fission cross section provided by the Monte-Carlo method is not too small, and greater than its statistical error printed in the output. At low incident energies, for reactions on light preactinide nuclei, the fission cross sections calculated by the Monte-Carlo method may be too small (or even zero), with big statistical errors. In such cases, users should instead use the fission probabilities and cross sections calculated by the statistical weight-function method.

We would like to emphasize that all results provided by CEM03.03 for reactions induced by bremsstrahlung γ 's are normalized to the so-called "equivalent γ quanta", as is usually done in the literature (see details on bremsstrahlung reactions in [19]). This point is mentioned in the CEM03.03 output, but the situation is different from all other types of reactions considered by the code, so we wish to again remind users about this. The output and the figures for Example 10 in the Appendices illustrate this difference.

Finally, for the input parameters mspec, mpyld, mdubl, and mang having the value 2 instead of 1, the spectra, particle multiplicities and the mean kinetic energies printed in the output are for events with fission only but not for all simulated events.

Acknowledgments

The authors of this Manual and all co-authors of the CEM03.xx codes are grateful to Dr. Shiori Furihata for providing to us her Generalized Evaporation/fission Model code GEM2 which we have incorporated into CEM03.xx, several useful discussions, and allowing us to use GEM2 in our codes and to distribute it further to other users without needing further permission. We thank Dr. Helder Duarte for providing us with numerical values of experimental cross sections from his collection, useful discussions, and help. We thank Dr. Kumataro Ukai for providing us with numerical values of single-pion photoproduction cross sections from their compilation [131]. We thank Prof. Koh Sakamoto and Drs. Hiroshi Matsumura, Hiromitsu Haba, and Yasuji Oura for providing us with their publications and numerical tables of their measured data, as well as for useful discussions, help in creating several figures for us, and their interest in our modeling. We thank Dr. Igor Pshenichnov for sending us the $\gamma - p$ and $\gamma - n$ event generators from their Moscow photonuclear reaction INC [70]; we use a small portion of a large data file developed for this code in CEM03.xx.

We thank Drs. Arjan Koning, Nathalie Marie-Nourry, Valentin Blideanu, Alain Letourneau, Yury Titarenko, Vechaslav Batyaev, Vitaly Pronskikh, Carmen Villagrasa-Canton, Alexander Prokofiev, Anatoly Ignatyuk, and Satoshi Chiba for providing us with tabulated values of many of their measurements and experimental data by other authors from their collections, which we have used while developing CEM03.xx.

We thank Dr. Forrest Brown for providing to us his MCNP5 random number generator and allowing us to use it in our codes and to distribute it further to other users without needing further permission.

Last but not least, we express our gratitude to many colleagues, in particular Drs. Mark Chadwick, Tony Gabriel, Franz Gallmeier, Tim Goorley, Gerry Hale, Alexandra Heath, Robert Little, Marcus Mendenhall, Igor Moskalenko, Jerry Nolen, Richard Olsher, Jerry Peterson, Laurie Waters, Robert Webster, and Robert Weller for useful discussions, interest in and support of our work.

This work was carried out under the auspices of the National Nuclear Security Administration of the U.S. Department of Energy at Los Alamos National Laboratory under Contract No. DE-AC52-06NA25396.

We ask users of CEM03.03 to contact us (specifically, SGM and AJS) using the E-mail addresses provided on the first page in case of questions on our code. We thank them in advance for comments and information about possible problems in using the code or "bugs" they will find.

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Appendix 1

CEM03.03 Input Example 1

p500Ni6	5.inf	/File name for diagnostic output. (<31 char.)
p500Ni6	.res	/File name for results of calculation. (<31 char.)
prot	/pname/	projectile particle name:
	p	rot - proton, neut - neutron, pipl - pi+, pimi - pi-, pize - pi0,
	ga	amm - gamma with fixed energy, gamb - bremss. gamma, stop - no more calc.
500.0	/tOmev/	minimum (initial) projectile kinetic energy in MeV; [tgmin for gamb]
58.	/anucl/	target mass number
28.	/znucl/	target atomic number
10000	/limc/	total number of inelastic events, normally 2000-500000
-20.	/dt0/	projectile kinetic energy step-size in MeV [Only 1 energy if <0.]
500.5	/tOmax/	maximum (final) projectile kinetic energy in MeV, [tgmax for gamb]
10.	/dteta/	<pre>step-size (degrees) in ejectile angular distributions [mang > 0]</pre>
0	/mspec/	(0/1,2) if ejectile energy spectra (are not/are) needed
1	/mpyld/	(0/1,2) if particle yield tables (are not/are) needed
0	/mchy/	(0/1) if particle channel yields (are not/are) needed
0	/misy/	(0/1,2,3) if isotope yields (are not/are) needed
1	/mdubl/	(0/1,2) if double differential spectra (are not/are) needed
0	/mang/	(0/1,2) if angular distributions (are not/are) needed
22/i	.par1,ipa	2/ range of ejectile types for spectrum calcs. [Below, ang. bins for mdubl > 0]
60.0 70	0.0 85.0 9	D5.0 115.0 125.0 165.0 -5.0 65.0 75.0 85.0 95.0 115.0 125.0 135.0 145.0 155.0 165.0
0. 22.	1. 22. 10	00. 3. 100. 500. 10. 500. 5000. 200. /tmin, tmax, dt, j=1-4/
6	/nevtype/	'number of evaporated particle types (see table in bldatgem.f).
1	/ityp/	Version of the random no. generator used; 1-7 OK; default 1
1	/nh/	Lines of text (<11) to be read in; printed on results file (line 2).
Example	e No. 1: H	Proton spectra from 500 MeV p + Ni58; 10,000 events.
stop		

pim5000	Cu.inf	/File name for diagnostic output. (<31 char.)
pim5000	Cu.res	/File name for results of calculation. (<31 char.)
pimi	/pname/	projectile particle name:
	p	rot - proton, neut - neutron, pipl - pi+, pimi - pi-, pize - piO,
	ga	amm - gamma with fixed energy, gamb - bremss. gamma, stop - no more calc.
500.0	/tOmev/	minimum (initial) projectile kinetic energy in MeV; [tgmin for gamb]
64.	/anucl/	target mass number
29.	/znucl/	target atomic number
10000	/limc/	total number of inelastic events, normally 2000-500000
-20.	/dt0/	projectile kinetic energy step-size in MeV [Only 1 energy if <0.]
500.5	/tOmax/	maximum (final) projectile kinetic energy in MeV, [tgmax for gamb]
10.	/dteta/	step-size (degrees) in ejectile angular distributions [mang > 0]
1	/mspec/	(0/1,2) if ejectile energy spectra (are not/are) needed
1	/mpyld/	(0/1,2) if particle yield tables (are not/are) needed
0	/mchy/	(0/1) if particle channel yields (are not/are) needed
0	/misy/	(0/1,2,3) if isotope yields (are not/are) needed
1	/mdubl/	(0/1,2) if double differential spectra (are not/are) needed
0	/mang/	(0/1,2) if angular distributions (are not/are) needed
88 /i	par1,ipa	r^2 range of ejectile types for spectrum calcs. [Below, ang. bins for mdubl > 0]
25.0 35	5.0 45.0 \$	55.0 65.0 75.0 -55.0 165.0 55.0 65.0 75.0 85.0 95.0 105.0 115.0 125.0 135.0 145.0 155.0 165.0
0. 500	10. 500	. 600. 10. 600. 700. 10. 700. 5000. 20. /tmin, tmax, dt, j=1-4/
6	/nevtype,	/ number of evaporated particle types (see table in bldatgem.f).
2	/ityp/	Version of the random no. generator used; 1-7 OK; default 1
1	/nh/	Lines of text (<11) to be read in; printed on results file (line 2).
Example	e No. 2: j	piO spectra from 500 MeV pi- + Cu64; 10,000 events.
stop		

n562Cu.inf /File name for diagnostic output. (<31 char.) n562Cu.res /File name for results of calculation. (<31 char.) neut /pname/ projectile particle name: prot - proton, neut - neutron, pipl - pi+, pimi - pi-, pize - pi0, gamm - gamma with fixed energy, gamb - bremss. gamma, stop - no more calc. 562.5 /tOmev/ minimum (initial) projectile kinetic energy in MeV; [tgmin for gamb] 64. /anucl/ target mass number /znucl/ target atomic number 29. 10000 /limc/ total number of inelastic events, normally 2000-500000 -10. /dt0/ projectile kinetic energy step-size in MeV [Only 1 energy if <0.] 600.5 /tOmax/ maximum (final) projectile kinetic energy in MeV, [tgmax for gamb] /dteta/ step-size (degrees) in ejectile angular distributions [mang > 0] 10. /mspec/ (0/1,2) if ejectile energy spectra (are not/are) needed 0 /mpyld/ (0/1,2) if particle yield tables (are not/are) needed 1 0 /mchy/ (0/1) if particle channel yields (are not/are) needed (0/1,2,3) if isotope yields (are not/are) needed 0 /misy/ 1 /mdubl/ (0/1,2) if double differential spectra (are not/are) needed (0/1,2) if angular distributions (are not/are) needed 0 /mang/ 9 9 /ipar1,ipar2/ range of ejectile types for spectrum calcs. [Below, ang. bins for mdubl > 0] 25.0 35.0 55.0 65.0 75.0 85.0 115.0 125.0 -5.0 65.0 75.0 85.0 95.0 105.0 115.0 125.0 135.0 145.0 155.0 165.0 0. 500. 20. 500. 600. 20. 600. 700. 20. 700. 5000. 20. /tmin, tmax, dt, j=1-4/ $\,$ /nevtype/ number of evaporated particle types (see table in bldatgem.f). 6 /ityp/ Version of the random no. generator used; 1-7 OK; default 1 4 Lines of text (<11) to be read in; printed on results file (line 2). /nh/ 1 Example No. 3: pi+ spectra from 562.5 MeV n + Cu64; 10,000 events. stop

pip1_5F	'e.inf	/File name for diagnostic output. (<31 char.)
pip1_5F	e.res	/File name for results of calculation. (<31 char.)
pipl	/pname/	projectile particle name:
	pı	cot - proton, neut - neutron, pipl - pi+, pimi - pi-, pize - pi0,
	ga	amm - gamma with fixed energy, gamb - bremss. gamma, stop - no more calc.
1500.0	/tOmev/	minimum (initial) projectile kinetic energy in MeV; [tgmin for gamb]
56.	/anucl/	target mass number
26.	/znucl/	target atomic number
10000	/limc/	total number of inelastic events, normally 2000-500000
-10.	/dt0/	projectile kinetic energy step-size in MeV [Only 1 energy if <0.]
1600.5	/tOmax/	maximum (final) projectile kinetic energy in MeV, [tgmax for gamb]
10.	/dteta/	<pre>step-size (degrees) in ejectile angular distributions [mang > 0]</pre>
0	/mspec/	(0/1,2) if ejectile energy spectra (are not/are) needed
1	/mpyld/	(0/1,2) if particle yield tables (are not/are) needed
0	/mchy/	(0/1) if particle channel yields (are not/are) needed
0	/misy/	(0/1,2,3) if isotope yields (are not/are) needed
1	/mdubl/	(0/1,2) if double differential spectra (are not/are) needed
0	/mang/	(0/1,2) if angular distributions (are not/are) needed
11 /i	par1,ipar	$c^2/$ range of ejectile types for spectrum calcs. [Below, ang. bins for mdubl > 0]
25.0 35	5.0 85.0 9	95.0 145.0 155.0 125.0 -5.0 65.0 75.0 85.0 95.0 105.0 115.0 125.0 135.0 145.0 155.0
0. 10.	1. 10. 10	00. 10. 100. 1500. 100. 1500. 5000. 200. /tmin, tmax, dt, j=1-4/
6 /	'nevtype/	number of evaporated particle types (see table in bldatgem.f).
3	/ityp/	Version of the random no. generator used; 1-7 OK; default 1
1	/nh/	Lines of text (<11) to be read in; printed on results file (line 2).
Example	e No. 4: N	Neutron spectra from 1.5 GeV pi+ + Fe56; 10,000 events.
stop		

```
nAu6.inf
                                /File name for diagnostic output. (<31 char.)
nAu6.res
                                /File name for results of calculation. (<31 char.)
neut
       /pname/ projectile particle name:
             prot - proton, neut - neutron, pipl - pi+, pimi - pi-, pize - pi0,
             gamm - gamma with fixed energy, gamb - bremss. gamma, stop - no more calc.
30.0 /tOmev/ minimum (initial) projectile kinetic energy in MeV; [tgmin for gamb]
197.
        /anucl/ target mass number
       /znucl/ target atomic number
79.
10000 /limc/ total number of inelastic events, normally 2000-500000
10.
      /dt0/
               projectile kinetic energy step-size in MeV [Only 1 energy if <0.]
300.5 /tOmax/ maximum (final) projectile kinetic energy in MeV, [tgmax for gamb]
      /dteta/ step-size (degrees) in ejectile angular distributions [mang > 0]
10.
      /mspec/ (0/1,2) if ejectile energy spectra (are not/are) needed
0
      /mpyld/ (0/1,2) if particle yield tables (are not/are) needed
1
0
      /mchy/
               (0/1) if particle channel yields (are not/are) needed
0
                (0/1,2,3) if isotope yields (are not/are) needed
      /misy/
0
      /mdubl/ (0/1,2) if double differential spectra (are not/are) needed
               (0/1,2) if angular distributions (are not/are) needed
0
      /mang/
2 2 /ipar1,ipar2/ range of ejectile types for spectrum calcs. [Below, ang. bins for mdubl > 0]
25.0 35.0 45.0 55.0 65.0 75.0 -55.0 165.0 55.0 65.0 75.0 85.0 95.0 105.0 115.0 125.0 135.0 145.0 155.0 165.0
0. 500. 10. 500. 600. 10. 600. 700. 10. 700. 5000. 20. /tmin, tmax, dt, j=1-4/ \,
     /nevtype/ number of evaporated particle types (see table in bldatgem.f).
6
       /ityp/ \, Version of the random no. generator used; 1-7 OK; default 1 \,
2
                Lines of text (<11) to be read in; printed on results file (line 2).
2
        /nh/
Example No. 5: Fission cross section of Au197 bombarded with
neutrons from 30 to 300 MeV with a step of 10 MeV; 10,000 events.
stop
```

p62_9Pb6.inf	/File name for diagnostic output. (<31 char.)
p62_9Pb6.res	/File name for results of calculation. (<31 char.)
prot /pname/ p	rojectile particle name:
pro	t - proton, neut - neutron, pipl - pi+, pimi - pi-, pize - pi0,
gam	n - gamma with fixed energy, gamb - bremss. gamma, stop - no more calc.
62.9 /tOmev/ mi	nimum (initial) projectile kinetic energy in MeV; [tgmin for gamb]
208. /anucl/	target mass number
82. /znucl/ t	arget atomic number
10000 /limc/ t	otal number of inelastic events, normally 2000-500000
-10. /dt0/	projectile kinetic energy step-size in MeV [Only 1 energy if <0.]
200.5 /tOmax/ m	aximum (final) projectile kinetic energy in MeV, [tgmax for gamb]
10. /dteta/ s	tep-size (degrees) in ejectile angular distributions [mang > 0]
1 /mspec/ (0/1,2) if ejectile energy spectra (are not/are) needed
1 /mpyld/ (0/1,2) if particle yield tables (are not/are) needed
0 /mchy/ (0/1) if particle channel yields (are not/are) needed
0 /misy/ (0/1,2,3) if isotope yields (are not/are) needed
1 /mdubl/ (0/1,2) if double differential spectra (are not/are) needed
1 /mang/ (0/1,2) if angular distributions (are not/are) needed
1 6 /ipar1,ipar2	/ range of ejectile types for spectrum calcs. [Below, ang. bins for mdubl $>$ 0]
22.5 27.5 52.5 57	.5 72.5 77.5 92.5 97.5 112.5 117.5 152.5 157.5 -5.0 105.0 115.0 125.0 135.0 145.0 155.0 165.0
0. 22. 1. 22. 120	. 2. 120. 400. 10. 400. 1000. 20. /tmin, tmax, dt, j=1-4/
6 /nevtype/ n	umber of evaporated particle types (see table in bldatgem.f).
6 /ityp/	Version of the random no. generator used; 1-7 OK; default 1
2 /nh/	Lines of text (<11) to be read in; printed on results file (line 2).
Example No. 6: En	ergy, angular, and double-differential spectra
of n to He from 6	2.9 MeV p + Pb208; 100,000 events;
stop	

```
p800Au6.inf
                               /File name for diagnostic output. (<31 char.)
p800Au6.res
                               /File name for results of calculation. (<31 char.)
prot
       /pname/ projectile particle name:
             prot - proton, neut - neutron, pipl - pi+, pimi - pi-, pize - pi0,
             gamm - gamma with fixed energy, gamb - bremss. gamma, stop - no more calc.
800.0 /tOmev/ minimum (initial) projectile kinetic energy in MeV; [tgmin for gamb]
197.
       /anucl/ target mass number
       /znucl/ target atomic number
79.
10000 /limc/ total number of inelastic events, normally 2000-500000
-20.
       /dt0/
               projectile kinetic energy step-size in MeV [Only 1 energy if <0.]
1000.5 /tOmax/ maximum (final) projectile kinetic energy in MeV, [tgmax for gamb]
      /dteta/ step-size (degrees) in ejectile angular distributions [mang > 0]
10.
       /mspec/ (0/1,2) if ejectile energy spectra (are not/are) needed
0
      /mpyld/ (0/1,2) if particle yield tables (are not/are) needed
1
0
      /mchy/
               (0/1) if particle channel yields (are not/are) needed
      /misy/
                (0/1,2,3) if isotope yields (are not/are) needed
1
0
      /mdubl/
               (0/1,2) if double differential spectra (are not/are) needed
               (0/1,2) if angular distributions (are not/are) needed
0
      /mang/
2 2 /ipar1,ipar2/ range of ejectile types for spectrum calcs. [Below, ang. bins for mdubl > 0]
25.0 35.0 45.0 55.0 65.0 75.0 -55.0 165.0 55.0 65.0 75.0 85.0 95.0 105.0 115.0 125.0 135.0 145.0 155.0 165.0
0. 500. 10. 500. 600. 10. 600. 700. 10. 700. 5000. 20. /tmin, tmax, dt, j=1-4/ \,
     /nevtype/ number of evaporated particle types (see table in bldatgem.f).
6
        /ityp/ \, Version of the random no. generator used; 1-7 OK; default 1 \,
7
                Lines of text (<11) to be read in; printed on results file (line 2).
2
        /nh/
Example No. 7: xsec and kinetic energy of all products measured
at GSI in inverse kinematics for 800 MeV p + Au197; 10,000 events.
stop
```

p1000Fe	e6.inf	/File name for diagnostic output. (<31 char.)
p1000Fe	6.res	/File name for results of calculation. (<31 char.)
prot	/pname/	projectile particle name:
	p	rot - proton, neut - neutron, pipl - pi+, pimi - pi-, pize - pi0,
	g	amm - gamma with fixed energy, gamb - bremss. gamma, stop - no more calc.
1000.0	/tOmev/	minimum (initial) projectile kinetic energy in MeV; [tgmin for gamb]
56.	/anucl/	target mass number
26.	/znucl/	target atomic number
10000	/limc/	total number of inelastic events, normally 2000-500000
-20.	/dt0/	projectile kinetic energy step-size in MeV [Only 1 energy if <0.]
1000.5	/tOmax/	maximum (final) projectile kinetic energy in MeV, [tgmax for gamb]
10.	/dteta/	step-size (degrees) in ejectile angular distributions $[mang > 0]$
0	/mspec/	(0/1,2) if ejectile energy spectra (are not/are) needed
1	/mpyld/	(0/1,3) if particle yield tables (are not/are) needed
0	/mchy/	(0/1) if particle channel yields (are not/are) needed
3	/misy/	(0/1,2,3) if isotope yields (are not/are) needed
0	/mdubl/	(0/1,2) if double differential spectra (are not/are) needed
0	/mang/	(0/1,2) if angular distributions (are not/are) needed
22/i	.par1,ipa	r^{2} range of ejectile types for spectrum calcs. [Below, ang. bins for mdubl > 0]
25.0 35	5.0 45.0	55.0 65.0 75.0 -55.0 165.0 55.0 65.0 75.0 85.0 95.0 105.0 115.0 125.0 135.0 145.0 155.0 165.0
0. 500.	10. 500	. 600. 10. 600. 700. 10. 700. 5000. 20. /tmin, tmax, dt, j=1-4/
6 /	'nevtype/	number of evaporated particle types (see table in bldatgem.f).
5	/ityp/	Version of the random no. generator used; 1-7 OK; default 1
3	/nh/	Lines of text (<11) to be read in; printed on results file (line 2).
Example	e No. 8: '	Yields, mean kinetic energy, angles of emission,
and muc	h more (the most complete output) of all products measured
at GSI	in inver	se kinematics for 1000 MeV p + Fe56; 10,000 events.
stop		

```
g300Cu.inf
                               /File name for diagnostic output. (<31 char.)
g300Cu.res
                               /File name for results of calculation. (<31 char.)
gamm
       /pname/ projectile particle name:
             prot - proton, neut - neutron, pipl - pi+, pimi - pi-, pize - pi0,
             gamm - gamma with fixed energy, gamb - bremss. gamma, stop - no more calc.
300.0 /tOmev/ minimum (initial) projectile kinetic energy in MeV; [tgmin for gamb]
64.
       /anucl/ target mass number
       /znucl/ target atomic number
29.
10000 /limc/ total number of inelastic events, normally 2000-500000
-5.
      /dt0/
               projectile kinetic energy step-size in MeV [Only 1 energy if <0.]
300.5 /tOmax/ maximum (final) projectile kinetic energy in MeV, [tgmax for gamb]
      /dteta/ step-size (degrees) in ejectile angular distributions [mang > 0]
10.
      /mspec/ (0/1,2) if ejectile energy spectra (are not/are) needed
0
      /mpyld/ (0/1,2) if particle yield tables (are not/are) needed
1
0
      /mchy/
               (0/1) if particle channel yields (are not/are) needed
0
      /misy/
               (0/1,2,3) if isotope yields (are not/are) needed
1
      /mdubl/ (0/1,2) if double differential spectra (are not/are) needed
               (0/1,2) if angular distributions (are not/are) needed
0
      /mang/
2 2 /ipar1,ipar2/ range of ejectile types for spectrum calcs. [Below, ang. bins for mdubl > 0]
42.5 47.5 87.5 92.5 132.5 137.5 -55.0 165.0 55.0 65.0 75.0 85.0 95.0 105.0 115.0 125.0 135.0 145.0 155.0 165.0
0. 22. 1. 22. 120. 2. 120. 400. 10. 400. 1000. 20. /tmin, tmax, dt, j=1-4/
     /nevtype/ number of evaporated particle types (see table in bldatgem.f).
6
       /ityp/ \, Version of the random no. generator used; 1-7 OK; default 1 \,
3
                Lines of text (<11) to be read in; printed on results file (line 2).
2
       /nh/
Example No. 9: Proton spectra from monochromatic 300 MeV gamma + 64Cu;
10,000 events.
stop
```

b1000Au6.inf /File name for diagnostic output. (<31 char.)
b1000Au6.res /File name for results of calculation. (<31 char.)
gamb /pname/ projectile particle name:
prot - proton, neut - neutron, pipl - pi+, pimi - pi-, pize - pi0,
gamm - gamma with fixed energy, gamb - bremss. gamma, stop - no more calc.
30.0 /tOmev/ minimum (initial) projectile kinetic energy in MeV; [tgmin for gamb]
197. /anucl/ target mass number
79. /znucl/ target atomic number
10000 /limc/ total number of inelastic events, normally 2000-500000
-20. /dt0/ projectile kinetic energy step-size in MeV [Only 1 energy if <0.]
1000.0 /tOmax/ maximum (final) projectile kinetic energy in MeV, [tgmax for gamb]
10. /dteta/ step-size (degrees) in ejectile angular distributions [mang > 0]
0 /mspec/ (0/1,2) if ejectile energy spectra (are not/are) needed
1 /mpyld/ (0/1,2) if particle yield tables (are not/are) needed
0 /mchy/ (0/1) if particle channel yields (are not/are) needed
3 /misy/ (0/1,2,3) if isotope yields (are not/are) needed
0 /mdubl/ (0/1,2) if double differential spectra (are not/are) needed
0 /mang/ (0/1,2) if angular distributions (are not/are) needed
2 2 /ipar1,ipar2/ range of ejectile types for spectrum calcs. [Below, ang. bins for mdubl > 0]
25.0 35.0 45.0 55.0 65.0 75.0 -55.0 165.0 55.0 65.0 75.0 85.0 95.0 105.0 115.0 125.0 135.0 145.0 155.0 165.0
0. 500. 10. 500. 600. 10. 600. 700. 10. 700. 5000. 20. /tmin, tmax, dt, j=1-4/
6 /nevtype/ number of evaporated particle types (see table in bldatgem.f).
3 /ityp/ Version of the random no. generator used; 1-7 OK; default 1
4 /nh/ Lines of text (<11) to be read in; printed on results file (line 2).
Example No. 10: Yields, mean kinetic energy, emission angles,
neutron multiplicities, Forward/Backward ratios, and much more
(the most complete output) of all products from E_max = 1000 MeV
bremsstrahlung gammas + 197Au; 10,000 events.
stop

Appendix 2

To save space, parts of long tables shown in all 10 examples of the CEM03.03 output are deleted and replaced here with dashed lines, keeping only the first two and the last two lines of each table. We also deleted the Copyright Notice and the information about the level-density parameter used in the preequilibrium calculations in all examples except the first one.

CEM03.03 Output Example 1

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Wed Feb 1 10:54:48 2012

Example No. 1: Proton spectra from 500 MeV $\rm p$ + Ni58; 10,000 events. Number of types of evaporated particles = 6

M TO A Z Q B limc idel 0.9383 0.5000 58. 28. 1 1 10000 1

dt0 = -20.0, t0max = 500.5, dteta = 10.0

rOm = 1.2, & cevap = 12.0.

 Theta1
 Theta2
 Theta3
 Theta4
 Theta5
 Theta6

 60.0
 70.0
 85.0
 95.0
 115.0
 125.0
 155.0
 165.0
 -5.0
 65.0
 75.0
 85.0

Theta7Theta8Theta9Theta1095.0105.0115.0125.0135.0145.0155.0165.0

Tmin, Tmax, dT{1};Tmin, Tmax, dT{2};Tmin, Tmax, dT{3};Tmin, Tmax, dT{4}. 0.0 22.0 1.00 22.0 100.0 3.00 100. 500. 10.0 500. 5000. 200.

lim = 100000 .

Geometrical cross section = 1387.69 mb. Calculation takes into account fission and evaporation processes using Furihata-s GEM2 code. The following level density parameters were used in the preequilibrium part of this calculation: a(Z,N,E) was calculated with Moller, Nix, Myers & Swiatecki microscopic corrections; [Atomic Data Nucl. Data Tables, 59, 185 (1995)]; Level density is from a shifted Fermi-gas formula, with the shift given by 0, delta-p, delta-n, or delta-p + delta-n, for odd-odd; odd-n, even-p; odd-p, even-n; and even-even nuclei, respectively for the compound nucleus, and similarly using deltaM-p and deltaM-n for the saddle point. delta-n and delta-p are tabulated by Moller, Nix & Kratz and deltaM-n and deltaM-p are 4.80 MeV * Bs * {1/N**(1/3) or 1/Z**(1/3)}. Bs is the surface area of the saddle-point shape with respect to a sphere.

Inelastic cross section used here = 684.39 mb

Monte Carlo inelastic cross section = 700.61 mb

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500.0 MeV (Z = 1, A = 1) + (Z = 28., A = 58.)

Number of inelastic interactions = 10000. Number of elastic interactions = 9807.

Reaction cross section = 684.39 mb. Elastic cross section = 671.18 mb.

The mean excitation energy, charge, mass, and angular momentum of the 10000 nuclei after the cascade and before preequilibrium decay are: E*av = 84.4 +/- 74.1 MeV; E*min = -3.4; E*max = 459.1 Zav = 27.2 +/- 1.1; Zmin = 21.; Zmax = 30. Aav = 55.8 +/- 1.6; Amin = 49.; Amax = 58. Lav = 5.3 +/- 3.4 h-bar; Lmin = 0.; Lmax = 22. The mean charge, mass, and angular momentum of the 526 residual nuclei with less than 3 MeV of excitation energy after the cascade are: Zav = 28.1 +/- 0.2; Zmin = 28.; Zmax = 29. Aav = 57.1 +/- 0.4; Amin = 55.; Amax = 58. Lav = 2.4 +/- 1.6 h-bar; Lmin = 0.; Lmax = 10. The mean excitation energy, charge, mass, and angular momentum of the $$9474\ nuclei$ after preequilibrium decay and before the start of statistical decay are: E*av = 72.5 +/- 62.0 MeV; E*min = 0.8; E*max = 444.7Zav = 26.7 +/- 1.3; Zmin = 18.; Zmax = 30. Aav = 55.0 +/- 2.3; Amin = 42.; Amax = 58. Lav = 6.1 +/- 4.2 h-bar; Lmin = 0.; Lmax = 34.

 The mean kinetic energy, charge, mass, and angular momentum of the
 10000 residual nuclei are:

 Ekav =
 2.6 +/- 3.4 MeV; Ekmin =
 0.0; Ekmax =
 41.7

 Zav =
 23.7 +/- 3.4; Zmin =
 9.; Zmax =
 29.

 Aav =
 49.5 +/- 6.8; Amin =
 18.; Amax =
 58.

 Lav =
 5.9 +/- 4.2 h-bar; Lmin =
 0.; Lmax =
 34.

 Number of coalesced d, t, He3, He4 = 2371 423 159 364 Mean multiplicities, yields, and mean energies of ejected particles: (Notation: T - all production mechanisms, C - cascade, P - pre-equilibrium, Sp - from spallation residues, Pf - from nuclei before fission, F - from fission fragments, E - total evaporation = Sp + Pf + F, Co - Coalescence from cascade; Values which are identically zero are not printed.
 Part.
 Multiplicities
 Yields
 [mb]
 <TKE>
 [MeV]
 2.4439 +/- 0.0156 1672.582 +/- 10.699 1.0507 +/- 0.0103 719.089 +/- 7.015 0.1001 +/- 0.0032 68.507 +/- 2.165 Tn 55.04 C n P n 120.68 17.56 1.2931 +/- 0.0114 1.2931 +/- 0.0114 884.985 +/- 7.783 884.985 +/- 7.783 4.61 Sp n Ēn 4 61 ***** ***** ****** ***** 2392.355 +/- 12.796 964.580 +/- 8.125 3.4956 +/- 0.0187 1.4094 +/- 0.0119 Тр 66.35 С р 151.94 0.1493 +/- 0.0039 102.180 +/-Рp 2.644 19.94 1325.596 +/-1.9369 +/- 0.0139 9.525 Spp 7.65 1.9369 +/- 0.0139 1325.596 +/- 9.525 Е р 7.65 ***** ***** ***** ***** T d 0.5766 + / - 0.0076394.620 +/- 5.197 31.13 60.500 +/-0.0884 +/- 0.0030 P d 2.035 22.84 0.2511 +/- 0.0050 171.850 +/-3.429 9.71 Sp d Ēd 0.2511 +/- 0.0050 171.850 +/-3,429 9 71 0.2371 +/- 0.0049 162.269 +/-Co d 3.332 56.91 ****** ****** ***** T t 0.1015 + / - 0.003269.466 +/- 2.180 24.01 0.0250 +/- 0.0016 17.110 +/-27.67 P t 1.082 0.0342 +/- 0.0018 23.406 +/-1.266 10.44 Sp t Ēt 0.0342 + / - 0.001823.406 + / -1.266 10.44 28.950 +/-0.0423 +/- 0.0021 Co t 1.408 32.81 ****** ***** ****** 76.241 +/- 2.284 0.1114 +/- 0.0033 T He3 26.77

0.0462 +/- 0.0021 0.0462 +/- 0.0021 31.619 +/- 1.471 E He3 13.61 0.0364 +/- 0.0019 CoHe3 24.912 +/-1.306 40.53 ****** ***** ***** 0.4321 +/- 0.0066 295.725 +/- 4.499 T He4 14.08 0.0233 +/- 0.0015 P He4 15.946 +/-1.045 40.15 SpHe4 0.3929 + / - 0.0063268.897 +/-4,290 11.91 E He4 0.3929 +/- 0.0063 268.897 +/-4.290 11.91 CoHe4 0.0159 +/- 0.0013 10.882 +/-0.863 29.37 14.646 +/- 1.001 40.447 +/- 1.664 0.0214 +/- 0.0015 pi-49.69 pi0 0.0591 +/- 0.0024 55 57 0.0439 + / - 0.002130.045 +/- 1.434 pi+ 0.0439 +/- 0.0021 30.045 +/- 1.434 50.98

19.710 +/-

31.619 +/-

1.161

1.471

30.48

13.61

Double differential cross sections [mb/MeV/sr];

0.0288 +/- 0.0017

P He3

SpHe3

Lab. angle = 60.0 to 70.0 degrees.

 Tp
 [MeV]
 Total
 Cascade
 Precompound
 Total Evaporation

 1.0 2.0
 6.895E-02 +/- 6.89E-02 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00
 6.895E-02 +/- 6.89E-02

 2.0 3.0
 2.068E+00 +/- 3.78E-01
 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00
 2.068E+00 +/- 3.78E-01

 290.0 300.0
 6.895E-03 +/- 6.89E-03 6.895E-03 +/- 6.89E-03
 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00

 300.0 310.0
 6.895E-03 +/- 6.89E-03 6.895E-03 +/- 6.89E-03
 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00

Integrated: 2.155E+02 +/- 3.85E+00 9.680E+01 +/- 2.58E+00 9.653E+00 +/- 8.16E-01 1.090E+02 +/- 2.74E+00

Double differential cross sections [mb/MeV/sr]; Lab. angle = 85.0 to 95.0 degrees.

 Tp
 [MeV]
 Total
 Cascade
 Precompound
 Total Evaporation

 1.0 2.0
 4.374E-01 +/- 1.65E-01 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 4.374E-01 +/- 1.65E-01

 2.0 3.0
 2.562E+00 +/- 4.00E-01 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 2.562E+00 +/- 4.00E-01

 220.0 230.0
 6.249E-03 +/- 6.25E-03 6.249E-03 +/- 6.25E-03 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00

 230.0 240.0
 6.249E-03 +/- 6.25E-03 6.249E-03 +/- 6.25E-03 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00

Integrated: 1.521E+02 +/- 3.08E+00 3.868E+01 +/- 1.55E+00 8.186E+00 +/- 7.15E-01 1.052E+02 +/- 2.56E+00

Double differential cross sections [mb/MeV/sr]; Lab. angle = 115.0 to 125.0 degrees.

 Tp
 [MeV]
 Total
 Cascade
 Precompound
 Total
 Evaporation

 1.0 2.0
 2.886E-01 +/- 1.44E-01
 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00
 2.886E-01 +/- 1.44E-01

 2.0 3.0
 4.113E+00 +/- 5.45E-01
 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00
 4.100E+00
 4.113E+00 +/- 5.45E-01

 140.0 150.0
 1.443E-02 +/- 1.02E-02
 1.443E-02 +/- 1.02E-02
 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00

 150.0 160.0
 7.216E-03 +/- 7.22E-03
 7.216E-03 +/- 7.22E-03
 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00

Integrated: 1.123E+02 +/- 2.85E+00 1.306E+01 +/- 9.71E-01 6.133E+00 +/- 6.65E-01 9.308E+01 +/- 2.59E+00

Double differential cross sections [mb/MeV/sr]; Lab. angle = 155.0 to 165.0 degrees.

 Tp
 [MeV]
 Total
 Cascade
 Precompound
 Total
 Evaporation

 1.0 2.0
 5.481E-01 +/- 3.16E-01 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 5.481E-01 +/- 3.16E-01
 2.0 3.0
 6.212E+00 +/- 1.07E+00 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 6.212E+00 +/- 1.07E+00

 120.0 130.0
 1.827E-02 +/- 1.83E-02 1.827E-02 +/- 1.83E-02 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00
 1.00E+00 0.000E+00 +/- 0.00E+00

 130.0 1.427E-02 +/- 1.83E-02 1.827E-02 +/- 1.83E-02 0.000E+00 +/- 0.00E+00 +/- 0.00E+00
 1.00E+00 0.000E+00 +/- 0.00E+00

Integrated: 1.063E+02 +/- 4.41E+00 8.222E+00 +/- 1.23E+00 2.375E+00 +/- 6.59E-01 9.574E+01 +/- 4.18E+00

Elapsed cpu time = 0. min and 4.440 sec.

CEM03.03 Output Example 2

Wed Feb 1 10:55:24 2012 Example No. 2: pi0 spectra from 500 MeV pi- + Cu64; 10,000 events. Number of types of evaporated particles M TO A Z Q B limc idel 0.1396 0.5000 64. 29. -1 0 10000 1 dt0 = -20.0, t0max = 500.5, dteta = 10.0mspec mpyld mchy misy mdubl mang ipar1 ipar2 1 1 0 0 0 8 rOm = 1.2, & cevap = 12.0. Theta4 Theta2 Theta3 Theta1 Theta5 Theta6 25.0 35.0 45.0 55.0 65.0 75.0 -55.0 165.0 55.0 65.0 75.0 85.0 Theta8 Theta9 Theta7 Theta10 95.0 105.0 115.0 125.0 135.0 145.0 155.0 165.0 Tmin, Tmax, dT{1};Tmin, Tmax, dT{2};Tmin, Tmax, dT{3};Tmin, Tmax, dT{4}.
0.0 500.0 10.00 500.0 600.0 10.00 600. 700. 10.0 700. 5000. 20. lim = 100000 . Geometrical cross section = 1445.99 mb. Inelastic cross section used here = 667.09 mb Monte Carlo inelastic cross section = 667.09 mb Wed Feb 1 10:55:24 2012 500.0 MeV (Z = -1, A = 0) + (Z = 29., A = 64.)Number of inelastic interactions = 10000. Number of elastic interactions = 11676, Reaction cross section = 667.09 mb, Elastic cross section = 778.90 mb. The mean excitation energy, charge, mass, and angular momentum of the 10000 nuclei after the cascade and before preequilibrium decay are:

 Exav = 104.5 +/- 91.2 MeV; E*min = -3.7; E*max = 525.5

 Zav = 27.7 +/- 1.2; Zmin = 21.; Zmax = 30.

 Aav = 60.8 +/- 2.4; Amin = 52.; Amax = 64.

 Lav = 5.1 +/- 3.1 h-bar; Lmin = 0.; Lmax = 23.

 The mean charge, mass, and angular momentum of the 465 residual nuclei with less than 3 MeV of excitation energy after the cascade are: Zav = 28.9 +/- 0.3; Zmin = 27.; Zmax = 29. Aav = 62.9 +/- 0.5; Amin = 59.; Amax = 64. Lav = 2.5 +/- 1.5 h-bar; Lmin = 0.; Lmax = 11. The mean excitation energy, charge, mass, and angular momentum of the 9535 nuclei after preequilibrium decay and before the start of statistical decay are: decay and before the start of statistical decay are: E*av = 90.7 +/- 79.7 MeV; E*min = 0.4; E*max = 523.5 Zav = 27.2 +/- 1.6; Zmin = 19; Zmax = 30. Aav = 59.9 +/- 3.1; Amin = 45.; Amax = 64. Lav = 6.2 +/- 4.1 h-bar; Lmin = 0.; Lmax = 39. The mean kinetic energy, charge, mass, and angular momentum

 The mean kinetic energy, charge, mass, and angular moment

 of the
 10000 residual nuclei are:

 Ekav =
 2.4 +/- 3.1 MeV; Ekmin = 0.0; Ekmax = 29.4

 Zav =
 24.7 +/- 3.8; Zmin = 8.; Zmax = 29.

 Aav =
 53.2 +/- 8.5; Amin = 16.; Amax = 64.

 Lav =
 6.0 +/- 4.1 h-bar; Lmin = 0.; Lmax = 39.

 2275 Number of coalesced d, t, He3, He4 = 674 151 161 Mean multiplicities, yields, and mean energies of ejected particles: (Notation: T - all production mechanisms, C - cascade, P - pre-equilibrium, Sp - from spallation residues, Pf - from nuclei before fission, F - from fission fragments, E - total evaporation = Sp + Pf + F, Co - Coalescence from cascade; Values which are identically zero are not printed. <TKE> [MeV] Yields [mb] Part. Multiplicities
 4.9853 +/- 0.0223
 3325.656 +/ 14.895

 1.9559 +/- 0.0140
 1304.766 +/ 9.330

 0.1723 +/- 0.0042
 114.940 +/ 2.769

 2.8571 +/- 0.0169
 1905.950 +/ 11.276
 27.27 T n C n 60.84 Рп 18.00 4.84 Sp n En 4.84 30.42 92.48

22.11

Sp p	1.250	3 +/- 0.0	0112	834	. 399	+/-	7.461	8.40	
Ер	1.250	3 +/- 0.0	0112	834	. 399	+/-	7.461	8.40	
******** T d	****** 0 675	***************************************	******	****** 450	**** 487	***** +/-	5 480	********** 04 57	****
P d	0.096	5 +/- 0.0	0031	430	.374	+/-	2.072	24.00	
Sp d	0.354	3 +/- 0.0	060	236	.351	+/-	3.971	10.22	
E d	0.354	3 +/- 0.0	060	236	.351	+/-	3.971	10.22	
Co d	0.224	5 +/- 0.0	0047	149	.762	+/-	3.161	47.46	
******** T +	0 188	********* 5 +/- 0 (*****	******	****' 814	***** +/-	2 897	********** 22 22	****
Pt	0.043	5 +/- 0.0	021	29	.085	+/-	1.393	30.52	
Sp t	0.079	3 +/- 0.0	0028	52	. 900	+/-	1.879	10.59	
Ēt	0.079	3 +/- 0.0	028	52	.900	+/-	1.879	10.59	
Co t	0.065	7 +/- 0.0	026	43	. 828	+/-	1.710	32.64	
********	******	********	*****	******	***** 400	******	*******	**********	****
I He3	0.083	1 +/- 0.0	029	10	.435	+/-	1.923	24.21	
SpHe3	0.039	3 +/- 0.0	011	26	.217	+/-	1.322	14.42	
E He3	0.039	3 +/- 0.0	0020	26	.217	+/-	1.322	14.42	
CoHe3	0.014	7 +/- 0.0	0012	9	. 806	+/-	0.809	37.33	
*******	*****	*******	*****	******	****	*****	******	*******	****
T He4	0.446) +/- 0.0	0067	297	.523	+/-	4.455	14.22	
P ne4	0.025	± +/- 0.0	010	260	072	+/-	1.065	10.07	
E He4	0.404	7 +/- 0.0	004	269	.972	+/-	4.244	12.20	
CoHe4	0.015	+/- 0.0	0013	10	. 607	+/-	0.841	24.52	
******	*****	******	*****	*****	****	*****	*****	******	****
pi-	0.545	5 +/- 0.0	0074	363	. 966	+/-	4.927	281.44	
pi0	0.276	5 +/- 0.0	053	184	.518	+/-	3.508	152.66	
pi+	0.060	2 +/- 0.0	025	40	. 159	+/-	1.637	/8.51	
******	*****	*****	*****	****		*****	*****	*****	****
******	*****	*******	*****	*****	neut	t pion	s *****	******	*****
				Energy	Spee	ctrum	[mb/MeV]	
Energ	gy spe	ctrum fro	om O	.0 to	570	.0 MeV	(zero	values suj	ppressed).
Tri0[M	(eV]	Tota	= Ca	scade					
0.0-	10.0	8.072E-0)1 +/-	7.34E	-02				
10.0-	20.0	1.181E+0	00 +/-	8.88E	-02				
480.0-4	190.0	1.868E-0)1 +/-	3.53E	-02				
490.0- 5	500.0	2.001E-0)2 +/-	1.105	-02				
Integrat	ed:	1.845E+0)2 +/-	3.51E-	+00				
Double di	iffere	ntial cro	oss se	ctions	[mb,	/MeV/s	r];		
Lab. angl	Le = :	25.0 to	35.0	degree	з.				
Troi0 [M	[لام)	Tota	= Ca	ohene					
10.0-	20.0	8.527E-0	$\frac{1}{2} = 0a$	3.22E	-02				
20.0-	30.0	7.309E-0)2 +/-	2.98E	-02				
470.0- 4	180.0	1.096E-0)1 +/-	3.65E-	-02				
480.0- 4	190.0	3.655E-0)2 +/-	2.11E	-02				
Integrat	- od •	3 119F+0)1 +/-	1 956-	+00				
Incegrat	Jeu.	5.11561	/· ·/	1.355	00				
Double di	iffere	ntial cro	oss se	ctions	[mb,	/MeV/s	r];		
Lab. angl	Le =	15.0 to	55.0	degree	з.				
_	_								
Tpi0[M	leV]	Total	L = Ca	scade					
0.0-	10.0	5.566E-0)2 +/-	2.10E	-02				
	20.0	7.156E-0		2.39E	-02				
440.0- 4	150.0	1.590E-0)2 +/-	1.12E	-02				
460.0- 4	170.0	7.951E-0)3 +/-	7.95E	-03				
Integrat									
	ed:	1.558E+0	01 +/-	1.11E	+00				
Doublo di	ed:	1.558E+()1 +/-	1.11E	+00 [mb	/MoV/s	.		
Double di Lab, angl	ed: ifferen Le = 0	1.558E+0 ntial cro)1 +/- oss se 75.0	1.11E ctions degree	+00 [mb, s.	/MeV/s	r];		
Double di Lab. angl	ced: ifferen Le = 0	1.558E+(ntial cro 35.0 to)1 +/- oss se 75.0	1.11E ctions degrees	+00 [mb, s.	/MeV/s	r];		
Double di Lab. angl Tpi0[M	ced: ifferen Le = (MeV]	1.558E+(ntial cro 55.0 to Total	01 +/- oss se 75.0 L = Ca	1.11E ctions degrees scade	+00 [mЪ, s.	/MeV/s	r];		
Double di Lab. angl Tpi0[M 0.0-	ifferen Le = 0 MeV] 10.0	1.558E+(ntial cro 35.0 to Total 7.130E-(01 +/- oss se 75.0 L = Ca 02 +/-	1.11E ctions degrees scade 2.15E	+00 [mb, s.	/MeV/s	r];		
Double di Lab. angl Tpi0[M 0.0- 10.0-	ted: ifferen le = 0 10.0 20.0	1.558E+(ntial cro 55.0 to Total 7.130E-(9.723E-(01 +/- 055 Se 75.0 L = Ca 02 +/- 02 +/-	1.11E ctions degrees scade 2.15E 2.51E	+00 [mb, s. -02 -02	/MeV/s	r];		
Double di Lab. angl Tpi0[M 0.0- 10.0- 	<pre>sed: ifferen le = 0 feV] 10.0 20.0 </pre>	1.558E+(ntial cro 55.0 to Total 7.130E-(9.723E-(01 + / - 055 se 75.0 L = Ca 02 + / - 02 + / - 03 + / -	1.11E ctions degrees scade 2.15E 2.51E 	+00 [mb, s. -02 -02 	/MeV/s	r];		
Double di Lab. angl Tpi0[M 0.0- 10.0- 410.0- 4 420.0- 4	<pre>ced: ifferen le = (10.0 20.0 120.0 130.0</pre>	1.558E+(ntial cro 55.0 to Total 7.130E-(9.723E-(6.482E-(6.482E-(01 +/- 055 se 75.0 L = Ca 02 +/- 02 +/- 03 +/- 03 +/-	1.11E ctions degrees scade 2.15E 2.51E 6.48E 6.48E	+00 [mb, 3. -02 -02 -03 -03 -03	/MeV/s	r];		
Double di Lab. angl Tpi0[M 0.0- 10.0- 410.0- 420.0- 4	<pre>ced: ifferen le = 0 10.0 20.0 420.0 130.0</pre>	1.558E+(ntial cro 55.0 to Total 7.130E-(9.723E-(6.482E-(6.482E-(01 +/- 055 Se 75.0 L = Ca 02 +/- 02 +/- 03 +/- 03 +/-	1.11E- ctions degrees 2.15E- 2.51E- 6.48E- 6.48E-	+00 [mb, s. -02 -02 -03 -03	/MeV/s	r];		
Double di Lab. angl Tpi0[M 0.0- 10.0- 410.0- 420.0- 4 Integrat	<pre>ted: ifferent le = (10.0 20.0 120.0 130.0 ted:</pre>	1.558E+(ntial cro 55.0 to 7.130E-(9.723E-(6.482E-(6.482E-(1.458E+($\begin{array}{l} 01 + / - \\ 058 & \text{se} \\ 75.0 \\ - \\ 02 + / - \\ - \\ 03 + / - \\ 03 + / - \\ 01 + / - \end{array}$	1.11E- ctions degrees scade 2.15E- 2.51E- 6.48E- 6.48E- 9.72E-	+00 [mb, 5. -02 -02 -03 -03 -03	/MeV/s	r];		

Elapsed cpu time = 0. min and 5.228 sec.

CEM03.03 Output Example 3

Wed Feb 1 10:49:14 2012 Example No. 3: pi+ spectra from 562.5 MeV n + Cu64; 10,000 events. Number of types of evaporated particles = 6 M TO A Z Q B limc idel 0.9396 0.5625 64. 29. 0 1 10000 1 dt0 = -10.0, t0max = 600.5, dteta = 10.0mspec mpyld mchy misy mdubl mang ipar1 ipar2 0 1 0 0 0 1 9 rOm = 1.2, & cevap = 12.0. Theta2 Theta1 Theta3 Theta4 Theta5 Theta6 25.0 35.0 55.0 65.0 75.0 85.0 115.0 125.0 -5.0 65.0 75.0 85.0 Theta8 Theta7 Theta9 Theta10 95.0 105.0 115.0 125.0 135.0 145.0 155.0 165.0 Tmin, Tmax, dT{1};Tmin, Tmax, dT{2};Tmin, Tmax, dT{3};Tmin, Tmax, dT{4}. 0.0 500.0 20.00 500.0 600.0 20.00 600. 700. 20.0 700. 5000. lim = 100000 . Geometrical cross section = 1445.99 mb. Inelastic cross section used here = 807.64 mb Monte Carlo inelastic cross section = 759.57 mb Wed Feb 1 10:49:14 2012 562.5 MeV (Z = 0, A = 1) + (Z = 29., A = 64.)Number of inelastic interactions = 10000, Number of elastic interactions = 9037 Reaction cross section = 807.64 mb, Elastic cross section = 729.86 mb. The mean excitation energy, charge, mass, and angular momentum of the 10000 nuclei after the Cascade and before prequilibrium decay are: E*av = 94.2 +/- 79.6 MeV; E*min = -2.2; E*max = 491.0 Zav = 28.0 +/- 1.0; Zmin = 23.; Zmax = 30. Aav = 61.3 +/- 1.9; Amin = 52.; Amax = 65. Lav = 5.8 +/- 3.7 h-bar; Lmin = 0.; Lmax = 25. The mean charge, mass, and angular momentum of the 460 residual nuclei with less than $3~\ensuremath{\text{MeV}}$ of excitation energy after the cascade are: Zav = 29.0 +/- 0.0; Zmin = 28.; Zmax = 29. Aav = 63.0 +/- 0.3; Amin = 60.; Amax = 64. Lav = 2.3 +/- 1.3 h-bar; Lmin = 0.; Lmax = 8. The mean excitation energy, charge, mass, and angular momentum of the 9539 nuclei after preequilibrium decay and before the start of statistical decay are:

 Backary and berote the scale of scalestical decay are.

 E*av = 80.8 +/- 68.3 MeV; E*min = 0.2; E*max = 480.2

 Zav = 27.6 +/- 1.3; Zmin = 21.; Zmax = 30.

 Aav = 60.4 +/- 2.6; Amin = 47.; Amax = 65.

 Lav = 6.6 +/- 4.4 h-bar; Lmin = 0.; Lmax = 32.

 E*av = The mean kinetic energy, charge, mass, and angular momentum The mean kinetic energy, charge, mass, and angular moment of the 10000 residual nuclei are: Ekav = 2.6 +/- 3.6 MeV; Ekmin = 0.0; Ekmax = 42.0 Zav = 25.3 +/- 3.2; Zmin = 11.; Zmax = 30. Aav = 54.4 +/- 7.2; Amin = 24.; Amax = 64. Lav = 6.4 +/- 4.4 h-bar; Lmin = 0.; Lmax = 32. Number of coalesced d, t, He3, He4 = 2439 642 189 169 Mean multiplicities, yields, and mean energies of ejected particles: (Notation: T - all production mechanisms, C - cascade, P - pre-equilibrium, Sp - from spallation residues, Pf - from nuclei before fission, - from fission fragments, E - total evaporation = Sp + Pf + F, Co - Coalescence from cascade: Values which are identically zero are not printed. <TKE> [MeV] Part. Multiplicities Yields [mb] T n 59.51 C n 127.55 P n 17.19 4.45 Sp n 4.45 En *************
 2.0272 +/- 0.0142
 1637.242 +/- 11.499

 0.6935 +/- 0.0083
 560.096 +/- 6.726

 0.1168 +/- 0.0034
 94.332 +/- 2.760
 Т р 53.37 Сp 138.41

Рр

20.76

Sp p	1.2169 +/- 0.0110	982.814 +/-	8.909	8.04
Ер	1.2169 +/- 0.0110	982.814 +/-	8.909	8.04
******	*****	*****	******	*****
Τd	0.6273 +/- 0.0079	506.631 +/-	6.397	32.05
P d	0.0962 +/- 0.0031	77.695 +/-	2.505	23.06
Sp d	0.2875 +/- 0.0054	232.196 +/-	4.330	10.04
Ed	0.2875 +/- 0.0054	232.196 +/-	4.330	10.04
Co d	0.2436 +/- 0.0049	196.740 +/-	3.986	61.58
******	*****	*****	******	*****
Τt	0.1579 +/- 0.0040	127.526 +/-	3.209	24.01
Ρt	0.0369 +/- 0.0019	29.802 +/-	1.551	25.78
Sp t	0.0570 +/- 0.0024	46.035 +/-	1.928	10.59
Εt	0.0570 +/- 0.0024	46.035 +/-	1.928	10.59
Co t	0.0640 +/- 0.0025	51.689 +/-	2.043	34.94
******	*****	*****	******	*****
T He3	0.0769 +/- 0.0028	62.107 +/-	2.240	27.51
P He3	0.0268 +/- 0.0016	21.645 +/-	1.322	28.80
SpHe3	0.0312 +/- 0.0018	25.198 +/-	1.427	14.44
E He3	0.0312 +/- 0.0018	25.198 +/-	1.427	14.44
CoHe3	0.0189 +/- 0.0014	15.264 +/-	1.110	47.24
******	*****	*****	******	*****
T He4	0.4096 +/- 0.0064	330.808 +/-	5.169	14.61
P He4	0.0235 +/- 0.0015	18.979 +/-	1.238	38.69
SpHe4	0.3692 +/- 0.0061	298.180 +/-	4.907	12.50
E He4	0.3692 +/- 0.0061	298.180 +/-	4.907	12.50
CoHe4	0.0169 +/- 0.0013	13.649 +/-	1.050	27.08
******	*****	*****	******	*****
pi-	0.0908 +/- 0.0030	73.333 +/-	2.434	67.85
pi0	0.0869 +/- 0.0029	70.184 +/-	2.381	64.14
pi+	0.0162 +/- 0.0013	13.084 +/-	1.028	56.27
******	*****	*****	******	*****

Double differential cross sections [mb/MeV/sr]; Lab. angle = 25.0 to 35.0 degrees.

Tpi+[MeV] Total = Cascade 40.0- 60.0 7.374E-03 +/- 7.37E-03 100.0- 120.0 7.374E-03 +/- 7.37E-03 180.0- 200.0 1.475E-02 +/- 1.04E-02 220.0- 240.0 7.374E-03 +/- 7.37E-03

Integrated: 1.180E+00 +/- 4.17E-01

Double differential cross sections [mb/MeV/sr]; Lab. angle = 55.0 to 65.0 degrees.

Tpi+	[MeV]	Total = Cascade
0.0-	20.0	1.703E-02 +/- 8.51E-03
20.0-	40.0	1.277E-02 +/- 7.37E-03
40.0-	60.0	1.277E-02 +/- 7.37E-03
100.0-	120.0	4.257E-03 +/- 4.26E-03

Integrated: 9.366E-01 +/- 2.82E-01

Double differential cross sections [mb/MeV/sr]; Lab. angle = 75.0 to 85.0 degrees.

Тр	i+[MeV]	Total = Cascade				
0.	0- 20.0	1.123E-02 +/-	6.48E-03			
20.	0- 40.0	1.123E-02 +/-	6.48E-03			
120. 140.	0- 140.0 0- 160.0	3.744E-03 +/- 3.744E-03 +/-	3.74E-03 3.74E-03			

Integrated: 9.734E-01 +/- 2.70E-01

Double differential cross sections [mb/MeV/sr]; Lab. angle = 115.0 to 125.0 degrees.

Tpi+[MeV] Total = Cascade
 0.0 20.00
 2.980E-02
 +/ 1.13E-02

 20.0 40.0
 1.277E-02
 +/ 7.37E-03

 40.0 60.0
 8.515E-03
 +/ 6.02E-03

Integrated: 1.022E+00 +/- 2.95E-01

Elapsed cpu time = 0. min and 4.865 sec.

CEM03.03 Output Example 4

Wed Feb 1 10:55:39 2012 Example No. 4: Neutron spectra from 1.5 GeV pi+ + Fe56; 10,000 events. Number of types of evaporated particles = 6 M TO A Z Q B limc idel 0.1396 1.5000 56. 26. 1 0 10000 1 dt0 = -10.0, t0max = 1600.5, dteta = 10.0mspec mpyld mchy misy mdubl mang ipar1 ipar2 0 0 1 0 rOm = 1.2, & cevap = 12.0. Theta1 Theta2 Theta3 Theta4 Theta5 Theta6 25.0 35.0 85.0 95.0 145.0 155.0 -15.0 125.0 -5.0 65.0 75.0 85.0 Theta7 Theta8 Theta9 Theta10 95.0 105.0 115.0 125.0 135.0 145.0 155.0 165.0 Tmin, Tmax, dT{1};Tmin, Tmax, dT{2};Tmin, Tmax, dT{3};Tmin, Tmax, dT{4}. 0.0 10.0 1.00 10.0 10.0 10.00 100. 1500. 100.0 1500. 5000. 200. 100000 lim = Geometrical cross section = 1367.65 mb. Inelastic cross section used here = 678.77 mb Monte Carlo inelastic cross section = 678.77 mb Wed Feb 1 10:55:39 2012 1500.0 MeV (Z = 1, A = 0) + (Z = 26., A = 56.) Number of inelastic interactions = 10000. Number of elastic interactions = 10149, Reaction cross section = 678.77 mb, Elastic cross section = 688.88 mb. The mean excitation energy, charge, mass, and angular momentum of the 10000 nuclei after the cascade and before preequilibrium decay are:

 Exave and before preddillorium decay are:

 Exav = 214.9 +/-167.5 MeV;
 E*min = -3.5;
 E*max = 967.8

 Zav = 23.9 +/ 1.9;
 Zmin = 15.;
 Zmax = 29.

 Aav = 50.2 +/ 4.1;
 Amin = 35.;
 Amax = 56.

 Lav = 7.1 +/ 4.5 h-bar;
 Lmin = 0.;
 Lmax = 28.

 The program called Fermi breakup 620 times. The mean charge, mass, and angular momentum of the 172 residual nuclei with less than $3\ {\rm MeV}$ of excitation energy after the cascade are: Zav = 26.0 +/- 0.2; Zmin = 26.; Zmax = 27. Aav = 55.1 +/- 0.5; Amin = 52.; Amax = 56. Lav = 2.5 +/- 1.6 h-bar; Lmin = 0.; Lmax = 10. The mean excitation energy, charge, mass, and angular momentum of the 9828 nuclei after preequilibrium decay and before the start of statistical decay are:

 Exav = 192.0 +/- 161.7 MeV; E*min = 0.1; E*max = 963.4

 Zav = 23.4 +/- 2.1; Zmin = 14.; Zmax = 29.

 Aav = 49.1 +/- 4.5; Amin = 32.; Amax = 56.

 Lav = 8.2 +/- 5.3 h-bar; Lmin = 0.; Lmax = 33.

 The mean kinetic energy, charge, mass, and angular momentum The mean kinetic energy, charge, mass, and angular moment of the 10000 residual nuclei are: Ekav = 5.2 +/- 6.5 MeV; Ekmin = 0.0; Ekmax = 70.7 Zav = 17.8 +/- 5.7; Zmin = 6.; Zmax = 27. Aav = 37.6 +/- 12.3; Amin = 13.; Amax = 56. Lav = 8.1 +/- 5.3 h-bar; Lmin = 0.; Lmax = 33. Number of coalesced d, t, He3, He4 = 681 3844 360 217 Mean multiplicities, yields, and mean energies of ejected particles: (Notation: T - all production mechanisms, C - cascade, P - pre-equilibrium, Sp - from spallation residues, Pf - from nuclei before fission, F - from fission fragments, E - total evaporation = Sp + Pf + F, Co - Coalescence from cascade: Values which are identically zero are not printed. Yields [mb] <TKE> [MeV] Multiplicities Part.

Тр	5.0712	+/- 0	0.0225	3442.179	+/-	15.285	56.71
Ср	2.0809	+/- 0	0.0144	1412.453	+/-	9.791	120.52
Рр	0.1391	+/- 0	.0037	94.417	+/-	2.532	24.84
Spp	2.8512	+/- 0	0.0169	1935.309	+/-	11.461	11.70
Ер	2.8512	+/- 0	0.0169	1935.309	+/-	11.461	11.70
******	******	*****	******	********	****	******	****
Τd	1.7013	+/- 0	.0130	1154.792	+/-	8.853	22.21
Ρd	0.1097	+/- 0	.0033	74.461	+/-	2.248	26.40
Sp d	1.2169	+/- 0	0.0110	825.995	+/-	7.488	13.64
Ed	1.2169	+/- 0	0.0110	825.995	+/-	7.488	13.64
Co d	0.3747	+/- 0	.0061	254.335	+/-	4.155	48.84
******	******	*****	******	*******	****	******	****
Τt	0.3541	+/- 0	.0060	240.352	+/-	4.039	19.70
Ρt	0.0496	+/- 0	.0022	33.667	+/-	1.512	33.61
Spt	0.2377	+/- 0	0.0049	161.344	+/-	3.309	13.49
Ēt	0.2377	+/- 0	0.0049	161.344	+/-	3.309	13.49
Co t	0.0668	+/- 0	.0026	45.342	+/-	1.754	31.47
******	******	*****	******	********	****	******	****
T He3	0.3383	+/- 0	.0058	229.628	+/-	3.948	22.33
P He3	0.0533	+/- 0	.0023	36.178	+/-	1.567	35.83
SpHe3	0.2499	+/- 0	.0050	169.625	+/-	3.393	17.09
E He3	0.2499	+/- 0	.0050	169.625	+/-	3.393	17.09
CoHe3	0.0351	+/- 0	0.0019	23.825	+/-	1.272	39.15
******	******	*****	******	********	****	******	****
T He4	0.9033	+/- 0	.0095	613.133	+/-	6.451	14.59
P He4	0.0470	+/- 0	.0022	31.902	+/-	1.472	48.55
SpHe4	0.8353	+/- 0	0.0091	566.977	+/-	6.204	12.38
E He4	0.8353	+/- 0	.0091	566.977	+/-	6.204	12.38
CoHe4	0.0210	+/- 0	0.0014	14.254	+/-	0.984	26.27
******	******	*****	******	******	****	******	****
pi-	0.3448	+/- 0	0.0059	234.040	+/-	3.986	218.77
pi0	0.6841	+/- 0	.0083	464.347	+/-	5.614	235.69
pi+	0.7787	+/- 0	.0088	528.558	+/-	5.990	463.33
******	*******			*********		*******	************

Double differential cross sections [mb/MeV/sr]; Lab. angle = 25.0 to 35.0 degrees.

	Tn	[MeV]	Total	Cascade	Precompound	Total Evaporation
	0.0-	· 1.0	1.463E+01 +/- 1.35E+00	0.000E+00 +/- 0.00E+00	0.000E+00 +/- 0.00E+00	1.463E+01 +/- 1.35E+00
	1.0-	- 2.0	2.157E+01 +/- 1.64E+00	3.842E+00 +/- 6.90E-01	0.000E+00 +/- 0.00E+00	1.772E+01 +/- 1.48E+00
9	00.0-	1000.0	9.916E-03 +/- 3.51E-03	9.916E-03 +/- 3.51E-03	0.000E+00 +/- 0.00E+00	0.000E+00 +/- 0.00E+00
10	00.0-	1100.0	2.479E-03 +/- 1.75E-03	2.479E-03 +/- 1.75E-03	0.000E+00 +/- 0.00E+00	0.000E+00 +/- 0.00E+00

Integrated: 5.045E+02 +/- 7.91E+00 2.716E+02 +/- 5.80E+00 8.553E+00 +/- 1.03E+00 2.243E+02 +/- 5.27E+00

Double differential cross sections [mb/MeV/sr]; Lab. angle = 85.0 to 95.0 degrees.

 Tn
 [MeV]
 Total
 Cascade
 Precompound
 Total Evaporation

 0.0 1.0
 1.407E+01 +/- 9.34E-01 0.000E+00 +/- 0.00E+00
 6.198E-02 +/- 6.20E-02 1.401E+01 +/- 9.32E-01

 1.0 2.0
 2.262E+01 +/- 1.18E+00 3.842E+00 +/- 4.88E-01 1.40E+01 +/- 8.76E-02 1.865E+01 +/- 1.08E+00

 200.0 300.0
 2.541E-02 +/- 3.97E-03 2.541E-02 +/- 3.97E-03 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00

 400.0 500.0
 6.198E-04 +/- 6.20E-04 6.198E-04 +/- 6.20E-04 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00

Integrated: 2.974E+02 +/- 4.29E+00 1.147E+02 +/- 2.67E+00 7.003E+00 +/- 6.59E-01 1.757E+02 +/- 3.30E+00

Double differential cross sections [mb/MeV/sr]; Lab. angle = 145.0 to 155.0 degrees.

 Tn
 [MeV]
 Total
 Cascade
 Precompound
 Total Evaporation

 0.0 1.0
 1.240E+01 +/- 1.24E+00
 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00
 1.240E+01 +/- 1.24E+00

 1.0 2.0
 1.958E+01 +/- 1.56E+00
 2.479E+00 +/- 5.54E-01
 2.479E-01 +/- 1.56E+01 +/- 1.45E+00

 200.0 300.0
 3.719E-03 +/- 2.15E-03
 3.719E-03 +/- 2.15E-03
 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00

 300.0 400.0
 1.240E-03 +/- 1.24E-03
 1.240E-03 +/- 1.24E-03
 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00

Integrated: 2.372E+02 +/- 5.42E+00 7.425E+01 +/- 3.03E+00 4.710E+00 +/- 7.64E-01 1.583E+02 +/- 4.43E+00

The program called Fermi breakup 620 times.

Elapsed cpu time = 0. min and 8.309 sec.
Wed Feb 1 10:49:23 2012 Example No. 5: Fission cross section of Au197 bombarded with neutrons from 30 to 300 MeV with a step of 10 MeV; 10,000 events. Number of types of evaporated particles = 6 M TO A Z Q B limc idel 0.9396 0.0300 197. 79. 0 1 10000 1 dt0 = 10.0, t0max = 300.5, dteta = 10.0 mspec mpyld mchy misy mdubl mang ipar1 ipar2 ٥ ٥ 0 0 0 rOm = 1.2, & cevap = 12.0. lim = 100000 . Geometrical cross section = 2394.46 mb. Inelastic cross section used here = 2333.73 mb Monte Carlo inelastic cross section = 1647.94 mb Wed Feb 1 10:49:23 2012 30.0 MeV (Z = 0, A = 1) + (Z = 79., A = 197.) Number of inelastic interactions = 10000. Number of elastic interactions = 4530, Reaction cross section = 2333.73 mb. Elastic cross section = 1057.18 mb. The mean excitation energy, charge, mass, and angular momentum 10000 nuclei after the of the cascade and before preequilibrium decay are:

 Exave 34.9 +/- 5.0 MeV; E*min = -0.1; E*max = 36.3

 Zav = 79.0 +/- 0.0; Zmin = 78.; Zmax = 79.

 Aav = 197.9 +/- 0.3; Amin = 195.; Amax = 198.

 Lav = 6.2 +/- 2.4 h-bar; Lmin = 0.; Lmax = 11.

 The mean charge, mass, and angular momentum of the 34 residual nuclei with less than 3 MeV of excitation energy after the cascade are: Zav = 79.0 +/- 0.0; Zmin = 79.; Zmax = 79. Aav = 196.2 +/- 0.5; Amin = 195.; Amax = 197. Lav = 3.0 +/- 1.4 h-bar; Lmin = 1.; Lmax = 6. The mean excitation energy, charge, mass, and angular momentum of the 9965 nuclei after preequilibrium
 of the
 9965 nuclei after prequilibrium

 decay and before the start of statistical decay are:

 E*av = 28.2 +/ 10.4 MeV; E*min = 0.1; E*max = 36.2

 Zav = 79.0 +/ 0.2; Zmin = 77.; Zmax = 79.

 Aav = 197.6 +/ 0.6; Amin = 193.; Amax = 198.

 Lav =
 6.5 +/ 2.7 h-bar; Lmin =
 0.; Lmax = 20.
 Statistical Weight Functions Method: Fissility = 0.0000, Fission cross section = 1.35863E-02 mb. 1 Number of coalesced d, t, He3, He4 = 0 0 0 Mean multiplicities, yields, and mean energies of ejected particles: (Notation: T - all production mechanisms, C - cascade, P - pre-equilibrium, Sp - from spallation residues, Pf - from nuclei before fission, F - from fission fragments, E - total evaporation = Sp + Pf + F, Co - Coalescence from cascade; Values which are identically zero are not printed. Yields [mb] <TKE> [MeV] Part. Multiplicities
 3.1473 +/- 0.0177
 7344.961 +/- 41.402

 0.0758 +/- 0.0028
 176.897 +/- 6.425

 0.3028 +/- 0.0055
 706.655 +/- 12.842
 Τn 2.64 C n 9.24 0.3028 +/- 0.0055 2.7687 +/- 0.0166 706.655 +/- 12.842 P n 10.63 6461.409 +/- 38.832 6461.409 +/- 38.832 Sp n 1.59 1.59 2.7687 +/- 0.0166 Ēn ******* 77.713 +/- 4.259 1.167 +/- 0.522 0.0333 +/- 0.0018 Т р 17.02 C p 0.0005 +/- 0.0002 12.41 75.846 +/- 4.207 0.700 +/- 0.404 0.700 +/- 0.404 Р 0.0325 +/- 0.0018 17.16 0.0003 +/- 0.0002 Spp 9.15 9.15 Ēp 0.0003 +/- 0.0002 0.0092 +/- 0.0010 21.470 +/- 2.238 0.0091 +/- 0.0010 21.237 +/- 2.226 Τd 17.60 Ρd 17.72

Co d 0.0001 + / - 0.00010.233 + / - 0.2336.90 ****** 5.368 +/- 1.119 5.134 +/- 1.095 0.233 +/- 0.233 0.233 +/- 0.233 T t 13.92 Ρt 14.09 0.0001 +/- 0.0001 Sp t 10.11 0.0001 +/- 0.0001 Ēt 10.11 ****** ******* 6.534 +/- 1.235 5.368 +/- 1.119 1.167 +/- 0.522 1.167 +/- 0.522 T He4 0.0028 +/- 0.0005 23.84 0.0023 +/- 0.0005 0.0005 +/- 0.0002 P He4 24.64 SpHe4 20.16 0.0005 +/- 0.0002 E He4 20.16 **************** Geometrical cross section = 2394.46 mb. Inelastic cross section used here = 2222.93 mb Monte Carlo inelastic cross section = 1636.01 mb Wed Feb 1 10:49:33 2012 40.0 MeV (Z = 0, A = 1) + (Z = 79., A = 197.)Number of inelastic interactions = 10000, Number of elastic interactions = 4636. The mean total fission product kinetic energy after neutron emission is 138.49 MeV. Direct Monte Carlo Simulation Method: Fissility = 0.0002 +/- 0.0001, Fission cross section = 4.44586E-01 +/- 3.14E-01 mb. Statistical Weight Functions Method: Fissility = 0.0000, Fission cross section = 8.69020E-02 mb. 50.0 MeV (Z = 0, A = 1) + (Z = 79., A = 197.) Number of inelastic interactions = 10000. Number of elastic interactions = 5051. Reaction cross section = 2135.39 mb, Elastic cross section = 1078.58 mb. Statistical Weight Functions Method: Fissility = 0.0001, Fission cross section = 1.95606E-01 mb. 60.0 MeV (Z = 0, A = 1) + (Z = 79., A = 197.)Number of inelastic interactions = 10000, Number of elastic interactions = 5308, Reaction cross section = 2066.02 mb. Elastic cross section = 1096.64 mb. The mean total fission product kinetic energy after neutron emission is 139.48 MeV. Direct Monte Carlo Simulation Method: Fissility = 0.0003 +/- 0.0002, Fission cross section = 6.19806E-01 +/- 3.58E-01 mb. Statistical Weight Functions Method: Fissility = 0.0002, Fission cross section = 4.26275E-01 mb. 290.0 MeV (Z = 0, A = 1) + (Z = 79., A = 197.)Number of inelastic interactions = 10000, Number of elastic interactions = 6491. Reaction cross section = 1766.57 mb, Elastic cross section = 1146.68 mb. The mean total fission product kinetic energy after neutron emission is 130.08 MeV. Direct Monte Carlo Simulation Method: Fissility = 0.0132 +/- 0.0011, Fission cross section = 2.33187E+01 +/- 2.03E+00 mb. Statistical Weight Functions Method: Fissility = 0.0130,

Fission cross section = 2.30023E+01 mb.

 $300.0 \ \text{MeV} \ (Z = 0, \ \text{A} = 1) \ + \ (Z = 79., \ \text{A} = 197.)$ Number of inelastic interactions = 10000, Number of elastic interactions = 6420, Reaction cross section = 1767.01 mb, Elastic cross section = 1134.42 mb. The mean excitation energy, charge, mass, and angular momentum

The mean excitation energy, charge, mass, and angular moment of the 10000 nuclei after the cascade and before preequilibrium decay are: E*av = 100.8 +/-67.9 MeV; E*min = -0.4; E*max = 304.4Zav = 78.6 +/-0.6; Zmin = 76.; Zmax = 80.Aav = 195.0 +/-1.3; Amin = 190.; Amax = 198.Lav = 8.2 +/-4.7 h-bar; Lmin = 0.; Lmax = 33.

 The mean kinetic energy, charge, mass, and angular momentum of the
 9825 residual nuclei are:

 Ekav =
 0.8 +/ 0.9 MeV; Ekmin =
 0.0; Ekmax =
 8.0

 Zav =
 77.8 +/ 1.2; Zmin =
 71.; Zmax =
 80.

 Aav =
 186.8 +/ 6.0; Amin =
 166.; Amax =
 197.

 Lav =
 9.1 +/ 5.5 h-bar; Lmin =
 0.; Lmax =
 41.

The mean excitation energy, charge, mass, angular momentum, and fission barrier height of the 175 fissioning nuclei are: E*av = 140.8 +/- 43.9 MeV; E*min = 40.3; E*max = 282.0 Zav = 78.3 +/- 0.8; Zmin = 74.; Zmax = 80. Aav = 190.3 +/- 3.2; Amin = 179.; Amax = 196. Lav = 11.6 +/- 5.1 h-bar; Lmin = 1.; Lmax = 30.Bfav = 18.3 +/- 1.4 MeV; Bfmin = 14.3; Bfmax = 22.2

The mean total fission product kinetic energy after neutron emission is 127.84 MeV.

 $\label{eq:Direct Monte Carlo Simulation Method:} \\ Fissility = 0.0175 +/- 0.0013, \\ Fission cross section = 3.09227E+01 +/- 2.34E+00 \mbox{ mb}. \\ \end{cases}$

 $\label{eq:statistical Weight Functions Method:} Statistical Weight Functions Method: Fissility = 0.0147, \\ Fission cross section = 2.59347E+01 \mbox{ mb}.$

Number of coalesced d, t, He3, He4 = 1064 369 30 30

Mean multiplicities, yields, and mean energies of ejected particles: (Notation: T - all production mechanisms, C - cascade, P - pre-equilibrium, Sp - from spallation residues, Pf - from nuclei before fission, F - from fission fragments, E - total evaporation = Sp + Pf + F, Co - Coalescence from cascade; Values which are identically zero are not printed.

Par	ct.	Multiplicities		Yie	Lds [n	<tke> [MeV]</tke>	
***	******	******	******	********	*****	******	*****
Т	n	9.4214 .	+/- 0.0307	16647.712	+/-	54.237	17.93
С	n	2.4302 .	+/- 0.0156	4294.189	+/-	27.546	59.32
Ρ	n	0.4444 ·	+/- 0.0067	785.259	+/-	11.779	15.55
Sp	n	6.3099 ·	+/- 0.0251	11149.659	+/-	44.386	2.69
Ρf	n	0.0425 ·	+/- 0.0021	75.098	+/-	3.643	4.47
F	n	0.1944 ·	+/- 0.0044	343.507	+/-	7.791	3.73
Е	n	6.5468	+/- 0.0256	11568.264	+/-	45.212	2.73
***	******	******	******	********	*****	******	****
Т	q	0.6631 ·	+/- 0.0081	1171.705	+/-	14.389	53.23
С	p	0.2654 ·	+/- 0.0052	468.965	+/-	9.103	107.15
Ρ	p	0.1789 ·	+/- 0.0042	316.118	+/-	7.474	24.72
Sp	p	0.2136 ·	+/- 0.0046	377.433	+/-	8.167	11.18
Ρf	p	0.0020 ·	+/- 0.0004	3.534	+/-	0.790	11.65
F	p	0.0032 ·	+/- 0.0006	5.654	+/-	1.000	7.91
Е	p	0.2188 ·	+/- 0.0047	386.622	+/-	8.265	11.14
***	******	******	******	********	*****	******	****
Т	d	0.2507 ·	+/- 0.0050	442.990	+/-	8.847	28.52
Р	d	0.0919 ·	+/- 0.0030	162.388	+/-	5.357	26.02
Sp	d	0.0507 ·	+/- 0.0023	89.587	+/-	3.979	11.73
Ρf	d	0.0011 .	+/- 0.0003	1.944	+/-	0.586	13.07
F	d	0.0006 ·	+/- 0.0002	1.060	+/-	0.433	9.95
Е	d	0.0524 ·	+/- 0.0023	92.591	+/-	4.045	11.74
Co	d	0.1064 ·	+/- 0.0033	188.010	+/-	5.764	38.94
***	******	******	******	********	*****	******	****
Т	t	0.0847 ·	+/- 0.0029	149.666	+/-	5.143	22.94
Ρ	t	0.0234 ·	+/- 0.0015	41.348	+/-	2.703	29.50

Sp t	0.0238 +/- 0.0015	42.055 +/-	2.726	12.57
Pf t	0.0006 +/- 0.0002	1.060 +/-	0.433	14.04
Et	0.0244 +/- 0.0016	43.115 +/-	2.760	12.60
Co t	0.0369 +/- 0.0019	65.203 +/-	3.394	25.61
******	*****	*****	******	******
T He3	0.0127 +/- 0.0011	22.441 +/-	1.991	35.92
P He3	0.0093 +/- 0.0010	16.433 +/-	1.704	38.94
SpHe3	0.0004 +/- 0.0002	0.707 +/-	0.353	22.18
E He3	0.0004 +/- 0.0002	0.707 +/-	0.353	22.18
CoHe3	0.0030 +/- 0.0005	5.301 +/-	0.968	28.39
******	*****	*****	******	*****
T He4	0.1056 +/- 0.0032	186.596 +/-	5.742	23.11
P He4	0.0055 +/- 0.0007	9.719 +/-	1.310	40.19
SpHe4	0.0951 +/- 0.0031	168.043 +/-	5.449	22.21
PfHe4	0.0007 +/- 0.0003	1.237 +/-	0.468	22.32
F He4	0.0013 +/- 0.0004	2.297 +/-	0.637	17.04
E He4	0.0971 +/- 0.0031	171.577 +/-	5.506	22.14
CoHe4	0.0030 +/- 0.0005	5.301 +/-	0.968	23.01
******	*****	*****	*******	******
pi-	0.0098 +/- 0.0010	17.317 +/-	1.749	35.23
pi0	0.0036 +/- 0.0006	6.361 +/-	1.060	30.35
******	*****	*****	*****	******

Elapsed cpu time = 2. min and 49.225 sec.

_____ Wed Feb 1 10:55:00 2012 Example No. 6: Energy, angular, and double-differential spectra of n to He from 62.9 MeV p $\,$ + Pb208; 100,000 events; Number of types of evaporated particles = 6 M TO A Z Q B limc idel 0.9383 0.0629 208. 82. 1 1 10000 1 dt0 = -10.0, t0max = 200.5, dteta = 10.0mspec mpyld mchy misy mdubl mang ipar1 ipar2 0 0 1 1 rOm = 1.2, & cevap = 12.0. Theta1 Theta2 Theta3 Theta4 Theta5 Theta6 22.5 27.5 52.5 57.5 72.5 77.5 92.5 97.5 112.5 117.5 152.5 157.5 Theta7 Theta8 Theta9 Theta10 -5.0 105.0 115.0 125.0 135.0 145.0 155.0 165.0
 Tmin, Tmax, dT{1};Tmin, Tmax, dT{2};Tmin, Tmax, dT{3};Tmin, Tmax, dT{4}.

 0.0
 22.0
 1.00
 22.00
 120.0
 2.00
 120.
 400.
 1000.
 20.
 100000 lim = Geometrical cross section = 2457.28 mb. Inelastic cross section used here = 1962.23 mb Monte Carlo inelastic cross section = 1638.74 mb Wed Feb 1 10:55:00 2012 62.9 MeV (Z = 1, A = 1) + (Z = 82., A = 208.)Number of inelastic interactions = 10000, Number of elastic interactions = 4995. Reaction cross section = 1962.23 mb, Elastic cross section = 980.14 mb. The mean excitation energy, charge, mass, and angular momentum of the 10000 nuclei after the cascade and before preequilibrium decay are:

 Exave and before preddillorium decay are:

 Exav = 47.9 +/- 21.9 MeV; E*min = -1.6; E*max = 65.5

 Zav = 82.8 +/- 0.4; Zmin = 81.; Zmax = 83.

 Aav = 208.5 +/- 0.6; Amin = 206.; Amax = 209.

 Lav = 6.6 +/- 3.1 h-bar; Lmin = 0.; Lmax = 16.

 The mean charge, mass, and angular momentum of the 363 residual nuclei with less than 3 MeV of excitation energy after the cascade are:

 Zav = 82.3 +/ 0.4;
 Zmin = 82.;
 Zmax = 83.

 Aav = 207.5 +/ 0.5;
 Amin = 206.;
 Amax = 208.

 Lav = 3.2 +/ 2.0 h-bar;
 Lmin = 0.;
 Lmax = 10.

 The mean excitation energy, charge, mass, and angular momentum of the 9636 nuclei after preequilibrium decay and before the start of statistical decay are:

 Exav = 33.8 +/ 18.9 MeV; E*min = 0.1; E*max = 65.1

 Zav = 82.5 +/ 0.6; Zmin = 80.; Zmax = 83.

 Aav = 207.8 +/ 0.8; Amin = 203.; Amax = 209.

 Lav = 7.7 +/ 3.9 h-bar; Lmin = 0.; Lmax = 28.

 The mean kinetic energy, charge, mass, and angular momentum

 In a word window of the 9943 residual nuclei are:

 Ekav = 0.3 +/- 0.2 MeV; Ekmin = 0.0; Ekmax = 2.1

 Zav = 82.5 +/- 0.6; Zmin = 80.; Zmax = 83.

 Aav = 204.8 +/- 1.6; Amin = 200.; Amax = 208.

 Lav = 7.5 +/- 4.0 h-bar; Lmin = 0.; Lmax = 28.

 The mean excitation energy, charge, mass, angular momentum, and fission barrier height of the 57 fissioning nuclei are: E*av = 56.8 +/- 8.5 MeV; E*min = 34.3; E*max = 65.1 Zav = 83.0 +/- 0.0; Zmin = 83.; Zmax = 83. Aav = 208.2 +/- 0.8; Amin = 206.; Amax = 209. Large = 7.0 +/- 2.0 Applet.
 Law
 208.2
 +/ 0.8;
 Amin
 206.;
 Amax
 209.

 Law
 7.9
 +/ 2.9
 h-bar;
 Lmin
 1.;
 Lmax
 14.

 Bfav
 23.6
 +/ 0.2
 MeV;
 Bfmin
 23.3;
 Bfmax
 23.9
 The mean total fission product kinetic energy after neutron emission is 140.36 MeV. Direct Monte Carlo Simulation Method: Fissility = 0.0057 +/- 0.0008, Fission cross section = 1.11847E+01 +/- 1.48E+00 mb. Statistical Weight Functions Method: Fissility = 0.0052,

Fission cross section = 1.01828E+01 mb.

Number of coalesced d, t, He3, He4 = 27 0 0 0

Mean multiplicities, yields, and mean energies of ejected particles: (Notation: T - all production mechanisms, C - cascade, P - pre-equilibrium, Sp - from spallation residues, Pf - from nuclei before fission, F - from fission fragments, E - total evaporation = Sp + Pf + F, Co - Coalescence from cascade; Values which are identically zero are not printed.

Part.	Multiplicities	Yields [mb] <tke> [MeV]</tke>			

T n	3.6353 +/- 0.0191	7133.305 +/-	37.413 4.64			
C n	0.2867 +/- 0.0054	562.572 +/-	10.507 21.87			
Рп	0.2982 +/- 0.0055	585.138 +/-	10.715 14.77			
Sp n	3.0175 +/- 0.0174	5921.037 +/-	34.086 2.02			
Pf n	0.0030 +/- 0.0005	5.887 +/-	1.075 2.11			
Fn	0.0299 +/- 0.0017	58.671 +/-	3.393 2.26			
En	3.0504 +/- 0.0175	5985.595 +/-	34.271 2.02			
*******	******	*****	*****			
τp	0.4624 +/- 0.0068	907.336 +/-	13.343 29.61			
Ср	0.2436 +/- 0.0049	478.000 +/-	9.685 32.60			
Рр	0.2100 +/- 0.0046	412.069 +/-	8.992 26.94			
Spp	0.0088 +/- 0.0009	17.268 +/-	1.841 10.66			
Ер	0.0088 +/- 0.0009	17.268 +/-	1.841 10.66			
*******	*****	*****	*****			
Τd	0.0409 +/- 0.0020	80.255 +/-	3.968 26.14			
Ρd	0.0378 +/- 0.0019	74.172 +/-	3.815 26.84			
Sp d	0.0004 +/- 0.0002	0.785 +/-	0.392 11.65			
Ed	0.0004 +/- 0.0002	0.785 +/-	0.392 11.65			
Co d	0.0027 +/- 0.0005	5.298 +/-	1.020 18.53			
*******	******	*****	*****			
т t	0 0127 +/- 0 0011	24 920 +/-	2 211 23 22			
P t	$0.0126 \pm - 0.0011$	24 724 +/-	2 203 23 32			
Snt	$0.0001 \pm 7 0.0001$	0.196 +/-	0 196 10 90			
F +	0.0001 + / - 0.0001	0 196 +/-	0 196 10 90			
	0.0001 // 0.0001	0.100 .1	·····			
T U-2	$0.0016 \pm l_{-} 0.0004$	-/+ 01/0 c	0 705 20 00			
D Uo2	0.0016 +/- 0.0004	2 140 +/-	0.705 32.32			
r neo	0.0018 +/- 0.0004	3.140 +/-	0.785 32.92			
******	**********************	***********	******			
T He4	0.0074 +/- 0.0009	14.521 +/-	1.688 31.27			
P He4	0.0062 +/- 0.0008	12.166 +/-	1.545 33.12			
SpHe4	0.0012 +/- 0.0003	2.355 +/-	0.680 21.72			
E He4	0.0012 +/- 0.0003	2.355 +/-	0.680 21.72			
*******	******	******	******			

----- Energy Spectrum [mb/MeV] -----

Energy spectrum from 0.0 to 64.0 MeV (zero values suppressed).

 Tn [MeV]
 Total
 Cascade
 Precompound
 Total Evaporation

 0.0 1.0
 1.833E+03
 +/ 1.90E+01
 0.00E+00
 +/ 0.326E+00
 +/ 1.33E+03
 +/ 1.89E+01

 1.0 2.0
 1.847E+03
 +/ 1.90E+01
 2.374E+01
 +/ 2.16E+00
 1.923E+01
 +/ 1.94E+03
 +/ 1.884E+01

 56.0 58.0
 5.396E+00
 +/ 7.28E-01
 5.004E+00
 +/ 7.01E-01
 3.924E-01
 +/ 0.00E+00
 +/ 0.

Integrated: 7.133E+03 +/- 3.74E+01 5.626E+02 +/- 1.05E+01 5.851E+02 +/- 1.07E+01 5.986E+03 +/- 3.43E+01

-----Normalized Energy Probability Spectrum [1/MeV] ------

Energy spectrum from 0.0 to 64.0 MeV (zero values suppressed).

Integrated: 1.000E+00 +/- 5.24E-03 7.887E-02 +/- 1.47E-03 8.203E-02 +/- 1.50E-03 8.391E-01 +/- 4.80E-03

----- Angular Distributions [mb/sr] ------

Ang.n	Total	Cascade	Precompound	Total Evaporation
5.0 15.0	6.845E+02 +/- 3.75E+01 7.691E+02 +/- 2.31E+01	8.223E+01 +/- 1.30E+01 1.405E+02 +/- 9.86E+00	1.089E+02 +/- 1.50E+01 1.059E+02 +/- 8.56E+00	4.934E+02 +/- 3.18E+01 5.226E+02 +/- 1.90E+01
165.0 175.0	4.853E+02 +/- 1.83E+01 4.872E+02 +/- 3.16E+01	6.922E-01 +/- 6.92E-01 0.000E+00 +/- 0.00E+00	2.077E+01 +/- 3.79E+00 1.233E+01 +/- 5.04E+00	4.638E+02 +/- 1.79E+01 4.749E+02 +/- 3.12E+01
Integ.	7.133E+03 +/- 3.74E+01	5.626E+02 +/- 1.05E+01	5.851E+02 +/- 1.07E+01	5.986E+03 +/- 3.43E+01
Double	differential cross section	ons [mb/MeV/sr];		

Lab. angle = 22.5 to 27.5 degrees.

 Tn
 [MeV]
 Total
 Cascade
 Precompound
 Total
 Evaporation

 0.0 1.0
 1.262E+02 +/ 1.03E+01
 0.00E+00 +/ 0.00E+00 +/ 0.00E+00 +/ 0.262E+02 +/ 1.03E+01

 1.0 2.0
 1.457E+02 +/ 1.11E+01
 5.929E+00 +/ 2.24E+00
 1.64E+00
 +/ 1.381E+02 +/ 1.08E+01

 56.0 58.0
 4.235E+00 +/ 1.34E+00
 4.235E+00 +/ 1.34E+00
 0.00E+00 +/ 0.00E+00 +/-

Integrated: 7.573E+02 +/- 2.53E+01 1.762E+02 +/- 1.22E+01 8.809E+01 +/- 8.64E+00 4.930E+02 +/- 2.04E+01

Double differential cross sections [mb/MeV/sr];

Lab. angle = 52.5 to 57.5 degrees.

 Tn
 [MeV]
 Total
 Cascade
 Precompound
 Total Evaporation

 0.0 1.0
 1.403E402 +/- 7.83E+00 0.000E+00 +/- 0.00E+00 8.740E-01 +/- 6.18E-01 1.394E+02 +/- 7.81E+00
 1.394E+02 +/- 7.81E+00

 1.0 2.0
 1.385E+02 +/- 7.78E+00 3.059E+00 +/- 1.16E+00 2.185E+00 +/- 9.7TE-01 1.33E+02 +/- 7.63E+00

 54.0 56.0
 8.740E-01 +/- 4.37E-01 8.740E-01 +/- 4.37E-01 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00

 56.0 58.70E-01 +/- 2.19E-01 2.185E-01 +/- 2.19E-01 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00

Integrated: 6.241E+02 +/- 1.65E+01 7.997E+01 +/- 5.91E+00 7.254E+01 +/- 5.63E+00 4.715E+02 +/- 1.44E+01

Double differential cross sections [mb/MeV/sr]; Lab. angle = 72.5 to 77.5 degrees.

 Tn
 [MeV]
 Total
 Cascade
 Precompound
 Total
 Evaporation

 0.0 1.0
 1.583E+02
 +/ 7.66E+00
 0.000E+00
 +/ 0.00E+01
 +/ 3.71E-01
 1.579E+02
 +/ 7.65E+00

 1.0 2.0
 1.523E+02
 +/ 7.51E+00
 1.853E+00
 +/ 8.29E-01
 1.482E+00
 +/ 7.41E-01
 1.490E+02
 +/ 7.43E+00

 48.0 50.0
 1.853E-01
 +/ 1.852E-01
 0.000E+00
 +/ 0.00E+00
 +/-

Integrated: 5.841E+02 +/- 1.47E+01 4.373E+01 +/- 4.03E+00 4.744E+01 +/- 4.19E+00 4.929E+02 +/- 1.35E+01 Double differential cross sections [mb/MeV/sr]:

Lab. angle = 92.5 to 97.5 degrees.

 Tn
 [MeV]
 Total
 Cascade
 Precompound
 Total Evaporation

 0.0 1.0
 1.445E+02 +/- 7.20E+00
 0.000E+00 +/- 0.00E+00
 7.187E-01 +/- 5.08E-01
 1.437E+02 +/- 7.19E+00

 1.0 2.0
 1.520E+02 +/- 7.39E+00
 1.797E+00 +/- 8.04E-01
 1.437E+00 +/- 7.19E-01
 1.488E+02 +/- 7.31E+00

 34.0 36.0
 3.593E-01 +/- 2.54E-01
 0.000E+00 +/- 0.00E+00
 3.593E-01 +/- 2.64E-01
 0.000E+00 +/- 0.00E+00

 38.0 40.0
 1.797E-01 +/- 1.80E-01
 0.000E+00 +/- 0.00E+00
 1.797E-01 +/- 1.80E-01
 0.00E+00 +/- 0.00E+00

Integrated: 5.365E+02 +/- 1.39E+01 1.725E+01 +/- 2.49E+00 3.450E+01 +/- 3.52E+00 4.848E+02 +/- 1.32E+01

Double differential cross sections [mb/MeV/sr];

Lab. angle = 112.5 to 117.5 degrees.

 Tn
 [MeV]
 Total
 Cascade
 Precompound
 Total
 Evaporation

 0.0 1.0
 1.540±+02 +/- 7.80±+00 0.000±+00 +/- 0.00±+00 0.000±+00 +/- 0.00±+00 1.540±+02 +/- 7.80±+00
 1.0 2.0
 1.544±+02 +/- 7.81±+00 1.580±+00 +/- 7.90±+01 2.370±+00 +/- 9.68±-01 1.505±+02 +/- 7.71±+00

 30.0 32.0
 1.975±-01 +/- 1.97±-01 0.000±+00 +/- 0.00±+00 1.975±-01 +/- 1.97±-01 0.000±+00 +/- 0.00±+00
 3.50±-01 +/- 1.97±-01 0.000±+00 +/- 0.00±+00

 34.0 36.0
 3.50±-01 +/- 2.79±-01 0.000±+00 +/- 0.00±+00 3.950±-01 +/- 1.97±-01 0.000±+00 +/- 0.00±+00

Integrated: 5.159E+02 +/- 1.43E+01 7.900E+00 +/- 1.77E+00 2.686E+01 +/- 3.26E+00 4.811E+02 +/- 1.38E+01

Double differential cross sections [mb/MeV/sr]; Lab. angle = 152.5 to 157.5 degrees.

----- Energy Spectrum [mb/MeV] -----

Energy spectrum from 0.0 to 64.0 MeV (zero values suppressed).

 THe4[MeV]
 Total
 Coalescence
 Precompound
 Total Evaporation

 19.0 20.0
 3.924E-01 +/- 2.78E-01
 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00
 3.924E-01 +/- 2.78E-01

 20.0 21.0
 7.849E-01 +/- 3.92E-01
 0.000E+00 +/- 0.00E+01
 1.962E-01 +/- 1.96E-01
 5.87E-01 +/- 3.40E-01

 54.0 56.0
 9.811E-02 +/- 9.81E-02
 0.000E+00 +/- 0.00E+00
 9.811E-02 +/- 9.81E-02
 0.000E+00 +/- 0.00E+00

 56.0 58.0
 9.811E-02 +/- 9.81E-02
 0.000E+00 +/- 0.00E+00
 9.811E-02 +/- 9.81E-02
 0.000E+00 +/- 0.00E+00

Integrated: 1.452E+01 +/- 1.69E+00 0.000E+00 +/- 0.00E+00 1.217E+01 +/- 1.55E+00 2.355E+00 +/- 6.80E-01

----- Angular Distributions [mb/sr] -----

 Ang.He4
 Total
 Coalescence
 Precompound
 Total Evaporation

 [deg.]
 15.0
 2.769E+00 +/- 1.38E+00 0.000E+00 +/- 0.00E+00 2.769E+00 +/- 1.38E+00 0.000E+00 +/- 0.00E+00
 2.542E+00 +/- 1.04E+00 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 1.04E+00 0.000E+00 +/- 0.00E+00

 165.0
 6.922E-01 +/- 6.92E-01 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00 +/- 0.00E+00

 175.0
 2.056E+00 +/- 1.04E+00 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00 +/- 0.00E+00
 0.00E+00 +/- 0.00E+00 +/- 0.00E+00

Integ. 1.452E+01 +/- 1.69E+00 0.000E+00 +/- 0.00E+00 1.217E+01 +/- 1.55E+00 2.355E+00 +/- 6.80E-01

Double differential cross sections [mb/MeV/sr]; Lab. angle = 22.5 to 27.5 degrees.

THe4[]	MeV]	Total	Coalescence	Precompound	Total Evaporation
26.0-	28.0	4.235E-01 +/- 4.24E-01	0.000E+00 +/- 0.00E+00	4.235E-01 +/- 4.24E-01	0.000E+00 +/- 0.00E+00

Integrated: 8.471E-01 +/- 8.47E-01 0.000E+00 +/- 0.00E+00 8.471E-01 +/- 8.47E-01 0.000E+00 +/- 0.00E+00

Double differential cross sections [mb/MeV/sr]; Lab. angle = 52.5 to 57.5 degrees.

 THe4[MeV]
 Total
 Coalescence
 Precompound
 Total Evaporation

 22.0 24.0
 2.185E-01 +/- 2.19E-01 0.000E+00 +/- 0.00E+00 2.185E-01 +/- 2.19E-01 0.000E+00 +/- 0.00E+00
 2.185E-01 +/- 2.19E-01 0.000E+00 +/- 0.00E+00
 2.185E-01 +/- 2.19E-01 0.000E+00 +/- 0.00E+00

 24.0 28.0
 2.185E-01 +/- 2.19E-01 0.000E+00 +/- 0.00E+00
 2.185E-01 +/- 2.19E-01 0.000E+00 +/- 0.00E+00

 26.0 28.0
 2.185E-01 +/- 2.19E-01 0.000E+00 +/- 0.00E+00
 2.185E-01 +/- 2.19E-01 0.000E+00 +/- 0.00E+00

 30.0 32.0
 2.185E-01 +/- 2.19E-01 0.000E+00 +/- 0.00E+00
 2.185E-01 +/- 2.19E-01 0.000E+00 +/- 0.00E+00

Integrated: 1.748E+00 +/- 8.74E-01 0.000E+00 +/- 0.00E+00 1.748E+00 +/- 8.74E-01 0.000E+00 +/- 0.00E+00

Double differential cross sections [mb/MeV/sr]; Lab. angle = 72.5 to 77.5 degrees.

 THe4[MeV]
 Total
 Coalescence
 Precompound
 Total
 Evaporation

 26.0 28.0
 1.853E-01 +/- 1.85E-01
 0.000E+00 +/- 0.00E+00
 1.853E-01 +/- 1.85E-01
 0.000E+00 +/- 0.00E+00

 36.0 38.0
 1.853E-01 +/- 1.85E-01
 0.000E+00 +/- 0.00E+00
 1.853E-01 +/- 1.85E-01
 0.000E+00 +/- 0.00E+00

Integrated: 7.412E-01 +/- 5.24E-01 0.000E+00 +/- 0.00E+00 7.412E-01 +/- 5.24E-01 0.000E+00 +/- 0.00E+00

Double differential cross sections [mb/MeV/sr]; Lab. angle = 92.5 to 97.5 degrees.

 THe4 [MeV]
 Total
 Coalescence
 Precompound
 Total
 Evaporation

 22.0 24.0
 1.797E-01 +/- 1.80E-01
 0.000E+00 +/- 0.00E+00
 1.797E-01 +/- 1.80E-01
 0.000E+00 +/- 0.00E+00

 24.0 26.0
 1.797E-01 +/- 1.80E-01
 0.000E+00 +/- 0.00E+00
 1.797E-01 +/- 1.80E-01
 0.000E+00 +/- 0.00E+00

 30.0 32.0
 1.797E-01 +/- 1.80E-01
 0.000E+00 +/- 0.00E+00
 1.797E-01 +/- 1.80E-01
 0.000E+00 +/- 0.00E+00

Integrated: 1.078E+00 +/- 6.22E-01 0.000E+00 +/- 0.00E+00 1.078E+00 +/- 6.22E-01 0.000E+00 +/- 0.00E+00

Elapsed cpu time = 0. min and 5.669 sec.

Wed Feb 1 10:55:15 2012 Example No. 7: xsec and kinetic energy of all products measured at GSI in inverse kinematics for 800 MeV p + Au197; 10,000 events. Number of types of evaporated particles = 6M TO A Z Q B limc idel 0.9383 0.8000 197. 79. 1 1 10000 1 dt0 = -20.0, t0max = 1000.5, dteta = 10.0mspec mpyld mchy misy mdubl mang ipar1 ipar2 1 0 1 0 0 rOm = 1.2, & cevap = 12.0. lim = 100000 . Geometrical cross section = 2394.46 mb. Inelastic cross section used here = 1629.65 mb Monte Carlo inelastic cross section = 1590.37 mb Wed Feb 1 10:55:15 2012 800.0 MeV (Z = 1, A = 1) + (Z = 79., A = 197.) Number of inelastic interactions = 10000. Number of elastic interactions = 5056, Reaction cross section = 1629.65 mb. Elastic cross section = 823.95 mb. The mean excitation energy, charge, mass, and angular momentum 10000 nuclei after the of the cascade and before preequilibrium decay are:

 E*av = 226.4 +/-15.04 MeV; E*min = -1.1; E*max = 717.9

 Zav = 78.6 +/- 1.1; Zmin = 73.; Zmax = 82.

 Aav = 192.7 +/- 2.6; Amin = 182.; Amax = 198.

 Lav = 12.7 +/- 7.5 h-bar; Lmin = 0.; Lmax = 46.

 The mean charge, mass, and angular momentum of the 165 residual nuclei with less than 3 MeV of excitation energy after the cascade are: Zav = 79.2 +/- 0.4; Zmin = 79.; Zmax = 80. Aav = 196.1 +/- 0.4; Amin = 194.; Amax = 197. Lav = 3.5 +/- 2.2 h-bar; Lmin = 0.; Lmax = 13. The mean excitation energy, charge, mass, and angular momentum of the 9835 nuclei after preequilibrium
 of the
 9835 nuclei after preequilibrium

 decay and before the start of statistical decay are:

 E*av = 181.2 +/- 128.5 MeV; E*min = 1.0; E*max = 683.9

 Zav = 77.7 +/- 1.7; Zmin = 69.; Zmax = 81.

 Aav = 190.6 +/- 4.2; Amin = 172.; Amax = 197.

 Lav = 14.8 +/- 9.0 h-bar; Lmin = 0.; Lmax = 66.

 The mean kinetic energy, charge, mass, and angular momentum of the
 9566 residual nuclei are:

 Ekav =
 2.1 +/- 2.6 MeV; Ekmin = 0.0; Ekmax = 23.6

 Zav = 74.9 +/- 3.9;
 Zmin = 60.; Zmax = 81.

 Aav = 175.8 +/- 13.3;
 Amin = 132.; Amax = 197.

 Lav =
 14.5 +/- 9.1 h-bar; Lmin = 0.; Lmax = 66.

 The mean excitation energy, charge, mass, angular momentum, and The sour operators of the grad firstoning nuclei are: E*av = 254.1 +/-102.0 MeV; E*min = 48.9; E*max = 551.3 Zav = 76.9 +/- 2.4; Zmin = 66.; Zmax = 81. Aav = 184.0 +/- 6.3; Amin = 160.; Amax = 196. Lav = 17.7 +/- 8.2 h-bar; Lmin = 2.; Lmax = 51. Bfav = 15.9 +/- 2.8 MeV; Bfmin = 8.3; Bfmax = 25.7 The mean total fission product kinetic energy after neutron emission is 121.75 MeV. Direct Monte Carlo Simulation Method: Fissility = 0.0434 +/- 0.0021, Fission cross section = 7.07268E+01 +/- 3.39E+00 mb. Statistical Weight Functions Method: Fissility = 0.0418, Fission cross section = 6.81368E+01 mb. 3566 1169 Number of coalesced d, t, He3, He4 = 221 235 Mean multiplicities, yields, and mean energies of ejected particles: (Notation: T - all production mechanisms, C - cascade, P - pre-equilibrium, Sp - from spallation residues, Pf - from nuclei before fission, F - from fission fragments, E - total evaporation = Sp + Pf + F, Co - Coalescence from cascade; Values which are identically zero are not printed. Part. Multiplicities Yields [mb] <TKE> [MeV]

Сп				4//.	531	''	61.	.855	20.12		
	3.1900 -	+/- 0.0	0179 5	198.	580	+/-	29	.106	76.92		
Рп	0.5990 -	+/- 0.0	0077	976.	160	+/-	12.	.613	19.31		
Spn	9.8640 -	+/- 0.0	0314 16	074.	857	+/-	51.	. 182	3.82		
Pf n	0.1216 -	+/- 0.0	0035	198.	165	+/-	5.	. 683	6.80		
Fn	0.6319 -	+/- 0.0	0079 1	029.	775	+/-	12.	.954	4.83		
En	10.6175 -	+/- 0.0	0326 17	302.	797	+/-	53.	.101	3.92		
******	********	*****	*******	****	****	*****	***	******	******	****	
Тр	2.3550 -	+/- 0.0	0153 3	837.	823	+/-	25.	.009	80.49		
Ср	0.8762 -	+/- 0.0	0094 1	427.	898	+/-	15.	.254	189.69		
Рр	0.3430 -	+/- 0.0	0059	558.	970	+/-	9.	.544	28.11		
Sp p	1.0578 -	+/- 0.0	0103 1	723.	843	+/-	16.	.761	12.15		
Pfp	0.0179 -	+/- 0.0	0013	29.	171	+/-	2.	. 180	14.39		
Fn	0.0601 -	+/- 0.0	0025	97	942	+/-	3	995	9.75		
	1 1358 -	+/- 0 (107 1	850	055	+/-	17	368	12.06		
******* - h	1.1000			····			. ۱۱ د بد بد ر		12.00	دددد	
******* T J	1 0705	. / /	100 1	****	F 20	. /	10	001		*****	
	1.0705 -	+/- 0.0	0103 1	144.	539	+/-	10.	.001	36.49		
Ρα	0.2514 -	+/- 0.0	0050	409.	694	+/-	8.	. 1 / 1	31.55		
Sp d	0.4316 -	+/- 0.0	0066	703.	356	+/-	10.	.706	13.31		
Pf d	0.0136 -	+/- 0.0	0012	22.	163	+/-	1.	. 900	14.79		
Fd	0.0182 -	+/- 0.0	0013	29.	660	+/-	2.	.199	11.71		
Ed	0.4634 -	+/- 0.0	0068	755.	179	+/-	11.	.094	13.29		
Co d	0.3557 -	+/- 0.0	0060	579	666	+/-	9	719	70 19		
*******	*********	··/ ···		****		· /				****	
******* T +	0 2004	. / /		~~~~		. /	10	100	07 60	****	
1 U	0.3694 -	+/- 0.0	0062	034.	505	+/-	10.	.169	21.60		
Ρt	0.0802 -	+/- 0.0	0028	130.	698	+/-	4.	.615	36.83		
Sp t	0.1773 -	+/- 0.0	0042	288.	937	+/-	6.	.862	14.13		
Pf t	0.0065 -	+/- 0.0	8000	10.	593	+/-	1.	.314	14.93		
Ft	0.0087 -	+/- 0.0	0009	14.	178	+/-	1.	.520	13.03		
Εt	0.1925 -	+/- 0.0	0044	313.	707	+/-	7	.150	14.11		
Co t	0 1167 -	+/- 0 0	0034	190	180	+/-	5	567	43 50		
*******	**************************************	·/ U.(100	• / • • • • • • •			-10.00	*****	
	····**************	** -/- ^ ·	~~~~***** \/\21	~~*** 1 F 77	004	~~~*** +/	·***	060	*******		
і нез	0.0965 -	+/- 0.0	1031	15/.	201	+/-	ь.	.062	44.27		
Р НеЗ	0.0555 -	+/- 0.0	0024	90.	446	+/-	3.	.839	44.17		
SpHe3	0.0173 -	+/- 0.0	0013	28.	193	+/-	2.	. 143	23.59		
PfHe3	0.0007 -	+/- 0.0	0003	1.	141	+/-	0.	.431	25.34		
F He3	0.0009 -	+/- 0.0	0003	1.	467	+/-	0.	489	21.30		
E He3	0.0189 -	+/- 0.0	0014	30.	800	+/-	2.	.240	23.55		
CoHe3	0 0221 -	+/- 0 0	0015	36	015	+/-	2	423	62 24		
*******	********	******		****				. <u>120</u>		*****	
T II-4	0 5045	. / /	077	000	000	. /	40		05 20		
п печ	0.5945 -	+/- 0.0		900.	020	+/-	12.	. 505	25.30		
P He4	0.0343 -	+/- 0.0	0019	55.	897	+/-	3.	.018	54.99		
SpHe4	0.5015 -	+/- 0.0	0071	817.	269	+/-	11.	. 541	23.24		
PfHe4	0.0085 -	+/- 0.0	0009	13.	852	+/-	1.	. 502	25.69		
F He4	0.0269 -	+/- 0.0	0016	43.	838	+/-	2.	.673	17.93		
E He4	0.5369 -	+/- 0.0	0073	874.	959	+/-	11.	941	23.01		
CoHe4	0.0233 -	+/- 0.0	0015	37.	971	+/-	2.	488	36.26		
******	*********	******	******	****	****	*****	***	*****	******	*****	
ni-	0 1051	+/- 0 0	1032	171	276	+/-	5	283	72 /10		
pi ni0	0.1001	+/_ 0.0	0012	200	606	+/-	7	002	04 07		
	0.1007 -	-/- 0.0	043	302.	770	+/-		.023	04.07		
P10	0 0070	. / ^ /	029	141.	119	+/-	4.	.807	115.45		
pi+	0.0870 -	+/- 0.0									
pi+ ******	0.0870 - *********	+/- 0.(*****	******	****	****	****	***	******	******	****	
pi+ ******	0.0870 - *********	+/- 0.(*****	******	****	****	*****	***	******	******	*****	
pi+ ********	0.0870 - ********** ******* 1	+/- 0.(******* Nuclide	********* e yields	**** [mb]	**** (z	*****	*** alu	****** 1es supj	****** pressed	*****) ***	· ·*********
pi+ ******* ******	0.0870 - ********** ******* 1	+/- 0.(******* Nuclide	********* e yields	**** [mb]	**** (z	***** ero v	alu	****** 1es supj	******* pressed	*****) ***	· ·*********
pi+ ******* ******	0.0870 - ********** ******** 1	+/- 0.0 ******* Nuclide Z = 8:	********* 9 yields 1.	**** [mb]	***** (z	***** ero v Z =	**** ralı 80.	******* 1es supj	****** pressed	*****) *** Z =	**************************************
pi+ ******** ********	0.0870 - ********* ******** 1 2 0.000E+00	+/- 0.(****** Nuclide Z = 8: 0 +/- (********* e yields 1. 0.00E+00 1	**** [mb] 5.54	***** (z 1E+0	***** ero v Z = 0 +/-	**** ralı 80. 9.	******** 1es supj .50E-01	******* pressed 4.889E	*****) *** Z = +00 +	*********** * 79. •/- 8.93E-01
PIO pi+ ******** ******* A = 197 A = 196	0.0870 - ********* ******** 1 0.000E+00 0.000E+00	+/- 0.(****** Nuclide Z = 8: D +/- (D +/- (********* e yields 1. 0.00E+00 = 0.00E+00 =	**** [mb] 5.54 8.80	***** (z 1E+0 00E+0	***** ero v Z = 0 +/- 0 +/-	**** valu 80. • 9. • 1.	******** 1es supj .50E-01 .20E+00	******** pressed 4.889E 3.748E	*****) *** Z = +00 + +01 +	************ 79. -/- 8.93E-01 -/- 2.47E+00
A = 197 A = 196	0.0870 - ************************************	+/- 0.(******* Nuclide Z = 8: 0 +/- (0 +/- (********* e yields 1. 0.00E+00 = 0.00E+00 =	**** [mb] 5.54 8.80	***** (z 1E+0 00E+0 	z = 0 +/- 0 +/-	**** alu 80. 9. 1.	******** les supj .50E-01 .20E+00	******* pressed 4.889E 3.748E	*****) *** Z = +00 + +01 + 	************ 79. -/- 8.93E-01 -/- 2.47E+00
A = 197 A = 196 A = 197	0.0870 - ************************************	+/- 0.(******* Nuclide Z = 8: D +/- (D +/- (D +/- (<pre>************************************</pre>	**** [mb] 5.54 8.80 1 63	(z 1E+0)0E+0 	z = 0 +/- 0 +/- 	**** ralu 80. 9. 1.	********* les supj .50E-01 .20E+00 	******* pressed 4.889E 3.748E	*****) *** Z = +00 + +01 + 	<pre>************************************</pre>
A = 197 A = 196 A = 182 A = 182	0.0870 - ************************************	+/- 0.(******* Nuclide Z = 8: D +/- (D +/- (D +/- (0 +/- (**************************************	**** [mb] 5.54 8.80 1.63	(z 1E+0) 0E+0 30E-0	<pre>****** Z = 0 +/- 0 +/ 1 +/- 1 +/-</pre>	**** alu 80. 9. 1. 1.	********* les supj .50E-01 .20E+00 .63E-01	4.889E 3.748E 0.000E	*****) *** Z = +00 + +01 + +00 +	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00
A = 197 A = 196 A A = 182 S = 16	0.0870 - ************************************	+/- 0.(****** Nuclide Z = 8: D +/- (D +/- (D +/- (1 +/- 3	********** • yields 1. 0.00E+00 0.00E+00 0.00E+00 3.64E-01	**** [mb] 5.54 8.80 1.63 7.69	(z 1E+0) 0E+0 30E-0 2E+0	<pre>***** ero v Z = 0 +/- 0 +/ 1 +/- 1 +/-</pre>	**** 80. 9. 1. 1. 3.	********* les supj	******* pressed 4.889E 3.748E 0.000E 1.974E	*****) *** Z = +00 + +01 + +00 + +02 +	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00
A = 197 A = 196 A = 182 S = 16	0.0870 - ************************************	+/- 0.(******* Vuclide Z = 8: 0 +/- (0 +/- (0 +/- (1 +/- 3	**************************************	**** [mb] 5.54 8.80 1.63 7.69	(z 1E+0) 00E+0 30E-0 2E+0	<pre>***** ero v Z = 0 +/- 0 +/ 1 +/- 1 +/-</pre>	**** 80. 9. 1. 1. 3.	********* les supj .50E-01 .20E+00 .63E-01 .54E+00	******* pressed 4.889E 3.748E 0.000E 1.974E	*****) *** Z = +00 + +01 + +00 + +02 +	
A = 197 A = 196 A = 182 S = 16	0.0870 - ************************************	+/- 0.0 +/- 0.0 Nuclide Z = 8: D +/- 0 D +/- 0 D +/- 0 D +/- 0 D +/- 3 Z = 78	**************************************	**** [mb] 5.54 8.80 1.63 7.69	(z 11E+0 00E+0 30E-0 92E+0	<pre>***** ero v Z = 0 +/- 0 +/ 1 +/- 1 +/- Z =</pre>	**** ralu 80. 9. 1. 1. 3. 77.	*********	******** pressed 4.889E 3.748E 0.000E 1.974E	*****) *** Z = +00 + +01 + +00 + +02 + Z =	- 79. - 8.93E-01 - 2.47E+00
A = 197 A = 197 A = 182 S = 16 A = 182 S = 16 A = 196	0.0870 - ************************************	+/- 0.(******* Nuclide Z = 8: D +/- (D +/- (D +/- (D +/- (1 +/- 3 Z = 78 1 +/- 2	********** • yields 1. 0.00E+00 0.00E+00 0.00E+00 3.64E-01 3. 2.30E-01	**** [mb] 5.54 8.80 1.63 7.69 0.00	(z 1E+0 00E+0 30E-0 2E+0 00E+0	<pre>x***** xero v Z = 0 +/- 0 +/ 1 +/- 1 +/- Z = 0 +/-</pre>	***** ralu 80. 9. 1. 1. 1.	**************************************	******** pressed 4.889E 3.748E 0.000E 1.974E 0.000E	*****) *** C = +00 + +01 + +00 + +02 + Z = +00 +	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 - 76. -/- 0.00E+00
A = 197 A = 196 A = 182 S = 16 A = 196 A = 196 A = 196 A = 195	0.0870 - ************************************	+/- 0.0 ******** Nuclide Z = 8: 0 +/- 0 0 +/- 0 1 +/- 3 Z = 78 1 +/- 2 1 +/- 2	<pre>********** a yields 1. 0.00E+00 2.00E+00 3.64E-01 3. 2.30E-01 1.35E+00 </pre>	**** [mb] 5.54 8.80 1.63 7.69 0.00 0.00	(z 1E+0 00E+0 30E-0 92E+0 00E+0 00E+0	<pre>***** ero v Z = 0 +/- 0 +/ 1 +/- 1 +/- Z = 0 +/- 0 +/- 0 +/-</pre>	***** ralu 80. 9. 1. 1.	********* 105 SUPJ 50E-01 20E+00 63E-01 54E+00 00E+00 .00E+00	******** pressed 4.889E 3.748E 0.000E 1.974E 0.000E 0.000E	*****) *** C = +00 + +01 + +00 + +02 + Z = +00 + +00 +	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 - 76. -/- 0.00E+00
A = 197 A = 197 A = 196 A = 182 S = 16 A = 196 A = 195 	0.0870 - ************************************	+/- 0.0 ***********************************	**************************************	**** [mb] 5.54 8.80 1.63 7.69 0.00 0.00	(z 1E+0 00E+0 30E-0 92E+0 00E+0 00E+0 	<pre>x***** ero v Z = 0 +/- 0 +/ 1 +/- 1 +/- Z = 0 +/- 0 +/</pre>	***** ralu 80. 9. 1. 1. 1.	**************************************	******* oressed 4.889E 3.748E 0.000E 1.974E 0.000E 0.000E	*****) *** Z = +00 + +01 + +00 + +02 + Z = +00 + +00 + 	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 - 76. -/- 0.00E+00 -/- 0.00E+00
A = 197 A = 196 A = 196 A = 182 S = 16 A = 196 A = 196 A = 196 A = 195 	0.0870 - ************************************	+/- 0.0 ******* Nuclide Z = 8: 0 +/- (0 +/- (0 +/- (1 +/- 3 Z = 78 1 +/- 2 1 +/- 2 0 +/- (0 +/- (1 +/- 3) 	<pre>*********** a yields 1. 0.00E+00 0.00E+00 3.64E-01 3. 2.30E-01 1.35E+00 0.00E+00 </pre>	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 	(z 11E+0 00E+0 30E-0 22E+0 00E+0 00E+0 00E+0 00E+0 00E+0 00E+0 00E+0	<pre>***** ero v Z = 0 +/- 0 +/ 1 +/- 1 +/- Z = 0 +/- 0 +/ 0 +/- 0 +/-</pre>	***** ralu 80. 9. 1.	**************************************	<pre>******* pressed 4.889E 3.748E 0.000E 1.974E 0.000E 0.000E 4.889E</pre>	<pre>*****) *** Z = +00 + +01 + +00 + +02 + Z = +00 + +00 +01 +</pre>	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 - 0.00E+00 -/- 0.00E+00
A = 197 A = 197 A = 196 A = 182 S = 16 A = 196 A = 196 A = 195 A = 195 A = 195 A = 29	0.0870 - ************************************	+/- 0.0 ******* Nuclide Z = 8: 0 +/- (0 +/- (0 +/- (1 +/- 3) Z = 78 1 +/- 2 1 +/- 2 1 +/- 2 0 +/- (0 +/- (<pre>********* * yields 100E+0000E+0000E+00 3.64E-01 3. 2.30E-01 1.35E+000E+00 0.00E+0035E+00 0.00E+0035E+00 0.00E+0035E+00 0.00E+0035E+00 0.00E+00 0.00E</pre>	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 0.000	(z 11E+0 00E+0 30E-0 22E+0 00E+0	<pre>x***** ero v Z = 0 +/- 0 +/ 1 +/- 1 +/- Z = 0 +/- 0 +/- 0 +/ 0 +/- 2 +/-</pre>	***** ralu 80. 9. 1. 1. 1.	**************************************	******* 0 - essed 4 - 889E 3 - 748E 0 - 000E 0 - 000E 0 - 000E 0 - 000E 4 - 889E 4 - 889E 1 - 57E	*****) *** Z = +00 + +01 + +00 + +02 + Z = +00 + -01 + +02 +	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.382E-01 -/- 2.82E-01
A = 197 $A = 197$ $A = 196$ $A = 196$ $S = 16$ $A = 196$ $A = 195$ $A = 195$ $A = 168$ $S = 29$	0.0870 - 	+/- 0.0 ******* Nuclide Z = 8: 0 +/- 0 0 +/- 0 1 +/- 3 Z = 78 1 +/- 3 Z = 78 1 +/- 2 1 +/- 2 0 +/- 0 2 +/- 6	<pre>********* * yields 100E+00 0.00E+00 3. 0.00E+00 3. 4. 2.30E+01 1. 3. 0.00E+00 5.43E+00 </pre>	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 0.00 1.37	<pre>***** (z 1E+0) 0E+0 30E-0 2E+0 00E+0 00E+0 00E+0 '1E+0</pre>	Z = 0 +/- 0 +/- 1 +/- 1 +/- Z = 0 +/- 0 +/- 0 +/- 2 +/-	***** ralu 80. 9. 1. 1. 1. 1.	**************************************	4.889E 3.748E 0.000E 1.974E 0.000E 0.000E 4.889E 1.657E	*****) *** Z = +00 + +01 + +00 + +02 + Z = +00 + -01 + +02 +	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 - 76. -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 5.20E+00
A = 197 $A = 197$ $A = 196$ $A = 196$ $A = 196$ $A = 196$ $A = 195$ $$ $A = 168$ $S = 29$	0.0870 - ************************************	+/- 0.(************************************	**************************************	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 0.00 0.00 1.37	(z 41E+0 00E+0 30E-0 02E+0 00E+0 00E+0 00E+0 11E+0	<pre>x***** xero v Z = 0 +/- 0 +/ 1 +/- 1 +/- Z = 0 +/- 0 +/ 0 +/- 2 +/- 7 =</pre>	***** ralu 80. 9. 1. 1. 1. 0. 0. 0. 0. 4. 74	********** 1005 Supp 50E-01 20E+00 63E-01 54E+00 00E+00 00E+00 00E+00 73E+00	******* pressed 4.889E 3.748E 0.000E 1.974E 0.000E 0.000E 0.000E 4.889E 1.657E	*****	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 5.20E+00
A = 197 A = 196 A = 196 A = 182 S = 16 A = 196 A = 195 A = 195 A = 168 S = 29	0.0870 - 	+/- 0.(+/- 0.(+****** Nuclide Z = 8: 0 +/- (0 +/- (1 +/- 3) Z = 78 1 +/- 2 1 +/- 2 1 +/- 2 0 +/- (2 +/- 6 Z = 75	<pre>********* * yields 1</pre>	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 0.00 0.00 1.37	(z (z (z (z (z) (z) (z) (z) (z) (z) (z)	Z = 0 +/- 0 +/- 1 +/- 1 +/- Z = 0 +/- 0 +/- 2 +/- Z =	***** ralu 80. 9. 1. 1. 1. 1. 0. 0. 0. 0. 4. 74.	**************************************	4.889E 3.748E 0.000E 0.000E 0.000E 0.000E 0.000E 1.974E 1.657E	****** Z = +00 + +01 + +00 + +02 + Z = -01 + +02 + Z = Z =	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 - 73.
A = 197 A = 196 A = 182 S = 16 A = 196 A = 195 A = 168 S = 29	0.0870 - 	+/- 0.(************************************	<pre>********* * yields 1</pre>	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 0.00 1.37	(z (z (z (z (z) (z) (z) (z) (z) (z) (z)	***** ero v Z = 0 +/- 0 +/- 1 +/- 1 +/- Z = 0 +/- 0 +/- 0 +/- 0 +/- 2 +/- Z = 2 +/- Z =	***** valu * 9. * 1. * 3. 77. * 0. * 0. * 0. * 4. 74.	**************************************	******* 9 ressed 4.889E 3.748E 0.000E 1.974E 0.000E 0.000E 1.974E 1.974E 1.657E	****** Z = +00 + +01 + +00 + +02 + Z = +00 + +02 + Z = -01 + +02 + Z = -	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 5.20E+00
A = 197 A = 196 A = 196 A = 196 A = 182 S = 16 A = 182 S = 16 A = 195 A = 168 S = 29	0.0870 - 	+/- 0.(++/- 0.(++++++++++++++++++++++++++++++++++++	**************************************	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 0.00 1.37 	<pre>(z (z (</pre>	<pre>***** ero v Z = 0 +/- 0 +/- 1 +/- 1 +/- Z = 0 +/- 0 +/- 0 +/- 2 +/- Z =</pre>	<pre>***** valu 80. 9. 1. 3. 77. 3. 77. 0. 4. 74. 74.</pre>	**************************************	4.889E 3.748E 0.000E 1.974E 0.000E 4.889E 1.657E	****** 2 = +00 + +01 + +00 + +02 + Z = +00 + +00 + +02 + Z = 	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 5.20E+00 -73.
A = 197 A = 196 A = 182 S = 16 A = 182 S = 16 A = 196 A = 195 A = 168 S = 29	0.0870 - 	$\begin{array}{rrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrrr$	<pre>> yields 1</pre>	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 0.00 1.37 	(z (z (z (z (z) (z) (z) (z) (z) (z) (z)	****** ero v Z = 0 +/- 0 +/- 1 +/- 1 +/- 2 = 0 +/- 0 +/- 0 +/- 2 +/- Z = 	***** ***** * alu * 9. * 1. * 1. * 3. * 77. * 0. * 0. * 0. * 0. * 4. 74. * -	**************************************	4.889E 3.748E 0.000E 1.974E 0.000E 0.000E 1.657E	****** Z = +00 + +01 + - - +00 + +02 + Z = +00 + +00 + - -01 + +02 + Z = - -	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 - 73.
A = 197 A = 196 A = 196 A = 186 A = 182 S = 16 A = 185 A = 168 S = 29	0.0870 - 	+/- 0.(******* Nuclide Z = 8: D +/- (D +/- (D +/- (1 +/- 3: T	**************************************	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 0.00 1.37 	<pre>****** (z (z (z 000E+0 000E+0 000E+0 000E+0 000E+0</pre>	****** ero v Z = 0 +/- 0 +/- 1 +/- 1 +/- Z = 0 +/- 0 +/- 0 +/- 0 +/- 2 +/- Z = 	<pre>***** ralu 80. 9. 1 1 1 1 3 77 0</pre>	**************************************	4.889E 3.748E 0.000E 1.974E 0.000E 4.889E 1.657E	******) *** Z = +00 + +01 + +00 + +02 + Z = +00 + +00 + +02 + Z = -01 + +02 + Z = 	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 - 76. -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 5.20E+00 - 73.
A = 197 $A = 197$ $A = 197$ $A = 182$ $S = 16$ $A = 196$ $A = 196$ $A = 196$ $S = 29$ $$	0.0870 - 	+/- 0.(******* Nuclide Z = 8:2 0 +/- (0 +/- (0 +/- (1 +/- 3 I +/- 3 I +/- 2 I	**************************************	***** [mb] 5.548.80 1.63 7.69 0.00 0.00 - 0.00 1.37 	<pre>****** (z if t=+0 00E+0 00E+0 00E+0 00E+0 00E+0 00E+0 00E+0 '1E+0</pre>	****** ero v Z = 0 +/- 0 +/- 1 +/- 1 +/- 2 = 0 +/- 2 +/- 2 +/- Z = 	<pre>***** valu 80. 9. 1 1 1 3 77 0</pre>	**************************************	4.889E 3.748E 0.000E 0.000E 0.000E 0.000E 1.974E 0.000E 1.974E 0.000E 1.974E	******) *** Z = +00 + +01 + +00 + +02 + Z = -01 + +00 + Z = 	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 5.20E+00 - 73.
A = 197 A = 197 A = 196 A = 186 A = 185 A = 185 A = 168 S = 29	0.0870 - 	<pre>//- 0.(//</pre>	**************************************	***** [mb] 5.548.80 1.637.69 0.000 0.000 0.000 1.37 	<pre>****** (z i1E+0 00E+0 30E-0 92E+0 00E+0 00E+0 00E+0 00E+0 '1E+0</pre>	****** Z = 0 +/- 0 +/- 1 +/- 1 +/- 2 = 0 +/- 0 +/- 2 +/- Z = 	***** ralu 80. 9. 1. 1. 3. 77. 0. 0. 4. 74. 74. 	**************************************	4.889E 3.748E 3.748E 0.000E 1.974E 0.000E 0.000E 1.974E 1.657E	****** 2 = +00 + +01 + - - - - - - - - - - - - -	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 - 73.
A = 197 A = 196 A = 196 A = 186 A = 182 S = 16 A = 185 C =	0.0870 - 	<pre>// - 0.(// - 0.(// + + + + + + + + + + + + + + + + + +</pre>	**************************************	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 0.00 1.37 	<pre>****** (z l1E+0 00E+0 00E+0 00E+0 00E+0 00E+0 00E+0 00E+0</pre>	****** ero v Z = 0 +/- 0 +/- 1 +/- 1 +/- Z = 0 +/- 0 +/- 0 +/- 2 +/- Z = Z = 	***** ralu 80. 9. 1. 1. 3. 77. 0. 0. 4. 74. 74. 	**************************************	4.889E 3.748E 0.000E 1.974E 0.000E 0.000E 1.974E 1.657E	******) *** Z = +00 + +01 + +00 + +02 + Z = +00 + +02 + Z = -01 + +02 + Z = 	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 - 76. -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 - 73.
A = 197 A = 197 A = 197 A = 182 S = 16 A = 196 A = 196 A = 196 S = 29 	0.0870 - 	Xuclidation Z = 8: D +/- (i) D +/- (i) D +/- (i) Z = 7!	**************************************	***** [mb] 5.54 8.80 - 1.63 7.69 0.00 0.00 - - 0.00 1.37 - - - - - - - -	<pre>****** (z l1E+0 00E+0 30E-0 30E-0 30E+0 00E+0 00E+0 00E+0</pre>	****** ero v Z = 0 +/- 0 +/- 1 +/- 1 +/- Z = 0 +/- 0 +/- 0 +/- 2 +/- 2 +/- 2 +/- 2 = Z = Z =	***** valu 80. 9. 1. • 1. • 1. • 1. • 1. • 1. • 0.	**************************************	4.839E 0.000E 0.00E 0.000E 0.000E 0.000E 0.000E 0.000E 0.000E 0.000E 0.	******) *** Z = +00 + +01 + +00 + +02 + Z = +00 + +02 + Z = Z = Z =	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 -/- 2.82E-01 -/- 5.20E+00 -73.
A = 197 A = 196 A = 196 A = 186 S = 16 A = 185 A = 168 S = 29 	0.0870 - 	<pre>Vuclide Z = 8:</pre>	**************************************	***** [mb] 5.54 8.80 - - 1.63 7.69 0.00 0.00 - - 0.00 1.37 - - - - - - - - -	<pre>****** (z if 1E+0 00E+0 0</pre>	****** ero v Z = 0 +/- 0 +/- 1 +/- Z = 0 +/- 0 +/- 1 +/- Z = 0 +/- 0 +/- 2 - 0 +/- 2 - Z = Z = Z = Z = 	<pre>***** valu 80. 9. 1. 71. 1. 77. 0. 77. 74. 74. 74. 8. 0</pre>	**************************************	4.889E 3.748E 0.000E 0.000E 0.000E 1.657E	******) *** Z = +00 + +01 + +00 + +00 + Z = Z = 	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 - 73.
A = 197 $A = 197$ $A = 197$ $A = 182$ $S = 16$ $A = 196$ $A = 196$ $A = 168$ $S = 29$ $$ $A = 168$ $S = 29$ $$ $A = 144$ $S = 14$	0.0870	Nuclide Z = 8: 0 +/- 0. 0 +/- (0 +/- (0 +/- (1 +/- 2 Z = 7! 0 2 +/- (Z = 7! 2 2 2 0 +/- (2 +/- 8 2 2 0 +/- (2 +/- 8) 	**************************************	***** [mb] 5.544 8.80 1.63 7.69 0.00 0.00 0.00 1.37 0.000	****** (z 11E+0 00E+0 00E+0 00E+0 00E+0 00E+0 00E+0 11E+0 00E+0 00E+0	****** 2 = 0 +/- 0 +/- 1 +/- 1 +/- 2 = 0 +/- 0 +/- 2 +/- 2 = - 2 = - 2 = - 0 +/- 0 +/- 0 +/- 0 +/- 2 +/- 2 = - 2 = - 	***** valu 80. 9. 1. 9. 1. 71. 0. 0. 74. 74. 74. 80. 80. 99. 1 74. 90. 91. 92. 93. 94. 95. 96. 97.	**************************************	4.839E 3.748E 0.000E 0.000E 0.000E 0.000E 1.657E 1.657E 1.630E	****** 2 = +00 + +01 + - +00 + +02 + Z = +00 + +00 + - - - - - - - - - - - - -	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 - 73.
A = 197 A = 196 A = 196 A = 196 A = 185 S = 16 A = 168 S = 29 	0.0870 - 	Xuclidat Z = 8: 0 +/- 0: 0 +/- 0: 0 +/- 0: 0 +/- 0: 1 +/- 2: 1 +/- 2: 0 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2	**************************************	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 0.00 1.37 0.00 0.00 0.00 0.	<pre>****** (z L1E+0)00E+0 00E+0 00E+0 00E+0 00E+0 00E+0</pre>	****** ero v Z = 0 +/- 0 +/- 1 +/-1 2 = 0 +/- 0 +/- 2 +/- Z = Z = 0 +/- 0 +/- 0 +/- 2 +/- 0 +/- 0 +/- 0 +/- 2 +/- 0 +/- 0 +/- 0 +/- 0 +/- 1 +/- 2 = 0 +/- 0 +/- 2 = 0 +/- 0 +/- 2 = 0 +/- 0 +/- 2 = 0 +/- 0 +/- 0 +/- 0 +/- 2 = 0 +/- 0 +/- 0 +/- 2 = 0 +/- 0 +/- 0 +/- 0 +/- 0 +/- 0 +/- 2 = 0 +/- 0 +/- 0 +/- 2 = 0 +/- 0 +	<pre>**** valu 80. 9. 1. 30. 1. 3. 77. 3. 77. 3. 77. 4. 74. 74. 74. 74. 8. 74. 8. 74. 74. 74. 74. 74. 74. 74. 74. 74. 74</pre>	**************************************	4.889E 3.748E 3.748E 0.000E 0.000E 0.000E 1.974E 1.657E 1.657E 1.630E	******) *** Z = +00 + +01 + Z = Z = -	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 - 73.
A = 197 $A = 196$ $A = 196$ $A = 192$ $S = 16$ $A = 195$ $C =$	0.0870 - 	Y/- 0.0 Y/- 0.0 Waclidat Z = 8: 0 +/- () 0 +/- () 0 +/- () 1 +/- 2 Z = 7! 0 - 0 +/- () Z = 7! 0 - 0 +/- () Z = 7! 0 0 - 0 +/- () Z = 7!	**************************************	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 0.00 1.37 0.00 0.00	<pre>****** (z ill=+0 00E+0 00E+0 00E+0 00E+0 00E+0 00E+0 00E+0</pre>	****** 2 =	**** ralu 80. 9. 1. . 1. . <tr< td=""><td>**************************************</td><td>4.889E 3.748E 0.000E 0.000E 0.000E 1.974E 1.657E 1.657E 1.630E</td><td>******</td><td>- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 </td></tr<>	**************************************	4.889E 3.748E 0.000E 0.000E 0.000E 1.974E 1.657E 1.657E 1.630E	******	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00
A = 197 $A = 197$ $A = 197$ $A = 182$ $S = 16$ $A = 196$ $A = 196$ $A = 196$ $S = 29$ $$ $A = 168$ $S = 29$ $$ $A = 14$ $S = 14$ $S = 14$	0.0870 - 	Xuclidat Z = 8: D +/- 0(Z = 7! Z = 3! Z = 7! Z = 3! Z = 3! Z = 7! Z = 3!	**************************************	***** [mb] 5.548.80 1.63 7.69 0.00 0.00 0.00 1.37 0.00	<pre>****** (z t1E+0 000E+0 000E+0 00E+0 00E+0 '1E+0</pre>	****** ero v Z = 0 +/ 0 +/ 1 +/-1 1 +/ 2 = 0 +/ Z = Z = Z =	<pre>**** ralu 80. 9. 1. 9. 1. 3. 77. 3. 77. 74. 74. 74. 74. 74. 74. 74. 74. 74</pre>	**************************************	4.889E 3.748E 3.748E 0.000E 1.974E 0.000E 4.889E 1.657E 1.657E 1.630E	******) *** Z = +00 + +01 + +00 + +02 + Z = +00 + +02 + Z = -01 + -01 + -0	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 - 76. -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 5.20E+00 - 73. - 7. -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E-01
A = 197 $A = 196$ $A = 108$ $S = 29$ $$	0.0870 - 	Xuclidat Z = 8: 0 +/- () 1 +/- 2 2 0 +/- () 1 +/- 2 2 0 +/- () 1 +/- 2 2 0 +/- () 1 +/- 2 2	<pre>********* * yields 1</pre>	***** [mb] 5.548.80 1.63 7.69 0.00 0.00 0.00 1.37 0.00 0.00 9.68	****** (z (z (z (z (z (z) (z) (z) (z) (z) (z)	****** ero v Z = 0 +/ 0 +/ 1 +/ 1 +/ 2 = 0 +/ 2 +/ Z = Z = 0 +/ 0 +/ 2 +/ Z = Z = 2 +/ 2 = 2 +/ 2 +/	<pre>**** **** **** **** **** **** **** **</pre>	**************************************	4.889E 3.748E 0.000E 0.000E 0.000E 1.657E 1.657E 1.630E 0.000E	$\begin{array}{c} ******\\ & Z = \\ +00 + \\ +01 + \\ - \\ - \\ +00 + \\ +01 + \\ - \\ - \\ -01 + \\ +02 + \\ Z = \\ -01 + \\ +02 + \\ Z = \\ - \\ - \\ - \\ Z = \\ - \\ - \\ - \\ Z = \\ - \\ - \\ - \\ - \\ - \\ - \\ - \\ - \\ - \\$	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 - 73. - 7. -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E-01
A = 197 $A = 197$ $A = 197$ $A = 182$ $S = 16$ $A = 196$ $A = 196$ $A = 196$ $S = 29$ $$ $A = 168$ $S = 29$ $$ $A = 14$ $S = 14$ $S = 14$ $A = 14$ $A = 4$ $A = 3$	0.0870	Y/- 0.(.) Yuclidation Z = 8: 0 +/- (.) 0 +/- (.) 1 +/- 2 Z = 7: 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0	<pre>********* * yields 1</pre>	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 0.00 1.37 0.00 0.00 0.00 0.00 0.00 0.	<pre>****** (z li1E+0 000E+0 300E+0 000E+0 000</pre>	****** ero v Z = 0 +/ 1 +/- 1 +/- 1 +/- 2 = 2 = Z = Z = Z = Z = 2 = 2 = 2 = 2 = 2 = 2 = 2 = 	***** valu 80. 9. 1. • <tr< td=""><td>**************************************</td><td>4.889E 3.74E 3.74E 0.000E 0.000E 0.000E 1.974E 0.000E 1.657E 1.630E 1.630E 0.000E</td><td>******) *** Z = +00 + +01 + +00 + +02 + Z = +00 + +02 + Z = Z = Z = Z = Z = Z = Z = -</td><td>- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 -73. </td></tr<>	**************************************	4.889E 3.74E 3.74E 0.000E 0.000E 0.000E 1.974E 0.000E 1.657E 1.630E 1.630E 0.000E	******) *** Z = +00 + +01 + +00 + +02 + Z = +00 + +02 + Z = Z = Z = Z = Z = Z = Z = -	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 -73.
A = 197 $A = 196$ $A = 106$ $S = 29$ $$	0.0870 - 	Xuclidat Z = 8: 0 +/- 0: 0 +/- 0: 0 +/- 0: 1 +/- 2: 7: 1 +/- 2: 0 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 <	<pre>********* * yields 1</pre>	***** [mb] 5.54 8.80 1.63 7.69 0.00 0.00 0.00 1.37 0.00 0.00 0.00 0.00 0.00 0.00 0	<pre>****** (z ille+0 00E+0 00E+0 00E+0 00E+0 00E+0 00E+0 38E+0 03E+0 00E+0</pre>	****** ero v Z = 0 +/- 0 +/- 1 +/- 1 +/- 2 = 0 +/- 2 +/- Z = Z = 0 +/- 0 +/- 2 +/- Z = Z = 0 +/- 0 +/- 0 +/- 2 +/- - - - 	**** 80. 9. 1. 71. 0. 77. 0. 77. 0. 74. 74. 74. - 8. 0. 2. 0. 2. 0. 1. 5. 0. 2. 1. 5. 0.	**************************************	******** pressed 4.889E 3.748E 0.000E 0.000E 0.000E 1.974E 4.889E 1.657E 1.630E 0.000E 0.000E 0.634EE 1.745E	******) *** Z = +00 + +01 + -0 + +02 + Z = -01 + +00 + +02 + Z = -01 + -01 + Z = -01 + -01 + Z = +00 + +02 + Z = -01 + -01 + -02 + -02 + -02 + -02 +	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 - 73. - 7. -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E+01 -/- 1.02E+00 -/- 1.02E+00 -/- 1.02E+00
A = 197 $A = 197$ $A = 197$ $A = 182$ $S = 16$ $A = 196$ $A = 196$ $A = 168$ $S = 29$ $$ $A = 14$ $S = 14$ $A = 14$ $S = 1$ $A = 4$ $A = 3$ $A = 2$ $A = 1$	0.0870	Nuclidat Z = 8: 0 +/- 0: 1 +/- 2: 1 +/- 2: 2 +/- 0: 1 +/- 2: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 2 +/- 0: 0 +/- 0: 0 +/- 0: 0 +/- 0: 0 +/- 0: 0 +/- 0: 0 +/- 0: 0 +/- 0: 0 +/- 0: 0 +/- 0: 0 +/- 0: 0 +/- 0:	**************************************	***** [mb] 5.54 8.80 1.63 7.69 0.000 0.000 1.37 0.000 9.68 1.57 0.000	<pre>****** (z ill=+0 00E+0 0</pre>	****** ero v Z = 0 +/- 1 +/- 1 +/- Z = Z = Z = Z = Z = Z = - - - - - - - - - - - - - - - - -	***** valu 80. 9. 1. - 1. - 0. 0. 0. 0. 0. 0. 0. 0. 0. 0. 0. 0. 0. 0. 0. 0. 2. 0. 1. 5. 0. 1. 5. 0. 0.	**************************************	4.889E 3.748E 3.748E 0.000E 0.000E 0.000E 1.657E 1.630E 1.630E 0.000E 6.346E 1.745E	******) *** Z = = +00 + +01 + +00 + +02 + Z = Z = Z = Z = Z = +00 + +02 + +02 + +02 + +02 + - - Z = Z = - 	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 -/- 2.82E-01 -/- 5.20E+00 -/- 2.82E-01 -/- 5.20E+00
A = 197 $A = 197$ $A = 197$ $A = 196$ $A = 196$ $A = 196$ $A = 195$ $A = 168$ $S = 29$ $A = 168$ $S = 29$ $A = 14$ $S = 1$ $A = 14$ $S = 1$ $A = 4$ $A = 3$ $A = 2$ $A = 1$	0.0870	Xuclidation Z = 8: D +/- 0: Z = 78: Z + 74:	<pre>********* * yields 1</pre>	***** [mb] 5.54 8.80 0.00 0.00 0.00 1.37 0.00 0.00 0.00 0.00 0.00 0.00	<pre>***** (z (z (z (z (z (z (z</pre>	****** erro v Z = 0 +/- 1 +/- 1 +/- Z = Z = Z =	***** ralu 80. 9. 1. . 1. . <t< td=""><td>**************************************</td><td>4.889E 3.748E 3.748E 0.000E 0.000E 0.000E 1.974E 1.657E 1.657E 1.630E 0.000E 0.000E 1.630E 0.000E 0.000E 1.745E 3.838E 0.2177</td><td>******) **** Z = +00 + +00 + +00 + -01 + -00 + -01 + -01 + Z = -01 + -01 + Z = -01 + -01 + -01 + -02 +</td><td>- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 - 76. -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 - 73. - 7. -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E+01 -/- 1.69E+01 -/- 1.69E+01 -/- 3.8E+01 -/- 3.8E+01</td></t<>	**************************************	4.889E 3.748E 3.748E 0.000E 0.000E 0.000E 1.974E 1.657E 1.657E 1.630E 0.000E 0.000E 1.630E 0.000E 0.000E 1.745E 3.838E 0.2177	******) **** Z = +00 + +00 + +00 + -01 + -00 + -01 + -01 + Z = -01 + -01 + Z = -01 + -01 + -01 + -02 +	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 - 76. -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 - 73. - 7. -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E+01 -/- 1.69E+01 -/- 1.69E+01 -/- 3.8E+01 -/- 3.8E+01
A = 197 $A = 196$ $A = 196$ $A = 196$ $A = 196$ $A = 195$ $A = 168$ $S = 29$ $$ $A = 14$ $S = 11$ $A = 4$ $A = 3$ $A = 2$ $A = 1$ $S = 4$	0.0870 - 	Nuclidat Z = 8: 0 +/- () 1 +/- 2 2 0 +/- () 1 +/- 2 2 0 +/- () 1 +/- 2 2 0 +/- () 1 +/- 2 2 0 +/- () 2 + - 7: 2	<pre>********* * yields 1</pre>	***** [mb] 5.548.80 1.637.69 0.000 0.000 0.000 1.37 0.000000	<pre>***** (z (z (z (z (z (z (z</pre>	****** ero v Z = 0 0 +/- 0 	**** valu 80. 9. 1. 3. 77. 0. 0. 74. 74. 74. 74. 0. 0. 1. 0. 1. 0. 1. 0.	**************************************	4.889E 3.748E 3.748E 0.000E 0.000E 0.000E 1.657E 1.657E 1.630E 0.000E 6.34E 0.383E 6.217E	******) *** = 0 + +00 +	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 - 73. - 7. -/- 1.63E-01 -/- 1.63E-01 -/- 1.62E+01 -/- 1.02E+01 -/- 1.69E+01 -/- 1.69E+01 -/- 3.18E+01
A = 197 $A = 197$ $A = 197$ $A = 182$ $S = 16$ $A = 196$ $A = 196$ $A = 196$ $A = 195$ $A = 168$ $S = 29$ $A = 14$ $S = 14$ $A = 3$ $A = 14$ $S = 1$ $A = 4$ $A = 3$ $A = 2$ $A = 1$ $S = 4$	0.0870	<pre>\Lambda Viet (1) (1) (1) (1) (1) (1) (1) (1) (1) (1)</pre>	<pre>********* * yields 1</pre>	***** [mb] 5.544 8.80 0.00 0.00 0.00 1.37 0.00 0.00 0.00 0.00 0.0	<pre>****** (z ilite+0 i0e+0 i=- i0e+0 i0e+0 i0e+0 i0e+0 i1e+0 i1e+0 i1e+0 i2e+0 i0e+0 i2e+0 i0e+0 i3e+0 i3e+0 i0e+0 i3e+0 i0e+0 i3e+0 i0e+0 i3e+0 i0e+0 i3e+0 i3</pre>	****** Z = Z = Z = Z = Z = Z =	80. 90. 1. 77. 0. 1. 3. 77. 0. - 0. - 0. - 0. - - 0. - 0. - 8. 0. - 8. 0. - 8. 0. - 8. 0. -	**************************************	4.889E 3.748E 3.748E 0.000E 1.974E 0.000E 1.974E 4.889E 1.657E 1.657E 1.630E 1.630E 0.000E 6.346E 1.745E 3.838E 6.217E	$\begin{array}{c} ****** \\ ****** \\ & \\ & \\ & \\ & \\ & \\ &$	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 5.20E+00 -73.
A = 197 $A = 196$ $S = 29$ $$	0.0870 - 	Xuclidat Z = 8: 0 +/-0(0 +/-0(0 +/-0(1 +/-2 2 = 7: 1 +/-2 2 +/-0(2 +/-0(2 +/-0(2 +/-0(2 +/-0(2 2 +/-0(2	<pre>********* * yields 1</pre>	***** [mb] 5.544 8.80 1.63 7.69 0.00 0.00 1.37 0.00 0.00 0	<pre>***** (z (z (z (z (z (z (z</pre>	****** ero v Z = ' 0 +/ 1 +/- 1 1 +/- 2 2 = ' 0 +/ 0 +/ 0 +/ 0 +/ 2 +/- 2 Z = - Z = - - 	x*** xalu 80. 9. 1. 3. 77. 0. - 0. - 74. - 74. - 88. 0. - 8. 0. 2. 1. 5. 0. 2. 1. 5. 0. 1. 5. 0. 1. 5. 0. 1.	**************************************	******** pressed 4.889E 3.748E 0.000E 0.000E 0.000E 1.974E 0.000E 1.974E 1.657E 1.657E 1.630E 0.00E 0.00E	$\begin{array}{c} ******\\) & ***\\ Z &=\\ +00 &+\\ +01 &+\\\\ +00 &+\\ +00 &+\\ +00 &+\\ +00 &+\\ +00 &+\\ +00 &+\\ +00 &+\\ +00 &+\\\\1 &-\\\\\\\\\\\\\\\\\\\\$	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 2.82E-01 -/- 2.82E-01 -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E-01 -/- 1.63E+01 -/- 1.02E+00 -/- 1.02E+00 -/- 1.02E+01 -/- 2.50E+01 -/- 3.18E+01
A = 197 $A = 197$ $A = 197$ $A = 182$ $S = 16$ $A = 196$ $A = 196$ $A = 196$ $S = 29$ $$ $A = 14$ $S = 1$ $A = 4$ $A = 3$ $A = 2$ $A = 1$ $S = 4$ $A = 1$	0.0870 - ************************************	Y/- 0.(.) Xuelidation Z = 8: 0 +/-(.) 0 +/-(.) 1 +/-2 Z = 7! 0 +/-(.)	**************************************	***** [mb] 5.544 8.80 - - 1.63 7.69 0.00 0.00 1.37 - - - - - - - - - - - - - - - - - 0.00 0 0.000000	<pre>(z (z (</pre>	***** ero V Z = V 0 +/ 1 +/- 1 +/- 2 +/ 2 +/	**** *alu 80. 91. - 1. 3. 77. 0. - 0. - 0. - 74. - 74. - 74. - 8. 0. - 8. 0. 1. 5. 0. 1. 5. 0. 1. 5. 0. 1. 5. 0. 1. 5. 0. 1.	**************************************	4.889E 3.748E 3.748E 0.000E 0.000E 0.000E 1.974E 0.000E 1.974E 1.657E 1.657E 1.630E 1.630E 0.000E 6.346E 0.217E	******) *** = 0 +00 +00 +00 +00 +00 +00 +00 +	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 -/- 2.82E-01 -/- 5.20E+00 -/- 2.82E-01 -/- 5.20E+00
A = 197 $A = 197$ $A = 196$ $A = 196$ $A = 196$ $A = 196$ $A = 195$ $C =$	0.0870	Xuclide Z = 8: 0 +/- 0: 0 +/- 0: 0 +/- 0: 1 +/- 2: 2 + 7: 2 + 7: 2 + 7: 2 + 7: 2 + 7: 2 + 7: 2 + 7: 2 + 7: 2 + 7: 2 + 7: 2 0 + 7. 0 + 7. 0 + 7. 0 + 7. 0 0 + 7. 0 0 + 7. 0	********* • yields 1. 0.00E+00 0.00E+00 0.00E+00 3. 0.00E+00 3. 0.00E+00 5. 0.00E+00	***** [mb] 5.544 8.80 - - 1.63 7.69 0.00 0.00 1.37 - - - - - - - - - - - - - - - - - - -	<pre>***** (z (z (z (z (z (z (z</pre>	****** ero v Z = / / / / / / / / / / / / / / / / / /	**** 80. 9. 1. - 1. - 1. - 0. - 0. - 0. - 0. - - 0. - 8. 0. - 8. 0. 1. 5. 0. 1. 5. 0. 1. 5. 0. 1. 5. 0. 1. 5. 0. 1. 5. 0. 1. 5. 0. 1. 5. 0. 1. 5. 6. 6. 7. 7. <t< td=""><td>**************************************</td><td>******** pressed 4.889E 3.748E 0.000E 0.000E 1.974E 0.000E 1.974E 1.657E 1.657E 1.657E 0.000E 0.000E 1.630E 0.000E 0.000E 1.974E 1.630E 1.630E 0.000E 0.000E 1.974E 0.000E 0.00</td><td>$\begin{array}{c} ******\\ & \\ & \\$</td><td>- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 - 76. -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 -/- 73. </td></t<>	**************************************	******** pressed 4.889E 3.748E 0.000E 0.000E 1.974E 0.000E 1.974E 1.657E 1.657E 1.657E 0.000E 0.000E 1.630E 0.000E 0.000E 1.974E 1.630E 1.630E 0.000E 0.000E 1.974E 0.000E 0.00	$\begin{array}{c} ******\\ & \\ & \\ & \\ & \\ & \\ & \\ & \\ & \\$	- 79. -/- 8.93E-01 -/- 2.47E+00 -/- 0.00E+00 -/- 5.67E+00 - 76. -/- 0.00E+00 -/- 0.00E+00 -/- 2.82E-01 -/- 2.82E-01 -/- 5.20E+00 -/- 73.

End of nuclide yields.

Mass yield [mb] and the mean and variance of the kinetic energy $[{\tt MeV}]$ of residual nuclei:

A = 197 1.043E+01 +/- 1.30E+00 1.227E-01 +/- 1.28E-01 A = 196 4.661E+01 +/- 2.76E+00 7.283E-02 +/- 9.35E-02 A = 1 2.732E+04 +/- 6.67E+01 2.912E+01 +/- 8.56E+01 S = 156 3.252E+04 +/- 7.28E+01 2.829E+01 +/- 7.98E+01 Charge yield [mb] and the mean and variance of the kinetic energy [of residual nuclei:

Charge yield [mb] and the mean and variance of the kinetic energy [MeV] of residual nuclei: Z = 81 8.148E-01 +/- 3.64E-01 6.399E-01 +/- 4.53E-01 Z = 80 7.692E+01 +/- 3.54E+00 4.428E-01 +/- 4.24E-01 Z = 0 2.348E+04 +/- 6.19E+01 2.072E+01 +/- 6.90E+01 S = 69 3.252E+04 +/- 7.28E+01 2.829E+01 +/- 7.98E+01

Elapsed cpu time = 0. min and 8.608 sec.

Wed Feb 1 10:54:39 2012

Example No. 8: Yields, mean kinetic energy, angles of emission, and much more (the most complete output) of all products measured at GSI in inverse kinematics for 1000 MeV p + Fe56; 10,000 events. Number of types of evaporated particles = 6

M TO A Z Q B limc idel 0.9383 1.0000 56. 26. 1 1 10000 1

dt0 = -20.0, t0max = 1000.5, dteta = 10.0mspec mpyld mchy misy mdubl mang ipar1 ipar2 0 3 0 0 2 1 rOm = 1.2, & cevap = 12.0. lim = 100000 . Geometrical cross section = 1367.65 mb. Inelastic cross section used here = 735.39 mb Monte Carlo inelastic cross section = 744.87 mb Wed Feb 1 10:54:39 2012 1000.0 MeV (Z = 1, A = 1) + (Z = 26., A = 56.) Number of inelastic interactions = 10000, Number of elastic interactions = 8361, Reaction cross section = 735.39 mb, Elastic cross section = 614.86 mb. The mean excitation energy, charge, mass, and angular momentum of the 10000 nuclei after the Cascade and before prequilibrium decay are: E*av = 148.9 +/-119.9 MeV; E*min = -3.0; E*max = 707.8 Zav = 24.7 +/- 1.4; Zmin = 18.; Zmax = 28. Aav = 52.1 +/- 2.8; Amin = 40.; Amax = 56. Lav = 6.6 +/- 4.3 h-bar; Lmin = 0.; Lmax = 33. The program called Fermi breakup 46 times. The mean charge, mass, and angular momentum of the 319 residual nuclei with less than 3 MeV of excitation energy after the cascade are: Zav = 26.1 +/- 0.3; Zmin = 26.; Zmax = 27. Aav = 55.1 +/- 0.4; Amin = 54.; Amax = 56. Lav = 2.2 +/- 1.3 h-bar; Lmin = 0.; Lmax = 7. The mean excitation energy, charge, mass, and angular momentum of the 9681 nuclei after preequilibrium
 of the
 9681 nuclei
 atter preequilibrium

 decay and before the start of statistical decay are:

 E*av = 128.7 +/- 108.8 MeV; E*min = 1.3;
 E*max = 689.5

 Zav = 24.1 +/- 1.8;
 Zmin = 16.;
 Zmax = 28.

 Aav = 51.0 +/- 3.4;
 Amin = 35.;
 Amax = 56.

 Lav = 7.7 +/- 5.1 h-bar;
 Lmin = 0.;
 Lmax = 33.

 The mean kinetic energy, charge, mass, and angular momentum of the
 10000 residual nuclei are:

 Ekav =
 4.4 +/ 5.7 MeV; Ekmin =
 0.0; Ekmax =
 52.6

 Zav =
 19.8 +/ 4.8; Zmin =
 6.; Zmax =
 27.

 Aav =
 41.7 +/ 10.4; Amin =
 13.; Amax =
 56.

 Lav =
 7.5 +/ 5.1 h-bar; Lmin =
 0.; Lmax =
 33.
 Number of coalesced d, t, He3, He4 = 4275 847 493 Mean multiplicities, yields, and mean energies of ejected particles: (Notation: T - all production mechanisms, C - cascade, P - pre-equilibrium, Sp - from spallation residues, Pf - from nuclei before fission, F - from fission fragments, E - total evaporation = Sp + Pf + F, Co - Coalescence from cascade: Values which are identically zero are not printed. <TKE> [MeV] Part. Multiplicities Yields [mb] T n 66.38 C n 155.67 P n 20.23 Sp n 6.27 En 6.27 ******* 3.8522 +/- 0.0196 2832.854 +/- 14.433 1.5569 +/- 0.0125 1144.922 +/- 9.176 0.1437 +/- 0.0038 105.675 +/- 2.788 Т р 85.07 C p 195.22 Рр 23.69 Sp p 2.1516 + / - 0.01471582.256 +/- 10.787 1582.256 +/- 10.787 9.46 2.1516 +/- 0.0147 E p 2.1516 +/- 0.0147 1582.256 +/- 10.787 9.46 35.69 T d P d

Sp d

26.83

11.68

490

Ed	0.7037 +/-	0.0084	517.491	+/-	6.169	11.68
Co d	0.4262 +/-	0.0065	313.422	+/-	4.801	77.74
******	********	*****	*******	******	*******	*****
т +	0 2611 +/-	0.0051	192 009	+/-	3 758	25 40
P +	0.0443 +/-	0.0021	32 578	+/-	1 548	30.68
	0.1202 +/-	0.0021	07 202	+/-	0 675	11 70
spi	0.1323 +/-	0.0030	91.292	+/-	2.015	11.72
Et	0.1323 +/-	0.0036	97.292	+/-	2.675	11.72
Co t	0.0845 +/-	0.0029	62.140	+/-	2.138	44.04
*******	*********	******	*******	******	*******	*****
T He3	0.2164 +/-	0.0047	159.138	+/-	3.421	29.46
P He3	0.0474 +/-	0.0022	34.857	+/-	1.601	34.40
SpHe3	0.1198 +/-	0.0035	88.099	+/-	2.545	15.67
E He3	0.1198 +/-	0.0035	88.099	+/-	2.545	15.67
CoHe3	0.0492 +/-	0.0022	36.181	+/-	1.631	58.26
******	******	*****	*******	******	******	*****
m 11 4	0 7000 . /	0.0005	507 400		0.007	45.40
I He4	0.7308 +/-	0.0085	537.420	+/-	6.287	15.18
P He4	0.0419 +/-	0.0020	30.813	+/-	1.505	43.49
SpHe4	0.6400 +/-	0.0080	470.647	+/-	5.883	12.11
E He4	0.6400 +/-	0.0080	470.647	+/-	5.883	12.11
CoHe4	0.0489 +/-	0.0022	35.960	+/-	1.626	30.99
******	******	*****	*******	******	******	*****
ni-	0 1301 +/-	0.0036	95 674	+/-	2 652	108 73
P-	0.0607 1/	0.0051	102 001	.,	2.002	115 41
p10	0.263/ +/-	0.0051	193.921	+/-	3.110	115.41
pi+	0.1871 +/-	0.0043	137.591	+/-	3.181	157.10
******	*********	******	*******	******	*******	*****

A = 55 0.000E+00 +/- 0.00E+00 1.691E+00 +/- 3.53E-01 3.250E+01 +/- 1.55E+00 _ _ _ . A = 52 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 1.030E+00 +/- 2.75E-01 S = 5 0.000E+00 +/- 0.00E+00 5.883E+00 +/- 6.58E-01 5.817E+01 +/- 2.07E+00 Z = 25. Z = 24. Z = 23. A = 55 1.633E+01 +/- 1.10E+00 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 A = 54 1.302E+01 +/- 9.78E-01 4.412E-01 +/- 1.80E-01 0.000E+00 +/- 0.00E+00 A = 46 0.000E+00 + - 0.00E+00 0.000E+00 + - 0.00E+00 1.030E+00 + - 2.75E-01S = 10 7.619E+01 +/- 2.37E+00 7.802E+01 +/- 2.40E+00 5.045E+01 +/- 1.93E+00 Z = 22. Z = 21. Z = 20. -----Z = Z = Z = 4. 3. $\begin{array}{rcl} A &=& 7 & 7.354E-02 & +/- & 7.35E-02 & 0.000E+00 & +/- & 0.00E+00 & 0.000E+00 & +/- & 0.00E+00 \\ A &=& 6 & 0.000E+00 & +/- & 0.00E+00 & 2.206E-01 & +/- & 1.27E-01 & 0.000E+00 & +/- & 0.00E+00 \\ A &=& 4 & 0.000E+00 & +/- & 0.00E+00 & 0.000E+00 & +/- & 0.00E+00 & 5.374E+02 & +/- & 6.29E+00 \\ \end{array}$ $\begin{array}{rcl} A &=& 3 & 0.000E+00 & +/- & 0.00E+00 & 0.000E+00 & +/- & 0.00E+00 & 1.591E+02 & +/- & 3.42E+00 \\ S &=& 4 & 7.354E-02 & +/- & 7.35E-02 & 2.206E-01 & +/- & 1.27E-01 & 6.966E+02 & +/- & 7.16E+00 \\ \end{array}$ 7. = 1. 7. = 0 A = 3 1.920E+02 +/- 3.76E+00 0.000E+00 +/- 0.00E+00A = 2 9.156E+02 +/- 8.21E+00 0.000E+00 +/- 0.00E+00 A = 1 2.833E+03 +/- 1.44E+01 3.468E+03 +/- 1.60E+01 S = 3 3.940E+03 +/- 1.70E+01 3.468E+03 +/- 1.60E+01 End of nuclide vields. Mass yield [mb] and the mean and variance of the kinetic energy [MeV] $% \left[M_{\rm e} \right]$ of residual nuclei: $A = 56 \ 8.751E+00 \ +/- \ 8.02E-01 \ 3.434E-01 \ +/- \ 3.79E-01$ A = 55 5.052E+01 +/- 1.93E+00 2.156E-01 +/- 3.13E-01 A = 1 6.301E+03 +/- 2.15E+01 7.478E+01 +/- 1.62E+02 S = 53 8.840E+03 +/- 2.55E+01 5.937E+01 +/- 1.41E+02 Charge yield [mb] and the mean and variance of the kinetic energy [MeV] of residual nuclei: Z = 27 5.883E+00 +/- 6.58E-01 4.566E-01 +/- 3.71E-01 Z = 26 5.817E+01 +/- 2.07E+00 3.842E-01 +/- 5.73E-01 Z = 0 3.468E+03 +/- 1.60E+01 6.638E+01 +/- 1.53E+02 S = 28 8.840E+03 +/- 2.55E+01 5.937E+01 +/- 1.41E+02

********** Nuclide yields [mb] in forward direction (theta_lab < 90) ********** (zero values suppressed)</pre>

Z = 3. 7. = 4. 7. = A = 7 7.354E-02 +/- 7.35E-02 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 A = 6 0.000E+00 +/- 0.00E+00 1.471E-01 +/- 1.04E-01 0.000E+00 +/- 0.00E+00 A = 4 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 3.214E+02 +/- 4.86E+00 A = 3 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 1.057E+02 +/- 2.79E+00 S = 4 7.354E-02 +/- 7.35E-02 1.471E-01 +/- 1.04E-01 4.272E+02 +/- 5.60E+00 A = 3 1.293E+02 + - 3.08E+00 0.000E+00 + - 0.00E+00A = 25.944E+02 + - 6.61E+00 0.000E+00 + - 0.00E+00A = $1 \cdot 1.892E+03 +/- 1.18E+01 \cdot 2.264E+03 +/- 1.29E+01$ S = $3 \cdot 2.616E+03 +/- 1.39E+01 \cdot 2.264E+03 +/- 1.29E+01$ End of nuclide yields (forward direction). Mass yield [mb] and the mean and variance of the kinetic energy [MeV] of residual nuclei in the forward direction: A = 56 8.751E+00 +/- 8.02E-01 3.434E-01 +/- 3.79E-01 A = 55 3.015E+01 +/- 1.49E+00 2.417E-01 +/- 2.89E-01 1 4.157E+03 +/- 1.75E+01 1.050E+02 +/- 1.92E+02 S = 53 5.870E+03 +/- 2.08E+01 8.170E+01 +/- 1.67E+02Charge yield [mb] and the mean and variance of the kinetic energy [MeV] of residual nuclei in the forward direction: Z = 27 5.148E+00 +/- 6.15E-01 4.243E-01 +/- 7.18E-01 Z = 26 3.765E+01 +/- 1.66E+00 4.450E-01 +/- 1.07E+00 Z = 0 2.264E+03 +/- 1.29E+01 9.405E+01 +/- 9.16E+01 S = 28 5.870E+03 +/- 2.08E+01 8.170E+01 +/- 1.67E+02 ********* Nuclide yields [mb] in backward direction (theta_lab > 90) ******** (zero values suppressed) Z = 28. Z = 27. 26. S = 4 0.000E+00 +/- 0.00E+00 7.354E-01 +/- 2.33E-01 2.052E+01 +/- 1.23E+00 Z = 25. Z = 24. Z = 23. Z = Z = Z = 3. A = 6 0.000E+00 +/- 0.00E+00 7.354E-02 +/- 7.35E-02 0.000E+00 +/- 0.00E+00 A = 4 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 2.160E+02 +/- 3.99E+00 A = A = 3 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 5.339E+01 +/- 1.98E+00 S = 3 0.000E+00 +/- 0.00E+00 7.354E-02 +/- 7.35E-02 2.694E+02 +/- 4.45E+00Z = $\begin{array}{rcl} A &=& 3 & 6.273 \pm 01 & +/- & 2.15 \pm 00 & 0.000 \pm 10 & +/- & 0.00 \pm 100 \\ A &=& 2 & 3.211 \pm 02 & +/- & 4.86 \pm 100 & 0.000 \pm 00 & +/- & 0.00 \pm 100 \\ \end{array}$ A = 1 9.406E+02 +/- 8.32E+00 1.204E+03 +/- 9.41E+00S = 3 1.324E+03 +/- 9.87E+00 1.204E+03 +/- 9.41E+00 End of nuclide yields (backward direction). Mass yield [mb] and the mean and variance of the kinetic energy [MeV] Also pictal nuclei in the backward direction: A = 55 2.037E+01 +/- 1.22E+00 1.769E-01 +/- 3.41E-01 A = 54 1.052E+01 +/- 8.79E-01 4.401E-01 +/- 5.32E-01 A = 1 2.145E+03 +/- 1.26E+01 1.615E+01 +/- 2.71E+01S = 50 2.970E+03 +/- 1.48E+01 1.524E+01 +/- 2.47E+01 Charge yield [mb] and the mean and variance of the kinetic energy [MeV] of residual nuclei in the backward direction: ------Z = 0 1.204E+03 +/- 9.41E+00 1.435E+01 +/- 2.58E+01 S = 27 2.970E+03 +/- 1.48E+01 1.524E+01 +/- 2.47E+01 ******* Nuclide average kinetic energies [MeV] (zero yield suppressed) ******** 7 = 28. 7 = 27. 7 = 26. A = 56 0.000E+00 +/- 0.00E+00 4.148E-01 +/- 3.61E-01 2.879E-01 +/- 3.84E-01 A = 55 0.000E+00 +/- 0.00E+00 4.777E-01 +/- 3.17E-01 2.224E-01 +/- 3.00E-01 Z = 25. Z = 24. Z = 2 Z = 23. - - - - - - - - -Z = 3. Z = 4. Z = 2.

 $\begin{array}{rcl} {\tt A} &=& 7 \ 9.753{\tt E}{+}00 \ +/- \ 0.00{\tt E}{+}00 \ 0.000{\tt E}{+}00 \ +/- \ 0.00{\tt E}{+}00 \ 0.000{\tt E}{+}00 \ +/- \ 0.00{\tt E}{+}00 \ \\ {\tt A} &=& 6 \ 0.000{\tt E}{+}00 \ +/- \ 0.00{\tt E}{+}00 \ 1.124{\tt E}{+}01 \ +/- \ 9.67{\tt E}{+}00 \ 0.000{\tt E}{+}00 \ +/- \ 0.00{\tt E}{+}00 \ \\ {\tt A} &=& 4 \ 0.000{\tt E}{+}00 \ +/- \ 0.00{\tt E}{+}00 \ 0.000{\tt E}{+}00 \ +/- \ 0.00{\tt E}{+}00 \ 1.518{\tt E}{+}01 \ +/- \ 1.42{\tt E}{+}01 \ \\ {\tt A} &=& 3 \ 0.000{\tt E}{+}00 \ +/- \ 0.00{\tt E}{+}00 \ 0.000{\tt E}{+}00 \ +/- \ 0.00{\tt E}{+}00 \ 1.84{\tt E}{+}01 \ +/- \ 3.2{\tt E}{+}01 \ \\ {\tt S} &=& 4 \ 9.753{\tt E}{+}00 \ +/- \ 0.00{\tt E}{+}00 \ 1.124{\tt E}{+}01 \ +/- \ 9.67{\tt E}{+}00 \ 1.844{\tt E}{+}01 \ +/- \ 2.29{\tt E}{+}01 \ \\ \end{array}$ Z = Ζ = 0. S = 3 7.069E+01 +/- 1.50E+02 6.638E+01 +/- 1.53E+02 End of nuclide average kinetic energies. Mass yield [mb] and the mean and variance of the emission angle [deg.] of residual nuclei: $\begin{array}{l} A = 56 & 8.751 \pm 00 \ +/- \ 8.02 \pm -01 \ 8.315 \pm +01 \ +/- \ 1.74 \pm +00 \ \\ A = 55 \ 5.052 \pm 01 \ +/- \ 1.93 \pm +00 \ 8.380 \pm +01 \ +/- \ 3.26 \pm +01 \ \end{array}$ The mean and variance of the z velocity [v/c] of residual nuclei, and the forward/backward ratio: A = 56 + 4.003E - 04 + 7 - 2.63E - 041.000E+00 +/- 0.00E+00 A = 55 3.056E-04 +/- 1.09E-03 1.480E+00 +/- 1.62E-01

Charge yield [mb] and the mean and variance of the emission angle [deg.] of residual nuclei:

Z = 27	5.883E+00 +/- 6.58E-01	8.332E+01 +/- 1.23E+01
Z = 26	5.817E+01 +/- 2.07E+00	8.288E+01 +/- 3.12E+01
Z = 0	3.468E+03 +/- 1.60E+01	7.434E+01 +/- 4.04E+01
S = 28	8.840E+03 +/- 2.55E+01	7.382E+01 +/- 3.97E+01

The mean and variance of the z velocity [v/c] of residual nuclei,

anu the	IOIWalu/Dackwalu latio.		
Z = 27	3.928E-04 +/- 1.01E-03	4.348E+00 +/- 2.46E+00	
Z = 26	5.406E-04 +/- 1.56E-03	2.993E+00 +/- 5.41E-01	
Z = 0	1.000E-01 +/- 2.16E-01	1.232E+01 +/- 3.27E-02	
S = 28	8.698E-02 +/- 1.99E-01	1.977E+00 +/- 1.68E-02	

Mass distributions of nuclei:

					at start of ev	ap, which:	just prior to
			after cascade	after preeq	evap. only	fission	fission
	A =	56	3.910E-02	3.160E-02	3.160E-02	0.000E+00	0.000E+00
	A =	55	1.908E-01	1.468E-01	1.468E-01	0.000E+00	0.000E+00
	A =	35	0.000E+00	1.000E-04	1.000E-04	0.000E+00	0.000E+00
	<a>	=	5.202E+01	5.102E+01	5.102E+01	0.000E+00	0.000E+00
St	Dv A	=	2.745E+00	3.445E+00	3.445E+00	0.000E+00	0.000E+00
	norm	=	1.000E+00	1.000E+00	1.000E+00	0.000E+00	0.000E+00

Charge distributions of nuclei:

				at start of evap, which:				
			after cascade	after preeq	evap. only	fission	fission	
	Z =	28	1.900E-03	1.600E-03	1.600E-03	0.000E+00	0.000E+00	
	Z =	27	5.630E-02	4.010E-02	4.010E-02	0.000E+00	0.000E+00	
	Z =	16	0.000E+00	3.000E-04	3.000E-04	0.000E+00	0.000E+00	
	<z></z>	=	2.463E+01	2.413E+01	2.413E+01	0.000E+00	0.000E+00	
St	Dv Z	=	1.445E+00	1.779E+00	1.779E+00	0.000E+00	0.000E+00	
	$\verb"norm"$	=	1.000E+00	1.000E+00	1.000E+00	0.000E+00	0.000E+00	

Excitation energy distributions [1/MeV] of nuclei:

			at start of ev	ap, which:	just prior to
E*(MeV)	after cascade	after preeq	evap. only	fission	fission
0 10.	4.380E-03	5.010E-03	5.010E-03	0.000E+00	0.000E+00
10 20.	3.630E-03	5.070E-03	5.070E-03	0.000E+00	0.000E+00
680 690.	1.000E-05	1.000E-05	1.000E-05	0.000E+00	0.000E+00
< <u>E</u> *> =	1.496E+02	1.286E+02	1.286E+02	0.000E+00	0.000E+00
St dev E* =	1.151E+02	1.086E+02	1.086E+02	0.000E+00	0.000E+00
norm =	1.000E+00	1.000E+00	1.000E+00	0.000E+00	0.000E+00

Linear momentum distributions [1/MeV/c] of nuclei: at start of evap, which: just prior to P(MeV/c) after cascade after preeq evap. only fission 0.- 20. 4.000E-05 4.500E-05 4.500E-05 0.000E+00 20.- 40. 3.050E-04 3.100E-04 3.100E-04 0.000E+00 fission 0.000E+00 0.000E+00 - - - - - -1940.-1960. 1.000E-05 1.000E-05 1.000E-05 0.000E+00 0.000E+00 <P> = 5.097E+02 5.052E+02 5.052E+02 0.000E+00 0.000E+00 St dev P = 3.426E+02 3.376E+02 3.376E+02 0.000E+00 0.000E+00 1.000E+00 1.000E+00 1.000E+00 0.000E+00 0.000E+00 norm =

Angular momentum distributions [1/hbar] of nuclei: at start of evap, which: just prior to

L	after cascade	after preeq	evap. only	fission	fission	
0 1.	4.300E-03	3.800E-03	3.800E-03	0.000E+00	0.000E+00	
1 2.	4.780E-02	4.010E-02	4.010E-02	0.000E+00	0.000E+00	

33 34.	1.000E-04	2.000E-04	2.000E-04	0.000E+00	0.000E+00
<l> =</l>	6.771E+00	7.700E+00	7.700E+00	0.000E+00	0.000E+00
St dv L =	4.291E+00	5.092E+00	5.092E+00	0.000E+00	0.000E+00
norm =	1.000E+00	1.000E+00	1.000E+00	0.000E+00	0.000E+00

Neutron-multiplicity probability:

	Nn		Total	Cascade	Preequil.	Evap. res.	Pre-fiss.	Post-fiss.
	0		3.250E-02	1.602E-01	8.701E-01	1.393E-01	0.000E+00	0.000E+00
	1		8.940E-02	3.230E-01	1.167E-01	1.943E-01	0.000E+00	0.000E+00
	15		1.000E-04	0.000E+00	0.000E+00	0.000E+00	0.000E+00	0.000E+00
	<n></n>	=	4.716E+00	1.884E+00	1.444E-01	2.688E+00	0.000E+00	0.000E+00
St	dv n	=	2.695E+00	1.500E+00	3.939E-01	2.069E+00	0.000E+00	0.000E+00
	norm	=	1.000E+00	1.000E+00	1.000E+00	1.000E+00	0.000E+00	0.000E+00

The program called Fermi breakup 46 times.

Elapsed cpu time = 0. min and 6.201 sec.

```
Wed Feb 1 10:49:05 2012
Example No. 9: Proton spectra from monochromatic 300 MeV gamma + 64Cu;
10.000 events
Number of types of evaporated particles = 6
M TO A Z Q B limc idel
0.0000 0.3000 64. 29. 0 0 10000 1
dt0 = -5.0, t0max = 300.5, dteta = 10.0
  mspec mpyld mchy misy mdubl mang ipar1 ipar2
                   0 0 1
                                           0
                                                    2
    0
            1
 rOm = 1.2, & cevap = 12.0.
    Theta1
                   Theta2
                                   Theta3
                                                    Theta4
                                                                    Theta5
                                                                                    Theta6
  42.5 47.5 87.5 92.5 132.5 137.5 -55.0 165.0 55.0 65.0 75.0 85.0
                   Theta8
                                    Theta9
    Theta7
                                                  Theta10
  95.0 105.0 115.0 125.0 135.0 145.0 155.0 165.0
Tmin, Tmax, dT{1};Tmin, Tmax, dT{2};Tmin, Tmax, dT{3};Tmin, Tmax, dT{4}.
0.0 22.0 1.00 22.0 120.0 2.00 120. 400. 10.0 400. 1000. 20.
           6000000
lim =
Geometrical cross section = 1445.99 mb.
       Inelastic cross section used here = 26.58 mb
Monte Carlo inelastic cross section = 28.02 mb
      Wed Feb 1 10:49:06 2012
                    300.0 \text{ MeV} (Z = 0, A = 0) + (Z = 29., A = 64.)
 Number of inelastic interactions =
                                                        10000.
                                                    505970,
 Number of elastic interactions =
Reaction cross section = 26.58 mb, Elastic cross section = 1344.95 mb.
  The mean excitation energy, charge, mass, and angular momentum of the 10000 nuclei after the
  cascade and before preequilibrium decay are:

      Exav = 86.6 +/- 52.5 MeV; E*min = 1.3; E*max = 299.2

      Zav = 28.3 +/- 0.9; Zmin = 24.; Zmax = 30.

      Aav = 61.8 +/- 1.4; Amin = 56.; Amax = 64.

      Lav = 4.3 +/- 2.5 h-bar; Lmin = 0.; Lmax = 17.

  The mean charge, mass, and angular momentum
  of the
                13 residual nuclei with less than
  3 MeV of excitation energy after the cascade are:

      Zav = 28.0 +/- 0.0;
      Zmin = 28.;
      Zmax = 28.

      Aav = 62.0 +/- 0.0;
      Amin = 62.;
      Amax = 62.

      Lav = 3.6 +/- 1.5 h-bar;
      Lmin = 2.;
      Lmax = 7.

  The mean excitation energy, charge, mass, and angular momentum
  of the 9986 nuclei after preequilibrium

        Of the
        9986 nuclei arter preequilibrium

        decay and before the start of statistical decay are:

        E*av = 66.1 +/- 44.2 MeV; E*min = 0.7; E*max = 298.3

        Zav = 27.8 +/- 1.1; Zmin = 22.; Zmax = 30.

        Aav = 60.9 +/- 2.0; Amin = 51.; Amax = 64.

        Lav = 5.4 +/- 3.4 h-bar; Lmin = 0.; Lmax = 35.

  Number of coalesced d, t, He3, He4 =
                                                       758
                                                                    95
                                                                               13
                                                                                          4
 Mean multiplicities, yields, and mean energies of ejected particles:
 (Notation: T - all production mechanisms, C - cascade, P -
                                                                              pre-equilibrium,
Sp - from spallation residues, Pf - from nuclei before fission,
F - from fission fragments, E - total evaporation = Sp + Pf + F,
 Co - Coalescence from cascade;
Values which are identically zero are not printed.
                                             Yields [mb] <TKE> [MeV]
            Multiplicities
 Part.
 C n 1.3350 .,
P n 0.2486 +/- 0.0050
Sp n 2.5207 +/- 0.0159 67.004 +/- 0.422 3.79
E n 2.5207 +/- 0.0159 67.004 +/- 0.422 3.79
```

Ср	0.7029 +/-	0.0084	18.684 +/-	0.223	64.50					
Рр	0.1849 +/-	0.0043	4.915 +/-	0.114	21.64					
Spp	1.0838 +/-	0.0104	28.809 +/-	0.277	7.43					
Ер	1.0838 +/-	0.0104	28.809 +/-	0.277	7.43					
******	******	*****	*****	******	******					
T d	0.3705 +/-	0.0061	9.848 +/-	0.162	16.76					
P d	0.1206 +/-	0.0035	3.206 +/-	0.092	23.68					
Sp d	0.1742 +/-	0.0042	4.631 +/-	0.111	8.99					
Ed	0.1742 +/-	0.0042	4.631 +/-	0.111	8.99					
Co d	0.0757 +/-	0.0028	2.012 +/-	0.073	23.61					
******	*******	*****	******	******	******					
T t	0.0705 +/-	0.0027	1.874 +/-	0.071	17.86					
Ρt	0.0320 +/-	0.0018	0.851 +/-	0.048	25.79					
Sp t	0.0290 +/-	0.0017	0.771 +/-	0.045	9.23					
Et	0.0290 +/-	0.0017	0.771 +/-	0.045	9.23					
Co t	0.0095 +/-	0.0010	0.253 +/-	0.026	17.50					
******	******									
T He3	0.0357 +/-	0.0019	0.949 +/-	0.050	23.34					
P He3	0.0217 +/-	0.0015	0.577 +/-	0.039	29.28					
SpHe3	0.0127 +/-	0.0011	0.338 +/-	0.030	12.94					
E He3	0.0127 +/-	0.0011	0.338 +/-	0.030	12.94					
CoHe3	0.0013 +/-	0.0004	0.035 +/-	0.010	25.82					
******	******	*****	*****	******	******					
T He4	0.3421 +/-	0.0058	9.094 +/-	0.155	13.38					
P He4	0.0162 +/-	0.0013	0.431 +/-	0.034	38.47					
SpHe4	0.3255 +/-	0.0057	8.652 +/-	0.152	12.12					
E He4	0.3255 +/-	0.0057	8.652 +/-	0.152	12.12					
CoHe4	0.0004 +/-	0.0002	0.011 +/-	0.005	18.83					
******	******	*****	*****	*****	*****					
pi-	0.1220 +/-	0.0035	3.243 +/-	0.093	55.35					
pi0	0.2211 +/-	0.0047	5.877 +/-	0.125	58.90					
pi+	0.0802 +/-	0.0028	2.132 +/-	0.075	48.01					
******	**************************************									

Double differential cross sections [mb/MeV/sr]; Lab. angle = 42.5 to 47.5 degrees.

 Tp
 [MeV]
 Total
 Cascade
 Precompound
 Total Evaporation

 3.0 4.0
 1.509E-01 +/- 3.22E-02 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 1.509E-01 +/- 3.22E-02
 4.0 5.0
 4.389E-01 +/- 5.49E-02 6.858E-03 +/- 6.86E-03 6.858E-03 +/- 6.86E-03 4.252E-01 +/- 5.40E-02

 220.0 230.0
 1.372E-03 +/- 9.70E-04 1.372E-03 +/- 9.70E-04 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00

 230.0 240.0
 6.858E-04 +/- 6.86E-04 +/- 6.86E-04 +/- 6.86E-04 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00

Integrated: 5.219E+00 +/- 1.89E-01 2.188E+00 +/- 1.22E-01 5.212E-01 +/- 5.98E-02 2.510E+00 +/- 1.31E-01

Double differential cross sections [mb/MeV/sr]; Lab. angle = 87.5 to 92.5 degrees.

 Tp
 [MeV]
 Total
 Cascade
 Precompound
 Total
 Evaporation

 2.0 3.0
 1.940E-02 +/- 9.70E-03
 0.000E+00 +/- 0.00E+00
 0.000E+00 +/- 0.00E+00
 1.940E-02 +/- 9.70E-03

 3.0 4.0
 2.861E-01 +/- 3.72E-02
 0.000E+00 +/- 0.00E+00
 4.849E-03 +/- 4.85E-01 +/- 3.859E-02
 1.969E-02 +/- 9.70E-03

 170.0 180.0
 4.849E-04 +/- 4.85E-04 4.849E-04 +/- 4.85E-04
 0.00E+00 +/- 0.00E+00
 -0.00E+00
 0.00E+00 +/- 0.00E+00

 180.0 190.0
 4.849E-04 +/- 4.849E-04 +/- 4.85E-04
 0.00E+00 +/- 0.00E+00
 0.00E+00 +/- 0.00E+00
 0.00E+00 +/- 0.00E+00

Integrated: 3.884E+00 +/- 1.37E-01 1.014E+00 +/- 7.01E-02 3.686E-01 +/- 4.23E-02 2.502E+00 +/- 1.10E-01

Double differential cross sections [mb/MeV/sr]; Lab. angle = 132.5 to 137.5 degrees.

 Tp
 [MeV]
 Total
 Cascade
 Precompound
 Total Evaporation

 2.0 3.0
 3.429E-02 +/- 1.53E-02 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 3.429E-02 +/- 1.53E-02
 3.0 4.0
 3.223E-01 +/- 4.70E-02 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 3.223E-01 +/- 4.70E-02

 130.0 140.0
 2.743E-03 +/- 1.37E-03 2.743E-03 +/- 1.37E-03 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00
 0.00E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00

 140.0 150.0
 6.588E-04 +/- 6.868E-04 +/- 6.868E-04 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00
 0.00E+00 +/- 0.00E+00

Integrated: 3.381E+00 +/- 1.52E-01 8.710E-01 +/- 7.73E-02 3.018E-01 +/- 4.55E-02 2.208E+00 +/- 1.23E-01

Elapsed cpu time = 0. min and 16.136 sec.

```
Wed Feb 1 10:48:33 2012
Example No. 10: Yields, mean kinetic energy, emission angles,
neutron multiplicities, Forward/Backward ratios, and much more
(the most complete output) of all products from E_max = 1000 MeV
bremsstrahlung gammas + 197Au; 10,000 events.
 Number of types of evaporated particles =
 M TO A Z Q B limc idel 0.0000 0.4595 197. 79. 0 0 10000 1
 dt0 = -20.0, t0max = 1000.0, dteta = 10.0
   mspec mpyld mchy misy mdubl mang ipar1 ipar2
                                       3
                                                  0
 rOm = 1.2, & cevap = 12.0.
 lim = 6000000 .
 Geometrical cross section = 2394.46 mb.
   Number of equivalent gamma quanta = 3.220898E-01
  Inelasic cross section per eqqv = 1.389253E+02
Averaged absorption cross section = 4.474643E+01
   Results are normalized to eqqv.
 Inelastic cross section used here = 138.93 mb
Monte Carlo inelastic cross section = 35.37 mb
        Wed Feb 1 10:48:33 2012
                            459.5 \text{ MeV} (Z = 0, A = 0) + (Z = 79., A = 197.)
 Number of inelastic interactions =
                                                                            10000
 Number of elastic interactions =
                                                                           666883.
 Reaction cross section = 138.93 mb, Elastic cross section = 9264.69 mb.
   The mean excitation energy, charge, mass, and angular momentum
   of the 10000 nuclei after the cascade and before preequilibrium decay are:

      Cascade and before preddifiorium decay are:

      E*av = 126.4 +/-104.9 MeV;
      E*min = 1.2;
      E*max = 78.6.4

      Zav = 78.6 +/-
      0.8;
      Zmin = 73.;
      Zmax = 81.

      Aav = 194.7 +/-
      2.2;
      Amin = 182.;
      Amax = 197.

      Lav = 6.0 +/-
      4.6 h-bar;
      Lmin = 0.;
      Lmax = 40.

   The mean charge, mass, and angular momentum
                          2 residual nuclei with less than
   of the
  of the 2 residual nuclei with less than

3 MeV of excitation energy after the cascade are:

Zav = 79.0 +/- 0.0; Zmin = 79.; Zmax = 79.

Aav = 196.0 +/- 0.0; Amin = 196.; Amax = 196.

Lav = 3.5 +/- 2.5 h-bar; Lmin = 1.; Lmax = 6.
   The mean excitation energy, charge, mass, and angular momentum
   of the 9998 nuclei after preequilibrium decay and before the start of statistical decay are:

      Exav = 92.9 +/-
      87.4 MeV; E*min = 0.2; E*max = 713.4

      Zav = 78.1 +/-
      1.2; Zmin = 70.; Zmax = 81.

      Aav = 193.5 +/-
      3.2; Amin = 175.; Amax = 197.

      Lav = 8.1 +/-
      6.0 h-bar; Lmin = 0.; Lmax = 61.

   The mean kinetic energy, charge, mass, and angular momentum

      In a word window of the 9799 residual nuclei are:

      Ekav = 0.6 +/- 1.1 MeV; Ekmin = 0.0; Ekmax = 21.9

      Zav = 77.3 +/- 2.5; Zmin = 60.; Zmax = 81.

      Aav = 185.3 +/- 9.4; Amin = 134.; Amax = 196.

      Lav = 8.0 +/- 6.0 h-bar; Lmin = 0.; Lmax = 61.

   The mean excitation energy, charge, mass, angular momentum, and fission barrier height of the 201 fissioning nuclei are:
  The mean excitation energy, charge, mass, angular momentum, fission barrier height of the 201 fissioning nuclei are:

Exav = 203.9 +/- 96.6 MeV; E*min = 46.2; E*max = 487.2

Zav = 77.0 +/- 2.5; Zmin = 68.; Zmax = 81.

Aav = 186.8 +/- 6.6; Amin = 165.; Amax = 196.

        Aav = 186.8 +/- 6.6;
        Amin = 165.;
        Amax = 196.

        Lav = 10.8 +/- 7.1 h-bar;
        Lmin = 1.;
        Lmax = 40.

        Bfav = 18.4 +/- 2.2 MeV;
        Bfmin = 12.7;
        Bfmax = 24.8

 The mean total fission product kinetic energy after neutron emission is 122.90 MeV.
                          Direct Monte Carlo Simulation Method:
 Fissility = 0.0201 +/- 0.0014,
Fission cross section = 2.79240E+00 +/- 1.97E-01 mb.
                          Statistical Weight Functions Method:
 Fissility = 0.0170,
 Fission cross section = 2.36377E+00 mb.
  Number of coalesced d, t, He3, He4 =
                                                                        1051
                                                                                          333
                                                                                                            49
                                                                                                                           40
```

Mean multiplicities, yields, and mean energies of ejected particles: (Notation: T - all production mechanisms, C - cascade, P - pre-equilibrium, Sp - from spallation residues, Pf - from nuclei before fission, F - from fission fragments, E - total evaporation = Sp + Pf + F, Co - Coalescence from cascade; Values which are identically zero are not printed.

Pai	t.	Multi	plicit	ies	Yie	lds	[mb]	<tke> [MeV</tke>]
***	*****	******	*****	*****	**********	****	*******	**********	****
Т	n	9.1639	+/- 0	.0303	1273.098	+/-	4.206	10.78	
С	n	1.6086	+/- 0	.0127	223.475	+/-	1.762	41.52	
Ρ	n	0.5664	+/- 0	.0075	78.687	+/-	1.046	19.26	
Sp	n	6.6590	+/- 0	.0258	925.104	+/-	3.585	2.93	
Ρf	n	0.0553	+/- 0	.0024	7.683	+/-	0.327	6.19	
F	n	0.2746	+/- 0	.0052	38.149	+/-	0.728	4.37	
Е	n	6.9889	+/- 0	.0264	970.935	+/-	3.673	3.01	
***	*****	******	*****	*****	*****	****:	******	*****	****
т	n	0.9682	+/- 0	.0098	134.507	+/-	1.367	44.12	
ĉ	P D	0 3384	+/- 0	0058	47 012	+/-	0 808	88 78	
D	P	0.0001	+/- 0	0054	40.538	+/-	0.000	20.23	
1 (7m)	P	0.2310	+/- 0	0054	40.000	÷/_	0.750	11 01	
ър	P	0.3100	+/- 0	00000	44.202		0.704	14 54	
PI	р	0.0048	+/- 0	.0007	0.667	+/-	0.096	14.54	
F	р	0.0146	+/- 0	.0012	2.028	+/-	0.168	9.23	
Е	р	0.3380	+/- 0	.0058	46.957	+/-	0.808	11.74	
***	*****	*****	*****	*****	********	****:	******	******	****
Т	d	0.3494	+/- 0	.0059	48.541	+/-	0.821	28.84	
Ρ	d	0.1196	+/- 0	.0035	16.615	+/-	0.480	31.04	
Sp	d	0.1179	+/- 0	.0034	16.379	+/-	0.477	12.92	
Ρf	d	0.0042	+/- 0	.0006	0.583	+/-	0.090	15.72	
F	d	0.0036	+/- 0	.0006	0.500	+/-	0.083	11.02	
Е	d	0.1257	+/- 0	.0035	17.463	+/-	0.493	12.95	
Co	d	0.1041	+/- 0	.0032	14.462	+/-	0.448	45.50	
***	*****	******	*****	*****	*********	****:	******	******	****
т	t	0.1152	+/- 0	0034	16.004	+/-	0.472	22.79	
Þ	+	0.0284	+/- 0	0017	3 945	+/-	0.234	33 13	
	+	0.0201	+/- 0	00022	7 071	÷/_	0.201	12 67	
ъ£	ι +	0.0009	+/- 0	0023	0.000	+/-	0.313	12.07	
F1	ι -	0.0015	+/- 0	0004	0.208		0.054	10.19	
r	τ.	0.0016	+/- 0	.0004	0.222	+/-	0.056	12.10	
E	t	0.0540	+/- 0	.0023	7.502	+/-	0.323	13.63	
Co	t	0.0328	+/- 0	.0018	4.557	+/-	0.252	28.91	
***	*****	*****	*****	*****	*********	****:	*****	******	****
Τŀ	le3	0.0261	+/- 0	.0016	3.626	+/-	0.224	42.07	
Ρŀ	le3	0.0168	+/- 0	.0013	2.334	+/-	0.180	47.29	
Spł	le3	0.0042	+/- 0	.0006	0.583	+/-	0.090	23.53	
PfH	le3	0.0001	+/- 0	.0001	0.014	+/-	0.014	25.09	
FΗ	le3	0.0001	+/- 0	.0001	0.014	+/-	0.014	13.66	
Εŀ	le3	0.0044	+/- 0	.0007	0.611	+/-	0.092	23.34	
CoF	le3	0.0049	+/- 0	.0007	0.681	+/-	0.097	40.97	
***	*****	******	,	*****	**********	, ****:	*******	*********	****
ΤF	le4	0.1774	+/- 0	.0042	24,645	+/-	0.585	24.18	
p L	104	0 0074	+/- 0	0000	1 028	+/-	0 120	56 90	
I I Gri	101	0 1569	+/- 0	0040	21.020	+/-	0.120	20.30	
Dfi	104	0.1000	+/- 0	00040	21.703	+/-	0.000	22.73	
PII	104	0.0041	+/- 0	0000	0.570	+/-	0.069	25.74	
r 1	164	0.0051	+/- 0	.0007	0.709	+/-	0.099	17.39	
Εŀ	le4	0.1660	+/- 0	.0041	23.062	+/-	0.566	22.64	
Coł	le4	0.0040	+/- 0	.0006	0.556	+/-	0.088	27.64	
***	*****	*****	*****	*****	*****	****	******	*****	****
pi-	-	0.1199	+/- 0	.0035	16.657	+/-	0.481	100.65	
pi)	0.1332	+/- 0	.0036	18.505	+/-	0.507	89.72	
pi+	+	0.0513	+/- 0	.0023	7.127	+/-	0.315	130.85	
***	*****	******	*****	*****	**********	****:	******	*******	****
************** Nuclide yields [mb] (zero values suppressed) ********************									
			Z =	81.		Z =	80.		Z =
	100	0.000	~~ · /	0 001			/ 0 00F	00 4 4465	o1 . /

196 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 4.446E-01 +/- 7.86E-02 A = 196 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 4.446E-01 +/- 7.86E-02 A = 195 0.000E+00 +/- 0.00E+00 1.667E-01 +/- 4.81E-02 2.987E+00 +/- 2.04E-01 A = 183 0.000E+00 +/- 0.00E+00 1.389E-02 +/- 1.39E-02 1.389E-02 +/- 1.39E-02 $S = 14 \ 1.389E-02 \ +/- \ 1.39E-02 \ 3.779E+00 \ +/- \ 2.29E-01 \ 4.232E+01 \ +/- \ 7.67E-01$

Z = 78. Z = 77. Z = 76. A = 196 2.362E-01 +/- 5.73E-02 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 A = 195 1.320E+00 +/- 1.35E-01 1.389E-02 +/- 1.39E-02 0.000E+00 +/- 0.00E+00 A = 169 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 1.389E-02 +/- 1.39E-02 S = 28 4.303E+01 +/- 7.73E-01 1.595E+01 +/- 4.71E-01 1.136E+01 +/- 3.97E-01

Z = 74. Z = 75. Z = 73.

End of nuclide yields.

Mass vield [mb] and the mean and variance of the kinetic energy [MeV] of residual nuclei: A = 196 6.807E-01 +/- 9.72E-02 1.629E-01 +/- 9.21E-02 A = 195 4.487E+00 +/- 2.50E-01 1.524E-01 +/- 1.40E-01 _ _ _ _ _ _ _ _ _ - - - -- - - - - - - - -Charge vield [mb] and the mean and variance of the kinetic energy [MeV] of residual nuclei: Z = 81 1.389E-02 +/- 1.39E-02 2.362E-01 +/- 0.00E+00 Z = 80 3.779E+00 +/- 2.29E-01 3.332E-01 +/- 3.29E-01 $Z = 0 \ 1.273E+03 \ +/- \ 4.21E+00 \ 1.078E+01 \ +/- \ 2.84E+01$ $S = 61 \ 1.642E+03 \ +/- \ 4.78E+00 \ 1.376E+01 \ +/- \ 3.26E+01$ ********** Nuclide yields [mb] in forward direction (theta_lab < 90) ********* (zero values suppressed) 7 = 81. Z = 80. Z = 79. A = 196 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 1.806E-01 +/- 5.01E-02 A = 195 0.000E+00 +/- 0.00E+00 1.667E-01 +/- 4.81E-02 1.514E+00 +/- 1.45E-01 - - - -A = 183 0.000E+00 +/- 0.00E+00 1.389E-02 +/- 1.39E-02 1.389E-02 +/- 1.39E-02 S = 14 1.389E-02 +/- 1.39E-02 3.459E+00 +/- 2.19E-01 2.613E+01 +/- 6.03E-01 Z = 78. Z = 77. Z = 76. A = 196 4.168E-02 +/- 2.41E-02 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 A = 195 4.862E-01 +/- 8.22E-02 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 A = 169 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 1.389E-02 +/- 1.39E-02 S = 27 2.455E+01 +/- 5.84E-01 9.878E+00 +/- 3.70E-01 7.669E+00 +/- 3.26E-01 Z = 75. 7 = 74. 7 = 73. 7 = 3 7. = 2 7. = S = 4 0.000E+00 +/- 0.00E+00 1.496E+01 +/- 4.56E-01 1.217E+02 +/- 1.30E+00 Ζ = End of nuclide yields (forward direction). Mass yield [mb] and the mean and variance of the kinetic energy [MeV] of residual nuclei in the forward direction: A = 196 2.223E-01 +/- 5.56E-02 2.082E-01 +/- 1.25E-01 A = 195 2.167E+00 +/- 1.74E-01 1.471E-01 +/- 1.28E-01 Charge yield [mb] and the mean and variance of the kinetic energy [MeV]
 Z = 81
 1.389E-02
 +/ 1.39E-02
 2.362E-01
 +/ 4.25E-01

 Z = 80
 3.459E+00
 +/ 2.19E-01
 3.408E-01
 +/ 7.41E-01
 $Z = 0 \ 6.777E+02 \ +/- \ 3.07E+00 \ 1.329E+01 \ +/- \ 8.47E+00$ $S = 54 \ 9.031E+02 \ +/- \ 3.54E+00 \ 1.693E+01 \ +/- \ 3.97E+01$ ********* Nuclide yields [mb] in backward direction (theta_lab > 90) ********* (zero values suppressed) 7. = 81. 7. = 80. Z. = A = 196 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 2.640E-01 +/- 6.06E-02 A = 195 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 1.473E+00 +/- 1.43E-01 7 = 78. 7 = 77. 7 = 76. A = 196 1.945E-01 +/- 5.20E-02 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 A = 195 8.336E-01 +/- 1.08E-01 1.389E-02 +/- 1.39E-02 0.000E+00 +/- 0.00E+00 Z = 75. Z = 74. Z = 73.

7 = 3 7 = 2 7 = 1. 4 0.000E+00 +/- 0.00E+00 1.179E+01 +/- 4.05E-01 0.000E+00 +/- 0.00E+00 A = A = 3 0.000E+00 +/- 0.00E+00 1.514E+00 +/- 1.45E-01 6.460E+00 +/- 3.00E-01 A = 2 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 2.030E+01 +/- 5.31E-01 A = 1 0.000E+00 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 5.060E+01 +/- 8.38E-01 4 0.000E+00 +/- 0.00E+00 1.331E+01 +/- 4.30E-01 7.735E+01 +/- 1.04E+00 S = 0. A = 1 5.954E+02 +/- 2.88E+00S = 1 5.954E+02 +/- 2.88E+00 End of nuclide yields (backward direction). Mass yield [mb] and the mean and variance of the kinetic energy [MeV] of residual nuclei in the backward direction: A = 196 4.585E-01 +/- 7.98E-02 1.409E-01 +/- 5.97E-02 A = 195 2.320E+00 +/- 1.80E-01 1.574E-01 +/- 1.50E-01 1 6.460E+02 +/- 3.00E+00 9.552E+00 +/- 2.02E+01 A = $S = 123 \ 7.390E+02 \ +/- \ 3.20E+00 \ 9.892E+00 \ +/- \ 2.00E+01$ Charge vield [mb] and the mean and variance of the kinetic energy [MeV] of residual nuclei in the backward direction: Z = 80 3.195E-01 +/- 6.66E-02 2.509E-01 +/- 2.70E-01 Z = 79 1.618E+01 +/- 4.74E-01 1.447E-01 +/- 1.72E-01 Z = 0 5.954E+02 +/- 2.88E+00 7.918E+00 +/- 1.80E+01 S = 56 7.390E+02 +/- 3.20E+00 9.892E+00 +/- 2.00E+01 ******* Nuclide average kinetic energies [MeV] (zero yield suppressed) ******* Z = 80. A = 195 0.000E+00 +/- 0.00E+00 2.419E-01 +/- 2.74E-01 1.210E-01 +/- 8.70E-02 A = 183 0.000E+00 +/- 0.00E+00 2.226E-01 +/- 0.00E+00 2.385E+00 +/- 0.00E+00 S = 14 2.362E-01 +/- 0.00E+00 3.332E-01 +/- 3.29E-01 1.754E-01 +/- 2.32E-01 Z = 77. Z = 78. 76. A = 196 1.484E-01 + - 6.31E-02 0.000E+00 + - 0.00E+00 0.000E+00 + - 0.00E+00A = 195 2.103E-01 +/- 1.81E-01 3.451E-01 +/- 0.00E+00 0.000E+00 +/- 0.00E+00 $A = 169 \ 0.000E+00 \ +/- \ 0.00E+00 \ 0.000E+00 \ +/- \ 0.00E+00 \ 2.814E+00 \ +/- \ 0.00E+00$ S = 28 4.145E-01 +/- 4.36E-01 7.161E-01 +/- 6.47E-01 9.505E-01 +/- 8.63E-01 7. = 74 7 = 75 7 = 73 З 7 = 2. $A = 4 \ 0.000E+00 \ +/- \ 0.00E+00 \ 2.418E+01 \ +/- \ 9.69E+00 \ 0.000E+00 \ +/- \ 0.00E+00$

 A = 3
 0.000E+00
 +/-0.00E+00
 4.207E+01
 +/-2.47E+01
 2.72E+01
 +/-1.96E+01

 A = 2
 0.000E+00
 +/-0.00E+00
 0.000E+00
 +/-0.00E+00
 2.884E+01
 +/-2.93E+01

 A = 1
 0.000E+00
 +/-0.00E+00
 0.000E+00
 +/-0.00E+00
 4.412E+01
 +/-5.92E+01

 S = 4 0.000E+00 +/- 0.00E+00 2.648E+01 +/- 1.40E+01 3.868E+01 +/- 5.17E+01 7. = 0 A = 1 1.078E+01 +/- 2.84E+01S = 1 1.078E+01 +/- 2.84E+01 End of nuclide average kinetic energies. Mass yield [mb] and the mean and variance of the emission angle [deg.] of residual nuclei: A = 196 6.807E-01 +/- 9.72E-02 9.892E+01 +/- 4.05E+01 A = 195 4.487E+00 +/- 2.50E-01 9.217E+01 +/- 4.52E+01 - - - - - - - -The mean and variance of the z velocity [v/c] of residual nuclei, A = 196 -9.863E-05 +/- 8.57E-04 4.848E-01 +/- 2.06E-01 A = 195 -3.324E-05 +/- 8.47E-04 9.341E-01 +/- 1.47E-01 A = 1 1.708E-02 +/- 1.04E-01 1.179E+00 +/- 1.05E-02 S = 140 1.571E-02 +/- 9.91E-02 1.222E+00 +/- 1.01E-02 Charge vield [mb] and the mean and variance of the emission angle [deg.] Z = 81 1.389E-02 +/- 1.39E-02 3.843E+01 +/- 0.00E+00 Z = 80 3.779E+00 +/- 2.29E-01 4.682E+01 +/- 3.04E+01 Z = 0 1.273E+03 +/- 4.21E+00 8.689E+01 +/- 3.98E+01 S = 61 1.642E+03 +/- 4.78E+00 8.516E+01 +/- 4.02E+01 The mean and variance of the z velocity [v/c] of residual nuclei, and the forward/backward ratio: Z = 81 1.283E-03 +/- 0.00E+00 1.000E+00 +/- 0.00E+00 Z = 80 1.127E-03 +/- 8.82E-04 1.471E+01 +/- 7.72E+00 Z = 0 1.170E-02 +/- 9.07E-02 1.910E+00 +/- 5.65E-03

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S = 61 1.571E-02 +/- 9.91E-02 1.222E+00 +/- 1.01E-02

Mass distributions of nuclei: At start of evap, which: just prior to after cascade after preeq evap. only fission A = 197 2.070E-01 8.720E-02 8.710E-02 1.000E-04 0.000E+00 A = 196 2.637E-01 2.248E-01 2.236E-01 1.200E-03 4.000E-04 ----. - - - - - - -- - - - -- - - - - -- - - - - 0.000E+00 0.000E+00 0.000E+00 0.000E+00 1.947E+02 1.935E+02 1.935E+02 1.912E+02 2.205E+00 3.212E+00 3.187E+00 3.606E+00 1.000E+00 1.000E+00 9.799E-01 2.010E-02 A = 165 1.000E-04 <A> = 1.868E+02 St Dv A = 6.608E+00 norm = 2.010E-02 Charge distributions of nuclei: at start of evap, which: just prior to after cascade after preeq evap. only fission fission Z = 81 1.100E-03 4.000E-04 3.000E-04 1.000E-04 1.000E-04 Z = 80 5.090E-02 3.660E-02 3.490E-02 1.700E-03 1.400E-03 -----1.000E-04 Z = 68 0.000E+00 0.000E+00 0.000E+00 0.000E+00 <Z> = 7.858E+01 7.809E+01 7.809E+01 7.790E+01 Dv Z = 7.861E-01 1.235E+00 1.227E+00 1.558E+00 7.696E+01 St Dv Z = 7.646E+01 1.000E+00 1.000E+00 9.799E-01 2.010E-02 norm = 2.010E-02 Excitation energy distributions [1/MeV] of nuclei: at start of evap, which: just prior to E*(MeV) after cascade after preeq evap. only fission fission 0.000E+00 0.000E+00 0.000E+00

0 10.	3.4006-04	2.3/0E-03	2.3/06-03	0.0005+00	0.0005+00	
10 20.	2.210E-03	6.260E-03	6.260E-03	0.000E+00	0.000E+00	
780 790.	1.000E-05	0.000E+00	0.000E+00	0.000E+00	0.000E+00	
< <u>E</u> *> =	1.260E+02	9.288E+01	8.946E+01	2.593E+02	2.039E+02	
St dev E* =	1.045E+02	8.738E+01	8.321E+01	1.186E+02	9.656E+01	
norm =	1.000E+00	1.000E+00	9.799E-01	2.010E-02	2.010E-02	

Linear momentum distributions [1/MeV/c] of nuclei:

			at start of ev	vap, which:	just prior to
P(MeV/c)	after cascade	after preeq	evap. only	fission	fission
0 10.	1.000E-05	0.000E+00	0.000E+00	0.000E+00	0.000E+00
10 20.	4.000E-05	5.000E-05	5.000E-05	0.000E+00	0.000E+00
20702080.	0.000E+00	0.000E+00	0.000E+00	0.000E+00	1.000E-05
<p> =</p>	2.999E+02	3.357E+02	3.318E+02	5.253E+02	5.774E+02
St dev P =	2.208E+02	2.254E+02	2.229E+02	2.644E+02	2.924E+02
norm =	1.000E+00	1.000E+00	9.799E-01	2.010E-02	2.010E-02

Angular momentum distributions [1/hbar] of nuclei:

			at start of	evap, which:	just prior to
L	after cascade a	after preeq	evap. only	fission	fission
0 1.	2.420E-02	1.340E-02	1.340E-02	0.000E+00	0.000E+00
1 2.	1.379E-01	8.070E-02	8.050E-02	2.000E-04	2.000E-04
61 62.	0.000E+00	1.000E-04	1.000E-04	0.000E+00	0.000E+00
<l> =</l>	5.997E+00	8.098E+00	8.043E+00	1.082E+01	1.082E+01
St dv L =	4.626E+00	6.045E+00	6.008E+00	7.133E+00	7.133E+00
norm =	1.000E+00	1.000E+00	9.799E-01	2.010E-02	2.010E-02

Neutron-multiplicity probability:

	Nn	Total	Cascade	Preequil.	Evap. res.	Pre-fiss.	Post-fiss.
	0	2.500E-03	2.878E-01	5.879E-01	1.340E-02	5.200E-03	0.000E+00
	1	1.400E-02	3.172E-01	2.960E-01	4.520E-02	3.800E-03	0.000E+00
	37	1.000E-04	0.000E+00	0.000E+00	0.000E+00	0.000E+00	0.000E+00
	<n> =</n>	9.164E+00	1.609E+00	5.664E-01	6.796E+00	2.751E+00	1.366E+01
St	dv n =	5.816E+00	1.704E+00	8.036E-01	4.392E+00	2.830E+00	4.707E+00
	norm =	= 1.000E+00	1.000E+00	1.000E+00	9.799E-01	2.010E-02	2.010E-02

Elapsed cpu time = 0. min and 23.331 sec.





Figure 4: Experimental proton spectra from 500 MeV p + Ni [132] compared with CEM03.03 results obtained using the input shown in Example 1 of Appendix 1 (the corresponding output is shown in Example 1 of Appendix 2). In contrast to the input file shown in Appendix 1, the results shown in this figure are for one million simulated inelastic events (limc=1000000). The option considering 66 types of possible evaporated particles (nevtype=66) requires 18 min 34 sec of computing time on an UltraSPARC 1.3 GHz Sunstation, while the nevtype=6 option requires only 6 min 51 sec, providing almost the same results.

Example 2



Figure 5: Experimental π^0 spectra from 500 MeV π^- + Cu [133, 27] compared with CEM03.03 results obtained using the input shown in Example 2 of Appendix 1 (the corresponding output is shown in Example 2 of Appendix 2). In contrast to the input file shown in Appendix 1, the results shown in this figure are for one million simulated inelastic events (**limc=1000000**). As pions are produced by CEM03.03 only at the INC stage of reactions, calculated pion spectra do not depend on the value of **nevtype**; this calculation was done using only the **nevtype=6** option in the input and it took 8 min 11 sec on an UltraSPARC 1.3 GHz Sunstation.

Example 3



Figure 6: Experimental π^+ spectra from 562.5 MeV n + Cu [134] compared with CEM03.03 results obtained using the input shown in Example 3 of Appendix 1 (the corresponding output is shown in Example 3 of Appendix 2). In contrast to the input file shown in Appendix 1, the results shown in this figure are for one million simulated inelastic events (**limc=1000000**). This calculation was done using only the **nevtype=6** option in the input, for the same reason as discussed for Example 2, and it took 7 min 33 sec on an UltraSPARC 1.3 GHz Sunstation.

Example 4



Figure 7: Experimental neutron spectra from 1.5 GeV π^+ + Fe [135] compared with CEM03.03 results obtained using the input shown in Example 4 of Appendix 1 (the corresponding output is shown in Example 4 of Appendix 2). In contrast to the input file shown in Appendix 1, the results shown in this figure are for one million simulated inelastic events (limc=1000000). The nevtype=66 option requires 38 min 17 sec of computing time on an UltraSPARC 1.3 GHz Sunstation, while the nevtype=6 option requires only 13 min 9 sec, providing almost the same results.



Figure 8: Experimental neutron-induced fission cross section of 197 Au [136]–[138] compared with CEM03.03 results obtained using the input shown in Example 5 of Appendix 1 (the corresponding output is shown in Example 5 of Appendix 2). The results shown in this figure are the **Direct Monte Carlo Simulation Method** fission cross sections from the CEM03.03 output. These calculations were done at neutron energies from 30 MeV (t0mev=30.0) to 500 MeV (t0max=500.5) with a step of 10 MeV (dt0=10.0) and, in contrast to the input file shown in Appendix 1, use 100000 simulated inelastic events for each energy point (limc=100000). The nevtype=66 option requires 1 hr 34 min 11 sec of computing time on an UltraSPARC 1.3 GHz Sunstation, while the nevtype=6 option requires only 43 min 29 sec, providing almost the same results.



Figure 9: Experimental angle-integrated energy spectra (upper left plot), energy-integrated angular distributions (upper right plot), and double-differential spectra of nucleons and complex particles [139] compared with CEM03.03 results obtained using the input shown in Example 6 of Appendix 1 (the corresponding output is shown in Example 6 of Appendix 2). In contrast to the input file shown in Appendix 1, the results shown in this figure are for ten million simulated inelastic events (limc=10000000). The nevtype=66 option requires 1 hr 51 min 8 sec of computing time on an UltraSPARC 1.3 GHz Sunstation, while the nevtype=6 option requires only 1 hr 26 min 20 sec, providing practically the same results, indistinguishable within the scale of this figure.

Figure 10: The measured [140] mass and charge distributions of the product yields from the reaction 800 MeV/A ¹⁹⁷Au+p and of the mean kinetic energy of these products, and the mass distributions of the cross sections for the production of thirteen elements with the charge Z from 20 to 80 (open symbols) compared with CEM03.03 results obtained using the input shown in Example 7 of Appendix 1 (the corresponding output is shown in Example 7 of Appendix 2). In contrast to the input file from Appendix 1, the results shown in this figure are for ten million simulated inelastic events (**limc=10000000**). The **nevtype=66** option requires 9 hr 7 min 23 sec of computing time on an UltraSPARC 1.3 GHz Sunstation, while the **nevtype=6** option requires only 2 hr 14 min 45 sec, providing almost the same results for the spallation and fission products. The fragment (2 < Z < 13, 6 < A < 29) results are very different, therefore we need to use the option **nevtype=66** when we are interested in fragment production.

Figure 11: Experimental mass distributions of the yields of eight isotopes from Na to Mn [141] and of all light fragments from Li to O [142] from the reaction 1 GeV/A ⁵⁶Fe+p and the mass number- and charge-distributions of the product yield compared with CEM03.03 results obtained using the input shown in Example 8 of Appendix 1 (the corresponding output is shown in Example 8 of Appendix 2). Predictions of CEM03.03 for the mean kinetic energy, mean production angle Θ , mean parallel velocity v_z , and of the F/B ratio of the forward product cross sections to the backward ones of all isotopes in the laboratory system are given as well. In contrast to the input file from Appendix 1, the results shown in this figure are for ten million simulated inelastic events (**limc=10000000**). The **nevtype=66** option requires 4 hr 57 min 52 sec of computing time on an UltraSPARC 1.3 GHz Sunstation, while the **nevtype=6** option requires only 1 hr 38 min 23 sec, providing almost the same results for the spallation products. The yields of light fragments, especially of Li and Be, differ by several orders of magnitude, therefore we need to use the option **nevtype** > **6** when we are interested in light-fragment production.

Example 9

Figure 12: Proton spectra at 45°, 90°, and 135° from the reaction 300 MeV γ + Cu. Symbols are experimental data from [143] and histograms are CEM03.03 results obtained using the input shown in Example 9 of Appendix 1 (the corresponding output is shown in Example 9 of Appendix 2). In contrast to the input file from Appendix 1, the results shown in this figure are for one million simulated inelastic events (**limc=1000000**). The **nevtype=66** option requires 35 min 18 sec of computing time on an UltraSPARC 1.3 GHz Sunstation, while the **nevtype=6** option requires only 26 min 23 sec, providing almost the same results.

Figure 13: Results from CEM03.03 for the product yield of all isotopes and of their mean laboratory kinetic energy as functions of the product mass-number A and charge Z from interactions of bremsstrahlung gammas with a maximum energy of 1 GeV with Au. In contrast to the input file from Appendix 1, whose output we show in Example 10 of Appendix 2, we use ten million simulated inelastic events (limc=10000000). The nevtype=66 option requires 8 hr 58 min 58 sec of computing time on an UltraSPARC 1.3 GHz Sunstation, while the nevtype=6 option requires only 6 hr 17 min 31 sec, providing almost the same results for the spallation and fission products but underestimating the yields of light fragments by more than two orders of magnitude. The experimental cross sections shown for comparison by circles are from the review [144] and their tabulated values were kindly sent us by by Dr. Hiroshi Matsumura.