

IAEA Photonuclear Data Library 2019

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Photo-induced reaction cross section data are of importance for a variety of current or emerging applications, such as radiation shielding design and radiation transport analyses, calculations of absorbed dose in the human body during radiotherapy, physics and technology of fission reactors (influence of photo-reactions on neutron balance) and fusion reactors (plasma diagnostics and shielding), activation analyses, safeguards and inspection technologies, nuclear waste transmutation, medical isotope production and astrophysical applications.

To address these data needs the IAEA Photonuclear Data library was produced in 1999, containing evaluated photo-induced cross sections and neutron spectra for 164 nuclides which were deemed relevant for the applications.

Since the release of the IAEA Photonuclear Data Library however, new experimental data as well as new methods to assess the reliability of experimental cross sections have become available. Theoretical models and input parameters used to evaluate photo-induced reactions have improved significantly over the years. In addition, new measurements of partial photoneutron cross sections using mono-energetic photon beams and advanced neutron detection systems have been performed allowing for the validation of the evaluations and assessments of the experimental data. Furthermore, technological advances have led to the construction of new and more powerful gamma-beam facilities, therefore new data needs are emerging.

We report our coordinated efforts to address these data needs and present the results of the new up-to-date evaluations included in the new updated IAEA Photonuclear Data Library consisting of 219 nuclides. The new library includes 188 new evaluations produced by the CRP evaluators, and one evaluation taken from the JENDL/PD-2016 library, while 20 evaluations were retained from the previous 1999 IAEA Photonuclear Data Library. In most of the cases, the photon energy goes up to 200 MeV. A total of 55 nuclides are new in this library reflecting the progress in measurements but also the developing data needs. In this paper we discuss the new assessment method and make recommendations to the user community in cases where the experimental data are discrepant

and the assessments disagree. In addition, in the absence of experimental data, we present model predictions for photo-induced reaction cross section on nuclides of potential interest to medical radioisotope production.

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I. INTRODUCTION

Photonuclear data describing interactions of photons with atomic nuclei are important for a range of applications such as (i) radiation shielding and radiation transport analyses, (ii) calculation of absorbed doses in the human body during radiotherapy, (iii) activation analyses, (iv) safeguards and inspection technologies, (v) nuclear waste transmutation, (vi) fission and fusion reactor technologies, and (vii) astrophysical nucleosynthesis.

Photons can be produced as bremsstrahlung radiation by electron accelerators which are commonly used in hospitals, industries and laboratories. Significant technological advances led to the production of quasi-monochromatic beams using the positron annihilation in flight technique at the national facilities of the Lawrence Livermore National Laboratory (LLNL, USA) and Centre d'Études Nucléaires de Saclay (France) in the 1960's. As a result, a large number of measurements of photonuclear cross sections was performed at these two facilities for almost three decades (1962 – 1987). This pioneering

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work was captured by B.L. Berman in his comprehensive compilation of photonuclear cross sections “Atlas of Photoneutron Cross-sections obtained by Monoenergetic Photons” published in 1975 [1], and later in the follow-up produced in 1988 [2]. Note that because these institutes changed their names several times, we herewith abbreviate them simply to Livermore and Saclay.

Despite those experimental efforts, there was still a need for evaluated photonuclear data since (a) it is not possible to produce a complete photonuclear data files based on measured cross sections alone, (b) often the experimental data suffer from systematic discrepancies which are not easy to resolve, and (c) there is a lack of data in a number of cases. These deficiencies in the experimental data can be addressed and often resolved by performing an evaluation which consists of three systematic operations: compilation of experimental data, critical assessment of the measurement techniques used and theoretical calculations based on reliable nuclear models. As a result of the evaluation, one can obtain production cross sections as well as energy and angular distributions of the emitted particle for a wide range of incident and outgoing energies, which is useful for the applications.

To address the growing needs for photonuclear data, the IAEA held a Coordinated Research Project (CRP) under the title Compilation and Evaluation of Photonuclear Data for Applications between 1996 and 1999. This CRP produced the IAEA Photonuclear Data Library which is described in the Handbook on Photonuclear Data for Applications [3] prepared by M.B. Chadwick (chair), A.I. Blokhin, T. Fukahori, Y.-O. Lee, M.N. Martins, V.V. Varlamov, B. Yu, Y. Han, S.F. Mughabghab, J. Zhang, and P. Obložinský, and is available at the IAEA web-site (<http://www-nds.iaea.org/photonuclear>). The library includes photon absorption data, total and partial photoneutron reaction cross sections and neutron spectra for 164 isotopes, primarily for structural, shielding, biological and fissionable materials. In this paper we refer this previous library to IAEA 1999.

The list of 164 isotopes included in the 1999 Photonuclear Data Library [3] can be broken down in four categories:

- Structural, shielding and bremsstrahlung target materials: Be, Al, Si, Ti, V, Cr, Fe, Co, Ni, Cu, Zn, Zr, Mo, Sn, Ta, W, and Pb;
- Biological materials: C, N, O, Na, S, P, Cl, and Ca;
- Fissionable materials: Th, U, Np, and Pu; and
- Other materials: H, K, Ge, Sr, Nb, Pd, Ag, Cd, Sb, Te, I, Cs, Sm, and Tb.

Of the above four categories, the most important for the applications are the 40 major isotopes of the 29 elements in the first three groups.

Although this database has been extremely useful to a broad user community, it has become evident that it needs to be revised since

- Some of the experimental data measured with quasi-monochromatic photon beams are unreliable and discrepant;
- Data have been measured for 37 isotopes that have not been evaluated;
- Improved methods to resolve experimental discrepancies are available; and
- New data measured with modern techniques have been published in recent years.

New experimental facilities, such as HIγS at the Triangle University National Laboratory (USA) and the Laser Compton Scattering Facility (NewSUBARU) at University of Hyogo (Japan), offering highly monoenergetic photon beams in combination with advancements in neutron detector technologies have opened the field to new possibilities: new measurements of photoneutron cross sections with better accuracy that are expected to help resolve the long-standing discrepancies observed between the data measured using quasi-monoenergetic beams.

Furthermore, the needs for evaluated photonuclear data are growing. In the field of medical isotope production, photonuclear reactions are being explored for the production of medical radionuclides. With the advent of new facilities producing brilliant photon beams with extremely high activity, photo-production of some important radionuclides could become a competitive alternative to the traditional methods using neutrons produced at highly-enriched Uranium reactors or charged-particle beams [4–7]. A list of radionuclides proposed for potential diagnostic and therapeutic applications in nuclear medicine which have also been identified as suitable candidates for production via photonuclear reactions is given below:

- $^{166,170}\text{Er}$ for the production of $^{165,169}\text{Er}$,
- ^{187}Re for the production of ^{186}Re ,
- ^{226}Ra for the production of ^{225}Ra which further decays to ^{225}Ac ,
- ^{98}Ru for the production of ^{97}Ru ,
- ^{194}Pt for the production of $^{193\text{m}}\text{Pt}$,
- ^{132}Xe for the production of ^{131}I ,
- ^{162}Dy for the production of ^{161}Tb , and
- ^{178}Hf for the production of ^{177}Lu .

Other important candidates such as ^{100}Mo (for the production of ^{99}Mo via (γ, n)), ^{48}Ca , ^{52}Cr , ^{65}Cu , ^{48}Ti and ^{46}Ti are already included in the existing photonuclear data library.

In view of the above developments, it was timely to update the existing IAEA Photonuclear Data Library to reflect the progress in the field and the emerging demands

for photonuclear data. A coordinated research project (CRP) was endorsed by the International Nuclear Data Committee at the 2014 meeting in Vienna and was initiated by the IAEA with the title “Updating the Photonuclear Data Library.” All the 164 isotopes in the existing library were revisited and evaluated by considering new data, the results of experimental-based evaluations as well as the new Giant Dipole Resonance (GDR) parameters from the recently updated Atlas of GDR parameters [8] and improved reaction models. In addition to these 164 isotopes, evaluations were performed for 37 isotopes for which experimental data are available, as well as for the 9 isotopes identified in the above-mentioned list as relevant for medical applications. In some cases where no experimental information is available, a model prediction is given as the evaluation. In total, 219 isotopes were evaluated. All available experimental data for photo-absorption cross sections, photoneutron production cross sections and yields, as well as for partial photoneutron cross sections and photo-charged-particle cross sections that were available in the EXFOR database [9] up to the cut-off date of April 2019 were considered. In addition, all the new measurements obtained with the new direct neutron-multiplicity sorting technique [10] that was fully developed and implemented for the CRP, were also taken into account in the evaluations. The evaluations were extended to energies of 200 MeV for use in accelerator-driven transmutation technologies, to complement the neutron and proton high-energy libraries that are being developed for radiation transport simulation codes. Although the highest energy of 200 MeV exceeds the pion production threshold, its small production cross section can be ignored.

The CRP has a second branch under the title “Generating a Reference Database for Photon Strength Functions” which has led to a new database of experimental and calculated photon strength functions [11]. The evaluations performed for the updated photonuclear data library have formed the basis for the recommended photonuclear photon strength functions in Ref. [11]. In this paper we adopt the same definition of the photon strength functions as in Ref. [11].

In this report, we present the results of both experimental and evaluation efforts carried out within the CRP to update the IAEA Photonuclear Data Library. The measurements of reliable photoneutron production cross sections using the new neutron multiplicity sorting technique as well as an overview of all available experimental data can be found in Sec. III. In Sec. IV, we present the nuclear reaction models that are used to describe photonuclear reactions as well as the different codes employed in this coordinated effort. In Sec. V, we present the different evaluations and highlight some comparisons with experimental data. The contents of the new IAEA photonuclear data library are described in Sec. VI. Our conclusions are given in Sec. VII. The new Atlas of GDR parameters is provided in the Appendix.

II. DEFINITIONS

Experimental photonuclear reaction data are usually obtained by directly counting the number of emitted particles or by measuring the residual nucleus activity. For energies above the multi-particle emission threshold, more than one combination of emitted light-particles can accompany the same number of produced neutrons or can lead to the same residual nucleus, respectively. Often, authors do not mention clearly what it is they measure which can lead to confusion when comparing experimental data with evaluations. For example, an experimental data set that is reported as the (γ, n) cross section could have charged particles emitted concurrently with one-neutron emission, *i.e.* $(\gamma, 1n) + (\gamma, 1np) + (\gamma, n2p) + \dots$ depending on the energy. This can lead to significant differences, especially for light nuclei, where photo-charged-particle emission is quite strong. Here we define several cross sections relevant to the photonuclear data library.

When the charged-particle emission is negligible, the measured one-neutron emission cross section is identical to the cross section for the production of $(Z, A - 1)$ nucleus, and the two-neutron emission is equal to the production of $(Z, A - 2)$, and so on. However, when charged-particle emission is non-negligible, then the measured one-neutron emission cross section σ_{1nX} , alternatively $\sigma(\gamma, 1nX)$, reads

$$\begin{aligned}\sigma_{1nX} &\equiv \sigma(\gamma, 1nX) \\ &= \sigma(\gamma, 1n) + \sigma(\gamma, np) + \sigma(\gamma, n\alpha) + \dots, \quad (1)\end{aligned}$$

and *ditto* for $\sigma(\gamma, 2nX)$, $\sigma(\gamma, 3nX)$, *etc.* Thus, $\sigma(\gamma, inX)$ is understood to be the inclusive *i*-neutron emission cross section, where X stands for anything except for *i*-neutrons. These cross sections are not given explicitly in the evaluated photonuclear data library, instead one has to reconstruct σ_{1nX} by summing each term in Eq. (1).

The photoneutron production cross section is defined in two ways, either involving the neutron multiplicity or not. We denote the photoneutron yield (production) cross section σ_{xn} , alternatively $\sigma(\gamma, xn)$,

$$\begin{aligned}\sigma_{xn} &\equiv \sigma(\gamma, xn) \\ &= \sigma(\gamma, 1n) + \sigma(\gamma, np) + \sigma(\gamma, n\alpha) + \dots \\ &+ 2\sigma(\gamma, 2n) + 2\sigma(\gamma, 2np) + 2\sigma(\gamma, 2n\alpha) + \dots \\ &+ 3\sigma(\gamma, 3n) + 3\sigma(\gamma, 3np) + 3\sigma(\gamma, 3n\alpha) + \dots, \\ &= \sum_i i\sigma_{inX}, \quad (2)\end{aligned}$$

and the total photoneutron cross section σ_{Sn} , alternatively $\sigma(\gamma, Sn)$,

$$\begin{aligned}\sigma_{Sn} &\equiv \sigma(\gamma, Sn) \\ &= \sigma(\gamma, 1n) + \sigma(\gamma, np) + \sigma(\gamma, n\alpha) + \dots \\ &+ \sigma(\gamma, 2n) + \sigma(\gamma, 2np) + \sigma(\gamma, 2n\alpha) + \dots \\ &+ \sigma(\gamma, 3n) + \sigma(\gamma, 3np) + \sigma(\gamma, 3n\alpha) + \dots \\ &= \sum_i \sigma_{inX}. \quad (3)\end{aligned}$$

When the photo-charged-particle reaction cross sections — $\sigma(\gamma, p)$, $\sigma(\gamma, \alpha)$, *etc.* — are negligible, and no photo-fission occurs, then σ_{Sn} is the same as the photo-absorption cross section σ_{abs} . Otherwise, the sum of this cross section, σ_{Sn} , with the photo-charged-particle cross section gives the total photo-absorption cross section σ_{abs} , alternatively $\sigma(\gamma, abs)$,

$$\begin{aligned}\sigma_{abs} &\equiv \sigma(\gamma, abs) \\ &= \sigma_{Sn} + \sigma(\gamma, p) + \sigma(\gamma, 2p) + \dots + \\ &\quad + \sigma(\gamma, d) + \sigma(\gamma, dp) + \dots + \sigma(\gamma, \alpha) + \dots\end{aligned}\quad (4)$$

Usually σ_{xn} is explicitly given in the photonuclear data library as the neutron multiplicity, while σ_{Sn} is implicit. When the target photo-fissions, Eqs. (2) and (3) have an extra term

$$\begin{aligned}\sigma_{xn} &= \sigma(\gamma, n) + 2\sigma(\gamma, 2n) + \dots + \bar{\nu}\sigma(\gamma, f), \quad (5) \\ \sigma_{Sn} &= \sigma(\gamma, n) + \sigma(\gamma, 2n) + \dots + \sigma(\gamma, f), \quad (6)\end{aligned}$$

where $\bar{\nu}$ is the average number of neutrons per fission.

Varlamov *et al.* [12] proposed an F_i value,

$$F_i = \frac{\sigma(\gamma, inX)}{\sigma_{xn}}, \quad (7)$$

to facilitate the assessment and evaluation of the experimental $\sigma(\gamma, inX)$ cross sections. F_i has a maximum value for each i-neutron reaction channel,

- $F_1 < 1$ for $i = 1$,
- $F_2 < 1/2$ for $i = 2$, and
- $F_3 < 1/3$ for $i = 3$.

When F_i exceeds these limits, then there are issues in the measurement of $\sigma(\gamma, inX)$ data by the same group. Since the above-mentioned limits are automatically satisfied in the theoretical calculations, it is essential to apply a model-based approach in the evaluation of the available photonuclear experimental data to avoid such inconsistencies.

III. EXPERIMENTAL METHODS AND AVAILABLE EXPERIMENTAL DATA

The experimental photonuclear reaction data have been obtained in various types of measurements, with bremsstrahlung and quasi-monoenergetic photons from positron annihilation in flight and more recently from laser Compton scattering (LCS). While the photoneutron yield cross sections are obtained by counting the total number of neutrons emitted, the determination of partial photoneutron cross sections requires neutron-multiplicity sorting. Below we provide a brief description of the experimental procedures most frequently used in the past [3] and most recently developed.

A. Experiments

To determine the photonuclear reaction cross section, one needs to measure on an absolute scale both the flux of the incident photons and the number of reaction products. The incident photon flux can be determined directly with an ionization chamber, scintillator or solid-state detector or indirectly by normalizing to known reaction cross sections (for bremsstrahlung beams). The number of reaction products is determined either by counting the emitted particles or by measuring the activity of radioactive residual nuclei. We summarize the characteristics of various γ -ray sources and photonuclear cross section measurements below.

1. Bremsstrahlung

Bremsstrahlung beams were used in the first measurements of photonuclear cross sections. The continuous bremsstrahlung spectra were produced by striking a radiator target with an electron beam from an accelerator (initially betatrons and synchrotrons, and later, linear accelerators). Several laboratories around the world (mostly in Russia, Canada, Australia, and Germany) developed this type of photon beams.

As the spectrum of photon energies is continuous, only the yield of the reaction can be measured,

$$Y(E_0) = N_R \int_{E_{th}}^{E_0} \frac{\sigma(E_\gamma)}{E_\gamma} W(E_0, E_\gamma) dE_\gamma, \quad (8)$$

where E_γ is the photon energy, $\sigma(E_\gamma)$ is the reaction cross section, $W(E_0, E_\gamma)$ is the bremsstrahlung energy spectrum, E_0 is the end-point energy of the bremsstrahlung spectrum which is equal to the electron beam energy, E_{th} is the threshold energy, and N_R is the normalization coefficient. Changing E_0 in small steps allows one to measure a yield curve and then by applying an “unfolding” procedure to obtain the photonuclear reaction cross section.

Several unfolding methods for spectrum-averaged cross sections have been developed, among which the most widely used ones are:

- The Photon Difference Method [13] in which the difference of two bremsstrahlung spectra near the slightly differing end-point energies is interpreted as an almost quasi-monoenergetic spectrum; a modified version uses the linear combination of three bremsstrahlung spectra [14] which improves the shape of the resulting spectrum;
- The Penfold-Leiss method [15] in which the integral Eq. (8) is replaced by a set of linear equations for finite analysis bins; some modifications of this method [16] vary the analysis bin depending on the yield accuracy;
- The regularization method [17] in which the mean square difference between the reaction cross section

and the model value is minimized assuming different approaches for smoothing the cross section; modifications of this method are based on various regularizers [18–20].

The typical apparatus functions (effective photon spectra) for these methods have different line shapes localized in photon energy though not close to a Gaussian line shape, allowing experimentalists to obtain the information on the reaction cross section at energy E_γ with energy resolution dependent on the data processing.

The advantage of bremsstrahlung measurements is the large photon beam intensity, which allows one to obtain reasonable counting statistics even for relatively small reaction cross sections. However, there are several disadvantages of using such a technique. First, one needs to know the bremsstrahlung spectrum sufficiently well for all electron energies. Second, measuring a reaction yield curve in small energy steps requires a stable accelerator and large counting statistics. Third, the process of subtracting the yield curves in the unfolding procedure may introduce correlations between the experimental data points that can lead to unphysical fluctuations in the unfolded cross sections.

2. Positron Annihilation in Flight

While the unfolding of the experimental reaction yield resulting from bremsstrahlung measurements may involve a “mathematical” method of obtaining the “quasi-monoenergetic” photons, positron annihilation in flight offered an “apparatus method” of producing them with variable energies. The method [21] was realized in several laboratories around the world, mostly in LLNL (USA) and Saclay (France) [22]. An intense beam of high-energy electrons from a linear accelerator hits a thick high- Z converter and produces bremsstrahlung which undergoes electron-positron pair production in the converter. The fast positron beam then impinges on a thin, low- Z target and produces annihilation photons accompanied by positron bremsstrahlung. The Livermore and Saclay facilities were nearly the same except that while in the Livermore system positrons from the converter were re-accelerated before undergoing annihilation, in the Saclay system the positrons were simply separated without re-acceleration.

To separate the bremsstrahlung component from the annihilation photons, the measurement proceeded in the following three steps:

1. Similar to the bremsstrahlung measurement, the reaction yield, $Y^+(E_0)$, is obtained with the γ -ray spectrum that consists of the positron annihilation and bremsstrahlung components. See Ref. [23] for the 15.6 MeV positrons on the 0.0060 in. thick LiH target. The peak energy of the annihilation γ -rays is given by $E_0 + 3mc^2/2$ with the kinetic energy of positrons E_0 and the electron rest mass energy mc^2 ;

2. The reaction yield $Y^-(E_0)$ is given by bremsstrahlung of electrons with the energy E_0 ; and
3. The reaction cross section is deduced as the difference between the positron and electron reaction yields assuming that the bremsstrahlung spectra from positrons and electrons are identical to the first order approximation; the subtraction is supposedly subject to the stability of the accelerator parameters.

Hence the photonuclear cross section reads

$$\sigma(E_\gamma) = Y^+(E_0) - Y^-(E_0) . \quad (9)$$

The effective photon spectrum for this method is ideally the positron-annihilation line shape. The disadvantage of this method, as compared to bremsstrahlung, is the low intensity of the photon beam attributed to the small cross sections of the electron-positron pair production and positron annihilation processes.

3. Bremsstrahlung Tagging

As an alternative to the complex task of unfolding the reaction cross section from the experimental reaction yield by solving the inverse integral equation of Eq. (8), the method of producing (quasi)monoenergetic photons by tagging was proposed [24]. It was implemented at the University of Illinois (USA) using the beam from a high duty-cycle electron linear accelerator. The idea was to produce photons by letting electrons impinge on a very thin radiator with an energy E_0 and then to tag the photons with scattered electrons with an energy E_{scatt} . Thus, photons are tagged with an energy E_γ ,

$$E_\gamma = E_0 - E_{\text{scatt}} . \quad (10)$$

The scattered electron is deflected by the magnetic spectrometer to an array of detectors placed in its focal plane. The energy spread (resolution) of the photon beam depends on the number of detectors mounted in focal plane. A thin-sliced part of the bremsstrahlung spectrum represents the typical effective photon spectrum from this method, which is close to a Gaussian line shape. partial reaction cross sections were obtained using the tagged bremsstrahlung photons.

The disadvantage of this method is the low-intensity photon beam extracted from the very thin bremsstrahlung-producing radiator that makes coincidence measurements difficult.

4. Laser Compton-scattering

Following the development of high duty-cycle electron accelerators and high-power lasers, new quasi-monoenergetic γ -ray beams have been produced in collisions of laser photons with relativistic electrons which

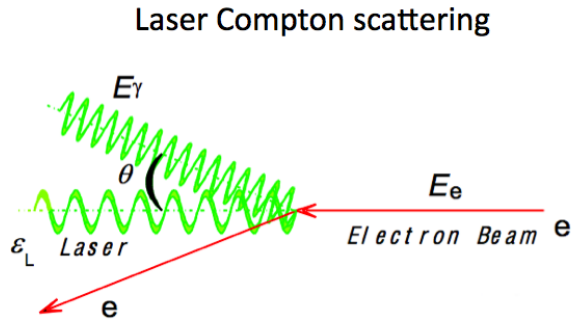


FIG. 1. (Color online) Laser Compton scattering of laser photons from relativistic electrons.

is referred to as laser Compton scattering (LCS) [25–28] as illustrated in Fig. 1. The LCS γ -ray beam-line of the NewSUBARU synchrotron radiation facility [28] is shown in Fig. 2. An electron beam at 1.0 GeV in nominal energy is injected from a linear accelerator to the storage ring and either decelerated to 0.5 GeV or accelerated to 1.5 GeV.

Pulse γ -ray beams are produced with Q-switch lasers; low-energy beams are produced typically at 20 kHz with the INAZUMA laser (1064 nm wavelength and 60 ns pulse width) for $(\gamma, 1n)$ cross section measurements below $(\gamma, 2n)$ threshold, while high-energy beams at 1 kHz with the Talon laser (532 nm, 40 ns) for partial cross section measurements above $(\gamma, 2n)$ threshold. The frequency and pulse width of the electron beam in the NewSUBARU storage ring are 500 MHz and 60 ps, respectively, so that the frequency and width of the γ -ray pulse are the same as those of the laser.

The electron beam energy has been calibrated with the accuracy of the order of 10^{-5} [29] by using low-energy LCS γ -ray beams produced with a grating-fixed CO₂ laser with the central wavelength of the P(20) master transition ($\lambda = 10.5915 \mu\text{m} \pm 3 \text{ \AA}$) with the bandwidth 1.3 $\text{\AA}$ in the full width at half maximum. The γ -ray beam energy is determined from the calibrated electron beam energy.

The flux of pulsed γ -rays is determined with the pile-up or Poisson-fitting method [30–32]. Large numbers of laser photons and electrons are involved in the collision with a small probability of the laser Compton scattering. As a result, the number of photons involved in a γ -ray pulse follows the Poisson distribution [30]. Figure 3 shows an experimental multi-photon (pile-up) spectrum for 34-MeV γ -rays measured with a $8'' \times 12''$ NaI(Tl) detector along with the best-fit Poisson distribution. The average number of photons per γ -pulse is determined with the intrinsic accuracy less than 0.1% [32]. The γ -ray flux is determined with the experimental formula [31, 32] for a

product ($m^{\text{exp}} \times N_{\gamma}^{\text{pulse}}$) of the average number,

$$m^{\text{exp}} = \frac{\langle N_m \rangle}{\langle N_s \rangle}, \quad (11)$$

and the number of γ -ray pulses

$$N_{\gamma}^{\text{pulse}} = \sum_i N_m(i), \quad (12)$$

where $N_m(i)$ and $N_s(i)$ are the number of photons at channel i in the multi- and single-photon spectra, respectively, and $\langle N \rangle$ stands for their average channel number [32].

The energy-profile of the LCS γ -rays is determined by best reproducing response functions of an energy-profile monitor with a Monte Carlo code GEANT4 that incorporates the kinematics of the laser Compton scattering, transportation of LCS γ -rays through the collimators (C1 and C2) to the energy-profile monitor mounted in the experimental hutch GACKO (GAMMA Collaboration hutch of KOnan university) in the γ -ray beam line of the NewSUBARU facility, and electromagnetic interactions of γ -rays inside the monitor detector. Figure 4 shows experimental γ -ray spectra for the 3 mm C1 and 2 mm C2 collimators measured with a $3.5'' \times 4.0''$ LaBr₃(Ce) detector, best-fit Monte Carlo simulations, and incident γ -ray spectra at 6, 10, 20, 30 and 39 MeV. As shown in Fig. 4, the energy-profile of the LCS γ -rays produced at the NewSUBARU facility is characterized by a sharp cut-off at the maximum energy corresponding to the head-on collision and a low-energy tail with an energy spread determined by the electron beam divergence and the collimator size. The energy spread is a few % in the full width at half maximum.

The synchrotron radiation accompanies the LCS γ -rays as background which is considerably lower in both intensity and energy than the positron bremsstrahlung which accompanied the positron-annihilation-in-flight γ -rays. The laser was turned on and off at 10 Hz for 80 ms and 20 ms, respectively, to measure the background neutrons of the synchrotron-radiation and cosmic-ray origin during the beam off.

B. Partial Reaction Measurement

One can determine the neutron yield cross section from the total number of neutrons detected using Eq. (2) virtually in all measurements with the different γ -ray sources summarized in Sec. III A. However, it is not straightforward at all to determine partial photoneutron cross sections, $(\gamma, 1n)$, $(\gamma, 2n)$, $(\gamma, 3n)$, *etc.* In this section, we describe how partial photoneutron cross sections are derived by using the various experimental techniques, and then focus our discussions on the long-standing discrepancy between the Livermore and Saclay data and the new direct neutron-multiplicity sorting technique, which constitutes a major topic of the present CRP.

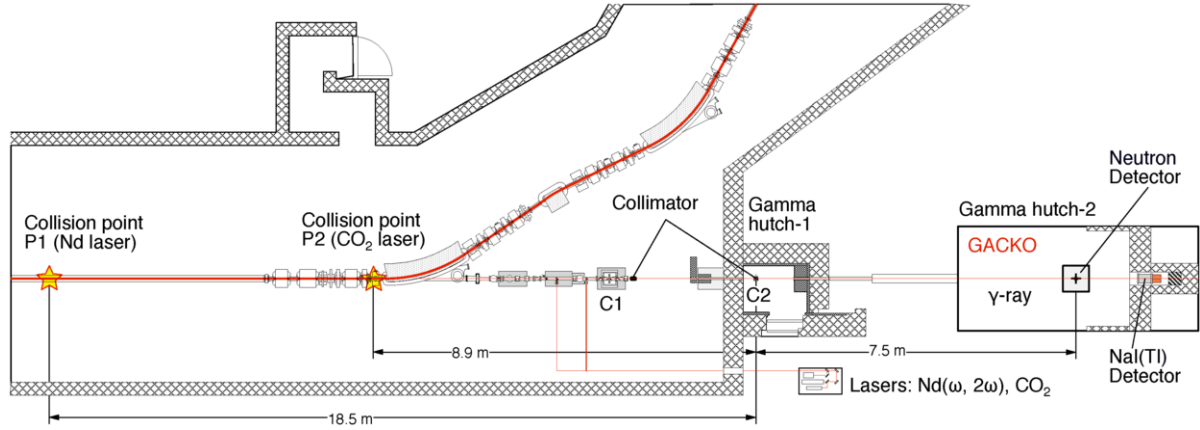
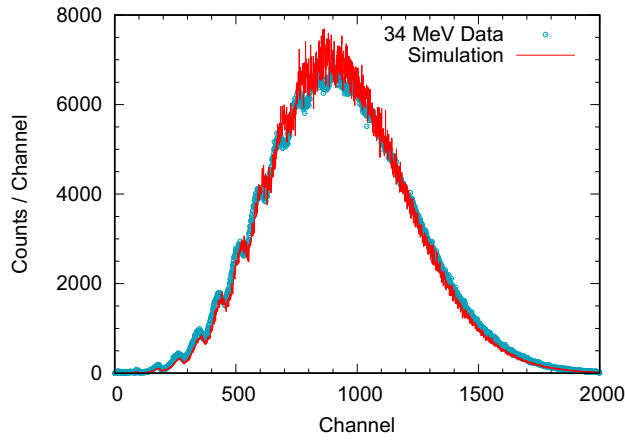
FIG. 2. (Color online) The γ -ray beam-line BL01 of the NewSUBARU synchrotron radiation facility.

FIG. 3. (Color online) Experimental multi-photon spectrum at 34 MeV in comparison with the best-fit Poisson distribution.

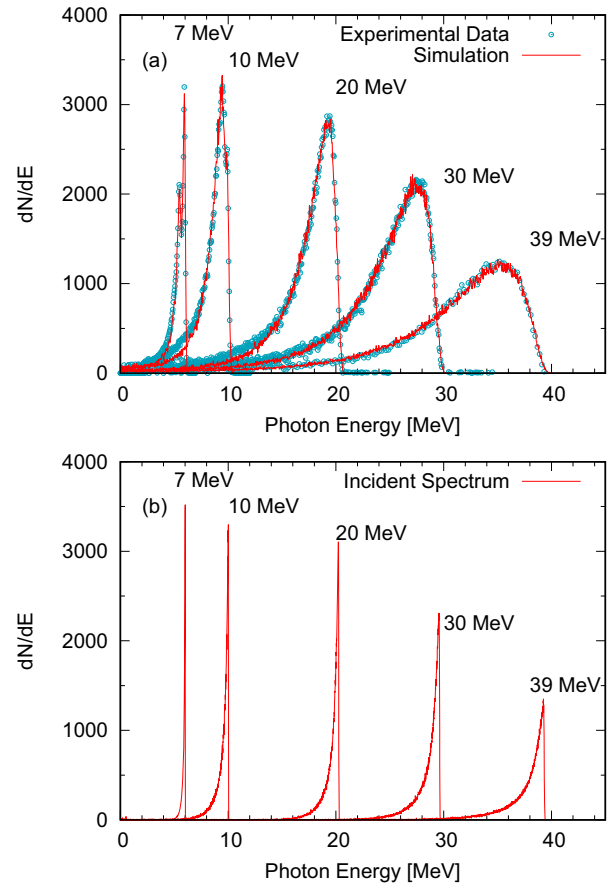
1. Bremsstrahlung

Since the effective photon spectrum is continuous, a special correction that is based on the statistical theory of nuclear reactions is applied to $\sigma(\gamma, in)$ to obtain the total photoneutron reaction cross section of Eq. (3). After the correction is applied, $(\gamma, 2n)$ and $(\gamma, 1n)$ cross sections below the $(\gamma, 3n)$ reaction threshold B_{3n} , can be obtained by subtraction as follows:

$$\sigma(\gamma, 2n) = \sigma_{xn} - \sigma_{Sn} , \quad (13)$$

$$\sigma(\gamma, 1n) = \sigma_{Sn} - \sigma(\gamma, 2n) , \quad (14)$$

$$\sigma(\gamma, 1n) = \sigma_{xn} - 2\sigma(\gamma, 2n) . \quad (15)$$

FIG. 4. (Color online) (a) Experimental and simulated response functions of a $3.5'' \times 4.0''$ LaBr₃(Ce) detector to LCS γ -ray beams, and (b) energy spectra of incident LCS γ -ray beams obtained by Monte Carlo simulations with the GEANT4 code. The energy spread is a few % in the full width at half maximum: 68, 247, 325, 578, and 965 keV for 6.0, 10, 20, 30, and 39 MeV LCS γ -ray beams, respectively.

2. Quasi-monoenergetic Annihilation Photons

Highly-efficient 4π neutron detectors for measuring multi-neutron coincidences and neutron multiplicity sorting techniques for separating partial reactions were developed in Livermore [22], Saclay, and other laboratories, to handle the competition between various partial reactions $(\gamma, 1n)$, $(\gamma, 2n)$, $(\gamma, 3n)$, *etc.* over a wide range of energies.

Large arrays of $^{10}\text{BF}_3$ tubes embedded in a paraffin or polyethylene matrix in several concentric rings were used at Livermore. For example, the array of 48 $^{10}\text{BF}_3$ tubes in a paraffin moderator had a neutron detection efficiency of 45 – 30% in the energy range up to 5 MeV [33]. The ring-ratio technique was developed to determine the average neutron kinetic energy [33]. The ring-ratio, i.e. the ratio of the neutron count of the outer-ring detector to that of the inner-ring detector, is a monotonically-increasing function with neutron energy. It allows to distinguish, for example, between $(\gamma, 2n)$ and two $(\gamma, 1n)$ events. Thus, it became possible to directly determine/probe the total photoneutron (Eq. (3)) and neutron yield (Eq. (2)) cross sections.

A large tank of Gd-loaded liquid scintillator was used at Saclay. The high neutron detection efficiency of over 90% was achieved in the energy range up to 5 MeV, allowing direct measurements of partial photoneutron reaction cross sections, from which the total and neutron-yield cross sections, Eqs. (2) and (3), were determined. The discrimination of various partial reaction cross sections was more complex and less reliable with the Livermore detector than with the Saclay detector, even with the help of the ring-ratio technique. The reason was that the detection efficiency of the Livermore detector was lower and varied more largely with energy than that of the Saclay detector. The Saclay detector, on the other hand, suffered from pile-up and background events as well as dead-time caused by a γ -flash from the target [22].

The results obtained at the two laboratories for the same nuclei show intricate discrepancies in magnitude not only for the total but for partial cross sections as well. For example, it was found [34–36] that, in general, the Saclay $(\gamma, 1n)$ cross sections are larger than the Livermore data, whereas the Saclay $(\gamma, 2n)$ cross sections are smaller than the corresponding Livermore data. This is discussed in detail in Sec. V A.

3. Bremsstrahlung Activation

A series of activation measurements was performed by using high-intensity bremsstrahlung. The partial photoneutron reaction cross sections $(\gamma, 1n)$, $(\gamma, 2n)$, $(\gamma, 3n)$, *etc.* were determined by measuring the activity of the residual unstable nuclei.

After irradiating a target by bremsstrahlung with various end-point energies, γ -ray counting was carried out for residual nuclei. The γ -lines unique to residual nuclei were clearly identified in the γ -ray spectra measured

with an energy- and efficiency-calibrated high-purity germanium (HPGe) detector [37–39]. Photonuclear reaction cross sections were deduced from the γ -lines.

4. Direct Neutron-multiplicity Sorting with a Flat-efficiency Detector

The partial photoneutron cross section $\sigma(\gamma, in)$ with neutron multiplicity $i = 1, 2, 3, \dots$ can typically be determined experimentally from the number of (γ, in) reactions N_i by

$$N_i = N_\gamma N_T \sigma(\gamma, in), \quad (16)$$

where N_γ is the number of γ -rays incident on a target, and N_T is the number of target nuclei per unit area.

The problem with using Eq. (16) lies in the fact that N_i is not a direct experimental observable. In general, the neutron detection efficiency of moderator-based neutron detectors depends on the neutron kinetic energy. The ring-ratio technique [33] was developed at Livermore to determine the average neutron energy. However, this technique cannot be applied to N_i , but to the experimental observable, i.e. multi-neutron coincidence events. This may be a source of uncertainties associated with the Livermore partial photoneutron cross sections.

A novel technique [10] was developed to overcome the shortcomings of the neutron-multiplicity sorting of Livermore. This technique, referred to as direct neutron-multiplicity sorting with a flat-efficiency detector (FED), combined with the LCS γ -ray beam provides an experimental opportunity to obtain partial photoneutron cross sections with improved reliability.

In the case of neutron detection, for example, between $(\gamma, 3n)$ and $(\gamma, 4n)$ thresholds, using a pulsed γ -ray beam, j -fold events for single ($j = 1$), double ($j = 2$), and triple ($j = 3$) neutron coincidences are observed.

The single neutron event corresponds to observing only one neutron during the time interval of two successive γ -ray pulses. There are three contributions from $(\gamma, 1n)$, $(\gamma, 2n)$, and $(\gamma, 3n)$ reactions to the single neutron event as

$$\begin{aligned} N_s = & N_1 \varepsilon(E_1) \\ & + N_2 {}_2C_1 \varepsilon(E_2) \{1 - \varepsilon(E_2)\} \\ & + N_3 {}_3C_1 \varepsilon(E_3) \{1 - \varepsilon(E_2)\}^2. \end{aligned} \quad (17)$$

The first term means simply that one neutron emitted in the $(\gamma, 1n)$ reaction is observed with detection efficiency $\varepsilon(E_1)$ for neutron kinetic energy E_1 . The second term means that one of two neutrons emitted in the $(\gamma, 2n)$ reaction is observed with detection efficiency $\varepsilon(E_2)$ for neutron kinetic energy E_2 and that the other neutron is not observed with unobserved efficiency $(1 - \varepsilon(E_2))$. The third term corresponds to the observation of one of three neutrons emitted in the $(\gamma, 3n)$ reaction. It is noted that there is no way to know E_1 , E_2 , and E_3 because the ring-ratio technique is applied

to the experimental observable N_s , not to the number of reactions N_1 , N_2 , and N_3 , individually. Furthermore, the neutron kinetic energy depends on the emission order from an excited nucleus. Therefore, the second term of Eq. (17) should be written as

$$\begin{aligned} & N_2 {}_2C_1 \varepsilon(E_2) \{1 - \varepsilon(E_2)\} \\ &= N_2 \varepsilon(E_{21}) \{1 - \varepsilon(E_{22})\} \\ &+ N_2 \varepsilon(E_{22}) \{1 - \varepsilon(E_{21})\}, \end{aligned} \quad (18)$$

using kinetic energies E_{21} and E_{22} of the first neutron and second neutron emitted, respectively.

The concept of the novel technique is to make the detection efficiency independent of neutron kinetic energies. Thus, using a constant efficiency ε , we can rewrite Eq. (17) as

$$\begin{aligned} N_s &= N_1 \varepsilon \\ &+ N_2 {}_2C_1 \varepsilon(1 - \varepsilon) \\ &+ N_3 {}_3C_1 \varepsilon(1 - \varepsilon)^2. \end{aligned} \quad (19)$$

Similarly, the double and triple neutron coincident events are written as

$$N_d = N_2 \varepsilon^2 + N_3 {}_3C_2 \varepsilon^2(1 - \varepsilon), \quad (20)$$

and

$$N_t = N_3 \varepsilon^3, \quad (21)$$

respectively.

One can solve the set of Eqs. (19) – (21) with known ε to obtain N_1 , N_2 , and N_3 from which then the partial cross sections $(\gamma, 1n)$, $(\gamma, 2n)$, and $(\gamma, 3n)$ are determined as in Eq. (16).

The FED consists of three concentric rings of 4, 9, and 18 ${}^3\text{He}$ counters embedded in a 46 cm (horizontally) $\times$ 46 cm (vertically) $\times$ 50 cm (along the beam axis) polyethylene moderator at the distances of 5.5, 13.0 and 16.0 cm from the γ -ray beam axis, respectively. The moderator is shielded by additional 5 cm-thick borated polyethylene plates for background neutron suppression. Figure 5 shows the total detection efficiency and efficiencies of the individual rings of the FED. Results of the calibration with a ${}^{252}\text{Cf}$ source are shown by the filled symbols. Results of the MCNP Monte Carlo simulations for monochromatic neutrons [10] are shown by the broken lines, while those for the neutron-evaporation spectra by the solid lines.

There are two issues that make the data reduction to obtain partial photoneutron cross sections rather complicated. First, double firings of photo-reactions can be induced by multi-photons involved in a single γ -ray pulse. For example, double firings of $(\gamma, 1n)$ reactions can be identified as a $(\gamma, 2n)$ reaction. Indeed, a small amount of non-zero $(\gamma, 2n)$ events attributable to such double firings was often observed below $(\gamma, 2n)$ threshold. Second, the electromagnetic interactions (pair production, Compton scattering, photo-electric absorption) of high-energy

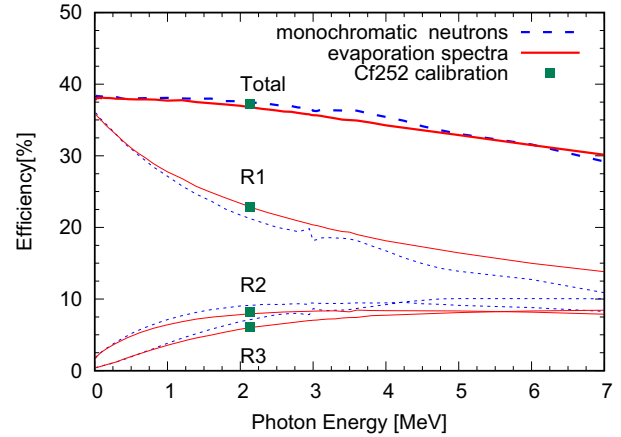


FIG. 5. (Color online) The total detection efficiency and efficiencies of three rings.

γ -rays in thick high- Z target material can generate secondary γ -rays which can induce the giant dipole resonance most effectively in the peak region which is mostly governed by the $(\gamma, 1n)$ channel. Thus, the secondary gamma rays produce extra neutrons which may mistakenly be assigned to reaction neutrons of the $(\gamma, 1n)$ channel associated with the primary gamma rays. Such effect was observed when a 10mm-thick ${}^{209}\text{Bi}$ target was irradiated with a 40 MeV LCS γ -ray beam.

The procedure of unfolding the photoneutron cross section which is convoluted with the energy distribution of the LCS γ -ray beam (Fig. 4) is rather straightforward because of the monochromaticity of the LCS γ -ray beam with low background. Previously, the Taylor expansion method [40] and the least-squares fitting method [41] were developed. These methods were used to mostly deduce $(\gamma, 1n)$ cross sections at the average γ -ray energy below the $(\gamma, 2n)$ reaction threshold [42–45]. More recently, a new unfolding method has been developed [46] that deduces partial photoneutron cross sections at the maximum γ -ray energy corresponding to the maximum γ -ray yield (Fig. 4) over a wide energy range. The unfolded cross sections resulting from the Taylor expansion method and the new method were cross-checked in the case of ${}^{209}\text{Bi}$ [47] partial cross sections and were found to be in good agreement.

C. Experimental Databases and Resources

Reference [3] includes a large collection of experimental information available at that time. Here we expand the list by including newly available information. For completeness we briefly provide older resources as well.

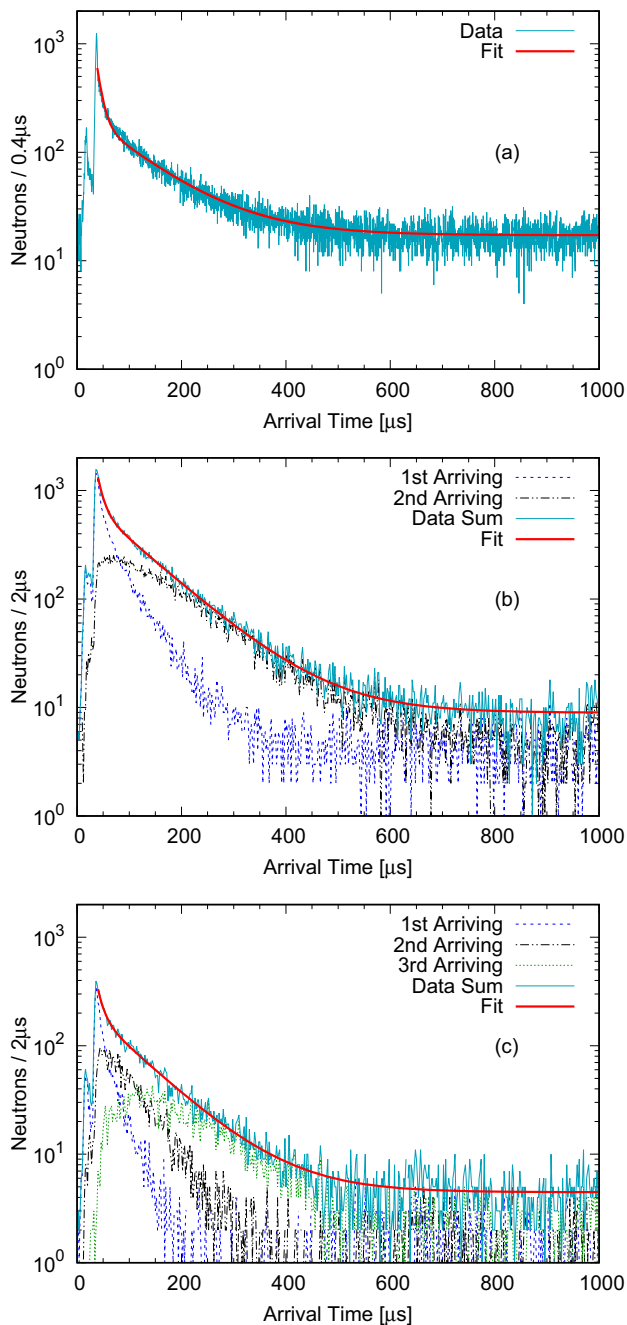


FIG. 6. (Color online) Neutron arrival-time distributions for (a) the single, (b) double, and (c) triple neutron events in the $^{209}\text{Bi}(\gamma, in)$ reactions at 34 MeV.

1. Bibliographic Data

The list of photonuclear reaction bibliographies has remained unchanged since the publication of the previous IAEA Photonuclear Data Library in 1999 and can be found in Ref. [3]. The list includes mainly compilations of

total and partial reaction cross sections for photo-induced disintegration and fission, photon absorption and scattering processes up to 1999. The list includes, for example, “photonuclear reactions,” 10th IAEA Bibliographical Series at IAEA [48], “Photonuclear data - abstract sheets 1955 – 1982,” at National Institute of Standards and Technology (NIST) [49], “Photonuclear data index 1976 – 1995,” at Centr Dannykh Fotoyadernykh Eksperimentov (CDFE) [50], and “Bibliographic Index to photonuclear reaction data (1955 – 1992)” at Japan Atomic Energy Research Institute (JAERI) [51]. More detailed information is given in Ref. [3]. These indices are available at CDFE [52].

2. Compilation of Photonuclear Cross Sections and GDR Parameters

The photonuclear cross sections and GDR parameters compiled prior to this CRP, have been reviewed in Ref. [3]. Below we briefly summarize the important publications and the most recent compilation of GDR parameters performed within this CRP.

- “Handbook on Nuclear Activation Data” compiled by Forkman and Petersson and published in 1987 by IAEA [53];
- Dietrich and Berman [2] published “Atlas of photoneutron cross sections obtained with monoenergetic photons” in 1988, which contains the photonuclear data measured with quasi-monoenergetic photons by annihilation in flight of fast positrons and by bremsstrahlung tagging;
- “Plots of the Experimental and Evaluated Photonuclear Cross-Sections” by Blokhon and Nasyrova [54], published in 1991, gives graphs of experimental and evaluated photoneutron reaction data for some selected nuclides;
- Varlamov *et al.* [55] published “Atlas of GDR” in 1999, which is a compilation of many photonuclear reaction data. This report covers all the reaction data compiled by Antonescu [48], and is given in the Annex of previous IAEA photonuclear data library report [3];
- The GDR parameters for heated atomic nuclei are determined from the γ -decay data. Compilation and parametrization of the GDR resonances built on excited states are given by Schiller and Thoennessen [56] in 2007; and
- The most recent comprehensive databases of the GDR parameters with their uncertainties from ground-state photo-absorption are compiled by Plujko *et al.* [8, 57] in 2011 and 2018.

In the recent databases [8, 57], Plujko *et al.* applied the least-squares technique to the updated experimental

databases to obtain the GDR parameters and their uncertainties. The theoretical photo-absorption cross section, which is fitted to the experimental data, consists of both the GDR component with the SLO and SMLO models [58] and the QD contribution, which are given later. These databases [8, 57] were constructed based on the experimental data reported up to January 2010 and June 2017, respectively. Further updated GDR parameter tables, where the data published up to December 2018 are involved, are given in Appendix B.

Another source of experimental information of potential relevance to the evaluation of the light elements is the evaluated nuclear structure database by Ajzenberg-Selove and Lauritsen [59], now maintained and updated at TUNL (Triangle Universities Nuclear Laboratory, <http://www.tunl.duke.edu/nucldata>).

3. EXFOR Database

The EXFOR database [9, 60], maintained by the international network of Nuclear Reaction Data Centers (NRDC) [61], is the main foundation for producing evaluated nuclear data libraries nowadays. The database as well as the Web data retrieval system [62] facilitates access to published experimental data in a computer-readable format. It is fair to say that, as a result of the concerted compilation effort performed during the CRP, the database is at present complete with respect to photonuclear data relevant to applications.

IV. NUCLEAR MODELS AND CODES

The models described below can be applied virtually to all nuclei in the IAEA Photonuclear Data Library 2019, from ${}^6\text{Li}$ through ${}^{241}\text{Pu}$, except for ${}^2\text{H}$ and ${}^3\text{H}$.

A. Theory of Photonuclear Reactions

1. Photo-absorption Cross Section

To calculate and evaluate the photonuclear reaction cross sections in the 1 – 200 MeV photon energy range, we adopt the independent hypothesis of a compound nucleus (CN) reaction, namely that the decay of the CN is independent of how it is formed, whereas the energy, spin, and parity are conserved. A photon is absorbed by a nucleus through two distinct nuclear reaction mechanisms; the Giant Dipole Resonance (GDR) and the quasi-deuteron (QD) photo-absorption [63, 64]. While GDR is the dominant process in the energy range of typical photonuclear data applications ($E_\gamma = 10 \sim 20$ MeV), the QD model describes the photo-absorption for $E_\gamma > 30$ MeV or so. The experimentally observed photo-absorption cross sec-

tion σ_{abs} includes both of these mechanisms

$$\sigma_{\text{abs}}(E_\gamma) = \sigma_{\text{GDR}}(E_\gamma) + \sigma_{\text{QD}}(E_\gamma), \quad (22)$$

where σ_{GDR} and σ_{QD} are the absorption cross sections for the GDR and QD mechanism, respectively.

We often apply a phenomenological parameterization to describe the GDR part of the photonuclear reaction cross sections. When experimental data show isolated resonance-like structure, which is typical for light elements, these peaks can be represented by the Breit-Wigner shape. For the medium and heavy nuclei, several models can be used to represent the measured absorption cross section σ_{GDR} , such as Lorentzian, Breit-Wigner or Gaussian functions. However the most simple and suitable function that has been confirmed by experimental photo-absorption data [22] is a Lorentzian shape of

$$\sigma_{\text{GDR}}(E_\gamma) = \sigma_R \frac{E_\gamma^2 \Gamma_R^2}{(E_R^2 - E_\gamma^2)^2 + E_\gamma^2 \Gamma_R^2}, \quad (23)$$

where σ_R is the GDR peak cross section, E_R is the resonance energy, and Γ_R is the width. In the case of deformed nuclei, the GDR peak splits into two, where each corresponds to the major or minor axis of ellipsoid, and the photo-absorption cross section is given by the sum of two GDRs. This is often called the Standard Lorentzian (SLO) model for the photon strength function [58], which is also known as the Brink-Axel Lorentzian.

While the standard Lorentzian (SLO) parameterization of Eq. (23) is adequate to describe GDR around the peak, it fails to reproduce the data at lower energies around the neutron separation energy. Modified versions of the SLO, such as the generalized Lorentzian [65–67] and the Simple Modified Lorentzian (SMLO) [8, 57] were developed to account for data both at the GDR peak and at lower energies around the neutron separation energies. In addition to the phenomenological GDR models, significant effort has been devoted to developing more microscopic approaches such as the quasiparticle random phase approximation for the description of giant multipole resonances [68, 69] and even the shell model has been applied to electromagnetic excitations of light nuclei (see Ref. [11] for an overview).

For the purpose of updating the photonuclear evaluations in the IAEA Photonuclear Data Library, the SLO, SMLO and GLO models available in RIPL [58] were used to describe the GDR cross sections. The GDR parameters of these models have been fitted to available experimental data, and the systematic behavior of these parameters, as a function of Z and A numbers, E_γ , and/or nuclear deformation, has been extracted [58, 70]. When we assume the multipolarity of E1 is the largest contribution to the photo-absorption, the formed CN state will have the spin J of $|I - 1| \leq J \leq I + 1$, where I is the target nucleus spin, and the parity flips. This will be the initial configuration in the statistical model calculation.

The QD model of Chadwick *et al.* [71] is commonly implemented in the model codes we employ, where the

Levinger constant L is taken to be 6.5. This is a scaling constant that has been adjusted to the experimental photo-absorption cross section σ_{QD} by [71], and is seldom fine-tuned in practical calculations. The model reads

$$\sigma_{\text{QD}} = L \frac{NZ}{A} \sigma_d p_b, \quad (24)$$

where the photo-disintegration cross section σ_d is parameterized as

$$\sigma_d = \begin{cases} 61.2(E_\gamma - B_d)^{3/2} E_\gamma^{-3} \text{ mb} & E_\gamma > B_d \\ 0 & \text{otherwise} \end{cases}, \quad (25)$$

where the energies are in MeV, and B_d is the deuteron binding energy (2.22452 MeV). In the energy range $20 \leq E_\gamma \leq 140$ MeV, the Pauli blocking factor P_b is given by a polynomial,

$$P_b = 0.083714 - 0.0098343 E_\gamma + 4.1222 \times 10^{-4} E_\gamma^2 - 3.4762 \times 10^{-6} E_\gamma^3 + 9.3537 \times 10^{-9} E_\gamma^4, \quad (26)$$

and it is extrapolated to energies beyond this range as

$$P_b = \begin{cases} \exp(-73.3/E_\gamma) & E_\gamma < 20 \text{ MeV} \\ \exp(-24.2348/E_\gamma) & E_\gamma > 140 \text{ MeV} \end{cases}. \quad (27)$$

Assuming the QD component is negligible in the GDR energy range, the determination of the GDR parameters σ_R , E_R , and Γ_R can be performed by taking $\sigma_{\text{abs}} \simeq \sigma_{\text{GDR}}$.

2. Pre-equilibrium Particle Emission

Nucleon or composite particle emission during the pre-equilibrium process has not been studied extensively for photonuclear reactions. In the GDR photo-absorption mechanism, the initial nuclear excitation can be interpreted as a superposition of particle-hole excitations in the shell model space, whereby the $1p-1h$ configuration initiates the pre-equilibrium chain in the Q -space as classified in the theory of Feshbach, Kerman, and Koonin [72], and eventually one nucleon is scattered into the $\mathcal{P}$ -space. The QD model directly leads to the pre-equilibrium chain by creating the $2p-2h$ configuration.

However, the nuclear reaction codes that were employed to create the photonuclear data library — EMPIRE [73, 74], TALYS [75], CCONE [76], MEND-G [77], GLUNF [78], and CoH₃ [79] — have implemented relatively simple pre-equilibrium models for photonuclear reactions. In some cases, the exciton model for a neutron-induced reaction is considered as a surrogate for the photo-induced precompound process, and the initial configuration of $1p-0h$ or $2p-1h$ is assumed.

The pre-equilibrium process implemented in CoH₃ is different from the other codes. A fraction of the pre-equilibrium process in the GDR absorption is restricted

to the bound $1p-1h$ configuration,

$$\sigma_a^{\text{PE}} = R \sigma_{\text{GDR}}, \quad (28)$$

$$R = \frac{\omega_B(1p, 1h)}{\omega(1p, 1h)} \simeq \frac{S_n}{E_\gamma - \Delta}, \quad (29)$$

where S_n is the neutron separation energy, Δ is the pairing energy, ω is the $1p-1h$ state density, while ω_B is $1p-1h$ but with all particles in the bound state. When we insert the state density of Williams' [80] for ω , and Betak and Dobes [81] for ω_B , the right-hand-side of Eq. (29) becomes simply $S_n/(E_\gamma - \Delta)$. This is analogous to a correction factor in the multi-step compound strength [82, 83]. The QD process creates a $2p-2h$ initial configuration, and CoH₃ makes $1p-1h$ configuration in both the neutron and proton shells. On the other hand, the GDR $1p-1h$ is created in the neutron shell. Because we have two different initial configurations, CoH₃ calculates the pre-equilibrium process twice and sums the results.

To illustrate the impact on the pre-equilibrium calculations of the different treatment of the initial configuration as well as the pre-equilibrium damping factor in Eq. (29), we calculate the $(\gamma, 1n)$ and $(\gamma, 2n)$ cross sections of ¹⁸¹Ta with the initial configuration of $1p-0h$, $1p-1h$, and $2p-1h$. The results are shown by the ratio to the CoH₃ default calculation, which is the $1p-1h$ for GDR and $2p-2h$ for QD, in Fig. 7. Because $\sigma_{\text{abs}} \simeq \sigma_{\text{QD}}$ above 20 MeV, the difference increases rapidly as the photon energy increases. However, at these higher energies we are comparing very small cross sections obtained from Eq. (24). The uncertainty in σ_{QD} could be comparable to the pre-equilibrium model deficiency.

For the light elements, at present, there is no model available to calculate proper pre-equilibrium emission other than the approximate QD model, therefore, the results need to be treated with caution as the model deficiencies may be larger.

3. Decay of Compound Nucleus

The decay of the CN state formed by the photo-absorption reaction is treated in the usual Hauser-Feshbach model [84], the only difference being the relatively narrow distribution of compound nucleus J compared to the particle-induced CN reactions. We define a population $P(E_x, J, \Pi)$ at the excitation energy of E_x , which is formed by the incoming photon $E_\gamma = E_x$, and is normalized to $\sigma_{\text{abs}}(E_\gamma)$ as

$$\sigma_{\text{abs}}(E_\gamma) = \sum_{J, \Pi} P(E_x, J, \Pi). \quad (30)$$

The decay of the CN state (E_x, J, Π) into a final outgoing channel is governed by the branching ratio for that channel which is calculated using the particle and γ -ray transmission coefficients. For a nucleon or a composite particle, the transmission coefficient is denoted by $T_{lj}(\epsilon)$, where l is the orbital angular momentum, spin j , and the

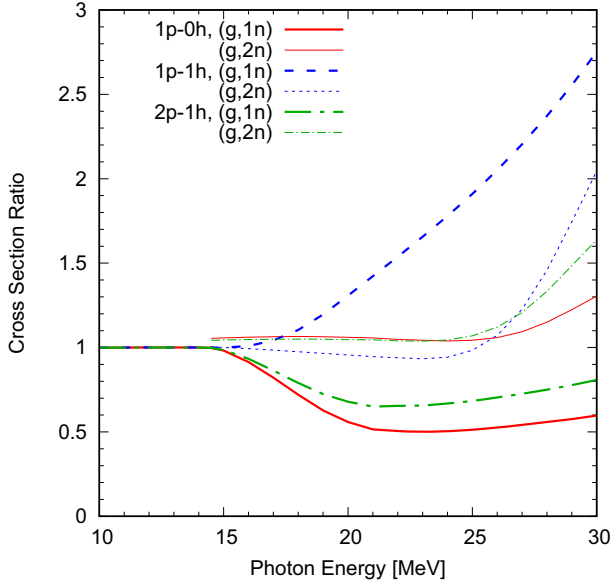


FIG. 7. (Color online) Calculated $^{181}\text{Ta}(\gamma, n)$ and $(\gamma, 2n)$ cross section when the initial exciton configuration is $1p-0h$ (solid), $1p-1h$ (dashed), and $2p-1h$ (dot-dashed). The calculated cross sections are shown by the ratios to the CoH3 default calculation.

center-of-mass energy ϵ . The transmission coefficient for the γ -ray emission is specified by the multipolarity XL , where $X = E$ or M . Hereafter we drop Π since the parity conservation is trivial.

To simplify, we consider a case where no charged particles are emitted, and all the final states are in the continuum. A population of (E'_x, J') formed after the γ -ray transition is written as

$$P(E'_x, J') = \frac{P(E_x, J)}{N} \sum_{XL} T_{XL}(E_\gamma) \times \rho_0(E'_x, J') dE, \quad (31)$$

$$E_\gamma = E_x - E'_x, \quad (32)$$

where N is the normalization shown later, $\rho_0(E, J)$ is the level density of the same nucleus as the target, and $|J - J'| \leq L \leq J + J'$ is fulfilled in the summation. When a neutron is emitted, the population of (E'_x, J') in the $(Z, A - 1)$ nucleus is

$$P(E'_x, J') = \frac{P(E_x, J)}{N} \sum_{lj} T_{lj}(E_n) \times \rho_1(E'_x, J') dE, \quad (33)$$

$$E_n = E_x - E'_x - S_n, \quad (34)$$

where S_n is the neutron separation energy, the level density $\rho_1(E'_x, J')$ is for the residual $(Z, A - 1)$, and the triangular relation reads $|J - J'| \leq j \leq J + J'$. The normalization factor N is given by integrating all the possible

final states,

$$N = \int_0^{E_x} \sum_{XL} T_{XL}(E_\gamma) \rho_0(E'_x, J') dE + \int_0^{E_x - S_n} \sum_{lj} T_{lj}(E_n) \rho_1(E'_x, J') dE. \quad (35)$$

The neutron transmission coefficient T_{lj} is obtained by solving the Schrödinger equation for a complex one-body (optical) potential. The γ -ray transmission coefficients are calculated from the photon-strength function $f_{XL}(\epsilon)$ for the γ -decay as

$$T_{XL}(\epsilon) = 2\pi\epsilon^{2L+1} f_{XL}(\epsilon). \quad (36)$$

When the final state is in the discrete part of the level scheme, the integration in Eq. (35) is replaced by a proper summation over branching ratios to these discrete levels which are often taken from the nuclear structure database [58]. An extension of these formulas to the case where charged-particle channels are involved should be straightforward. These calculations are repeated until the initial excitation energy E_x is exhausted by the emitted particle and γ -ray energies, as well as the particle binding energies.

One of the theoretical problems in the light element analysis is the replacement of the photon decay width by the gamma-ray strength function, where an energy-average procedure is involved. This is because there is no way to obtain an exact photon decay width in the CN reaction, hence the best we can do is to use the strength function, which leads to the Hauser-Feshbach theory. By doing that, the width fluctuation would be involved, but we ignore this as the elastic cross section is very small as discussed in Sec. IV B.

4. Energy and Angular Distributions of Secondary Particles

The secondary particle energy and angular distributions are less known experimentally, hence this type of information relies strongly on the nuclear reaction models that are used to calculate the cross sections. The angle-integrated particle emission spectra consists of two components — the pre-equilibrium contribution and the evaporation component. Many of the reaction codes employ the exciton model and the Hauser-Feshbach statistical model to calculate the angle-integrated energy spectra. As mentioned above, different pre-equilibrium exciton models are implemented in the codes, therefore the hardness of the calculated energy spectra can be different. These differences are not expected to be significant, however, since the most important energy range is near GDR, where the CN decay dominates giving an isotropic evaporation spectrum and all the codes describe this part similarly.

A forward-peaked angular distribution is expected at higher incident photon energies due to the pre-equilibrium process. This is not so extensively studied

for the photo-induced reaction case, and there is no general description for the evaluation of the angular distribution. A practical workaround [85], which was also applied to the previous IAEA photonuclear data library, is to emulate the neutron-induced reaction, and adopt the double-differential data systematics by Kalbach [86]. As an example, Fig. 8 shows the energy and angular distribution for the emitted neutron from ^{56}Fe at the incident photon energy of 20 MeV.

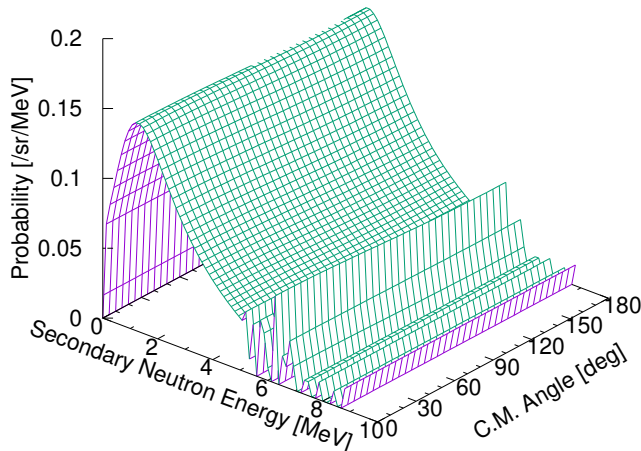


FIG. 8. (Color online) Energy and angular distribution of emitted neutron for the $^{56}\text{Fe}(\gamma, n)$ reaction at $E_\gamma = 20$ MeV.

5. Model Parameters in Photonuclear Reaction Calculation

As mentioned in the previous section, there are some differences in the pre-equilibrium models implemented in the nuclear reaction codes that have been employed to evaluate the photonuclear reaction cross sections. These differences are, however, notable in the high energy region only. The dominant photonuclear reaction part is in the GDR energy range, and the differences in the model calculations mainly come from the Hauser-Feshbach model parameters. These parameters include the neutron and charged-particle optical potentials, the level density, and the γ -ray strength function. When the photo-absorption cross section is estimated within the reaction code itself, the built-in GDR parameter library can also be a source of difference between the various calculations. The global optical potential of Koning and Delaroche [87] is widely used in these type of calculations, therefore the differences in the calculations are mostly due to the level density models. Furthermore, the probability of γ -ray emission is much smaller than particle emission, so differences among the γ -ray strength functions have a very modest impact on the photonuclear reaction calculations.

B. Nuclear Reaction Codes for Photonuclear Data Evaluation

Since it is almost impossible to obtain complete information for photonuclear reactions from experiments only, the nuclear reaction codes play a significant role in creating the evaluated data library. For example, the only possible way to generate the evaluated energy and angular distributions of the emitted particle is to use the model calculations, which, of course, should consistently describe the available cross section data. In the previous release of the IAEA photonuclear data library [3], the evaluation was done using several reaction model codes available at the participating institutes: GNASH [88, 89] at LANL and KAERI, ALICE-F [90] and MCPHOTO [91] at JAERI, GUNF [92] and GLUNF [78] at CIAE, and XGFISS [93] at IPPE.

The modern Hauser-Feshbach nuclear reaction codes, EMPIRE [73, 74], TALYS [75], CCONE [76], and CoH₃ [79], which are often utilized for nucleon-induced reaction calculations, are also capable of calculating photo-induced reaction cross sections. As a result, they are widely used in the evaluation of general purpose nuclear data libraries, and are combined with the advanced-capabilities utility codes to create ENDF-6 format data files. Albeit these codes are built on common well-established nuclear reaction models to describe the various reaction mechanisms, the implementation of the nuclear reaction models in each code may give rise to differences in the calculated results which are inherent to the code and do not depend on the model or model parameters. It is therefore important to understand the characteristics of each code before comparing evaluations performed with the codes. Recently, an inter-comparison of the codes EMPIRE, TALYS, CCONE, and CoH₃ was performed for neutron-induced reactions on actinides [94] by fixing all the input parameters as consistently as possible, so that any observed difference would be due to the model implementation. It was concluded that in terms of model implementation, the implementation of the width fluctuation correction might give rise to differences in the calculated cross sections by up to $\approx 15\%$. The ambiguity related to the width fluctuation correction is not expected to affect the photonuclear reaction case, since the (γ, n) channel is much larger than the photon elastic scattering (the elastic enhancement does not impact (γ, n) [95].) However, as already discussed in Sec. IV A, the pre-equilibrium models employed in the different codes are different therefore the pre-equilibrium component is one of the sources of ambiguity among the codes.

The newly evaluated photonuclear data were produced with the codes EMPIRE, TALYS, CCONE, GLUNF and MEND-G. Some data files were produced with ALICE-F as was the case in the previous IAEA photonuclear data library. CoH₃ has not been used to produce any evaluations, since it focuses mainly on low-energy nuclear reactions. However, CoH₃ has been considered in the

code comparison. The CPNRM (Combined Photonuclear Reaction Model) code [96, 97] developed at SINP/MSU is also included in the code comparison as it has been used extensively to correct the experimental data as described in Sec. V A.

1. Special Notes on Each Model Code

As described in Sec. IV A, photonuclear reaction calculations consist of three stages, the photo-absorption process, the pre-equilibrium particle emission, and the statistical Hauser-Feshbach decay. The absorption cross section is unambiguously determined by providing a set of GDR parameters (energy, width and peak cross section) using a standard Lorentzian or variants such as the Generalized Lorentzian or Simple Modified Lorentzian. Alternatively the absorption cross section can be pre-calculated and stored in an external file.

For the pre-equilibrium emission component, the codes adopt the one-component or two-component exciton model, however the initial particle-hole configuration used in each code may be different. CoH₃ assumes the initial configuration is $1p\text{-}1h$ created in the neutron shell for GDR, and $2p\text{-}2h$ ($1p\text{-}1h$ in both neutron and proton shells) for the QD process. It also includes the damping factor of Eq. (29). EMPIRE begins the calculation with $1p\text{-}0h$, unless specified otherwise. TALYS also assumes $1p\text{-}0h$. When the two-component exciton model is selected, the particle-hole pair is given in the neutron shell. In MEND-G, $2p\text{-}2h$ is the initial configuration for the QD contribution. CCONE assumes $1p\text{-}0h$ as the initial configuration. The particle-hole pair in the two-component model is given in the neutron shell for photo-reaction.

In the statistical decay stage, the particle transmission coefficients are calculated using an optical potential. Table I summarizes the optical potentials employed in each code for the evaluation. Since often the potential parameters are adjusted to better reproduce experimental data, we provide a default set of potentials when not specified. CoH₃ does not offer a default option for the potential. However, those given in the table are typical choices for the photonuclear reaction calculations.

2. Photonuclear Reaction Code Inter-comparison

The F_i values in Eq. (7) were calculated with CPNRM, TALYS, CCONE, CoH₃, EMPIRE, and MEND-G, for the photo-induced reactions on ¹⁸¹Ta. We chose all default model parameters in each code. The results are shown in Fig. 9, together with the evaluated experimental data by Varlamov *et al.* [109]. Large discrepancies amongst the model codes are seen in the high energy region, where pre-equilibrium emission is the dominant process, yet it should be noted that the cross sections in this region are tiny compared to GDR. In general, CPNRM gives the largest pre-equilibrium contribution, while the

TABLE I. Optical potential used in the photonuclear cross section calculations. The abbreviations are, KD03 (Koning and Delaroche [87]), KD03-f (folding KD03 potential by technique of Watanabe [98]), AV94 (Avrigneanu *et al.* [99]), AV09 (Avrigneanu *et al.* [100]), AV10 (Avrigneanu *et al.* [101]), AV14 (Avrigneanu *et al.* [102]), AC06 (An and Cai [103]), BO88 (Bjowald *et al.* [104]), BG69 (Becchetti and Greenlees) [105, 106], HA06 (Han *et al.*) [107], and XU11 (Xu *et al.*) [108].

	n	p	α	d	t	³ He
EMPIRE	KD03	KD03	AV94	AC06	BG69	BG69
TALYS	KD03	KD03	AV14	KD03-f	KD03-f	KD03-f
CCONE	KD03	KD03	AV10	HA06	KD03-f	XU11
MEND-G	KD03	KD03	BG69	BG69	BG69	BG69
CoH ₃	KD03	KD03	AV09	BO88	BG69	BG69

pre-equilibrium component of CoH₃ is the weakest. This is understood as a result of applying the correction factor of Eq. (29) in CoH₃. The F_1 and F_2 values by CPNRM increase above 30 MeV, which is not seen in the other codes.

From Fig. 9 it is clear that not only do the codes give different values of F_i , but that these differences vary with energy. The theoretical F_i values are determined by the competition between the neutron and the γ -ray emissions in the statistical decay of CN. Obviously they depend on the neutron and γ -ray transmission coefficients, as well as the level density of the final states. The conclusion is that if the theoretical F_i s are used to correct problematic or inconsistent experimental data, as proposed in Sec. V A, then the uncertainties in F_i due to different codes and model parameters need to be considered. Moreover, since these uncertainties are shown to vary with energy, the introduction of an overall constant uncertainty may not be adequate.

V. EVALUATION

The new evaluations for the updated Photonuclear Data Library were performed at five institutes: SINP/MSU (Skobeltsyn Institute of Nuclear Physics, Lomonosov Moscow State University), NDC/KAERI (Nuclear Data Center, Korean Atomic Energy Research Institute), IFIN-HH (Horia Hulubei National Institute for Physics and Nuclear Engineering), NDC/JAEA (Nuclear Data Center, Japan Atomic Energy Agency), and CNDC/CIAE (China Nuclear Data Center, China Institute of Atomic Energy). In the following sections, a summary of the evaluation procedure, adoption of experimental data, and models used by each institute is given along with highlights of the resulting evaluations. The complete list of evaluations with graphical comparisons will be made available from an IAEA web interface.

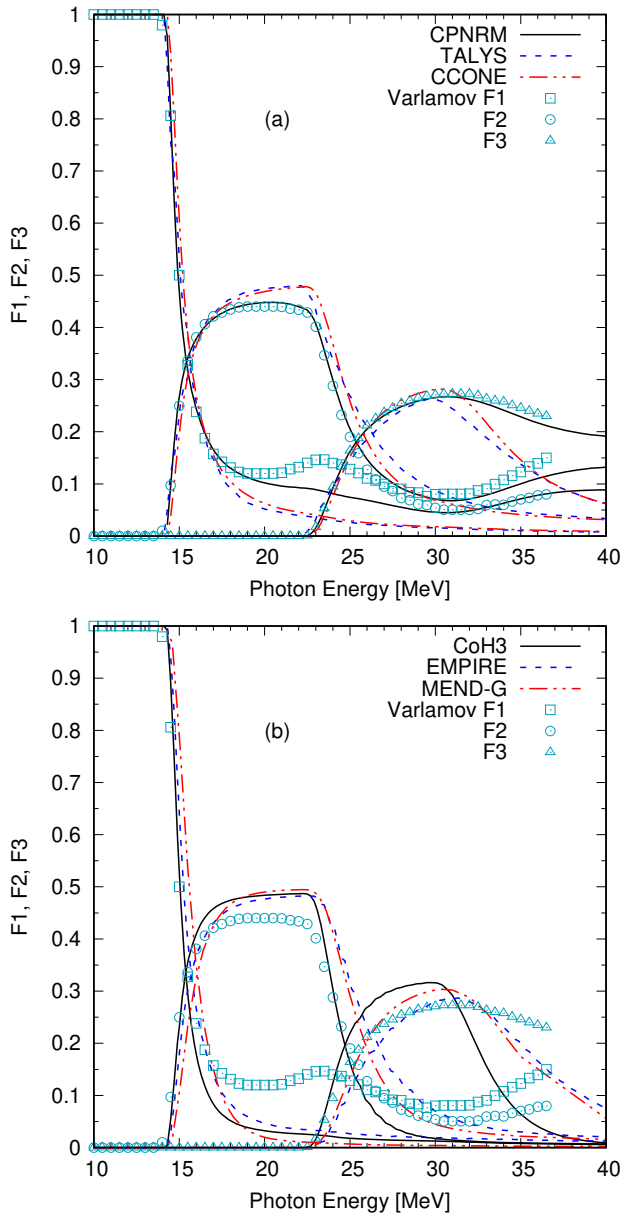


FIG. 9. (Color online) Calculated F_i values for $^{181}\text{Ta}(\gamma, inX)$ cross sections with (a) CPNRM, TALYS, and CCONE, and (b) CoH3, EMPIRE, and MEND-G. The symbols are evaluated experimental data by Varlamov *et al.*

A. Evaluation of Experimental Data

The majority of available experimental total and partial photoneutron reaction cross sections were obtained by the quasi-monoenergetic annihilation photon beam and the photoneutron multiplicity sorting technique at Livermore and Saclay [1, 2, 22]. Significant systematic disagreements in the experimental $(\gamma, 1n)$ and $(\gamma, 2n)$ data from these two institutes have been reported for

19 nuclei [35, 110, 111]: ^{51}V , ^{75}As , ^{89}Y , ^{90}Zr , ^{115}In , $^{116,117,118,120,124}\text{Sn}$, ^{127}I , ^{159}Tb , ^{181}Ta , ^{197}Au , ^{208}Pb , ^{232}Th , and ^{238}U . It was found that the $(\gamma, 1n)$ cross sections from Saclay tend to be larger than the Livermore data by up to 100%, while Livermore gives larger $(\gamma, 2n)$ cross sections by up to 100%. An average overall estimate of the differences is given by the ratio of the experimental integrated cross section from Saclay σ_S^{int} to that from Livermore σ_L^{int} . The experimental integrated cross sections are obtained from

$$\sigma_{S,L}^{\text{int}} = \int_{E_{\text{th}}}^{E_{\text{max}}} \sigma_{S,L}(E_\gamma) dE_\gamma, \quad (37)$$

where E_{th} is the threshold energy of $1n$ or $2n$ reaction and E_{max} is the highest photon energy in the experiments. The average value $\langle \sigma_S^{\text{int}} / \sigma_L^{\text{int}} \rangle$ is found to be 1.07 for the $(\gamma, 1n)$ reaction, and 0.84 for the $(\gamma, 2n)$ reaction. The ratio $\sigma_S^{\text{int}} / \sigma_L^{\text{int}}$ fluctuates considerably between 0.76 and 1.34 for the $(\gamma, 1n)$ cross section and between 0.71 and 1.22 for all the 19 nuclei mentioned above. Given the large fluctuations in $\sigma_S^{\text{int}} / \sigma_L^{\text{int}}$, the discrepancies between the Livermore and Saclay data cannot be rectified by simply applying a constant normalization factor common to all the 19 nuclei.

On the other hand, the neutron yield cross-section data from these two institutes (Eq. (2)) also disagree by up to 10% on average [111]. This implies that the systematic inconsistency observed in the measured partial reaction cross sections may be due to a shortcoming of the neutron multiplicity sorting method. In order to understand the underlying systematic uncertainties in the experimental data, Varlamov *et al.* [12] introduced the F_i function of Eq. (7). This function provides objective criteria for assessing the consistency and reliability of the experimental partial reaction cross-sections obtained in a given measurement, and allows one to investigate the systematic uncertainties. As has been mentioned in Sec. II, since F_1 is defined as the ratio of $\sigma(\gamma, 1n)$ to the photoneutron yield cross section $\sigma(\gamma, xn)$, it can never be larger than 1; analogously F_2 cannot exceed 1/2, F_3 cannot exceed 1/3, and so on. In addition, F_i should always be positive. These constraints on $F_i(E)$ values allow one to identify a potential problem in the way the neutron yield cross section has been partitioned into the individual one, two, three, or even more neutron emission components.

Varlamov *et al.* [12] has shown that the constraints imposed by F_i are not satisfied in many cases [12, 109–116]. The problematic experimental data, where negative cross-sections are reported, include the measurements of photoneutron reactions on ^{80}Se , $^{91,94}\text{Zr}$, ^{115}In , $^{112-124}\text{Sn}$, ^{133}Cs , ^{138}Ba , ^{159}Tb , ^{181}Ta , $^{186-192}\text{Os}$, ^{197}Au , ^{208}Pb , ^{209}Bi , and some others. In these cases, the corresponding F_1^{exp} values and/or F_2^{exp} values are found to be larger than the aforementioned upper limits, or negative as well. A typical example of the energy dependent $F_i^{\text{exp}}(E)$ values for ^{65}Cu [117] is given in Fig. 10, where it is compared with the theoretical $F_i^{\text{th}}(E)$ curves [115] obtained for partial reactions (γ, in) with $i = 1$ and 2

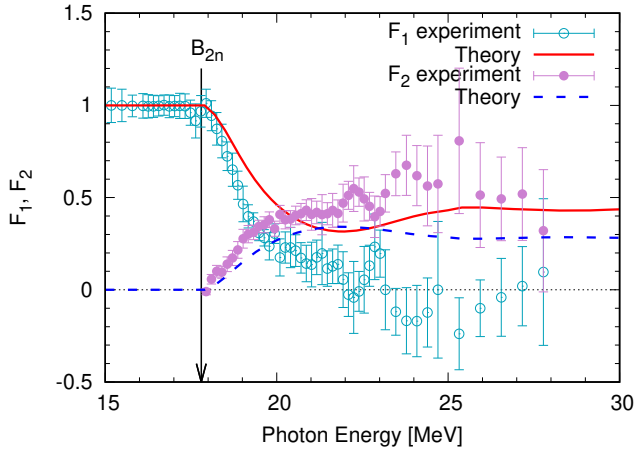


FIG. 10. (Color online) Comparison of the ratios F_i^{exp} [115] using the experimental data (triangles; Ref. [117]) and the theoretical F_i^{th} functions for ^{65}Cu .

within the framework of the CPNRM code [96, 97].

The partial photoneutron reactions cross sections should be evaluated under the condition that the inherent issues in the neutron multiplicity sorting method as well as the limitations of the statistical model do not form a bias. In other words, the systematic experimental uncertainties as well as model-dependent uncertainties need to be considered in the evaluation. The evaluation method proposed by Varlamov *et al.* [12], combines the experimental photoneutron yield data $\sigma^{\text{exp}}(\gamma, xn)$ (Eq. (2)) and the theoretical estimate of F_i^{th} obtained with the CPNRM code [96, 97] and is applicable to nuclei in the medium to heavy mass range. In this method, the physical reliability criteria mentioned above are automatically satisfied because the competition between the partial reaction cross sections is imposed by the model calculations performed with the CPNRM code. The evaluated partial photoneutron reaction cross section $\sigma^{\text{eval}}(\gamma, in)$ is obtained by multiplying the experimental photoneutron yield cross section $\sigma^{\text{exp}}(\gamma, xn)$ given in Eq. (2) by the theoretical F_i^{th} functions computed with the CPNRM code for neutron multiplicity $i = 1, 2, 3, \dots$

$$\begin{aligned}\sigma^{\text{eval}}(\gamma, in) &= F_i^{\text{th}} \sigma^{\text{exp}}(\gamma, xn) \\ &= \frac{\sigma^{\text{th}}(\gamma, in)}{\sigma^{\text{th}}(\gamma, xn)} \sigma^{\text{exp}}(\gamma, xn) .\end{aligned}\quad (38)$$

The former term in Eq. (38) satisfies the reliability criteria as already mentioned, while the second term is not affected by the issues associated with the experimental neutron multiplicity sorting. The resulting derived cross section data $\sigma^{\text{eval}}(\gamma, in)$ should therefore be free from the systematic experimental uncertainties of the Livermore and Saclay data.

In many cases, the evaluated cross-section data $\sigma^{\text{eval}}(\gamma, in)$ obtained from Eq. (38) form a substantial correction to the experimental data obtained by the neutron multiplicity sorting method [12, 109–116]. A typical ex-

ample of this evaluation method is given in Fig. 11, where the results obtained for ^{59}Co are compared with the experimental data.

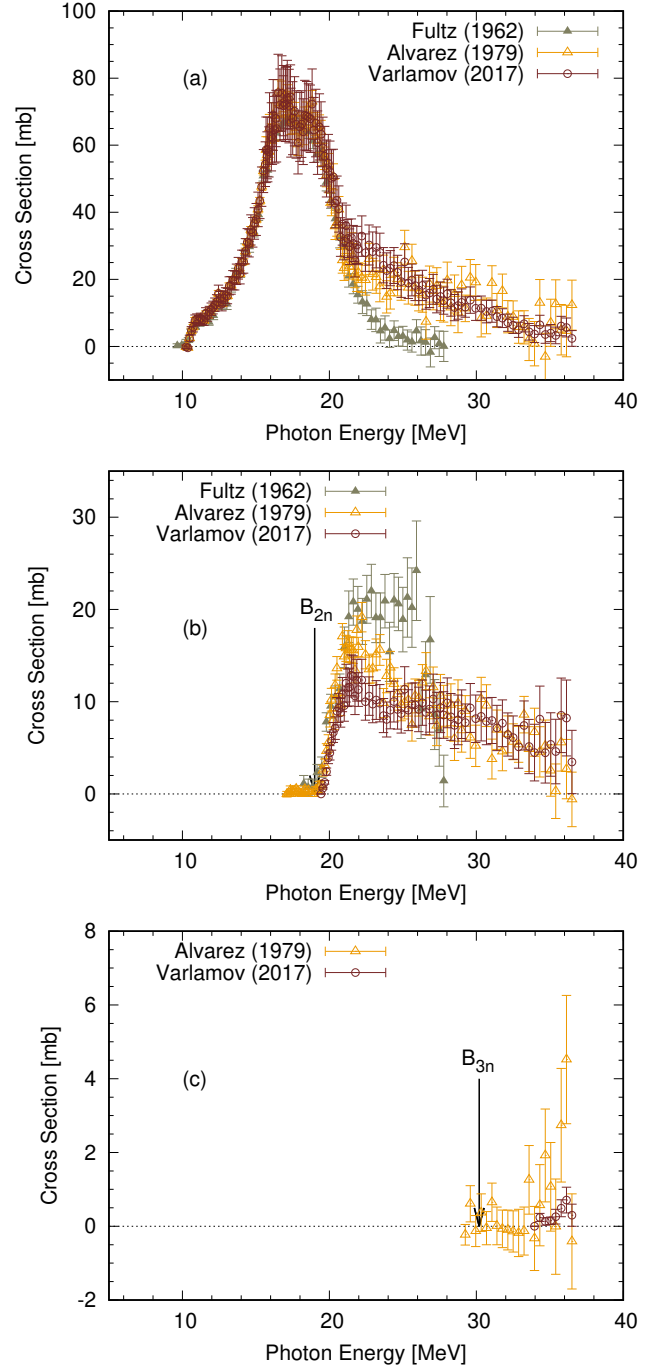


FIG. 11. (Color online) Comparison of the evaluated (open circles with error bars) [119] and experimental (filled triangles [120] and open triangles [121]) partial photoneutron cross sections for ^{59}Co . (a) $\sigma(\gamma, 1n)$, (b) $\sigma(\gamma, 2n)$, and (c) $\sigma(\gamma, 3n)$.

The neutron multiplicity sorting technique sometimes

cannot distinguish $1n$ and $2n$ events clearly, or $2n$ and $3n$ events. As a result, the partial cross sections $\sigma^{\text{eval}}(\gamma, in)$ fluctuate although the total photoneutron yield cross section $\sigma^{\text{exp}}(\gamma, xn)$ seems to be reasonable. This can be seen clearly by calculating the differences between the experimental and evaluated integrated cross sections $\sigma^{\text{int,exp}}$ and $\sigma^{\text{int,eval}}$ for the $(\gamma, 1n)$ reaction,

$$\Delta\sigma_1 = \sigma^{\text{int,eval}}(\gamma, 1n) - \sigma^{\text{int,exp}}(\gamma, 1n), \quad (39)$$

and for the $(\gamma, 2n)$ reaction,

$$\Delta\sigma_2 = \sigma^{\text{int,eval}}(\gamma, 2n) - \sigma^{\text{int,exp}}(\gamma, 2n). \quad (40)$$

An example of the differences is shown for ^{92}Zr [33, 118] in Fig. 12. From the figure, it is evident that some of the $(\gamma, 1n)$ events are recorded as $(\gamma, 2n)$ events. It has been shown [38, 111, 122] that this problem is due to the fact that neutrons were sorted into each reaction channel according to the measured neutron kinetic energy. However, the neutron energy spectra for different partial reactions may overlap with each other leading to cross-talks between different partial cross sections, whereby some of the neutrons originating from the $(\gamma, 1n)$ reaction are mistakenly assigned to those from the reaction $(\gamma, 2n)$ and vice versa.

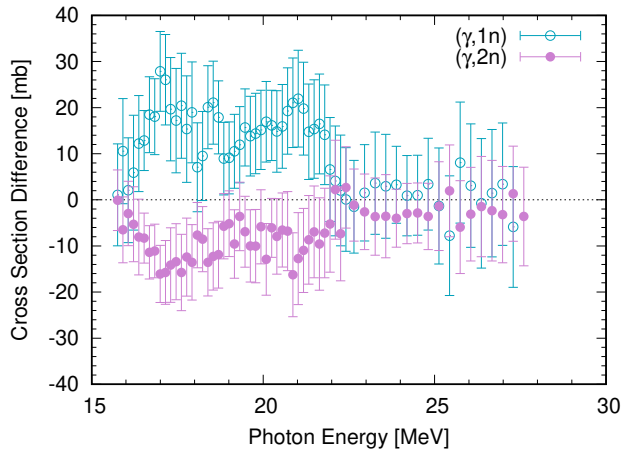


FIG. 12. (Color online) Differences between the evaluated [118] and the experimental [33] photonuclear cross sections for ^{92}Zr . The open circles are for the $(\gamma, 1n)$ reaction ($\Delta\sigma_1$ in Eq. (39)), and filled circles are for $(\gamma, 2n)$ reaction ($\Delta\sigma_2$ in Eq. (40)).

To test the evaluation method described above, partial photoneutron reaction cross sections evaluated for ^{181}Ta [109] and ^{209}Bi [116] were compared with those obtained in activation measurements of reaction yields with bremsstrahlung [38, 122]. In the activation measurement, cross sections of various partial reactions were directly determined by identifying the radioactivity from the residual nuclei. Ishkhanov *et al.* [122] and Belyshev *et al.* [38] obtained (γ, in) cross sections with $i = 1 -$

6 for ^{181}Ta and ^{209}Bi as a function of the bombarding photon energy based on the experimental photoneutron yields and the CPNRM calculations. The evaluated data for the photoneutron multiplicity sorting obtained agree well with the activation data. A further confirmation was the agreement found by making a detailed comparison [123] of the evaluated ^{197}Au $(\gamma, 1n)$ and $(\gamma, 2n)$ data with those obtained in activation measurements with bremsstrahlung [39].

Therefore, it is advisable to compare the F_i^{exp} with the from F_i^{th} and check whether the following listed conditions are satisfied. In case they are not satisfied, one should investigate possible issues in the measurement that could be responsible for unreliable partial photoneutron cross sections:

- F_i^{exp} do not exceed the definitive upper limits;
- $\sigma^{\text{exp}}(\gamma, in)$ and corresponding F_i^{exp} are positive; and
- the differences between F_i^{exp} and F_i^{th} are not significant.

Considering the above criteria, the partial and total photoneutron reaction cross sections were evaluated using the method described above for the following nuclei; ^{59}Co , $^{63,65}\text{Cu}$, ^{75}As , $^{76,78,80,82}\text{Se}$, ^{89}Y , $^{90,91,92,94}\text{Zr}$, ^{98}Mo , ^{103}Rh , ^{115}In , $^{116,117,118,119,120,122,124}\text{Sn}$, ^{133}Cs , ^{138}Ba , ^{139}La , $^{140,142}\text{Ce}$, ^{141}Pr , $^{145,148}\text{Nd}$, ^{153}Eu , ^{159}Tb , ^{160}Gd , ^{165}Ho , ^{181}Ta , ^{186}W , $^{188,189,190,192}\text{Os}$, ^{197}Au , ^{208}Pb , and ^{209}Bi . The derived cross-section data were compiled into the international EXFOR database as “evaluated” data and are available to the nuclear data community.

B. Data Evaluation at KAERI

30 nuclides in the new IAEA photonuclear data library were evaluated by KAERI. The experimental data were mainly taken from the EXFOR database [9] and were used as reference for the evaluation work. The F_i -corrected data and the new measurements produced under this CRP, if available, were used preferentially over the other existing data. The evaluation was made with the nuclear reaction model code TALYS [75], combined with a newly developed automatic model parameter tuning system to facilitate the present evaluation work. In the automated parameter tuning procedure, the optical model potential parameters, the GDR parameters, the level density-related parameters, and pre-equilibrium model parameters were adjusted — most of them were within 20% from their default values, while sometimes they differed by up to 50% — to reproduce the experimental data or the F_i -corrected data. For the level densities and the γ -ray strength function, the constant temperature and Fermi gas models and the Brink-Axel Lorentzian, respectively, were used in the present evaluations. For the pre-equilibrium reaction, the exciton

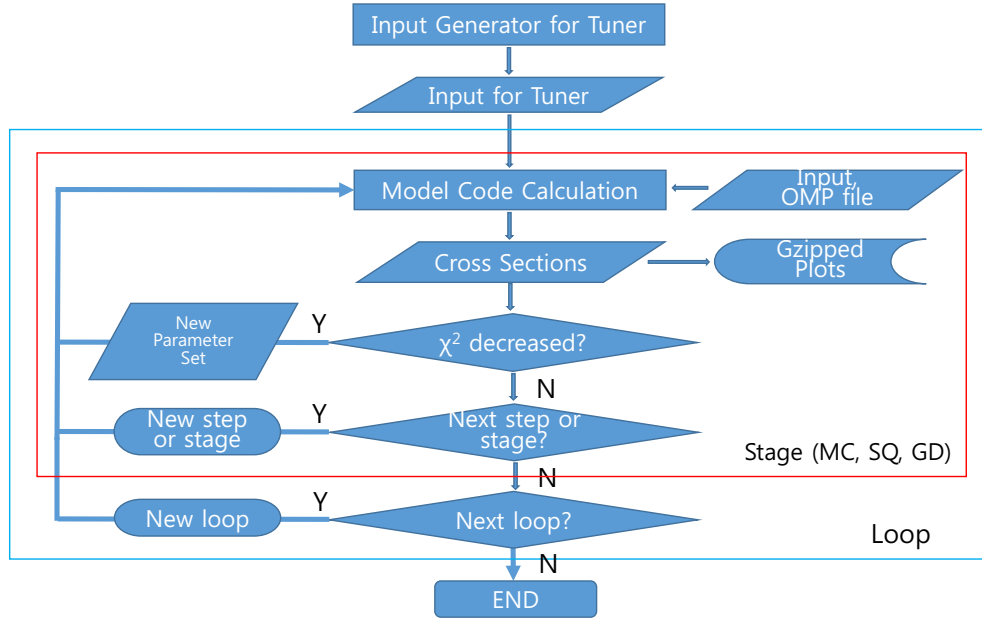


FIG. 13. (Color online) Schematic flow of the automated model parameter tuning at KAERI.

model was used. The list of adjusted model parameters is summarized below.

- **Optical model:** local parameters for neutron-induced reactions based on the global optical model parameters in the Koning-Delaroche form [87]. 19 model parameters are included.
- **Level density related parameters:** the level density parameter, the asymptotic level density, the damping parameter for the shell effects, the pairing correction, the temperature in the Gilbert-Cameron formula for the target and some residual nuclei. About 35 parameters in total for each evaluation are included.
- **GDR parameters:** the peak cross section, energy and width of GDR are adjusted. The γ -ray transmission coefficient calculated from GDR can be re-normalized by a scaling factor.
- **Pre-equilibrium model parameters:** in TALYS the strength of the exciton model is determined by the average squared matrix element M^2 . This is parameterized as

$$M^2 = \frac{C_1}{A^3} \left\{ 7.48C_2 + \frac{4.62 \times 10^5}{\left[\frac{E_{\text{tot}}}{n} + 10.7C_3 \right]^3} \right\}, \quad (41)$$

where n is the number of excitons, C_1 , C_2 , and C_3 are the adjustable parameters included. Since the two-component exciton model in TALYS distinguishes an exciton to be a neutron or a proton, the interaction strength between neutron (ν)

and proton (π) can be adjusted by introducing the scaling factors $M_{\pi\pi}^2 = R_{\pi\pi}M^2$, $M_{\nu\nu}^2 = R_{\nu\nu}M^2$, $M_{\pi\nu}^2 = R_{\pi\nu}M^2$, and $M_{\nu\pi}^2 = R_{\nu\pi}M^2$. These R factors are also adjusted.

Figure 13 shows the schematic flow of our automated model parameter tuning process, which consists of a pre-processor and a tuning tool. The preprocessor automatically generates necessary input files for the tuning tool by reading an output file of TALYS default run, as well as relevant EXFOR entries. The tuning tool creates a TALYS input file, invokes the code, compares the calculated results with the experimental data, computes χ^2 , and repeats this procedure until user-provided convergence criteria are satisfied by using the gradient search technique, grid search and/or random search. To reduce the computational time, multiple CPUs are employed simultaneously using MPI (Message Passing Interface). The whole process is terminated after a user-specified number of loops is completed. We can monitor the current status of fitting by checking automatically generated plots at each iteration. In the following we compare the KAERI evaluations with available experimental data for selected cases.

1. ^{94}Zr

The photonuclear data for ^{94}Zr in the previous IAEA 1999 library were evaluated by KAERI using the GUNF and GNASH code. That evaluation was based on the $(\gamma, 1nX)$, $(\gamma, 2nX)$, σ_{Sn} and σ_{xn} reaction cross sections of

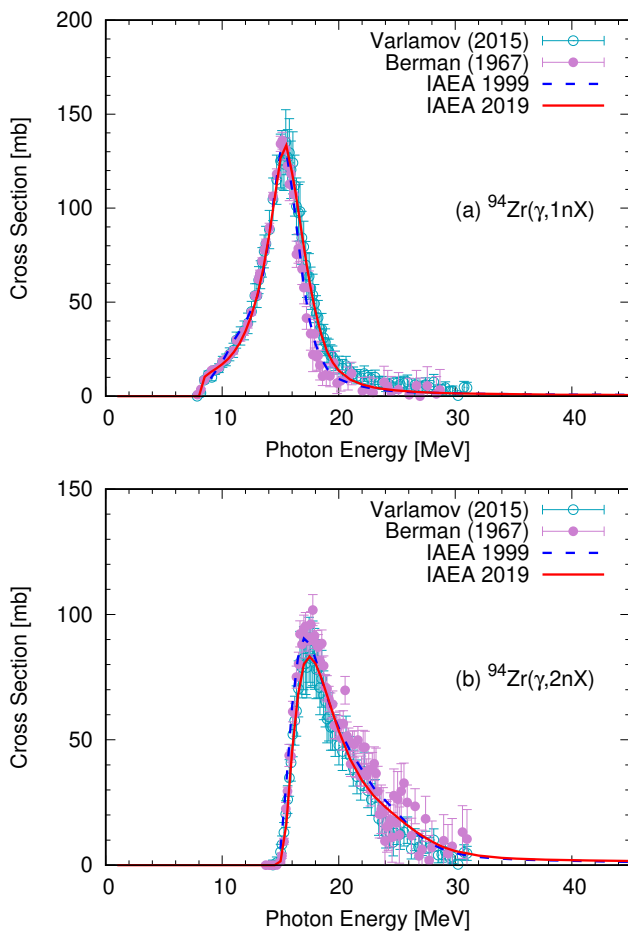


FIG. 14. (Color online) Comparison of evaluated and experimental cross sections for the γ -ray induced reaction on ^{94}Zr ; (a) $\sigma(\gamma, 1nX)$, (b) $\sigma(\gamma, 2nX)$.

Berman *et al.* [33]. The present evaluation was performed to reproduce the new $(\gamma, 1nX)$, $(\gamma, 2nX)$, $(\gamma, 3nX)$ and σ_{Sn} cross sections of Varlamov *et al.* [116] which are F_i -corrected ones. The $(\gamma, 1nX)$ cross sections of Varlamov are higher than those of Berman by 10% and the new $(\gamma, 2nX)$ cross sections of Varlamov are lower than those of Berman by 15% in the peak region.

The calculated photonuclear cross sections for ^{94}Zr are compared with the experimental and evaluated data [33, 113] in Figs. 14 and 15. As described above, the model parameters were adjusted to reproduce the evaluated experimental data of Varlamov *et al.*

2. ^{133}Cs

The photonuclear data for ^{133}Cs in the previous IAEA 1999 library were evaluated by KAERI using the GUNF and GNASH code. The previous evaluation was based on the $(\gamma, 1nX)$, $(\gamma, 2nX)$, σ_{Sn} and σ_{xn} reaction cross

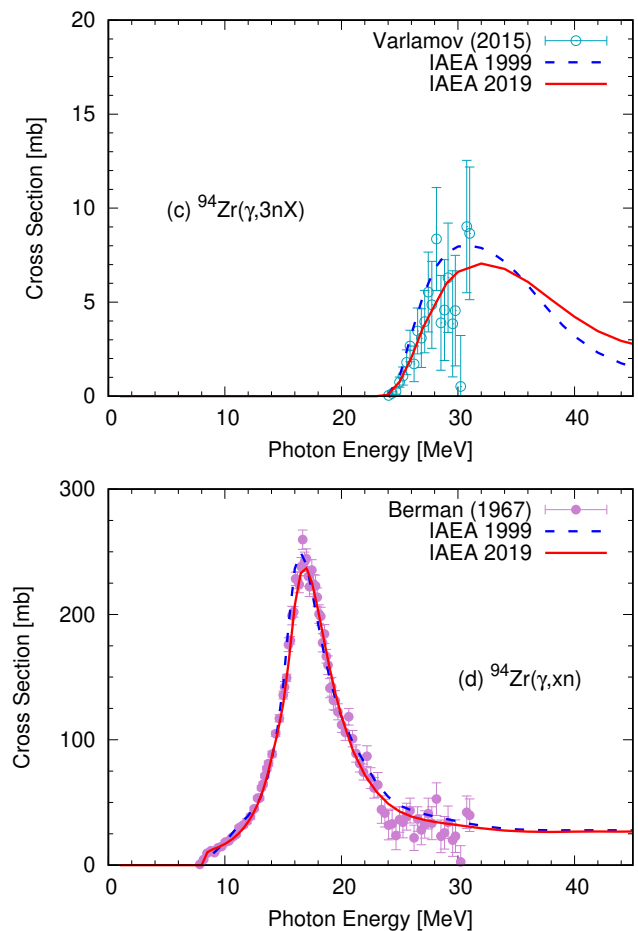


FIG. 15. (Color online) The same as the previous figure; (c) $\sigma(\gamma, 3nX)$, (d) σ_{xn} .

sections of Leprêtre *et al.* [124]. The present evaluation was performed to reproduce the new $(\gamma, 1nX)$, $(\gamma, 2nX)$, $(\gamma, 3nX)$ and σ_{Sn} cross sections of Varlamov *et al.* [116] which are F_i -corrected ones. The $(\gamma, 1nX)$ cross sections of Varlamov are similar to those of Leprêtre while the $(\gamma, 2nX)$ cross sections of Varlamov are much higher than those of Leprêtre by 30% in the peak region. The calculated photonuclear cross sections for ^{133}Cs are compared with the evaluated and experimental data [116, 124, 125] in Figs. 16 and 17.

3. ^{138}Ba

The photonuclear data for ^{138}Ba were not included in the previous IAEA 1999 library. The present evaluation was performed to reproduce the new $(\gamma, 1nX)$, $(\gamma, 2nX)$, $(\gamma, 3nX)$ and σ_{Sn} cross sections of Varlamov *et al.* [116] which are F_i -corrected ones. The calculated photonuclear cross sections for ^{138}Ba are compared with the experimental and evaluated data [116, 126] in Fig. 18.

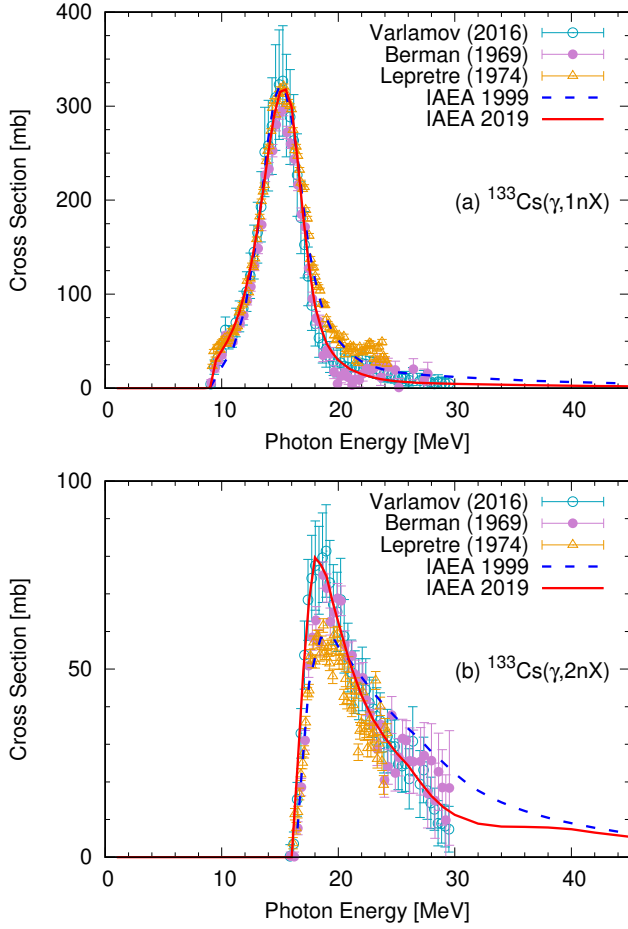


FIG. 16. (Color online) Comparison of evaluated and experimental cross sections for the γ -ray induced reaction on ^{133}Cs ; (a) $\sigma(\gamma, 1nX)$, (b) $\sigma(\gamma, 2nX)$.

4. ^{142}Ce

The photonuclear data for ^{142}Ce were not included in the previous IAEA 1999 library. The present evaluation was performed to reproduce the new $(\gamma, 1nX)$, $(\gamma, 2nX)$ and σ_{xn} cross sections of Lepretre *et al.* [127]. The calculated photonuclear cross sections for ^{142}Ce are compared with the experimental data [127] in Fig. 19.

C. Data Evaluation at IFIN-HH

Photoneutron reaction cross sections for ^{59}Co , ^{89}Y , ^{103}Rh , ^{159}Tb , ^{165}Ho , ^{169}Tm and ^{181}Ta have been recently measured in the GDR region using quasi-monochromatic LCS γ -ray beams at the NewSUBARU facility. The partial photoneutron cross sections, σ_{inX} with $i = 1$ to 4, were determined with the direct neutron multiplicity sorting method described in Sec. III B 4, which is based on a

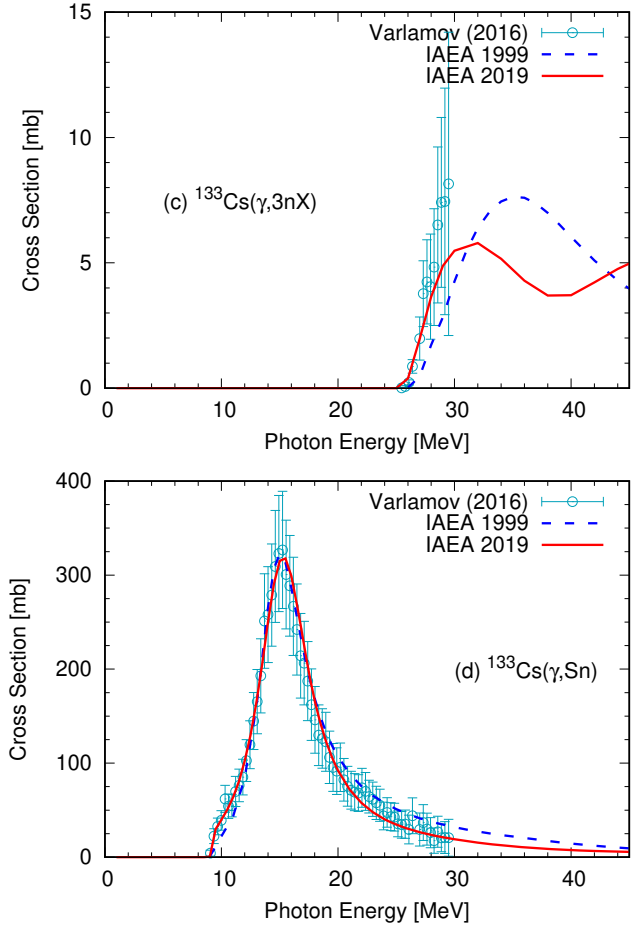


FIG. 17. (Color online) The same as the previous figure; (c) $\sigma(\gamma, 3nX)$, (d) σ_{Sn} .

slow response flat efficiency neutron detector. The average neutron emission energy corresponding to the total neutron-yield cross section was also provided using the ring-ratio method, as described in Sec. III B 2. The measured total σ_{Sn} and partial σ_{inX} ($i = 1 - 4$) cross sections were communicated among the CRP members prior to publication for use in the new evaluations. The final published results are not expected to differ substantially. We evaluated the newly measured NewSUBARU data for the seven nuclei listed above in an attempt to resolve the long-standing discrepancies between the Livermore and Saclay measurements detailed in Sec. V A.

The EMPIRE statistical model code [74] has been employed for the evaluations. For each evaluated nucleus, we have investigated which of the MLO1, MLO2, SLO and SMLO closed-forms for E1 γ -ray strength functions [58] implemented in EMPIRE reproduces best the experimental data. We also implemented in EMPIRE a third SLO component, which was used where necessary. EMPIRE also includes the quasi-deuteron contribution to describe the photo-absorption cross section above the GDR region,

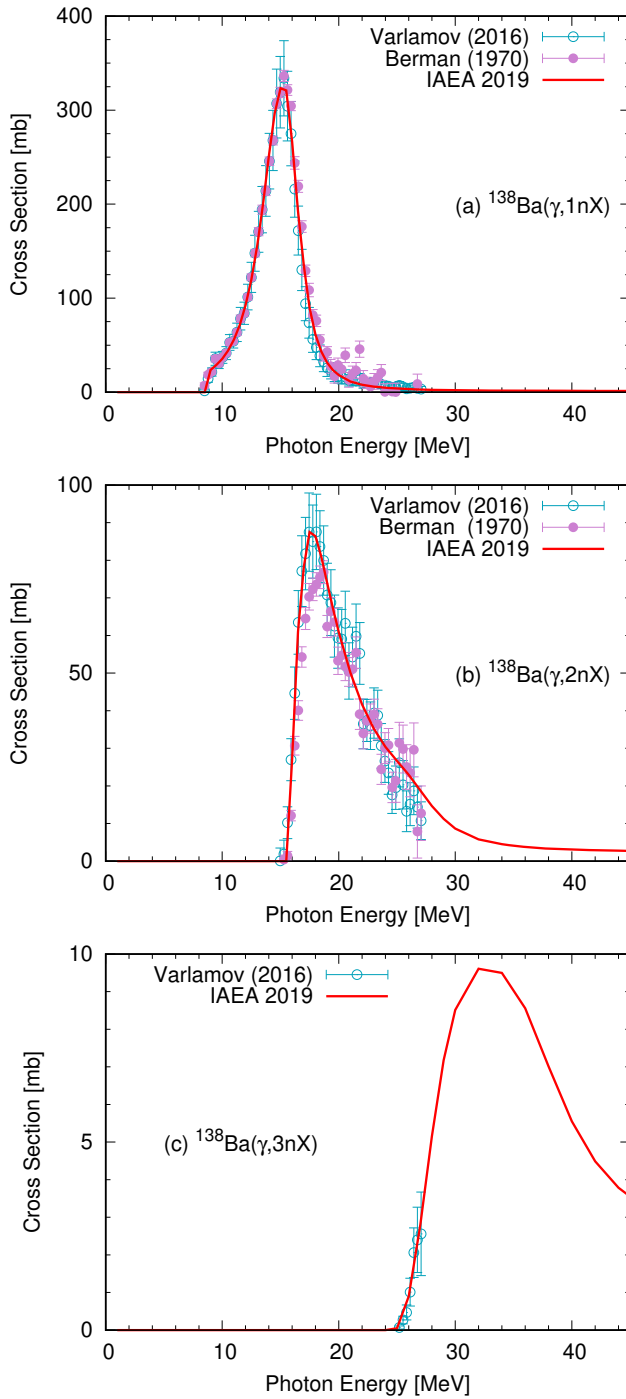


FIG. 18. (Color online) Comparison of evaluated and experimental cross sections for the γ -ray induced reaction on ^{138}Ba ; (a) $\sigma(\gamma, 1nX)$, (b) $\sigma(\gamma, 2nX)$, and (c) $\sigma(\gamma, 3nX)$.

up to 200 MeV.

In the NewSUBARU measurements, as also in the case of Saclay and Livermore experiments, the detection system is not sensitive to charged-particle emission from the

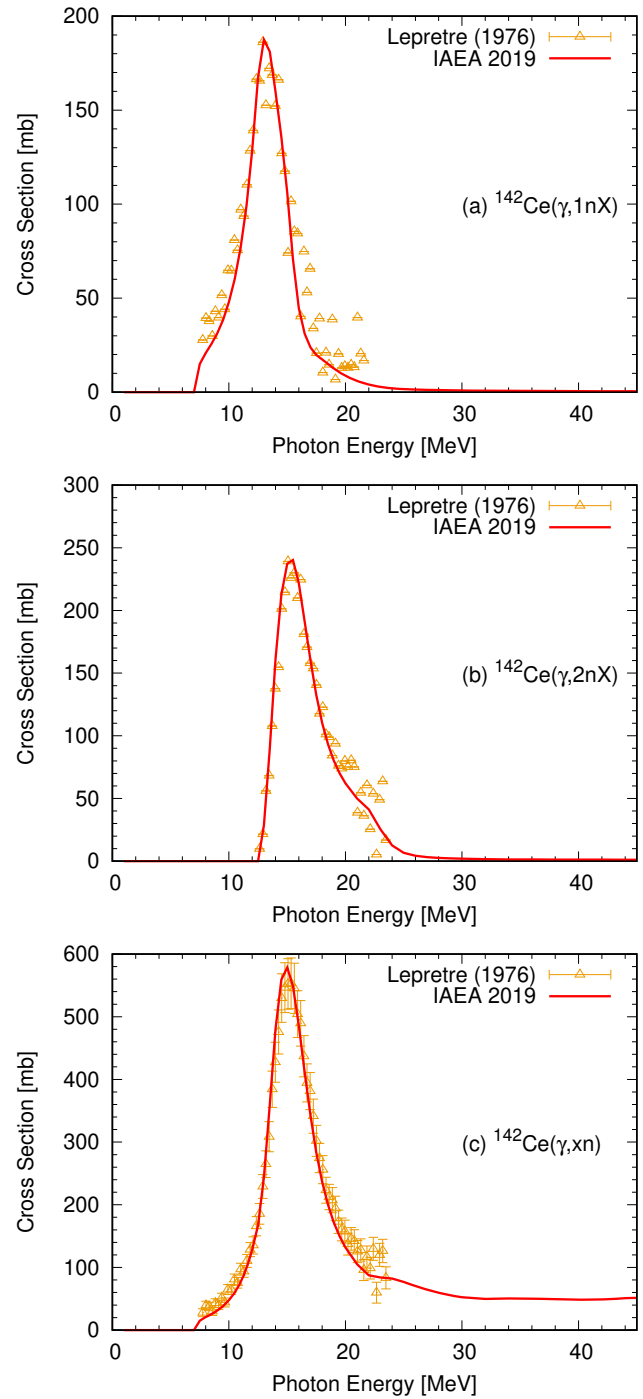


FIG. 19. (Color online) Comparison of evaluated and experimental cross sections for the γ -ray induced reaction on ^{142}Ce ; (a) $\sigma(\gamma, 1nX)$, (b) $\sigma(\gamma, 2nX)$, and (c) σ_{xn} .

target. The NewSUBARU experiments therefore, provide the sum of cross sections with i neutrons in the final state (γ, inX), where $i = 1$ to 4. The total photoneutron cross section σ_{Sn} was obtained as the sum $\sum_i \sigma_{inX}$. For

each nucleus, we adjusted the photo-absorption to reproduce the total photoneutron cross section σ_{Sn} . Finally we tuned the level density parameters, and the single particle level density parameters from the PCROSS pre-equilibrium model in order to reproduce the experimental σ_{inX} data with $i = 1 - 4$. The global Koning-Delaroche potentials [87] have been used to obtain the transmission coefficients for proton and neutron emission.

In order to compute complex σ_{inX} cross sections, one has to determine cross sections for different reaction chains which populate the same residual nucleus. For example, let us consider the ${}^A_{Z-1}X$ photo-disintegration to ${}^A_{Z-1}X$ nucleus, which can be produced by six different reaction chains (γ, nnp) , (γ, npn) , (γ, pnn) , (γ, nd) , (γ, dn) and (γ, t) . We compute the sum cross sections $(\gamma, nnp) + (\gamma, npn) + (\gamma, pnn)$ and $(\gamma, nd) + (\gamma, dn)$ separately, which contribute to the $(\gamma, 2nX)$ and $(\gamma, 1nX)$ reactions, respectively.

Because of limitations imposed by the available computing power and huge memory requirement, the EMPIRE statistical model code uses only one memory buffer per compound nucleus (CN) which is incremented by all reactions chains that populate the same CN. Thus, the code provides the population cross sections of each accessible CN disregarding the way it was produced and the emission cross section of each particle that can be emitted from the given CN. To obtain the cross sections for different reaction chains which populate the same residual nucleus, we developed a method of reconstructing all possible pathways by processing the statistical model code results. This treatment is particularly important for low mass nuclei which have a large contribution of charged-particle emission.

The method uses the additional information provided by EMPIRE concerning exclusive cross sections, which are defined as the double-differential cross sections / occupations for obtaining a nucleus in a certain state after having emitted a certain particle (proton, neutron, α -particle, ...). This cross section takes into account all emissions of that particular particle that lead to the given final state. However, the method does not provide exact results, because the information concerning spin and parity population distribution of the CN is taken into account only as an average. EMPIRE could easily avoid this approximation, but the calculation would grow by many orders of magnitude, an increase which would not be justified by an advantage of more precise splitting of inclusive spectra into their exclusive components. Also, obtaining cross sections for all pathways is possible up to a certain energy above which, due to the complexity of all the possible pathways, the information concerning exclusive population cross section alone is not anymore sufficient to solve very complex equation systems. The NewSUBARU measurements were performed at energies below this energy limit to take all possible pathways into account properly.

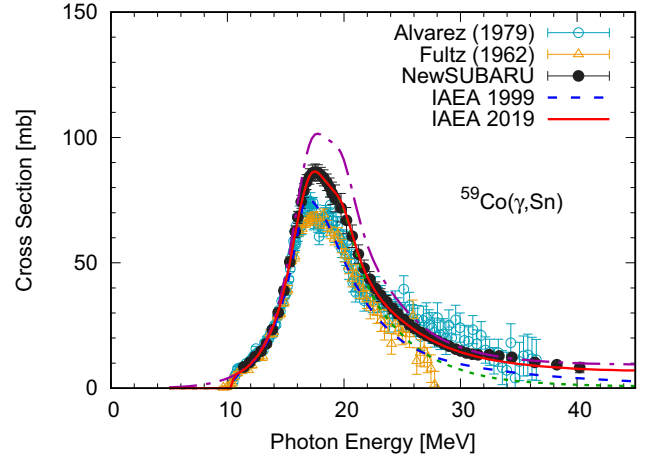


FIG. 20. (Color online) Comparison between experimental and evaluated (solid line) total cross sections σ_{Sn} for ${}^{59}\text{Co}$. Evaluated photo-absorption cross section including charged-particle channels is also shown by the dot-dashed line. The evaluated sum cross section $\sum_i \sigma_{in}$ ($i = 1 - 3$) of all channels without charged-particle emission is also shown by the dotted line.

1. ${}^{59}\text{Co}$

The previous ${}^{59}\text{Co}$ evaluation was performed by KAERI using the GNASH code [88, 89]. The KAERI evaluation adopted the $(\gamma, 1nX)$ and $(\gamma, 2nX)$ reaction cross sections of Alvarez *et al.* [121] and Fultz *et al.* [120], the $(\gamma, 3nX)$ reaction cross sections of Alvarez and the σ_{xn} data of Alvarez and Bazhanov *et al.* [128]. New σ_{Sn} , σ_{1nX} , σ_{2nX} and σ_{3nX} have been measured at NewSUBARU for ${}^{59}\text{Co}$ in the energy region between 10 and 40 MeV. The NewSUBARU data are shown by the filled circles in Figs. 20 and 21. The NewSUBARU (γ, Sn) cross section is about 20% higher than the Livermore data of Fultz and Alvarez and the peaks of the two GDR Lorentzians are not well separated. The MLO1 γ -ray strength function with two centroids reproduces best the (γ, Sn) cross sections. A third SLO has been introduced to reproduce the large width of the GDR region.

For this particular nucleus measured at NewSUBARU, the charged-particle emission contribution is stronger, as can be seen in Figs. 20 and 21. The EMPIRE photo-absorption and photoneutron cross sections are displayed in Fig. 20 by the dot-dashed and continuous lines, respectively, where the difference between the two cross sections is attributed to the charged-particle emission channel which is not accompanied by neutron emission. The sum cross section of all neutron-emission-only channels is shown with the dotted line. We observe that, because of the strong charged-particle contribution, the GDR parameters, namely the peak cross section and width, cannot be obtained by directly fitting the measured photoneutron cross section with a set of Lorentzians.

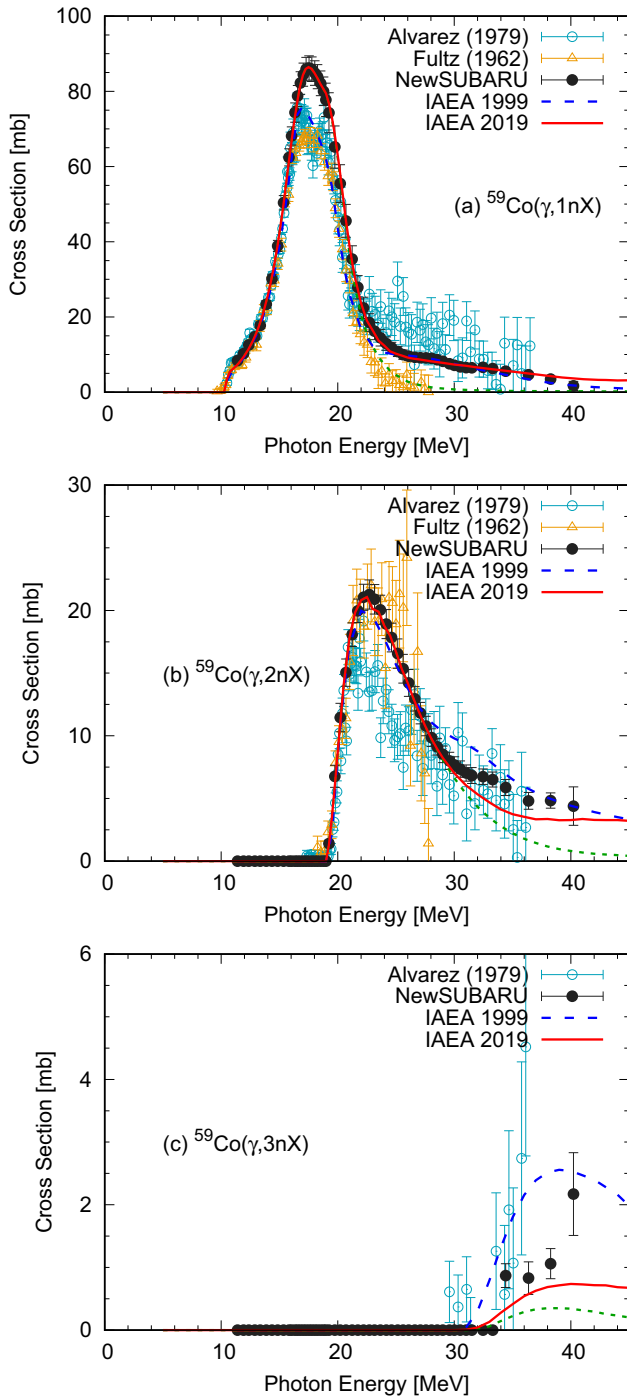


FIG. 21. (Color online) Comparison between experimental and evaluated (solid lines) partial cross sections for ^{59}Co ; (a) σ_{1nX} , (b) σ_{2nX} , and (c) σ_{3nX} . See Fig. 20 for the dotted lines.

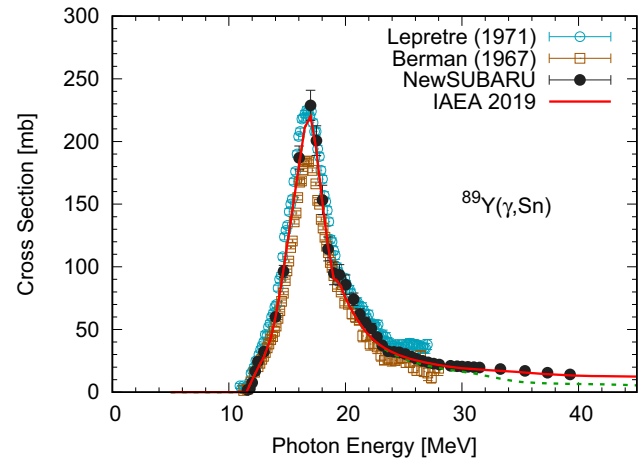


FIG. 22. (Color online) Comparison between evaluated (solid line) and experimental total cross sections σ_{Sn} for ^{89}Y . The evaluated sum cross section $\sum_i \sigma_{in}$ ($i = 1 - 3$) of all channels without charged-particle emission is also shown by the dotted line.

Figure 21 shows the comparison between (γ, inX) (solid lines) and (γ, in) (dotted lines) cross sections for $i = 1 - 3$, where the emission of accompanying charged particles increases with increasing energy.

2. ^{89}Y

^{89}Y was not included in the previous IAEA 1999 database. The $(\gamma, 1nX)$, $(\gamma, 2nX)$, (γ, Sn) reaction cross sections and the neutron yield σ_{xn} have been measured by Berman *et al.* [33] and by Leprêtre *et al.* [129] between 11 and 28 MeV excitation energy. New (γ, Sn) , $(\gamma, 1nX)$, $(\gamma, 2nX)$ and $(\gamma, 3nX)$ cross sections have been measured at NewSUBARU for ^{89}Y in the energy region between 11 and 40 MeV. The NewSUBARU and the Saclay cross sections of Leprêtre are in good agreement in the GDR peak energy region. The (γ, Sn) reaction cross section is best reproduced by using the SMLO model for the γ -ray strength function with two centroids. The evaluation is compared to the experimental data in Figs. 22 and 23.

3. ^{103}Rh

^{103}Rh was not included in the previous IAEA 1999 database. The $(\gamma, 1nX)$, $(\gamma, 2nX)$, (γ, Sn) reaction cross sections and the neutron yield σ_{xn} have been measured by Leprêtre *et al.* [124] between 9 and 26 MeV excitation energy. The (γ, Sn) cross section has been measured using bremsstrahlung beams by Parsons [130] and by Bogdankevich *et al.* [131], where Bogdankevich reported also the $(\gamma, 2nX)$ cross section. New (γ, Sn) , $(\gamma, 1nX)$, $(\gamma, 2nX)$ and $(\gamma, 3nX)$ cross sections were measured at

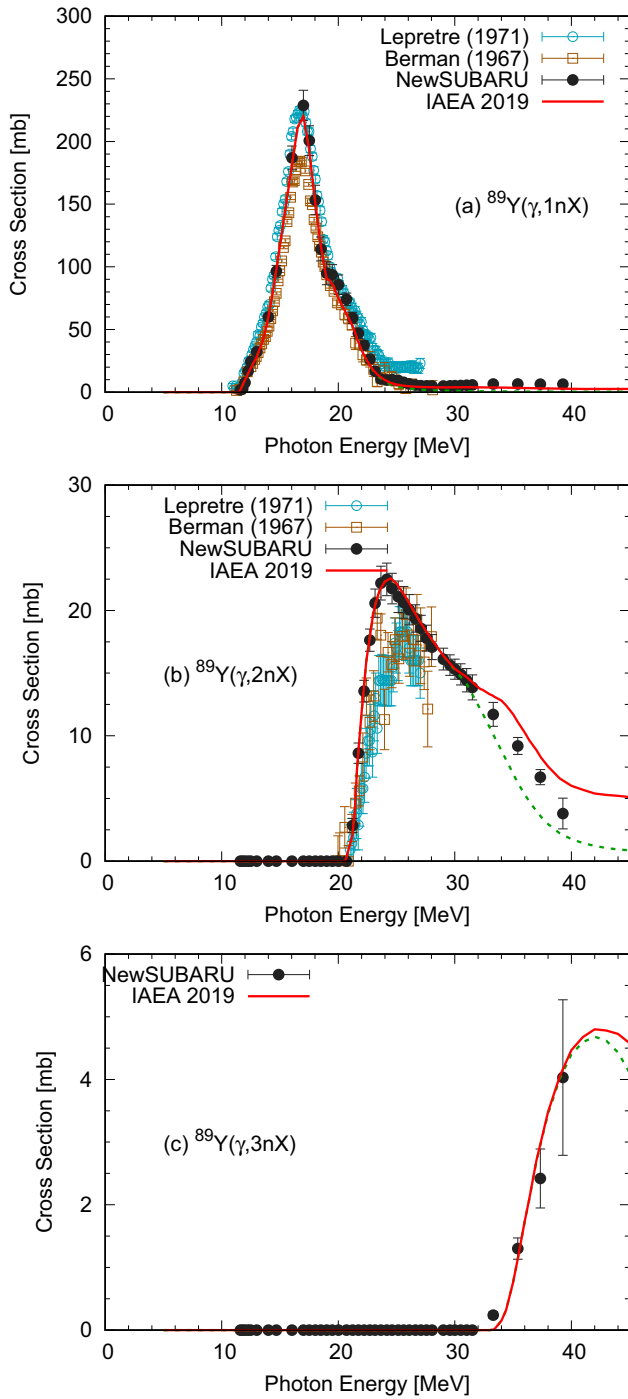


FIG. 23. (Color online) Comparison between experimental and evaluated (solid lines) partial cross sections for ^{89}Y : (a) σ_{1nX} , (b) σ_{2nX} , and (c) σ_{3nX} . See Fig. 22 for the dotted lines.

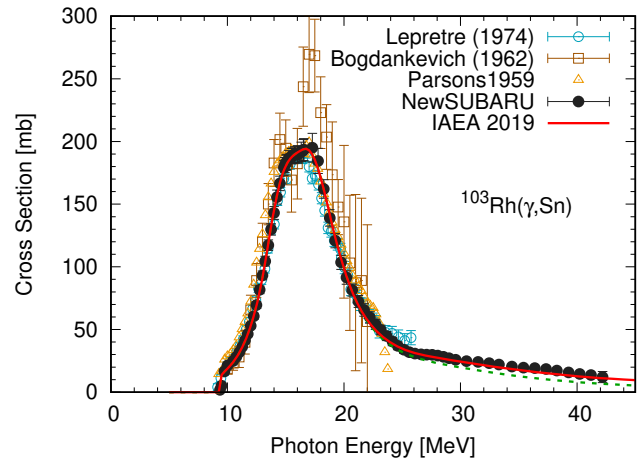


FIG. 24. (Color online) Comparison between experimental and evaluated (solid line) total cross sections σ_{Sn} for ^{103}Rh . The evaluated sum cross section $\sum_i \sigma_{in}$ ($i=1-3$) of all channels without charged-particle emission is also shown by the dotted line.

NewSUBARU for ^{103}Rh in the energy region between 9 and 42 MeV. The NewSUBARU and the Saclay cross sections of Lepretre are in overall good agreement in the entire investigated energy region. The MLO2 with two centroids reproduces best the (γ, Sn) cross sections. A third SLO has been introduced to reproduce the large width of the GDR peak region.

The evaluation is compared to the experimental data in Figs. 24 and 25. The comparison between the experimental and calculated average neutron emission energy corresponding to the neutron-yield cross section is displayed in Fig. 26. We note that the model parameters were adjusted to the experimental cross sections not to the average neutron energies.

4. ^{159}Tb

The previous ^{159}Tb evaluation has been performed by KAERI using the GNASH code. The $(\gamma, 1nX)$, $(\gamma, 2nX)$ and (γ, xn) cross sections have been measured both by Bramblett *et al.* [132] and by Bergère *et al.* [133]. Bergère also reported the $(\gamma, 3nX)$ and (γ, Sn) cross sections. The σ_{Sn} have been measured using bremsstrahlung beams by Bogdankevich *et al.* [131] and by Goryachev *et al.* [134]. The KAERI evaluation adopts the Bergère (γ, Sn) , $(\gamma, 1 \sim 3nX)$ cross sections. New (γ, Sn) , $(\gamma, 1nX)$, $(\gamma, 2nX)$, $(\gamma, 3nX)$ and $(\gamma, 4nX)$ cross sections were measured at NewSUBARU for ^{159}Tb in the energy region between 8 and 42 MeV. The NewSUBARU $(\gamma, 1nX)$ and $(\gamma, 3nX)$ cross sections are in agreement with the Saclay data of Bergère, while the new $(\gamma, 2nX)$ cross sections are in agreement with the Livermore data of Bramblett. The NewSUBARU (γ, Sn) reaction cross

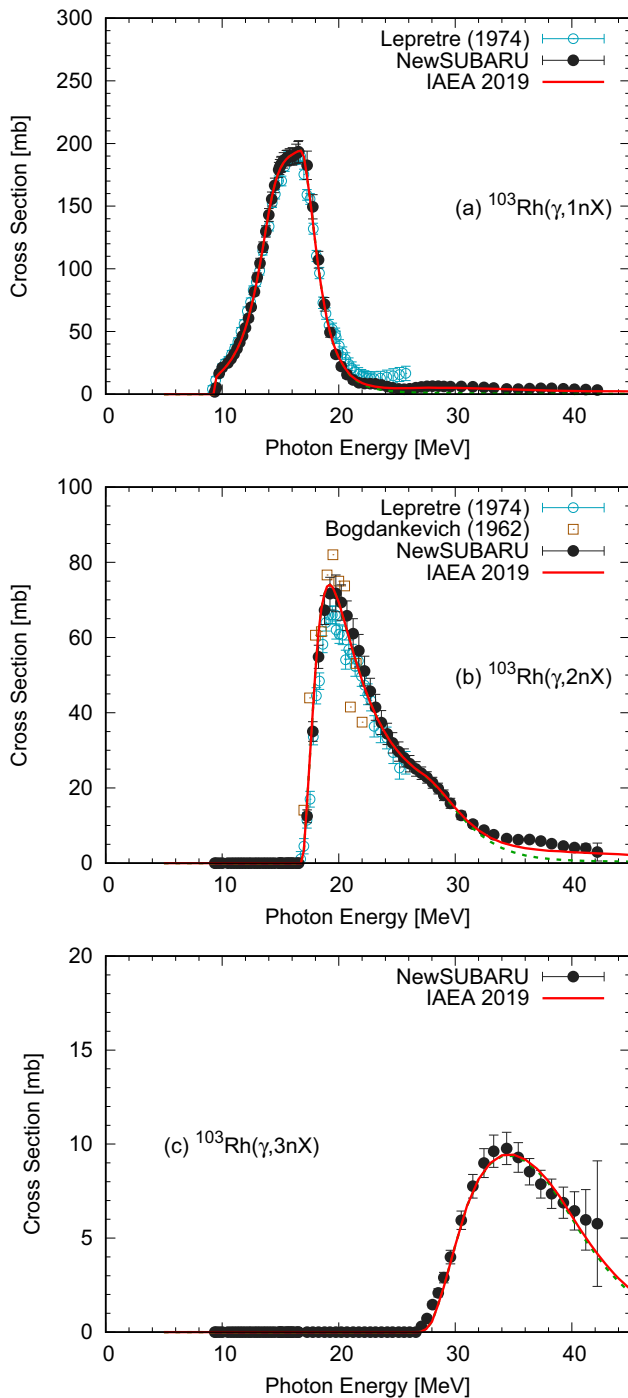


FIG. 25. (Color online) Comparison between experimental and evaluated (solid lines) partial cross sections for ^{103}Rh ; (a) σ_{1nX} , (b) σ_{2nX} , and (c) σ_{3nX} . See Fig. 24 for the dotted lines.

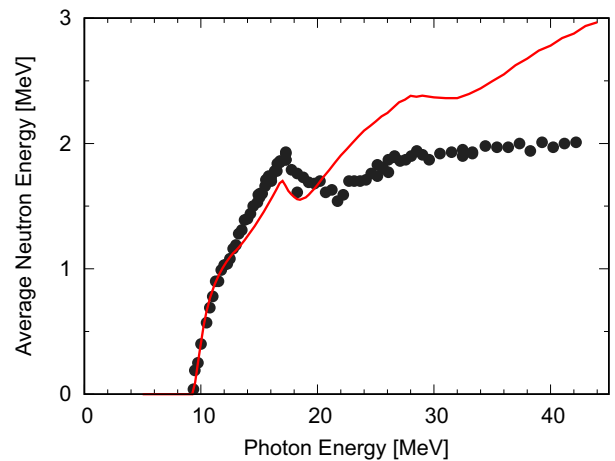


FIG. 26. (Color online) Comparison between evaluated and experimental values for average energy of neutrons emitted in photo-induced reactions on ^{103}Rh .

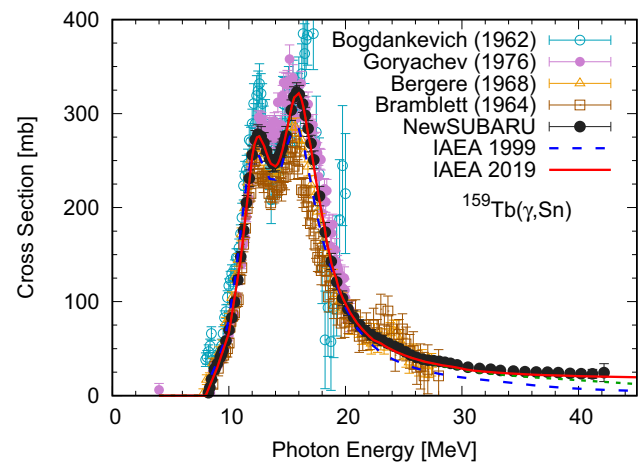


FIG. 27. (Color online) Comparison of experimental and evaluated total cross sections σ_{Sn} for ^{159}Tb . The solid line is the current evaluation, the dashed line is IAEA 1999 library [3]. The evaluated sum cross section $\sum_i \sigma_{in}$ ($i = 1 - 4$) without charged-particle emission is also shown by the dotted line.

section is best reproduced by using the SMLO model for the γ -ray strength function with two centroids.

The evaluation is compared to the experimental data in Fig. 27, and Figs. 28 and 29.

5. ^{165}Ho

The previous ^{165}Ho evaluation has been performed by KAERI using the GNASH code. The (γ, Sn) , $(\gamma, 1nX)$, $(\gamma, 2nX)$, $(\gamma, 3nX)$ cross sections have been measured both by Berman *et al.* [135] and by Bergère *et al.* [133].

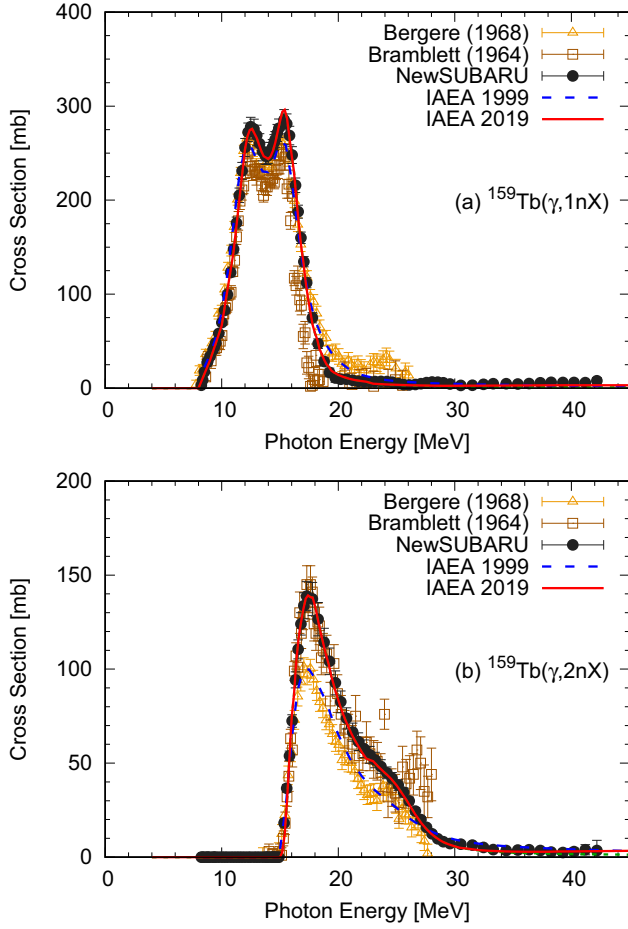


FIG. 28. (Color online) Comparison of experimental and evaluated (solid and dashed lines) partial cross sections for ^{159}Tb ; (a) σ_{1nX} , (b) σ_{2nX} . See Fig. 27 for the dotted lines.

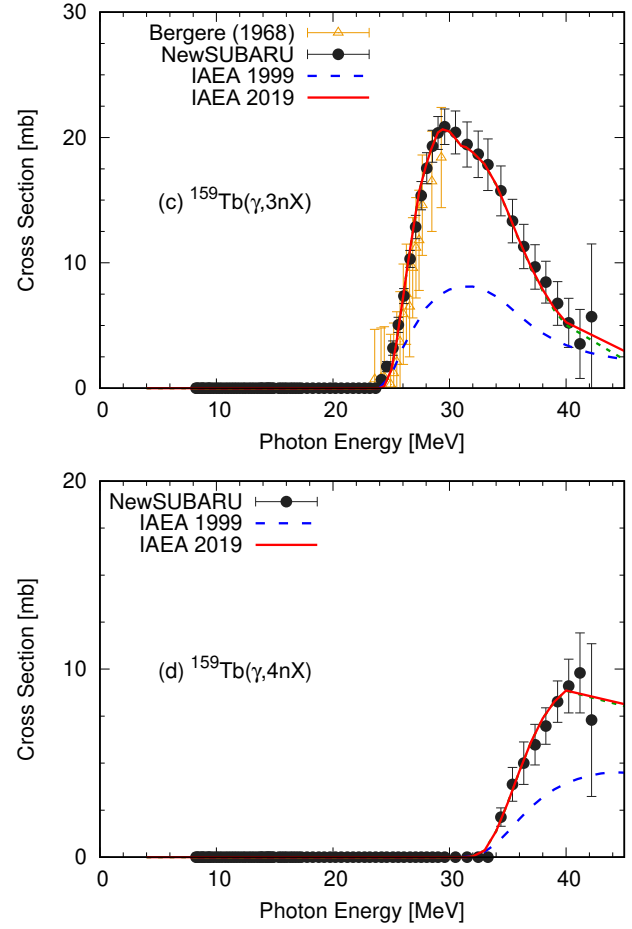


FIG. 29. (Color online) The same as the previous figure; (c) σ_{3nX} , and (d) σ_{4nX} .

The total nuclear photo-absorption cross section σ_{abs} has been measured by Gurevich *et al.* [136]. The σ_{Sn} and σ_{2nx} have also been reported by Goryachev *et al.* [134]. The KAERI evaluation adopted the Bergère cross sections. New (γ,Sn) , $(\gamma,1nX)$, $(\gamma,2nX)$, $(\gamma,3nX)$ and $(\gamma,4nX)$ cross sections were measured at NewSUBARU for ^{165}Ho in the energy region between 8 and 43 MeV. The NewSUBARU $(\gamma,1nX)$ cross sections are in better agreement with the Saclay measurement of Bergère while the new $(\gamma,2nX)$ cross sections are higher than both the Saclay and Livermore data of Berman by $\sim 20\%$ in the peak region. The NewSUBARU (γ,Sn) reaction cross section is best reproduced by using the SMLO model for the γ -ray strength function with two centroids.

The evaluation is compared to the experimental data in Fig. 30, and Figs. 31 and 32.

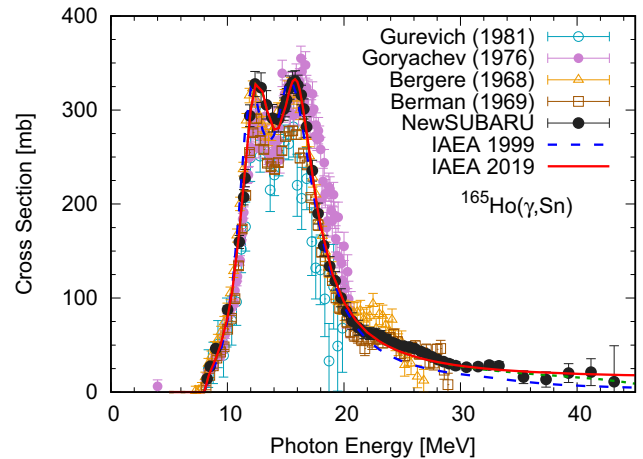


FIG. 30. (Color online) Comparison between experimental and evaluated (solid line) σ_{Sn} for ^{165}Ho . The evaluated sum cross section $\sum_i \sigma_{in}$ ($i = 1 - 4$) of all channels without charged-particle emission is also shown by the dotted lines.

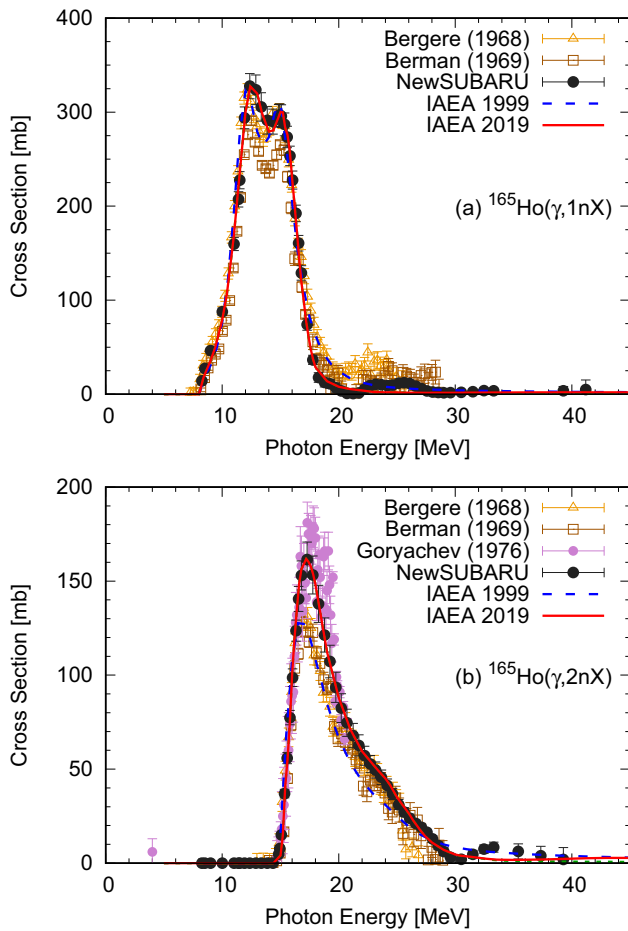


FIG. 31. (Color online) Comparison between experimental and evaluated (solid lines) partial cross sections for ^{165}Ho ; (a) σ_{1nX} , (b) σ_{2nX} . See Fig. 30 for the dotted lines.

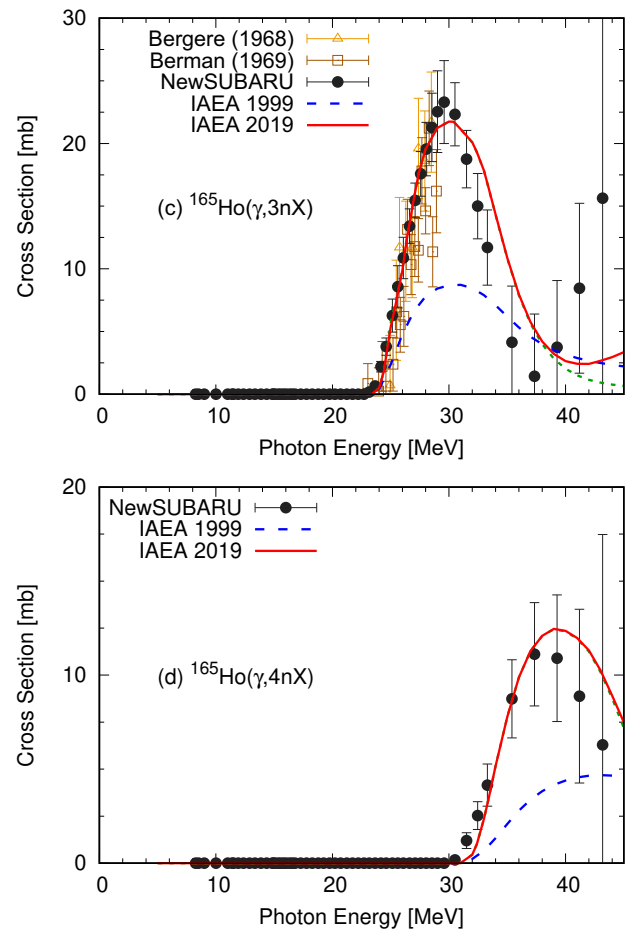


FIG. 32. (Color online) The same as the previous figure; (c) σ_{3nX} , and (d) σ_{4nX} . See Fig. 30 for the dotted lines.

6. ^{169}Tm

The (γ, Sn) , $(\gamma, 1nX)$, $(\gamma, 2nX)$, $(\gamma, 3nX)$ and $(\gamma, 4nX)$ cross sections have been measured at NewSUBARU for ^{169}Tm in the energy region between 8 and 40 MeV. No other data exist. The (γ, Sn) reaction cross section is best reproduced by using the SMLO model for the γ -ray strength function with two centroids. The evaluation is compared to the experimental data in Fig. 33 and Figs. 34, 35.

7. ^{181}Ta

The previous ^{181}Ta evaluation was taken from the JENDL Photonuclear Data Library [137]. The evaluation, produced with the ALICE-F [90] code, was based on existing (γ, Sn) , $(\gamma, 1nX)$ and $(\gamma, 2nX)$ cross sections measured by Bergère *et al.* [133] and by Bramblett *et al.* [138], where the latter Livermore cross sec-

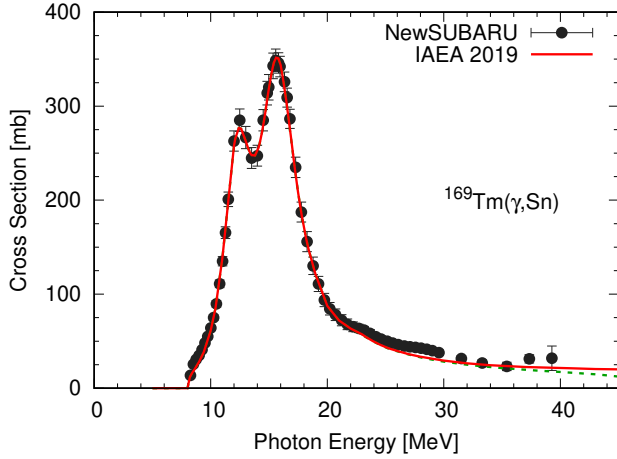


FIG. 33. (Color online) Comparison between evaluated and NewSUBARU σ_{Sn} for ^{169}Tm . The evaluated sum cross section $\sum_i \sigma_{in}$ ($i = 1 - 4$) of all channels without charged-particle emission is also shown by the dotted lines.

tions of Bramblett *et al.* were reconstructed based on a study performed at the Sao Paulo laboratory [34, 139]. The σ_{Sn} has been measured using bremsstrahlung beams by Belyaev *et al.* [140], Fuller *et al.* [141], Antropov *et al.* [142], Bogdankevich *et al.* [131] and Gurevich *et al.* [143]. Bogdankevich [131] reported also the $(\gamma, 2nX)$. Additional $(\gamma, 1nX)$ measurements have been performed using LCS γ -ray beams by Utsunomiya *et al.* [42] and Goko *et al.* [144]. New (γ, Sn) , $(\gamma, 1nX)$, $(\gamma, 2nX)$, $(\gamma, 3nX)$ and $(\gamma, 4nX)$ cross sections were measured at NewSUBARU for ^{181}Ta in the energy region between 12 and 41 MeV. The NewSUBARU $(\gamma, 1nX)$ cross sections are in better agreement with the Saclay measurement of Bergère, but have higher cross sections in the GDR peak region. The NewSUBARU results do not separate well the peaks of the two GDR Lorentzians, resembling better the Livermore data of Bramblett. The new $(\gamma, 2nX)$ cross sections are in better agreement with the Livermore data. The NewSUBARU (γ, Sn) reaction cross section is best reproduced by using the SMLO model for the γ -ray strength function with two centroids.

The evaluation is compared to the experimental data in Fig. 36 and Figs. 37, 38.

D. Data Evaluation at JAEA

The evaluations by JAEA are categorized into three groups (the complete list of nuclides is given in Appendix A). The first group includes ^3He , which was directly transferred from the JENDL photonuclear data file 2016 (JENDL/PD-2016) released in December 2017. Since no modification was made, the highest energy remains at 140 MeV.

The second group includes $^6,7\text{Li}$, ^{19}F and $^{40,44,48}\text{Ca}$.

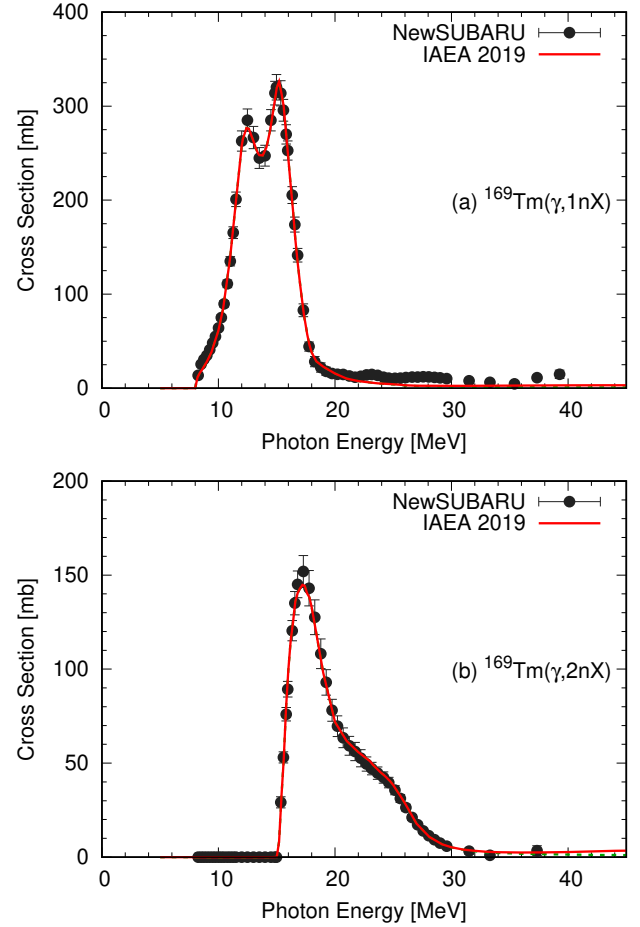


FIG. 34. (Color online) Comparison between evaluated and NewSUBARU partial cross sections for ^{169}Tm ; (a) σ_{1nX} , (b) σ_{2nX} . See Fig. 33 for the dotted lines.

The evaluated data were first produced by the method described by Murata *et al.* [145] using the ALICE-F code [90] for JENDL/PD-2016. In Murata's method, measured photo-absorption and $(\gamma, 1nX)$ reaction cross sections were reproduced by resonances and quasi-deuteron disintegration models. The other cross sections were calculated with ALICE-F, which provided cross section ratios of particle emission cross sections to the photo-absorption one. The energy-angle distributions of these nuclides were then replaced by CCONE [76] to obtain better prediction of particle emission. The highest energy is still 140 MeV. We denote this as JENDL(u) in Appendix A.

Finally, quite a few nuclides in the medium to heavy mass range in the IAEA 2019 library are upgrades of JENDL/PD-2016, in which newly available photo-induced reaction data were considered (photo-absorption cross section, photo-fission cross section, the average number of prompt and delayed fission neutrons, particle and photon emission cross sections and energy-angle

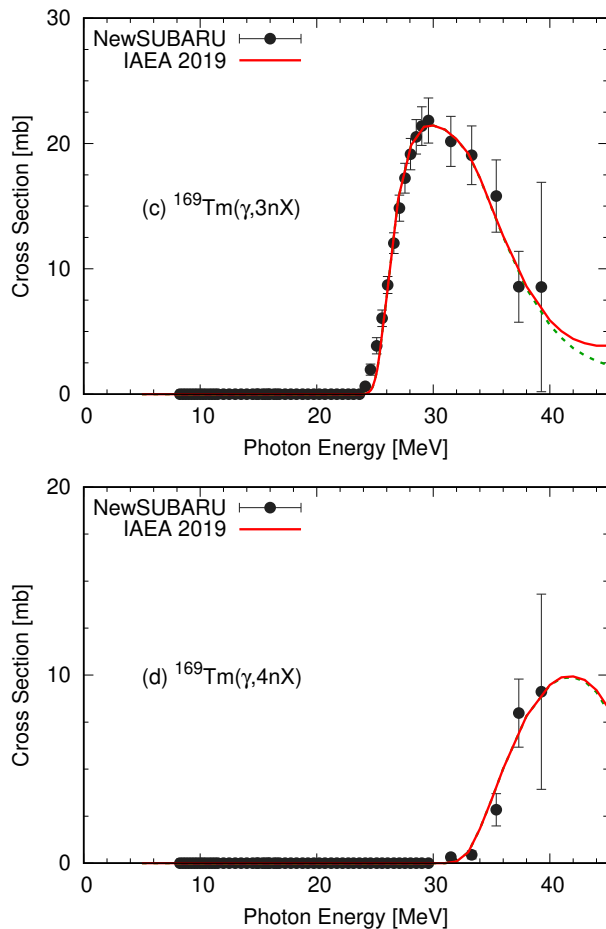


FIG. 35. (Color online) Comparison between evaluated and NewSUBARU partial cross sections for ^{169}Tm ; (c) σ_{3nX} , (d) σ_{4nX} . See Fig. 33 for the dotted lines.

distribution, residual cross sections, *etc.*) up to 200 MeV. The evaluation was made using the CCONE code [76]. In Appendix A, they are indicated as JAEA.

The CCONE code consists of the Hauser-Feshbach model for statistical decay and the two-component exciton model for pre-equilibrium decay. The deexcitation by γ -rays is expressed by the E1, M1 and E2 transitions. For the E1 radiation, the MLO1 model is basically utilized for the photon strength function. The default GDR parameters are taken from the RIPL-2 systematics [146]. Those parameters are modified so as to reproduce measured photo-absorption and/or photoneutron cross sections. For the M1 and E2 radiations the formulations of Kopecky and Uhl [66] are adopted. The discrete level information is taken from the RIPL-3 database [58]. The Gilbert-Cameron formula [147] is employed to describe the level density above the discrete levels, where the Fermi-gas model is replaced by the revised model of Mengoni and Nakajima [148]. In the statistical decay, we consider six particle-emission channels, namely neutron,

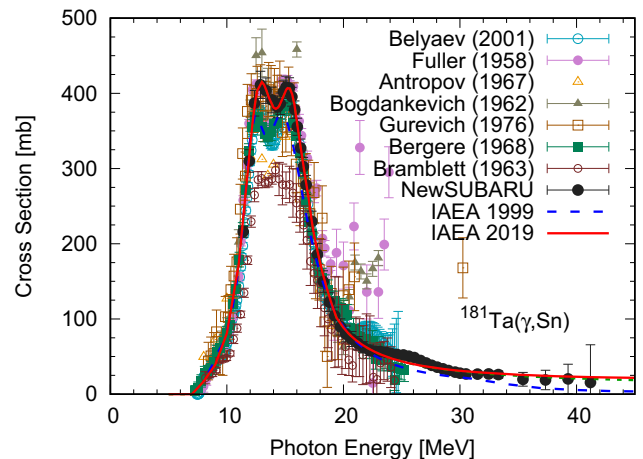


FIG. 36. (Color online) Comparison of experimental and evaluated σ_{Sn} for ^{181}Ta . The solid line is the current evaluation, the dashed line is IAEA 1999 library [3]. The evaluated sum cross section $\sum_i \sigma_{in}$ ($i = 1 - 4$) is shown by the dotted line.

proton, deuteron, triton, ^3He , and α -particle, as well as the γ -ray emission. For the particle emission, the optical model potential parameters of Koning-Delaroche [87] for neutrons and protons, Han *et al.* [107] for deuteron, folding potential generated from the neutron and proton potentials of Koning-Delaroche for triton, Xu *et al.* [108] for ^3He and Avrigeanu *et al.* [101] for α -particles are adopted. The photon incident energy region is extended up to 200 MeV. The lowest energy of 1 MeV is unchanged in the evaluations by CCONE. The quasi-deuteron disintegration model, which has an important contribution above the photon energy of 40 MeV, is also implemented. The Pauli-blocking function $P_b(E_\gamma)$ in the model is originally derived in the energy range of 20 to 140 MeV. The extension to lower and higher energies is necessary to produce nuclear data from 1 to 200 MeV. The function is extrapolated using Eq. (27).

The inclusive data form (MF/MT=6/5) in the ENDF-6 format is adopted in the JAEA evaluations. In this format, the $(\gamma, 1n)$ and $(\gamma, 2n)$ reaction cross sections are given as the production cross sections of residual nuclides with the mass numbers of $A - 1$ and $A - 2$ for a target with A . On the other hand, the experimental $(\gamma, 1nX)$ reaction cross section, which is the sum of $(\gamma, 1n)$, $(\gamma, 1np)$ and $(\gamma, 1n\alpha)$ cross sections and so on, cannot be separated into individual reaction cross sections. Thus, in the figures shown below, the present evaluated results for $(\gamma, 1n)$ and $(\gamma, 2n)$ reactions are used for comparisons with measured $(\gamma, 1nX)$ and $(\gamma, 2nX)$ reaction cross sections. Such a comparison is justified for nuclides with high atomic number, since photoneutron reactions with accompanying charged-particle emissions such as $(\gamma, 1np)$ and $(\gamma, 1n\alpha)$ are highly suppressed due to the large Coulomb barrier, hence the corresponding cross sections are very small.

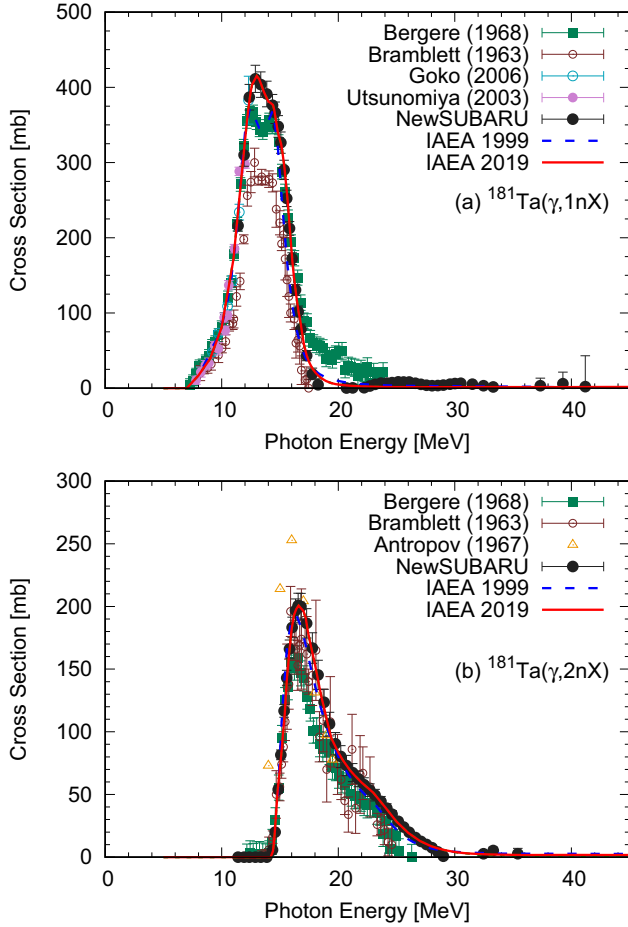


FIG. 37. (Color online) Comparison of evaluated and NewSUBARU partial cross sections for ^{181}Ta ; (a) σ_{1nX} , (b) σ_{2nX} . See Fig. 36 for the dotted lines.

In the case of nuclides with fission reaction channel, the average number of prompt neutrons per fission $\bar{\nu}_p$ is calculated using the linear functions with respect to photon energy fitted by Berman *et al.* [149] and Caldwell *et al.* [150], complemented with an exponential function [151] at the higher energies. The average number of delayed neutrons per fission $\bar{\nu}_d$ and the delayed neutron spectra, if available, are adopted from the neutron nuclear data in JENDL-4.0, if the compound nuclide is the same as the target in the photon-incident case.

1. ^{40}Ca

The ^{40}Ca file in the IAEA 1999 library [3] was produced by the GNASH code at LANL. In the updated library, we adopted the file from JENDL/PD-2016, but the energy-angular distributions were replaced by the CCONE calculation. We utilized available experimental data on photo-absorption, $(\gamma, 1nX)$, $(\gamma, 1pX)$, (γ, Sn) and (γ, xn) reac-

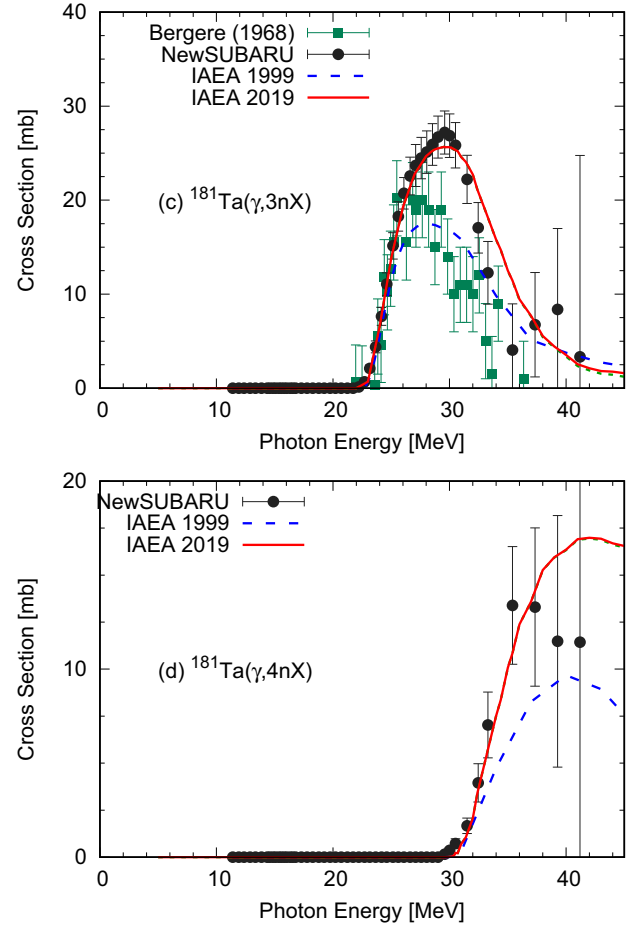


FIG. 38. (Color online) The same as the previous figure; (c) σ_{3nX} , (d) σ_{4nX} .

tions, to perform the evaluation. The photo-absorption cross section measured by Ahrens *et al.* [152] was reconstructed by 10 resonances, which also reproduced the $(\gamma, 1nX)$ reaction cross section of Veyssi re *et al.* [153]. The other particle emissions and residual production cross sections were calculated with ALICE-F.

Figure 39 shows comparison of the present evaluation with the previous evaluation [3] and the measured photo-absorption data, $(\gamma, 1nX)$ and $(\gamma, 1pX)$ cross sections. The measurements of photo-absorption cross sections are for the natural Ca target. The comparison of the present evaluation of ^{40}Ca with the measured data is physically meaningful, since the abundance of ^{40}Ca accounts for 96.9% of natural Ca. The present evaluation gives similar photo-absorption cross section as the IAEA 1999 library. The evaluated $(\gamma, 1p)$ reaction cross section is in good agreement with the experimental data of Goryachev *et al.* [154] and higher than the previous evaluation in the IAEA 1999 library by 45% at the peak cross section.

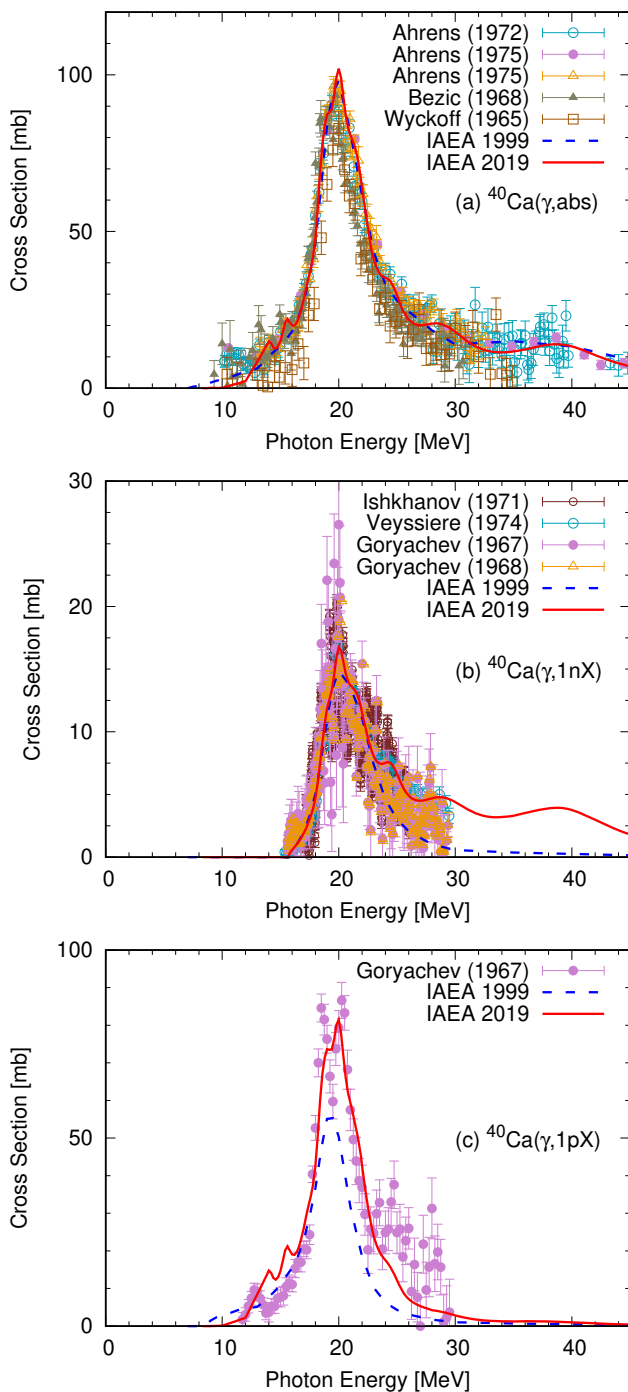


FIG. 39. (Color online) Comparisons of the present results with the evaluated (IAEA 1999) and experimental data for ^{40}Ca . The cross sections of (a) photo-absorption, (b) $(\gamma, 1nX)$ and (c) $(\gamma, 1pX)$ reactions are illustrated in each panel.

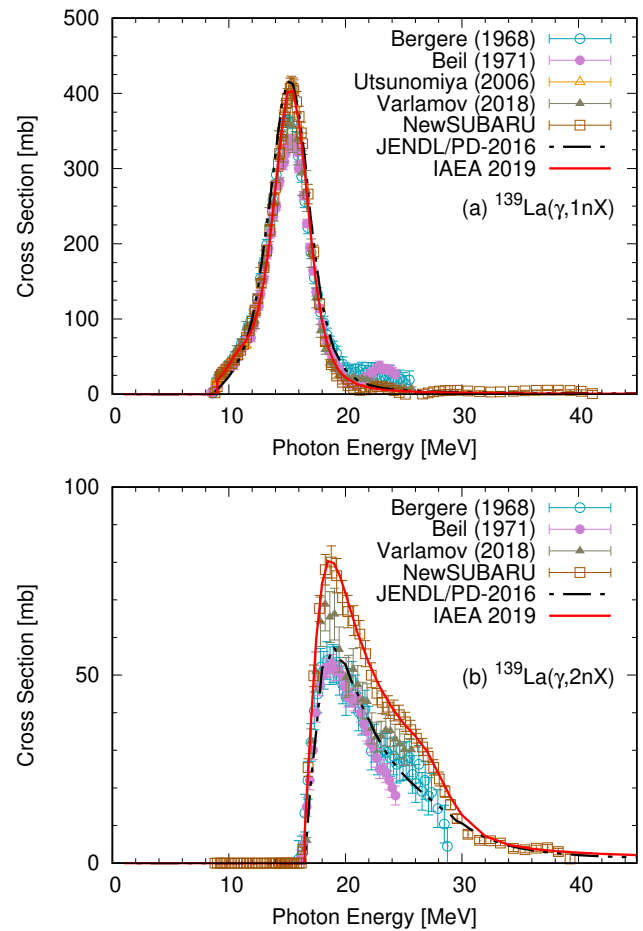


FIG. 40. (Color online) Comparisons of the present photoneutron cross sections with the evaluated (JENDL/PD-2016) and experimental data for ^{139}La ; (a) $(\gamma, 1nX)$, (b) $(\gamma, 2nX)$.

2. ^{139}La

The photonuclear data of ^{139}La were not included in the IAEA 1999 library. New measurements were carried out at the NewSUBARU facility and provided the cross sections of $(\gamma, 1nX)$, $(\gamma, 2nX)$, $(\gamma, 3nX)$ and $(\gamma, 4nX)$ reactions. The measured data are compared with previous measurements in Figs. 40 and 41. It is found that the $(\gamma, 1nX)$ and $(\gamma, 2nX)$ reaction cross sections of NewSUBARU are much higher than the other measurements. The present evaluation obtained with the SMLO model for the E1 photon strength function was based on the data measured at NewSUBARU.

The new evaluated cross sections for the $(\gamma, 1n)$, $(\gamma, 2n)$, $(\gamma, 3n)$ and $(\gamma, 4n)$ reactions are compared with JENDL/PD-2016 [155] in Figs. 40 and 41. The present evaluations are higher than JENDL/PD-2016, except for the $(\gamma, 1n)$ reaction cross section, where both evaluations agree since there is only one neutron emission below the threshold energy of $(\gamma, 2n)$ at 16.2 MeV. So, there is no

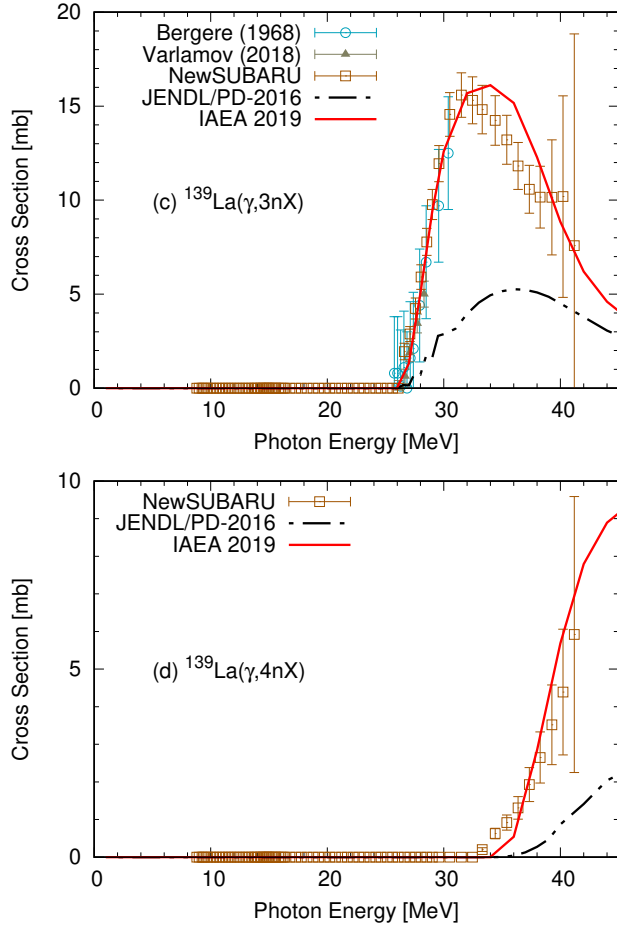


FIG. 41. (Color online) Comparisons of the present photoneutron cross sections with the evaluated (JENDL/PD-2016) and experimental data for ^{139}La ; (c) $(\gamma, 3nX)$, (d) $(\gamma, 4nX)$.

ambiguity in detecting 1n or 2n neutrons. The recent evaluated data of Varlamov *et al.* [156] have been obtained on the basis of the old data of Beil *et al.* [157] from Saclay and are also lower than those of NewSUBARU.

3. ^{239}Pu

The nuclear data of ^{239}Pu in the previous IAEA 1999 library was evaluated by IPPE. The energy region is limited to 20 MeV. The average number of prompt neutrons per fission $\bar{\nu}_p$ is given in the MF/MT=1/456 section (MF/MT combination defined in the ENDF-6 format), based on an estimation from the neutron nuclear data of ^{238}Pu . The MF/MT=3/5 format is especially used as the sum of $(\gamma, 1n)$ and $(\gamma, 1np)$ reaction cross sections. The current new library adopts the data from JENDL/PD-2016, but with the incident energy range extended to 200 MeV. The evaluation was made with CCONE. $\bar{\nu}_p$ and $\bar{\nu}_d$ were included in the evaluation.

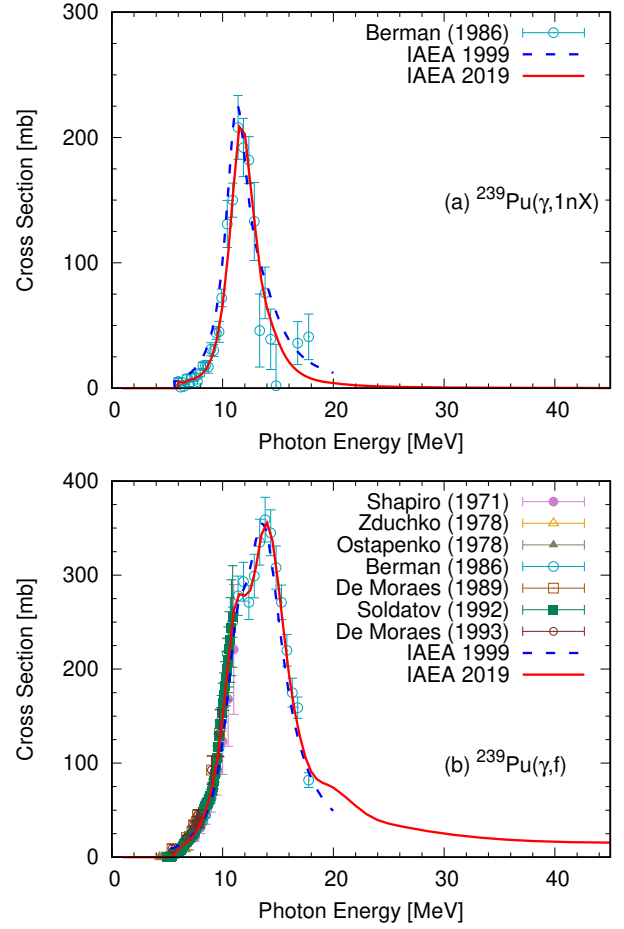


FIG. 42. (Color online) Comparisons of the present results with the evaluated (IAEA 1999) and experimental data for ^{239}Pu ; (a) $(\gamma, 1nX)$, (b) photo-fission.

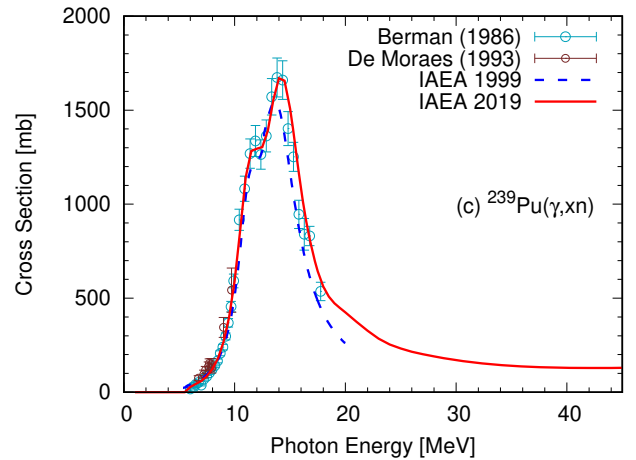


FIG. 43. (Color online) The same as the previous figure; (c) (γ, xn) .

Figures 42 and 43 show the comparison of the new evaluations with the previous IAEA 1999 library and the measured data for $(\gamma, 1nX)$, photo-fission and (γ, xn) reactions. The $(\gamma, 1nX)$ reaction cross section in the previous library shifts slightly to the lower energy region, relative to the measured data. The photo-fission cross sections are almost consistent with each other. The accuracy of this reaction cross section may be important for the development of inspection technology for nuclear fuel material. The small bump seen around 20 MeV comes from the contribution of third chance fission. This is not seen in the previous evaluation. The (γ, xn) reaction cross section has a contribution of fission neutrons as well as neutron emission cross sections multiplied by neutron multiplicity. The present (γ, xn) reaction cross section reproduces well the data of Berman *et al.* [149]. In contrast, the data in the previous library are lower than the measured data.

E. Data Evaluation at CIAE

The scheme used for evaluation of photonuclear data at CIAE/CNDC is shown in Fig. 44. First, we collect the available experimental data compiled in EXFOR for the following photonuclear reaction cross sections: (γ, abs) , (γ, xn) , $(\gamma, 1nX)$, $(\gamma, 2nX)$, $(\gamma, 3nX)$, $(\gamma, 1p)$, *etc.* We evaluate these experimental data and then use them to guide our theoretical calculations. The new photonuclear reaction calculation systems, GLUNF for the light nuclei (^9Be , $^{10,11}\text{B}$, ^{12}C , ^{14}N , and ^{16}O) and MEND-G for medium-heavy nuclei, were specially built for incident γ -ray energies E_γ below 200 MeV, in collaboration with Nankai University.

For the evaluation of an individual light element, a different reaction scheme is first constructed in GLUNF for each photonuclear reaction chain, since determination of all the competing particle-emission channels is most important in the light element case. Once the reaction chain is determined, the statistical theory of light-nucleus reactions [158, 159] is applied to calculate all the reaction cross sections. In the case of medium to heavy nuclei, MEND-G consists of the spherical optical model, and pre-equilibrium and equilibrium statistical models. Reaction channels with up to 18 particle emissions are considered.

We then analyze the available experimental data for (γ, abs) by fitting several empirical Lorentzian functions [11], known as SLO, MLO1, MLO2, MLO3, EGLO, GFL, and SMLO, and examine the best fit. Meanwhile, we also consider a semi-microscopic, relativistic QRPA model (RQRPA) [160], to further verify the GDR absorption. By adding the phenomenological QD model to the calculated GDR cross section, the photo-absorption cross section is extended to 200 MeV.

1. ^9Be

^9Be is a natural stable isotope of beryllium with an abundance of 100%. The evaluation in the previous IAEA

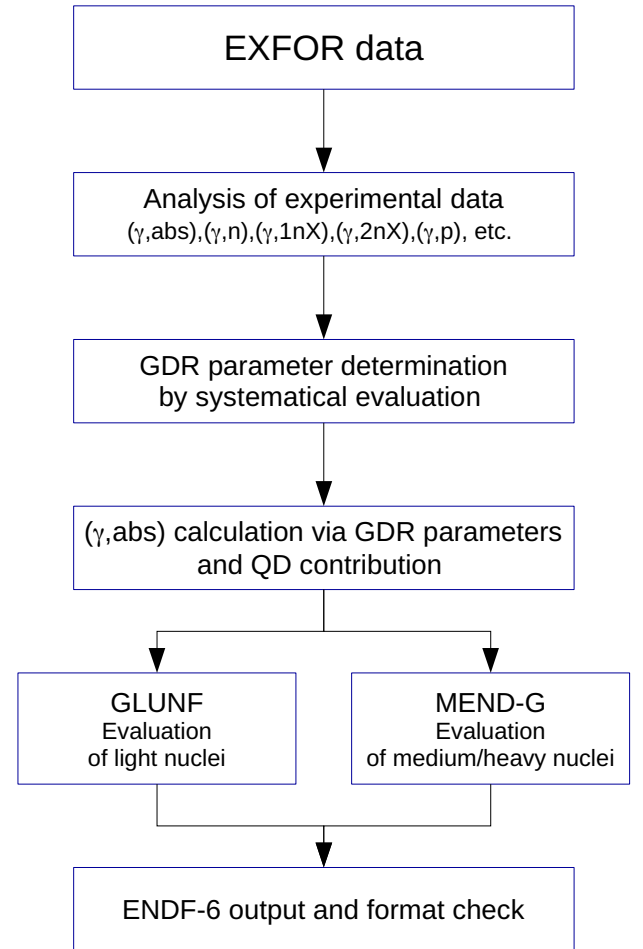


FIG. 44. Scheme of photonuclear data evaluation at CIAE/CNDC.

1999 library [3] was performed with the GLUNF code at CIAE/CNDC. In the GLUNF reaction scheme used in our present evaluation, the photo-absorption cross section σ_{abs} is split into the reaction channels of $(\gamma, 2n)$, $(\gamma, 2n\alpha)$, (γ, np) , $(\gamma, n2\alpha)$, (γ, nd) , (γ, nt) , $(\gamma, 2np)$, (γ, pd) , $(\gamma, d\alpha)$, (γ, p_0) , (γ, p_1) , (γ, d_0) , (γ, d_1) , (γ, t_0) , (γ, t_1) , $(\gamma, ^3\text{He}_0)$, and $(\gamma, ^3\text{He}_1)$, where (γ, p_0) stands for proton production leaving the residual in its ground state.

EXFOR contains experimental data on σ_{abs} , (γ, n) , $(\gamma, 2n)$, and the charged-particle emissions channels. The new NewSUBARU data for $(\gamma, n) + (\gamma, np)$ measured within this CRP are the main new experimental data used in our current update. Figure 45 compares our evaluated photo-absorption cross section with these experimental data. In the energy range 30–150 MeV, the evaluation of absorption cross section is based on the measurement of Ahrens *et al.* [152].

A new reaction chain scheme was developed in GLUNF

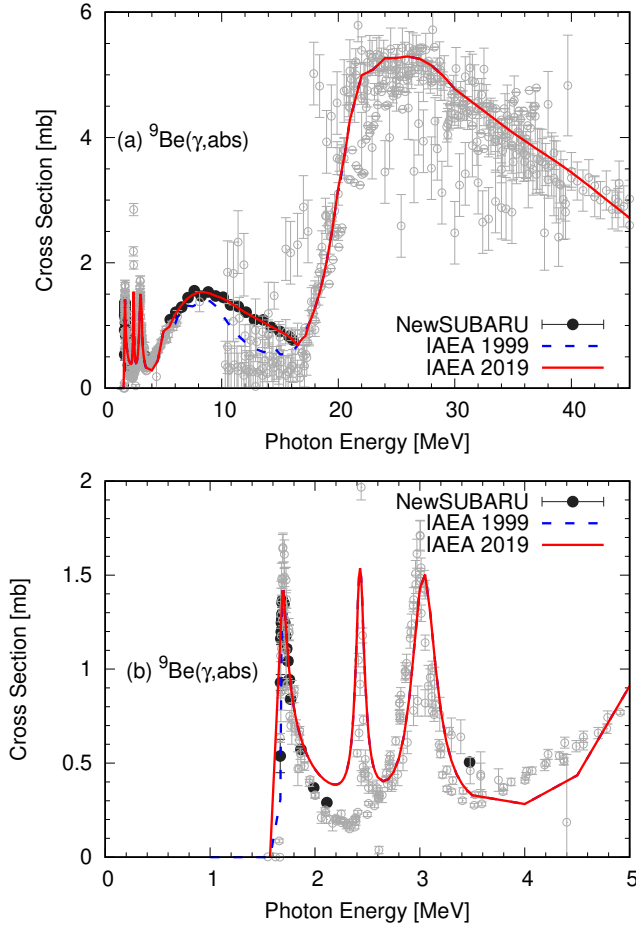


FIG. 45. (Color online) The evaluated photo-absorption cross section for ${}^9\text{Be}$. The bottom panel (b) is the zoom in the 1 – 5 MeV region. All the data are taken from EXFOR database, except for the NewSUBARU data shown by the filled circles.

for ${}^9\text{Be}$ to evaluate the photonuclear data below 150 MeV. The optical model parameters were adjusted to the experimental data adopted in our evaluation.

In Fig. 46, the calculated $(\gamma, n2\alpha)$ is compared with the available experimental data. The (γ, n) experimental data below 15 MeV were taken from EXFOR. Above 15 MeV, the measurement of Buchnea *et al.* [161] is compiled as the ${}^9\text{Be}(\gamma, n\alpha){}^4\text{He}$ reaction in EXFOR. The data shown by the filled circles are from the new measurement of Utsunomiya *et al.* reported in this CRP. The measurement of (γ, n) is represented by $(\gamma, n2\alpha)$. The results of $(\gamma, 2n)$ in our work are in good agreement with the experimental data of Utsunomiya as shown in Fig. 46. Whereas the ${}^9\text{Be}$ evaluation in the IAEA 1999 library, also performed at CIAE/CNDC with the GLUNF code, predicted almost zero cross section for $(\gamma, n2\alpha)$ at 30 MeV, our new evaluation has a more reasonable shape and natural extrapolation to the higher energy region.

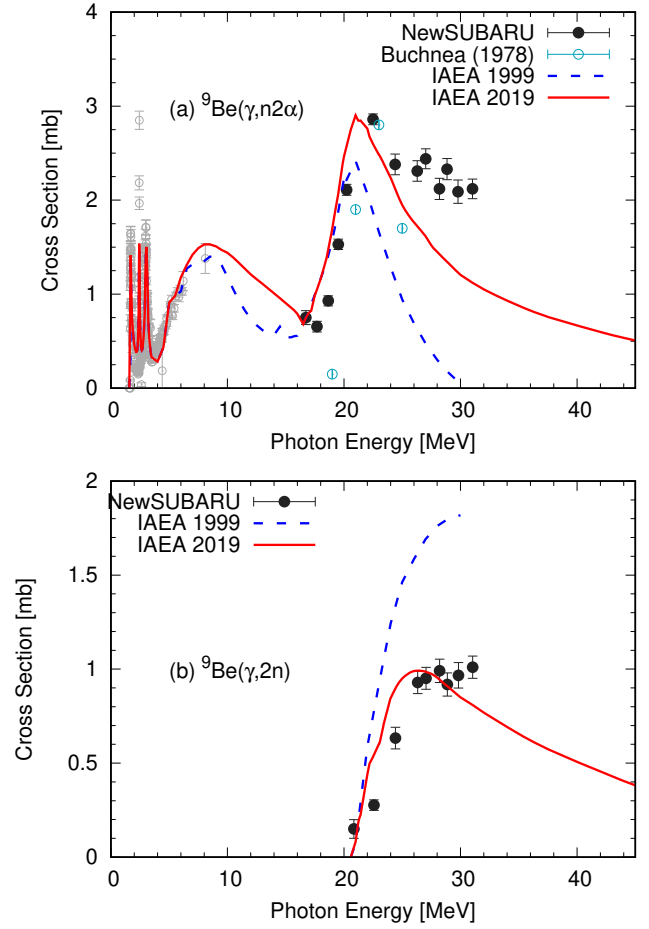


FIG. 46. (Color online) Comparison of the evaluated and experimental (a) ${}^9\text{Be}(\gamma, n2\alpha)$ and (b) $(\gamma, 2n)$ cross sections.

2. ${}^{14}\text{N}$

${}^{14}\text{N}$ is a natural stable isotope of nitrogen, which has the largest abundance of 99.64%. A new evaluation for the photo-induced reaction on ${}^{14}\text{N}$ has been performed at CIAE/CNDC. By setting up a reaction scheme in GLUNF, we calculated σ_{abs} , $(\gamma, 2n)$, $(\gamma, n3\alpha)$, (γ, np) , (γ, nd) , $(\gamma, n{}^3\text{He})$, $(\gamma, nd2\alpha)$, $(\gamma, nt2\alpha)$, $(\gamma, n2p)$, $(\gamma, n0)$, $(\gamma, 2\alpha)$, $(\gamma, 3\alpha)$, $(\gamma, 2p)$, $(\gamma, p\alpha)$, $(\gamma, t2\alpha)$, $(\gamma, d2\alpha)$, (γ, pd) , (γ, pt) , $(\gamma, d\alpha)$, as well as the proton, deuteron, triton, ${}^3\text{He}$, and α -particle emissions leaving the residuals in their discrete states.

The photo-absorption cross section is evaluated based on the measurement of Bezic *et al.* [162], which is shown in Fig. 47. The resonance at $E_\gamma = 9.172$ MeV is reported by Ajzenberg-Selove and Lauritsen [59]. We added this contribution by adopting the cross section in ENDF/B-VIII.0 [163]. Below 30 MeV, the $(\gamma, 1nX)$ cross section consists of the (γ, n) , (γ, np) and $(\gamma, np3\alpha)$ reactions in our calculation, and the comparison of evaluated and ex-

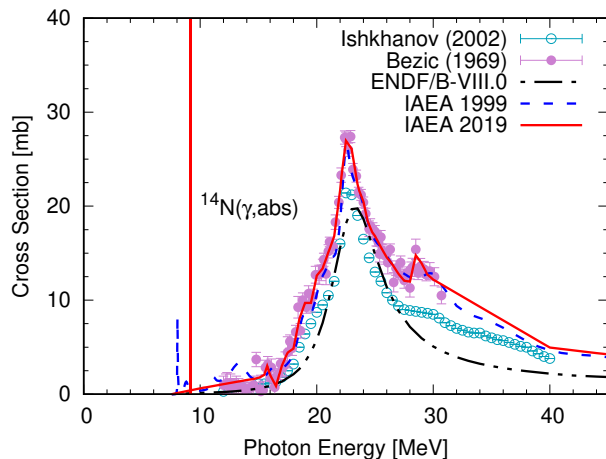


FIG. 47. (Color online) Comparison of evaluated and experimental data for the photo-absorption cross section of ^{14}N . The peak at 9.17 MeV, which is outside the plot, reaches 2.52 b.

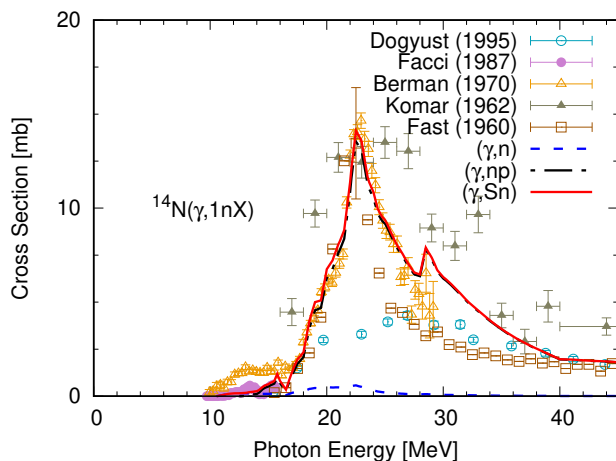


FIG. 48. (Color online) Comparison of evaluated and experimental data of neutron-production reaction cross sections, (γ, n) and (γ, np) for the photo-induced reaction on ^{14}N .

perimental data for $(\gamma, 1nX)$ is shown in Fig. 48. In Table I of Ref. [164], a relative yield of the (γ, np) reaction is reported to be 35%, while there are two channels given, 16% of (γ, n) and 11% of $(\gamma, 3\alpha pn)$, which produce one neutron. In addition, the residual of the (γ, p) reaction at higher energies, $^{13}\text{C}^*$ undergoes neutron-decay, with a branching ratio estimated to be 28%. Since the data of Komar are for (γ, np) only, we multiplied their data by $(28 + 35 + 16 + 11)/35 = 2.57$. The re-scaled data agree better with the other experimental data as well as with our model calculations, as shown in Fig. 48.

The excited $^{14}\text{N}^*$ emits a neutron and the residual nucleus ^{13}N continues to decay by emitting a proton. The first excited state in ^{13}N would be 100% proton-decay and

the probability of ^{13}N ground-state production would be very small. This is predicted by our code as we get a very small (γ, n_0) cross section. In addition, $(\gamma, 2n)$ is also included in GLUNF, although it has very small values, typically less than 0.01 mb.

3. ^{16}O

^{16}O is a natural stable isotope of oxygen with an abundance of 99.76%.

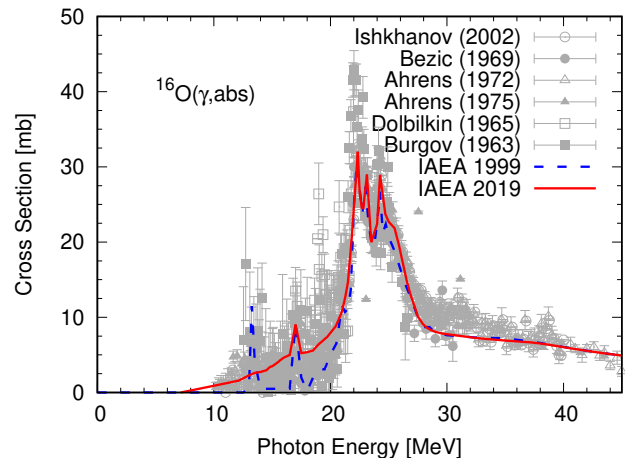


FIG. 49. (Color online) Comparison of theoretical results and experimental data for the photo-absorption cross section of ^{16}O .

We evaluated σ_{abs} and all of the exclusive cross sections for the $(\gamma, 2n)$, $(\gamma, n\alpha)$, $(\gamma, n^3\text{He})$, $(\gamma, 2np)$, $(\gamma, n_0) - (\gamma, n_6)$, $(\gamma, 3\alpha)$, $(\gamma, 2p)$, $(\gamma, p\alpha)$, $(\gamma, t^2\alpha)$, $(\gamma, d^2\alpha)$, (γ, pd) , (γ, pt) , and $(\gamma, d\alpha)$ reactions, as well as the discrete transitions for the proton, deuteron, triton, ^3He , and α -particle emission reactions. As examples, the evaluated photo-absorption cross section and $(\gamma, 2n)$ cross section are shown in Figs. 49 and 50, respectively. A notable difference between the evaluated photo-absorption cross sections of the previous IAEA 1999 library and the current one is seen in the energy range below 20 MeV. While the older IAEA 1999 data have a resonance near 13 MeV, the new data do not exhibit this peak. The new evaluation for the $(\gamma, 2n)$ reaction in Fig. 50 is significantly lower than the experimental data in the energy range above 40 MeV. We are unable to reproduce the data within our theoretical model framework, and we have no clear explanation for this. However, it should be noted that the previous evaluation, which employed a different statistical model, also shows the same tendency.

4. ^{27}Al

A new evaluation for ^{27}Al based on the MEND-G code calculation was adopted in the new photonuclear data

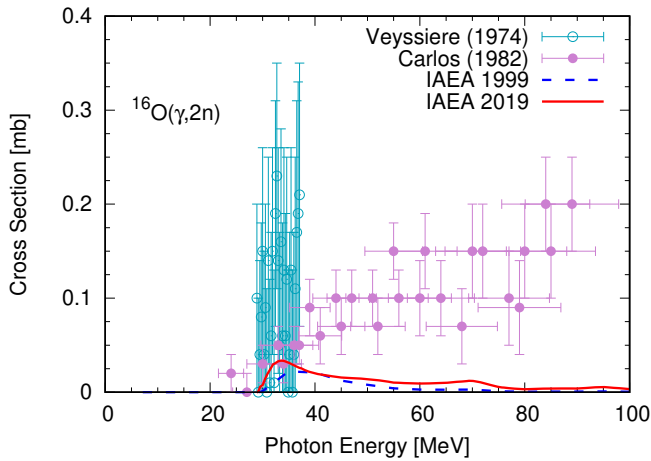


FIG. 50. (Color online) Comparison of evaluation and experimental data for the cross sections of $^{16}\text{O}(\gamma, 2n)^{14}\text{O}$.

library. The photo-absorption cross section is evaluated based on the experimental data of Ahrens *et al.* [152, 165], and the PSF calculation using the SMLO model. The evaluated photo-absorption cross section, shown in Fig. 51 (a), is very similar to the older evaluation up to the GDR peak of 20 MeV, then tends to be higher at higher energies and agrees better with the Ahrens data.

The evaluated reaction cross sections are compared with measurements of Fultz *et al.* [166], Veyssière *et al.* [153], and Shoda *et al.* [167] in Figs. 51, 52 (b)–(f). The new evaluations agree better with Fultz rather than Veyssière, although these data sets do not reveal critical discrepancies. In the case of the (γ, p) reaction shown in Fig. 52 (f), the experimental data of Shoda *et al.* are shifted to lower energies by 4 MeV to match the realistic reaction threshold energy.

5. ^{50}Cr

There are four natural stable isotopes for chromium, and all of them were evaluated by CIAE/CNDC in the former IAEA Photonuclear data library 1999. The new library includes an updated evaluation of ^{50}Cr - whose abundance is 4.35% - and updated $^{53,54}\text{Cr}$ data, while the ^{52}Cr was replaced by a new JAEA evaluation. The evaluation of the new ^{50}Cr data was performed with the MEND-G code. We used the EGLO model to produce the photo-absorption cross section of ^{50}Cr . Figure 53 compares the current evaluation with some available experimental data for the $(\gamma, 1n)$ reaction and the previous evaluation. The peak location of the new evaluation is slightly shifted toward the higher energy side, and the cross section drops rapidly above 25 MeV. This is more physical as the number of open channels increases at higher energies.

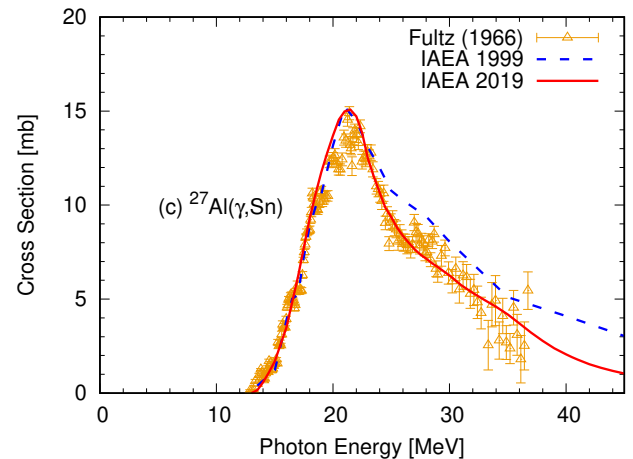
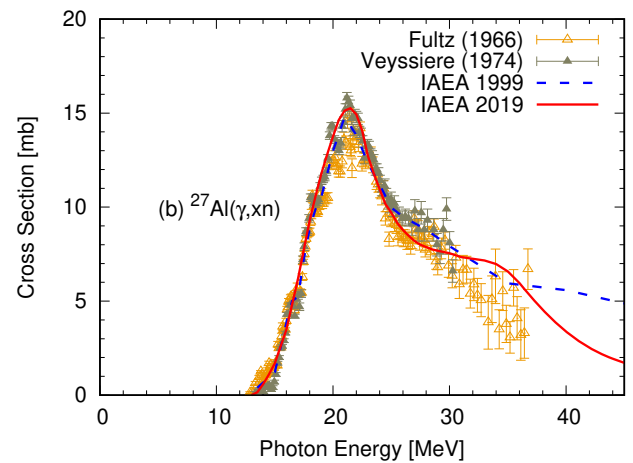
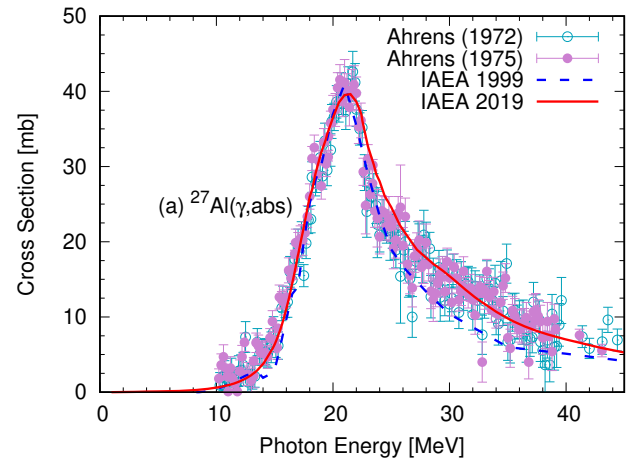


FIG. 51. (Color online) Comparison of evaluated and experimental data for ^{27}Al ; (a) photo-absorption, (b) σ_{xn} , (c) σ_{Sn} .

6. ^{90}Zr

Natural zirconium includes five stable isotopes, and CIAE/CNDC evaluated ^{90}Zr which has the largest abundance of 51.45%. The new evaluation was performed with

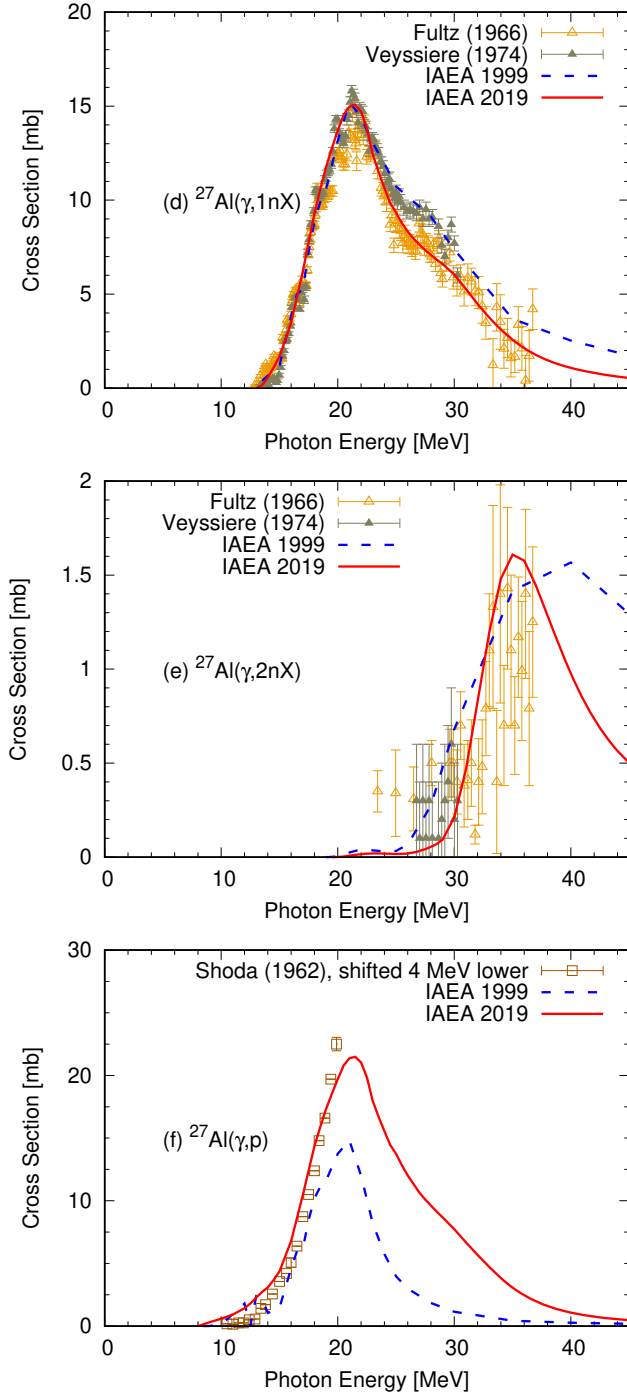


FIG. 52. (Color online) Comparison of evaluated and experimental data for ^{27}Al ; (d) $(\gamma, 1n) + (\gamma, 1np)$, (e) $(\gamma, 2n) + (\gamma, 2np)$, and (f) (γ, p) reactions. In the panel (f), the data of Shoda *et al.* [167] are lower shifted by 4 MeV.

the MEND-G code. The experimental data of Berman *et al.* [33] and Leprêtre *et al.* [129] are reported for ^{90}Zr . Varlamov *et al.* also published their evaluations

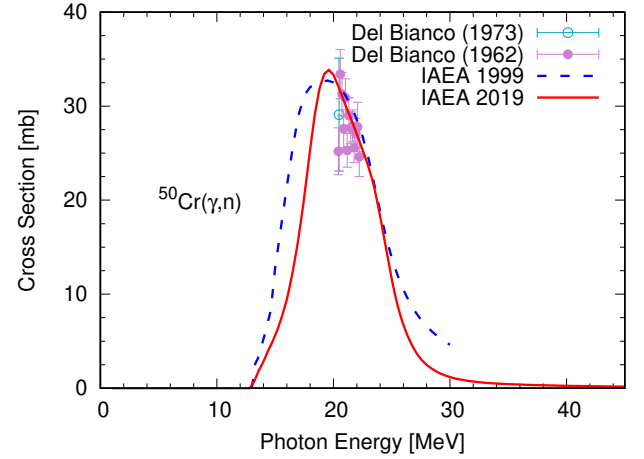


FIG. 53. (Color online) Comparison of evaluation and experimental data for $^{50}\text{Cr}(\gamma, 1n)$.

of the partial neutron production cross sections [118] in 2018. As has been mentioned in previous sections, the data of Berman (Livermore) and Leprêtre (Saclay) show some inconsistencies. In this case, after considering all the available data, we decided to adopt Berman's data for our evaluation. SMLO is used to produce the photo-absorption cross sections. The evaluated photo-absorption cross section and the related neutron emission cross sections are in good agreement with the experimental data of Berman as shown in Fig. 54. We concluded that the Leprêtre data for σ_{1nX} and σ_{2nX} could have a neutron mis-counting issue, hence the current evaluation does not follow the Leprêtre data.

7. ^{118}Sn

There are 10 stable isotopes in natural tin. The IAEA photonuclear data library adopted ^{120}Sn from JENDL/PD-2016 [155], which has the largest abundance of 32.6%. CIAE/CNDC undertook the evaluation of ^{118}Sn , whose abundance is the second largest of 24.22%. The experimental data of Fultz *et al.* [168] and Leprêtre [124] reported in 1960–1970s provide overall excitation functions of σ_{xn} and σ_{inX} for $i = 1, 2$ and 3. On the other hand, new neutron emission data by Utsunomiya *et al.* [169] were measured in 2011. Our evaluation strictly adopts the new data by Utsunomiya, which are compared with the other experimental data as well as the corrected data by Varlamov *et al.* in Figs. 55 and 56.

VI. CONTENTS OF THE LIBRARY

The new IAEA photonuclear data library contains photo-induced reaction data for 219 nuclides for photon

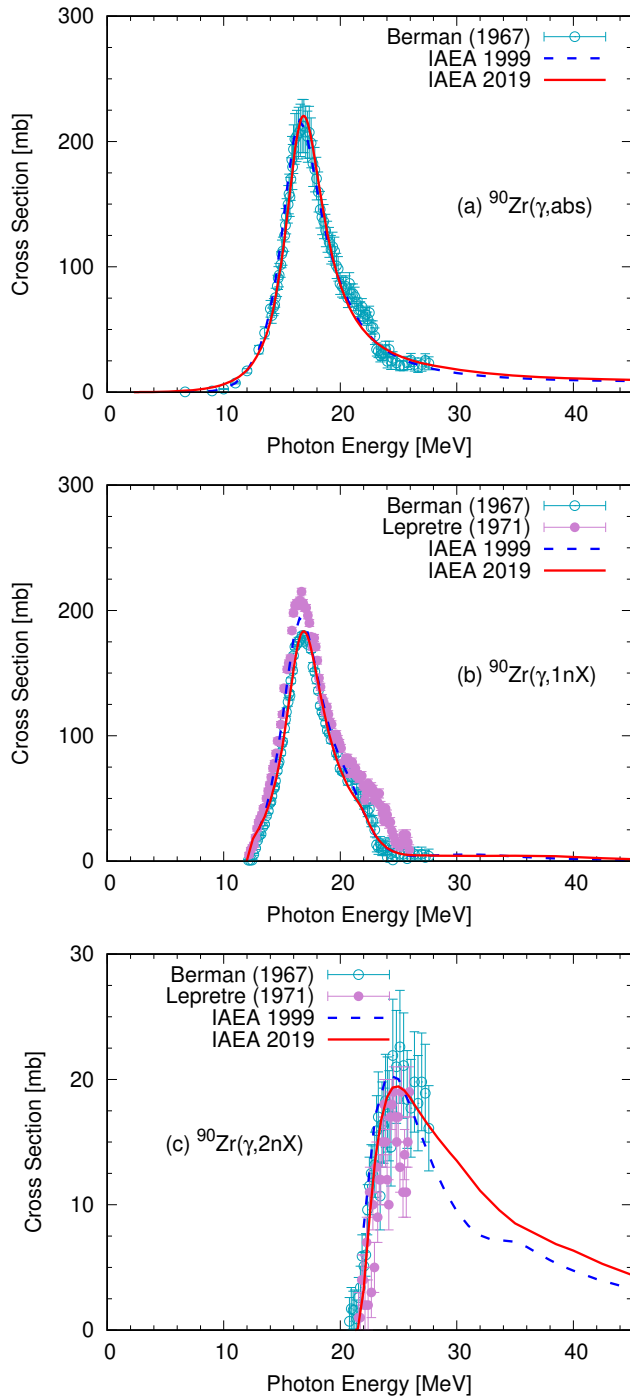


FIG. 54. (Color online) Comparison of evaluated and experimental data for ^{90}Zr ; (a) photo-absorption, (b) $(\gamma, 1n) + (\gamma, 1np)$, and (c) $(\gamma, 2n) + (\gamma, 2np)$ cross sections.

energies up to 200 MeV (some files go up to 140 MeV). The energy and mass ranges are expanded compared to the previous 1999 version [3] which included 164 nuclides with the energies mostly up to 140 MeV. Here we call

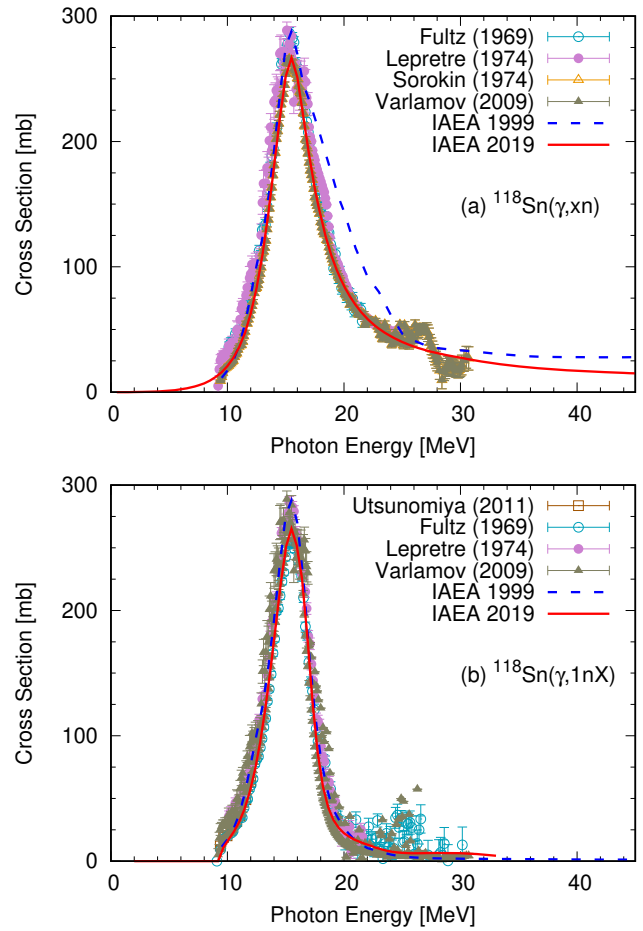


FIG. 55. (Color online) Comparison of evaluated and experimental data for ^{118}Sn ; (a) (γ, xn) , (b) $(\gamma, 1nX)$.

the old and new library IAEA 1999 and IAEA 2019, respectively. In Appendix A we summarize the sources of evaluated data for each isotope in both IAEA 1999 and IAEA 2019.

Many of the isotope evaluations included in IAEA 1999 originated from different evaluated nuclear data libraries maintained by different institutes. For example, the actinides data were taken from BOFOD [170], the JAERI evaluations were from JENDL photonuclear data file 2004 [137]. KAERI had significant contributions to both JENDL 2004 and IAEA 1999.

The sources of the IAEA 2019 library are classified into:

- New photonuclear data evaluations for IAEA 2019 at CIAE/CNDC, as well as special upgrades of their new upcoming CENDL (Chinese Evaluated Nuclear Data Library);
- JENDL Photonuclear Data File [155], its upgrade and high energy extension, as well as new evaluations specifically for IAEA 2019 at NDC/JAEA;

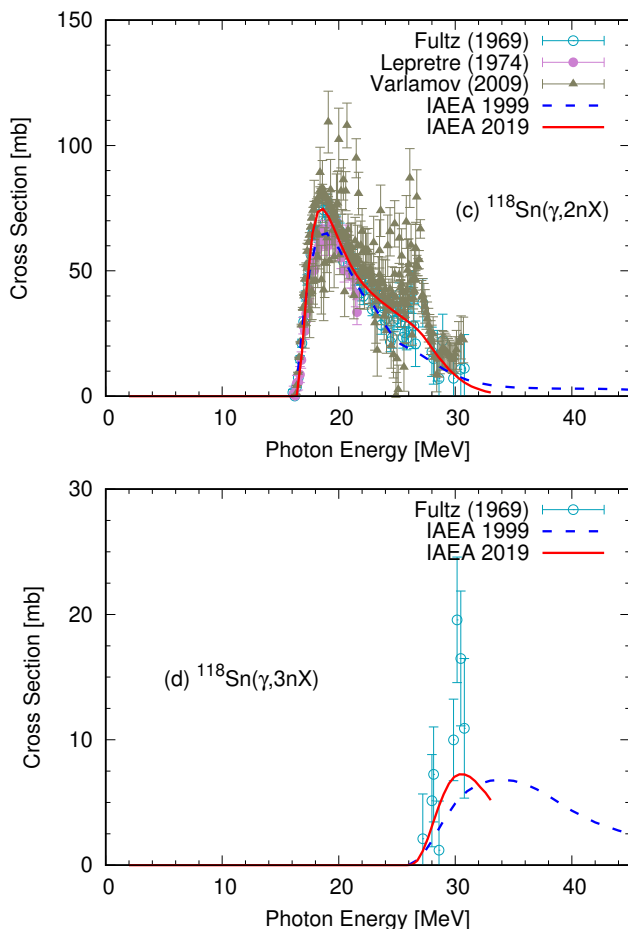


FIG. 56. (Color online) Comparison of evaluated and experimental data for ^{118}Sn ; (c) $(\gamma, 2nX)$, (d) $(\gamma, 3nX)$.

- New evaluations at IFIN-HH;
- New evaluations at NDC/KAERI; and
- Carry-over from IAEA 1999 photonuclear data library.

In some cases, the same isotopes were evaluated by more than one institutes. The inter-comparison of the different evaluations led to improvements in the evaluations performed by the individual institutes. They also triggered discussions on potential issues in the experimental data themselves, as most often, the differences in the evaluations were a result of adopting different experimental data sets. In the end, one set of evaluations was selected for these isotopes. The criteria for selection were based mainly on how well these evaluations reproduce the recommended available experimental data. A review team consisting of nuclear data evaluation experts from the CRP (T. Kawano (LANL), R. Capote (IAEA)) and external (O. Iwamoto (JAEA), Y.-O. Lee (KAERI)) reviewed the evaluations by comparing with available ex-

perimental data, and following discussions with the evaluators arrived at a consensus on a unique set of the data library. It should be emphasized that all the evaluated data files that were produced within this CRP, including those that were not included in the new IAEA library, are an improvement with respect to the previous evaluations as they are based on improved models and codes as well new and corrected experimental data, and could be available as part of other libraries such as CENDL, JENDL, *etc.* in the near future.

The new IAEA Photonuclear Data Library is produced in the ENDF-6 format. The ENDF-6 format adopted for the stored cross sections and energy spectra depends on the isotopes, due to a difficulty in applying a common evaluation technique and format for all the isotopes. For some nuclides, typically for light elements, exclusive nuclear reaction channels are explicitly given in MF=3 (MF: the file number defined in ENDF-6). While in many cases the MF=3 contains the total photo-absorption cross section only, and all individual information, such as the particle multiplicities and the residual nucleus production probabilities, is stored in MF=6. The MF=6 representation is the most common method since the library contains the high energy data part where the number of exclusive reaction channels exceeds the limitation of the ENDF-6 rule.

The new IAEA Photonuclear Data Library 2019 is available from the web interface of the IAEA Nuclear Data Section (<http://www-nds.iaea.org/>).

VII. CONCLUSIONS

The new 2019 IAEA Photonuclear Data Library contains 219 isotopes, including 55 newly added data files. This library is one of the major final products of the IAEA CRP “Updating the Photonuclear Data Library and Generating a Reference Database for Photon Strength Functions.” The photon energy range was extended to 200 MeV to cover many applications of photonuclear data for radiation transport calculations as well as isotope production. In this paper we reviewed the experimental techniques used to measure photonuclear reaction cross sections, and summarized possible issues in the reported experimental data due to deficiencies in the techniques. We discussed the method of assessing published partial photoneutron cross section data to identify neutron mis-counting issues that lurk in the neutron-multiplicity sorting technique. The method which is based on the F_i correction factors, has been applied to several new evaluations. However, the need to consider the model-dependent uncertainties when applying it to correct the experimental data was highlighted in this review. One important accomplishment of this CRP was the completion of the new direct multiplicity counting technique at the NewSUBARU facility that led to new measurements of (γ, inX) cross sections for seven isotopes. These new data were considered in the new eval-

uations performed for the updated library.

The statistical Hauser-Feshbach model calculations was heavily involved in the data evaluation; GLUNF and MEND-G codes at CIAE, CCONE at JAEA, TALYS at KAERI, and EMPIRE at IFIN-HH. Since an understanding of the characteristics of each Hauser-Feshbach code is crucial to estimate the uncertainties coming from the limitation in the evaluation technique, a code inter-comparison was performed. The comparison also included the CoH₃ code at LANL, as well as the CPNRM code at SINP/MSU, which was used for the F_i value correction to experimental data. We reported that the largest difference amongst the model codes is seen in the pre-equilibrium modeling in each code.

We assembled the new data library by collecting five main sources: new evaluations performed at CIAE along with some special upgrades of the new CENDL photodata library for the IAEA, JENDL/PD-2016 evaluations and new evaluations at JAEA, new evaluations at KAERI, and new evaluations at IFIN-HH based on the experimental data from NewSUBARU Laser Compton Scattering Facility, and some data files carried-over from the 1999 IAEA Photonuclear Data Library. Finally some selected comparisons of the newly evaluated data with available experimental data were given to demonstrate the improved evaluations compared to the previous IAEA data library.

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Appendix A: Contents of IAEA 1999 and 2019 Photonuclear Data Libraries

In the following table, we summarize the source of evaluated file for each isotope. In the IAEA 1999 column, an institute name that undertook the evaluation is given. When this column is empty, no evaluation is given in IAEA 1999. In the IAEA 2019 column, JENDL means these files were taken from JENDL Photonuclear Data File 2016 [155], the files denoted by JENDL(u) are the updated JENDL files, the = sign indicates the IAEA 1999 data were carried over, and the institution names stand for the new evaluations performed at each institute.

TABLE II. Contents of IAEA 1999 and 2019 photonuclear data libraries. The source of the data or leading evaluation institute is given. The = sign means the 2019 library is the same as 1999. JENDL and JENDL(u) are JENDL/PD-2016 ((u) stands for update).

	IAEA 1999	IAEA 2019	Author	Mon/Year
² H	JAERI	=	T.Murata	Aug95
³ He		JENDL	T.Murata	Feb03
⁶ Li		JENDL(u)	T.Murata	Aug13
⁷ Li		JENDL(u)	T.Murata	Aug13
⁹ Be	CIAE	CIAE	X.Tao	Dec18
¹² C	LANL	CIAE	X.Tao	Jun19
¹³ C	KAERI	=	Y.Han	Dec99
¹⁴ C		KAERI	Y.-S.Cho	Apr19
¹⁴ N	JAERI	CIAE	X.Tao	Sep19
¹⁵ N	KAERI	=	Y.Han	Dec99
¹⁶ O	LANL	CIAE	X.Tao	Jun19
¹⁷ O	KAERI	=	Y.Han	Dec99
¹⁸ O	KAERI	=	Y.Han	Dec99
¹⁹ F		JENDL(u)	T.Murata	Nov13
²³ Na	KAERI	=	Y.Han	Dec99
²⁴ Mg	KAERI	=	Y.Han	Dec99
²⁵ Mg	KAERI	=	Y.Han	Dec99
²⁶ Mg	KAERI	=	Y.Han	Dec99
²⁷ Al	LANL	CIAE	J.Wang	Apr19
²⁷ Si	KAERI	=	Y.Han	Dec99
²⁸ Si	KAERI	=	Y.Han	Dec99
²⁹ Si	KAERI	=	Y.Han	Dec99
³⁰ Si	KAERI	=	Y.Han	Dec99
³² S	KAERI	=	Y.Han	Dec99
³³ S	KAERI	=	Y.Han	Dec99
³⁴ S	KAERI	=	Y.Han	Dec99
³⁶ S	KAERI	=	Y.Han	Dec99
³⁵ Cl	KAERI	JAEA	N.Iwamoto	May18
³⁷ Cl	KAERI	JAEA	N.Iwamoto	May18
³⁶ Ar	KAERI	JAEA	N.Iwamoto	May18
³⁸ Ar	KAERI	JAEA	N.Iwamoto	May18
⁴⁰ Ar	KAERI	JAEA	N.Iwamoto	May18
³⁹ K	KAERI	JAEA	N.Iwamoto	Apr19
⁴⁰ K	KAERI	JAEA	N.Iwamoto	Apr19
⁴¹ K	KAERI	JAEA	N.Iwamoto	Apr19
⁴⁰ Ca	LANL	JENDL(u)	T.Murata	Nov13
⁴² Ca	KAERI	=	Y.Han	Dec99

TABLE II continued.

	IAEA 1999	IAEA 2019	Author	Mon/Year
⁴³ Ca	KAERI	=	Y.Han	Dec99
⁴⁴ Ca	KAERI	JENDL(u)	T.Murata	Nov13
⁴⁶ Ca	KAERI	=	Y.Han	Dec99
⁴⁸ Ca	KAERI	JENDL(u)	T.Murata	Nov13
⁴⁵ Sc		JAEA	N.Iwamoto	May18
⁴⁶ Ti	KAERI	JAEA	N.Iwamoto	May18
⁴⁷ Ti	KAERI	JAEA	N.Iwamoto	May18
⁴⁸ Ti	KAERI	JAEA	N.Iwamoto	May18
⁴⁹ Ti	KAERI	JAEA	N.Iwamoto	May18
⁵⁰ Ti	KAERI	JAEA	N.Iwamoto	May18
⁵⁰ V		JAEA	N.Iwamoto	May18
⁵¹ V	CIAE	JAEA	N.Iwamoto	May18
⁵⁰ Cr	CIAE	CIAE	X.Tao	Dec18
⁵² Cr	CIAE	JAEA	N.Iwamoto	May18
⁵³ Cr	CIAE	CIAE	X.Tao	Dec18
⁵⁴ Cr	CIAE	CIAE	X.Tao	Dec18
⁵⁵ Mn	KAERI	JAEA	N.Iwamoto	May18
⁵⁴ Fe	JAERI	JAEA	N.Iwamoto	May18
⁵⁶ Fe	JAERI	JAEA	N.Iwamoto	May18
⁵⁷ Fe	KAERI	JAEA	N.Iwamoto	May18
⁵⁸ Fe	KAERI	JAEA	N.Iwamoto	May18
⁵⁹ Co	KAERI	IFIN-HH	D.Filipescu	Apr19
⁵⁸ Ni	JAERI	JAEA	N.Iwamoto	May18
⁶⁰ Ni	KAERI	JAEA	N.Iwamoto	May18
⁶¹ Ni	KAERI	JAEA	N.Iwamoto	May18
⁶² Ni	KAERI	JAEA	N.Iwamoto	May18
⁶⁴ Ni	KAERI	JAEA	N.Iwamoto	May18
⁶³ Cu	LANL	JAEA	N.Iwamoto	May18
⁶⁵ Cu	JAERI	JAEA	N.Iwamoto	May18
⁶⁴ Zn	JAERI	KAERI	Y.-S.Cho	Apr19
⁶⁶ Zn	KAERI	JAEA	N.Iwamoto	Apr19
⁶⁷ Zn	KAERI	JAEA	N.Iwamoto	Apr19
⁶⁸ Zn	KAERI	JAEA	N.Iwamoto	Apr19
⁷⁰ Zn	KAERI	JAEA	N.Iwamoto	Apr19
⁷⁰ Ge	KAERI	JAEA	N.Iwamoto	Apr19
⁷² Ge	KAERI	JAEA	N.Iwamoto	Apr19
⁷³ Ge	KAERI	JAEA	N.Iwamoto	Apr19
⁷⁴ Ge	KAERI	JAEA	N.Iwamoto	Apr19
⁷⁶ Ge	KAERI	JAEA	N.Iwamoto	Apr19
⁷⁵ As		KAERI	Y.-S.Cho	Apr19
⁷⁶ Se		KAERI	Y.-S.Cho	Apr19
⁷⁸ Se		KAERI	Y.-S.Cho	Apr19
⁸⁰ Se		KAERI	Y.-S.Cho	Apr19
⁸² Se		KAERI	Y.-S.Cho	Apr19
⁸⁴ Sr	KAERI	JAEA	N.Iwamoto	Apr19
⁸⁶ Sr	KAERI	JAEA	N.Iwamoto	Apr19
⁸⁷ Sr	KAERI	JAEA	N.Iwamoto	Apr19
⁸⁸ Sr	KAERI	JAEA	N.Iwamoto	Apr19
⁹⁰ Sr	KAERI	JAEA	N.Iwamoto	Apr19
⁸⁹ Y		IFIN-HH	D.Filipescu	Apr19
⁹⁰ Zr	KAERI	CIAE	J.Wang	Jun18
⁹¹ Zr	CIAE	KAERI	Y.-S.Cho	Apr19
⁹² Zr	CIAE	JAEA	N.Iwamoto	Apr19
⁹³ Zr	KAERI	JAEA	N.Iwamoto	Apr19
⁹⁴ Zr	KAERI	KAERI	Y.-S.Cho	Apr19

TABLE II continued.

	IAEA 1999	IAEA 2019	Author	Mon/Year
⁹⁶ Zr	CIAE	JAEA	N.Iwamoto	Apr19
⁹³ Nb	KAERI	JAEA	N.Iwamoto	Apr19
⁹⁴ Nb	KAERI	JAEA	N.Iwamoto	Apr19
⁹² Mo	KAERI	JAEA	N.Iwamoto	Apr19
⁹⁴ Mo	KAERI	JAEA	N.Iwamoto	Apr19
⁹⁵ Mo	KAERI	JAEA	N.Iwamoto	Apr19
⁹⁶ Mo	KAERI	JAEA	N.Iwamoto	Apr19
⁹⁷ Mo	KAERI	JAEA	N.Iwamoto	Apr19
⁹⁸ Mo	KAERI	JAEA	N.Iwamoto	Apr19
¹⁰⁰ Mo	KAERI	JAEA	N.Iwamoto	Apr19
⁹⁸ Ru		JAEA	N.Iwamoto	Apr19
¹⁰³ Rh		IFIN-HH	D.Filipescu	May19
¹⁰² Pd	KAERI	JAEA	N.Iwamoto	Apr19
¹⁰⁴ Pd	KAERI	JAEA	N.Iwamoto	Apr19
¹⁰⁵ Pd	KAERI	JAEA	N.Iwamoto	Apr19
¹⁰⁶ Pd	KAERI	JAEA	N.Iwamoto	Apr19
¹⁰⁷ Pd	KAERI	JAEA	N.Iwamoto	Apr19
¹⁰⁸ Pd	KAERI	JAEA	N.Iwamoto	Apr19
¹¹⁰ Pd	KAERI	JAEA	N.Iwamoto	Apr19
¹⁰⁷ Ag	KAERI	JAEA	N.Iwamoto	Apr19
¹⁰⁸ Ag	KAERI	JAEA	N.Iwamoto	Apr19
¹⁰⁹ Ag	KAERI	JAEA	N.Iwamoto	Apr19
¹⁰⁶ Cd	KAERI	JAEA	N.Iwamoto	Apr19
¹⁰⁸ Cd	KAERI	JAEA	N.Iwamoto	Apr19
¹¹⁰ Cd	KAERI	JAEA	N.Iwamoto	Apr19
¹¹¹ Cd	KAERI	JAEA	N.Iwamoto	Apr19
¹¹² Cd	KAERI	JAEA	N.Iwamoto	Apr19
¹¹³ Cd	KAERI	JAEA	N.Iwamoto	Apr19
¹¹⁴ Cd	KAERI	JAEA	N.Iwamoto	Apr19
¹¹⁶ Cd	KAERI	JAEA	N.Iwamoto	Apr19
¹¹⁵ In		KAERI	Y.-S.Cho	Apr19
¹¹² Sn	KAERI	JAEA	N.Iwamoto	Apr19
¹¹⁴ Sn	KAERI	JAEA	N.Iwamoto	Apr19
¹¹⁵ Sn	KAERI	JAEA	N.Iwamoto	Apr19
¹¹⁶ Sn	KAERI	KAERI	Y.-S.Cho	Apr19
¹¹⁷ Sn	KAERI	JAEA	N.Iwamoto	Apr19
¹¹⁸ Sn	KAERI	CIAE	Y.Tian	Dec17
¹¹⁹ Sn	KAERI	JAEA	N.Iwamoto	Apr19
¹²⁰ Sn	KAERI	JAEA	N.Iwamoto	Apr19
¹²² Sn	KAERI	JAEA	N.Iwamoto	Apr19
¹²⁴ Sn	KAERI	JAEA	N.Iwamoto	Apr19
¹²¹ Sb	KAERI	JAEA	N.Iwamoto	Apr19
¹²³ Sb	KAERI	JAEA	N.Iwamoto	Apr19
¹²⁰ Te	KAERI	JAEA	N.Iwamoto	Apr19
¹²² Te	KAERI	JAEA	N.Iwamoto	Apr19
¹²³ Te	KAERI	JAEA	N.Iwamoto	Apr19
¹²⁴ Te	KAERI	JAEA	N.Iwamoto	Apr19
¹²⁵ Te	KAERI	JAEA	N.Iwamoto	Apr19
¹²⁶ Te	KAERI	JAEA	N.Iwamoto	Apr19
¹²⁸ Te	KAERI	JAEA	N.Iwamoto	Apr19
¹³⁰ Te	KAERI	JAEA	N.Iwamoto	Apr19
¹²⁷ I	KAERI	JAEA	N.Iwamoto	Apr19
¹²⁹ I	KAERI	JAEA	N.Iwamoto	Apr19
¹³² Xe		JAEA	N.Iwamoto	Apr19
¹³³ Cs	KAERI	KAERI	Y.-S.Cho	Apr19

TABLE II continued.

	IAEA 1999	IAEA 2019	Author	Mon/Year
¹³⁵ Cs	KAERI	JAEA	N.Iwamoto	Apr19
¹³⁷ Cs	KAERI	JAEA	N.Iwamoto	Apr19
¹³⁸ Ba		KAERI	Y.-S.Cho	Apr19
¹³⁹ La		JAEA	N.Iwamoto	Apr19
¹⁴⁰ Ce		KAERI	Y.-S.Cho	Apr19
¹⁴² Ce		KAERI	Y.-S.Cho	Apr19
¹⁴¹ Pr	KAERI	JAEA	N.Iwamoto	Apr19
¹⁴² Nd		KAERI	Y.-S.Cho	Apr19
¹⁴³ Nd		KAERI	Y.-S.Cho	Apr19
¹⁴⁴ Nd		KAERI	Y.-S.Cho	Apr19
¹⁴⁵ Nd		KAERI	Y.-S.Cho	Apr19
¹⁴⁶ Nd		KAERI	Y.-S.Cho	Apr19
¹⁴⁸ Nd		KAERI	Y.-S.Cho	Apr19
¹⁵⁰ Nd		KAERI	Y.-S.Cho	Apr19
¹⁴⁴ Sm	KAERI	JAEA	N.Iwamoto	Apr19
¹⁴⁷ Sm	KAERI	JAEA	N.Iwamoto	Apr19
¹⁴⁸ Sm	KAERI	JAEA	N.Iwamoto	Apr19
¹⁴⁹ Sm	KAERI	JAEA	N.Iwamoto	Apr19
¹⁵⁰ Sm	KAERI	JAEA	N.Iwamoto	Apr19
¹⁵¹ Sm	KAERI	JAEA	N.Iwamoto	Apr19
¹⁵² Sm	KAERI	JAEA	N.Iwamoto	Apr19
¹⁵⁴ Sm	KAERI	JAEA	N.Iwamoto	Apr19
¹⁵³ Eu		KAERI	Y.-S.Cho	Apr19
¹⁵⁶ Gd		JAEA	N.Iwamoto	Apr19
¹⁵⁷ Gd		JAEA	N.Iwamoto	Apr19
¹⁵⁸ Gd		JAEA	N.Iwamoto	Apr19
¹⁶⁰ Gd		JAEA	N.Iwamoto	Apr19
¹⁵⁸ Tb	KAERI	JAEA	N.Iwamoto	Apr19
¹⁵⁹ Tb	KAERI	IFIN-HH	D.Filipescu	May19
¹⁶² Dy		JAEA	N.Iwamoto	Apr19
¹⁶³ Dy		JAEA	N.Iwamoto	Apr19
¹⁶⁵ Ho	KAERI	IFIN-HH	D.Filipescu	May19
¹⁶⁶ Er		JAEA	N.Iwamoto	Apr19
¹⁷⁰ Er		JAEA	N.Iwamoto	Apr19
¹⁶⁹ Tm		IFIN-HH	D.Filipescu	Apr19
¹⁷⁵ Lu		KAERI	Y.-S.Cho	Apr19
¹⁷⁴ Hf		JAEA	N.Iwamoto	Apr19
¹⁷⁶ Hf		JAEA	N.Iwamoto	Apr19
¹⁷⁷ Hf		JAEA	N.Iwamoto	Apr19
¹⁷⁸ Hf		JAEA	N.Iwamoto	Apr19
¹⁷⁹ Hf		JAEA	N.Iwamoto	Apr19
¹⁸⁰ Hf		JAEA	N.Iwamoto	Apr19
¹⁸¹ Ta	JAERI	IFIN-HH	D.Filipescu	Apr19
¹⁸⁰ W	CIAE	CIAE	X.B.Ke	Dec17
¹⁸² W	JAERI	CIAE	X.B.Ke	Dec17
¹⁸³ W	CIAE	CIAE	X.B.Ke	Dec17
¹⁸⁴ W	LANL	CIAE	X.B.Ke	Dec17
¹⁸⁶ W	JAERI	CIAE	X.B.Ke	Dec17
¹⁸⁵ Re		JAEA	N.Iwamoto	Apr19
¹⁸⁷ Re		JAEA	N.Iwamoto	Apr19
¹⁸⁶ Os		KAERI	Y.-S.Cho	Apr19
¹⁸⁸ Os		KAERI	Y.-S.Cho	Apr19
¹⁸⁹ Os		KAERI	Y.-S.Cho	Apr19
¹⁹⁰ Os		KAERI	Y.-S.Cho	Apr19
¹⁹² Os		KAERI	Y.-S.Cho	Apr19

TABLE II continued.

	IAEA 1999	IAEA 2019	Author	Mon/Year
¹⁹⁴ Pt		JAEA	N.Iwamoto	Apr19
¹⁹⁷ Au	KAERI	JAEA	N.Iwamoto	Apr19
²⁰⁶ Pb	LANL	JAEA	N.Iwamoto	Apr19
²⁰⁷ Pb	LANL	JAEA	N.Iwamoto	Apr19
²⁰⁸ Pb	LANL	JAEA	N.Iwamoto	Apr19
²⁰⁹ Bi	CIAE	KAERI	Y.-S.Cho	Apr19
²²⁶ Ra		JAEA	N.Iwamoto	Apr19
²³² Th	IPPE	JAEA	N.Iwamoto	Apr19
²³³ U	IPPE	JAEA	N.Iwamoto	Apr19
²³⁴ U	IPPE	JAEA	N.Iwamoto	Apr19
²³⁵ U	IPPE	JAEA	N.Iwamoto	Apr19
²³⁶ U	IPPE	JAEA	N.Iwamoto	Apr19
²³⁸ U	IPPE	JAEA	N.Iwamoto	Apr19
²³⁷ Np		JAEA	N.Iwamoto	Apr19
²³⁸ Pu	IPPE	JAEA	N.Iwamoto	Apr19
²³⁹ Pu	IPPE	JAEA	N.Iwamoto	Apr19
²⁴¹ Pu	IPPE	JAEA	N.Iwamoto	Apr19

Appendix B: GDR ATLAS

Here we summarize the updated tables for the recommended experimental GDR parameters within the Standard Lorentzian (SLO) and the Simplified version of the modified Lorentzian (SMLO) approaches. In the case of double humped shape, two sets of GDR parameters are given. A complete list of all the references is given at <https://www-nds.iaea.org/CRP-photonuclear/>, and only abbreviations are given for the sake of simplicity.

TABLE III. Recommended experimental GDR parameters within the Standard Lorentzian (SLO) approach.

	E [MeV]	Γ [MeV]	σ [mb]	Range [MeV]	Ref.
⁶ Li	23.69	5.26	3.71	21.5 – 27.0	1986Var
⁷ Li	18.59	16.28	3.65	13.2 – 25.6	1985Ahr
⁹ Be	23.75	9.47	5.17	17.5 – 26.0	1975Ahr
¹⁰ B	21.72	9.08	4.64	8.5 – 24.9	1987Ahs
¹² C	22.86	3.61	21.30	20.1 – 25.0	1969Bez
¹³ C	24.60	8.43	12.71	14.5 – 29.0	2002Ish
¹⁴ C	15.41	5.82	7.49	14.5 – 30.0	2002Ish
	26.13	7.78	8.13		
^{nat} C	23.12	4.19	19.28	19.5 – 25.6	1985Ahr
¹⁴ N	23.05	6.95	22.95	18.2 – 28.0	1969Bez
¹⁵ N	24.78	12.82	13.82	14.5 – 28.0	2002Ish
¹⁶ O	23.70	5.36	27.96	18.1 – 26.0	1975Ahr
¹⁷ O	23.40	5.48	21.82	18.5 – 26.5	2002Ish
¹⁸ O	19.08	2.12	5.13	18.5 – 26.0	2002Ish
	24.10	5.25	13.48		
^{nat} O	23.60	5.82	27.07	18.9 – 27.9	1985Ahr
¹⁹ F	21.61	12.57	16.58	10.0 – 24.0	2002Ish
²³ Na	17.43	3.10	12.38	14.2 – 23.0	1981Ish
	21.13	4.51	26.98		
²⁴ Mg	19.51	2.71	21.40	16.5 – 27.0	1966Dol
	23.88	8.86	25.20		
²⁵ Mg	22.06	6.09	34.99	9.0 – 24.2	2002Ish
²⁶ Mg	17.38	2.15	16.53	16.1 – 26.5	2003Var
	23.64	7.25	39.12		
^{nat} Mg	22.55	7.97	20.88	15.1 – 26.6	1965Wyc
²⁷ Al	20.73	7.45	38.99	14.0 – 24.1	1985Ahr
²⁸ Si	19.81	2.56	38.75	16.7 – 23.0	2003Var
	21.81	3.15	40.23		
²⁹ Si	20.70	5.60	40.01	14.2 – 23.0	2002Ish
³⁰ Si	20.86	7.40	29.56	14.2 – 23.0	2002Ish
^{nat} Si	20.35	4.53	51.41	16.4 – 25.8	1975Ahr
	25.16	2.86	10.47		
³² S	19.51	4.83	35.43	14.7 – 23.0	1968Dol
³⁴ S	20.89	9.61	50.54	12.0 – 25.0	1986Ass
^{nat} S	20.31	5.48	47.84	17.2 – 23.6	1965Wyc
⁴⁰ Ar	19.86	9.12	56.89	10.5 – 25.0	2002Ish
^{nat} K	21.12	6.89	22.58	16.0 – 25.9	1974Ve1
⁴⁰ Ca	20.58	6.23	104.78	17.2 – 23.7	1966Dol
⁴² Ca	20.11	8.07	72.15	15.2 – 23.0	2003Ero
⁴⁴ Ca	19.60	11.33	63.70	15.5 – 26.0	2003Ero
⁴⁸ Ca	19.70	6.23	105.44	17.9 – 21.6	1987Oke
^{nat} Ca	20.06	4.89	94.13	15.1 – 24.0	1975Ahr
⁴⁶ Ti	19.96	6.92	78.94	13.2 – 25.0	2002Ish

TABLE III continued.

	E [MeV]	Γ [MeV]	σ [mb]	Range [MeV]	Ref.
^{48}Ti	19.78	8.42	63.74	14.5 – 23.0	2002Ish
^{51}V	17.90	4.55	60.32	14.1 – 22.9	1962Fu1
	21.26	4.37	26.38		
^{52}Cr	19.16	6.19	81.34	14.3 – 23.0	2002Ish
^{55}Mn	16.43	2.95	27.02	14.0 – 23.0	1979Al2
	19.77	8.61	52.03		
^{54}Fe	19.35	5.50	147.00	16.0 – 23.0	1978Nor
^{59}Co	16.43	2.73	28.28	14.0 – 20.9	1979Al2
	18.64	7.31	57.16		
^{58}Ni	18.78	5.57	87.90	14.1 – 22.0	2003Var
^{60}Ni	16.69	3.47	62.80	12.1 – 21.0	2003Var
	19.57	5.26	71.14		
^{63}Cu	16.43	4.84	79.79	14.0 – 21.0	2003Var
	20.15	5.52	49.39		
^{65}Cu	16.92	8.09	86.38	14.2 – 21.0	2003Var
$^{\text{nat}}\text{Cu}$	18.12	5.61	97.93	14.4 – 24.9	1965Wyc
^{64}Zn	16.23	3.25	41.21	14.0 – 20.8	1976Ca1
	19.16	5.91	54.90		
^{65}Zn	16.17	3.06	34.89	12.0 – 21.0	2003Rod
	19.04	6.50	55.53		
^{70}Ge	15.16	5.92	160.51	10.0 – 20.0	1975Mcc
^{72}Ge	17.88	5.71	167.57	10.0 – 24.0	1975Mcc
^{74}Ge	14.51	2.01	25.54	13.1 – 20.8	1976Ca1
	17.03	7.97	100.80		
^{76}Ge	15.48	4.37	61.70	13.1 – 20.8	1976Ca1
	18.87	10.99	71.09		
^{75}As	14.98	3.66	41.85	13.1 – 20.9	1969Be1
	17.59	7.12	75.35		
^{76}Se	15.67	6.33	151.59	13.1 – 19.7	1978Gur
^{78}Se	14.97	3.91	70.45	13.1 – 20.8	1976Ca1
	18.42	6.19	79.42		
^{80}Se	16.60	6.80	137.82	13.1 – 20.0	2016Va1
^{82}Se	16.00	5.68	175.06	13.1 – 19.9	1978Gur
$^{\text{nat}}\text{Rb}$	16.73	4.25	190.04	10.6 – 17.9	1971Lep
$^{\text{nat}}\text{Sr}$	16.79	4.32	205.99	10.9 – 17.9	1971Lep
^{89}Y	16.74	4.23	224.56	14.0 – 19.0	1971Le1
^{90}Zr	16.82	3.99	253.58	14.9 – 18.5	2003Var
^{91}Zr	16.58	4.17	183.17	14.0 – 18.9	1967Be2
^{92}Zr	16.26	4.64	164.72	14.0 – 18.9	1967Be2
^{94}Zr	16.21	5.25	159.83	14.0 – 18.9	1967Be2
^{93}Nb	16.58	4.95	200.25	14.0 – 19.0	1971Le1
^{92}Mo	17.16	4.68	239.77	14.4 – 19.0	2003Var
^{94}Mo	16.53	5.12	192.92	9.6 – 18.9	1974Be3
^{96}Mo	16.11	5.64	184.80	13.2 – 17.0	1974Be3
^{98}Mo	15.79	5.90	188.16	13.2 – 18.9	1974Be3
^{100}Mo	15.72	7.68	170.10	12.1 – 20.0	1974Be3
^{103}Rh	16.24	7.49	192.03	13.1 – 19.0	2003Var
^{108}Pd	14.97	5.42	143.49	10.0 – 19.0	1969Dea
	18.13	4.03	103.37		
$^{\text{nat}}\text{Pd}$	15.89	6.30	201.91	10.2 – 17.8	1971Lep
^{107}Ag	15.83	6.49	185.05	9.5 – 19.0	1969Ish
^{109}Ag	13.54	3.49	80.46	13.1 – 19.0	1969Ish
	16.62	4.41	113.46		
$^{\text{nat}}\text{Ag}$	16.06	7.33	197.91	13.2 – 18.9	1971Lep
$^{\text{nat}}\text{Cd}$	15.77	5.70	228.50	10.2 – 17.8	1971Lep

TABLE III continued.

	E [MeV]	Γ [MeV]	σ [mb]	Range [MeV]	Ref.
^{115}In	15.72	5.57	245.50	13.2 – 17.8	1974Le1
^{112}Sn	15.62	5.01	262.94	10.9 – 18.0	1974Sor
^{114}Sn	15.82	6.08	243.01	13.1 – 18.0	1975Sor
^{116}Sn	15.55	5.06	269.30	13.1 – 17.9	1974Le1
^{117}Sn	15.64	5.02	257.30	13.2 – 17.8	1974Le1
^{118}Sn	15.43	4.84	277.65	13.1 – 17.9	1974Le1
^{119}Sn	15.53	4.78	251.37	13.0 – 17.9	1969Fu1
^{120}Sn	15.37	5.08	284.00	13.1 – 17.9	1974Le1
^{122}Sn	15.34	4.73	266.89	13.1 – 18.0	1975Sor
^{124}Sn	15.31	4.94	270.63	13.1 – 18.0	2003Var
$^{\text{nat}}\text{Sb}$	15.48	5.01	279.38	10.2 – 17.8	1971Lep
^{124}Te	15.23	5.50	280.36	12.0 – 18.9	1976Le2
^{126}Te	15.15	5.36	295.55	12.0 – 18.9	1976Le2
^{128}Te	15.12	5.30	304.18	12.0 – 18.9	1976Le2
^{130}Te	15.11	4.98	319.24	12.0 – 18.9	1976Le2
^{127}I	14.77	4.09	229.29	12.0 – 20.0	1999Bel
	17.30	3.69	55.30		
^{133}Cs	15.33	5.28	315.25	12.0 – 19.0	1974Le1
^{138}Ba	15.13	4.51	317.33	12.1 – 18.7	2016Va2
$^{\text{nat}}\text{Ba}$	15.29	4.93	353.21	10.1 – 17.8	1971Be4
^{139}La	15.24	4.82	367.47	12.0 – 18.9	1972De1
^{140}Ce	15.03	4.39	381.89	12.0 – 18.9	1976Le2
^{142}Ce	14.85	5.08	331.25	12.0 – 18.9	1976Le2
^{141}Pr	15.19	4.23	342.68	12.1 – 16.9	1987Ber
^{142}Nd	14.95	4.46	360.54	13.1 – 18.0	2003Var
^{143}Nd	15.00	4.73	347.60	12.0 – 19.0	1971Ca1
^{144}Nd	15.04	5.25	315.51	12.0 – 18.9	1971Ca1
^{145}Nd	14.94	6.27	295.27	12.0 – 18.9	1971Ca1
^{146}Nd	14.73	5.74	309.04	12.0 – 18.9	1971Ca1
	12.78	4.03	110.23		
^{148}Nd	15.49	5.22	215.89	10.8 – 18.6	1971Ca1
	12.30	3.38	175.75		
^{150}Nd	16.03	5.12	220.50	10.8 – 18.6	1971Ca1
	15.31	4.42	381.69		
^{144}Sm	15.31	4.42	381.69	12.1 – 18.9	1974Ca5
^{148}Sm	14.82	5.06	337.78	12.1 – 18.9	1974Ca5
^{150}Sm	14.59	5.92	310.73	12.1 – 18.9	1974Ca5
^{152}Sm	12.39	2.99	176.80	10.9 – 18.8	1974Ca5
	15.73	5.15	232.25		
^{154}Sm	12.17	2.80	181.90	10.9 – 18.6	1981Gur
	15.63	5.89	209.70		
^{151}Eu	13.88	4.69	254.17	10.2 – 18.0	1971Vas
	14.45	0.75	65.45		
^{153}Eu	12.33	2.77	155.86	10.9 – 18.7	1969Be8
	15.78	5.76	219.21		
^{152}Gd	11.79	3.03	145.74	10.2 – 18.0	1971Vas
	14.72	3.16	253.50		
^{154}Gd	11.97	2.65	163.89	10.2 – 18.0	1971Vas
	15.05	3.35	245.22		
^{156}Gd	12.46	3.14	230.03	10.9 – 18.7	1981Gur
	15.79	4.56	215.94		
^{158}Gd	11.86	2.94	182.05	10.2 – 18.0	1971Vas
	15.16	3.31	245.62		
^{160}Gd	12.28	3.33	234.33	10.9 – 18.8	2003Var
	16.06	5.12	247.23		

TABLE III continued.

	E [MeV]	Γ [MeV]	σ [mb]	Range [MeV]	Ref.
^{159}Tb	12.42	2.71	171.70	11.1 – 19.0	1976Gor
	15.86	5.98	295.78		
^{165}Ho	12.38	2.59	220.67	11.1 – 18.7	1981Gur
	15.48	4.05	226.69		
^{168}Er	12.09	3.66	237.40	10.9 – 18.8	1981Gur
	15.54	3.99	252.64		
^{174}Yb	12.50	3.41	339.31	10.9 – 18.7	1981Gur
	15.68	3.74	291.85		
^{175}Lu	12.32	2.59	218.42	11.0 – 18.7	1969Be6
	15.47	4.64	284.83		
^{176}Hf	12.34	2.77	279.26	10.9 – 17.9	1977Gor
	15.67	4.72	275.10		
^{178}Hf	12.42	4.89	363.72	10.8 – 18.6	1981Gur
	15.70	3.13	234.94		
^{180}Hf	12.55	4.71	351.75	10.8 – 18.7	1981Gur
	15.61	3.27	243.23		
^{181}Ta	12.19	2.93	262.35	10.8 – 18.6	1981Gur
	14.99	5.13	317.52		
^{182}W	11.98	3.91	283.99	11.0 – 18.8	1981Gur
	14.94	5.16	259.40		
^{184}W	11.92	4.52	347.68	11.0 – 17.6	1981Gur
	15.05	3.87	233.17		
^{186}W	13.04	6.60	410.29	10.9 – 18.7	1981Gur
	14.89	2.12	69.05		
^{185}Re	12.59	2.22	221.75	10.2 – 18.0	1973Gor
	15.17	6.02	328.79		
$^{\text{nat}}\text{Re}$	14.11	6.52	470.27	10.2 – 18.0	1975Vey
^{186}Os	12.73	2.34	214.24	11.1 – 18.9	2015Var
	14.73	4.08	355.57		
^{188}Os	12.81	2.83	281.11	10.8 – 18.9	2014Var
	14.93	3.83	388.79		
^{189}Os	12.92	3.07	329.04	10.8 – 18.9	2014Var
	15.02	3.94	299.55		
^{190}Os	13.10	3.34	372.87	10.8 – 18.9	2015Var
	15.12	3.80	247.98		
^{192}Os	12.59	2.13	173.74	10.8 – 18.9	2015Var
	14.32	4.60	426.27		
^{191}Ir	12.72	2.08	183.14	11.0 – 16.8	1978Go1
	14.21	5.27	382.41		
^{193}Ir	12.86	1.90	229.81	11.0 – 16.8	1978Go1
	14.30	5.62	356.07		
$^{\text{nat}}\text{Ir}$	13.77	4.86	495.41	10.2 – 18.0	1975Vey
^{194}Pt	13.42	3.61	453.02	11.0 – 17.8	1978Go1
	15.97	6.16	111.63		
^{195}Pt	12.99	2.92	357.53	11.0 – 17.8	1978Go1
	14.90	4.85	253.95		
^{196}Pt	13.28	3.10	345.43	11.0 – 17.8	1978Go1
	14.81	7.51	192.99		
^{198}Pt	13.56	4.88	528.76	11.0 – 17.8	1978Go1
^{197}Au	13.58	5.32	522.88	11.1 – 17.0	1981Gur
^{203}Tl	14.04	3.77	436.99	9.0 – 17.9	1970Ant
^{205}Tl	14.46	2.95	478.97	10.5 – 17.9	1970Ant
^{206}Pb	13.58	3.83	512.45	10.0 – 17.0	1964Ha2
^{207}Pb	13.55	3.95	479.80	10.0 – 17.0	1964Ha2

TABLE III continued.

	E [MeV]	Γ [MeV]	σ [mb]	Range [MeV]	Ref.
^{208}Pb	13.37	3.93	645.49	10.9 – 18.8	2003Var
$^{\text{nat}}\text{Pb}$	13.48	3.99	637.69	12.1 – 16.9	1985Ahr
^{209}Bi	13.79	5.02	588.63	10.9 – 18.3	1976Gu2
^{232}Th	10.86	2.86	230.28	10.2 – 18.3	1976Gu1
	13.74	4.74	382.21		
^{233}U	10.98	1.53	140.38	9.4 – 17.8	1986Be2
	13.30	5.71	439.10		
^{234}U	11.18	2.41	379.61	9.4 – 17.8	1986Be2
	14.03	4.46	398.77		
^{235}U	10.82	3.88	302.02	9.5 – 18.4	1976Gu1
	13.80	4.54	319.35		
^{236}U	11.04	2.65	279.96	9.5 – 17.8	1980Ca1
	13.92	4.77	407.27		
^{238}U	11.06	2.95	283.53	9.2 – 18.8	1976Gu1
	14.26	4.80	347.16		
$^{\text{nat}}\text{U}$	10.73	2.47	301.11	10.2 – 17.9	1985Ahr
	13.72	5.04	405.91		
^{237}Np	10.99	2.20	310.02	9.4 – 17.8	1986Be2
	14.08	4.66	535.89		
^{239}Pu	11.07	3.29	227.77	9.3 – 18.7	1976Gu1
	14.00	5.51	358.18		

TABLE IV. Recommended experimental GDR parameters within the Simplified version of the modified Lorentzian (SMLO) approach.

	E [MeV]	Γ [MeV]	σ [mb]	Range [MeV]	Ref.
^6Li	23.75	5.33	3.70	21.5 – 27.0	1986Var
^7Li	20.21	21.81	3.47	13.2 – 25.6	1985Ahr
^9Be	24.18	10.86	5.09	17.5 – 26.0	1975Ahr
^{10}B	23.31	17.51	4.12	8.5 – 24.9	1987Ahs
^{12}C	22.90	3.69	21.21	20.1 – 25.0	1969Bez
^{13}C	25.02	10.72	11.65	14.5 – 29.0	2002Ish
^{14}C	15.69	6.51	7.67	14.5 – 30.0	2002Ish
	26.27	7.11	7.74		
$^{\text{nat}}\text{C}$	23.14	4.24	19.38	19.5 – 25.6	1985Ahr
^{14}N	23.19	7.09	22.95	18.2 – 28.0	1969Bez
^{15}N	26.29	19.45	12.85	14.5 – 28.0	2002Ish
^{16}O	23.78	5.67	27.70	18.1 – 26.0	1975Ahr
^{17}O	23.46	5.95	21.29	18.5 – 26.5	2002Ish
^{18}O	19.12	2.31	5.51	18.5 – 26.0	2002Ish
	24.19	5.24	13.38		
$^{\text{nat}}\text{O}$	23.64	5.88	27.13	18.9 – 27.9	1985Ahr
^{19}F	24.09	24.69	14.94	10.0 – 24.0	2002Ish
^{23}Na	17.57	3.75	13.62	14.2 – 23.0	1981Ish
	21.26	4.35	25.45		
^{24}Mg	19.46	2.80	22.51	16.5 – 27.0	1966Dol
	24.30	9.80	24.18		
^{25}Mg	22.73	8.44	33.04	9.0 – 24.2	2002Ish
^{26}Mg	17.39	2.36	17.88	16.1 – 26.5	2003Var
	23.83	7.57	38.38		
$^{\text{nat}}\text{Mg}$	22.32	8.72	21.08	15.1 – 26.6	1965Wyc

TABLE IV continued.

	E [MeV]	Γ [MeV]	σ [mb]	Range [MeV]	Ref.
^{27}Al	21.00	8.62	37.64	14.0 – 24.1	1985Ahr
^{28}Si	19.88	2.91	38.41	16.7 – 23.0	2003Var
	21.84	3.39	36.60		
^{29}Si	20.97	6.88	37.83	14.2 – 23.0	2002Ish
^{30}Si	21.32	9.16	28.52	14.2 – 23.0	2002Ish
$^{\text{nat}}\text{Si}$	20.45	4.85	51.22	16.4 – 25.8	1975Ahr
	25.24	2.10	8.02		
^{32}S	19.57	5.10	34.93	14.7 – 23.0	1968Dol
^{34}S	21.66	13.43	47.29	12.0 – 25.0	1986Ass
$^{\text{nat}}\text{S}$	20.42	5.74	47.22	17.2 – 23.6	1965Wyc
^{40}Ar	19.91	11.50	54.19	10.5 – 25.0	2002Ish
$^{\text{nat}}\text{K}$	21.28	7.40	22.28	16.0 – 25.9	1974Ve1
^{40}Ca	20.72	6.44	104.09	17.2 – 23.7	1966Dol
^{42}Ca	20.53	9.75	68.87	15.2 – 23.0	2003Ero
^{44}Ca	20.08	12.45	62.19	15.5 – 26.0	2003Ero
^{48}Ca	19.90	6.42	104.95	17.9 – 21.6	1987OKe
$^{\text{nat}}\text{Ca}$	20.09	5.07	93.50	15.1 – 24.0	1975Ahr
^{46}Ti	19.95	7.99	76.44	13.2 – 25.0	2002Ish
^{48}Ti	20.13	9.98	61.66	14.5 – 23.0	2002Ish
^{51}V	18.18	5.29	63.92	14.1 – 22.9	1962Fu1
	21.37	3.36	18.23		
^{52}Cr	19.16	6.70	79.20	14.3 – 23.0	2002Ish
^{55}Mn	16.43	2.91	21.84	14.0 – 23.0	1979Al2
	20.13	11.28	51.12		
^{54}Fe	19.39	5.69	145.35	16.0 – 23.0	1978Nor
^{59}Co	16.44	2.42	20.11	14.0 – 20.9	1979Al2
	18.68	8.63	59.76		
^{58}Ni	18.87	6.16	86.03	14.1 – 22.0	2003Var
^{60}Ni	16.59	3.17	39.23	12.1 – 21.0	2003Var
	19.39	7.82	77.98		
^{63}Cu	16.79	5.76	87.60	14.0 – 21.0	2003Var
	20.37	4.56	35.92		
^{65}Cu	17.23	8.38	86.10	14.2 – 21.0	2003Var
$^{\text{nat}}\text{Cu}$	17.91	5.24	99.60	14.4 – 24.9	1965Wyc
^{64}Zn	16.37	3.84	48.04	14.0 – 20.8	1976Ca1
	19.49	5.82	47.80		
^{65}Zn	16.06	2.26	21.02	12.0 – 21.0	2003Rod
	18.87	8.62	60.27		
^{70}Ge	15.31	7.19	151.53	10.0 – 20.0	1975Mcc
^{72}Ge	17.85	6.22	159.50	10.0 – 24.0	1975Mcc
^{74}Ge	14.42	2.46	32.43	13.1 – 20.8	1976Ca1
	17.47	8.37	96.06		
^{76}Ge	15.42	3.62	38.51	13.1 – 20.8	1976Ca1
	18.69	13.81	82.19		
^{75}As	15.25	4.73	59.84	13.1 – 20.9	1969Be1
	18.16	6.73	59.05		
^{76}Se	15.86	6.50	150.72	13.1 – 19.7	1978Gur
^{78}Se	15.23	4.67	82.67	13.1 – 20.8	1976Ca1
	18.76	5.68	66.04		
^{80}Se	16.84	7.45	135.22	13.1 – 20.0	2016Va1
^{82}Se	16.13	5.81	175.20	13.1 – 19.9	1978Gur
$^{\text{nat}}\text{Rb}$	16.92	4.93	185.68	10.6 – 17.9	1971Lep
$^{\text{nat}}\text{Sr}$	16.99	5.03	201.18	10.9 – 17.9	1971Lep
^{89}Y	16.82	4.42	222.48	14.0 – 19.0	1971Le1

TABLE IV continued.

	E [MeV]	Γ [MeV]	σ [mb]	Range [MeV]	Ref.
^{90}Zr	16.90	4.13	251.98	14.9 – 18.5	2003Var
^{91}Zr	16.64	4.32	181.76	14.0 – 18.9	1967Be2
^{92}Zr	16.34	4.70	165.00	14.0 – 18.9	1967Be2
^{94}Zr	16.35	5.52	157.58	14.0 – 18.9	1967Be2
^{93}Nb	16.70	5.18	198.63	14.0 – 19.0	1971Le1
^{92}Mo	17.28	5.05	236.19	14.4 – 19.0	2003Var
^{94}Mo	16.73	6.04	186.31	9.6 – 18.9	1974Be3
^{96}Mo	16.42	6.52	182.00	13.2 – 17.0	1974Be3
^{98}Mo	15.96	6.17	186.76	13.2 – 18.9	1974Be3
^{100}Mo	16.02	8.44	167.02	12.1 – 20.0	1974Be3
^{103}Rh	16.59	8.44	187.85	13.1 – 19.0	2003Var
^{108}Pd	16.15	8.26	169.74	10.0 – 19.0	1969Dea
	18.09	1.65	45.85		
$^{\text{nat}}\text{Pd}$	16.24	7.63	195.79	10.2 – 17.8	1971Lep
^{107}Ag	16.05	7.51	179.36	9.5 – 19.0	1969Ish
^{109}Ag	13.74	3.81	90.32	13.1 – 19.0	1969Ish
	16.76	4.17	103.59		
$^{\text{nat}}\text{Ag}$	16.39	8.03	194.95	13.2 – 18.9	1971Lep
$^{\text{nat}}\text{Cd}$	16.03	6.69	222.13	10.2 – 17.8	1971Lep
^{115}In	15.91	6.00	242.23	13.2 – 17.8	1974Le1
^{112}Sn	15.82	5.87	254.14	10.9 – 18.0	1974Sor
^{114}Sn	16.08	6.74	238.62	13.1 – 18.0	1975Sor
^{116}Sn	15.69	5.29	267.45	13.1 – 17.9	1974Le1
^{117}Sn	15.77	5.29	254.71	13.2 – 17.8	1974Le1
^{118}Sn	15.55	5.02	276.03	13.1 – 17.9	1974Le1
^{119}Sn	15.65	5.09	248.89	13.0 – 17.9	1969Fu1
^{120}Sn	15.50	5.26	282.55	13.1 – 17.9	1974Le1
^{122}Sn	15.45	4.98	262.09	13.1 – 18.0	1975Sor
^{124}Sn	15.41	5.06	269.85	13.1 – 18.0	2003Var
$^{\text{nat}}\text{Sb}$	15.62	5.59	273.38	10.2 – 17.8	1971Lep
^{124}Te	15.36	5.81	276.72	12.0 – 18.9	1976Le2
^{126}Te	15.27	5.62	292.05	12.0 – 18.9	1976Le2
^{128}Te	15.23	5.55	300.47	12.0 – 18.9	1976Le2
^{130}Te	15.21	5.19	316.19	12.0 – 18.9	1976Le2
^{127}I	15.05	4.76	242.61	12.0 – 20.0	1999Bel
	17.33	1.83	22.56		
^{133}Cs	15.44	5.50	312.95	12.0 – 19.0	1974Le1
^{138}Ba	15.16	4.68	312.33	12.1 – 18.7	2016Va2
$^{\text{nat}}\text{Ba}$	15.46	5.57	344.80	10.1 – 17.8	1971Be4
^{139}La	15.30	5.11	360.94	12.0 – 18.9	1972De1
^{140}Ce	15.09	4.51	378.93	12.0 – 18.9	1976Le2
^{142}Ce	14.95	5.24	328.39	12.0 – 18.9	1976Le2
^{141}Pr	15.33	4.49	340.06	12.1 – 16.9	1987Ber
^{142}Nd	15.02	4.41	361.62	13.1 – 18.0	2003Var
^{143}Nd	15.08	4.99	341.49	12.0 – 19.0	1971Ca1
^{144}Nd	15.17	5.56	311.38	12.0 – 18.9	1971Ca1
^{145}Nd	15.14	6.75	289.80	12.0 – 18.9	1971Ca1
^{146}Nd	14.88	6.04	304.19	12.0 – 18.9	1971Ca1
^{148}Nd	13.34	5.41	156.93	10.8 – 18.6	1971Ca1
	15.79	4.57	164.60		
^{150}Nd	12.49	3.93	194.54	10.8 – 18.6	1971Ca1
	16.23	4.85	197.33		
^{144}Sm	15.37	4.53	379.27	12.1 – 18.9	1974Ca5
^{148}Sm	14.91	5.15	336.69	12.1 – 18.9	1974Ca5

TABLE IV continued.

	E [MeV]	Γ [MeV]	σ [mb]	Range [MeV]	Ref.
^{150}Sm	14.76	6.01	309.77	12.1 – 18.9	1974Ca5
^{152}Sm	12.56	3.53	202.99	10.9 – 18.8	1974Ca5
	15.97	4.77	206.96		
^{154}Sm	12.31	3.27	208.97	10.9 – 18.6	1981Gur
	15.95	5.53	186.51		
^{151}Eu	13.43	5.18	168.67	10.2 – 18.0	1971Vas
	14.52	2.28	158.70		
^{153}Eu	12.47	3.26	180.63	10.9 – 18.7	1969Be8
	16.07	5.48	198.64		
^{152}Gd	11.96	3.49	161.00	10.2 – 18.0	1971Vas
	14.75	2.87	234.24		
^{154}Gd	12.09	3.08	176.27	10.2 – 18.0	1971Vas
	15.10	3.09	227.94		
^{156}Gd	12.63	3.61	252.40	10.9 – 18.7	1981Gur
	15.97	4.10	188.83		
^{158}Gd	12.00	3.37	193.78	10.2 – 18.0	1971Vas
	15.19	2.97	231.63		
^{160}Gd	12.47	3.85	258.30	10.9 – 18.8	2003Var
	16.26	4.69	218.92		
^{159}Tb	12.55	3.23	205.41	11.1 – 19.0	1976Gor
	16.16	5.74	270.81		
^{165}Ho	12.47	2.84	238.17	11.1 – 18.7	1981Gur
	15.59	3.78	209.10		
^{168}Er	12.33	4.18	258.17	10.9 – 18.8	1981Gur
	15.65	3.58	222.84		
^{174}Yb	12.73	3.95	367.77	10.9 – 18.7	1981Gur
	15.80	3.18	249.27		
^{175}Lu	12.44	2.99	245.80	11.0 – 18.7	1969Be6
	15.65	4.32	260.78		
^{176}Hf	12.46	3.13	303.74	10.9 – 17.9	1977Gor
	15.86	4.44	247.79		
^{178}Hf	12.59	4.95	372.55	10.8 – 18.6	1981Gur
	15.65	3.00	218.91		
^{180}Hf	12.74	4.93	364.50	10.8 – 18.7	1981Gur
	15.59	3.04	220.92		
^{181}Ta	12.36	3.41	311.29	10.8 – 18.6	1981Gur
	15.26	4.71	273.41		
^{182}W	13.08	7.29	380.79	11.0 – 18.8	1981Gur
	15.20	1.53	99.76		
^{184}W	12.27	5.17	373.92	11.0 – 17.6	1981Gur
	15.10	3.15	186.15		
^{186}W	13.01	6.25	381.26	10.9 – 18.7	1981Gur
	14.81	2.84	122.86		
^{185}Re	12.89	3.48	329.91	10.2 – 18.0	1973Gor
	15.84	4.37	252.98		
$^{\text{nat}}\text{Re}$	14.30	7.05	462.90	10.2 – 18.0	1975Vey
^{186}Os	13.00	3.09	309.47	11.1 – 18.9	2015Var
	15.09	3.39	274.49		
^{188}Os	13.14	3.59	382.50	10.8 – 18.9	2014Var
	15.22	2.98	294.24		
^{189}Os	13.18	3.64	416.27	10.8 – 18.9	2014Var
	15.32	2.95	214.75		
^{190}Os	13.21	3.59	427.44	10.8 – 18.9	2015Var
	15.28	2.93	206.88		

TABLE IV continued.

	E [MeV]	Γ [MeV]	σ [mb]	Range [MeV]	Ref.
^{192}Os	13.08	3.61	365.35	10.8 – 18.9	2015Var
	14.98	3.68	249.26		
^{191}Ir	13.16	3.75	418.50	11.0 – 16.8	1978Go1
	15.24	3.69	170.79		
^{193}Ir	12.85	1.64	160.61	11.0 – 16.8	1978Go1
	14.18	5.94	393.43		
$^{\text{nat}}\text{Ir}$	13.81	4.97	492.98	10.2 – 18.0	1975Vey
^{194}Pt	13.66	4.29	499.56	11.0 – 17.8	1978Go1
	16.70	4.55	34.46		
^{195}Pt	13.28	3.71	464.49	11.0 – 17.8	1978Go1
	15.44	3.56	137.09		
^{196}Pt	13.38	2.87	235.00	11.0 – 17.8	1978Go1
	14.18	6.94	275.78		
^{198}Pt	13.62	4.83	533.11	11.0 – 17.8	1978Go1
^{197}Au	13.72	5.43	522.61	11.1 – 17.0	1981Gur
^{203}Tl	14.06	3.95	435.91	9.0 – 17.9	1970Ant
^{205}Tl	14.47	2.93	482.88	10.5 – 17.9	1970Ant
^{206}Pb	13.61	4.01	504.02	10.0 – 17.0	1964Ha2
^{207}Pb	13.57	4.22	467.03	10.0 – 17.0	1964Ha2
^{208}Pb	13.34	3.64	662.00	10.9 – 18.8	2003Var
$^{\text{nat}}\text{Pb}$	13.58	4.05	635.27	12.1 – 16.9	1985Ahr
^{209}Bi	13.87	5.04	591.22	10.9 – 18.3	1976Gu2
^{232}Th	11.10	3.51	285.95	10.2 – 18.3	1976Gu1
	13.97	4.22	334.57		
^{233}U	11.01	2.23	209.80	9.4 – 17.8	1986Be2
	13.68	5.38	403.97		
^{234}U	11.30	2.83	428.01	9.4 – 17.8	1986Be2
	14.23	4.04	353.66		
^{235}U	11.11	4.52	345.36	9.5 – 18.4	1976Gu1
	13.92	3.88	269.50		
^{236}U	11.25	3.34	340.25	9.5 – 17.8	1980Ca1
	14.18	4.25	351.15		
^{238}U	11.24	3.54	324.58	9.2 – 18.8	1976Gu1
	14.45	4.26	305.65		
$^{\text{nat}}\text{U}$	10.86	2.83	353.45	10.2 – 17.9	1985Ahr
	13.96	4.69	368.18		
^{237}Np	11.10	2.72	366.61	9.4 – 17.8	1986Be2
	14.27	4.27	498.40		
^{239}Pu	11.47	4.50	303.03	9.3 – 18.7	1976Gu1
	14.37	4.76	278.70		