

# Atom Physics Review

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## Date 3.31

### Question 1

If one did the Franck-Hertz experiment on atomic hydrogen vapour, which lines would see if the maximum energy of the electrons were 12.5 eV? Give the  $\lambda$  values.

Solution:

Because

$$\Delta E = E_n - E_1 = \frac{E_1}{n^2} - E_1 = \frac{1 - n^2}{n^2} E_1 \leq 12.5 \text{ eV}$$

And

$$n = 1, E_1 = -13.6 \text{ eV}$$

Therefore we can get

$$n \leq 3$$

Thus  $n = 1, 2, 3$ , and there are three lines:

1. from 3rd to 2nd:

$$\lambda_{32} = \frac{hc}{E_3 - E_2} = 657 \text{ nm}$$

2. from 2nd to 1st:

$$\lambda_{21} = \frac{hc}{E_2 - E_1} = 122 \text{ nm}$$

3. from 3rd to 1st:

$$\lambda_{31} = \frac{hc}{E_3 - E_1} = 102 \text{ nm}$$

## Question 2

If an electron in hydrogen atom goes from the ground state to the state with  $n = 3$

1. Give the energy of the atom absorbed in this transition.
2. Which lines would one see when the atom transmits from the excited state with  $n = 3$  to the ground state?

Solution:

1.

$$\Delta E = E_3 - E_1 = \frac{E_1}{3^2} - E_1 = -13.6 \text{ eV} \left( \frac{1}{3^2} - 1 \right) = 12.09 \text{ eV}$$

2. There are three lines:

(a) from 3rd to 2nd:

$$\lambda_{32} = \frac{hc}{E_3 - E_2} = 657nm$$

(b) from 2nd to 1st:

$$\lambda_{21} = \frac{hc}{E_2 - E_1} = 122nm$$

(c) from 3rd to 1st:

$$\lambda_{31} = \frac{hc}{E_3 - E_1} = 102nm$$

# Date 5.7

## Question 1

Explain the symbols for the  $3^2D \Rightarrow 3^2P$  transition in sodium.  
How many lines can be expected in the spectrum?

Solution:

- for  $3^2D$

$$s = \frac{1}{2}, l = 2$$

so

$$j = \frac{5}{2}, \frac{3}{2} \Rightarrow 3^2D_{\frac{5}{2}, \frac{3}{2}}$$

- for  $3^2P$

$$s = \frac{1}{2}, l = 1$$

so

$$j = \frac{3}{2}, \frac{1}{2} \Rightarrow 3^2P_{\frac{3}{2}, \frac{1}{2}}$$

Because of **the selection rule for single electron:**

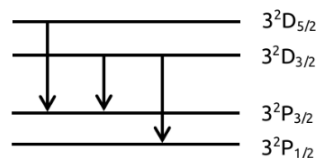
$$\begin{cases} \Delta l = \pm 1 \\ \Delta j = 0, \pm 1 \end{cases}$$

We can get three lines:

$$3^2D_{\frac{5}{2}} \Rightarrow 3^2P_{\frac{3}{2}}$$

$$3^2D_{\frac{3}{2}} \Rightarrow 3^2P_{\frac{3}{2}}$$

$$3^2D_{\frac{3}{2}} \Rightarrow 3^2P_{\frac{1}{2}}$$



## Question 2

Calculate the angle between the total and the orbital angular momenta in a  ${}^2F_{\frac{5}{2}}$  state.

Solution:

for  ${}^2F_{\frac{5}{2}}$  :

$$j = \frac{5}{2}, s = \frac{1}{2}, l = 3$$

$$L = \sqrt{l(l+1)} \frac{\hbar}{2\pi} = 2\sqrt{3}\hbar$$

$$S = \sqrt{s(s+1)} \frac{\hbar}{2\pi} = \frac{\sqrt{3}}{2}\hbar$$

$$J = \sqrt{j(j+1)} \frac{\hbar}{2\pi} = \frac{\sqrt{35}}{2}\hbar$$

because of **the law of cosines**:

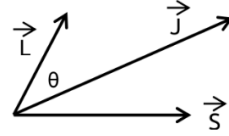
$$S^2 = L^2 + J^2 - 2LJ \cos \theta$$

then we can get

$$\begin{aligned} \cos \theta &= \frac{L^2 + J^2 - S^2}{2LJ} \\ &= \frac{12\hbar^2 + \frac{35}{4}\hbar^2 - \frac{3}{4}\hbar^2}{2 \times 2\sqrt{3} \times \frac{\sqrt{35}}{2}\hbar^2} \\ &= \frac{10}{\sqrt{105}} \end{aligned}$$

so

$$\theta = \arccos \frac{10}{\sqrt{105}}$$

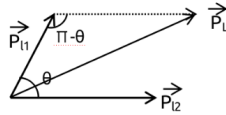


# Date 5.17

## Question 1

In a  $^3D$  state, that deduced from the electronic configuration 2p3d for the case of LS coupling.

1. Calculate the angle between  $\vec{P}_{l_1}$  and  $\vec{P}_{l_2}$
2. Calculate the angle between  $\vec{P}_{s_1}$  and  $\vec{P}_{s_2}$



1. for  $^3D$  (Note: you don't need to calculate  $J$ ):

$$S = 1, L = 2$$

for 2p:

$$s_1 = \frac{1}{2}, l_1 = 1$$

for 3d:

$$s_2 = \frac{1}{2}, l_2 = 2$$

$$P_L = \sqrt{L(L+1)}\hbar = \sqrt{6}\hbar$$

$$P_{l_1} = \sqrt{l_1(l_1+1)}\hbar = \sqrt{2}\hbar$$

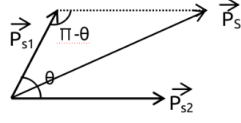
$$P_{l_2} = \sqrt{l_2(l_2+1)}\hbar = \sqrt{6}\hbar$$

$$\begin{aligned} P_L^2 &= P_{l_1}^2 + P_{l_2}^2 - 2P_{l_1}P_{l_2}\cos(\pi - \theta) \\ &= P_{l_1}^2 + P_{l_2}^2 + 2P_{l_1}P_{l_2}\cos\theta \end{aligned}$$

therefore

$$\cos\theta = \frac{P_L^2 - P_{l_1}^2 - P_{l_2}^2}{2P_{l_1}P_{l_2}} = -\frac{\sqrt{3}}{6}$$

$$\theta = \arccos\left(-\frac{\sqrt{3}}{6}\right)$$



2.

$$P_{s_1} = \sqrt{s_1(s_1 + 1)}\hbar = \frac{\sqrt{3}}{2}\hbar$$

$$P_{s_2} = \sqrt{s_2(s_2 + 1)}\hbar = \frac{\sqrt{3}}{2}\hbar$$

$$P_S = \sqrt{S(S + 1)}\hbar = \sqrt{2}\hbar$$

$$P_S^2 = P_{s_1}^2 + P_{s_2}^2 - 2P_{s_1}P_{s_2}\cos(\pi - \theta)$$

$$= P_{s_1}^2 + P_{s_2}^2 + 2P_{s_1}P_{s_2}\cos\theta$$

therefore

$$\cos\theta = \frac{P_S^2 - P_{s_1}^2 - P_{s_2}^2}{2P_{s_1}P_{s_2}} = \frac{1}{3}$$

$$\theta = \arccos\frac{1}{3}$$

## Question 2

In LS coupling, determine the number of possible terms of an excited carbon atom with the electronic configuration 2p3d? Which possible term is the ground state?

Solution:

$$\begin{array}{l} \text{for } 2p : s_1 = \frac{1}{2}, l_1 = 1 \\ \text{for } 3d : s_2 = \frac{1}{2}, l_2 = 2 \end{array} \Rightarrow \begin{cases} S = 1, 0 \\ L = 3, 2, 1 \end{cases}$$

$$\begin{cases} S = 0 \\ L = 3, 2, 1 \end{cases} \Rightarrow J = 3, 2, 1 \Rightarrow \begin{cases} {}^1F_3 \\ {}^1D_2 \\ {}^1P_1 \end{cases}$$

$$\begin{cases} S = 1 \\ L = 3 \end{cases} \Rightarrow J = 4, 3, 2 \Rightarrow {}^3F_{4,3,2}$$

$$\begin{cases} S = 1 \\ L = 2 \end{cases} \Rightarrow J = 3, 2, 1 \Rightarrow {}^3D_{3,2,1}$$

$$\begin{cases} S = 1 \\ L = 1 \end{cases} \Rightarrow J = 2, 1, 0 \Rightarrow {}^3P_{2,1,0}$$

So there are 12 possible terms.

According to **the Hund's rules**,  ${}^3F_2$  is the ground state.



## Date 5.22

### Question 1

Explain the symbols for the  $4^3F \Rightarrow 4^3D$  transition in sodium.  
How many lines can be expected in the spectrum?

Solution:

1. for  $4^3F$  :

$$s_1 = 1, l_1 = 3, j_1 = 4, 3, 2$$

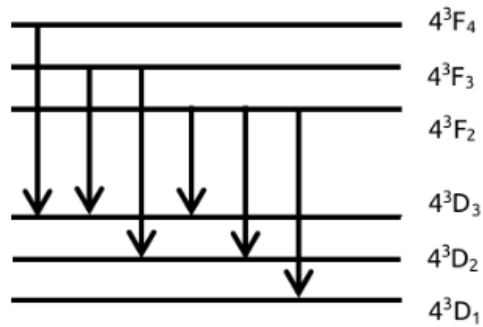
2. for  $4^3D$  :

$$s_2 = 1, l_2 = 3, j_2 = 3, 2, 1$$

According to **the selection rules for two-electron atoms:**

$$\begin{cases} \Delta S = 0 \\ \Delta L = 0, \pm 1 \\ \Delta J = 0, \pm 1 (J = 0 \not\Rightarrow J = 0) \end{cases}$$

We can get 6 lines:



## Question 2

1. Discuss a two-electron system with a 2p and a 3d electron for the case of **LS coupling** and show the number of possible state are the same as in **jj coupling**.
2. Calculate the ground state of the electron configuration 2p3d (in LS coupling).

Solution:

1. in LS coupling:

- for 2p

$$s_1 = \frac{1}{2}, l_1 = 1, j_1 = \frac{3}{2}, \frac{1}{2}$$

- for 3d

$$s_2 = \frac{1}{2}, l_2 = 2, j_2 = \frac{5}{2}, \frac{3}{2}$$

$$\Rightarrow \left\{ \begin{array}{l} S = 1, 0 \\ L = 3, 2, 1 \end{array} \right. \Rightarrow \left\{ \begin{array}{l} {}^1F_3, {}^1D_2, {}^1P_1 \\ {}^3F_{4,3,2} \\ {}^3D_{3,2,1} \\ {}^3P_{2,1,0} \end{array} \right.$$

So there are 12 states in LS coupling.

In jj coupling:

$$\left\{ \begin{array}{l} s_1 = s_2 = \frac{1}{2} \\ l_1 = 1 \\ l_2 = 2 \end{array} \right. \Rightarrow \left\{ \begin{array}{l} j_1 = \frac{1}{2}, \frac{3}{2} \\ j_2 = \frac{3}{2}, \frac{5}{2} \end{array} \right. \Rightarrow \left\{ \begin{array}{l} (\frac{1}{2}, \frac{3}{2})_{1,2} \\ (\frac{1}{2}, \frac{5}{2})_{2,3} \\ (\frac{3}{2}, \frac{3}{2})_{0,1,2,3} \\ (\frac{3}{2}, \frac{5}{2})_{1,2,3,4} \end{array} \right.$$

So there are also 12 states in jj coupling.

2. According to **the Hund's rules**, the ground state is  ${}^3F_2$  .

### Question 3

Ignoring spin-orbit coupling, determine the number of possible terms of an excited carbon atom with the electronic configuration  $1S^2 2S^2 2P^2 P$  ? Which possible term is the ground state?

Solution:

2p2p are **Equivalent Electrons**, we can calculate 2p3p first

$$\begin{matrix} s_1 = s_2 = \frac{1}{2}, \rightarrow S = 1, 0 \\ l_1 = l_2 = 1, \rightarrow L = 2, 1, 0 \end{matrix} \Rightarrow \begin{cases} {}^1D_2, {}^1P_1, {}^1S_0 \\ {}^3D_{3,2,1} \\ {}^3P_{2,1,0} \end{cases}$$

As for 2p2p, because of **the Paul Principle**: only the atomic states which satisfied

$$L + S = 2n(n = 0, 1, 2, \dots)$$

can exist. They are the following states:

$$\begin{cases} {}^1D_2 \\ {}^1S_0 \\ {}^3P_{2,1,0} \end{cases}$$

Finally, according to **the Hund's Rules**, ground state is  ${}^3P_0$

### Question 4

Calculate the effective magnetic moments of the states  $^1P_1$ ,  $^2P_{\frac{3}{2}}$ ,  $^4D_{\frac{1}{2}}$  and  $^2S_{\frac{1}{2}}$ .

Solution:

We need the following formulas:

$$g = 1 + \frac{j(j+1) - l(l+1) + s(s+1)}{2j(j+1)}$$

$$P_J = \sqrt{J(J+1)}\hbar$$

$$M_J = g \frac{e}{2m} P_J$$

1. for  $^1P_1$ :

$$s = 0, l = 1, j = 1$$

$$\Rightarrow g = 1, P_J = \sqrt{2}\hbar$$

$$\Rightarrow M_J = \frac{e\sqrt{2}}{2m}\hbar$$

2. for  $^2P_{\frac{3}{2}}$ :

$$s = \frac{1}{2}, l = 1, j = \frac{3}{2}$$

$$\Rightarrow g = \frac{4}{3}, P_J = \frac{\sqrt{15}}{2}\hbar$$

$$\Rightarrow M_J = \frac{\sqrt{15}e}{3m}\hbar$$

3. for  ${}^4D_{\frac{1}{2}}$ :

$$s = \frac{3}{2}, l = 2, j = \frac{1}{2}$$

$$\Rightarrow g = 0$$

$$\Rightarrow M_J = 0$$

4. for  ${}^2S_{\frac{1}{2}}$ :

$$s = \frac{1}{2}, l = 0, j = \frac{1}{2}$$

$$\Rightarrow g = 2, P_J = \frac{\sqrt{3}}{2}\hbar$$

$$\Rightarrow M_J = \frac{\sqrt{3}e}{2m}\hbar$$

## Date 6.17

### Question 1

1. Calculate the maximum components of the magnetic moments in the direction of the magnetic field for the vanadium ( $^4F$ ), manganese( $^6S$ ) and iron( $^5D$ ), if beams of these atoms are split into 6, 6, 9 parts in a Stern-Gerlach experiment.
2. What is the term symbol of the siglet state with a total splitting of  $\bar{\nu} = 1.4 \text{ cm}^{-1}$  in a magnetic field  $B_0 = 0.75$  tesla?

1. • for  $^4F$ :

$$S = \frac{3}{2}, L = 3$$

$$2J + 1 = 6 \Rightarrow J = \frac{5}{2}$$

$$g = 1 + \frac{J(J+1) - L(L+1) + S(S+1)}{2J(J+1)} = \frac{36}{35}$$

$$M_J = \pm\frac{1}{2}, \pm\frac{3}{2}, \pm\frac{5}{2}$$

$$M_{Jz} = M_J g \mu_B = \pm\frac{18}{35}\mu_B, \pm\frac{54}{35}\mu_B, \pm\frac{18}{7}\mu_B$$

$$M_{Jz}|_{max} = \frac{18}{7}\mu_B$$

- for  ${}^6S$ :

$$S = \frac{5}{2}, L = 0$$

$$2J + 1 = 6 \Rightarrow J = \frac{5}{2}$$

$$g = 2$$

$$M_J = \pm\frac{1}{2}, \pm\frac{3}{2}, \pm\frac{5}{2}$$

$$M_{Jz} = M_J g \mu_B = \pm\mu_B, \pm 3\mu_B, \pm 5\mu_B$$

$$M_{Jz}|_{max} = 5\mu_B$$

- for  ${}^5D$ :

$$S = 2, L = 2$$

$$2J + 1 = 9 \Rightarrow J = 4$$

$$g = \frac{3}{2}$$

$$M_J = 0, \pm 1, \pm 2, \pm 3, \pm 4$$

$$M_{Jz} = 0, \pm\frac{3}{2}\mu_B, \pm 3\mu_B, \pm\frac{9}{2}\mu_B, \pm 6\mu_B$$

$$M_{Jz}|_{max} = 6\mu_B$$

2. Because of the **singlet state**,  $2S + 1 = 1 \Rightarrow S = 0$

Therefore  $L = J$

$$g = 1 + \frac{J(J+1) - L(L+1) + S(S+1)}{2J(J+1)} = 1$$

As for total splitting:

$$\Delta E|_{max} = M_J|_{max} \mu_B B_0 = J \mu_B B_0$$

$$\Delta E|_{min} = M_J|_{min} \mu_B B_0 = -J \mu_B B_0$$

Before splitting, the energy is  $E_0$

After splitting:

$$(E_0 + \Delta E|_{max}) - (E_0 + \Delta E|_{min}) = 2J \mu_B B_0$$

Because

$$\Delta E = hc\bar{\nu}$$

Then

$$2J \mu_B B_0 = hc\bar{\nu}$$

$$J = L = 2$$

(Sorry I don't know how to calculate this  $\uparrow$ )

The state is  $^1D_2$

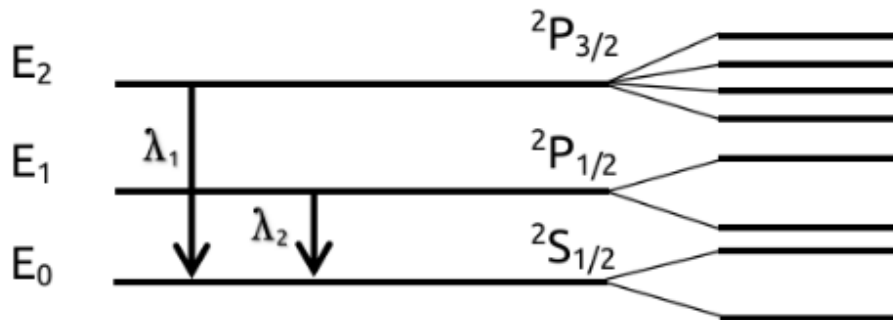


## Question 2

The special lines corresponding to the  $3p3s$  transition in sodium have the wavelengths  $\lambda_2 = 5895.9 \text{ \AA}$  and  $\lambda_1 = 5889.6 \text{ \AA}$

1. Calculate the magnetic field strength at which the lowest Zeeman level of the  ${}^2P_{3/2}$  term would coincide with the highest level of the  ${}^2P_{1/2}$  term. If the conditions for the anomalous Zeeman Effect were still fulfilled.
2. How large are the frequency differences between the outer two components of the two lines in a magnetic of the 1 tesla.

Solution:



1. • for  ${}^2P_{\frac{3}{2}}$ :

$$J_2 = \frac{3}{2}, S_2 = \frac{1}{2}, L_2 = 1, g_2 = \frac{4}{3}$$

- for  ${}^2P_{\frac{1}{2}}$ :

$$J_1 = \frac{1}{2}, S_1 = \frac{1}{2}, L_1 = 1, g_1 = \frac{2}{3}$$

- for  ${}^2S_{\frac{1}{2}}$

$$J_0 = \frac{1}{2}, S_0 = \frac{1}{2}, L_0 = 0, g_0 = 2$$

The energy of the lowest Zeeman Level of  ${}^2P_{\frac{3}{2}}$ :  $E_2 + \Delta E_2|_{min}$

The energy of the highest Zeeman Level of  ${}^2P_{\frac{1}{2}}$ :  $E_1 + \Delta E_1|_{max}$

According to the question:

$$E_2 + \Delta E_2|_{min} = E_1 + \Delta E_1|_{max}$$

- On the one hand

$$\begin{aligned} E_2 - E_1 &= \Delta E_1|_{max} - \Delta E_2|_{min} \\ &= M_{J_1} g_1 \mu_B B - M_{J_2} g_2 \mu_B B \\ &= [J_1 g_1 - (-J_2 g_2)] \mu_B B \\ &= \frac{7}{3} \mu_B B \end{aligned}$$

- On the other hand

$$\begin{aligned} E_2 - E_1 &= \frac{hc}{\lambda_1} - \frac{hc}{\lambda_2} \\ \Rightarrow B &= \frac{3hc}{7\mu_B} \left( \frac{1}{\lambda_1} - \frac{1}{\lambda_2} \right) \end{aligned}$$

2. (a)  ${}^2P_{\frac{3}{2}} \rightarrow {}^2S_{\frac{1}{2}}$

$$\begin{aligned}\Delta\tilde{\nu}_1 &= [M_{J_2} \cdot g_2 - M_{J_0} \cdot g_0]L \\ &= [\pm\frac{1}{3}, \pm\frac{3}{3}, \pm\frac{5}{3}]L \quad (\text{Rule } \Delta M_J = 0, \pm 1 \text{ on page 64})\end{aligned}$$

The outer two components:

$$\begin{aligned}\frac{1}{\lambda'_1} - \frac{1}{\lambda_1} &= \frac{5}{3}L \\ \frac{1}{\lambda'_2} - \frac{1}{\lambda_1} &= -\frac{5}{3}L\end{aligned}$$

The frequency difference:

$$\begin{aligned}\frac{c}{\lambda'_1} - \frac{c}{\lambda'_2} &= \frac{10c}{3}L \\ &= \frac{10c}{3} \cdot \frac{eB}{4\pi mc} \\ &= \frac{10B}{3} \cdot \frac{eh}{4\pi m} \cdot \frac{1}{h} \\ &= \frac{10B\mu_B}{3h} \\ &= \textcircled{9}\end{aligned}$$

(b)  ${}^2P_{\frac{1}{2}} \rightarrow {}^2S_{\frac{1}{2}}$

$$\begin{aligned}\Delta\tilde{\nu}_2 &= [M_{J_1} \cdot g_1 - M_{J_0} \cdot g_0]L \\ &= [\pm\frac{2}{3}, \pm\frac{4}{3}]L\end{aligned}$$

The outer two components:

$$\begin{aligned}\frac{1}{\lambda'_1} - \frac{1}{\lambda_2} &= \frac{4}{3}L \\ \frac{1}{\lambda'_2} - \frac{1}{\lambda_2} &= -\frac{4}{3}L\end{aligned}$$

The frequency difference:

$$\frac{c}{\lambda'_1} - \frac{c}{\lambda'_2} = \frac{8c}{3}L = \frac{8B\mu_B}{3h} = \textcircled{9}$$