Beta Decay of K^{47} , Ca^{50} , and $Sc^{50\dagger}$

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The isotopes K^{47} , Ca⁵⁰, and Sc⁵⁰ were formed by the bombardment of Ca⁴⁸ with 3.3-MeV tritons. Half-lives of 18.1 ± 1.0 , 13.9 ± 0.6 , and 102.5 ± 0.5 sec, respectively, were measured by following the decay of the γ rays emitted by these nuclides. The γ rays were observed with a Ge(Li) detector, and γ -ray energies and relative intensities were obtained. K⁴⁷ decays to the second and third excited states of Ca⁴⁷; Ca⁵⁰ to the fourth excited state of Sc⁵⁰; and Sc⁵⁰ to the second and third excited states of Ti⁵⁰. The results for K^{47} , together with previous results establish the spin-parity of the 2578- and 2599-keV levels of Ca⁴⁷ as $\frac{3}{2}$ ⁺ and $\frac{1}{2}$ ⁺, respectively. The results for Sc^{50} and Ca^{50} are consistent with previous work, but are more accurate.

I. INTRODUCTION

When Ca^{48} is bombarded with $3-MeV$ tritons the radioisotopes K^{47} , Ca⁵⁰, and Sc⁵⁰ are produced via the reactions $Ca^{48}(t, \alpha)K^{47}$ ($Q=4.006$ MeV), Ca^{48} - (t, p) Ca⁵⁰ ($Q = 3.017$ MeV), and Ca⁴⁸ (t, n) Sc⁵⁰ (Q =7.002 MeV). Sc⁵⁰ is also produced by $Ca^{50}(\beta^-)Sc^{50}$. Reported herein is a study of these activities produced via $Ca^{48}+t$.

The β^- emitter K^{47} was first found and studied via the $\text{Ca}^{48}(\gamma, p)\text{K}^{47}$ reaction.¹ It was observed to decay with a half-life of 17.5 ± 0.3 sec to the first excited state at 2.01 MeV (85%) and to the second excited state at 2.6 MeV (15%) with $\log ft$ values of 4.8 and 5.4, respectively. Since both these $\log ft$ values fall in the allowed region' it was concluded that the K^{47} ground state and the Ca⁴⁷ 2.01- and 2.6-MeV levels have the same parity. Subsequently, it has been established that the Ca^{47} 2.01-MeV level³ has $J^{\pi} = \frac{3}{2}^{-}$, and the K⁴⁷ ground state has⁴ $J^{\pi} = \frac{1}{2}^{+}$; while the 2.6 -MeV level of Ca^{47} is actually a doublet with excitation energies of 2580 ± 5 and 2601 \pm 5 keV and spin-parity assignments³ of $(\frac{3}{2}^+, \frac{5}{2}^+)$ and $\frac{1}{2}^+$, respectively. These findings are obviously in conflict with the K^{47} decay results of Kuroyanagi et al.¹ and so a reinvestigation of the β^- decay of K^{47} is in order.

The β ⁻ decay of Ca⁵⁰ has been most recently and thoroughly studied by Chase, McDonald, and Nightingale (CMN)' who observed that the decay proceeded to the 1848-keV fourth excited state of Sc^{50} rather than to the second excited state at 329 keV as had been first proposed.⁶ Half-lives of 14 ± 3 sec⁵ and 9 ± 2 sec⁶ have been reported. It was our intention to check the results of CMN and to provide more accurate energy measurements for the Sc⁵⁰ γ rays and a more accurate half-life for Ca⁵⁰.

 Sc^{50} has a half-life³ of 103.8 ± 1.7 sec and decays predominantly to the Ti⁵⁰ 3.20-MeV third excited state. There is evidence⁷ for a small branch (15) \pm 7)%] to the 2.68-MeV level, but other investigations' yield somewhat smaller, albeit even less well defined, values for this branch. It was our intention to determine more accurate energies for the Ti⁵⁰ γ rays and more accurate branching ratios for the decay of Sc^{50} to the 2.68- and 3.20-MeV levels of $Ti⁵⁰$.

II. EXPERIMENTAL PROCEDURE AND RESULTS

A.
$$
Sc^{50}(\beta^{r}) Ti^{50}
$$

A 30-cc Ge(Li) detector system with an energy resolution defined by a full width at half maximum (FWHM) of 2.1 keV for the $Co⁶⁰$ lines was used for investigation of the delayed γ rays from Ca⁴⁸ +t. The Ca^{48} target consisted of a 250-mg CaF , crystal with the Ca enriched to 96.6% in Ca⁴⁸. The CaF₂ was approximately in the form of a cylinder 1 cm in diameter and thick enough to stop the beam.^{8,9} $\mathbf{F_{2}}$
 \mathbf{cm} 8,9 The triton beam was provided by the BNL 3.5-MeV Van de Graaff accelerator.

The 1.7-min Sc^{50} activity was investigated first. The main problem was to obtain an accurate relative-intensity measurement of the three Ti⁵⁰ γ rays The main problem was to obtain an accurate relative-intensity measurement of the three Ti⁵⁰ γ randomserved $3.5-7$ in the β^- decay of Sc⁵⁰. A relative efficiency curve accurate to $\leq 2\%$ had already been determined for γ -ray photopeaks observed with the determined for γ -ray photopeaks observed with 30-cc Ge (Li) detector.¹⁰ Thus, γ -ray spectra were recorded under the same conditions as used in the determination of this curve (a source-to-detector-face distance of $\neg 6$ cm with no material between the detector housing and source).

The Ca⁴⁸ was placed under vacuum and bombarded for ² min with a 20-nA 3.3-MeV triton beam. It was then removed from the target chamber, carried to the shielded Ge(Li) detector situated in another room, and counted for ³ min after a 1.5-min wait. This procedure was repeated 12 times. The pertinent portion of the resulting Ge(Li) spectrum is shown in Fig. 1. The whole procedure was repeated once under different conditions of energy dispersion, counting rate, etc. The two spectra

 $\overline{2}$

FIG. 1. 30-cc Ge(Li) γ -ray spectrum observed following bombardment of Ca⁴⁸F₂ with a 3.3-MeV triton beam. The original spectrum has been compressed by a factor of 3 to facilitate display. The spectrum shown is the sum of 12 cycles consisting of 2-min bombardment at 20 nA, 1.5-min wait, and 3-min counting. The three lines ascribed to the decay of Sc⁵⁰ are indicated. The F²⁰ γ ray is due to F¹⁹(t,d)F²⁰(β)Ne²⁰(1+0). F²⁰ has $T_{1/2}$ =11 sec. The insert shows the origin of the three Sc⁵⁰ γ rays.

were analyzed to obtain relative intensities for the three Sc⁵⁰ γ rays indicated in Fig. 1. The two results were in good agreement yielding intensities for the 524- and 1554-keV lines, relative to 100 for the 1121-keV line, of 88 ± 3 and 102 ± 3 . The intensity of the 524-keV line is definitely less than that of the 1121-keV line so that there is a branch to the 4° , 2.68-MeV level as well as to the (6°) , 3.20-MeV level as indicated in the insert of Fig. ¹ and in Table I. Our result for the branch to the 2.68-MeV level $[(12 \pm 3)\%]$ is in excellent agreement with the value of $(15±7)$ % found by Chilosi et al.⁷

A search for other γ rays from Sc⁵⁰(β ⁻)Ti⁵⁰ was undertaken with 2.5 cm of Lucite inserted between the source and detector in order to stop electrons emanating from the source. This shielding decreased the background significantly, especially in the energy region above the 1554-keV γ -ray fullenergy peak. The main source of background for

 E_γ >1554 keV was the 8.8-min Ca⁴⁹ activity,¹¹ pro-
duced via the Ca⁴⁸(*t*, *d*)Ca⁴⁹ reaction (*Q* = -1.113 duced via the Ca⁴⁸ (t, d) Ca⁴⁹ reaction ($Q = -1.113$) MeV). No other γ rays were observed that could be ascribed to a 1.7-min activity. We can say that the intensity of any such γ rays with E_{γ} > 530 keV is less than 1% of the intensity of the 524-keV γ ray.

Accurate energies were obtained for the three Sc^{50} γ rays of Fig. 1 from a 4096-channel spectrum taken with mixed sources of Sc^{50} , Co^{60} , ThC", F^{20} , Ca^{49} , and Bi²⁰⁷. The peak positions were determined by Gaussian fits, and a least-squares fit to

$$
E = \sum_{n=0}^{m} a_n x^n
$$

was made to the γ rays of known energy, where x is the channel number and m is varied from 1 to 4. Four such spectra were taken and analyzed. In all cases the quality of the calibration improved

| Transition | Half-life (sec) | Branching ratio $(\%)$ | E_{β} (max) (Ref. a) (keV) | $\log ft$ | Conclusion |
|---|--------------------|-----------------------------|--|-----------------|---------------------|
| $K^{47}(\beta^-) Ca^{47}(1)$ | | <9.0 ^b | 4637 ± 13 | >6.03 | \cdots |
| $K^{47}(\beta^-) Ca^{47}(2)$ | 17.5 ± 0.3 | 14.6 ± 1.5 | 4072 ± 13 | 5.57 ± 0.05 | allowed |
| $K^{47}(\beta^-)Ca^{47}(3)$ | | 85.4 ± 5.0 | 4051 ± 13 | 4.79 ± 0.03 | allowed |
| $Ca^{50}(\beta^-)Sc^{50}(4)$ | 13.9 ± 0.6 | 100.0 | 3118 ± 17 | 4.13 ± 0.02 | allowed |
| $\text{Sc}^{50}(\beta^-) \text{Ti}^{50}(1)$ | | $< 5.0^{\circ}$ | 5336 ± 20 | >7.36 | \cdots |
| $\rm Sc^{50}(\beta^-)Ti^{50}(2)$ | 102.5 ± 0.5 | 12.0 ± 3.0 ^c | 4215 ± 20 | 6.51 ± 0.11 | $ \Delta J \leq 1$ |
| $Sc^{50}(\beta^-)Ti^{50}(3)$ | | 88.0 ± 3.0 ° | 3692 ± 20 | 5.39 ± 0.02 | allowed |

TABLE I. Half-lives, branching ratios, and log ft values for the β^- decays of K⁴⁷, Ca⁵⁰, and Sc⁵⁰.

From the present excitation energies and previous ground-state energy differences (see Ref. 3).

 b This branch was assumed to be zero in the calculation of other branching ratios and log ft values.

^c Note added in proof: These branching ratios are in serious disagreement with recently obtained values of $(24\pm1)\%$ and (76 ± 2) % [T. E. Ward and P. K. Kuvoda, Radiochim. Acta. 12, 217 (1969)].

drastically (i.e., χ^2 decreased markedly) when m was increased from ² to 3. This reflects the presence of monotonically varying nonlinearities in the
detector-amplifier-analyzer system.¹² Our Sc⁵⁰ detector-amplifier-analyzer system.¹² Our Sc⁵ γ -ray energies were taken from the calibration curve with $m=3$; the results are given in Table II, and the resulting excitation energies are listed in Table III.

The half-life of Sc^{50} was measured using a 5×6 in. NaI(Tl) γ -ray detector to view γ rays from the source. The $Nai(Tl)$ detector was shielded on the sides with Pb and had 2.5 cm of Lucite over its front face. An energy gate was set on the 1121 keV full-energy peak (Fig. 1), and the time decay of the γ -ray counts in this gate was observed using a Northern Scientific Inc. 4096-channel analyzer operating in a time scaling mode ("multiscaling") with 0.9 sec/channel. The decay was followed for 20 min after a 2-min bombardment and a 1-min wait. This procedure was repeated twice. The long-lived $(T_{1/2} \gg 1 \text{ min})$ background, mostly from the 8.8-min Ca⁴⁹ activity, was 0.6% of the initial Sc^{50} counting rate and thus introduced negligible error when taken into account. The half-life was extracted using several different computer fitting procedures and programs developed in a compreprocedures and programs developed in a compr
hensive study of β^{\pm} half-lives.¹³ The result was 102.5 ± 0.5 sec, where the uncertainty is mainly an estimate of the systematic errors - the statistical error being ± 0.2 sec. This result is in good agreement with the average³ of previous values, 103.8 $± 1.7$ sec.

B. $Ca^{50}(\beta^{2})Sc^{50}$

The study of the decay of both Ca^{50} and K^{47} was made using a pneumatic beam shutter about 5-m upstream from the Ca^{48} target. A cam system allowed continuous cycling with a 20-sec irradiation and a 80-sec counting period. A TMC 16384-channel analyzer was programmed to record four se-

TABLE II. Energies and assignments of γ rays from activities produced by $Ca^{48} + t$.

| E_{γ} (keV) | Assignment | Source |
|-----------------------|---------------------------------|---------------------------|
| 71.54 ± 0.20 | $\rm Sc^{50}$ 2 \rightarrow 1 | $Ca^{50}(\beta^-)$ |
| 256.94 ± 0.10 | $\rm Sc^{50}$ 1 \rightarrow 0 | $Ca^{50}(\beta^-)$ |
| 523.50 ± 0.10 | Ti^{50} 3 \rightarrow 2 | $\text{Sc}^{50}(\beta^-)$ |
| 564.74 ± 0.30 | Ca^{47} 2 \rightarrow 1 | $K^{47}(\beta^-)$ |
| 585.75 ± 0.30 | Ca^{47} 3 \rightarrow 1 | $K^{47}(\beta^-)$ |
| 1121.03 ± 0.10 | Ti^{50} 2 \rightarrow 1 | $\text{Sc}^{50}(\beta^-)$ |
| 1519.44 ± 0.30 | $\rm Sc^{50}$ 4 \rightarrow 2 | $Ca^{50}(\beta^-)$ |
| 1553.71 ± 0.20 | Ti^{50} 1 \rightarrow 0 | $\text{Sc}^{50}(\beta^-)$ |
| 1591.00 ± 0.30 | $\rm Sc^{50}$ 4 \rightarrow 1 | $Ca^{50}(\beta^-)$ |
| 2013.13 ± 0.30 | Ca^{47} 1 $\rightarrow 0$ | $K^{47}(\beta^-)$ |

TABLE III. Some excitation energies in Ca⁴⁷, Sc⁵⁰, and Ti⁵⁰.

 $^{\rm a}$ Transitions to or from the third excited state of $\rm Sc^{50}$

at 761 kev were not observed.

quential 4096-channel spectra, each of. 18-sec duration, during the 80-sec counting period. The 30 cc $Ge(Li)$ detector viewed the target in situ and was well shielded against the Van de Graaff accelerator by lead. The intensity of the 3.3-MeV triton beam was 60 nA throughout.

The principal features of the decay of Ca^{50} are illustrated in Fig. 2. The 72 -keV Sc^{50} second-tofirst excited state and 257 -keV Sc 50 first-to-zeroth excited-state transitions were studied first, Four high-dispersion (0.145 keV/channel) 4096- channel spectra were recorded as explained above with sources of ThC", $Co⁵⁷$, and Hg²⁰³ positioned to give energy calibration lines of reasonable intensity for use in the energy measurement of the 257 keV line. From these spectra, the 257-keV γ ray was determined to have an energy of 256.94 ± 0.10 keV and to decay with a half-life of 13.7 ± 0.6 sec. The energy of the 72-keV γ ray was measured relative to the accurately known energies of x rays from Pb^{208} and Bi^{208} emitted by the ThC" source. The portion of the spectra including the 72-keV line and the x rays is illustrated in Fig. 3 which shows the middle two of the four sequential spectra. The energy and area of the 72-keV line were obtained from a four-peak Gaussian fit to the 72-keV line and the three nearby x-ray peaks. The result was 71.54 ± 0.20 keV and a decay half-life of 15 ± 2 sec. No attempt was made to determine the relative intensities of the 72- and 257-keV lines.

An incidental result of this measurement was the observation of a γ ray of 139.78 \pm 0.06 keV and $T_{1/2}$ = 48.6 ± 3.5 sec. This we identify as the decay of the metastable first excited state of Ge^{75} , which is presumably formed by neutron capture on Ge⁷⁴

FIG. 2. Decay schemes of Ca⁵⁰ and Sc⁵⁰. All energies are in keV, the ground-state energies of Ca⁵⁰ and Ti⁵⁰ are relative to Sc^{50} . Uncertain assignments are in parentheses. The third excited state of Sc^{50} at 761 keV has been omitted.

 $(36.5\%$ abundant) in the Ge(Li) detector.

Decays to or from the 761-keV third excited state³ were not observed. The energies, half-lives, and relative intensities of the Sc⁵⁰ γ ray lines due to the decay of the 1848-keV fourth excited state to the first and second excited states were determined from spectra recorded in a similar fashion but with lower dispersion. Several different measurements were made. One is illustrated in Fig. 4 which shows the first of the four 18-sec spectra shifted by computer to match the conditions of Fig. 1. This spectrum was collected in -12 ^h with the principal aim of determining the relative intensities of the K^{47} lines. The detector was, at this time, beginning to show the effects of neutron damage as indicated by the low-energy tails on the peaks of Fig. 4. From these measurements, the half-lives of the 1519- and 1591-keV lines were determined to be consistent with each other and with those of the 72- and 257-keV lines. Our final adopted half-life for Ca^{50} is 13.9 ± 0.6 sec.

Our results for the energies of the 1519- and 1591-keV lines are included in Table II. The calibration lines used in the determination of those energies are the Sc⁵⁰, F^{20} , F^{21} , Na²⁴, and Ca⁴⁹ lines

FIG. 3. Portions of the 30 -cc γ -ray spectrum used in the determination of the half-life and energy of the 72 keV Sc⁵⁰ 2 \rightarrow 1 transition following the β ⁻ decay of Ca⁵⁰. The 72-keV Ca⁵⁰ γ ray is labeled as are various x rays from Pb and Bi. The two spectra shown are the middle two of the four sequential spectra as explained in the text. The energy resolution is 1.1 keV FWHM.

FIG. 4. 30-cc Ge(Li) γ -ray spectrum observed following bombardment of Ca⁴⁸F₂ with a 3.3-MeV triton beam. The original spectrum has been compressed and shifted to match the dispersion and bias of Fig. 1. The spectrum shown is the first of four obtained from the sum of 12 h of 100-sec cycles consisting of a 20-sec irradiation, four 18-sec counting intervals, and an 8-sec wait. γ rays are labeled by the parent activity and by their energies in keV. Two-escape peaks are labeled by energies (in keV) and by the symbol (2). The peak due to 511-keV annihilation radiation is not shown, since it overflowed. The Na²⁴ peaks are from Na²³(t, d)Na²⁴ induced in Na²³ impurities on the collimator system. The Al^{28} activity is from $Al^{27}(n, \gamma)$ in the detector housing, and the F^{20} and F^{21} activities are from $F^{19}+t$. The other peaks are discussed in the text.

indicated in Fig. 4. For the branching ratios of the Sc^{50} 1848-keV fourth excited state we find (42) $\pm 2\%$, (58 $\pm 2\%$, and <3% for branches to the first, second, and third excited states, respectively. These results are in good agreement with those of CMN⁵ which were $(37 \pm 7)\%$ and $(63 \pm 13)\%$ for the first two of these branches. No γ rays were observed which were attributable to any other $\beta^$ branches from Ca^{50} decay. However, the experiment was not very sensitive to weak branches and so we make no quantitative statements about limits on other branches but, for purposes of discussion, shall assume the decay proceeds 100% to the 1848 keV level.

C. $K^{47}(\beta^{\degree}) Ca^{47}$

The most intense γ ray attributable to the decay of K^{47} is that due to the Ca^{47} $1\rightarrow 0$ transition. This γ ray we find to have an energy of 2013.13 ± 0.30 keV and to decay with a half-life of 18.0 ± 1.0 sec (as compared with the expected values^{1,3} of 2016 ± 5 keV and 17.5 ± 0.3 sec, respectively). This is in agreement with Kuroyanagi et $al.$ ¹ who reported a 2.0-MeV γ ray from K⁴⁷. However, Kuroyanagi et al. reported a 2.6-MeV γ ray with an intensity of 15% relative to 85% for the 2.0-MeV γ ray, while we do not observe a 2.6-MeV γ ray and can set an intensity limit relative to that of the 2013 keV γ ray of <2% for such a radiation. We do observe two γ -ray peaks with energies of 564.74 ± 0.30 and 585.75 ± 0.30 keV and half-lives of 17.2 ± 2.1 and 18.1 ± 1.0 sec, respectively, which we attribute to the decay of the $Ca⁴⁷$ second and third

excited states to the first excited state following $K^{47}(\beta^-)Ca^{47}$. The deduced excitation energies for the Ca^{47} second and third excited states (see Table III and Fig. 5) are in good agreement with the previous values³ of 2580 ± 5 and 2601 ± 5 keV. We surmise that the 2.6-MeV peak observed by Kuroyanagi et al ¹ with a NaI(T1) detector was possibly due to summing of the Ca^{47} cascade radiations from the second and third excited states through the first excited state.

From the relative Ca⁴⁷ γ -ray intensities deduced from spectra such as Fig. 4, we deduce that the ground- state branches from the 2.58- and 2.60-

FIG. 5. Decay scheme of K^{47} . All energies are in keV. The spin-parity assignments are our final conclusions based on this and previous work as discussed in the text.

MeV levels are $\leq 28\%$ and $\leq 2.5\%$, respectively. The β^- branching ratios given in Table I are obtained assuming 100% cascade via the first excited state for both of these levels.

III. DISCUSSION

Our final results are summarized in Figs. 2 and 5 and in Tables I, II, and III. The results for the decay of Ca^{50} and Se^{50} are in good agreement with 5 and in Tables I, II, and III. The results for the decay of Ca^{50} and Sc^{50} are in good agreement with previous work,^{5,7} while those for the decay of K^{47} supercede the previous β ⁻ branching-ratio information¹ which is definitely in error.¹⁴ mation¹ which is definitely in error.¹⁴

Our results for $Ca^{50}(\beta^-)Sc^{50}$ add nothing new of theoretical significance. We note that the argument given by $CMN⁵$ in support of a 1⁺ assignment to the Sc^{50} 1848-keV level is unnecessarily weak. A stronger case for this assignment comes from the nature of the β^- -decay branch to this level. Since the transition to it is allowed and $0^+ \rightarrow 0^+$ transitions with $\Delta T = 1$ are forbidden, it necessaritransitions with ΔI =1 are iorpidated, it necessary has $J^{\pi} = 1^{+}$. The position of this 1⁺ level, the log ft value for decay to it, and its subsequent γ ray decay have been the subject of recent theoretiray decay have been the subject of recent theore
cal work.^{15,16} The theoretical predictions are in rather good agreement with experiment for the excitation energy^{15,16} and γ -ray decay modes,¹⁵ but
not so good for the β ⁻ decay rate.^{15,16} not so good for the β ⁻ decay rate.^{15,16}

As indicated in Figs. 1 and 2 the ground state and first three excited states of $Ti⁵⁰$ appear to form a $0^+, 2^+, 4^+, 6^+$ sequence.^{17,18} The spin-parity assignments for the first three are from various direct-reaction studies¹⁸ and appear to be quite firm. The 6' assignment for the third excited state is not as definite; however, it would be surstate is not as definite; however, it would be sur-
prising if it were incorrect.¹⁹ The indicated 5^+ assignment for the Sc^{50} ground state is also not definite. Since the β ⁻ branch of the Sc⁵⁰ ground state to the Ti⁵⁰ 3198-keV third excited state is allowed, these states have the same parity. Also, since the transition to the 4⁺ state has $|\Delta J| \le 1$, the Sc⁵⁰ ground state has $J^{\pi} = 5^{+}$ if the 3198-keV level has

 J^{π} = 6^+ . Both these latter assignments are pre-
dicted theoretically.^{15,16} Theoretical predictior dicted theoretically.^{15,16} Theoretical prediction for the β ⁻ decay of Sc⁵⁰ to the 4⁺ and 6⁺ states of for the β^- decay of Sc⁵⁰ to the 4⁺ and 6^+ states of Ti⁵⁰ are discussed by Hughes and Soga.¹⁵ Both the observed $\log ft$ values are considerably smaller than the predicted ones, but the ratio of the $\log ft$ values is in good agreement with the predictions.

The $K^{47}(\beta^-)Ca^{47}$ results summarized in Fig. 5 were expected. The fact that both decays are allowed and the K^{47} ground state⁴ is $\frac{1}{2}^+$ chooses for the 2578-keV level the $\frac{3}{2}$ ⁺ alternative allowed by the $l = 2$ assignment in the Ca⁴⁶ (d, p) Ca⁴⁷ reaction.³ If this $\frac{3}{2}$ ⁺ state were a pure $d_{3/2}$ -hole state, and the K^{47} ground state and Ca⁴⁷ 2599-keV level were pure $s_{1/2}$ -hole states, then the decay to the $\frac{3}{2}$ state (2578 keV) would be forbidden and the decay to the $\frac{1}{2}^+$ state (2599 keV) would have a log ft value roughg state (2599 keV) would have a log/t value roug
ly the same as that of $H^3(\beta^-)He^3$ (log/t = 3.0) or of
the $\frac{1}{2}^+$ $\rightarrow \frac{1}{2}^+$ decay, Ne¹⁹(β^-)F¹⁹ (log/t = 3.3). In acthe $\frac{1}{2}^+$ $\rightarrow \frac{1}{2}^+$ decay, Ne¹⁹(β^-)F¹⁹ (log *ft* = 3.3). In acthe $\frac{1}{2}$ – $\frac{1}{2}$ decay, Ne¹⁹(β ⁻)F¹⁹ (log*ft* = 3.3). In tual fact, the $\frac{1}{2}$ ⁺ $\rightarrow \frac{3}{2}$ ⁺ decay, although relativel weak, is not forbidden and the $\frac{1}{2}$ ⁺ $\rightarrow \frac{1}{2}$ ⁺ decay is some 30-60 times slower than the presumed $s_{1/2}$ + $s_{1/2}$ He³ and Ne¹⁹ transitions. Thus, we have definite evidence for the fragmentation of the basic $2s_{1/2}$ -hole states constructed by removing a $2s_{1/2}$
nucleon from a Ca⁴⁸ core (and also a $\,$ Sc⁴⁸ core in nucleon from a Ca^{48} core (and also a Sc^{48} core in the case of Ca^{47}). In particular, there would seem to be considerable $2s_{1/2}$ -hole strength at higher excitation energies in Ca^{47} and K^{47} . This conclusion is not inconsistent with the $Ca^{48}(d, t)Ca^{47}$ results of Yntema²⁰ and the Ca⁴⁸(d, He³)K⁴⁷ results of Newman and Hiebert.⁴

There seems to be some difficulty in understanding the properties of the low-lying even-parity ing the properties of the low-lying even-parity
states of mass $47.^{21}$ The β ⁻ decay rates given here should provide important data for the testing of any future predictions for these states.

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Neutron Emission from the Zn^{64} Compound Nucleus Formed in Two Ways: $p + Cu^{63}$ and $\alpha + Ni^{60}$ ⁺

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The Zn^{64} compound nucleus was formed at an excitation enerry of 33.3 MeV using two different target-projectile systems: 26.0-MeV protons on Cu⁶³ and 31.3-MeV α particles on Ni⁶⁰. Neutrons were detected with nuclear emulsions using the internal-radiator method. Exposures were made simultaneously at the following angles: 18, 33, 90, 147, and 162°. The c.m. energy spectra $(2-15 \text{ MeV})$ of the two systems are very nearly the same. For both systems symmetry about 90° in the c.m. angular distributions is seen up to about 6 MeV followed by a forward peaking at higher energies. The symmetric (compound-nuclear) parts of the angular distributions differ significantly in anisotropy. For the $p + Cu^{63}$ system the distribution is essentially isotropic while for the $\alpha + Ni^{60}$ system we have an average anisotropy of about 1.4. We also observe for the α + Ni⁶⁰ system a slight but significant increase in anisotropy with energy of the emitted neutron $(1.290 \pm 0.047 \text{ at } 2 \text{ MeV to } 1.649 \pm 0.119 \text{ at } 6 \text{ MeV})$. The changes in average anisotropy, as we change target-projectile systems, are basically consistent with multiple-emission calculations using a rigid-sphere moment of inertia $(r_0 = 1.22 \text{ F})$. The change in anisotropy with emission energy is not seen in the calculations. It is estimated that, for the $p + Cu^{63}$ and $\alpha + Ni^{60}$ systems, direct-reaction neutrons constitute 5 and 3% of the total observed neutron spectra.

I. INTRODUCTION

In order to isolate and study the role of angular momentum in the statistical decay of a compound nucleus, one often turns to a "Ghoshal-type experiment." Since Ghoshal did his classic work, in which he investigated the decay of the Zn^{64} compound nucleus formed with α particles and pro $tons¹$ many other systems have undergone similar analysis.² The typical output of such experiments is usually of the form of either excitation functions or particle spectra.

This work is focused on the Zn^{64} compound nu-

cleus. For this system the low-energy-reaction excitation functions have been well established for several projectile-target combinations, $1-9$ but the exact role of angular momentum is still uncertain.² Presented here are the results of a study in which the energy spectra (2-15 MeV) and angular distributions (18, 33, 90, 147, 162') of emitted neutrons were measured and calculated for the reaction systems $p + Cu^{63}$ and $\alpha + Ni^{60}$, both of which lead to the Zn^{64} compound nucleus at an excitation energy of 33.3 MeV.

In Sec. II we cover the experimental procedures used and present the results. In Sec. III the mea-