

Laser Spectroscopy Prelab

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Prelab Questions

1

$s = \frac{1}{2}$, always.

$5s^1$: $s \implies \ell = 0$ and since $j = |\ell - s|, \dots, \ell + s = \frac{1}{2}$.

Furthermore, $2s + 1 = 2$, so this gives us

$$^{2s+1}L_j \rightarrow 5^2S_{1/2}$$

$$5p^1: p \implies \ell = 1 \text{ and } j = |1 - \frac{1}{2}|, 1 + \frac{1}{2} = \frac{1}{2}, \frac{3}{2}$$

$$^{2s+1}L_j \rightarrow 5^2P_{1/2} \text{ and } 5^2P_{3/2}$$

2

The Hamiltonian associated with spin orbit coupling is given by equation (5):

$$H_{SO} = \zeta(r) \frac{\hbar^2}{2} [j(j+1) - \ell(\ell+1) - s(s+1)]$$

For the higher energy $5^2P_{3/2}$, $j = \frac{3}{2}, \ell = 1, s = \frac{1}{2}$, and we get an associated energy:

$$\begin{aligned} H_{SO} &= \zeta(r) \frac{\hbar^2}{2} \left[\frac{3}{2} \left(\frac{5}{2} \right) - 1(2) - \frac{1}{2} \left(\frac{3}{2} \right) \right] \\ &= \zeta(r) \frac{\hbar^2}{2} \left[\frac{15}{4} - \frac{8}{4} - \frac{3}{4} \right] \\ &= \zeta(r) \frac{\hbar^2}{2} \end{aligned}$$

For the lower energy $5^2P_{1/2}$, $j = \frac{1}{2}, \ell = 1, s = \frac{1}{2}$, and we get an associated energy:

$$\begin{aligned} H_{SO} &= \zeta(r) \frac{\hbar^2}{2} \left[\frac{1}{2} \left(\frac{3}{2} \right) - 1(2) - \frac{1}{2} \left(\frac{3}{2} \right) \right] \\ &= \zeta(r) \frac{\hbar^2}{2} [-2] \\ &= -\zeta(r) \hbar^2 \end{aligned}$$

To get the energy splitting, we take the difference between the two:

$$\begin{aligned} \Delta E &= \zeta(r) \frac{\hbar^2}{2} + \zeta(r) \hbar^2 \\ &= \zeta(r) \frac{3\hbar^2}{2} \end{aligned}$$

3

For ^{87}Rb , $I = \frac{3}{2}$, and $F = J + I$, with F taking on values $F = |J - I|, |J - I| + 1, \dots, J + I$

(1)

$5^2S_{1/2} \implies j = \frac{1}{2}$
So

$$F = \left| \frac{1}{2} - \frac{3}{2} \right|, \frac{1}{2} + \frac{3}{2} \\ = 1, 2$$

(2)

$5^2P_{1/2} \implies j = \frac{1}{2}$
So

$$F = \left| \frac{1}{2} - \frac{3}{2} \right|, \frac{1}{2} + \frac{3}{2} \\ = 1, 2$$

(3)

$5^2P_{3/2} \implies j = \frac{3}{2}$
So

$$F = \left| \frac{3}{2} - \frac{3}{2} \right|, \left| \frac{3}{2} - \frac{3}{2} \right| + 1, \left| \frac{3}{2} - \frac{3}{2} \right| + 2, \frac{3}{2} + \frac{3}{2} \\ = 0, 1, 2, 3$$

4

The presence of crossover resonances means that there are frequencies between the ‘standard’ transition frequencies of ^{87}Rb that can absorb light. I expect to observe 12 spectral lines in the transition from $5^2P_{3/2}$ to $5^2S_{1/2}$.

5

For the $5^2S_{1/2}$ state of ^{87}Rb , we have $I = \frac{3}{2}$, $J = \frac{1}{2}$, $F = 1$ or 2 . However, when we plug this into equation 10:

$$B \frac{\frac{3}{4}C(C+1) - I(I+1)J(J+1)}{2I(2I-1)J(2J-1)}$$

We notice that the denominator results in zero, due to the value of J : $2\frac{1}{2} - 1 = 0$. Since this term is a term in the total hyperfine interaction, we cannot have it diverge, or else we would have a divergent hyperfine transition! The only way to prevent this is if we force $B = 0$, regardless of our choice in $F = 1$ or 2 .