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COHERENT ELASTIC SCATTERING OFF NUCLEUS OF ⁵¹Cr NEUTRINOS

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Istituto Nazionale di Fisica Nucleare

OUTLINE

The talk is based on the published paper

- Motivations
- The idea
- The ⁵¹Cr source
- The flux measurement
- The detector
- Simulation results
- Conclusions

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Regular Article - Experimental Physics

Coherent elastic nuclear scattering of ⁵¹Cr neutrinos

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COHERENT ELASTIC NEUTRINO-NUCLEUS SCATTERING

It occurs if the condition is satisfied: $qR \cong 1$ $E_{\nu} < 50$ -100 MeV

The cross section is high

$$\frac{d\sigma_{\nu}}{dE_R} = \frac{G_F^2}{4\pi} Q_{\text{SM}}^2 m_N \left(1 - \frac{E_r m_N}{2E_{\nu}^2}\right) \underbrace{F^2(E_r)}_{\text{Form factor}}$$

$$Q_{\mathsf{SM}}^2 = [N - (1 - s_W^2)Z]^2 \simeq N^2$$

Maximum recoil energy

$$T_{\max} \le \frac{E_{\nu}}{1 + \frac{M_A}{2E_{\nu}}}$$

Total cross section

$$\sigma_{\rm SM} \simeq \frac{G_{\rm F}^2}{4\pi} N^2 E_{\nu}^2 \simeq 0.4 \times 10^{-44} N^2 E_{\nu}^2 \,{\rm cm}^2$$





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Both cross-section and maximum recoil energy increase with neutrino energy





with artificial sources:

Source	Half-life	Progeny	Production	$E_{ u}$
³⁷ Ar	35.04 days	³⁷ Cl	40 Ca(n, α) ³⁷ Ar	811 keV (90.2%), 813 keV (9.8%)
⁵¹ Cr	27.70 days	⁵¹ V	n capture on 50 Cr	747 keV (81.6%), 427 keV (9%), 752 keV (8.5%)
⁶⁵ Zn	244 days	⁶⁵ Cu	n capture on 64 Zn	1343 keV (49.3%), 227 keV (50.7%)



Magnificent CEvNS workshop

THE IDEA

THE SOURCE

→Electron capture decaying isotope ⁵¹Cr source
 →Half-life 27.7 days
 →5 MCi (single activation at reactor)
 →Neutrino energy 747 keV (81%) and 752 keV (9%)

CALORIMETRIC ACTIVITY MEASUREMENT per mill precision

LOW THRESHOLD DETECTOR

Recoil energy O(10) eV

Volume: 2000 cm³ Cryogenics phonon detectors (Germanium or sapphire) or CCD detectors

energy threshold: as low as possible

KEY POINTS FOR A PRECISE MEASUREMENT:

- \rightarrow Precise (<1%) knowledge of the neutrino spectrum and flux
- \rightarrow Background rejection



THE ⁵¹CR NEUTRINO SOURCE

The **neutrino** spectrum consists in four mono-energetic lines:



Associated gamma emission

- 320 keV
- Bremsstrahlung up to 780 keV from K capture branching ratio: 8 10⁻⁴ for gamma with E>320 keV

Impurities can be activated during the irradiation

Gamma spectrum from internal bremsstrahlung





THE SOURCE PRODUCTION

The GALLEX (INFN) sample:

Mass: 36 kg

with 3.6 g/cm³ effective density in metallic chips of 1-5 mm



Isotopic composition: 50 Cr 38.6% 52 Cr 60,7% 53 Cr 0,7% 54 Cr < 0,3%</th>enriched in 50 Cr and depleted in isotope 53 Cr (high neutron capture cross section)

Activation of the sample at reactor

GALLEX:

Siloé reactor in Grenoble with an estimated neutron flux 2 10¹⁴ neutrons cm⁻² s⁻¹ 23.8 Days of irradiation Final activity of ⁵¹Cr: 1.7 MCi

Challenging numbers: neutron flux 5 10¹⁴ neutrons cm⁻² s⁻¹ 24 days of irradiation Final activity of ⁵¹Cr: 3.5-7 MCi



THE IRRADIATION PROCESS

 $R = N \mid \varphi(E)\sigma(E)dE$

The activation rate

Source activity

$$A(t) = R(1 - e^{-\lambda t_{irr}})e^{-\lambda t}$$

depends on :

- $\varphi(E)$, the averaged thermal neutron flux (in GALLEX 5 10¹³ n/cm2 s)
- the average lifetime of neutron in the reactor
- the neutron absorption length for ⁵⁰Cr and ⁵³Cr (24% higher)

95% ⁵⁰Cr enriched : 0.7 cm

The ⁵¹Cr lifetime is reduced due to neutron capture (small effect)

We can improve with:

- \rightarrow Higher intense flux (new reactor)
- \rightarrow Optimized geometry and source
- \rightarrow More cycles (since ⁵¹Cr lifetime and neutron capture from ⁵¹Cr)

Suitable research reactors

- High Flux Isotope Reactor (HFIR) at Oak Ridge (USA), 85 MW power 1.2-2.5×10¹⁵ n/cm²s • To be investigated!!
- BR2 reactor at Mol (Belgium), 100 MW 0
- Jules Horowitz Reactor (France) under construction 0

Recently a new ⁵¹Cr source has been produced in Russia (Dimitrovgrad)!!!

By BEST experiment (3.2MCi) 5th July 2019



The efficiency depends on the geometry and on the source properties

THE BYOLOGICAL SHIELD

For gamma emitted

- from ⁵¹Cr decay (320 keV) BR 10%
- from internal Bremsstrahlung (max 750 keV)
- from the activated impurities ^{110m}Ag (max 1.5 MeV) 1 10¹⁰ gamma/s

Made of tungsten alloy (the SOX shield might be adapted)

- for temperature
- for higher density

If we consider the same activation factor of GALLEX

The total gamma flux must be reduced for:
 → the dosimetric issue (dose < 100 uSv at contact)
 8 cm are enough (attenuation factor of 2000)

→ reducing background (for maximizing the S/N ratio) more stringent requirement

	Nucleus	Meanlife	Energy [keV]	Source activity (GBq)		
				After ex	ktra	ction
	²⁴ Na	14.8 h	1368.5	0.10	±	0.02
			2753.9	0.07	±	0.01
			average:	0.08	±	0.01
2.0.1.016	°Sc	83.9 d	1121	0.13	±	0.02
3.8 10 ¹⁰ gamma/s	^{*°} Sc	43.7 h	983.5	0.05	±	0.03
1 0 1 0 1 2			1037.5	0.07	±	0.03
1.3 10 ¹³ gamma/s			1312.1	0.10	, ±	0.01
1 1 0 10	⁶⁰ C.		average:	0.095) <u>+</u>	0.01
1 10 ¹⁰ gamma/s	6		11/3.2	0.02	±	0.01
			1332.5	0.04	±	0.01
	⁶⁴ Cu	127h	1245 8	210	די גי	0.00 20
	⁷⁷ Ge	12.7 m 11 3 h	7347.3	0.5	т. +	20 02
ed)	⁷⁶ As	437h	1212.7	6.5	- +	3
			1216.0	3	+	1
			2096.3	1.5	±	0.2
			average:	1.6	±	0.2
	⁹⁷ Zr	17.0 h	1749.9	0.40	+	0.15
	^{10m} Ag	249.8 d	657.7	4	±	2
			763.9	10	±	3
LEX			818			
			884.7	4.3	±	0.5
			937.5	4.2	±	0.5
			1384.3	5.0	±	0.5
			1475.7	5.1	Ŧ	0.5
ct)			1505.0	4.3	±	0.5
			1562.3	4.0	<u>±</u>	0.5
of 2000)	¹²⁴ Sp	60.2.4	average:	4.33	Ť	0.25
	311	00.2 u	1308	04	+	0.15
			1691	0.36		0.05
ratio)			1919	0.20	-	0.05
i latioj			2039.6	0.4	±	0.03
			2091	0.51	±	0.08
			2185			
			2294			
			average:	0.40	±	0.03

GALLEX SAMPLE CONTAMINATION



THE ACTIVITY MEASUREMENT



The heat contribution from impurities is negligible

36.51 keV for each decay expected power 422 W for 5 MCi

$$P_g = \dot{m} \cdot [h(p, T_{out}) - h(p, T_{in})]$$

After the thermalization phase (2-3 days) the measured power follows:



$$P_m = P_g e^{-\frac{(t-\Delta t)}{\tau}} - P_{lost}$$

The precision depends only on

- P_{lost}
- the delay time Δ*t*, (minimized and measured in the calibration phase)



THE SOX CALORIMETER



Ref: K. Altenmüller et al., JINST 13(09), P09008 (2018)

It was calibrated and tested with an electrical heat source and **0.2% precision** was achieved,



THE DETECTOR

FOR LOW ENERGY NUCLEAR RECOIL

2000 cm³ of volume with a very low threshold

$$T_{\max} \leq \frac{E_{\nu}}{1 + \frac{M_A}{2E_{\nu}}}$$

The neutrino energy is not high- \rightarrow it is fundamental push the threshold down

→ Cryogenics phonon detector germanium, silicon, sapphire (Al₂O₃), ...

The minimum recoil threshold is related to:

mean energy fluctuations in the absorber related to T and the heat capacity
 temperature fluctuations

The heat capacity (C) depends on the mass and on temperature

$$\Delta T = \frac{E_{\rm R}}{C(T)} {\rm e}^{-t/\tau} \qquad \tau = C(T)/G(T),$$

\rightarrow CCD detectors:

very low noise, but the conversion efficiency has to be measured at low energy a threshold of **5 eV** have been demonstrated on 5 g (CONNIE experiment)



WHICH THRESHOLD VALUE?

• An energy threshold of **20 eV** has been already demonstrated on a 0.5 g

 $(5 \times 5 \times 5 \text{ mm}^3)$ sapphire Al₂O₃ target

NUCLEUS experiment PRD 96, 022009 (2017) CRESST experiment EPJC 77:63 (2017)

 60 eV have been demonstrated on a 33.4 g (20 mm × 20 mm) Germanium target EDELWEISS experiment PRD 99, 082003 (2019)



The challenge is scaling the detector mass to few kg! By realizing arrays of thousands of small mass detectors

New developments are expected in these years...the threshold might be pushed down!



THE PROPOSED LAYOUT

for maximizing the detected event



2000 cm³ volume (5-10 kg)

12 cm between source and detector

Ag gamma flux (GBq): reduced of 10⁻⁶

Cr Bremsstrahlung (PBq): reduced of 10⁻¹¹





THE SIMULATION RESULTS

Initial activity: 5 MCi Detector volume: 2000 cm³ Exposure: 55 days (2 half lives)

Neutrino flux in the detector



Average neutrino flux: 1 10¹³ v/cm²s

	Detector	Threshold [eV]	Counts in 55 days			
	Sapphire	20	900			
	Ge	8	3900			
			ost ost There are			
counts / 2000 cm ³ 55 4 d 5 MCi	10^4		Si Ge Al_2O_3 CaWO ₄ 40 50 60 $T_{\text{threshold}}$ [eV]			

THE BACKGROUND

- From gammas emitted by source impurities (^{110m}Ag, ..)
 - We extrapolated our Geant4 simulations to low energy (where they are not reliable)
 - It seems that the Ag impurities in the GALLEX sample should be reduced from ppm to ppb for not generate a background More precise simulations are necessary
- From environmental neutrons (if the measurement is performed not far from reactor)
 - hard to predict at these low energies
 - it will measured by upcoming reactor experiment
 - additional external absorber shield can be inserted in the design

THE BACKGROUND MUST BE CAREFULLY SIMULATED AND REDUCED



CONCLUSIONS

The proposed idea seems promising for achieving few percent cross section accuracy

Important advantages:

Very precise knowledge

- of the neutrino energy
- source activity

complementary information with antineutrino measurement from reactors

Critical points:

- source production
- background estimation

Next steps:

- deeper investigation for the source production
 - optimization for irradiation
 - impurities minimization)
- precise background simulation and estimation

