

DIRECT MEASUREMENT OF THE ¹²C+¹²C FUSION REACTION X.D. Tang^{1,2}

On behalf of the Carbon Fusion Experiment (CARFUSE) Collaboration

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CARbon FUSion Experiment (CARFUSE) @LEAF, IMP





RESONANCES IN C¹² ON CARBON REACTIONS

E. Almqvist, D. A. Bromley, and J. A. Kuehner Atomic Energy of Canada Limited, Chalk River Laboratories, Chalk River, Ontario, Canada (Received March 28, 1960)



The world's first tandem accelerator installed at Chalk River in 1959.



Molecular resonances in the ${}^{12}C+{}^{12}C$ fusion reaction measured by Almqvist et al., in 1960

Carbon burning in the universe

Nucleosynthesis in massive stars



Ignition conditions in type Ia supernovae





Uncertain cross section at stellar energies



- Large difference between
 THM and Hindrance→Highly uncertain rate
- INDIRECT: Corrected THM exhibits a trend similar to Hindrance by replacing PWIA with DWIA
- Unknow resonances: Need better selection T=0, J^π=0⁺,2⁺

 $^{12}C(^{12}C,p)^{23}Na$ (Q=2.24 MeV) $^{12}C(^{12}C,\alpha)^{20}Ne$ (Q=4.62 MeV) $^{12}C(^{12}C,n)^{23}Mg$ (Q=-2.62MeV)

Beck, Mukhamedzhanov and Tang, Eur. Phys. J. A (2020) 56:87 Mukhamedzhanov, Eur. Phys. J. A (2022) 58:71 Tang & Ru, EPJ Web of Conferences 260, 01002 (2022)

$$S^{*}(E) = \sigma E e^{(87.21/\sqrt{E} + 0.46E)}$$



I

Uncertainty in the reaction rate









Direct Measurement of ¹²C(¹²C,a_{0,1})²⁰Ne

Particle- γ coincidence at lower stellar energies



Particle- γ coincidence technique pushed the measurement down to sub-nb level Only detect p₁ and α_1 channels

Carbon fusion project at LUNA-MV

Massive lead shield and radon flushing \rightarrow push sensitivity to better than 100 reactions/day



$^{12}\mathrm{C}{+}^{12}\mathrm{C}$ - γ measurements



A. Best (SF III)

Direct measurement with charged particles



High Intensity+Time Projection Chamber



LINAC: High Intensity beam up to few hundreds of puA
 TPC: Ultra sensitive tracking detector
 Complementary to LUNA-MV and other experiments

Z.C.Zhang+ NIMA(2021) Doi: 10.1016/j.nima.2021.165740



- LEAF: 45GHz and 14.5GHz ECRs; 0.5MeV/u RFQ; 0.3MeV/u-0.7MeV/u DTL; Energy calibration by ¹²C(p,g), p(15N,a)12C*
- Maximum carbon beam intensity on target: ~200 puA (Spillane: 40puA); Energy Spread: <0.2%(s); Energy step size:~50 keV; beam time: ~1150 hr</p>
 Courtesy of Yao Yang (IMP)

Detector setup

Time Projection Chamber (TPC) + Si array - Particle identification



X-ray and other beam induced background







Using MicroMegas gas to control the Ion Back Flow(IBF)
 Alpha: Ar(5%)+CO2(5%)+He(90%); Proton: Ar(35%)+CO2(15%)+Kr(50%)

High Power Carbon with High Purity



- > Improving thermal conductivity using flexible graphene
- Carbon targets: Diamond like carbon(DLC), Heated graphite (5N), Highly Ordered Pyrolytic Graphite High (HOPG)

Analysis: E_{cm}=2.72MeV with Graphite (5N)



Clear identification with cuts in the energy loss and tracks in TPC

Accepted by NST

Degrading of the HOPG target

Wang et al., (2024), Tan et al., (2024)

First direct measurement of a₀ channel at energies below E_{cm}=2.5 MeV with a sensitivity better than the a₁ channel measured by Spillane et al. (2007)
 The thick target yields of direct measurement disagree with THM and mTHM

Error analysis using Monte Carlo Method

> 2.567MeV (0+) observed in THM does not appear in the direct measurement

> Spin/Parity assignment seems to agree with the Taniguchi&Kimura prediction of 0+/2+

Determination of J^{π} of E_{cm} =2.76 (MeV)

Algebraic approach to nuclear quasimolecular spectra

F. Iachello

A. W. Wright Nuclear Structure Laboratory, Yale University, New Haven, Connecticut 96511 and Kernfysisch Versneller Instituut, University of Groningen, Groningen, The Netherlands (Received 17 July 1980)

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VOLUME 23, NUMBER 6

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Rotational and vibrational excitations in nuclear molecular spectra

K. A. Erb and D. A. Bromley A. W. Wright Nuclear Structure Laboratory, Yale University, New Haven, Connecticut (Received 17 July 1980)

Prediction of 4⁺ state at 2.55 MeV is confirmed by our experiment!

Level density of the ¹²C+¹²C resonances

Correlation among the carbon isotope systems

For most energies, the ¹²C+¹²C cross sections are suppressed!

Only at resonant energies, the ${}^{12}C+{}^{12}C$ cross sections matches with those of ${}^{12}C+{}^{13}C$ and ${}^{13}C+{}^{13}C!$

Why?

0 0

0 0

M. Notani et al., Phys. Rev. C 85, 014607 (2012)

Correlation between carbon isotopes

Averaged S*

$$\sigma = \sum_{J} \sigma_{\rm CC}^{J} P_{J}$$

States for fusion

$$P_J = 1 - \exp(-2\pi\bar{\Gamma}_J/D_J)$$

 Γ : resonance width D: resonance spacing

Correlation between carbon isotopes

The ¹²C+¹²C resonance can be fragmented into small resonances due to the coupling to various decay channels, but the sum of the strength (after correcting for the penetration effect) should remain same

Sum rule of the molecular resonance

Extrapolation Towards lower energies Using statistical model approach

Faked resonances using Monte Carlo method

$Sum(\omega\gamma^*)$ vs. Er(MeV)

Reaction rate based on Monte carlo

T₉

Nature of the ¹²C+¹²C molecular resonance determines extrapolation

Resonant Energy, Strength, Penetration Factor

Gross structure (averaged xsec, Intermediate Structure Upper/lower limits) (structure of total fusion xsec)

Fine Structure (Structure in decay channels)

• Our direct measurement of ${}^{12}C+{}^{12}C$

 \rightarrow First direct measurement of a0 below 2.5 MeV indicates the needs of improving the THM/cTHM

- \rightarrow Our analysis suggests that statistical model is applicable at low energies
- Indirect (THM, 24Mg(a,a'))+Direct measurements will provide best extrapolation; Strongly coupled plasma screening needs to be studied
- Collaboration will end up with better science!

