

# Current status of multilayer neutron interferometry with gaseous samples at J-PARC

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Several few-body nuclear models, such as the Argonne  $v_{18}$  potential including three-nucleon forces and chiral effective field theories, predict the coherent scattering length ( $b_c$ ) for light nuclei. Reliable measurements of  $b_c$  therefore provide important tests of these models. We have developed a measurement system for determining  $b_c$  of gaseous samples using a multilayer-type neutron interferometer at J-PARC, and successfully observed the phase shifts of  $^3\text{He}$  and  $^4\text{He}$  gases. From the observed phase shifts, we obtained for  $^3\text{He}$   $b_c = 3.99 \pm 0.23$  (stat.)  $\pm 0.66$  (sys.) fm and for  $^4\text{He}$   $b_c = 2.95 \pm 0.11$  (stat.)  $\pm 0.67$  (sys.) fm (stat. and sys. denote statistical uncertainty and systematic uncertainty, respectively). To achieve higher precision, we are currently upgrading the experimental setup.

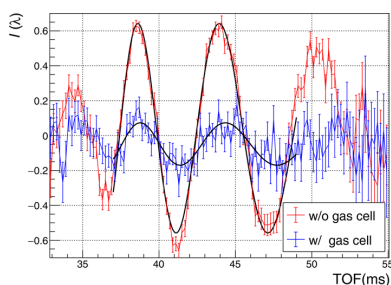
## 1. Introduction

The coherent scattering length ( $b_c$ ) describes neutron s-wave scattering and is a key parameter for understanding few-body nuclear systems. Few-body nuclei such as  $^3\text{He}$  and  $^4\text{He}$  are described by several phenomenological models, including the Argonne  $v_{18}$  (AV18) potential with three-nucleon interactions (3N), the  $R$  matrix approach (Hofmann & Hale, 2003) and chiral effective field theories (Bagnarol *et al.*, 2023), which predict  $b_c$ .

Therefore, measurements of the  $b_c$  of  $^3\text{He}$  and  $^4\text{He}$  (denoted hereafter as  $b_c^{^3\text{He}}$  and  $b_c^{^4\text{He}}$ ) can serve as benchmarks for studies of few-body nuclear systems.

Experimentally,  $b_c$  values have been measured using several methods, including scattering cross-section measurements (Haddock *et al.*, 2019), reflectometry (Kitchens *et al.*, 1974), diffraction (Shull & Shaw, 1973) and, most prominently, neutron interferometry (Ioffe *et al.*, 1998). For instance, the  $b_c$  values of H and D, obtained using neutron interferometers (NIs) with relative precisions of  $3.0 \times 10^{-4}$  (Koester & Nistler, 1975) and  $4.5 \times 10^{-4}$  (Schoen *et al.*, 2003), respectively, are consistent with the predictions of AV18 and AV18 with 3N. However, relative discrepancies of approximately  $2 \times 10^{-2}$  for  $b_c^{^3\text{He}}$  and  $7 \times 10^{-3}$  for  $b_c^{^4\text{He}}$  persist among the latest measurements using Si crystal NIs.

We aim to resolve these discrepancies with a relative precision of  $10^{-3}$  using pulsed neutrons and a multilayer-type NI. While  $b_c$  for solid samples has already been measured



using a multilayer-type NI, this paper reports its first application to gaseous samples with a newly fabricated gas cell. We present the results as a proof of principle for this method and discuss developments towards higher-precision experiments.

## 2. Experiment

A neutron interferometer splits a neutron wave into two paths and recombines them to produce interference. The potential difference between the two paths appears as a phase shift in the interference fringes. Since the first implementation in 1974 (Rauch *et al.*, 1974), NIs have found applications across diverse domains of physics (Colella *et al.*, 1975; Rauch *et al.*, 1975; Hasegawa *et al.*, 2003).

The phase shift  $\Delta\phi$  is described by

$$\Delta\phi = \frac{m\lambda L}{h^2} \Delta E, \quad (1)$$

where  $m$  is the neutron mass,  $\lambda$  is the neutron wavelength,  $L$  is the interaction length with the potential,  $\Delta E$  is the potential difference between the two paths and  $h$  is Planck's constant. The sensitivity of an NI is proportional to  $\lambda$  and  $L$ . Consequently, extending the path length of an NI and using a long wavelength enhances its sensitivity. A conventional NI made from an Si single crystal relies on Bragg scattering, which prevents its operation at  $\lambda$  longer than the Si lattice spacing. The interaction length  $L$  and the size of the NI are constrained by the dimensions over which high crystallinity can be maintained.

### 2.1. Multilayer neutron interferometer

We developed an NI using multilayer neutron mirrors. A multilayer neutron mirror consists of periodic layers of two materials (Ni/Ti) with different Fermi pseudo-potentials for neutrons. We can utilize cold neutrons which satisfy the reflection condition. Because the layer spacing is larger than the lattice spacing of an Si single crystal, neutrons with longer  $\lambda$  can be utilized than in a conventional NI. For the application of multilayer mirrors to a Jamin-type NI (shown in Fig. 1), we used a pair of beam-splitting etalons (BSEs) (Kitaguchi *et al.*, 2003; Seki *et al.*, 2010). Each BSE is composed of two multi-

layer mirrors on SiO<sub>2</sub> substrates, separated by an air gap of 211  $\mu\text{m}$ .

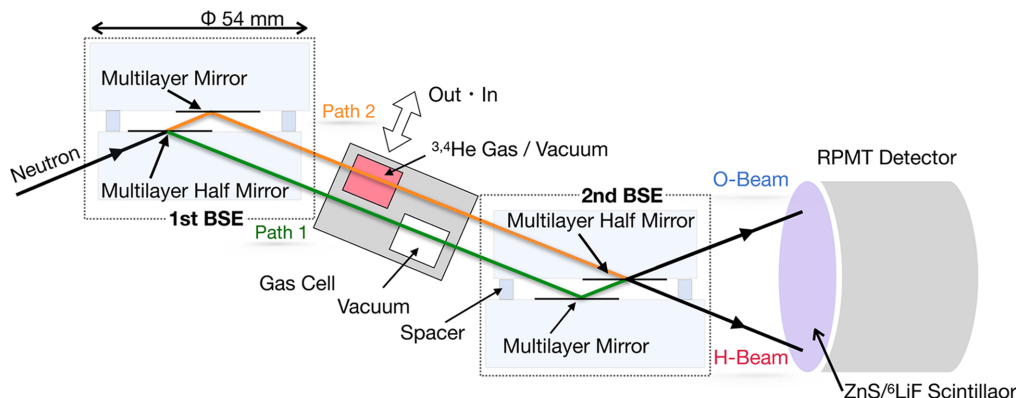
The NI with BSEs not only allows the extension of  $L$  but also enables the insertion of large samples that would be difficult to accommodate in a conventional NI, expanding the flexibility of the experimental setup. In the present measurement, a gas cell with a total width of 40 mm was inserted between the two paths, providing  $L = 8$  mm. The value of  $L$  is potentially scalable up to 1000 mm.

### 2.2. Experimental setup

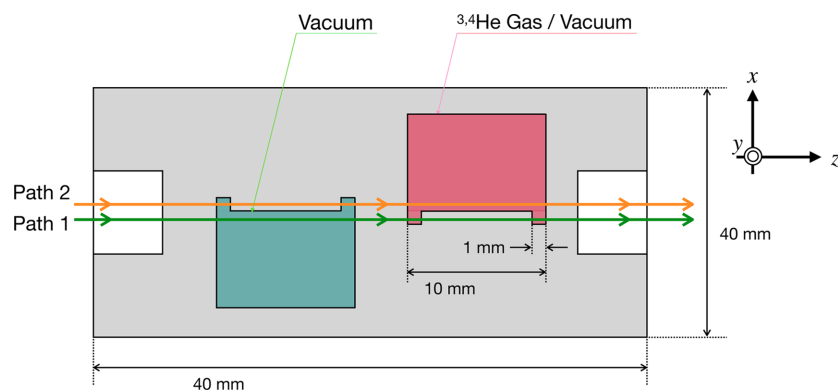
We conducted experiments on the low-divergence branch of the J-PARC beamline MLF BL05 (NOP) (Mishima *et al.*, 2009). The neutron wavelength used by a multilayer-type NI is determined by the time-of-flight (TOF) method, reducing time-varying disturbances in the analysis process. The multilayer-type NI and the sample insertion assembly were placed on a vibration isolation table inside a thermostatic chamber. To ensure the stability and alignment of the mirror substrates, the rotation of the entire NI and the relative angles between the two BSEs were precisely controlled and maintained using high-precision stepping motors. We observed an interferogram using pulsed neutrons (Fujie *et al.*, 2024). The bandwidth of  $\lambda$  is from 0.9 to 1.1 nm. The phase determination accuracy was comparable to that of a conventional NI. In our previous work, we had already measured  $b_c$  for several nuclei with solid samples and confirmed that these values were consistent with previous studies, except for V (Fujie *et al.*, 2024).

### 2.3. Gas cell

For measuring the  $b_c$  of gaseous samples, we designed a gas cell (shown in Fig. 2) which was machined from pure aluminium alloy (A1050) at RIKEN. In  $b_c$  measurements using an NI, one path passes through the sample while the other passes through a vacuum, thereby producing the phase shift. For a conventional NI, the centimetre-scale path separation allows the use of a two-chamber gas cell separated by a partition. However, in a multilayer-type NI, the beam separation is only 380  $\mu\text{m}$ , so neutrons pass through the corner radius formed during fabrication, making it difficult to determine  $L$  through the gas region accurately.



**Figure 1**  
Experimental setup of the Jamin-type NI employing beam-splitting etalons (BSEs).


**Figure 2**

Cross-section view of the gas cell. The red region contains the gaseous sample, while the green region is evacuated.

To address this problem, recess machining was performed to create surfaces perpendicular to the neutron path. By creating a 1 mm extension in the  $x$ -axis direction, neutrons are directed perpendicularly onto the gaseous sample. Neutrons pass through the red region, 2 mm in Path 1 and 10 mm in Path 2; therefore the effective  $L$  contributing to the phase shift is 8 mm.

#### 2.4. Experiment procedure and analysis

Measurements were repeated with and without the sample to minimize disturbances, as illustrated in Fig. 1, with measurements taken every 10 min. We used a position-sensitive detector with time resolution, consisting of a resistance-division photomultiplier tube (RPMT) and a ZnS<sup>6</sup>LiF scintillator. The detector was 0.3 m away from the interferometer setup (shown in Fig. 1). The details are described by Hirota *et al.* (2005).

The sample gases were filled at  $50256 \pm 6$  Pa at  $299.497 \pm 0.06$  K for  $^3\text{He}$  and at  $100017 \pm 10$  Pa at  $299.196 \pm 0.06$  K for  $^4\text{He}$ , measured by a piezoresistive transducer and a platinum resistance thermometer. The gas cell was evacuated by a turbo molecular pump to  $3.00 \times 10^{-3}$  Pa before filling, and the residual vacuum pressures before filling the gases were  $1.00 \times 10^{-2}$  Pa for  $^3\text{He}$  and  $1.17 \times 10^{-2}$  Pa for  $^4\text{He}$ . Table 1 sum-

marizes the assessment of sample contamination. The levels were below the experimental uncertainty and therefore negligible for these measurements.

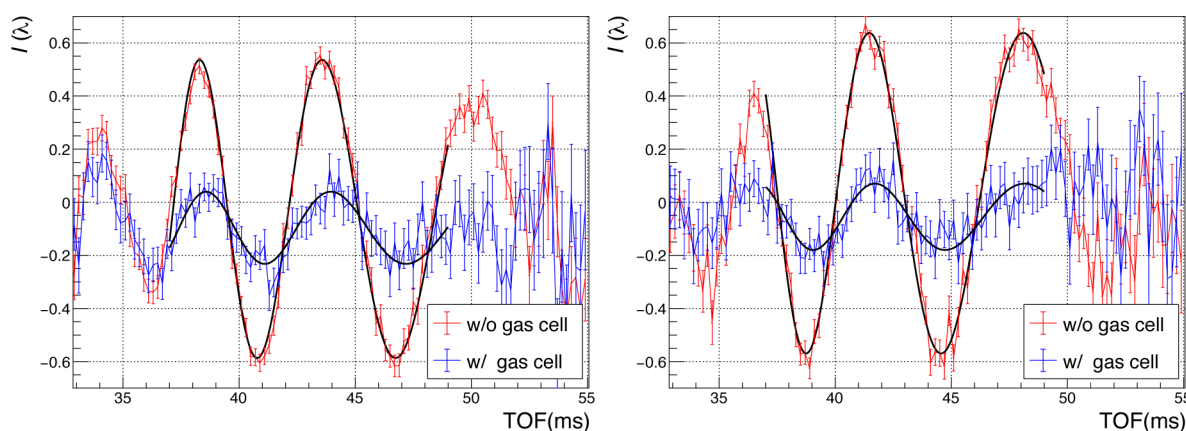
Figs. 3 and 4 show the interference fringes obtained over 10 min under conditions with the gas cell filled with  $^3\text{He}$ ,  $^4\text{He}$  and evacuated, respectively, and with the gas cell inserted and removed from the beam path. In Figs. 3 and 4, the vertical axis  $I(\lambda)$  is defined by

$$I(\lambda) = \frac{I_{\text{O}}/I_{\text{Ocd}} - I_{\text{H}}/I_{\text{Hcd}}}{I_{\text{O}}/I_{\text{Ocd}} + I_{\text{H}}/I_{\text{Hcd}}}, \quad (2)$$

where  $I_{\text{H}}$  and  $I_{\text{O}}$  are the intensities of the O and H beams, respectively, and  $I_{\text{Hcd}}$  and  $I_{\text{Ocd}}$  are those with a Cd block inserted. We fitted the obtained interference fringes using the fitting function

$$I(\lambda) = A \cos\left(\frac{P_{\text{L}}}{\lambda} + P_{\text{R}}\lambda - P_{\text{S}}\lambda\right) + B, \quad (3)$$

where  $A$  denotes the contrast of the interferogram. The coefficients  $P_{\text{L}}$  and  $P_{\text{R}}$  represent the inverse proportional and proportional terms, respectively, and they are derived by geometric optics (Fujiie *et al.*, 2024). Specifically,  $P_{\text{R}}$  accounts for the phase shift contributed by the SiO<sub>2</sub> substrate. The third term,  $P_{\text{S}} = Nb_{\text{c}}t$ , is the interaction with the sample, where  $N$  is


**Figure 3**

Interference fringes for (left)  $^3\text{He}$  and (right)  $^4\text{He}$ . Red and blue lines correspond to data with the cell out and in, respectively. Black lines indicate fits using equation (3).

**Table 1**  
Impurity list for samples.

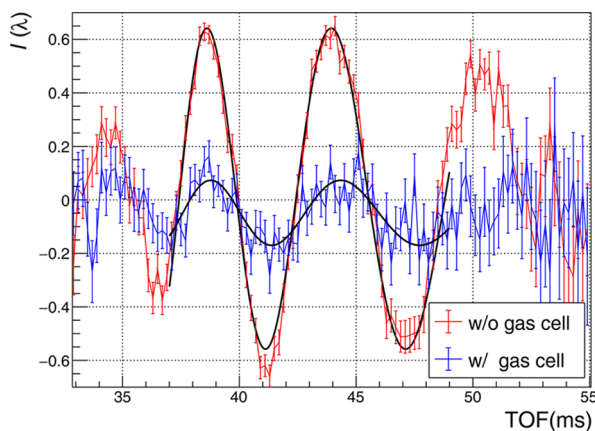
Sample	Impurity level
<sup>3</sup> He	<10 <sup>-3</sup>
<sup>4</sup> He	<10 <sup>-6</sup>

the number density of atoms and *t* is the sample thickness. The TOF region used for the fit was set from 37 to 49 ms, which corresponds to the reflecting momentum transfer range of the multilayer mirror, 0.232 < *Q* < 0.292 nm<sup>-1</sup>. Through relative measurements, *P<sub>S</sub>* was isolated as the difference between the wavelength-proportional terms with and without the sample. Scattering lengths were calculated from the phase shifts of each sample (shown in Fig. 3) after subtracting the value obtained in the vacuum measurement (shown in Fig. 4).

The extracted phase shifts were Δφ<sup>3He</sup> = 0.38 ± 0.022 (stat.) rad, Δφ<sup>4He</sup> = 0.57 ± 0.022 (stat.) rad and Δφ<sup>vac</sup> = 0.19 ± 0.015 (stat.) rad (where stat. means statistical uncertainty). Consequently, the results were *b<sub>c</sub><sup>3He</sup>* = 3.99 ± 0.23 (stat.) fm and *b<sub>c</sub><sup>4He</sup>* = 2.95 ± 0.11 (stat.) fm, with measurement times of 6 and 8 h, respectively. The relatively large statistical uncertainties were caused by a decrease in fringe contrast. The decrease in contrast caused by the insertion of the cell was attributed to small-angle scattering of neutrons by the 18 mm-thick (in total) aluminium wall (shown in Fig. 2).

**2.5. Uncertainties**

The systematic uncertainties are summarized in Table 2. The dominant systematic uncertainty is the phase distribution in the beam cross section, which is described below. In the condition with no cell, the phase of the interferogram in the TOF varied in the vertical position on the detector, indicating that the neutron phase was distributed along the *y* axis. We suspect that this effect was caused by a misalignment of the BSEs. We divided the data into three regions along the *y* axis and analysed each region to derive phase shifts, but the results were inconsistent. The systematic uncertainties arise from the weighted average taken for the three data points, expanding



**Figure 4**  
Interference fringes for the double-path vacuum measurement. Red and blue lines correspond to data with the cell out and in, respectively. Black lines indicate fits using equation (3).

**Table 2**  
A summary of systematic uncertainties for *b<sub>c</sub>*.

	Δ <i>b<sub>c</sub><sup>3He</sup></i> (fm)	Δ <i>b<sub>c</sub><sup>4He</sup></i> (fm)
Temperature	±0.00079	±0.00059
Pressure	±0.00063	±0.00023
Phase distribution	±0.66	±0.67

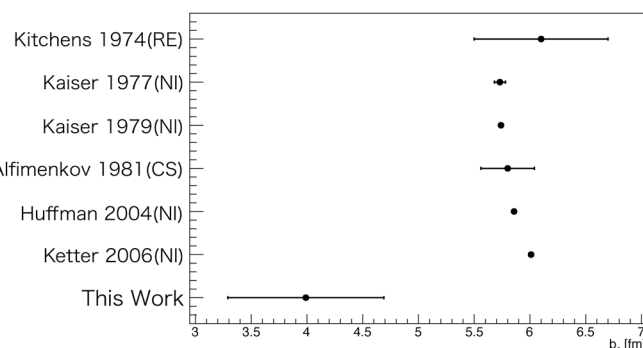
the error bar such that χ<sup>2</sup>/ndf = 1, where χ<sup>2</sup> is the chi-squared parameter and ndf the number of degrees of freedom.

**3. Results**

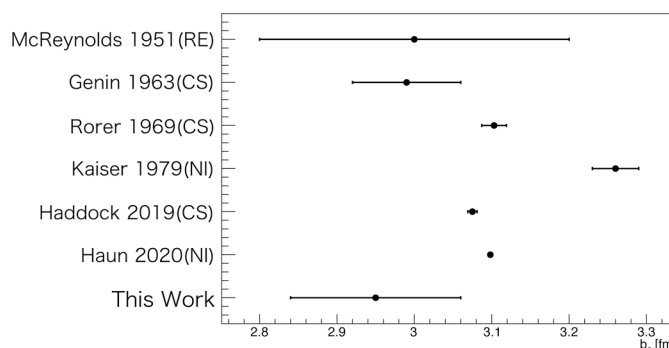
Figs. 5 and 6 compare our results with previous work (Kitchens *et al.*, 1974; Kaiser *et al.*, 1977; Kaiser *et al.*, 1979; Alfimenkov *et al.*, 1981; Huffman *et al.*, 2004; Ketter *et al.*, 2006; McReynolds, 1951; Genin *et al.*, 1963; Rorer *et al.*, 1969; Haddock *et al.*, 2019; Haun *et al.*, 2020). The methods used were cross section (CS), reflectivity (RE) and neutron interferometer (NI). Our results were *b<sub>c</sub><sup>3He</sup>* = 3.99 ± 0.23 (stat.) ± 0.66 (sys.) fm and *b<sub>c</sub><sup>4He</sup>* = 2.95 ± 0.11 (stat.) ± 0.67 (sys.) fm (where sys. means systematic uncertainty). In this first gaseous sample measurement, a significant phase shift was successfully observed, and several issues in the experimental setup for future measurements were identified.

**4. Discussion**

The experimental results revealed a vertical phase distribution and temporal instability, leading to reduced visibility and an



**Figure 5**  
Comparison of measured *b<sub>c</sub><sup>3He</sup>* with previous results.



**Figure 6**  
Comparison of measured *b<sub>c</sub><sup>4He</sup>* with previous results.

underestimation of the phase shift. We recently identified that this distribution stems primarily from the roll angle misalignment of the BSEs. Because these systematic effects have not yet been fully quantified, the actual uncertainty in these preliminary results is likely to be larger than that reported above. Furthermore, small-angle scattering from the aluminium degraded the contrast. To address these issues, we are upgrading the setup with high-rigidity stages and low-thermal-expansion materials.

## 5. Future prospects

To overcome the current limitations, we are preparing a new cell using SiO<sub>2</sub> crystals to eliminate contrast degradation from small-angle scattering. We will also correct the BSE roll angle misalignment by making adjustments, and improve the stability by introducing more rigid components. Furthermore, introducing supermirrors optimized for the J-PARC MLF BL05 spectrum is expected to increase the neutron flux by a factor of 20. With these upgrades,  $b_c^{3\text{He}}$  and  $b_c^{4\text{He}}$  can be measured with a relative precision of  $10^{-3}$  in 6 and 20 h, respectively.

## 6. Summary

To study the few-body system in nuclei, we performed measurements of  $b_c^{3\text{He}}$  and  $b_c^{4\text{He}}$  using a multilayer-type NI at J-PARC. The precision of  $b_c$  was limited by statistical uncertainties due to the small-angle scattering from aluminium and setup-related systematic uncertainties. By integrating SiO<sub>2</sub> cells, a high-rigidity stage and supermirror upgrades, we aim to achieve higher-accuracy measurements.

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## Conflict of interest

The authors declare that there are no conflicts of interest.

## Data availability

The data are available from the authors upon reasonable request.

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