

Intro to addition of angular momentum + spectroscopic notation

spin $1/2$: 2 electrons

how many ways can we combine spins?

$$|+\rangle|+\rangle \quad |+\rangle|-> \quad |->|+\rangle \quad |->|->$$

- in this representation, each spin is marked individually
 - can identify which electron is + or -
- we can ask: what is total spin?

$$\vec{S} = \vec{S}_1 + \vec{S}_2$$

as with general angular mom, total spin can be described with 2 quantum numbers

$$S, S_z$$

range of allowed S_z : $-S \dots +S$ w/ integer

S total: 0 or 1

w/ spacing

$$|S, S_z = 1, 1\rangle = |+\rangle|+\rangle$$

spins are parallel, both up

$$|1, -1\rangle = |->|->$$



two remaining:

$$|1, 0\rangle \text{ and } |0, 0\rangle$$

* we can't distinguish b/w particles

$$\rightarrow |1, 0\rangle = \frac{1}{\sqrt{2}} (|+\rangle|-> + |->|+\rangle) \text{ symmetric exchange}$$

$$|0, 0\rangle = \frac{1}{\sqrt{2}} (|+\rangle|-> - |->|+\rangle) \text{ antisymm.}$$

$|0, 0\rangle$ = "singlet state"

$|1, S_z\rangle$ = "triplet state"

general case:

$$\vec{J} = \vec{L}_1 + \vec{L}_2$$

J = total

\vec{L}_i = orbital or spin.

allowed ranges

$$j = |l_1 - l_2| \dots |l_1 + l_2| \text{ integer spacings.}$$

always > 0

$$j_z = -j \rightarrow +j \text{ integer spacings.}$$

examples:

$$l_1 = 1 \quad l_2 = l_2$$

$$\begin{array}{ll} j & j_z \\ \frac{1}{2} & \pm \frac{1}{2} \\ \frac{3}{2} & \pm \frac{3}{2}, \pm \frac{1}{2} \end{array}$$

$$l_1 = 2 \quad l_2 = 1$$

$$\begin{array}{ll} j & j_z \\ 1 & 0, \pm 1 \\ 2 & 0, \pm 1, \pm 2 \\ 3 & 0, \pm 1, \pm 2, \pm 3 \end{array}$$

each one of these states is a combination of individual $|l_1, l_{1z}\rangle |l_2, l_{2z}\rangle$ states.

* coeff: Clebsch-Gordan coeff.

How does spin state affect energy?

- electrons have spatial $\Psi(\vec{r})$

- symmetric combinations: e^- on avg closer \rightarrow higher energy.

Ex He $1s^2$ ground state

each e^- in same spatial Ψ : S

i. spin state must be antisymmetric
 \rightarrow Singlet $\rightarrow ^1S$

1st excited state

1s2s

can make symm or antisymm combinations
of spatial Ψ :

$$\Psi(r_1, r_2) = \frac{1}{\sqrt{2}} (\Psi_{1s}(\vec{r}_1) \Psi_{2s}(\vec{r}_2) + \Psi_{1s}(\vec{r}_2) \Psi_{2s}(\vec{r}_1))$$

overall state must be antisymmetric.

i.e. spin state can be singlet or triplet:

1S 3S

→ antisymmetric spatial

i.e. lower energy,