

$|s m\rangle$

Then, we have confirmed that indeed we form a **triplet** and a **singlet**, out of two spins $\frac{1}{2}$.

$$\left\{ \begin{array}{l} |1 1\rangle = \uparrow\uparrow \\ |1 0\rangle = \frac{1}{\sqrt{2}}(\uparrow\downarrow + \downarrow\uparrow) \\ |1 -1\rangle = \downarrow\downarrow \end{array} \right\} \quad s = 1 \quad \text{triplet}$$

$$\left\{ |0 0\rangle = \frac{1}{\sqrt{2}}(\uparrow\downarrow - \downarrow\uparrow) \right\} \quad s = 0 \quad \text{singlet}$$

Not in book (counting of states for 2 and 3 spins):

↑↑

↑↑

↑↓

$$\frac{1}{\sqrt{2}}(\uparrow\downarrow + \downarrow\uparrow)$$

↓↑

↓↓

↓↓

$$\frac{1}{\sqrt{2}}(\uparrow\downarrow - \downarrow\uparrow)$$

$2^2=4$ states
sort of
random

4 states grouped
as $S=1$ (3) and
 $S=0$ (1)

Note: after finding $S=1$, there was only 1 state left, thus had to be singlet and had to be orthogonal, thus fixing the "-"

↑↑↑

↑↑↓, ↑↓↑, ↓↑↑

↑↓↓, ↓↑↓, ↓↓↑

↓↓↓

$2^3 = 8$
states
sort of
random

↑↑↑, $S_{-}\uparrow\uparrow\uparrow$, $S^2_{-}\uparrow\uparrow\uparrow$, $S^3_{-}\uparrow\uparrow\uparrow$

4 states form S total $3/2$

The 4 states left form TWO
 S total $\frac{1}{2}$ states.

$3/2 \oplus 1/2 \oplus \frac{1}{2}$ (more about
this in a few pages)

Not in book (and FYI only):

FYI: spins can interact among themselves, not only with external magnetic fields.

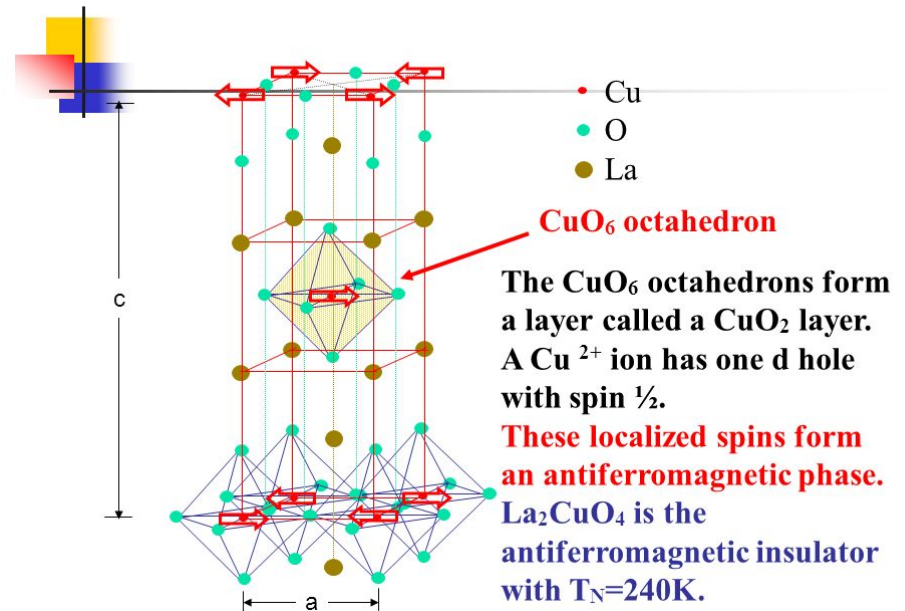
It is as if other spins "j" produce an **effective** magnetic field on the spin "i" you are looking at. This is the famous Heisenberg model:

$$H = J \sum_{\langle i,j \rangle} \mathbf{S}_i \cdot \mathbf{S}_j$$

Ground state? Number of states grows like 2^N (=2,4,8,16, ...)

Record done exactly $N \sim 40$.
 $2^{40} = 1,099,511,627,776$ states

Crystal structure of undoped La_2CuO_4



WITHOUT PROOF, this is what happens when you combine a spin s_1 and a spin s_2 (each individually $0, 1/2, 1, 3/2, \dots$).

The **total spin** s of the combination can be Eq.(4.182):

$$s = (s_1 + s_2), (s_1 + s_2 - 1), (s_1 + s_2 - 2), \dots, |s_1 - s_2|$$

Example 1: for $s_1=1/2$ and $s_2=1/2$, then s runs from $s_1+s_2=1$ to $|s_1-s_2|=0$, with nothing in between. We already confirmed this is true.

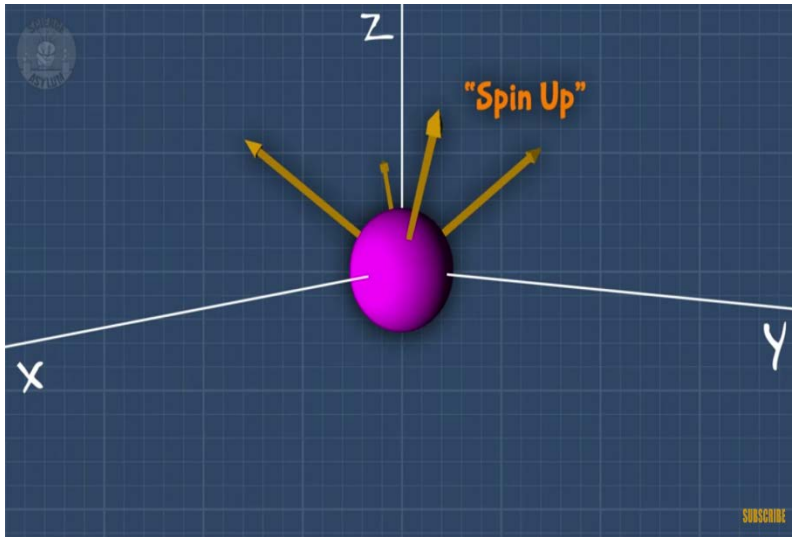
Example 2: for $s_1=3/2$ and $s_2=2$, then s runs from $s_1+s_2=7/2$ to $|s_1-s_2|=1/2$, with $5/2$ and $3/2$ in between.

Example 3: this unproven theorem holds also for the addition of **orbital angular momentum l** and **spin s** . For $l=2$ and $s=1/2$, then **total j** runs from $l+s=5/2$ to $|l-s|=3/2$, with nothing in between.

Example 4: if you have three particles with $s_1=1/2$, $s_2=1/2$, and $s_3=1/2$, then first you add two, such as s_1 and s_2 , finding $s_{\text{partial}} = 1, 0$ and then add s_{partial} with s_3 finding $3/2, 1/2$ (for $s_{\text{partial}}=1$) and another $1/2$ (for $s_{\text{partial}}=0$). So there are two independent combinations with total spin $\frac{1}{2}$. We already saw that in previous pages: $3/2 \oplus 1/2 \oplus 1/2$

We will NOT deal with the **Clebsch-Gordan coefficients**, but with the foundation given already, it should be easy for you to learn from the book.

Detour: Explanation of video about "spin"



$$\begin{pmatrix} 1 \\ 0 \end{pmatrix} = \frac{1}{\sqrt{2}} \cdot \frac{1}{\sqrt{2}} \begin{pmatrix} 1 \\ 1 \end{pmatrix} + \frac{1}{\sqrt{2}} \cdot \frac{1}{\sqrt{2}} \begin{pmatrix} 1 \\ -1 \end{pmatrix}$$

\uparrow \uparrow
 $+ \hbar/2$ $- \hbar/2$
 for S_x for S_x

$|\frac{1}{\sqrt{2}}|^2 = \frac{1}{2} = 50\%$ chance
 of measuring "up"
 when measuring S_x .

$$= \frac{1}{\sqrt{2}} \frac{1}{\sqrt{2}} \begin{pmatrix} 1 \\ i \end{pmatrix} + \frac{1}{\sqrt{2}} \frac{1}{\sqrt{2}} \begin{pmatrix} 1 \\ -i \end{pmatrix}$$

\uparrow \uparrow
 $+ \hbar/2$ $- \hbar/2$
 for S_y for S_y

50% chance of measuring
 "up" when measuring S_y .

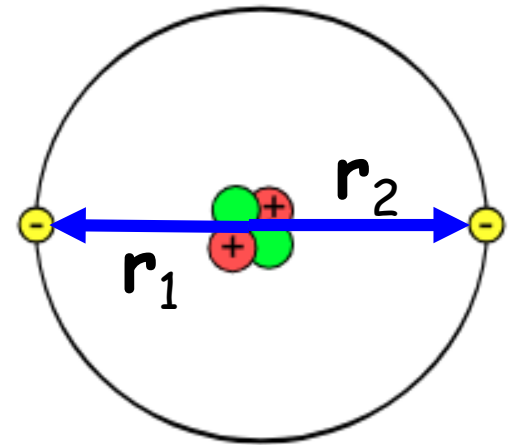
From
 HW
 4.32

Chapter 5: Identical Particles

For **one particle**, like one electron in the H atom, we simply need the wave function $\Psi(\mathbf{r}_1, t)$ where \mathbf{r}_1 is the coordinate of electron "1".

Consider now **two particles** as warm up.

For **two particles**, e.g. two electrons in the He atom, in QM we need the wave function $\Psi(\mathbf{r}_1, \mathbf{r}_2, t)$ where \mathbf{r}_1 and \mathbf{r}_2 are the coordinates of the two electrons.

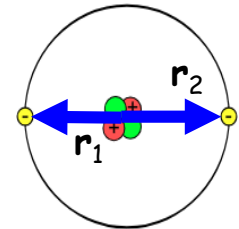


Mathematically, the Sch. Eq. $i\hbar \frac{\partial \Psi}{\partial t} = H\Psi$ has a more complicated Hamiltonian.

$$H = -\frac{\hbar^2}{2m_1} \nabla_1^2 - \frac{\hbar^2}{2m_2} \nabla_2^2 + V(\mathbf{r}_1, \mathbf{r}_2, t)$$

The potential V typically has terms like e-p attraction, but also e-e repulsion.

Example, for the He atom:



$$H = \left\{ -\frac{\hbar^2}{2m} \nabla_1^2 - \frac{1}{4\pi\epsilon_0} \frac{2e^2}{r_1} \right\} + \left\{ -\frac{\hbar^2}{2m} \nabla_2^2 - \frac{1}{4\pi\epsilon_0} \frac{2e^2}{r_2} \right\} + \frac{1}{4\pi\epsilon_0} \frac{e^2}{|\mathbf{r}_1 - \mathbf{r}_2|}$$

p-e₁ attraction is V(r₁)
p-e₂ attraction is V(r₂)
e₁-e₂ repulsion is V(r₁, r₂)