Absolute electromagnetic transition rates in semi-magic N = 50 isotones as a test for $(\pi g_{9/2})^n$ single particle calculations.

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- 1. Motivation
- 2. Fast timing using centroid difference methods
- 3. Experiments on 92 Mo and other N = 50 isotones
- 4. Conclusions

1. Motivation

Untruncated **numerical full shell model calculations** with the SR88MHJM interaction and all proton orbits between Z= 38 and Z=50.

Analytical single-j calculations with a seniority conserving interaction or with empirical two-body matrix elements. Example: 93 Tc as $(\pi 1g_{9/2})^3$

A de Shalit and I. Talmi, The Nuclear Shell Model (1963)

Recently, this approach was extended to electromagnetic quadrupole transition rates by Piet Van Isacker.

- Assumptions:
- Seniority is conserved.
- The effective charges in one-body E2 operator of the two-j nucleus can be state dependent.
- The effective charges in the quadrupole moment of the state with spin R are the same as those for $B(E2;R\rightarrow R-2) = B_R$ in the two-particle nucleus.

Then the following relation can be obtained:

$$
B(\mathrm{E2}; j^3[I]J \to j^3[I']J') = \left(\sum_R g_j(J, I, J', I', R)\sqrt{B_R}\right)^2
$$

First application to 135 l as $(\pi 1g_{7/2})^3$ P. Spagnoletti et al. Phys. Rev. C 95 (2017) 021302 Very succesful for ²¹¹At using ²¹⁰Po V. Karayonchev et al., Phys. Rev. C 106, (2022) 044321

2. Fast timing using Centroid Difference Methods

The Compton background has an important time dependence which needs to be corrected for.

Example time response using ⁶⁰Co.

See also H. Mach et al. Nucl. Phys. A 523 (1991) 197 section 2.3.

The Generalized Centroid Difference (GCD) method for g**-**g **fast timing arrays**

[J.M. Régis et al., NIM A 726 (2013) 191]

The generalisation of the MSCD method was done for the FATIMA@EXILL Array in Spring 2013. It holds for arrays holding the same type of scintillators, all at the same distance from the target. Then the averaged spectra still follow the MSCD relations.

$$
\Delta \overline{C}_{\text{decay}} (\Delta E) = \overline{C}_{\text{stop}} - \overline{C}_{\text{start}} =
$$

$$
\overline{PRD}(E_{\text{feeder}}) - \overline{PRD}(E_{\text{decay}}) + 2\tau
$$

The Symmetrized GCD method:

[J.M. Régis, M. Dannhoff, J. Jolie NIM A 897 (2018) 38]

Using the mirror symmetry between delayed and antidelayed time spectra, the latter can be mirrored and added to the first leading when T_0 is put to zero to: $\tau = C^D - TW$ (E_{feeder}, E_{decay}).

3. Experiments on 92Mo and other N = 50 isotones.

 $Z = 50$ In 98 In 99 In 100 $In 10$ In 102 In 103 114.818 $3.1 s$ $5.9 s$ $17s$ | $45m$ $16s$ $22.1 s$ 49 1004: 795 252: 750 Cd
112.411 Cd 97 **Cd 98** Cd 99 Cd 100 Cd 10⁻ Cd 102 1.2_m $16s$ $491s$ 48 343; 672;
583... 937; 140; 98; 1723
259; 925. Ag 97
25.3 s Ag 98
46.7 s Ag 99 Ag 100 Ag 101 $10.5 s$ $2.3 m$ $3.1 s$ 863; 679; **Pd 96 Pd 97 Pd 98 Pd 99 Pd 100** 2.0_m $17.7 m$ $21.4 m$ $3.7d$ 3.1_m ³⁺ 3.5...
_Y 265; 475; ⁺ 2.2...
136; 264; RIB $\frac{1}{6}$; $\frac{1}{6}$ + 0.7
(112; 663; **Rh 98 Rh 99 Rh 94 Rh 95 Rh 96 Rh 97** $15 m | 8.7 n$ Ru 99
12.76 $.93$ **Ru 94 Ru 95 Ru 96 Ru 97 Ru 98 Ru 92** 5.54 $2.9d$ 1.87 $3.65 m$ $51.8 m$ $1.65h$ $16:324$ **Tc 92 Tc 93** $.95$ **Tc 96 Tc 97** 4.4_m $92.2 d$ Mo 96 Mo 9. **Mo 90 Mo 91** 16.68 9.56 9.23 $5.7h$ $2.15n$ **Nb 92 Nb 89 Nb 90 Nb 91** 18.8 s | 14.6 $0.15 d$ 3.6 $\frac{Zr}{51.45}$ $Zr91$ _{11.22} $Zr92$
17.15 Zr 94 Zr 8 Zr 88 Zr 89 $\frac{21}{64.0}$ d $Z = 40$ $4.0 s$ 83.4 d $0.4; 1.1$
'57; 724. -0.014 0.049 $Y92$
3.54 Y 87 Y 88 Y 89 Y 91 Y 93 Y 94 $13 h$ 80.3h 106.6 d $16.0 s$ $3.19h$ 49.7_m $18.7r$ 48_m

 $N = 50$

After the success in the $N = 126$ isotones, it would be interesting to study similar isotones.

Candidates could be the N= 50 isotones above Z= 40 where the $\pi(1g_{9/2})$ orbit gets filled. Also here, the knowledge on lifetimes and B(E2) values is limited and often contradictory or unprecise.

The problem is to populate the isotones above 90Zr using stable or radioactive ion beams.

Here we report on the stable beam experiments performed recently in Cologne.

92Mo:

The main problem is the lifetime of the first 4⁺ state which is needed for the prediction of all other B(E2) values. Note B(E2) to first 2⁺ known via Coulex.

Recently, the lifetime of the 2^+ and 4^+ states were measured for the first time as 0.8(4) ps and 35.5(6) in a recoil distance experiment at GANIL. *R. M. Pérez-Vidal et al. Phys. Rev. Lett. 129 (2022) 112501.*

To verify the lifetime of the first 4⁺ state and reduce the statistical and systematic error two fusion evaporation reactions were used at the 10MV Tandem accelerator in Cologne.

EXP1: $90Zr(\alpha,2n)$ $92Mo$ @ 27 MeV on a 5.3mg cm⁻² 97.62% enriched target.

EXP2: $93Nb(p,2n)$ $92Mo$ @ 18 MeV on a 5.4mg cm⁻² monoisotopic target.

New:

Completely digital acquisition system (CAEN 500MHz digitisers) with digital CFD algorithm to reach timestamps with ps resolution*. A. Harter et al. NIM A 1053 (2023) 168279.* Symmetrized Analysis*. J. M. Régis et al. NIM A 897 (2018) 3.* Remeasurement of PRD relevant lifetime in 152Gd. *L. Knafla et al. NIM A 1052 (2023), 168279.* The HORUS array was equipped with 8 Ge detectors and 9 LaBr3(Ce) scintillators of which 6 with BGO shields. $LaBr+BGO$

esiduum [ps]

600

400

800

 E_{γ} [keV]

1000

1200

Time Walk (TW) is obtained from a 152Eu source using the new value for the first excited 2^+ state in 152 Gd.

L. Knafla et al. NIM A 1052 (2023), 168279.

$$
C_{PP} = C_{exp} + \tilde{t}_{cor},
$$
\n
$$
\tilde{t}_{cor} = \frac{P/B(E_f) t_{cor}(E_i) + P/B(E_i) t_{cor}(E_f)}{P/B(E_i) + P/B(E_f)},
$$
\n
$$
t_{cor} = \frac{C_{exp} - C_{BG}(E)}{P/B(E)},
$$
\n
$$
Exp1: \tau = 22.5(11) \text{ ps}
$$
\n
$$
Exp2: \tau = 23(2) \text{ ps}
$$

GANIL: $\tau = 35.5(6)$ ps

$J_i^{\pi_i} \rightarrow J_f^{\pi_f}$	$\tau_{\rm EXP1}$ ps	$\tau_{\rm EXP2}$ ps	τ_{adopted} ps	Multipolarity	$B(\sigma \lambda; J_i^{\pi_i} \rightarrow J_f^{\pi_f})$ adopted	$B(\sigma \lambda; J_i^{\pi_i} \rightarrow J_f^{\pi_f})$ literature
$2^+_1 \rightarrow 0^+_1$	≤ 3	≤ 8	\leqslant 3	E ₂	\geqslant 35 e^2 fm ⁴	$207(12)$ e^2 fm ⁴ [30,31]
$4^+_1 \rightarrow 2^+_1$	22.5(11)	$23(2)^{a}$	22.5(11)	E2	132^{+7}_{-6} e ² fm ⁴	84.3(14) e^2 fm ⁴ [5]
$6^+_1 \rightarrow 4^+_1$ \rightarrow 5 ⁻	2200(20)	2220(70)	2200(20)	$E2^{\rm b}$ $E1^{\rm b}$	$81(2)$ e^2 fm ⁴ $5.3(6) \times 10^{-5} e \text{ fm}^2$	80(3) e^2 fm ⁴ [30,32] $5.3(7) \times 10^{-5} e$ fm ² [30]
$8^+_1 \rightarrow 6^+_1$	$310(3) \times 10^{3c}$		$310(3) \times 10^3$	E2	28.6(3) e^2 fm ⁴	$32(1)$ e^2 fm ⁴ [30,33–37]
$5^{-}_{1} \rightarrow 4^{+}_{1}$	2270(30)	2250(60)	2270(30)	$E1^d$ M2 ^d	\geqslant 1.88(3)×10 ⁻⁵ e fm ² \leqslant 93 µN ² fm ⁴	$1.91(5) \times 10^{-5} e$ fm ² [30,38] $\leq 98 \mu N^2$ fm ⁴ [30]
$7^{-}_{1} \rightarrow 5^{-}_{1}$	\leqslant 5	\leqslant 7	\leqslant 5	E2	\geqslant 101 e^2 fm ⁴	
$9^{-}_{1} \rightarrow 7^{-}_{1}$	37(11)	29(7)	$31(6)^e$	E2	271^{+65}_{-44} e ² fm ⁴	

TABLE I. Summary of the measured mean lifetimes of the states $J_i^{\pi_i}$ and the respective reduced transition probabilities.

^a Averaged value from feeder-decay cascades 244–773 and 330–773 calculated using a Monte Carlo method.

^bThe branching ratio for the $6₁⁺$ level was derived using the intensities from Ref. [29].

^cDetermined using Ge-LaBr timing.

^dMixing ratio $\delta \leq 0.05$ from Ref. [39].

^eWeighted average from EXP1 and EXP2.

5: R. M. Pérez-Vidal et al. Phys. Rev. Let. 129, 112501 (2022)

M. Ley, L. Knafla, J. Jolie, A. Esmaylzadeh, A. Harter, A. Blazhev, C. Fransen, A Pfeil, J.-M. Regis, and P. Van Isacker, PRC108 (2023) 064313.

TABLE II. Experimental and calculated $B(E2)$ values in ⁹²Mo.

				$B(E2; J_i^{\pi} \to J_f^{\pi}) (e^2 \text{ fm}^4)$			
$v_{\rm i}$	J_i^{π}	$v_{\rm f}$	$J_{\rm f}^{\pi}$	Exp	$\hat{T}_1(E2)$	$\hat{T}'_1(E2)$	
2	2^{+}_{1}	0	0^{+}	207(12)	89	207(12)	
2	4^{+}_{1}		2^{+}_{1}	132^{+7}_{-6}	103	132^{+7}_{-6}	
2	6^{+}_{1}		4^{+}_{1}	81(2)	71	81(2)	
2	8^{+}_{1}		6^{+}_{1}	28.6(3)	28	28.6(3)	

The effective charge in the one-body operator $T_1(E2)$ is obtained from the quadrupole moment of the $9/2$ ⁺ ground state of $91Nb$, with the experimental value and uncertainty $Q(9/2^+) = -25(3)$ e fm2.

The operator **T´**1(E2) has effective charges that depend on the two-nucleon states $(\pi 1g_{9/2})^2$ and which will be used to predict transition rates in $(\pi 1g_{9/2})^n$ states with $n>2$.

TABLE IV. Experimental and calculated $B(E2)$ values in ⁹⁴Ru.

a: R. M. Pérez-Vidal et al. Phys. Rev. Let. 129, 112501 (2022)

τ_{4} = 87(8) ps from plunger measurement at GANIL

b: B. Das et al. Phys. Rev. C 105, L031304 (2022)

τ_{4} = 32(11) ps from fast timing measurement at GSI

93Tc:

Very few absolute transition rates are known in this three valence proton nucleus:

the B(E2; 17/2₁⁺ \rightarrow 13/2₁⁺)= 88(18) e²fm⁴ and the B(E2; 21/2₁⁺ \rightarrow 17/2₁⁺)*= 66(2) e²fm⁴

A fast timing experiment was performed in Cologne using the $90Zr$ ^{[6}Li, 3n)⁹³Tc @ 31MeV reaction on a : 5.3mg/cm^{2 90}Zr (98% enriched) target.

Results:

94Ru:

Fast Timing experiment at Cologne Tandem

 92 Mo(⁴He, 2n)⁹⁴Ru @ 28MeV on 5.5mg/cm² 92Mo (98% enriched)

Results:

 $\times 100$

 \mathbf{a}

 $b)$

 4_{1}^{+}

gat es:

 4^{+1} 2^{+1}

 \vert LaBr(311)

gat es:

 2^{+}_{1}

 $0^{+\frac{1}{2}}$

 $LaBr(756)$

 \vert Ge(1431)

 $4^{+}_{1} \rightarrow 2^{+}_{1}$

 $p/b = 24$

 $p/b = 42$

20

16

12

8

8

 $\begin{array}{c}\n\text{const/keV} \\
\text{12}\n\end{array}$

M. Ley, J. Jolie, L. Knafla, A. Blazhev, A. Esmaylzadeh, C. Fransen, A Pfeil, J.-M. Regis, and P. Van Isacker, Phys.Rev C 110 (2024) 034320.

 $\tau = 66(2)$ ps or B(E2; 4+ \rightarrow 2+)= 50(2) e²fm⁴ while the single-j prediction is B(E2; 4+ \rightarrow 2+)= 7.8(7) e²fm⁴. Also here the main problem is the lifetime of the first 4⁺ state. Two recent RIB experiments at FAIR Phase 0 and GANIL yielded contradictory results: $\tau = 32(11)$ ps ¹ and $\tau = 87(8)$ ps ². We got To obtain an understanding of this disagreement, it is essential to consider that for four particles or four holes in a $j = 9/2$ orbit two 4⁺ levels with $v = 2$ and $v = 4$ occur close in energy. Given that the $\Delta v = 2$ transition is ~14 times faster than the one with $\Delta v = 0$, a small admixture of $v = 4$ in the first 4⁺ state can considerably alter the B(E2; $4^+ \rightarrow 2^+$) value.

100 Seniority constructive interference However, the $v = 4.4$ ⁺ state is solvable for *any* interaction in $a_j = 9/2$ orbital, which means 80 that in order to mix with the $v = 2$ state $B(E2)$ [e²fm⁴] 60 necessarily must involve components outside Seniority mixed regime Seniority conserved regime the 0*g*9*/*2 space.20 $4, 2$ $\bf{0}$ Seniority destructive interferenc $1 B.$ Das et al. Phys. Rev. C 105, L031304 (2022) Fast Timing 0.7 0.8 0.9 $1₃$ 2 R. M. Pérez-Vidal et al. Phys. Rev. Let. 129, 112501 (2022) RDDM

Assuming an ad hoc mixed structure of the first 4+ and 6+ state:

$$
|4_{1}^{+}\rangle = \alpha_{4} |4_{v=2}^{+}\rangle + \beta_{4} |4_{v=4}^{+}\rangle,
$$

$$
|6_{1}^{+}\rangle = \alpha_{6} |6_{v=2}^{+}\rangle + \beta_{6} |6_{v=4}^{+}\rangle,
$$

Two almost equally good solutions ξ_1 and ξ_2 are obtained.

The values are compared to a large scale shell model calculation using a 1g,2d,3s configuration with up to 4p-4h excitations across $Z = 50$ [5].

[5] H. Mach et al., Phys. Rev. C 95, 014313 (2017).

4. Conclusions and outlook

Excellent agreement with the single-j predictions for the $B(E2)$ values in ²¹¹At was obtained when using the $B(E2)$ values in ²¹⁰Po as input.

In order to perform the same for the $N = 50$ isotones, precise lifetimes in 92 Mo were determined to serve as input for the calculations with more than two protons.

A fast timing experiment in 93 Tc yields promising results but more B(E2) values are needed, especially for low spin states.

The B(E2; $4+\rightarrow 2^+$) in ⁹⁴Ru was measured to solve contradictory results from RIB experiments, but it still disagrees with the single-j predictions and LSSM calculations.

Much more stable and RIB experiments are needed.