Time-resolved laser-induced fluorescence measurements of Rydberg states in LuI and comparison with theory

Dai, Zhenwen; Jiang, ZK; Xu, Huailiang; Zhang, Zhiguo; Svanberg, Sune; Biemont, E; Lefebvre, PH; Quinet, P

Published in:
Journal of Physics B: Atomic, Molecular and Optical Physics

DOI:
10.1088/0953-4075/36/3/306

Published: 2003-01-01

Citation for published version (APA):

General rights
Copyright and moral rights for the publications made accessible in the public portal are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

• Users may download and print one copy of any publication from the public portal for the purpose of private study or research.
• You may not further distribute the material or use it for any profit-making activity or commercial gain
• You may freely distribute the URL identifying the publication in the public portal

Take down policy
If you believe that this document breaches copyright please contact us providing details, and we will remove access to the work immediately and investigate your claim.
Time-resolved laser-induced fluorescence measurements of Rydberg states in Lu I and comparison with theory

This article has been downloaded from IOPscience. Please scroll down to see the full text article.
(http://iopscience.iop.org/0953-4075/36/3/306)

View the table of contents for this issue, or go to the journal homepage for more

Download details:
IP Address: 130.235.188.104
The article was downloaded on 01/07/2011 at 08:05

Please note that terms and conditions apply.
Time-resolved laser-induced fluorescence measurements of Rydberg states in Lu I and comparison with theory

Zhenwen Dai\(^1\), Jiang Zhankui\(^1\), Huailiang Xu\(^1\), Zhang Zhiguo\(^1,3\), S Svåberg\(^1\), E Biémont\(^4,5\), P H Lefèvre\(^4\) and P Quinet\(^4,5\)

\(^1\) Department of Physics, Lund Institute of Technology, PO Box 118, S-221 00 Lund, Sweden
\(^2\) Department of Physics, Jilin University, Changchun 130023, China
\(^3\) The Centre for Condensed Matter and Technology, Harbin Institute of Technology, Harbin 150001, China
\(^4\) IPNAS, Bâtiment B15, Université de Liège, Sart Tilman, B-4000 Liège, Belgium
\(^5\) Astrophysique et Spectroscopie, Université de Mons-Hainaut, 15, Rue de la Halle, B-7000 Mons, Belgium

Received 23 August 2002, in final form 13 December 2002
Published 23 January 2003
Online at stacks.iop.org/JPhysB/36/479

Abstract
Time-resolved laser-induced fluorescence measurements have been performed for ten odd Rydberg states of neutral lutetium, belonging to the \(6s^2 (1S)np(n = 8–9)\) and \(6s^2 (1S)nf(n = 5–8)\) series. For \(6s^2 (1S)8p\) and \(6s^2 (1S)7f\), the experimental lifetimes corresponding to the two \(J\) values within the doublet differ substantially. Comparison with theoretical values, calculated with extensive configuration interaction and core-polarization effects included, shows that the experimental trends are adequately reproduced both for \(np\) \((n = 8–9)\) and \(nf\) \((n = 5–8)\) states.

1. Introduction

Lutetium \((Z = 71)\) is the heaviest atom in the group of the lanthanides. \(^{175}\)Lu is by far (97\%) the dominant of the two stable isotopes present in the solar system, \(^{176}\)Lu representing only 3\%. Lu has been identified in the solar photosphere, and its abundance, which is one of the lowest of the rare-earth group, is deduced from two Lu II lines, located at 339.706 and 622.187 nm. The last revision of the solar lutetium abundance \((A_{Lu} = 0.06 \pm 0.07, \text{in the usual logarithmic scale, where the hydrogen abundance is equal to 12.00})\) is due to Bord et al (1998), and is in good agreement with the meteoritic result, \(A_{Lu} = 0.12 \pm 0.01, \text{according to the compilation of Anders and Grevesse (1989). This result, based on theoretical } f\text{-values, needs to be assessed by accurate experimental results.}\)

Despite its low cosmic abundance, lutetium has been found in different Ap stars in the form of Lu I or Lu II (see e.g. Jaschek and Brandi 1972 or Poli et al 1987), and the analysis of the stellar spectra requires accurate atomic data, the final aim being an improved knowledge...
of the chemical composition of the different types of stars and, in fine, input to the model of star formation.

The analysis of the high-resolution stellar spectra requires information about the atomic transition probabilities but also about the hyperfine structure (HFS) constants (Kurucz 1993), both isotopes of Lu showing HFS patterns (the nuclear spin being $I = 7/2$). The HFS and isotope effects in neutral lutetium have been investigated by different groups in the past: in particular, data have been published by Lin et al (1994), Reddy and Rao (1989), Kuhnert et al (1983) and Göbel (1970a, 1971).

Experimental lifetimes along the Rydberg series, particularly for higher-energy states, are a useful test of the theoretical models, the configuration interaction becoming increasingly complex when the myriad of close-lying high-energy levels is involved.

Transition probabilities or lifetimes in singly ionized lutetium have been the subject of a limited number of experimental or theoretical investigations in the 1970s: beam–foil spectroscopy analyses were published by Andersen and Sorensen (1974) and by Andersen et al (1975). More recently, experimental work based on selective laser excitation has been published by Den Hartog et al (1998) and by Fedchak et al (2000). On the theoretical side, a multiconfiguration Dirac–Fock study of the $6s^2\,1S_0$–$6s6p\,3P^\pi_0$, $1P^\pi_1$ transitions in the Yb isoelectronic sequence has been reported by Migdalek and Baylis (1987). Time-resolved laser-induced fluorescence studies in doubly ionized lutetium are also due to Biémont et al (1999) and to Fedchak et al (2000). Dipole polarizability and core-polarization effects have been studied by the same authors and by Migdalek (1980, 1982).

The transition probabilities in neutral lutetium are more poorly known and are still extremely scarce, despite the fact that some efforts have been devoted, during the last 30 years, to the investigation of this atom. A review paper on the resonance transitions was published by Doidge (1995a). The compilation of the Lu I lifetimes published by Blagoev and Komarovskii (1994) contains 42 values, but they concern only the levels of the configurations $5d6s6p$, $6s^2$ and $6s^2nd\ (n \geq 19)$. A double-resonance investigation of excited states is due to Göbel (1970b). Laser spectroscopic studies were reported by Kwiatkowski et al (1980), and Rydberg states were studied by laser multistep resonance ionization spectroscopy (Vidolova-Angelova et al 1992). Lifetime measurements in Lu I are due to Gorkshkov et al (1984) and Vidolova-Angelova (1992), but the most detailed and recent analysis has been published by Fedchak et al (2000), who performed time-resolved laser-induced fluorescence measurements on a slow beam of lutetium atoms. In the latter case, 22 odd-parity levels and four even-parity levels were measured.

For the different reasons stated above, we have decided to perform a new experimental investigation of some Rydberg states of neutral lutetium (i.e. the $6s^2(1S)n\ell (n = 8–9)$ and $6s^2(1S)n\ell (n = 5–8)$ states), using a laser-induced fluorescence technique. The measurements were compared to calculations performed with a relativistic Hartree–Fock (HFR) method (Cowan 1981) in order to assess the reliability of the theoretical model.

2. The atomic structure of neutral lutetium

The atomic spectrum of neutral lutetium, although partly based on old studies, is fairly well known (Martin et al 1978). In fact the NIST compilation is firstly based on levels deduced by Meggers and Scribner (1930, 1937), who made arc and spark observations in the 250–1077 nm region. Extensions of the analysis are due to Klinkenberg (1954) and Camus and Tomkins (1972), who made observations in the 228–370 nm region. Additional contributions are due to Pinnington (1963).

Neutral lutetium has a rather simple electronic structure, with three electrons outside a closed shell, formed by the 14 $f$ electrons. It appears, therefore, as an interesting case for
testing the accuracy of the theoretical models. The identified even and odd configurations are 5d6s2, 5d26s, 6s2n (n = 7–14), 6s2nd (n = 6–34), 5d6s7s and 6s2np (n = 6–27), 5d6s6p, 5d26p and 6s2nf (n = 5–35), respectively.

The NIST energy levels (Martin et al 1978) were used in the fitting procedure of the present work (see section 3).

3. Lifetime measurements

The lifetimes of nine levels of Lu I were measured in the present work, with the time-resolved laser-induced fluorescence technique. One-step excitation was used to excite these states due to the opposite parity of the ground configuration 5d6s2. The levels of interest have been excited according to the schemes summarized in table 1.

<table>
<thead>
<tr>
<th>Statea</th>
<th>Lower level (cm⁻¹)</th>
<th>Upper level (cm⁻¹)</th>
<th>Excitation</th>
<th>Schemeb</th>
<th>Detection</th>
</tr>
</thead>
<tbody>
<tr>
<td>6s²1(S)8p²P⁹/₁₂</td>
<td>0.0</td>
<td>36 808.76</td>
<td>271.67</td>
<td>2ω + AS</td>
<td>271.7</td>
</tr>
<tr>
<td>6s²1(S)8p²P⁷/₁₂</td>
<td>1993.92</td>
<td>37 131.38</td>
<td>284.59</td>
<td>2ω + AS</td>
<td>284.6</td>
</tr>
<tr>
<td>6s²1(S)6p²Po</td>
<td>1993.92</td>
<td>39 424.68</td>
<td>267.15</td>
<td>3ω + 2S</td>
<td>267.2</td>
</tr>
<tr>
<td>6s²1(S)5f²Fo</td>
<td>1993.92</td>
<td>36 633.31</td>
<td>288.68</td>
<td>2ω + AS</td>
<td>273.0</td>
</tr>
<tr>
<td>6s²1(S)5f²Fo</td>
<td>1993.92</td>
<td>36 644.12</td>
<td>288.59</td>
<td>2ω + AS</td>
<td>288.6</td>
</tr>
<tr>
<td>6s²1(S)6p²Po</td>
<td>0.0</td>
<td>39 212.61</td>
<td>255.02</td>
<td>3ω + 2S</td>
<td>255.0</td>
</tr>
<tr>
<td>6s²1(S)6p²Po</td>
<td>1993.92</td>
<td>39 220.17</td>
<td>268.62</td>
<td>3ω + 2S</td>
<td>268.6</td>
</tr>
<tr>
<td>6s²1(S)7f²Fo</td>
<td>0.0</td>
<td>40 626.82</td>
<td>246.14</td>
<td>3ω + 2S</td>
<td>246.1</td>
</tr>
<tr>
<td>6s²1(S)7f²Fo</td>
<td>1993.92</td>
<td>40 619.01</td>
<td>258.89</td>
<td>3ω + 2S</td>
<td>258.9</td>
</tr>
<tr>
<td>6s²1(S)8f²Fo</td>
<td>0.0</td>
<td>41 456.26</td>
<td>241.21</td>
<td>3ω + S</td>
<td>241.2</td>
</tr>
</tbody>
</table>

b 2ω means frequency doubling; 3ω means frequency tripling; AS, S and 2S stand for the anti-Stokes and the first- and the second-order Stokes components, respectively.

The experimental set-up used in our experiment is illustrated in figure 1. Lu atoms were produced in a laser-induced plasma, using 532 nm wavelength laser pulses emitted from a 10 Hz repetition rate and 10 ns duration Nd:YAG laser (Continuum Surelite), with variable pulse energy. Pulse energies in the range 2–10 mJ were normally used. The pulses were focused on the surface of a Lu foil, rotating in a vacuum chamber. In order to obtain the required excitation, another Q-switched and seeded injection Nd:YAG laser (Continuum NY-82), with a pulse duration of 8 ns, was combined with a stimulated Brillouin scattering compressor to shorten the pulses down to 1 ns. The laser was used to pump a dye laser (Continuum Nd-60), in which a DCM dye was used in the experiments. According to the excitation requirements, the second- or third-order harmonics of the dye laser were obtained employing a non-linear optical system, including a KDP crystal, a retarding plate and a BBO crystal. The measured levels were excited by the different Raman-shifted components of the harmonics, obtained in a cell with hydrogen at 10 bar, when the harmonics were focused into the cell. In the experiments, the excitation light was isolated with a prism and focused by a lens at a distance of about 1 cm above the foil. The two Nd:YAG lasers were triggered by a digital delay generator (Stanford Research System, model 535), and the delay between the ablation and excitation pulses was adjusted by the generator. The atoms produced by the ablation pulse were excited selectively
by the excitation laser beam crossing the plasma horizontally. The fluorescence, emitted from
the excited levels, was imaged by a fused-silica lens and focused on the entrance slit of a 1/8 m
monochromator. A Hamamatsu 1564U micro-channel plate was used for the detection.

The time-resolved signal was recorded and averaged with a digital transient recorder
(Tektronix model DSA 602), and the fluorescence decay curve was sent to a personal computer
for lifetime determination. The lifetime evaluation was performed by an exponential fit.

In order to be sure that the Lu I transitions of interest were indeed studied, the modification
of the fluorescence signal as a function of the delay time was investigated, the occurrence of a
maximum fluorescence signal appearing at longer delay times for Lu I than for various Lu ions.
This is because atoms have considerably lower speeds than ions in their flight from the ablation
site. The lower speed makes atoms less vulnerable to so-called flight-out-of-view effects. If
the detection efficiency is altered due to changing geometry during the signal observation time
interval, a pure exponential decay will not be observed. By using a wide monochromator slit
and optimizing its geometrical position, detrimental effects can be eliminated.

In order to obtain a sufficiently high signal-to-noise ratio, a decay curve was obtained by
averaging fluorescence photons from more than 2000 pulses.

For each level measured, more than seven fluorescence decay curves were recorded under
different experimental conditions, the delay time and intensities of the ablation and excitation
laser pulses being varied. Thus, the measurements were performed for different concentrations
and temperatures of the plasma. The evaluated lifetimes from the different curves were found
to agree well. This showed that there were no saturation, reabsorption or collisional effects
occurring, and the final result was formed by averaging the lifetime values. The nine lifetimes
measured are reported in the third column of table 2. The error bars reflect the statistical
scattering, but also a conservative estimate of the possible remaining systematic errors, adjusted
in such a way that an overall accuracy of about 10% is stated for each lifetime studied.
Table 2. Theoretical and experimental lifetime values in Lu I, and comparison with previous results. (c)–(e) HFR calculations with a scaling factor of 0.85, 0.60 and 0.50, respectively, for the $R^2$ integrals between $6s^2np$ ($n = 8$–9) and $5d6s6p$.

<table>
<thead>
<tr>
<th>Levels$^a$</th>
<th>Upper level (cm$^{-1}$)</th>
<th>This experiment</th>
<th>Previous experiments</th>
<th>Theoretical lifetimes (ns)</th>
</tr>
</thead>
<tbody>
<tr>
<td>$6s^2(1S)7p^2P_{1/2}^o$</td>
<td>29 430.90</td>
<td>9.2(5)$^b$, 8.6(8)$^c$</td>
<td>8.1</td>
<td>8.3</td>
</tr>
<tr>
<td>$6s^2(1S)7p^2P_{3/2}^o$</td>
<td>30 488.62</td>
<td>5.9(3)$^b$, 7.4(6)$^c$</td>
<td>6.3</td>
<td>6.5</td>
</tr>
<tr>
<td>$6s^2(1S)8p^2P_{1/2}^o$</td>
<td>36 808.76</td>
<td>44(4)</td>
<td>22</td>
<td>35</td>
</tr>
<tr>
<td>$6s^2(1S)8p^2P_{3/2}^o$</td>
<td>37 131.38</td>
<td>20.5(2.0)</td>
<td>9</td>
<td>12</td>
</tr>
<tr>
<td>$6s^2(1S)9p^2P_{1/2}^o$</td>
<td>39 321.96</td>
<td>82(10)</td>
<td>44</td>
<td>84</td>
</tr>
<tr>
<td>$6s^2(1S)9p^2P_{3/2}^o$</td>
<td>39 424.68</td>
<td>32(10)</td>
<td>138</td>
<td>142</td>
</tr>
<tr>
<td>$6s^2(1S)10p^2P_{1/2}^o$</td>
<td>40 661.02</td>
<td>89</td>
<td>95</td>
<td>96</td>
</tr>
<tr>
<td>$6s^2(1S)10p^2P_{3/2}^o$</td>
<td>40 735.33</td>
<td>89</td>
<td>95</td>
<td>96</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Levels$^a$</th>
<th>Upper level (cm$^{-1}$)</th>
<th>This experiment</th>
<th>Previous experiments</th>
<th>Theoretical lifetimes (ns)</th>
</tr>
</thead>
<tbody>
<tr>
<td>$6s^2(1S)5f^2F_{3/2}^o$</td>
<td>36 633.31</td>
<td>33.1(3.0)</td>
<td>30.2(1.5)$^b$, 31(2)$^b$</td>
<td>30</td>
</tr>
<tr>
<td>$6s^2(1S)5f^2F_{5/2}^o$</td>
<td>36 644.12</td>
<td>30.7(3.0)</td>
<td>30.3(1.5)$^b$, 29(2)$^b$</td>
<td>32</td>
</tr>
<tr>
<td>$6s^2(1S)6d^2F_{5/2}^o$</td>
<td>39 212.61</td>
<td>52(5)</td>
<td>52</td>
<td>52</td>
</tr>
<tr>
<td>$6s^2(1S)6d^2F_{7/2}^o$</td>
<td>39 220.17</td>
<td>50(5)</td>
<td>56</td>
<td>56</td>
</tr>
<tr>
<td>$6s^2(1S)7f^2F_{5/2}^o$</td>
<td>40 626.82</td>
<td>54(5)</td>
<td>68</td>
<td>68</td>
</tr>
<tr>
<td>$6s^2(1S)7f^2F_{7/2}^o$</td>
<td>40 619.01</td>
<td>66(7)</td>
<td>71</td>
<td>71</td>
</tr>
<tr>
<td>$6s^2(1S)8f^2F_{5/2}^o$</td>
<td>41 456.26</td>
<td>104(10)</td>
<td>127</td>
<td>127</td>
</tr>
<tr>
<td>$6s^2(1S)8f^2F_{7/2}^o$</td>
<td>41 460.08</td>
<td>133</td>
<td>133</td>
<td>133</td>
</tr>
<tr>
<td>$6s^2(1S)9g^2F_{5/2}^o$</td>
<td>42 000.72</td>
<td>83</td>
<td>83</td>
<td>83</td>
</tr>
<tr>
<td>$6s^2(1S)9g^2F_{7/2}^o$</td>
<td>42 006.00</td>
<td>87</td>
<td>87</td>
<td>87</td>
</tr>
</tbody>
</table>


4. The calculations

The theoretical HFR approach used was described previously by Fedchak et al (2000). Configuration interaction and core-polarization effects were taken into account in a detailed way. Configuration interaction was considered among the configurations $5d6s^2$, $6s^2ns$ ($n = 7$–10), $6s^2nd$ ($n = 6$–10), $6s^2ng$ ($n = 5$–10), $6s6p^2$, $5d^2ns$ ($n = 6$–10), $5d6ns$ ($n = 7$–10), $5d6nd$ ($n = 6$–10), $5d6np$ ($n = 6$–10), $5d^2nf$ ($n = 5$–10), $5d6np$ ($n = 6$–10) and $5d6nf$ ($n = 5$–10) (odd parity). These configuration lists extend considerably those considered in our previous work (Fedchak et al 2000). More precisely, we have added the even-parity configurations $6s^2ng$ ($n = 5$–10), $5d^2ns$ ($n = 8$–10), $5d^2nd$ ($n = 8$–10), $5d^2nf$ ($n = 8$–10) and $5d6nd$ ($n = 8$–10) and the odd-parity configurations $5d^2np$ ($n = 8$–10), $5d6np$ ($n = 8$–10) and $5d6nf$ ($n = 8$–10) in order to better represent the interaction along the $6s^2ns$, $6s^2np$, $6s^2nd$ and $6s^2nf$ Rydberg series up to $n = 10$.

The core-polarization effects were considered in the same way as in the HFR(A) approach of Fedchak et al (2000), since this model was found to lead to a very good agreement with the experimental lifetimes in the cases of $6s^27s$, $6s^27p$ and $6s^25f$ levels. More precisely, the core-polarization potential and the correction to the dipole operator, as described by Quinet et al (1999), were included in the model using the dipole polarizability of the ionic core, $\alpha_d$. 
it appears that the calculation of radiative lifetimes of the fine-structure variation of the lifetimes is correctly reproduced. Looking into the details, \( n \) (calculation (c)) and experiment (this work) is nearly constant along the mixing with the 5d6s

\[
\begin{align*}
\text{Slater integrals (taken from Vergès and Wyart (1978) and from the NIST compilation (Martin et al. 1976)) for the Lu IV ion, and a cut-off radius, } r_c, \text{ equal to 1.406 } a_0, \text{ which corresponds to the expectation value of } r \text{ for the outermost core orbital (5p)}^6, \text{ as calculated with Cowan’s codes.}
\end{align*}
\]

Using a well established least-squares fitting procedure, the radial parameter values were adjusted to obtain the best agreement between the calculated and the experimental energy levels taken from Vergès and Wyart (1978) and from the NIST compilation (Martin et al. 1978). The Slater integrals (\( F^j, G^k \) and \( R^l \)), not optimized semi-empirically, were scaled down by a factor equal to 0.85 (see, however, the discussion of section 5).

5. Results and discussion

The calculated lifetimes are given in the fifth, sixth and seventh columns of table 2, where they are compared with the experimental results obtained in the present work and with previous experimental data. The three sets of results, (c), (d) and (e), correspond to different scaling factors applied to some configuration interaction integrals (see the notes to table 2). When comparing these values (calculation (c)) with the laser measurements of the present work, a rather good agreement (from 0 to 30%) is observed along the 6s2np (\( n = 6–8 \)) Rydberg series. The calculated values remain a factor of two smaller than the experimental lifetimes for the 6s2np (\( n = 8, 9 \)) levels. It is interesting to observe that the ratio between theory (calculation (c)) and experiment (this work) is nearly constant along the np series, and that the fine-structure variation of the lifetimes is correctly reproduced. Looking into the details, it appears that the calculation of radiative lifetimes of the np (\( n = 8, 9 \)) levels is very sensitive to the mixing with the 5d6s(3D)6p 2P states. Indeed, when slightly changing the admixture of

<table>
<thead>
<tr>
<th>SF (^a)</th>
<th>Configuration</th>
<th>( J )</th>
<th>Composition (%)</th>
<th>Lifetime (ns)</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.85</td>
<td>6s28p</td>
<td>1/2</td>
<td>88.5 8p (^2)P + 4.5 5d6s(3D)6p (^2)P</td>
<td>21.6</td>
</tr>
<tr>
<td></td>
<td>6s28p</td>
<td>3/2</td>
<td>76.5 8p (^2)P + 11.6 5d6s(3D)6p (^2)P</td>
<td>8.9</td>
</tr>
<tr>
<td>0.75</td>
<td>6s28p</td>
<td>1/2</td>
<td>94.7 9p (^2)P + 0.8 5d6s(3D)6p (^2)P</td>
<td>82.1</td>
</tr>
<tr>
<td></td>
<td>6s28p</td>
<td>3/2</td>
<td>93.2 9p (^2)P + 1.7 5d6s(3D)6p (^2)P</td>
<td>44.2</td>
</tr>
<tr>
<td>0.65</td>
<td>6s28p</td>
<td>1/2</td>
<td>90.2 8p (^2)P + 3.7 5d6s(3D)6p (^2)P</td>
<td>25.5</td>
</tr>
<tr>
<td></td>
<td>6s28p</td>
<td>3/2</td>
<td>79.4 8p (^2)P + 10.0 5d6s(3D)6p (^2)P</td>
<td>9.8</td>
</tr>
<tr>
<td>0.60</td>
<td>6s28p</td>
<td>1/2</td>
<td>95.1 9p (^2)P + 0.6 5d6s(3D)6p (^2)P</td>
<td>101.6</td>
</tr>
<tr>
<td></td>
<td>6s28p</td>
<td>3/2</td>
<td>94.0 9p (^2)P + 1.3 5d6s(3D)6p (^2)P</td>
<td>56.4</td>
</tr>
<tr>
<td>0.50</td>
<td>6s28p</td>
<td>1/2</td>
<td>91.7 8p (^2)P + 2.6 5d6s(3D)6p (^2)P</td>
<td>31.1</td>
</tr>
<tr>
<td></td>
<td>6s28p</td>
<td>3/2</td>
<td>82.3 8p (^2)P + 8.3 5d6s(3D)6p (^2)P</td>
<td>11.2</td>
</tr>
<tr>
<td>0.45</td>
<td>6s28p</td>
<td>1/2</td>
<td>95.5 9p (^2)P + 0.4 5d6s(3D)6p (^2)P</td>
<td>126.8</td>
</tr>
<tr>
<td></td>
<td>6s28p</td>
<td>3/2</td>
<td>96.7 9p (^2)P + 0.7 5d6s(3D)6p (^2)P</td>
<td>73.3</td>
</tr>
<tr>
<td>0.40</td>
<td>6s28p</td>
<td>1/2</td>
<td>92.4 8p (^2)P + 2.4 5d6s(3D)6p (^2)P</td>
<td>34.9</td>
</tr>
<tr>
<td></td>
<td>6s28p</td>
<td>3/2</td>
<td>83.8 8p (^2)P + 7.4 5d6s(3D)6p (^2)P</td>
<td>12.2</td>
</tr>
<tr>
<td>0.35</td>
<td>6s28p</td>
<td>1/2</td>
<td>95.7 9p (^2)P + 0.3 5d6s(3D)6p (^2)P</td>
<td>142.0</td>
</tr>
<tr>
<td></td>
<td>6s28p</td>
<td>3/2</td>
<td>95.0 9p (^2)P + 0.8 5d6s(3D)6p (^2)P</td>
<td>84.2</td>
</tr>
<tr>
<td>0.30</td>
<td>6s28p</td>
<td>1/2</td>
<td>93.8 8p (^2)P + 1.6 5d6s(3D)6p (^2)P</td>
<td>45.7</td>
</tr>
<tr>
<td></td>
<td>6s28p</td>
<td>3/2</td>
<td>87.0 8p (^2)P + 5.6 5d6s(3D)6p (^2)P</td>
<td>15.2</td>
</tr>
<tr>
<td>0.25</td>
<td>6s28p</td>
<td>1/2</td>
<td>95.9 9p (^2)P + 0.2 5d6s(3D)6p (^2)P</td>
<td>178.3</td>
</tr>
<tr>
<td></td>
<td>6s28p</td>
<td>3/2</td>
<td>95.5 9p (^2)P + 0.5 5d6s(3D)6p (^2)P</td>
<td>112.9</td>
</tr>
</tbody>
</table>

\(^{a}\) Scaling factor applied to the configuration interaction integrals, \( R^k \), between the 6s2np (\( n = 8–9 \)) and 5d6s6p configurations.
this latter state in the eigenvector compositions of 6s²8p and 6s²9p levels (simply by changing the scaling factor applied to the $R^k$ integrals connecting these configurations, the cases (c), (d) and (e) corresponding to scaling factors of 0.85, 0.60 and 0.50, respectively), it appears that the calculated lifetimes for 8p and 9p are considerably modified, this variation even reaching a factor of two when the scaling factor is varied from 0.85 to 0.50. This corresponds in fact to a change of a few per cent in the 6s²8p and 6s²9p purities. The present discussion is clearly illustrated in table 3 and in figure 2, which show the variation of the lifetimes as a function of the eigenvector compositions.

It should be emphasized that the 6s²(1S)nt²F₉/₅,7/₅ levels are not dependent upon the choice of the scaling factor, as shown in table 2. The lifetimes of the 5d6s(3D)6p²P° levels, previously measured by Fedchak et al (2000), are much less sensitive than the 5d6s(3D)np²P° levels ($n = 8, 9$) to the choice of this scaling factor: indeed, using 0.50 instead of 0.85 would lead to $\tau = 6.0$ and $3.0 \text{ ns}$ for the levels 5d6s(3D)6p²P°⁰⁰/₅ (32 058.10 cm⁻¹) and 5d6s(3D)6p²P°⁵/₅ (34 436.49 cm⁻¹), values which are compatible with the published experimental results, $\tau = 4.9(0.2)$ and $3.7(0.2) \text{ ns}$, respectively.

An experimental lifetime of the 9p²P°⁰⁰/₅ state would also have been very valuable in view of the above considerations. However, in spite of a thorough search, laser-induced fluorescence could not be detected in the present experiments.
The theoretical lifetimes of the present work are also compared in table 2 with previous experimental measurements for the 7p and 5f levels, due to Fedchak et al (2000) and to Gorshkov et al (1984). The agreement observed is good for the four levels, the HFR result being located in between the two experimental results for 7p $^2P_3/2$.

Considering the experimental lifetimes and the theoretical branching fractions as obtained in the present work, it has been possible to deduce oscillator strengths for the transitions depopulating the levels of interest. In view of space limitations, the detailed results are not presented here. The extensive tables will be available in the database of astrophysical interest, DREAM, which is progressively created at the address http://www.unh.ac.be/~astro/dream.shtml.

Acknowledgments

This work was financially supported by the Swedish Natural Science Research Council and by the EU-IHP Access to Large-Scale Infrastructures Programme (contract HPRI-CT-1999-00041). Financial support from the Belgian FNRS is also acknowledged by EB and PQ. PHL is an FRRIA fellow. Financial support from the National Natural Science Foundation of China is acknowledged by Z-WD, JZ-K, H-LX and ZZ-G.

References

Blagoev K B and Komarovski V A 1994 At. Data Nucl. Data Tables 56 1
Camus P and Tomkins F S 1972 J. Physique 33 197
Dodge P S 1995b Spectrochim. Acta B 50 1421
Göbel L H 1970a Z. Naturf. A 26 611
Göbel L H 1971 Z. Naturf. A 26 1559
Jaschek M and Brandi E 1972 Astron. Astrophys. 20 233
Klinkenberg P F A 1954 Physica 21 53
Kurucz R L 1993 Phys. Scr. T 47 110
Martin W C, Zalubas R and Hagan L 1978 Atomic Energy Levels—The Rare Earth Elements NSRDS-NBS Circular 49

Z-W Dai et al