

# THE SHELL MODEL

22.02

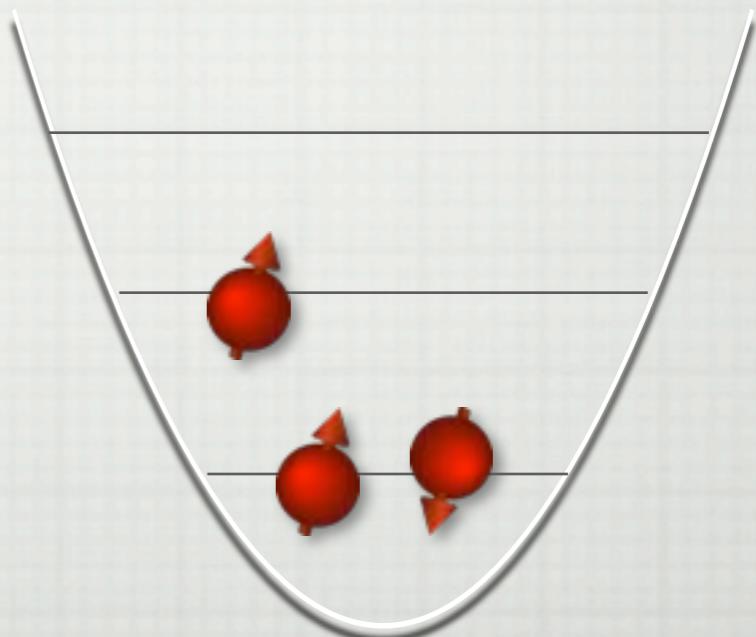
Introduction To Applied Nuclear Physics

Spring 2012

# Atomic Shell Model

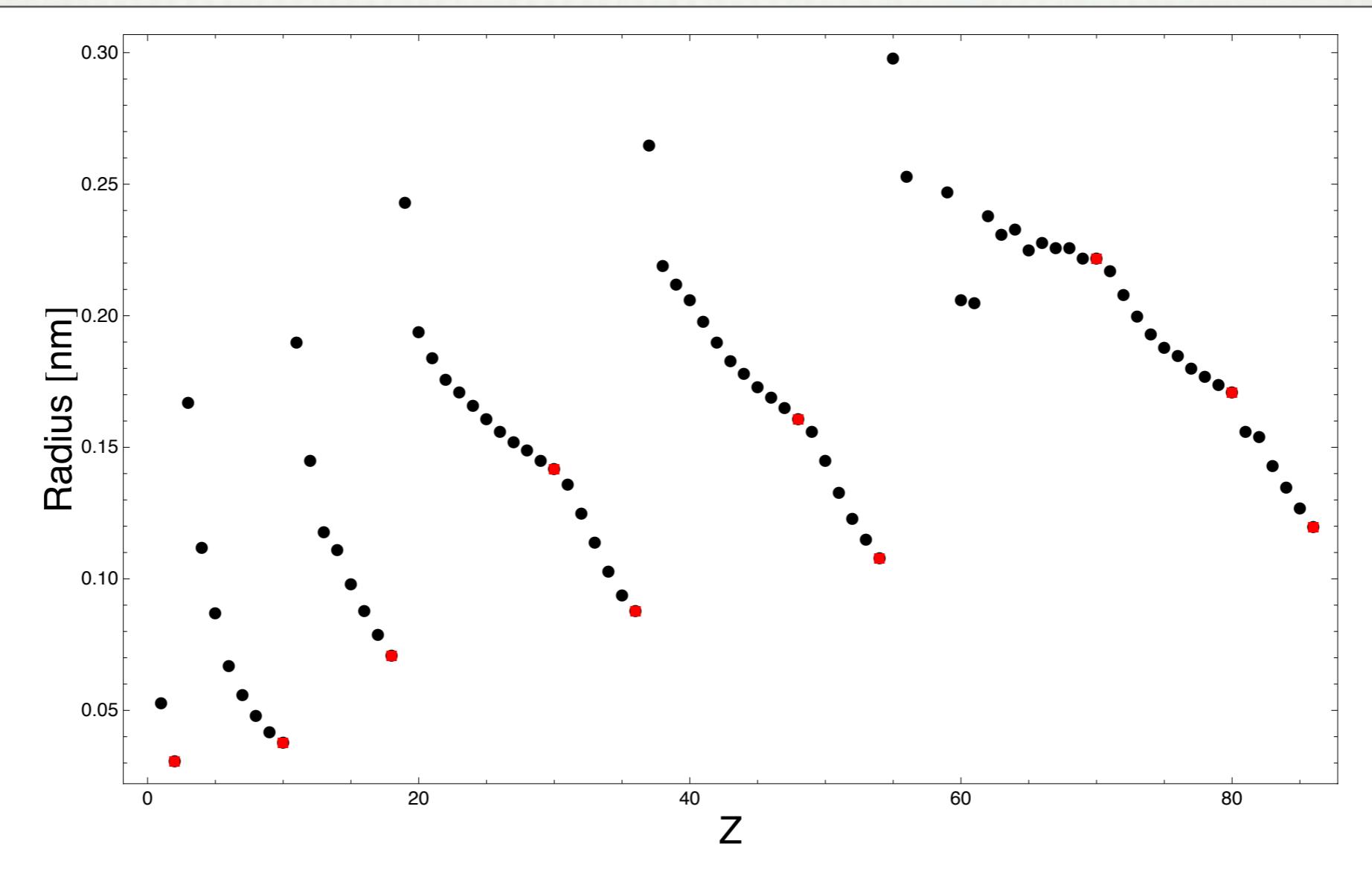
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- Chemical properties show a periodicity
- Periodic table of the elements
- Add electrons into shell structure

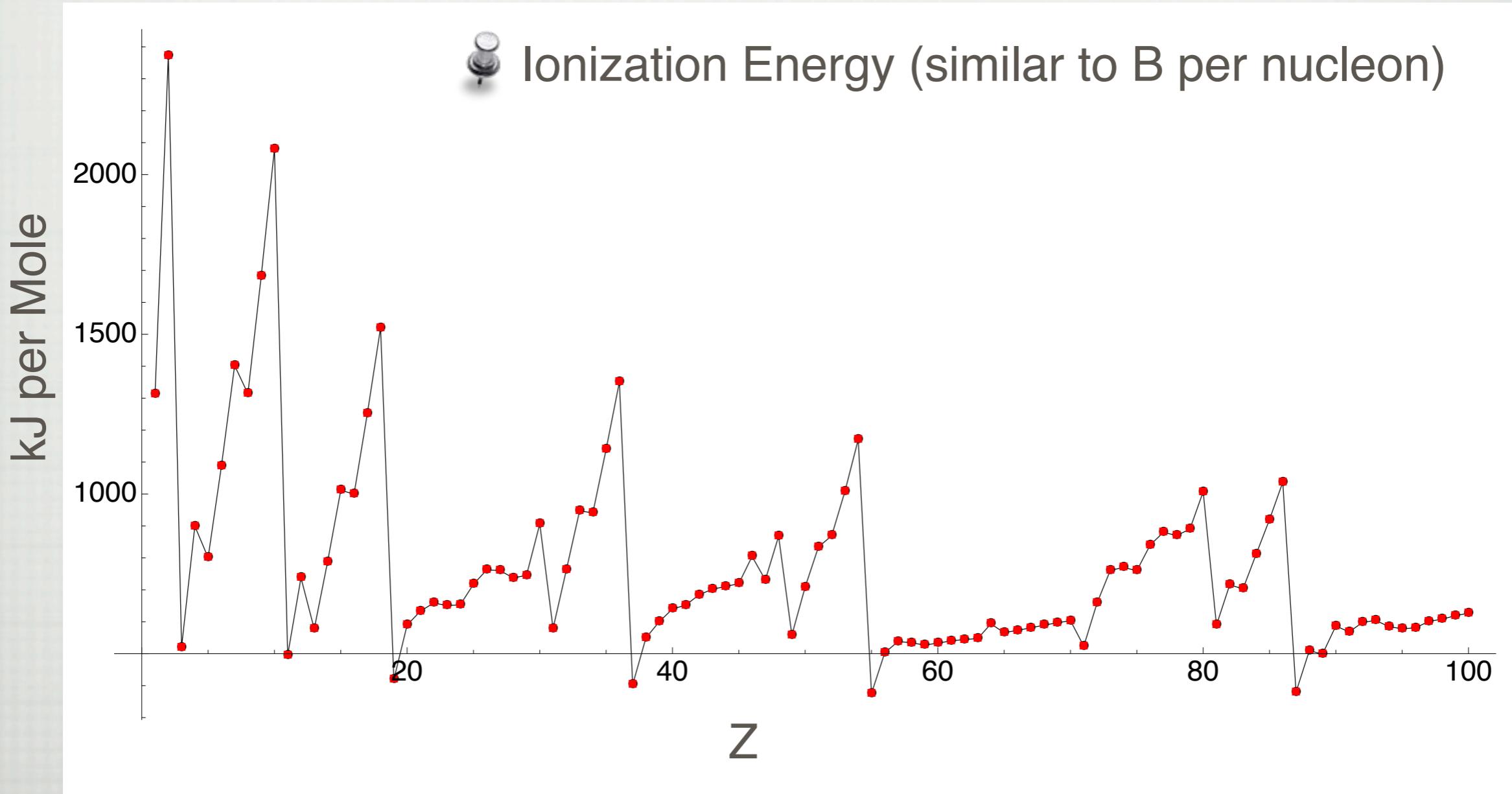


# Atomic Radius

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# Ionization Energy



# ATOMIC STRUCTURE

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- The atomic wavefunction is written as

$$|\psi\rangle = |n, l, m\rangle = R_{n,l}(r)Y_l^m(\vartheta, \varphi)$$

- where the labels indicate :

- n : principal quantum number
- l : orbital (or azimuthal) quantum number
- m: magnetic quantum number

- The degeneracy is

$$\mathcal{D}(l) = 2(2l + 1) \rightarrow \mathcal{D}(n) = 2n^2$$

# AUFBAU PRINCIPLE

- The orbitals (or shells) are then given by the n-levels (?)

1	0	1	2	3	4	5	6
Spectroscopic notation	s	p	d	f	g	h	i
$\mathcal{D}(l)$	2	6	10	14	18	22	26

historic structure      heavy nuclei

n	$\mathcal{D}(n)$	$e^-$ in shell
1	2	2
2	6	8
3	18	28

# ATOMIC PERIODIC TABLE

# AUFBAU PRINCIPLE

- The orbitals (or shells) are then given by close-by energy-levels

1	0	1	2	3	4	5	6
Spectroscopic notation	s	p	d	f	g	h	i
$\mathcal{D}(l)$	2	6	10	14	18	22	26

historic structure      heavy nuclei

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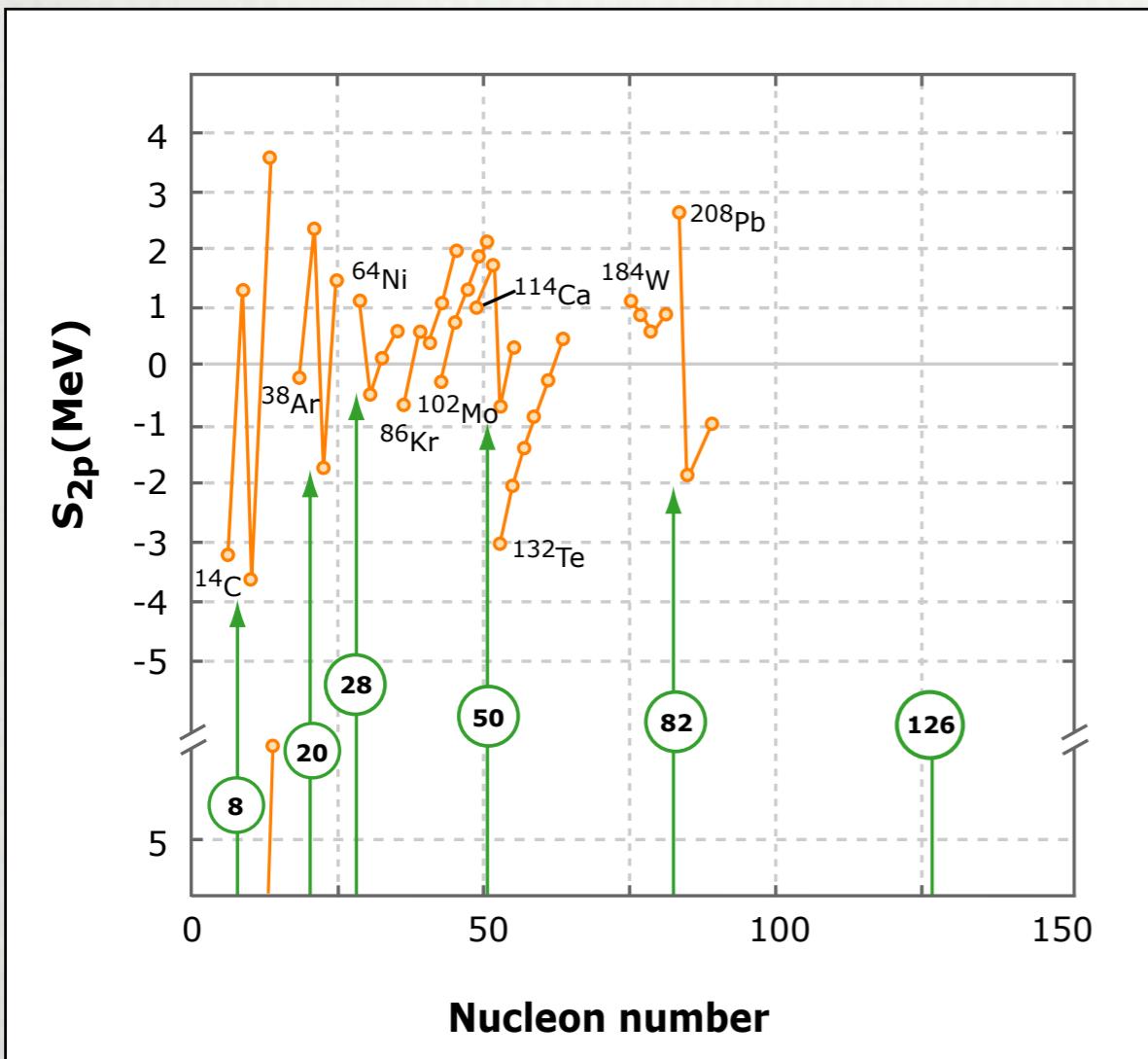
3s+3p form one level with # 10  
4s is filled before 3d

# Nuclear Shell Model

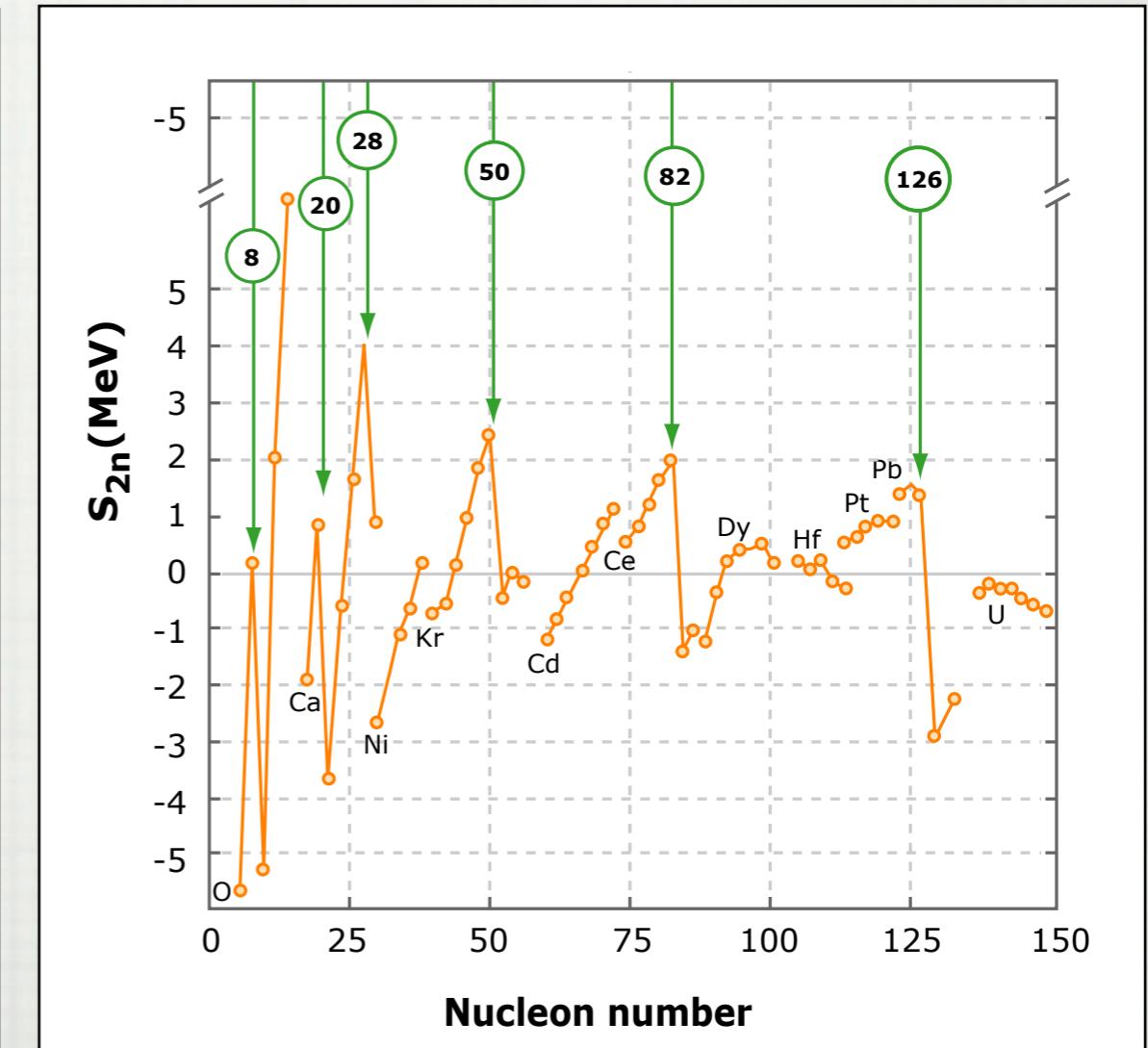
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- Picture of adding particles to an external potential is no longer good: each nucleon contributes to the potential
- Still many evidences of a shell structure

# Separation Energy



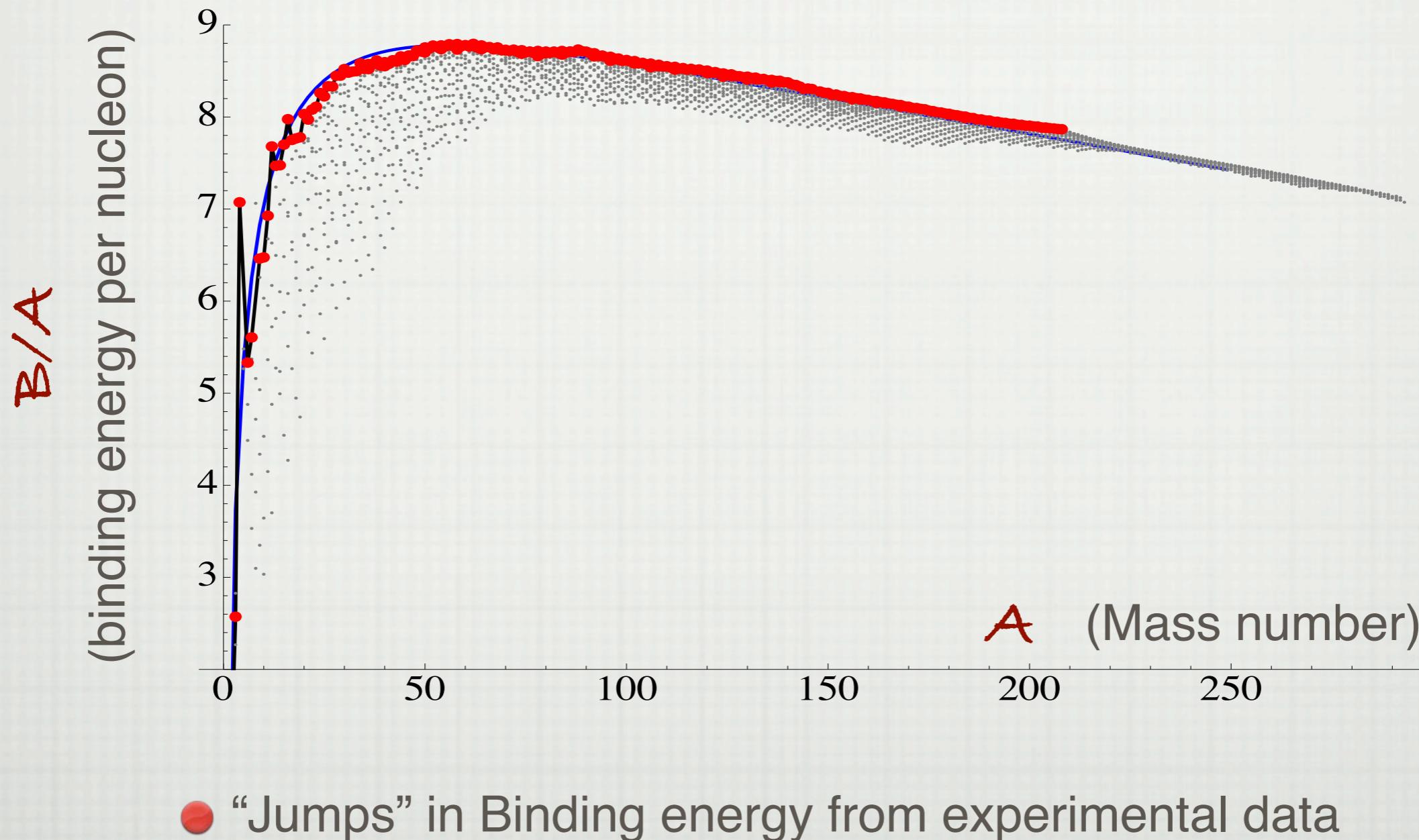
PROTON



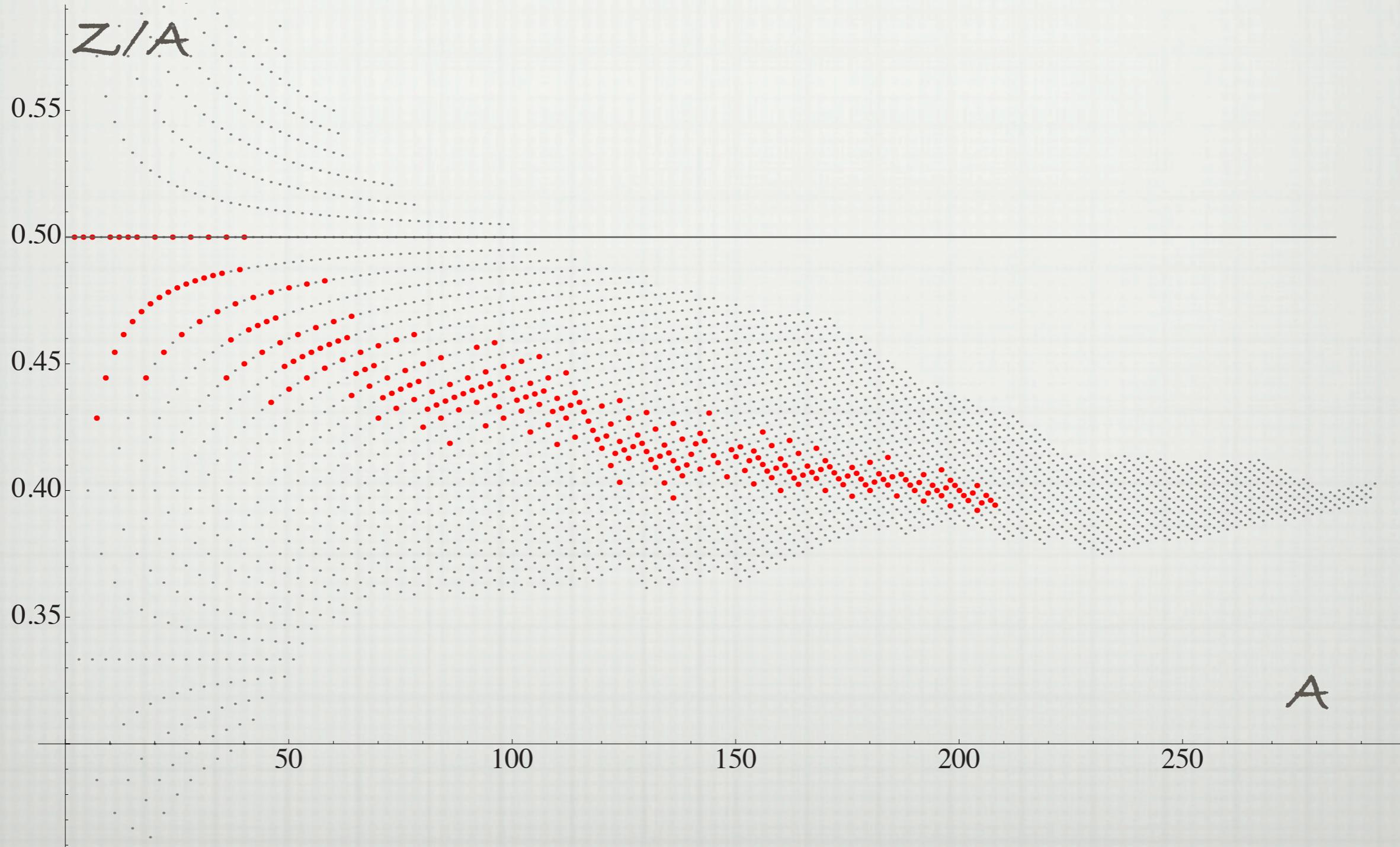
NEUTRON

Image by MIT OpenCourseWare. After Krane.

# B/A: JUMPS



# CHART of NUCLIDES (Z/A vs. A)



# CHART OF NUCLIDES



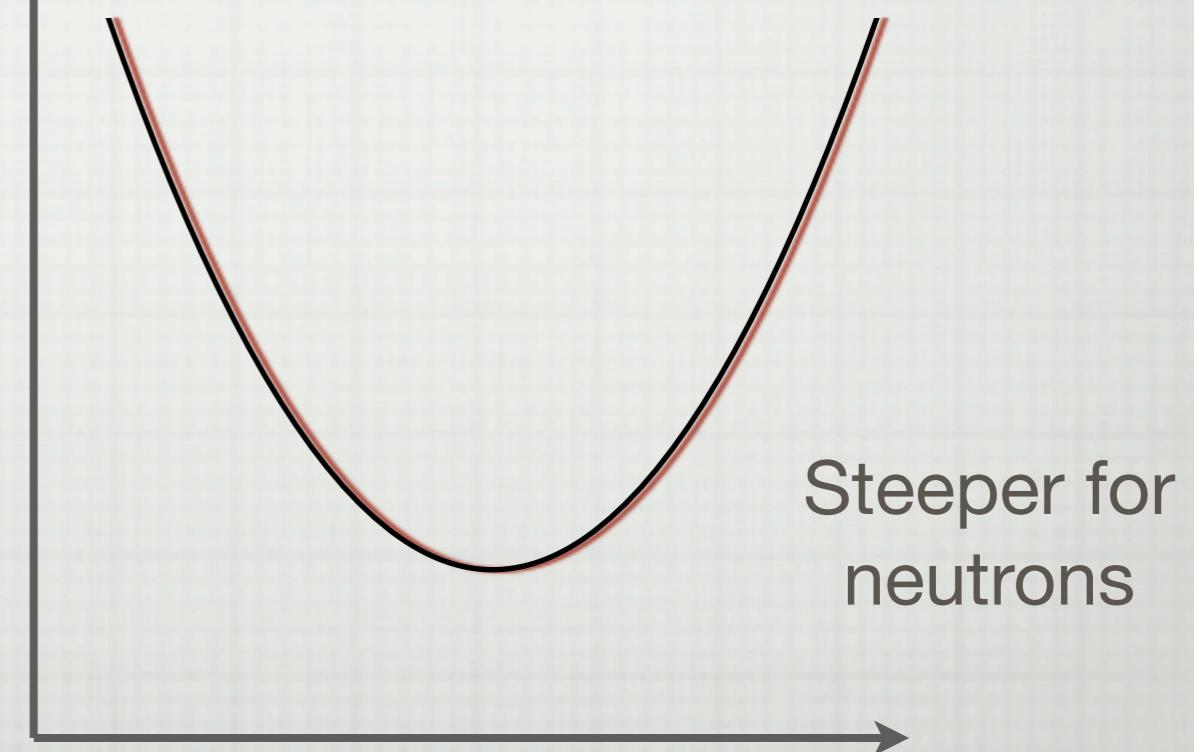
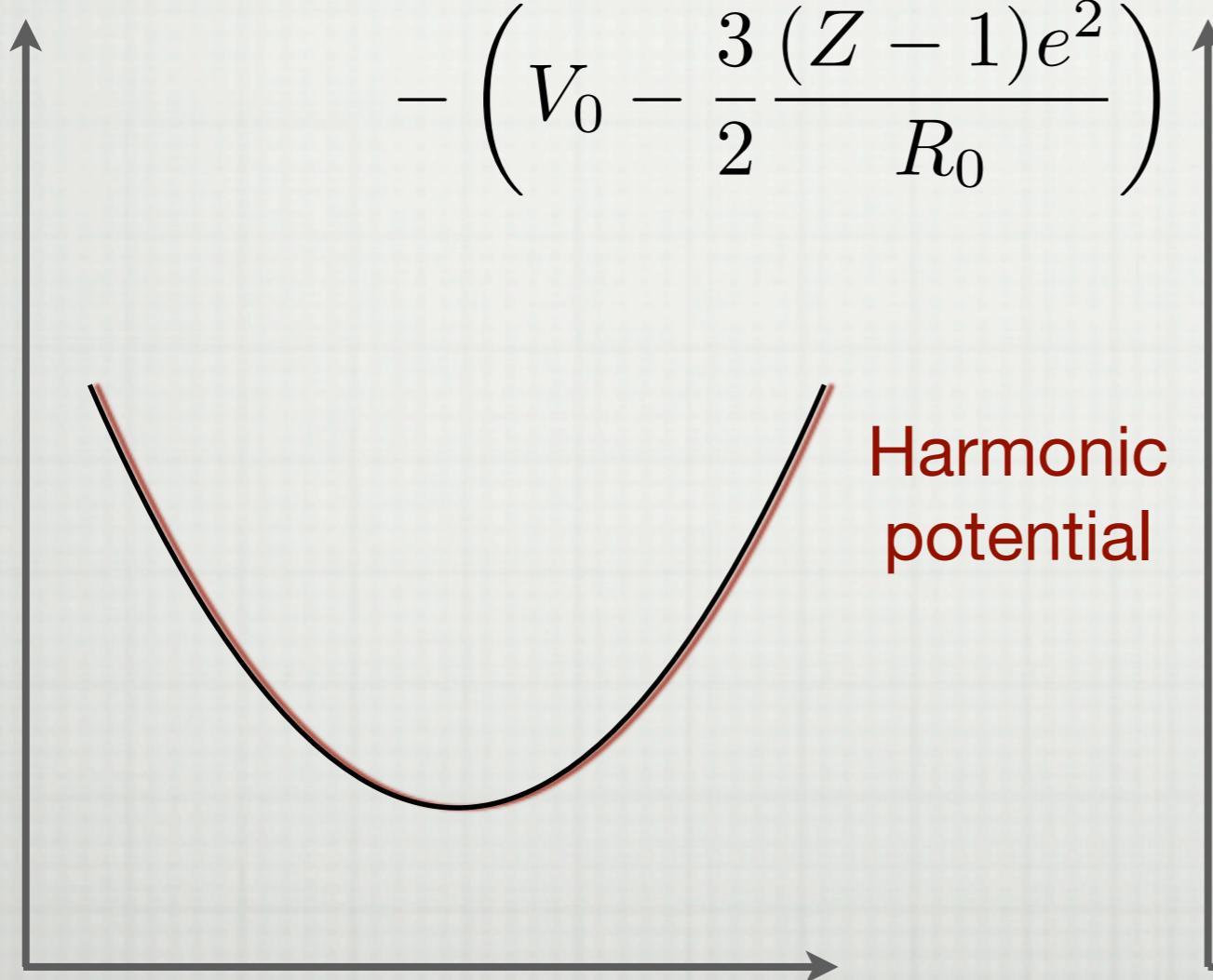
*“Periodic”, more complex properties → nuclear structure*

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# NUCLEAR POTENTIAL

$$V_p = r^2 \left( \frac{V_0}{R_0^2} - \frac{(Z-1)e^2}{2R_0^3} \right)$$

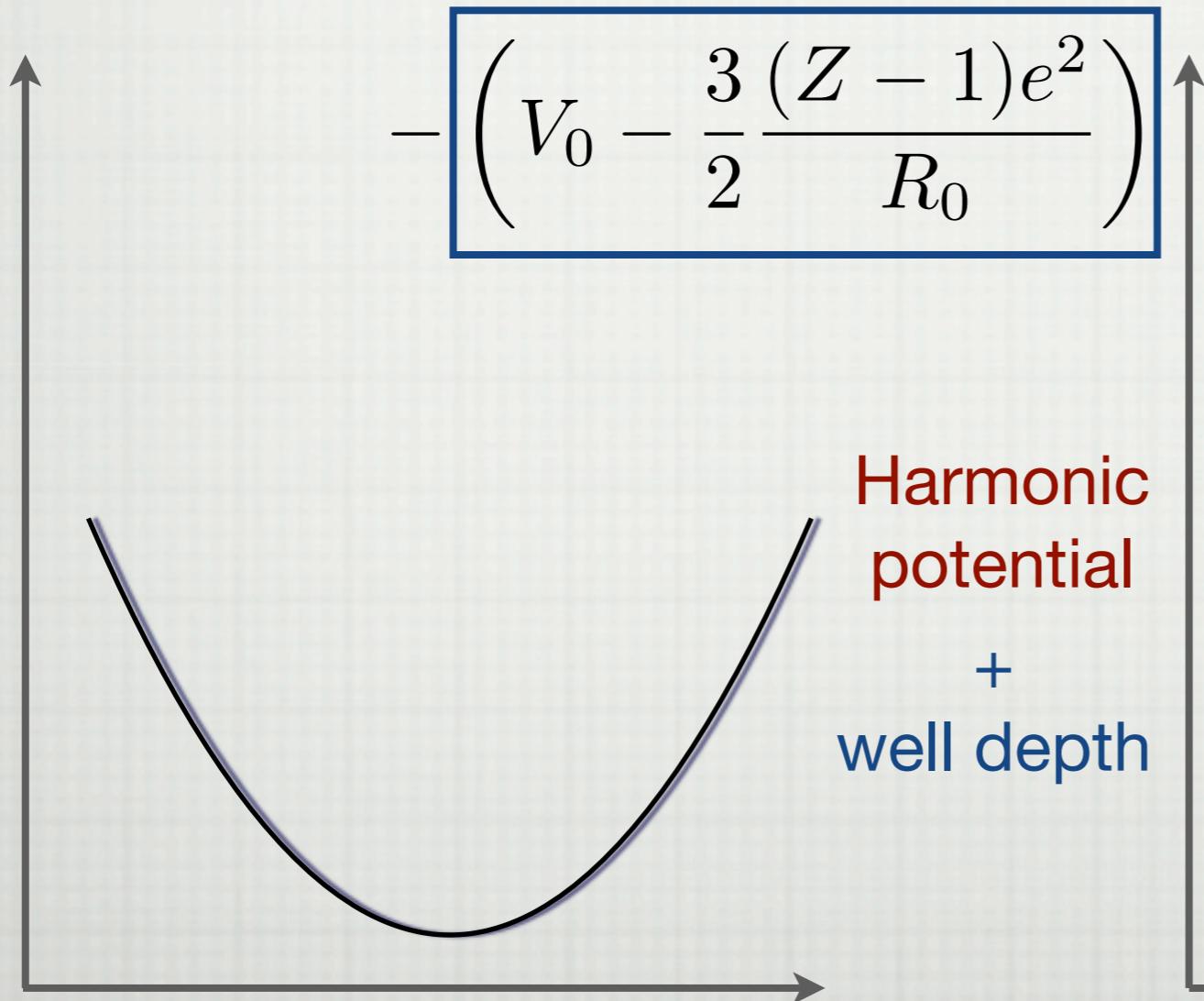
$$V_n = r^2 \left( \frac{V_0}{R_0^2} \right) - (V_0)$$



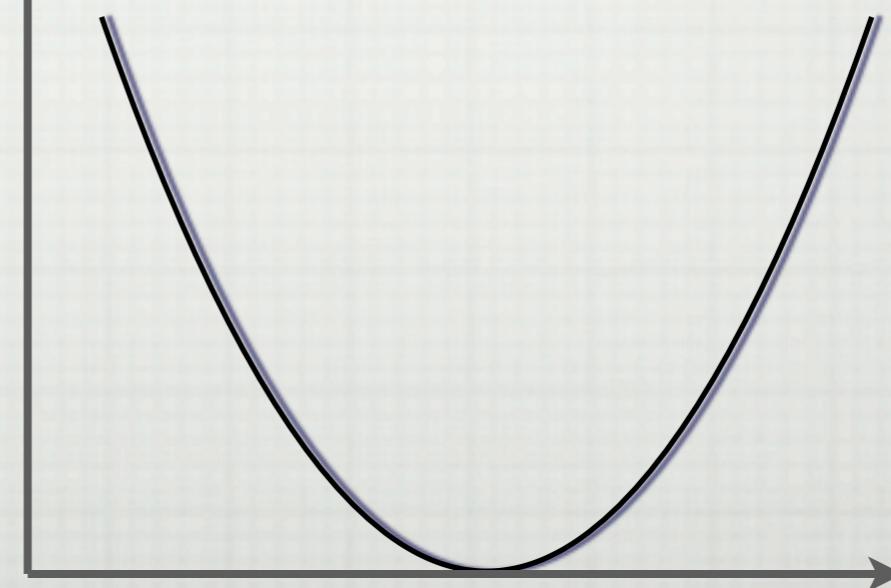
# NUCLEAR POTENTIAL

$$V_p = r^2 \left( \frac{V_0}{R_0^2} - \frac{(Z-1)e^2}{2R_0^3} \right)$$

$$V_n = r^2 \left( \frac{V_0}{R_0^2} \right) - (V_0)$$



Steeper and Deeper  
for neutrons



# Shell Mode

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- 📌 Harmonic oscillator: solve (part of) the radial equation
  - including the angular momentum (centrifugal force term) we obtain the usual principal quantum number  $n = (N-l)/2+1$

N	$l$	Spectroscopic Notation	$\mathcal{D}(N)$	$2\mathcal{D}(N)$	Cumulative #
0	0	1 s	1	2	2
1	1	1 p	3	6	8
2	0,2	2s,1d	6	12	20
3	1,3	2p,1f	10	20	40
4	0,2,4	3s,2d,1g	15	30	70

# Spin-Orbit Coupling

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- The spin-orbit interaction is given by  $V_{SO} = \frac{1}{\hbar^2} V_{so}(r) \hat{\vec{l}} \cdot \hat{\vec{s}}$
- We can calculate the dot product

$$\langle \hat{\vec{l}} \cdot \hat{\vec{s}} \rangle = \frac{1}{2} (\hat{j}^2 - \hat{l}^2 - \hat{s}^2) = \frac{\hbar^2}{2} [j(j+1) - l(l+1) - \frac{3}{4}]$$

- Because of the addition rules,  $j = l \pm \frac{1}{2}$

$$\langle \hat{\vec{l}} \cdot \hat{\vec{s}} \rangle = \begin{cases} l \frac{\hbar^2}{2} & \text{for } j=l+\frac{1}{2} \\ -(l+1) \frac{\hbar^2}{2} & \text{for } j=l-\frac{1}{2} \end{cases}$$

# Spin-Orbit Coupling

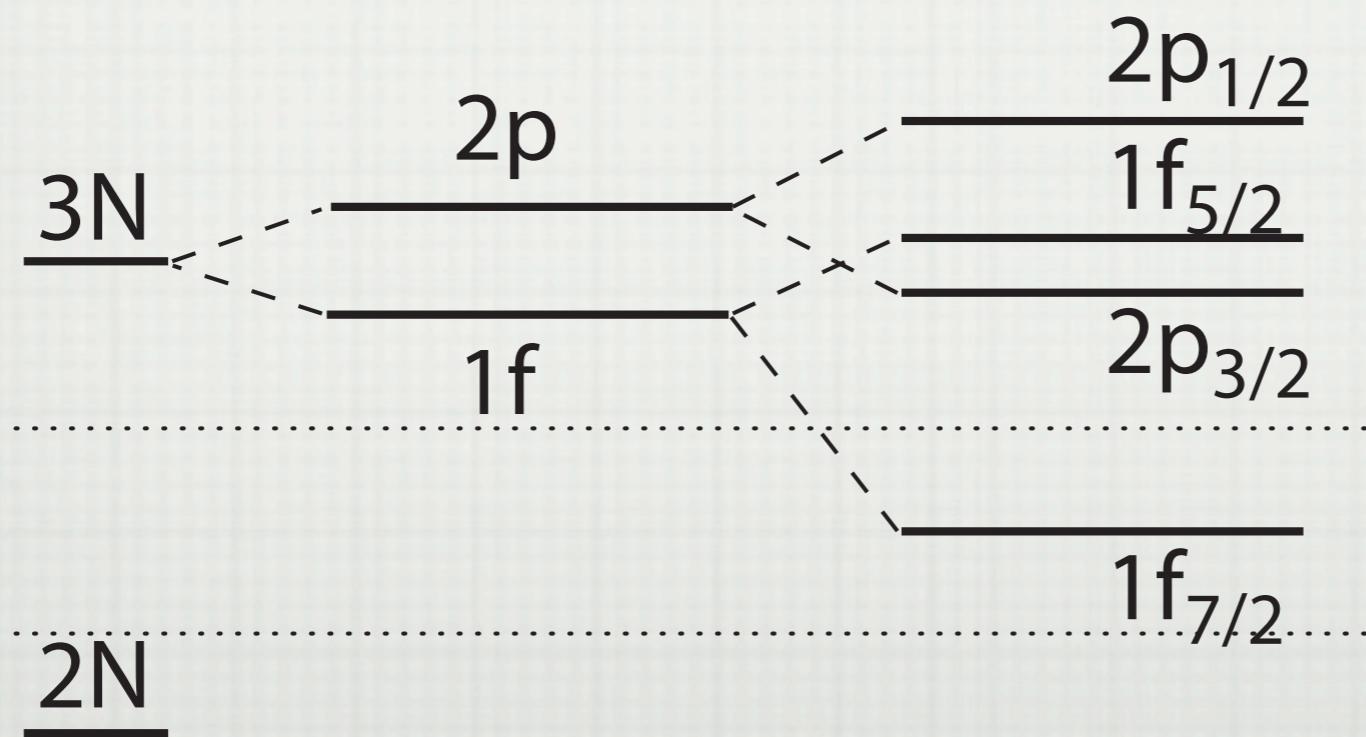
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- when the spin is **aligned** with the angular momentum  $j = l + \frac{1}{2}$  the potential becomes more negative,  
i.e. the well is deeper and the state more tightly bound.
- when spin and angular momentum are **anti-aligned**  $j = l - \frac{1}{2}$  the system's energy is higher.
- The difference in energy is  $\Delta E = \frac{V_{so}}{2}(2l + 1)$   
Thus it increases with  $l$ .

# Example

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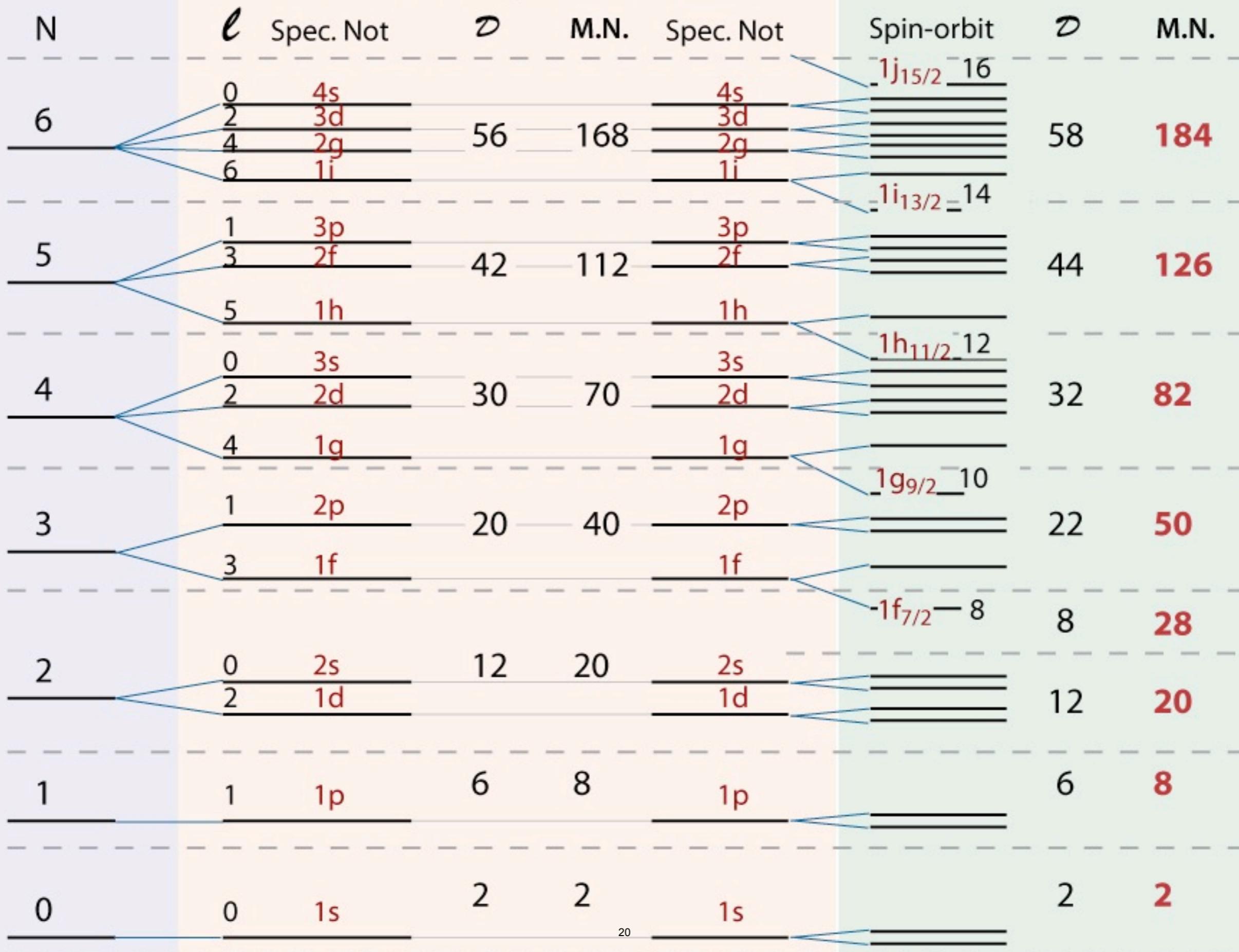
- $3N$  level, with  $l=3$  ( $1f$  level)  $j=7/2$  or  $j=5/2$ 
  - Level is pushed so down that it forms its own shell

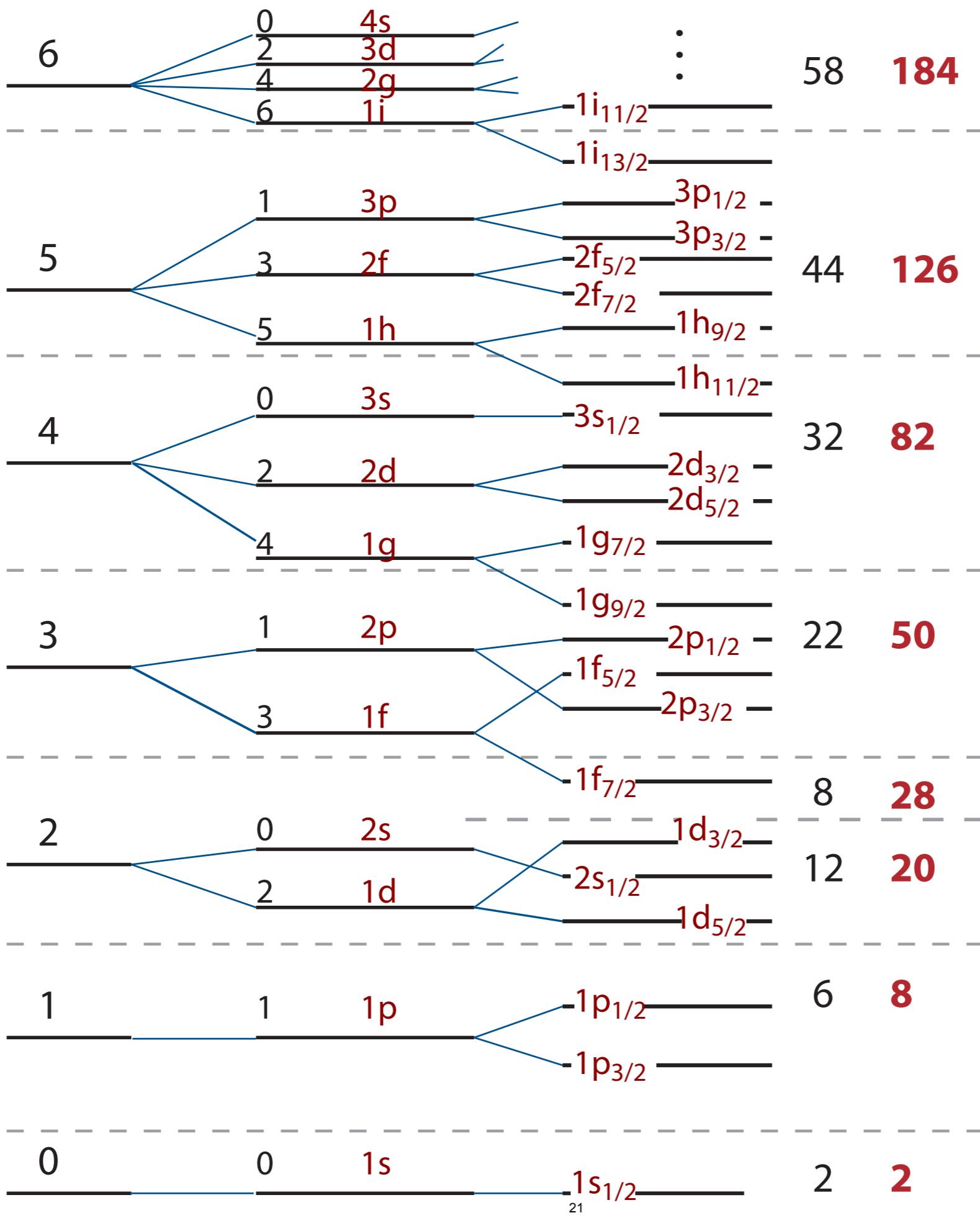


Harmonic  
Oscillator

Harmonic oscillator  
with angular momentum

Spin-Orbit  
Potential





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