

# Magic Numbers:

The binding energies predicted by the Liquid Drop Model underestimate the actual binding energies of "magic nuclei" for which either the number of neutrons N = (A - Z) or the number of protons, Z is equal to one of the following "**magic numbers**" 2, 8, 20, 28, 50, 82, 126.

This is particularly the case for "doubly magic" nuclei in which both the number of neutrons and the number of protons are equal to magic numbers.

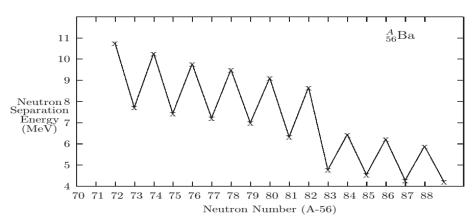
For example for 56 28Ni (nickel) the Liquid Drop Model predicts a binding energy of 477.7 MeV, whereas the measured value is 484.0 MeV. Likewise for 132 50 Sn (tin) the Liquid Drop model predicts a binding energy of 1084 MeV, whereas the measured value is 1110 MeV.

# **Nuclear Shell Model:**

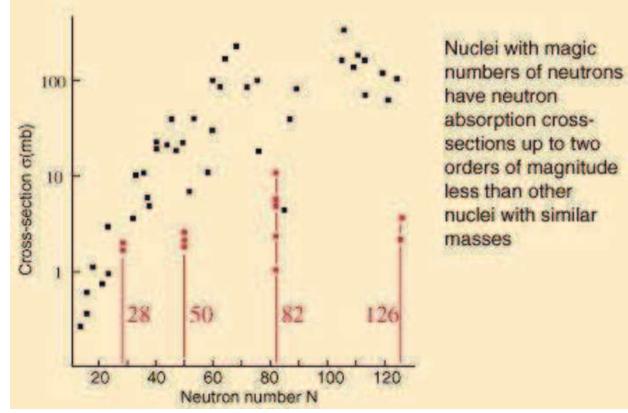
It has been found that the nuclei with proton number or neutron number equal to certain numbers 2,8,20,28,50,82 and 126 behave in a different manner when compared to other nuclei having neighboring values of Z or N. Hence these numbers are known as magic numbers. This is found to be in accordance with the observed nature of elements with filled shells. Thus Physicists looked at such a possibility in case of filling of nucleons in the nucleus. Thus a new model of nucleus has emerged. This model is known as the Shell model.

### **Experimental evidences for the existence of magic numbers;**

- 1. The binding energy of magic numbered nuclei is much larger than the neighboring nuclei. Thus larger energy is required to separate a single nucleon from such nuclei.
- Number of stable nuclei with a given value of Z and N corresponding to the magic number are much larger than the number of stable nuclei with neighboring values of Z and N. For example, Sn with Z=50 has 10 stable isotopes, Ca with Z=20 has six stable isotopes.
- 3. The neutron (proton) separation energies (the energy required to remove the last neutron (proton)) peaks if N (Z) is equal to a magic number.



4. If N is magic number then the cross-section for neutron absorption is much lower than for other nuclides.



- 5. Three naturally occurring radioactive series decay to the stable end product Pb with Z=82 in three isotopic forms having N=126 for one of them.
- 6. Nuclei with the value of N just one more than the magic number spontaneously emit a neutron (when excited by preceding beta-decay) E.g., O-17, K-87 and Xe-137.
- 7. Electric quadrupole moment of magic numbered nuclei is zero indicating the spherical symmetry of nucleus for closed shells.
- 8. Nuclei with magic numbers of neutrons or protons have their first excited states at higher energies than in cases of the neighboring nuclei.

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# Independent particle model

#### Harmonic oscillator potential:

 $\hbar^2$ 

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Consider a nucleon moving independently in the harmonic oscillator potential which is spherically symmetric. The Schrodinger equation given below can be solved in the Cartesian coordinate system as well as in the spherical coordinate system.

with

$$V(\mathbf{r}) = rac{1}{2}m\omega^2 r^2 = rac{1}{2}m\omega^2(x^2+y^2+z^2).$$

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In the Cartesian coordinate system, it can be separated into three linear harmonic oscillators and consequently, the energy of the three-dimensional harmonic oscillator is the sum of the energies of the three linear harmonic oscillators.

$$E = \hbar\omega\left(n_x + n_y + n_z + \frac{3}{2}\right) = \hbar\omega\left(N + \frac{3}{2}\right),$$
.....(2)

where N is the principal quantum number which will assume positive integral values including zero. From Eq. (2), we observe that, the energy depends only on the quantum number N and not on  $n_x$ ,  $n_y$  and  $n_z$ . So, there is a degeneracy of energy levels. Since the proton (or neutron) has spin

 $1/2\hbar$ , there are two possible spin orientations for each state specified by a set of quantum numbers  $n_x$ ,  $n_y$  and  $n_z$ . Applying the Pauli exclusion principle, we obtain the number of protons or neutrons that can have a particular energy as shown in Table I.

Table I: Number ( $N_N$ ) of protons or neutrons in each state with the principal quantum Number N. The multiplicative factor 2 in the last column is due to two spin states.

N	$n_x$	$n_y$	$n_z$	$\mathcal{N}_N$
0	0	0	0	1  imes 2 = 2
1	1	0	0	
	0	1	0	
	0	0	1	$3 \times 2 = 6$
2	1	1	0	
	1	0	1	
	0	1	1	
	2	0	0	
	0	2	0	
-	0	0	2	$6 \times 2 = 12$

In general

$$\mathcal{N}_N = (N+1)(N+2).$$
 ----(3)

However, it is found not convenient to work in Cartesian coordinate system. Since the potential is spherically symmetric, one can attempt to solve the Schrodinger equation in spherical coordinates.

Let us write  $\nabla^2$  in spherical coordinates.

$$\nabla^2 = \frac{1}{r^2} \frac{\partial}{\partial r} \left( r^2 \frac{\partial}{\partial r} \right) + \frac{1}{r^2 \sin \theta} \frac{\partial}{\partial \theta} \left( \sin \theta \frac{\partial}{\partial \theta} \right) + \frac{1}{r^2 \sin^2 \theta} \frac{\partial^2}{\partial \phi^2}$$
  
$$= \frac{1}{r^2} \frac{\partial}{\partial r} \left( r^2 \frac{\partial}{\partial r} \right) - \frac{L^2}{r^2 \hbar^2},$$
 (4)

where  $L^2$  is the square of the angular momentum operator

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$$\boldsymbol{L}^{2} = -\hbar^{2} \left[ \frac{1}{\sin\theta} \frac{\partial}{\partial\theta} \left( \sin\theta \frac{\partial}{\partial\theta} \right) + \frac{1}{\sin^{2}\theta} \frac{\partial^{2}}{\partial\phi^{2}} \right],$$
(5)

which has the spherical harmonics  $Y_{Im}(\theta, \phi)$  as eigen functions with eigen values  $l(l+1)\hbar^2$ .

$$L^{2}Y_{lm}(\theta,\phi) = l(l+1)\hbar^{2}Y_{lm}(\theta,\phi).$$
(6)

Substituting (4) in (1) we get,

$$\left\{\frac{1}{r^2}\frac{\partial}{\partial r}\left(r^2\frac{\partial}{\partial r}\right) - \frac{L^2}{r^2\hbar^2} + \frac{2m}{\hbar^2}\left(E - \frac{1}{2}m\omega^2r^2\right)\right\}\psi(\mathbf{r}) = 0.$$
(7)

Writing the solution of  $\psi(r)$  of eq. (7) as a product of radial and angular functions

$$\psi(\mathbf{r}) = R(\mathbf{r}) Y_{lm}(\theta, \phi), \tag{8}$$

And using the eigen value equation (6), we obtain the radial equation

$$\left\{\frac{1}{r^2}\frac{\partial}{\partial r}\left(r^2\frac{\partial}{\partial r}\right) - \frac{l(l+1)}{r^2} + \frac{2m}{\hbar^2}\left(E - \frac{1}{2}m\omega^2r^2\right)\right\}R(r) = 0.$$
(9)

Applying the bound state boundary condition that the radial function should vanish at infinity, we obtain the discrete energy levels of the three dimensional harmonic oscillator.

$$E = \left(2n+l+\frac{3}{2}\right)\hbar\omega, \qquad n = 0, 1, 2, \cdots.$$
(10)

Comparing (10) with the energy levels (2) obtained in the Cartesian coordinate system,

$$E_N = \left(N + \frac{3}{2}\right)\hbar\omega = \left(2n + l + \frac{3}{2}\right)\hbar\omega,$$
------(11)

One can find

$$N=2n+l, \qquad n=0,1,2,\cdots.$$

The eigenvalues and eigenfunctions in spherical coordinate system depend on two quantum numbers: the radial quantum number n and the orbital quantum number l. It follows that for a given total quantum number N, the orbital quantum number l can take only even values if N is even or only odd values if N is odd.

-(12)

$$l = N - 2n$$
  
= N, N - 2, N - 4, ..., 0 or 1.

The additional quantum number m occurring in Eq. (6.8) is the magnetic quantum number. It may be remarked that the accidental degeneracy in the states with a given *I* but with different m having the same energy, occurs in any spherical potential. However, in the case of the oscillator potential, there is a special degeneracy as well, i.e., states with different *I* values but with the same *N* have the same energy. Accordingly, the number of protons or neutrons with a given value of *N* is

$$\mathcal{N}_N = \sum_l 2(2l+1),$$

where the factor 2 is due to the two spin states. Equation (6.12) gives the different I values allowed for a given N. The permitted values of I are all even or odd and consequently, the parity (given by (-1)') of all the nucleons with a given quantum number N is the same.

The three dimensional harmonic oscillator energy levels are labelled by a pair of quantum numbers (n, l). Also the spectroscopic notation s, p, d, f, g, h, ..., is used to denote states with l = 0,1,2,3,4,5,...'.

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(n, l) : Os; Op; (Od, Is); (Of, Ip); (Og, Id, 2s);

(*Oh*, *If*, 2*p*); (Oi, *Ig*, 2*d*, 3*s*); (*OJ*, *Ih*, 2*f*, 3*p*). -----(14) The bracketed levels on the right hand side are degenerate. We also give below the normalized radial functions for a few (*n*, *l*) values.

$$R_{nl}(r) = N_{nl} \, \alpha^{3/2} e^{-\alpha^2 r^2/2} (\alpha r)^l f(r^2),$$

where

 $\alpha = \sqrt{\frac{m\omega}{\hbar}}$ 

and

$$f(r^2) = 1, \qquad \text{for } n = 0;$$
  
=  $\frac{2l+3}{2} - \alpha^2 r^2, \qquad \text{for } n = 1;$   
=  $\frac{1}{\sqrt{2}} \left\{ \frac{(2l+3)(2l+5)}{4} - (2l+5)(\alpha r)^2 + (\alpha r)^4 \right\}, \quad \text{for } n = 2.$ 

The normalization factor  $N_{nl}$  is given by

$$N_{nl} = \left\{ \frac{2^{n+l+2}}{\sqrt{\pi}(2n+2l+1)!!} \right\}^{1/2},$$

such that

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$$\int_0^\infty |R_{nl}(r)|^2 r^2 dr = 1.$$

In literature on nuclear shell model<sup>2</sup>, a slightly different radial quantum number  $n_r$  (to distinguish it from the radial quantum number n, we use the notation  $n_r$ ) is also used such that

 $N = 2n_r + l - 2$  and  $n_r = n + 1$ ,  $n_r = 1, 2, 3, \cdots$ .

In general usage, the oscillator energy levels are labelled by a pair of quantum numbers (n, l) as described in (6.14) or  $(n_r, l)$  as shown below.

$$\begin{array}{rl} (n_r,l) &:& 1s; 1p; (1d,2s); (1f,2p); (1g,2d,3s); \\ && (ih,2f,3p); (1i,2g,3d,4s); (1j,2h,3f,4p). \end{array}$$

The radial quantum number nr denotes that it is the nth time that the l value occurs in the scheme. Alternatively, nr can be interpreted as the number of nodes that occurs in the radial wave function including the one at infinity. The bracketed levels on the right hand side of (6.20) are degenerate. We shall use the notation (n, I) or (nr, I) as the situation demands.

In Table II, we present the number of protons or neutrons that can occupy each oscillator state with total quantum number N.

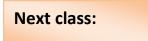
Table II: Number of protons or neutrons occupying each state with total quantum number N and the orbital substates with quantum numbers  $n_r$ , l. The energy E is in units of  $\hbar\omega$ .

N	E	Orbitals $(n_{\tau}, l)$	$ \mathcal{N}_N = (N+1)(N+2) \\ = \sum_l 2(2l+1) $	$\sum_N N_N$
0	3/2	1s	2	2
1	5/2	1p	6	8
2	7/2	1d, 2s	12	20
3	9/2	1f, 2p	20	40
4	11/2	1g, 2d, 3s	30	70
5	13/2	1h, 2f, 3p	42	112
6	15/2	1i, 2g, 3d, 4s	56	168

The numbers in the last column should correspond to the closed shell nuclei and hence to the magic numbers but the discrepancy arises after the first three shells. Further, there is a degeneracy of states with different I values. Such a degeneracy does not exist in the case of square well potential. Empirically, let us assume a level split of different I orbitals with the energy given by

$$E_{N,l} = \hbar\omega\left(N + \frac{3}{2}\right) + Dl(l+1),$$

assuming an additional force which lowers the states of larger *I*. A value  $D = -0.0225/\hbar\omega$  is found to be satisfactory.



• Spin – orbit potential

### **References :**

- 1. Nuclear physics by D. C.TAYAL
- 2. Lecture points of Dr.H R Sreepad