



Nondestructive technique for identifying nuclides using neutron resonance transmission analysis at CSNS Back-n

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Received: 7 July 2023 / Revised: 24 November 2023 / Accepted: 7 December 2023

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Abstract

Nondestructive and noninvasive neutron assays are essential applications of neutron techniques. Neutron resonance transmission analysis (NRTA) is a powerful nondestructive method for investigating the elemental composition of an object. The back-streaming neutron line (Back-n) is a newly built time-of-flight facility at the China Spallation Neutron Source (CSNS) that provides neutrons in the eV to 300 MeV range. A feasibility study of the NRTA method for nuclide identification was conducted at the CSNS Back-n via two test experiments. The results demonstrate that it is feasible to identify different elements and isotopes in samples using the NRTA method at Back-n. This study reveals its potential future applications.

Keywords CSNS · White neutron beam · NRTA · Nuclide identification · Nondestructive method

1 Introduction

Neutrons are unique probes for analyzing the interior properties of materials in a nondestructive manner owing to their strong penetration power. Various neutron techniques based on the interactions between neutrons and materials have been successfully developed and applied in different fields. Neutron resonance analysis (NRA) is a technique that uses

neutron resonance absorption to identify and quantify elements and isotopes in a sample [1–3]. Its basic principle is that the absorption of neutrons by nuclei as a function of neutron energy shows specific resonance peaks and dips based on which elemental composition of the material can be determined [4]. Neutron resonance transmission analysis (NRTA) is an NRA method based on the transmission of a neutron beam through a sample, in which the resonances are observed as dips as a function of neutron energy.

The first demonstration of NRTA as a nondestructive method for determining the isotopic abundance of spent nuclear fuel pins was by Priesmeyer and Harz [3]. ^{235,238}U and ²³⁹Pu contents were obtained in their study. Subsequently, NRTA for isotopic assay was used as a nondestructive technique to characterize special nuclear materials [5–7]. The results obtained using NRTA were consistent

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This work was supported by the National Natural Science Foundation of China (No. 12035017), Youth Innovation Promotion Association CAS (No. 2023014), and Guangdong Basic and Applied Basic Research Foundation (Nos. 2020A1515010360 and 2022B1515120032).

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with those obtained using mass spectrometry and neutron activation methods [8]. NRTA has also been applied in other fields, such as the characterization of reference materials for nuclear data measurement [9] and cultural heritage objects and artifacts [10]. Moreover, NRTA has also been used with neutron imaging, known as neutron resonance transmission imaging, at the J-PARC facility [11], ISIS spallation neutron source [12], and KURRI-LINAC [13]. The NRTA method was investigated extensively by a group from EC-JRC Geel at the GELINA facility [14–16], which is a white neutron source that provides neutrons from 10 meV to 20 MeV [17]. A white neutron beam covering the energy range from eV to MeV has significant advantages over thermal neutron beams because a broad energy range covers almost all the resonance regions of the elements, whereas the thermal beam is mainly used for heavy elements. The back-streaming neutron line (Back-n) at China Spallation Neutron Source (CSNS) is a newly built white neutron beam covering the range from 0.5 eV to 300 MeV [18, 19]. The time-of-flight (TOF) technique was used at Back-n to accurately determine the neutron energy. The energy resolution at Back-n end station 2 (ES#2) was lower than ~2% for neutrons below 1 MeV. Because of these favorable beam properties, Back-n is suitable for the study and application of the NRTA technique.

This paper describes the first test experiments using the NRTA method at the CSNS Back-n facility. The experimental results for two samples containing light or medium elements are presented. Data analysis shows that element/isotope identification using the NRTA method is feasible at Back-n.

2 Experimental setup of NRTA at CSNS Back-n

Located in Dongguan, Guangdong, the multi-disciplinary CSNS facility has been operational since 2018. It generates neutrons via a spallation reaction by impinging 1.6 GeV protons onto a massive tungsten target. The nominal proton beam power of CSNS Phase I is 100 kW. In the forthcoming CSNS Phase II upgrading project, the beam power will be increased to 500 kW, and several more spectrometers [20, 21] and experimental stations [22] will be built. Although most neutron beamlines at CSNS deliver thermal neutrons for neutron scattering applications, Back-n beamline delivers neutrons within a very broad energy range (from 0.5 eV to 300 MeV) mainly for nuclear data measurements [19, 23, 24] and nuclear technology applications [25, 26].

Since its inception, the Back-n beam's time characteristics have been a focal point of investigation [27] due to their crucial role in determining neutron energy and influencing energy resolution. The energy resolution of a TOF facility, considering the relativistic effect, is calculated as

$$\frac{\Delta E}{E} = \gamma(\gamma + 1) \sqrt{\left(\frac{\Delta T}{T}\right)^2 + \left(\frac{\Delta L}{L}\right)^2} \quad (1)$$

$$\gamma = \frac{1}{\sqrt{1 - \left(\frac{v}{c}\right)^2}} \quad (2)$$

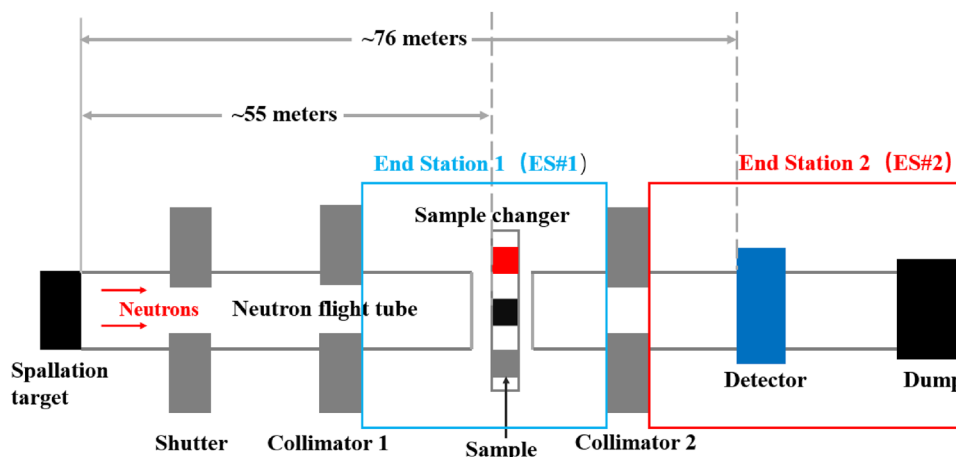
where γ is the Lorentz factor, v is the neutron velocity, and c is the speed of light. T and L are the neutron flight time and flight path length, respectively, and ΔT and ΔL are their respective uncertainties. ΔT is mostly from the incident proton pulse width, whose full-width at half-maximum was 60 ns during the measurements in this study. ΔL is caused by the neutron moderation and scattering in the spallation target before they enter the neutron tube. The moderation process has been investigated via Monte Carlo simulations [28, 29], and the uncertainties of the moderation distance (ΔL) at different energies were obtained. Table 1 lists the ΔL values at different energies and the energy resolutions calculated using Eq. (1), which indicates that the energy resolution is a function of the neutron energy.

In this study, we implemented NRTA measurements using a neutron total cross-section spectrometer. This spectrometer comprised a multilayer fast fission chamber (FIXM) and a sample changer [30]. The FIXM is a neutron detector that uses ^{235}U and ^{238}U samples as neutron converters. Figure 1 shows the typical setup for the neutron total cross-section measurements. The samples to be measured were placed on a sample changer at end station 1 (ES#1), and the detector was set up at ES#2. The beam spot at the sample position is typically turned to be $\Phi 60$ mm, whereas that at the detector position is typically configured to be approximately $\Phi 40$ mm. This configuration ensures that the neutron beam at ES#2 is entirely covered by a neutron converter with a diameter of $\Phi 50$ mm. Total cross-section measurements typically include sample-out and sample-in measurements. Sample-out measurements involve characterizing a neutron beam without any sample in the beam, whereas sample-in measurements

Table 1 Energy resolutions of CSNS Back-n at ES#2

E_n (eV)	ΔT (ns)	ΔL (cm)	$\Delta E_n/E_n$
1	60	12.2	3.14×10^{-3}
10	60	13.6	3.50×10^{-3}
10^2	60	24.0	6.18×10^{-3}
10^3	60	20.2	5.25×10^{-3}
10^4	60	18.0	5.10×10^{-3}
10^5	60	15.4	7.84×10^{-3}
10^6	60	8.8	2.15×10^{-2}
10^7	60	10.0	6.82×10^{-2}

Fig. 1 Layout for neutron total cross-section measurements



involve placing the sample in the beam at ES#1. The long distance (approximately 20 m) and the collimator between the sample and detector minimally suppress the multiscattering neutron background. The transmission spectrum is the ratio of the normalized neutron counts of the sample-in to the sample-out, from which the total cross section can be determined. The neutron total cross sections of ^{nat}C [30], ^{nat}Li [31], 9Be , ^{27}Al [32], ^{nat}Fe , ^{nat}Pb , ^{nat}Cr , and ^{209}Bi [33] have been measured using this setup at Back-n.

The NRTA experiment at Back-n utilized the same setup as that used for the total cross-section measurements. Photographs of the sample changer and neutron detector are shown in Fig. 2. The resonances were observed as absorption dips in the transmission spectrum as a function of the neutron energy, from which the nuclides in the sample could be identified because the resonances of each nuclide are as unique as fingerprints. The energy positions of the resonance dips in the transmission spectrum provide qualitative information about the composition, whereas the profiles of the resonance dips present quantitative details about the composition. In our first NRTA experiments, we resolved the qualitative information about the samples. Quantification of the elemental composition is more complicated and will be our subsequent task.

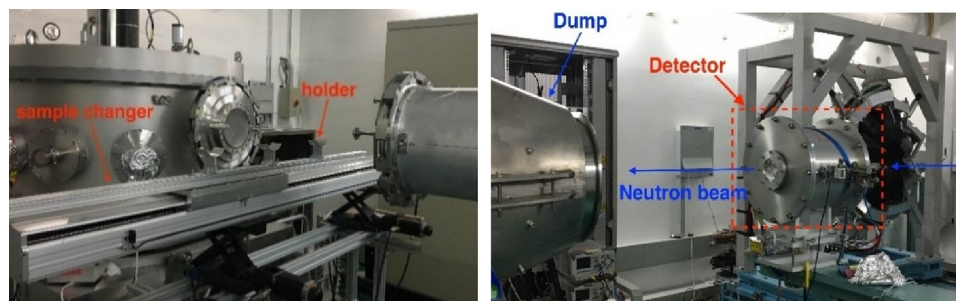
3 Results and discussion

The NRTA method is centered on measuring and analyzing neutron transmission spectra. The TOF technique was used to determine the neutron energy. Further details of the TOF method at Back-n can be found in our previous publications [18, 34]. Another crucial aspect of neutron energy determination is the double-bunch effect, in which two identical proton bunches have well-defined intervals (410 ns) in an incident pulse. Double-bunch unfolding is necessary for energy regions higher than 10 keV, as the 410 ns uncertainty is no longer negligible. An unfolding code based on Bayes' theorem was developed to unfold the TOF spectrum [35, 36]. The systematic uncertainty introduced by the unfolding process can be estimated using the unfolding code, which is highly dependent on statistical uncertainties. The uncertainty of the unfolding is typically 1.5–2.5 times that of the statistical uncertainty.

3.1 Experiment using imitated lunar soil

As shown in Fig. 3, an artificially simulated lunar soil sample was measured to verify the feasibility of the NRTA at Back-n. The sample was a cylinder with a diameter of 27 mm

Fig. 2 Sample changer at Back-n ES#1 (left) and detector at ES#2 (right)



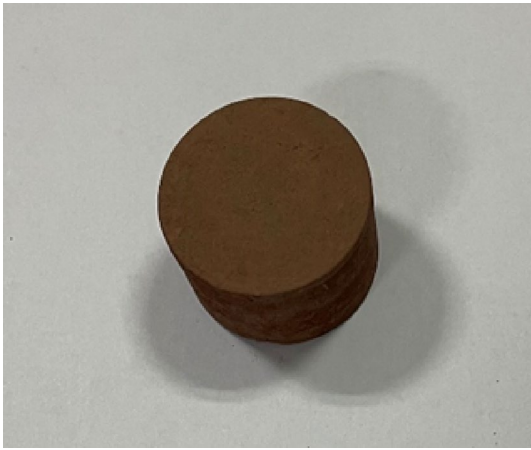


Fig. 3 Photograph of imitated lunar soil sample. (Color figure online)

and a height of 21 mm, and comprised uniformly distributed elements. The details of its composition are listed in Table 2.

Figure 4 compares the measured fission rate spectra of the sample-in and the sample-out. The sample-in spectrum (the red curve in Fig. 4) was attenuated by the scattering and absorption by the sample. Figure 5 shows the transmission of the neutron beam through the sample, which is the ratio of sample-in to sample-out measurements. Resonance absorption dips were observed in the transmission spectrum and could be used to determine the existing isotopes in the sample.

Figure 6 shows the transmission spectra in different energy regions to provide clear profiles of the absorption dips. All results are presented in 100 bins/decade (bpd), and the bin center is adopted as the energy of each bin. The resonances of sodium, aluminum, magnesium, iron, silicon, and oxygen were recognized. Table 3 presents a comparison of the measured resonance energy values and the corresponding resonance energy values in the ENDF/B-VIII.0 library [37]. The differences between the values of the measurements and library, and the energy resolution at each energy level are also presented. The differences at most points are comparable to the energy resolution, except at 88.11 keV, which is assumed to be the overlap of the resonances of ^{24}Mg and ^{56}Fe at 83.45 and 84.84 keV, respectively. The overlap of the two resonances may increase the deviation between the measured and evaluated data. The differences vary with energy because, as discussed in Sect. 2, the energy resolution is a function of the neutron energy. Binning is another reason for the differences in the resonance energy

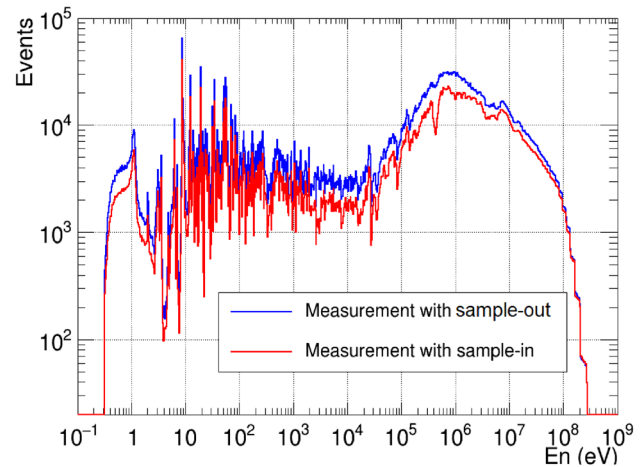


Fig. 4 Comparison of sample-out and sample-in measurements of imitated lunar soil

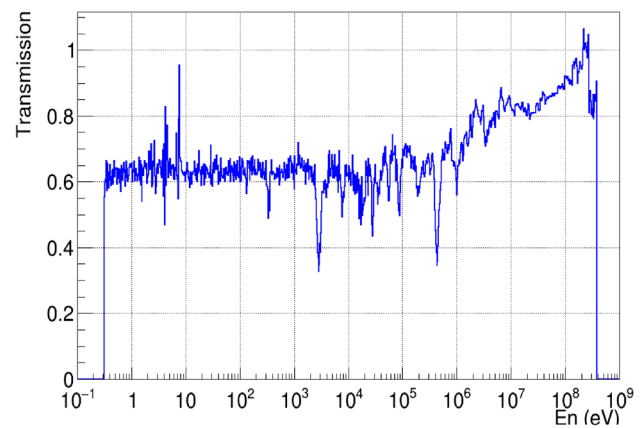


Fig. 5 Transmission spectrum of imitated lunar soil

positions between the measurement and library. The energy bin center depends on the bin width, and smaller energy binning should provide a more accurate resonance energy; however, it increases the statistical uncertainties. This issue can be addressed using long beam-time measurements in the future. In this study, a 100 bpd was used as the final option. In addition, although the unfolding method was used in a good order for the region higher than 10 keV, the double-bunch commission inevitably introduced systematic uncertainties.

Through qualitative analysis, most elements listed in Table 2 were identified, except for calcium, potassium, and titanium. Their absorption was not evident, probably

Table 2 Composition of imitated lunar soil sample

Composition	SiO ₂	Al ₂ O ₃	CaO	Fe ₂ O ₃	K ₂ O	MgO	Na ₂ O	TiO ₂
Mass fraction (%)	49.6	13.8	9.6	14.8	0.48	7.2	2.07	0.922

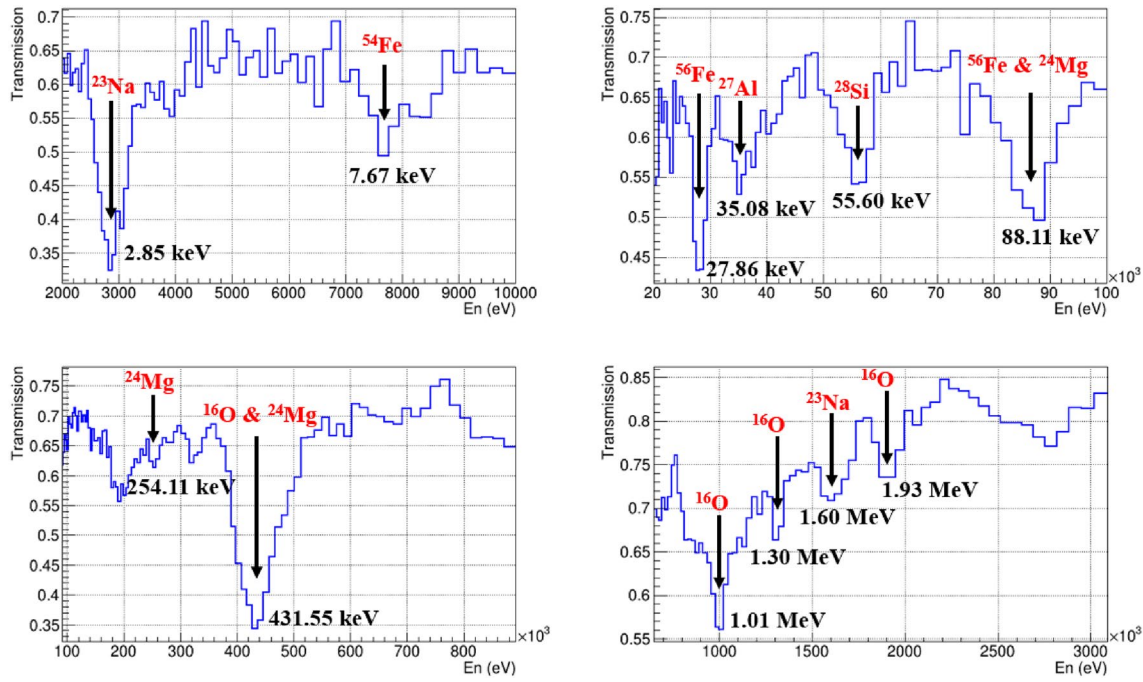


Fig. 6 Expanded transmission spectra of imitated lunar soil in different energy regions

Table 3 Resonance energy of imitated soil sample

Isotope	Resonance energy (keV)		Difference (%)	Energy resolution (%)
	Measurement	ENDF B-VIII.0		
²³ Na	2.85	2.81	1.4	0.63
²³ Na	1.60×10^3	1.60×10^3	0	2.4
²⁷ Al	35.08	35.06	0.06	0.59
²⁴ Mg	88.11	83.45	5.58	0.75
²⁴ Mg	254.11	257.19	1.20	1.02
²⁴ Mg	431.55	433.08	0.35	1.29
⁵⁴ Fe	7.67	7.82	1.96	0.80
⁵⁶ Fe	27.86	27.92	0.22	0.56
⁵⁶ Fe	88.11	84.84	3.85	0.75
²⁸ Si	55.60	55.80	0.36	0.65
¹⁶ O	431.55	434.00	0.57	1.29
¹⁶ O	1.01×10^3	1.00×10^3	1.0	2.19
¹⁶ O	1.30×10^3	1.31×10^3	0.77	2.31
¹⁶ O	1.93×10^3	1.90×10^3	1.58	2.63

because of the significant statistical fluctuations and their low abundance. In the above analysis, we only selected apparent dips and did not attempt to determine all the

resonances of each nuclide. This is feasible for a qualitative study of the sample.

3.2 Experiment using relic obtained from South China Sea

The NRTA at Back-n also investigated an underwater relic found in the South China Sea. Figure 7 shows a block of silicate and calcium carbonate embedded in many copper coins. It had an irregular shape, with an approximate size of 25 cm × 10 cm × 10 cm. The experimental setup was the same as that used to measure the imitated lunar soil. However, this object contained nonuniformly distributed elements and had an irregular shape. An associated experiment using a CMOS camera was also conducted to capture transmission images of the sample using the total transmission and resonance-selected methods [38].

Figure 8 displays the spectra for both sample-in and sample-out measurements. The sample-in measurements significantly attenuated because of the large sample size. Figure 9 shows the transmission of the neutron beam through the sample, from which the isotopes in the sample is examined. Figure 10 shows zoomed-in graphs of the transmission spectra in different energy regions. The resonances of copper, calcium, sodium, silicon, and iron were observed. These recognized isotopes were consistent with our expectations. ⁶³Cu and ⁶⁵Cu originated from the coins in the object. ²⁸Si, ⁴⁰Ca, ⁵⁶Fe, and ¹⁶O originated from silicate and calcium carbonate,

Fig. 7 Photographs of underwater relic found in South China Sea. (Color figure online)

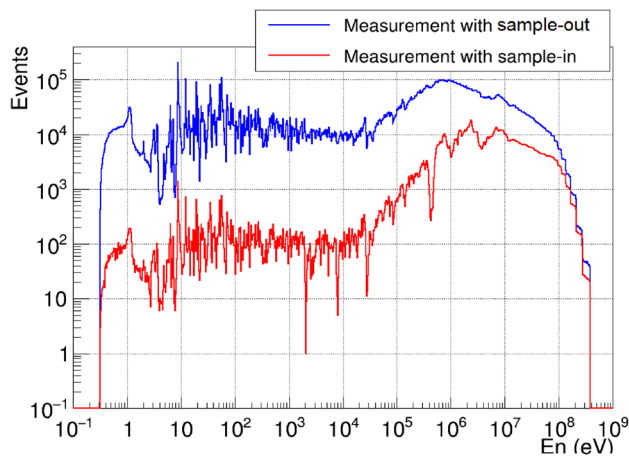
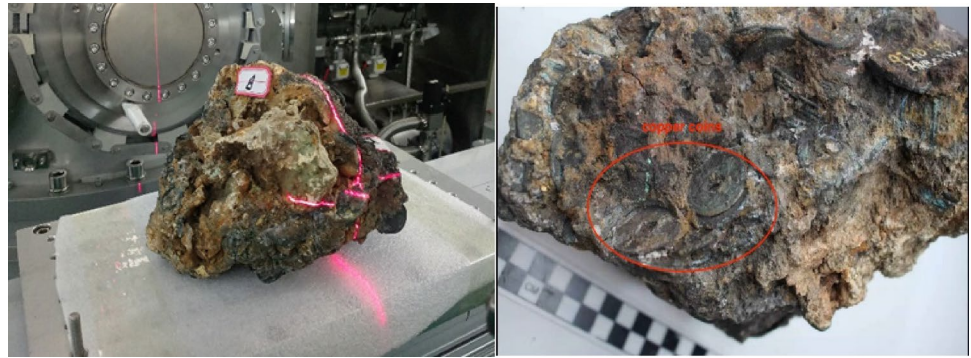


Fig. 8 Comparison of sample-out and sample-in measurement results of underwater relic

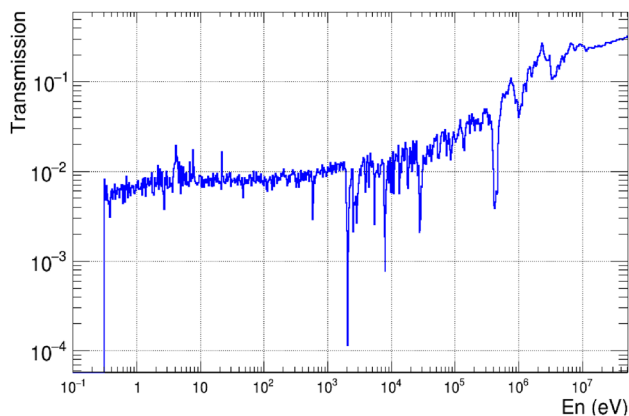


Fig. 9 Transmission spectrum of underwater relic

which are the main components of the object. The presence of ^{23}Na is normal because sodium is abundant in the sea.

We compared the measured resonance energy positions with corresponding values in the ENDF/B-VIII.0 library (Table 4). The energy binning was 100 bpd. The differences in the resonance energy positions between the measurement and the library were comparable to the energy resolution. In addition to the reasons mentioned in Sect. 3.1 for such differences, the irregular shape and massive volume of the sample could also increase the deviation between the measured and library values.

4 Conclusions and prospect

In 2018, CSNS introduced a new white neutron beamline, Back-n. Its beam characteristics (high neutron flux, broad energy range, and good energy resolution) motivated us to launch a feasibility study of NRTA for nuclide identification. Experiments on two different samples, an imitated lunar soil sample and an underwater relic, were performed for the first time at Back-n. The existence of ^{16}O , ^{28}Si , ^{27}Al , $^{54,56}\text{Fe}$, ^{24}Mg , and ^{23}Na in the simulated lunar soil sample and $^{63,65}\text{Cu}$, ^{16}O , ^{28}Si , ^{56}Fe , ^{40}Ca , and ^{23}Na in the underwater relic were confirmed using the NRTA method. A simple qualitative analysis demonstrated that CSNS Back-n could identify nuclides using the NRTA method. The beam characterization and results of this study demonstrate that the CSNS Back-n is a promising platform for nuclide identification.

FIXM serves as the primary neutron detector for transmission measurements at Back-n. It helps in monitoring the flux based on $^{235}\text{U}(n, f)$ and $^{238}\text{U}(n, f)$ reactions. However, the resonances of $^{235}\text{U}(n, f)$ cross section below

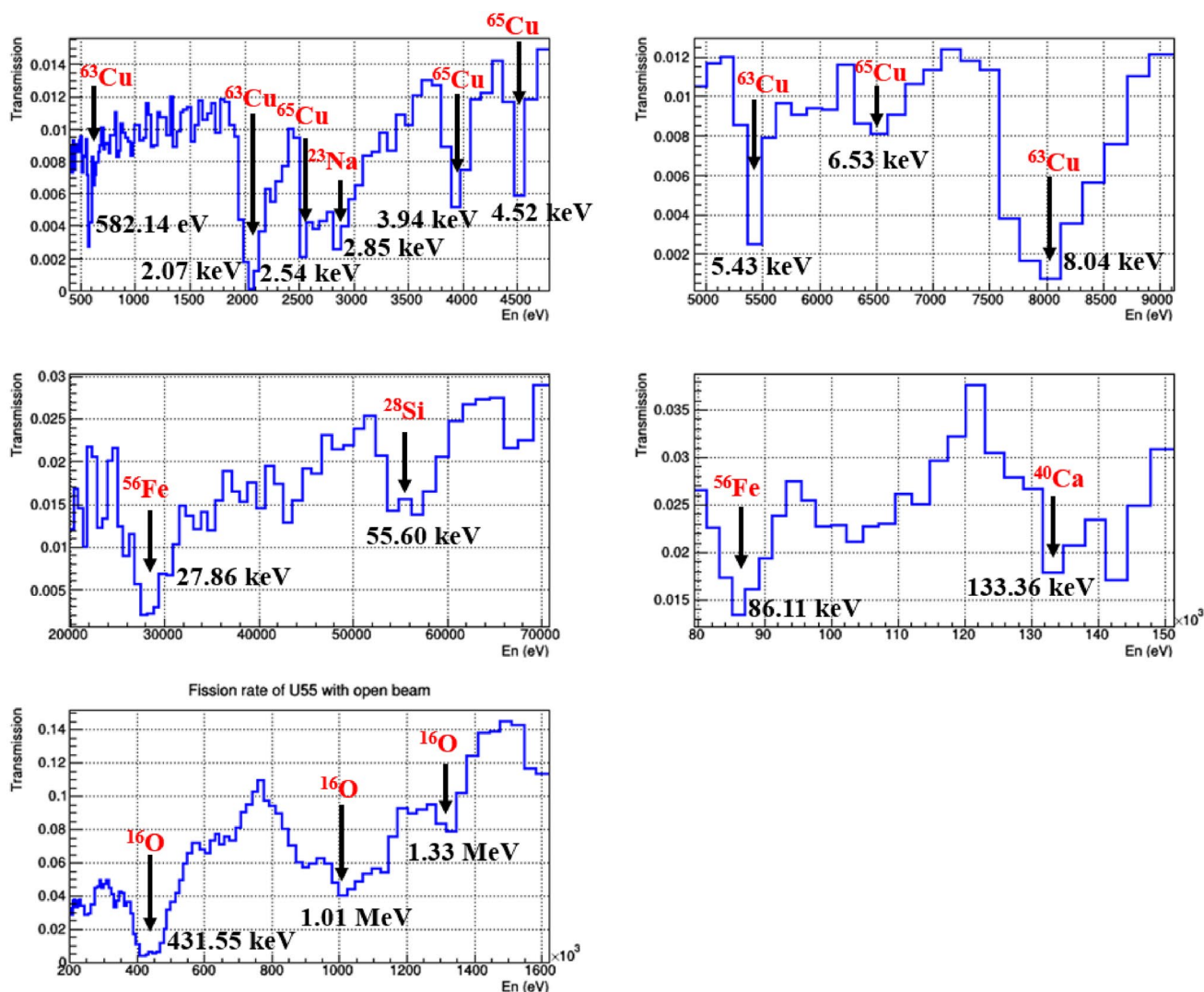


Fig. 10 Expanded transmission spectra of underwater relic in different energy regions

the ~ keV region are somewhat intense, which might induce fluctuations in the transmission spectrum. Lithium glass can be a valuable compensation for the fission chamber in the low-energy region because of the large and smooth cross section of the $^6\text{Li}(n, t)$ reaction [39]. A lithium glass monitor is planned for future transmission measurements at Back-n, which is characterized by its quick response, high efficiency, and smooth transmission spectrum. However, because the lithium glass scintillator is sensitive to γ -rays,

n/γ discrimination [40] and γ -rays background must be investigated thoroughly.

In this paper, we present a simple qualitative analysis focused on confirming the presence of specific isotopes. Accurate quantification analysis for resolving the elemental abundance in a sample is significantly more complicated. Quantification analysis is feasible by performing sophisticated resonance analysis, which must include the

Table 4 Resonance energy of imitated soil sample

Isotope	Resonance energy (keV)		Difference (%)	Energy resolution (%)
	Measurement	ENDF B-VIII.0		
²⁸ Si	55.60	55.80	0.36	0.65
²³ Na	2.85	2.81	1.4	0.63
⁵⁶ Fe	27.86	27.92	0.21	0.56
⁵⁶ Fe	86.11	84.84	1.5	0.74
⁶³ Cu	0.582	0.578	0.7	0.59
⁶³ Cu	2.07	2.05	0.98	0.61
⁶³ Cu	5.43	5.38	0.93	0.72
⁶³ Cu	8.04	7.95	1.13	0.80
⁶⁵ Cu	2.54	2.54	0	0.62
⁶⁵ Cu	3.94	3.92	0.5	0.67
⁶⁵ Cu	4.52	4.49	0.67	0.69
⁶⁵ Cu	6.53	6.45	1.2	0.75
⁴⁰ Ca	133.36	133.02	0.23	0.83
¹⁶ O	431.55	434.00	0.57	1.29
¹⁶ O	1.01 × 10 ³	1.00 × 10 ³	1.0	2.19
¹⁶ O	1.33 × 10 ³	1.31 × 10 ³	1.5	2.32

experimental effects, resolution function, sample properties, and detector characteristics. This is the subject of our subsequent study.

Author contribution All authors contributed to the study conception and design. Material preparation, data collection and analysis were performed by Sheng-Da Tang, Yong-Hao Chen, Jing-Yu Tang, Rui-Rui Fan, Qiang Li, Gong Li, Dong Liu, Zheng-Yao Jin, Xing-Zhu Cui, Tian-Xiang Chen, Yi-Wei Yang, Rong Liu, Han Yi, Yang Li and Zhen Yang. The first draft of the manuscript was written by YHC and all authors commented on previous versions of the manuscript. All authors read and approved the final manuscript.

Data availability The data that support the findings of this study are openly available in Science Data Bank at <https://cstr.cn/31253.11.sciencedb.j00186.00352> and <https://www.doi.org/https://doi.org/10.57760/sciencedb.j00186.00352>.

Declarations

Conflict of interest Jing-Yu Tang is an editorial board member for Nuclear Science and Techniques and was not involved in the editorial review, or the decision to publish this article. All authors declare that there are no competing interests.

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