Magnetic moments and shape coexistence in the light Br isotopes

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Low temperature nuclear orientation measurements have been made to study the nuclear magnetic dipole moments of ⁷²Br^g,^m, ^{73,74}Br^m, ^{75,76}Br^g, and ⁷⁷Br^g implanted into Fe. The analysis of the data has been performed within the framework of a two, nonzero field, site model to describe the occupation of Br nuclei in the host Fe lattice. The two sites were associated with magnetic hyperfine fields of $\pm 81.38(6)$ and $\pm 26(2)$ T. A spin of $\frac{1}{2}$ for the ⁷³Br ground state is deduced and magnetic dipole moments are determined for the remaining nuclides. In addition, conversion electrons were observed for ⁷²Br, confirming the multipolarity of the isomeric transition. An interpretation of the ground-state configurations and shapes of these isotopes is given using particle-rotor calculations and it is shown that the Br nuclides can be better described as a hole in the more deformed Kr nuclides rather than a particle beyond the Se cores.

46

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I. INTRODUCTION

Among the nuclei off the line of stability, those in the $Z \sim N \sim 40$ region are of particular interest, exhibiting large quadrupole deformations, possible triaxiality, and shape coexistence [1-5]. The nuclear ground-state spins of the odd-A bromine nuclides $^{75-87}$ Br with Z = 35 are all $\frac{3}{2}^{-1}$. Nilsson orbital calculations show that the last odd proton occupies the $[301]\frac{3}{2}$ orbital for deformations $\varepsilon < 0.22$ and the $[312]\frac{3}{2}$ orbital at larger prolate deformations. The magnetic dipole moments of these two configurations differ by a factor of 3 and thus clearly identify the single-particle configuration and hence the approximate deformation of the ground state.

With this aim static low temperature nuclear orientation (LTNO) measurements have been performed on the light bromine isotopes ⁷²Br^{g,m}, ^{73,74}Br^m, ^{75,76}Br^g, and ⁷⁷Br^g. These data are analyzed in terms of the best currently available models and supersede all preliminary results quoted in earlier reports [6,7]. A detailed theoretical interpretation of the data is also discussed.

II. EXPERIMENTAL DETAILS

Sources of ⁷⁶Br^g ($t_{1/2} = 16$ h) and ⁷⁷Br^g ($t_{1/2} = 57$ h), produced in the reaction of 600-MeV protons on a target of niobium powder, were implanted at room temperature into polycrystalline iron foils at the ISOLDE facility, CERN. These were subsequently studied in Oxford at temperatures down to 2 mK using a ³He-⁴He dilution refrigerator in conjunction with a PrNi₅ demagnetization stage [8]. Data were collected from two Ge(Li) detectors positioned axially and equatorially relative to the axis of orientation, defined by a 1.0-T polarizing field. The source to detector distance was 12 cm.

The isotopes ${}^{72}\text{Br}{}^{g,m}$ ($t_{1/2} = 1.3$ min, 10.1 s, respectively), ${}^{73}\text{Br}$ ($t_{1/2} = 3.4$ min), ${}^{74}\text{Br}{}^m$ ($t_{1/2} = 46$ min), and ${}^{75}\text{Br}$ ($t_{1/2} = 97$ min) were produced on-line using the reaction of 150-MeV ${}^{28}\text{Si}$ on an ${}^{54}\text{Fe}$ target which formed part of the FEBIAD ion source of the Daresbury On-Line Isotope Separator (DOLIS). After acceleration to 60 keV, the selected bromine ions were implanted into a polycrystalline iron foil soldered to the cold finger of the dilution refrigerator and studied at temperatures down to 8 mK. Production rates were low for the lighter isotopes, but well resolved gamma spectra were recorded in four Ge(Li) detectors, two in both the axial and equatorial positions. The detector to source distance was 8 cm. The iron foil was polarized by a 0.7-T magnetic field in a direction parallel to its surface and perpendicular to the implanted beam.

III. DATA ANALYSIS

The implanted Br nuclei are oriented at low temperatures through the large magnetic hyperfine interaction that they experience in an iron host lattice. The resulting angular distribution of gamma radiation from such nuclei is given by

 $W(\theta) = 1 + \sum f_i \sum Q_k B_k(\mu B_i / IkT) U_k A_k P_k(\cos\theta) ,$

the symbols having their usual meaning [9]. The index i

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labels the various lattice sites which may be occupied by the implanted nuclei. The strength of the magnetic hyperfine interaction $\mu B_i/I$, and in particular the magnetic moment μ may then be determined from the temperature dependence of the anisotropy of the gamma-ray angular distribution, defined as [W(0)/W(90)-1]%.

Within certain model limitations this interaction may be extracted from the data with relatively little dependence upon the fractional populations f_i which, at moderately low temperatures, largely determine the absolute magnitude of the anisotropy but not its temperature dependence. This point will be discussed in more detail below. To facilitate the data analysis, two different models describing the lattice site occupation of the bromine nuclei in the iron matrix will be assumed. The first model, henceforth known as model 1, is that most applicable to soluble implants where a fraction f of nuclei experience the full substitutional hyperfine field B_{sub} , while the remaining 1-f occupy lattice sites with zero hyperfine field. As an alternative, and to account in a simple way for the relative insolubility of bromine in iron, we consider a second model, hereafter referred to as model 2. Here, as before, a fraction f of nuclei experience the full substitutional hyperfine field B_{sub} , but now it is assumed that the remaining 1-f occupy a second lattice site subject to a single nonzero hyperfine field B_{low} .

Herzog et al. [10], using NMR-ON techniques for ⁸²Br, have measured the substitutional magnetic hyperfine field of Br<u>Fe</u> to be $B_{sub} = +81.38(6)$ T. Using this field it was found that the magnetic moment extracted from a model 1 fit to the integral LTNO data, as obtained from a low temperature implanted sample, differed by 10% from the known moment. A satisfactory fit could, however, be made using model 2 with $B_{\text{low}} = \pm 26(2)$ T. A subsequent study for both room and low temperature implanted samples over a wide range of doses showed that it was generally necessary to invoke model 2 in order to reproduce the magnetic moment [11]. The variation in the value of B_{low} extracted from the various runs was no greater than would be expected from the statistical error of the value quoted above. No further complexity of site distribution can be meaningfully tested by the integral LTNO method. Extensive channeling studies by Alexander et al. [12], in the BrFe system at room temperature and relatively high implantation dose, support this multiple site picture. They concluded that there was indeed a second, well defined, nonsubstitutional site and that this location was independent of the local bromine concentration over a wide range. Building upon this work, Callaghan et al. [13] performed an integral LTNO experiment on ${}^{82}BrFe$ in order to evaluate the two hyperfine fields. On the reasoned assumption that the second field contributed little to the experimental anisotropy and could therefore be neglected, they determined the substitutional hyperfine field to be $\pm 84(12)$ T with an associated population of 36(5)%. Thus even using the crude model 1 they were able to predict the substitutional hyperfine field to within 15%.

On the basis of these results it can be expected that the model uncertainties will only influence the extracted moments at the level of 10-15%. Fitted moments obtained

from both of the above models will be quoted in this work. The values ultimately adopted, however, will be those of model 2 after assigning uncertainties which reflect the model site distribution uncertainties as well as the statistical uncertainties.

IV. RESULTS

A. Results for ^{76,77}Br^g

Both of these nuclei were studied prior to the publication of Refs. [10,11] and so it was not anticipated that the shortcomings of model 1 would be significant. Since the magnetic moment of ⁷⁶Br (1⁻) had already been measured [14] it was considered sufficient, and indeed desirable, to perform a less full temperature dependence for ⁷⁶Br in order to extract the $U_2 A_2$ coefficients which were not known in advance. To this end a PrNi5 demagnetization stage was used to cool the nuclei to 2 mK, as measured by a ⁶⁰Co<u>Ni</u> thermometer, which resulted in the saturation of the gamma-ray anisotropies. Unfortunately, these anisotropies were generally rather small, even close to saturation. Of the strongest lines, only the 1130-, 1854-, and 2951-keV transitions gave anisotropies with sufficiently favorable proportional errors to produce reasonable temperature dependences. The data on the 1130- and 2951-keV transitions are illustrated in Fig. 1.

Within the framework of model 1 a fit was performed to the experimental axial-equatorial anisotropies with both μB_{sub} and $f U_2 A_2$ as variable parameters. From



FIG. 1. The temperature dependence of the anisotropies of the 1130- and 2950-keV γ rays in the decay of ⁷⁶Br^g and the anisotropies of the 304- and 575-keV γ rays in the decay of ⁷⁷Br^g. Both model 1 (dashed) and model 2 (solid) fitted curves are shown for comparison.

these data the average hyperfine interaction was found to be $\mu B_{sub} = 35.0(25)\mu_N T$. This is to be compared with the expected value (corrected for applied field) of $\mu B_{sub} = 45.16(3)\mu_N T$.

This 25(6)% difference highlights once again the failure of model 1 to adequately describe the Br<u>Fe</u> system as first noted by Herzog *et al.* [10]. Following their lead therefore, the more realistic model 2 is invoked. Here, with μ required to reproduce the known value, $U_2 A_2$, f, and B_{1ow} remain as variable parameters. In order to reduce the number of unknowns, B_{1ow} was taken to be 26(2) T as discussed previously. In this way we obtain a site distribution parameter of f = 58(9)%.

In order to understand how model 2 is able to correctly reproduce the substitutional hyperfine interaction it is instructive to consider the individual contributions from the various lattice sites, as shown in Fig. 2. The anisotropy data from the 1130-, 1854-, and 2951-keV transitions have been inverted to yield $\sum_i f B_2 U_2 A_2$ and then normalized to the 14-mK temperature points, which for this purpose have been combined, and finally summed. The resulting data therefore represent the relative orientation of the ⁷⁶Br nuclei as a function of temperature. The orientation predicted by model 1, shown dashed, arises solely from the hyperfine interaction of nuclei in substitutional sites. It is clear that the true substitutional hyperfine interaction, $\mu B_{sub} = 45.2 \mu_N T$, increasingly underestimates the experimental orientation as the temperature is reduced. This gap between the observed orientation and that due to the fully substitutional lattice site may be bridged by introducing a second lattice site which experiences a nonzero hyperfine interaction. The individual contributions of the substitutional and low field sites, weighted by their relative populations, to the total model 2 orientation are illustrated by the solid lines. The discrepancy between the magnetic moments extracted using the two models may be attributed to the proportion of the total orientation arising from this second nonsub-



FIG. 2. The contributions to the total orientation from the model lattice sites. The data are taken from transitions in the decay of $^{76}Br^{m,g}$. The model 1 and model 2 fitted curves are distinguished by dashed and solid lines, respectively.

stitutional hyperfine interaction. In the present case this proportion is as high as 35% at the lowest temperature datum, which results in the large disparity between the two extracted moments. The magnetic moments of the other nuclei studied in this work have all been extracted from data taken at temperatures *above* 6 mK, where the relative contribution from the second field site is greatly reduced. The model differences and hence the ambiguity in the extracted moment will therefore be correspondingly smaller.

Having determined the site distribution parameter f = 58(9)% of ⁷⁶Br<u>Fe</u> it is assumed that the same value may be applied to the case of ⁷⁷Br. This is justified since both were implanted under the same conditions and with similar dosage. Further, it is likely that the experimental error in f is sufficiently large to take account of any minor variations between the samples.

In contrast to ⁷⁶Br^g, the magnetic moment of ⁷⁷Br $(\frac{3}{2}^{-})$ had not previously been measured and therefore we performed a full temperature dependence down to 6 mK as deduced from the anisotropy of a ⁶⁰Co<u>Co</u> thermometer.

In order to analyze these data, we take the model 1 variable parameters to be μ and fU_2A_2 , and the model 2 variable parameters to be μ and $U_2 A_2$, considering f and $B_{\rm low}$ as known. It should be noted that the site distribution parameter f is *not* the same in the two models. The temperature dependences of the 304- and 575-keV transitions are shown in Fig. 1. Using data from the ten strongest transitions we obtain an average model 1 moment of $\mu = 0.82(3)\mu_N$, while for model 2 we obtain a moment of $\mu = 0.92(5)\mu_N$. Notice that as a result of the higher temperatures there is only an 11(7)% disagreement between the two models. Comparing the χ^2 values of the two sets of model fits, which are to all intents and purposes identical, we find that even with data of such relatively good quality the integral LTNO method is unable to distinguish between the two models. In recent studies in which ⁷⁷Br has been studied by NMR methods, a more precise value of $0.9738(5)\mu_N$ has been reported, a value at the edge of the uncertainty of the integral LTNO method [15].

B. Results for ⁷²⁻⁷⁵Br

These isotopes were implanted on-line into a cold iron foil at the DOLIS-COLD facility where temperatures as low as 8 mK were achieved, as deduced from the anisotropy of a ⁵⁷Co<u>Fe</u> thermometer. Unlike the previous case, there is no control isotope with a known moment with which to determine f. Therefore within the framework of model 2 there are at least four unknown parameters, μ , f, $U_{\lambda}A_{\lambda}$, and B_{1ow} . In order to be able to extract reliable values for the magnetic moment from the fitting procedure it is desirable to reduce this number to two. Since the $U_{\lambda}A_{\lambda}$ coefficients are not generally known *a priori*, they must be considered as variable parameters. Therefore not only B_{1ow} but also a range of values for the site distribution parameter f is assumed.

The variation in f as a function of implantation conditions of ${}^{82}\text{Br}\underline{Fe}$ has also been studied [11]. It is apparent from this work that the value of f is considerably dependent upon implantation dose and also the method of production. However, the values for different on-line implantations were all found to lie within $58\% \le f \le 75\%$. Although some dependence upon the method of preparation of the iron foil might also be expected, this range will be considered to be valid for the present work. Since all iron foils used in the present experiments were prepared in an identical manner, the dependence upon the method of preparation of the iron foils will not lead to any problems with internal self-consistency. Thus the likely variation in f will be spanned if a maximum value of 75%, model 2a, and a minimum value of 58%, model 2b, is taken. It should be noted that the choice of 58% as the minimum value for the site distribution parameter in online samples coincides with the value deduced for the offline prepared samples ^{76,77}Br. That the site distribution parameter is probably lower for off-line than for on-line prepared samples is apparent from the data of Ref. [11] and is in line with our work on implanted BrFe [13].

1. Results for ⁷⁵Br

The presence in the DOLIS output of an inactive contaminant beam at this mass required that the 75 Br source be built up by implanting for 2.5 h at approximately 50 mK. The procedure was then followed by a period of cooling down to temperatures of 9 mK with the isotope separator closed off. The cycle was repeated several times.

With the exception of the 377-keV transition, which gave data of good statistical accuracy, the resulting anisotropies were rather small. The temperature dependence of this peak is illustrated in Fig. 3. In addition, and uniquely among all the isotopes studied, the $U_2 A_2$ coefficient can be calculated for this transition with sufficient accuracy to enable some constraint to be placed on the fitting procedure. The use of the previously known mixing ratio for the 377-keV transition, $\delta(377) = -0.73(45)$ [16], leads to the inequality $+0.62 \le U_2 A_2 \le +0.81$. The model 2 fitted $U_2 A_2$ coefficient is compatible with this result to within one standard deviation for all site distribution parameters $f \ge 44\%$, which greatly exceeds the likely range of variation given above. Therefore the limiting values 75% (model 2a) and 58% (model 2b) are retained, and with fthus fixed, the inequality for $U_2 A_2$ is used to limit the error on the fitted moment. In this way we find moments of $(0.73(9)\mu_N, 0.75(15)\mu_N, \text{ and } 0.78(17)\mu_N \text{ for models } 1, 2a,$ and 2b, respectively. [For comparison we give the unconstrained values of $0.73(9)\mu_N$, $0.75(19)\mu_N$, and $0.78(24)\mu_N$, respectively. Of course in model 1 this constraint has no effect on the fitted moment and merely serves to provide the inequality $f \ge 53\%$.]

Note that despite the large difference between the site distribution parameters of models 2a and 2b there is only a *very slight* difference in the fitted moments.

2. Results for ⁷⁴Br^m

The heavy ion reaction of 150-MeV ²⁸Si on ⁵⁴Fe preferentially produced the high-spin isomer of ⁷⁴Br. The pro-



FIG. 3. The temperature dependance of the anisotropies of the 377-, 728-, 862-, and 101-keV γ rays in the decays of ⁷⁵Br, ⁷⁴Br^m, ⁷²Br^m, and ⁷²Br^g, respectively. The model 1 and model 2 fitted curves coincide.

portion of the ground state present, as deduced from the intensity ratio of the 615- to 634-keV transitions, was less than 5% and therefore essentially negligible. Large an-isotropies with small statistical errors were observed for the 728-keV transition, shown in Fig. 3, and also for the 615-, 634-, 839-, 1201-, 1250-, and 1269-keV transitions.

The isomeric spin has been deduced by atomic beam resonance methods to be I=4 [17], although the assignment of negative parity to this state, made on the basis of proposed first unique forbidden β^+ /EC decays to levels in ⁷⁴Se, is rather tentative. Partly as a result of this condition, but mainly due to the complex decay scheme, no $U_{\lambda}A_{\lambda}$ coefficients are known or can be calculated *a priori*. Further, due to the relatively large parent spin, these coefficients involve two terms corresponding to $\lambda=2$ and 4.

As a first step towards dealing with this extra parameter the data were fitted to model 2a ignoring fourth rank terms. The weighted mean of these moments, for the seven transitions given above, is $1.64(5)\mu_N$ for which at 10 mK the ratio of the high field orientation parameters is $B_4(B_{sub})/B_2(B_{sub}) \sim 30\%$. Although this ratio is likely to become smaller when allowance is made for the relative effects of the two $U_\lambda A_\lambda$ terms, the fourth rank terms possibly constitute a sizable effect. We therefore also include these terms as a third variable in fits to the temperature dependences of the statistically dominant 634- and 728-keV transitions, from which we obtain moments of $1.63(6)\mu_N$, $1.59(8)\mu_N$, and $1.77(11)\mu_N$ for models 1, 2a, and 2b, respectively. Once again the relatively small

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difference between the model 2a and 2b moments demonstrates their limited sensitivity to the assumed site distribution parameter. Recent NMR studies have yielded a somewhat more precise value of 1.82(1) that is independent of the assumptions about site occupancy [15].

3. Results for ⁷³Br

Anisotropies were observed for 16 of the strongest transitions in the decay of ⁷³Br. These are listed in Table I. The anisotropy of the ⁵⁷Co<u>Fe</u> thermometer corresponds to an average temperature of 12 mK. All the ⁷³Br effects are consistent with zero orientation. Since the iron foil in which ⁷³Br was studied was the same as that used to successfully orient ^{72,74}Br^m in the same on-line run, there can be no possibility of a non-nuclear origin for this lack of anisotropy.

The ground-state spin of ⁷³Br had been limited to $(\frac{1}{2}, \frac{3}{2}, \frac{5}{2})^-$ as a consequence of its allowed β^+ /EC decay to the 26-keV $\frac{3}{2}^-$ level in ⁷³Se [18]. On the basis of systematics in the neighboring odd-*A* isotopes this spin has often been taken as $\frac{3}{2}^-$. However, the lack of anisotropy is at variance with this hypothesis.

If the ground state were $\frac{3}{2}^{-}$ and had a moment similar to that of ⁷⁵Br then the relaxation time would be of the order of 5 s (see below). When compared to the 3.4-m half-life it is clear that incomplete relaxation cannot be responsible for reducing the orientation to the virtually zero levels observed.

It must therefore be concluded that the lack of observed anisotropy is due to the presence of one or more unoriented spin- $\frac{1}{2}$ levels. The transitions in ⁷³Se originate from many different levels which are independently populated in the decay. A recent spectroscopic study of the decay of ⁷³Br [19] has provided spin assignments for some of the levels assuming that the parent spin is $\frac{3}{2}^{-}$. While most of these are tentative it is clear that not all of the levels which give rise to the observed gamma transitions are likely to be of spin $\frac{1}{2}$. It is therefore most probable

TABLE I. The gamma-ray angular distribution anisotropies observed in the decay of oriented 73 Br at 12 mK.

Energy (keV)	A (%) (σ)	Anisotropy/ <i>o</i>
65	0.6(0.6)	1.0
275	0.8(4.9)	0.2
336	0.8(1.6)	0.5
374	-1.6(5.2)	-0.3
401	-0.8(2.0)	-0.4
490	7.0(8.4)	0.8
540	-0.3(3.7)	-0.1
550	4.1(7.3)	0.6
615	-6.5(4.4)	-1.5
639	-6.5(10.3)	0.6
700	1.9(1.4)	1.4
788	-10.0(11.0)	-0.9
849	-0.3(1.1)	-0.3
870	6.6(4.8)	1.4
914	3.1(2.9)	1.1
931	1.5(1.5)	1.0

that at least a few of the relevant levels in ⁷³Se should be capable of orientation. In principle the lack of orientation might also be due to unfortunate values of the gamma multipole mixing ratios which lead to near zero angular distribution coefficients. This possibility is also inherently unlikely to be simultaneously true for several transitions. Two levels in ⁷³Se (other than the ground state) are in fact known to have spins other than $\frac{1}{2}$ [18]. Unfortunately neither figure prominently in this work. The 151-keV $\frac{5}{2}$ level gives rise only to the 126-keV transition which lies between the 122- and 136-keV peaks of the thermometer, ⁵⁷Co<u>Fe</u>. Although the triplet is resolved, the background is discontinuous making its subtraction difficult and ambiguous. However, a crude manual background subtraction showed no significant anisotropy for this transition. The second level at 26 keV $(\frac{3}{2})$, 40 min) is isometric and therefore orients in its own

The decay of the high-spin states of ⁷³Br has recently been studied by Heese *et al.* [20,21]. They have been able to provide a satisfactory interpretation of the level structure of ⁷³Br using a $\frac{1}{2}^{-}$ ground-state spin and parity assignment. Moreover, another recent study of the levels of ⁷¹Br has provided evidence for a $\frac{5}{2}^{-}$ ground state and a $\frac{1}{2}^{-}$ level at 9 keV in that nuclide [22]. Consequently, we conclude that it is in fact the parent nucleus, ⁷³Br, which has a ground-state spin $\frac{1}{2}^{-}$. Although such an assignment differs from the spin- $\frac{3}{2}^{-}$ ground states of all the heavier odd isotopes up to ⁸⁷Br, it is fully consistent with all known features of the decay, and with the decay of the higher-spin levels and the structure of ⁷¹Br.

4. Results for $^{72}Br^{g, m}$

The anisotropy of the strongest line in the decay of ⁷²Br^g, at 862 keV, is shown in Fig. 3. In this nuclide, there occurs the added complication that the relatively short half-life (1.3 m) may not allow sufficient time for the attainment of thermal equilibrium. Relaxation measurements by Herzog [10] on ⁸²Br<u>Fe</u> have yielded the result $g^2C_k = 0.032(2)$, where g is the spin g factor and C_k the Korringa constant [9]. The product g^2C_k is an isotope independent quantity which allows a self-consistent calculation to be performed on the data of ⁷²Br, taking into account the effects of imperfect relaxation on the observed anisotropies. The fitted line in Fig. 3 is calculated for a moment of 0.55(21) μ_N , for which the anisotropy attenuation at 12.5 mK is about 15%. This result is virtually independent of the choice of the model, the small differences lying well within the statistical uncertainties.

Finally, we consider the 101-keV transition from the 10.1-s isomeric state, to the ground state, which was assumed to be a 1⁻ to 3⁺ transition by Garcia-Bermudez et al. [23]. This spin parity change is confirmed by present γ -conversion electron coincidence measurements [24]. These data yield a conversion coefficient $\alpha_K = 1.4(3)$, implying an almost pure M2 transition, with less than 13% E3 admixture. Analysis of the NO data, shown in Fig. 3, was difficult owing to the large Compton background associated with the photopeak and the shape

2233

of the temperature dependence was ill defined, resulting in a fitted moment of $1.3^{+10.0}_{-0.5}\mu_N$. Again the model dependence is well within the uncertainties. Taking the measured anisotropy of -11(3)% at 15.1 mK and assuming the maximum magnitudes for f and A_2 of 100% and -0.2778, respectively, then we can set a probable lower limit on the moment of $0.7\mu_N$.

V. EXPERIMENTAL CONCLUSIONS

LTNO measurements on the light bromine nuclei have yielded the nuclear magnetic dipole moments of $^{72}\text{Br}^{g,m}$, $^{74}\text{Br}^m$, ^{75}Br , and $^{76,77}\text{Br}^g$. In addition, the ground-state spin of ^{73}Br has been established as $\frac{1}{2}^{-}$.

The experimental data have been analyzed within the framework of the two models described in Sec. III. The results of this analysis are summarized in Table II. It has been shown, both here for the case of ⁷⁶Br and by Herzog *et al.* [10,11] for the case of ⁸²Br, that model 1 fails to yield the correct moment. Therefore it is necessary to adopt the more flexible model 2. This approach requires an extra parameter, B_{10w} and also causes f and $U_{\lambda}A_{\lambda}$ to become independent. Since the $U_{\lambda}A_{\lambda}$ coefficients are generally not known *a priori* for the isotopes studied, a limited range of values for f has been assumed in order to reduce the number of free parameters to be extracted from the experimental data.

For ⁷⁷Br, f is taken to be the same as that deduced from the data on the control isotope ⁷⁶Br since the two were produced under similar conditions. For ⁷²⁻⁷⁵Br no such control exists and it is necessary to adopt a range of likely values for f, taken to be 58% $\leq f \leq$ 75% (models 2a and 2b, respectively). It has been shown that within this range the model 2 extracted moments have only a limited sensitivity to the value of f.

In addition, it is in principle necessary to consider also the effects of a variable low field B_{low} , since B_{low} need not correspond to a well defined field but may, instead, represent some average over many fields. However, the work of Herzog *et al.* [10,11] has given some evidence that B_{low} varies by no more than 2 T between different warm or cold implantations The value of f, on the other hand, varied by 20% between warm and cold implantations and so it is clear that f and not B_{low} should be treated as the variable parameter.

In view of the proven inability of model 1 to correctly

describe the experimental data the most reliable moments are taken to be those derived from model 2. In the case of ⁷⁵Br, where both models 2a and 2b have been applied, the mean of the two results is taken with errors chosen so as to encompass the range of values emerging from the two models. Effectively this procedure corresponds to the assumption that f = 67(9)%. The resulting increase in the uncertainty is relatively modest despite realistically reflecting the site distribution uncertainties. These moments along with the values from NMR are listed in Table II. The more precise values are of assistance in giving credence to the integral measurements and do not change significantly the nuclear structure information that can be derived from the whole set of data.

VI. DISCUSSION

A. The odd-mass Br nuclides

The odd-A bromine nuclei $^{75-87}$ Br, with atomic number 35, are all assigned to have ground-state spins of $\frac{3}{2}$. In-beam spectroscopic studies of ⁷³Br [3,20,21,25,26] have revealed a band sequence which bears a remarkable resemblance to the ground-state band of ⁷⁵Br. However, the last stretched E2 transition terminates not on the ground state but instead on an excited level at 178 keV which has therefore been assigned as $\frac{3}{2}^{-}$. Recent data on ⁷¹Br show a similar situation with the band terminating at a level at 207 keV [22,27]. From Fig. 4 it can be seen that for prolate nuclei the 35th proton is indeed expected to occupy the $[301]\frac{3}{2}$ orbital for quadrupole deformations up to $\varepsilon < 0.22$ and the $[312]\frac{3}{2}$ orbital at larger deformations in the range $0.30 < \varepsilon < 0.40$. As these levels are associated with theoretical magnetic dipole moments of approximately +2.3 μ_N and +0.75 μ_N , respectively, the experimental magnetic moments of the $\frac{3}{2}$ ground states give a clear signature of the orbitals involved and hence the magnitude of the nuclear deformation.

Because the orbitals of interest to the odd-A bromine ground states have the same component of angular momentum and can therefore intermix, the transition between high and low moment is not sharp, but instead has a definite deformation dependence. This variation of the moment with deformation is most sensitive in the region $0.22 < \varepsilon < 0.30$ where, with increasing deformation, the Fermi level shifts from the $[301]\frac{3}{2}$ to the $[312]\frac{3}{2}$ orbital

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Nuclide						
	Model 1	Model 2a $(f=75\%)$	Model 2b $(f=58\%)$	Adopted integral value	NMR momen	
⁷⁷ Br	0.82(2)		0.92(5)	0.92(5)	0.9738(5) ^b	
⁷⁶ Br	0.43(3)		0.55	$\pm 0.55^{a}$	$0.5482(1)^{a}$	
⁷⁵ Br	0.73(9)	0.75(15)	0.78(17)	0.76(18)		
74 Br ^m	1.63(6)	1.59(8)	1.77(11)	1.68(18)	$1.82(1)^{b}$	
⁷² Br ^g	0.55(21)	0.55(21)	0.55(21)	0.55(21)		
72 Br ^m	> 0.7	> 0.7	> 0.7	>7		

TABLE II. A summary of the fitted moments of $^{72-77}$ Br.

^aReference [14].

^bReference [15].



FIG. 4. Proton single-particle levels in the $A \sim 75$ region as a function of quadrupole deformation ε . The approximate position of the Fermi level in the Br nuclides is indicated by the heavy line. The Nilsson modified oscillator parameters $\kappa = 0.068$ and $\mu = 0.38$ are used. The dotted lines are the positive parity levels and the solid lines are the negative parity levels.

via the $[440]\frac{1}{2}$ state. As a consequence, the dominant single-particle contribution to the total wave function changes between the two configurations and is accompanied by a smooth and rather rapid drop in the magnetic moment from $+2.3\mu_N$ to $+0.75\mu_N$. The variation of the ground-state moment with deformation is illustrated in Fig. 5. Only the positions of the negative parity $[301]\frac{3}{2}$, $[312]\frac{3}{2}$, and $[303]\frac{7}{2}$ bandheads, together with the $\frac{1}{2}^{-}$ and $\frac{3}{2}^{-}$ members of the $[310]\frac{1}{2}$ band, are shown.

To develop these ideas of ground-state evolution in Br isotopes, we performed two types of model calculation including low-energy states.

In the first type of calculation, the Lund particletriaxial rotor model [28] with the modified oscillator single-particle potential was used to fit experimental data



FIG. 5. Deformation dependence of the ground-state magnetic dipole moment in the odd-*A* Br isotopes. The positions of the $[301]\frac{3}{2}^{\dagger}$, $[312]\frac{3}{2}^{\ddagger}$, and $[303]\frac{7}{2}$ bandheads, together with the $\frac{1}{2}$ and $\frac{3}{2}$ (*) members of the $[310]\frac{1}{2}$ band, are indicated. The magnetic moments of these three lowest lying $\frac{3}{2}^{-}$ states are given in μ_N .

on the energy levels and electromagnetic properties of negative parity low-energy levels in $^{71-81}$ Br in order to extract deformation parameters ε_2 and γ . The original version of this calculation VMI(A) [29] was performed taking the parameters $\kappa = 0.068$ and $\mu = 0.38$, equal for all oscillator shells. A variable moment of inertia was used with parameters J_0 and C that were calculated using the experimental 2_1^+ , 4_1^+ , and 2_2^+ level energies in the neighboring even-even Se nuclides [30]. We subsequently performed the same calculation with so-called "standard parameters" κ and μ [31] both with a constant moment of inertia [28] (PTR-MO) and with a variable moment of inertia [VMI(S)]. In the VMI(S) calculation, J_0 was estimated using deformation extracted from total Routhian surfaces (TRS) (see below) and C was obtained from the best fit to energy levels in odd-A Br isotopes. ε_2 and γ were the only deformation parameters employed in calculations using the modified oscillator potential. The effect of including ε_4 on the quality of the fit was examined and found negligible.

In the second type of calculation, we used the particletriaxial rotor model with Woods-Saxon single-particle potential (PTR-WS) and a constant irrotational moment of inertia in order to calculate level energies and electromagnetic properties of $^{71-81}$ Br with deformation parameters β_2 , β_4 , and γ extracted from TRS [32]. TRS were calculated using the deformed mean field approach with the Woods-Saxon potential, Strutinsky shell corrections, and a monopole pairing force [33].

In all of the above calculations, pairing was treated using the standard BCS approximation. In the determination of electromagnetic properties the effective g_s factor was taken to be $0.65g_{s\,free}$. Quadrupole moments of the core were calculated macroscopically and $g_R = Z/A$ was used. No attenuation of the Coriolis matrix elements was applied. There were thus no variable parameters in the PTR-WS calculation, only two variable parameters, ε_2 and γ , in the PTR-MO and VMI(A) calculations, and three, ε_2 , γ , and C in the VMI(S) calculation.

The TRS calculations give collective minima for $^{70-74}$ Se and $^{72-78}$ Kr and the corresponding deformation



FIG. 6. Experimental structures of the even-even Se nuclides.

parameters β_2 , β_4 , and γ as a function of rotational frequency ω . The experimental energy level structure of these nuclides is shown in Figs. 6 and 7. For all these nuclei the calculations predict, for ω close to zero, coexisting prolate and oblate shapes having different deformations (see Table III). No collective minimum is predicted in heavier Se (A = 76-80) and Kr (A = 80-82). Using the parameters from the TRS calculations, we made calculations for $^{71-75}$ Br for both Se and Kr cores and 77 Br for a Kr core only. For ^{79,81}Br, where no TRS data are available, we performed the PTR-WS calculation, fitting the deformation parameters to energy levels and the ground-state magnetic dipole and electric quadrupole moments.

Theoretical results are summarized in Table III showing the best fit deformation parameters from different calculations for odd-A Br and the TRS results for even-even Se and Kr. Table IV displays experimental and calculated ground-state magnetic dipole and electric quadrupole moments together with $B(E2; \frac{7}{2})$ to $\frac{3}{2}$ and $B(E2; \frac{13}{2})$ to $\frac{9}{2}$) values in the $\frac{3}{2}$ yrast bands. Table V illustrates the calculated structure of the $\frac{3}{2}^{-}$ bandhead in $^{71-81}$ Br.

Our results point to a gradual increase in nuclear ground-state deformation from about $\varepsilon_2 = 0.20$, $\gamma = 0$ in ⁸¹Br to $\epsilon_2 = 0.33$, $\gamma = 0$ in ⁷⁵Br. The yrast $\frac{3}{2}^-$ prolate band in $^{\overline{71},73}$ Br still has strong deformation, $\varepsilon_2 > 0.30$. This trend is accompanied by a drop in the experimental magnetic dipole moment from $+2.3\mu_N$ to $+0.75\mu_N$ for $^{75-81}$ Br very well reproduced by theory (see Table IV). This trend indicates a transition of the single-particle Fermi level from the $[301]\frac{3}{2}$ to the $[312]\frac{3}{2}$ orbital. In Table V we see that the ground state of 81 Br is dominated by the $[301]\frac{3}{2}$ orbital, ⁷⁹Br is transitional, while the ground states of ^{77,75}Br, the 178-keV excited state of ⁷³Br, and the 207-keV excited state of ⁷¹Br are almost entirely of $[312]\frac{3}{2}$ character. We note that the asymptotic quantum numbers are used in this work only to label particular Nilsson orbitals. Therefore the quoted results represent the relative contributions from the two Nilsson levels and do not necessarily reflect the exact singleparticle configurations.

The $\frac{5}{2}^{-}$ ground state of ⁷¹Br [22] and the proposed $\frac{1}{2}^{-}$ ground state in ⁷³Br appear to arise from a fundamentally different structure. As can be seen from Fig. 8, there is no orbital lying near the position of the Fermi surface in bromine which would give rise to such a ground state at any prolate deformation. In particular, the $[310]\frac{1}{2}$ orbital cannot be placed on the Fermi surface without significant



 ${}^{72}_{36}\text{Kr}_{36} {}^{74}\text{Kr}_{38} {}^{76}\text{Kr}_{40} {}^{78}_{36}\text{Kr}_{42} {}^{80}_{36}\text{Kr}_{44} {}^{82}_{36}\text{Kr}_{46} {}^{84}_{36}\text{Kr}_{48} {}^{86}_{36}\text{Kr}_{50} {}^{88}_{38}\text{Kr}_{52} {}^{99}_{38}\text{Kr}_{54} {}^{92}_{36}\text{Kr}_{58} {}^{94}_{36}\text{Kr}_{58}$

0+ 2789

0 0+ 0

FIG. 7. Experimental structures of the even-even Kr nuclides.

<u>46</u>

TABLE III. Deformation parameters ε_2 . For each mass number, the first line shows prolate deformation (*P*) and the second line oblate deformation (*O*). For TRS ε_2 has been recalculated from β_2 in the original calculation.

A	TRS (Kr)	TRS (Se)	PTR-MO	PTR-WS	VMI(S)	VMI(A)
81 P			0.20	0.18	0.21	0.22
79 P			0.22	0.20	0.24	0.24
78 P	0.28					
0	0.21					
77 P			0.30		0.29	0.27
76 P	0.31					
0	0.23					
75 P			0.325		0.32	0.32
74 P	0.32	0.27				
0	0.27	0.21				
73 P		0.34			0.32	0.32
0		0.215				
72 P	0.31	0.29				
0	0.28	0.23				
71 P			0.31		0.31	
0			0.20			
70 P		0.23				
0		0.24				

rearrangement of the $3p_{1/2}$ and $3f_{5/2}$ subshells which then prevents the shift of the Fermi level between the $[301]\frac{3}{2}$ and $[312]\frac{3}{2}$ orbitals, which is crucial to the interpretation of the ground-state magnetic moments of $^{75-81}$ Br. The band arising from the $[440]\frac{1}{2}$ orbital is well known to have a $\frac{5}{2}^+$ bandhead and so cannot produce a spin- $\frac{1}{2}$ ground state.

In Fig. 9 we show the results of calculations, using the PTR-MO with constant moment of inertia, of the lowest energy levels with $I^{\pi} = \frac{1}{2}^{-}, \frac{3}{2}^{-}, \frac{5}{2}^{-}$ in ^{71,73}Br, as a function of γ , for three representative values of ε_2 . The calculation was performed for a large number of deformation values covering the region $0.20 \le \varepsilon_2 \le 0.34$. The results show that, for ⁷³Br, the $\frac{1}{2}^{-}$ ground state arises in the whole region of deformation, provided $\gamma \ge 30^{\circ}$. With decreasing ε_2 , γ approaches 60°. In ⁷¹Br, the $\frac{5}{2}^{-}$ ground state is first calculated for ε_2 =0.21 and $\gamma > 30^{\circ}$. Howev-

er, recent experimental data [21,22] provided another criterion to be satisfied, namely close proximity of a $\frac{5}{2}^{-}$ state (27 keV) to the $\frac{1}{2}^{-}$ ground state in ⁷³Br and of a $\frac{1}{2}^{-}$ state (9 keV) to the $\frac{5}{2}^{-}$ ground state in ⁷¹Br. Figure 6 shows that this level structure can be achieved in this deformation region for both nuclei only for γ close to 60° and that the two lowest-energy states in ^{71,73}Br can be best understood as pure oblate states. The structure of these states is very similar in both nuclides. The $\frac{1}{2}^{-}$ state has two major components, 75% (83%) [310] $\frac{1}{2}$ and 23% (15%) [321] $\frac{1}{2}$ in ⁷¹Br (⁷³Br). The makeup of the $\frac{5}{2}^{-}$ state is more complicated. It results from strong Coriolis interaction among orbitals of $2f_{5/2}$ and $3p_{3/2}$ spherical origin. The main component is 43% (45%) [312] $\frac{3}{2}$ with contributions from $\frac{5}{2}^{-}$ members of bands built on [321] $\frac{3}{2}$, [321] $\frac{1}{2}$, and [310] $\frac{1}{2}$ states 26% (24%), 18% (16%), and 10% (14%), in ⁷¹Br (⁷³Br). The calculated magnetic mo-

TABLE IV. Experimental and calculated magnetic dipole moments, electric quadrupole moments, and values of $B(E2; \frac{7}{2}^- \rightarrow \frac{3}{2}^-)$ and $B(E2; \frac{13}{2}^- \rightarrow \frac{9}{2}^-)$ for the $\frac{3}{2}^-$ band in $^{73-81}$ Br. Experimental data obtained from the current Nuclear Data Sheets and the present work. (a) $B(E2; \frac{7}{2}^- \rightarrow \frac{3}{2}^-)$ (W.u.); (b) $B(E2; \frac{13}{2}^- \rightarrow \frac{9}{2}^-)$ (W.u.).

A	exp	PTR-MO	PTR-WS (Kr)	PTR-WS (Se)	VMI(S)	VMI(A)
81 μ (μ_N)	+2.27	2.28	2.13 ^a		2.25	2.28
Q (eb)	+0.276(4)	0.26	0.27 ^a		0.28	0.28
79 μ (μ_N)	+2.11	2.14	2.01ª		2.16	2.11
Q (e b)	+0.331(4)	0.33	0.29 ^a		0.34	0.33
77 μ (μ_N)	+0.92(5)	0.93	0.70		1.02	0.93
(a)	35(9)	52	40		47	33
(b)	68(5)	106	80		99	81
75 μ (μ_N)	+0.76(18)	0.86	0.68	0.77	0.89	0.75
(a)	61(5)	64	53	37	61	60
(b)	148(+66, -34)	128	104	73	125	120
73 (a)	45(6)	72	57	42	60	60
(b)	168(+28,-21)	143	111	86	121	121

^aDeformation not available from TRS calculations (see text).

2237

TABLE V. Structure of the $\frac{3}{2}^{-}$ bandhead in $^{71-81}$ Br. Components of the total wave function are given in percent. (i) $[312]\frac{3}{2}$; (ii) $[301]\frac{3}{2}$.

A	PTR-MO	PTR-WS (Kr)	VMI(S)	VMI(A)
81 (i)	25	6	18	8
(ii)	67	88	74	84
79 (i)	46	20	47	33
(ii)	50	74	46	58
77 (i)	98	86	98	95
(ii)	< 1	1	< 1	4
75 (i)	98	92	98	98
(ii)	<1	<1	< 1	1
73 (i)	98	98	98	98
(ii)	< 1	<1	< 1	1
71 (i)	98	98	98	
(ii)	< 1	< 1	< 1	

ment of the $\frac{5}{2}^{-}$ ground state in ⁷¹Br for the proposed oblate shape is about $1.5\mu_N$.

Examination of Table III shows that the deformation parameters for prolate shapes fitted using PTR-MO (with both the constant and variable moment of inertia) agree very well with the corresponding TRS values for Kr cores, but show poorer agreement for Se cores. This result suggests that the actual core of the odd-A Br (for prolate shapes) is quite close to the shape of the neighboring even-even Kr nucleus and there is very little core polarization. Deformations found for oblate shapes in 71,73 Br are very similar to those calculated for oblate Se cores and are about 20% smaller than those for an oblate Kr core. The TRS calculations do not predict any deviation from axial symmetry in even-even Se and Kr, which is consistent with our findings that the negative parity states in odd-A Br can be described without a need for γ different from 0° or 60°. In summary, our finding of large prolate deformation of lighter odd-A Br isotopes starting to develop at ⁷⁷Br agrees very well with predictions of TRS calculations. Our fits suggest that the deformation of the oblate coexisting states in ^{71,73}Br is considerably smaller than that of the prolate states. Differences in extracted deformations for each odd-A nuclide in question are very small and reflect the effect of different models. It is quite remarkable how self-consistent are the results of all models used. The choice of moment of inertia is not expected to influence ground-state properties in the first approximation, but even the alternative choice of the single-particle parameters k, μ in the particle-rotor calculation makes very little difference in predictions of electromagnetic properties.

As a result of our analysis, a coherent description for the low-energy levels in $^{71-81}$ Br can be drawn. Going from 81 Br to 75 Br, we find a smooth change in prolate ground-state structure from $[301]\frac{3}{2}$ to almost pure $[312]\frac{3}{2}$ configuration. The structure in 71,73 Br can be understood in terms of shape coexistence between a less deformed oblate ground state and a more deformed prolate $\frac{3}{2}^{-}$ yrast band. In both the neighboring Se and Kr even-even nuclides, TRS calculations predict similar coexistence. It follows that there is a prolate to oblate ground-state shape transition between 75 Br and 73 Br.

B. The odd-odd Br nuclides

1. Structure of ⁷⁶Br

We now turn to the odd-odd bromine nuclei. The static electromagnetic moments of the 1⁻ ground state of ⁷⁶Br have been previously measured to be $\mu = \pm 0.5482(1)\mu_N$ and $Q = \pm 0.27(1)$ b [14]. The quadrupole moment implies a deformation of $\varepsilon = 0.31$, $\gamma = 0$



FIG. 8. Proton single-particle levels in the $A \sim 75$ mass region in the (ε, γ) plane. The Nilsson modified oscillator parameters $\kappa = 0.068$ and $\mu = 0.38$ are used.



FIG. 9. The relative positions of the lowest $\frac{1}{2}^{-}$, $\frac{3}{2}^{-}$, and $\frac{5}{2}^{-}$ bandheads in ^{71,73}Br calculated with the PTR-MO model with constant moment of inertia for three different values of ε_2 as a function of γ .

which fits in well with the systematics of the odd-A nuclear deformations given in Table IV. With this deformation, the particle-rotor model indeed leads to a $1^{-} \pi$ $[312]\frac{3}{2}\nu[422]\frac{5}{2}$ K=1 ground-state configuration, with predicted moments of $-0.77\mu_N$ and Q=+0.27 b. These values are consistent with the previous configuration assignment of Ekstrom *et al.* [17]. (Note that the $\pi[301]\frac{3}{2}\nu[422]\frac{5}{2}$ K=1 configuration is associated with a much larger magnetic moment of $-1.5\mu_N$ and, according to the Gallagher-Moszkowski selection rule, could not give rise to a 1^{-} ground state [34].)

2. Structure of ⁷⁴Br

As was mentioned previously, the parity assignment to the spin-4 isomeric state in ⁷⁴Br is not fully established. Negative parity has been proposed on the basis of supposed first unique forbidden β^+ /EC transitions to ⁷⁴Se [35] while positive parity has been suggested from a comparison with the 4⁺ and 4⁻ band structures in ⁷⁶Br [36,37]. In view of the fact that the corresponding neutron Fermi levels are virtually coincident in the region $0.3 < \varepsilon < 0.4$, as can be seen from Fig. 4, such similarities between the states of ^{74,76}Br might well be expected.

Considering first the case of negative parity, likely candidates for a 4⁻ isomeric state include the $\pi[312]\frac{3}{2}\nu[422]\frac{5}{2}$, $\pi[440]\frac{1}{2}\nu[301]\frac{3}{2}$, and $\pi[440]\frac{1}{2}\nu[303]\frac{5}{2}$ orbitals. However, the calculated moments of these configurations $-0.16\mu_N$, $+3.93\mu_N$, and $+4.20\mu_N$, respectively, are all incompatible with the experimental moment of $\pm 1.82(1)\mu_N$. The tentatively proposed $\pi[301]_2^3\nu[422]_2^5$ orbital [17], with a theoretical moment of $\pm 1.71\mu_N$, never comes close enough to the Fermi level to be a viable possibility.

By contrast, the moment of the positive parity $\pi[431]_{\frac{3}{2}}v[422]_{\frac{5}{2}}$ configuration is calculated to be $+1.77\mu_N$, in excellent agreement with the experimental value of $1.82(1)\mu_N$. Therefore the value of the measured magnetic moment strongly favors a 4⁺ assignment for the isomeric state. The positive parity assignment and the indication that the orbitals are both of positive parity are also favored by Holcomb *et al.* [37], on the basis of the fast *E*2 transitions that they observe, as well as on the similarity of the overall behavior of the band that is built on this isomer to those better defined bands in ⁷⁶Br [38,39].

3. Structure of ⁷²Br

Recent in-beam gamma spectroscopy studies on ⁷²Br have revealed three rotational bands [40]. Electric transition strengths within the band terminating on the 1⁻ isomeric state indicate a nuclear deformation in the region $|\varepsilon| \sim 0.30$. Shape coexistence at low spin is a real possibility, and interestingly, no band structure clearly associated with the 3⁺ ground state has been observed, reminiscent of the neighboring nucleus ⁷³Br. With this structure in mind, a search for a possible 3⁺ ground state was carried out for a mesh of prolate, oblate, and triaxial deformation points in the range $0.05 \le \varepsilon \le 0.40$, $\gamma = 0^{\circ}$, 15°, 25°, 35°, 45°, and 60°. Despite this extensive search no such ground state was found, nor did any low energy 3^+ states have moments consistent with the small measured value of $\pm 0.55(21)\mu_N$. A similar search for an isomeric 1^- state at large deformation also failed to produce an obvious candidate. At lower deformation, however, the above-mentioned $\pi [301] \frac{3}{2} \nu [422] \frac{5}{2}$ orbital [17] with a calculated moment of $\pm 1.71\mu_N$ is a possibility, but it is not favored by the Gallagher-Moszkowski rules.

On the other hand, examination of the structures of the adjacent odd-mass nuclides ⁷¹Br and ⁷¹Se [41] indicates a much less deformed shape for the possible 3⁺ ground state of ⁷²Br. As noted above, the observed structure of ⁷¹Br can be described by a nearly pure oblate structure calculation. Likewise, the calculations of Zhao support an oblate structure for the levels of ⁷¹Se [42]. Alternately, these states could be even less deformed and have shell model $f_{5/2}$ and $p_{1/2}$ configurations. The latter single-particle structure would offer some explanation as to why the deformed particle-rotor model could not provide an adequate description for the structure of ⁷²Br.

At small deformation where the multiplet splitting for odd-odd nuclides can be described by the long range quadrupole interaction and longer range spin-dipole interaction as suggested by Paar [43], there would be a low-energy $\pi f_{5/2} \nu f_{5/2}$ multiplet with spins ranging from 0^+ to 5^+ and two 2^+ , 3^+ doublets, the $\pi f_{5/2} \nu p_{1/2}$ and the other $\pi p_{1/2} v f_{5/2}$. For both of these doublets, the low-energy state would be the 3⁺ level. Empirical moments for these configurations are $2.14\mu_N$ and $0.74\mu_N$, respectively, using a $\pi f_{5/2}$ moment of $1.6\mu_N$ from the study of moments in odd-mass As nuclides [44-46]. Even using $\pi f_{5/2}$ the Schmidt limit value of 0.84 μ_N only brings the calculated $\pi f_{5/2} \nu p_{1/2}$ moment down to 1.42 μ_N . The measured moment of 0.55(21) μ_N is decisive between these two configurations in favor of the $\pi p_{1/2} v f_{5/2}$ configuration. The empirical moment for the 3^+ member of the $\pi f_{5/2} \nu f_{5/2}$ multiplet is 1.49 μ_N , indicating little contribution from that configuration. For the 1⁻ level, it can be seen that various combinations involving the $\frac{5}{2}^+$ level at 670 keV in ⁷¹Br and $\frac{3}{2}^-$ levels in ⁷¹Se would produce a 1^{-} level and that level would show a larger moment owing to the involvement of the positive parity $\frac{5}{2}^+$ level, whatever its origin. It is not at all clear why this possibility would be isomeric, however, and why it should lie at a lower energy than the 2^+ member of the fp doublets. It seems much more likely that the 1⁻ band is considerably more deformed than the 3^+ band and not well described by the Paar model.

VII. CONCLUSION

In the odd-A nuclei it has been shown that there is a transformation in the character of the ground-state configuration from the less deformed $[301]\frac{3}{2}$ orbital in ^{79,81}Br to the more deformed $[312]\frac{3}{2}$ orbital in ^{75,77}Br. The measured magnetic dipole moments are well reproduced by the theory and are fully consistent with the de-

formations suggested by the electric quadrupole data. While these magnetic moments are only compatible with prolate deformations, the ground-state spin of ⁷³Br can only be interpreted in terms of an oblate nuclear shape. This fact provides strong evidence in the bromine nuclei for the conjectured prolate-oblate shape transition [1]. (In fact the transition was predicted to occur as high as ⁷⁸Br which we have shown not to be the case.) The presence in ⁷³Br of a low-energy $\frac{3}{2}^{-1}$ rotational band, whose nature is similar to that of the ground-state band in ⁷⁵Br, indicates the occurrence of coexisting oblate and prolate shapes.

In the odd-odd nuclei, the moment of the ⁷⁶Br ground state demonstrates the involvement of a $\pi[312]_{\frac{3}{2}}^{\frac{3}{2}}$ orbital as observed in the neighboring odd-*A* nuclei, ^{75,77}Br. The magnetic moment of ⁷⁴Br^m also suggests a positive parity assignment for the spin-4 isomeric state and the transition rates in the band favor a combination of positive parity proton and neutron orbitals from the $g_{9/2}$ single-particle orbitals as opposed to combinations of the negative parity orbitals. The structure and moments for ⁷²Br suggest that the maximum deformation in Br lies in the region of ^{74,75,76}Br_{39,40,41} and that the lower mass Br nuclides revert to oblate and more spherical shape for $N \leq 38$.

The spectacular change in the moments for the $\frac{3}{2}^{-}$ ground states of the $_{35}$ Br nuclides, together with the large quantity of in-beam and decay data, aid in attributing nuclear deformation to the occupancy of the low-K downsloping positive parity orbitals from the $g_{9/2}$ proton and neutron orbitals. For those structures where there is little occupancy of either the proton or neutron $g_{9/2}$ orbitals as found for 68,70,71 Se_{34,36,37} deformation at low energy appears to be small and largely oblate in character. The odd-Z $_{33}$ As nuclides are an example of the fact that the moments for the $f_{5/2}$ levels increase as N approaches 40 where there is $g_{9/2}$ neutron orbitals are either full or empty, the moments move back to lower values nearer the Schmidt limits.

Thus, it is possible to observe that the Br nuclides lie exactly in the transition region between weakly deformed structures observed in the As and Se nuclides and the more deformed Kr and Rb nuclides. Our new particle-rotor calculations show that the addition of the 35th proton tips the balance at midshell toward deformation and that the N = 39,40,41 Br nuclides can be best described as a hole in the more deformed Kr nuclides rather than a particle coupled to the less deformed Se nuclides.

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