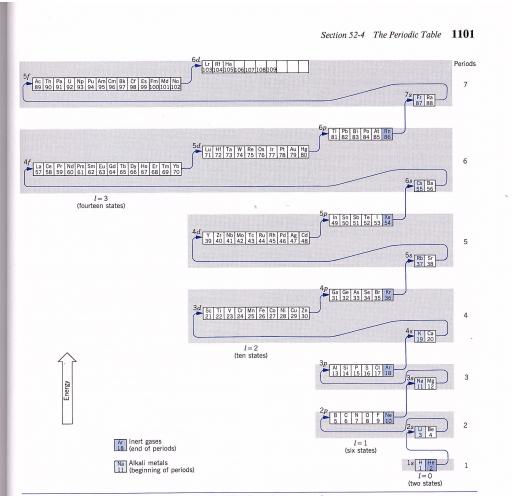
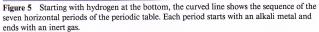
Group — ↓ Period	• 1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18
1	1 H																	2 He
2	3 Li	4 Be											5 B	6 C	7 N	8 O	9 F	10 Ne
3	11 Na	12 Mg											13 Al	14 Si	15 P	16 S	17 Cl	18 Ar
4	19 K	20 Ca	21 Sc	22 Ti	23 V	24 Cr	25 Mn	26 Fe	27 Co	28 Ni	29 Cu	30 Zn	31 Ga	32 Ge	33 As	34 Se	35 Br	36 Kr
5	37 Rb	38 Sr	39 Y	40 Zr	41 Nb	42 Mo	43 Tc	44 Ru	45 Rh	46 Pd	47 Ag	48 Cd	49 In	50 Sn	51 Sb	52 Te	53 I	54 Xe
6	55 Cs	56 Ba		72 Hf	73 Ta	74 W	75 Re	76 Os	77 Ir	78 Pt	79 Au	80 Hg	81 Tl	82 Pb	83 Bi	84 Po	85 At	86 Rn
7	87 Fr	88 Ra		104 Rf	105 Db	106 Sg	107 Bh	108 Hs	109 Mt	110 Ds	111 Rg	112 Cn	113 Uut	114 Uuq	115 Uup	116 Uuh	117 Uus	118 Uuo
Lanthanides		57 La	58 Ce	59 Pr	60 Nd	61 Pm	62 Sm	63 Eu	64 Gd	65 Tb	66 Dy	67 Ho	68 Er	69 Tm	70 Yb	71 Lu		
Actinides				89 Ac	90 Th	91 Pa	92 U	93 Np	94 Pu	95 Am	96 Cm	97 Bk	98 Cf	99 Es	100 Fm	101 Md	102 No	103 Lr





in this figure. States with the same value of *l* have been displaced to the left for clarity and grouped into columns according to their *l* value.

Before we look more closely at this table, we introduce a new notation for the angular momentum quantum number *l*. For historical reasons* the values of *l* have been

* The letters *s*, *p*, *d*, *f* stand for *sharp*, *principal*, *diffuse*, and *fundamental*, which were early spectroscopic designations of spectral lines. Beyond *f*, the states are labeled in alphabetic order.

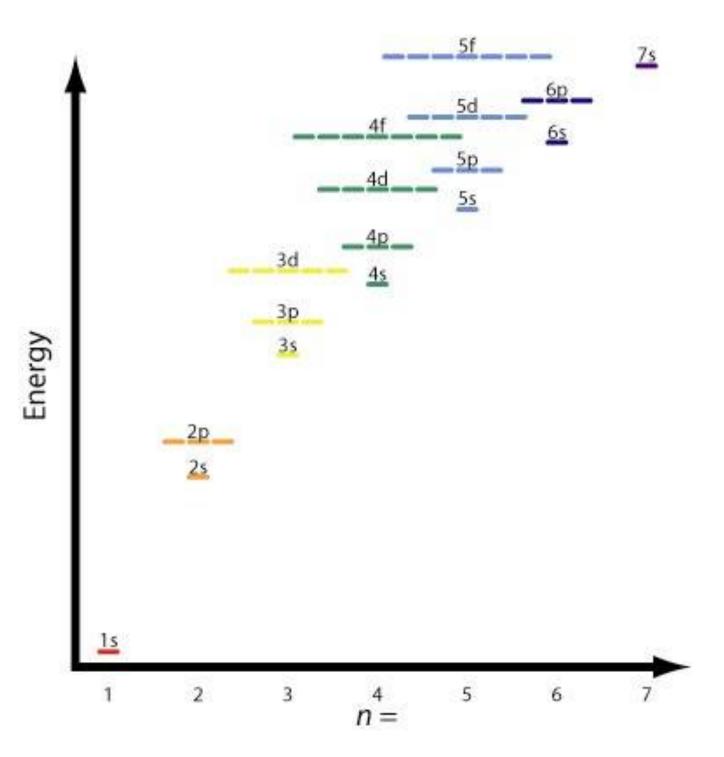
given letter equivalents, according to this scheme:

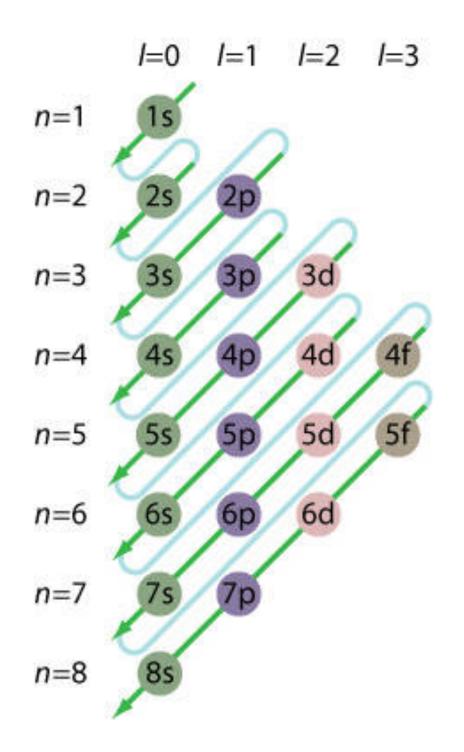
$$l \qquad 0 \quad 1 \quad 2 \quad 3 \quad 4 \quad 5 \dots$$

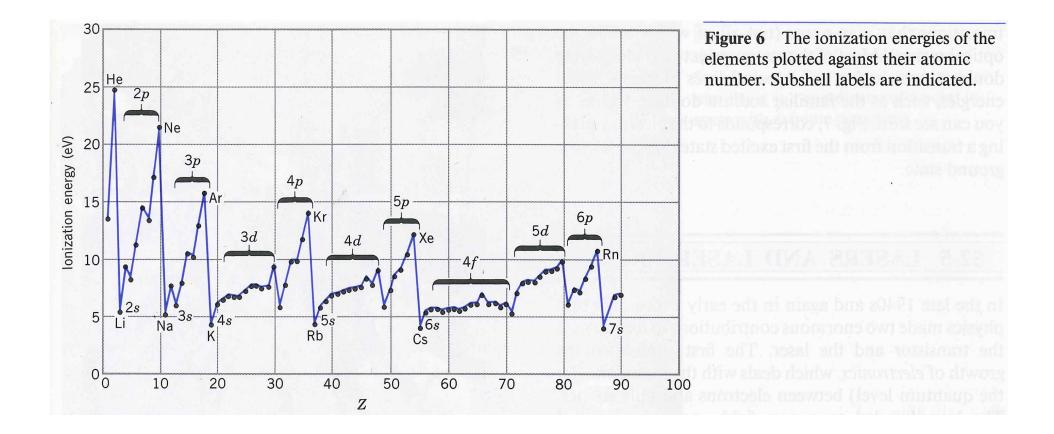
Symbol
$$s \ p \ d \ f \ g \ h \dots$$

In this notation, a state with n = 1 and l = 0 is called a "1s" state. Similarly, a state with n = 4 and l = 3 is called a "4f" state, and so on. These states are also known as *subshells*.

The dependence of energy on l is a dominant feature of Fig. 5. Look, for example, at the sequence of states 4s, 4p,

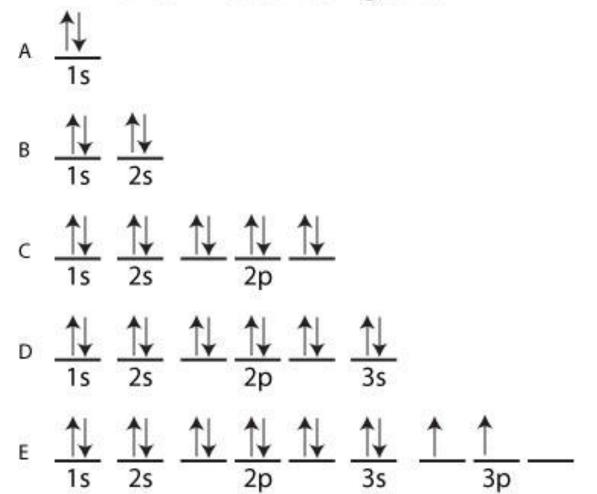


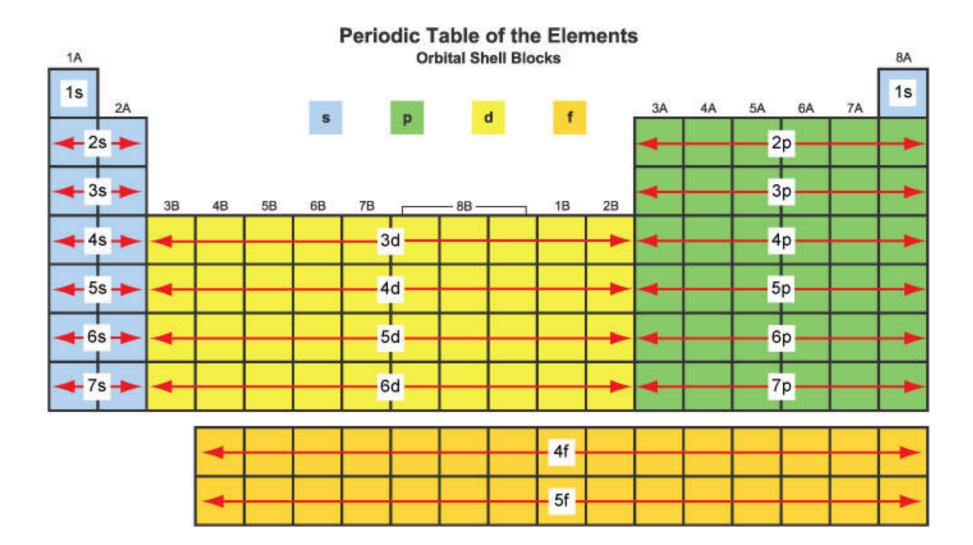




Hund's Rule (He/Be/Ne/Mg/Si)

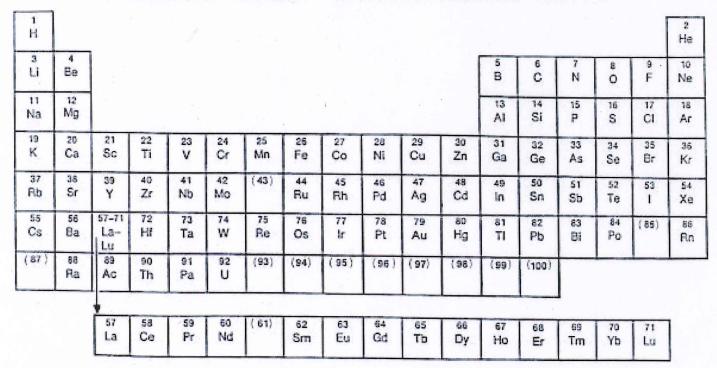
Silicon Electron Configuration





Pre-1941 Whoops!

PERIODIC TABLE - BEFORE WORLD WAR II



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Figure 1: Periodic table before World War II. Fermi irradiated U with neutrons and found products which are chemically similar to Mn and Re; he thought he had found element 93. Before Meitner proposed the theory of nuclear fission, she called the speculated elements 93, 94, 95 and 96 eka-rhenium, eka-osmium, eka-iridium and eka-platinum, respectively. Segre was convinced that element 93 would be chemically similar to Re and Mn, and missed the discovery of element 93, which is actually chemically similar to U.

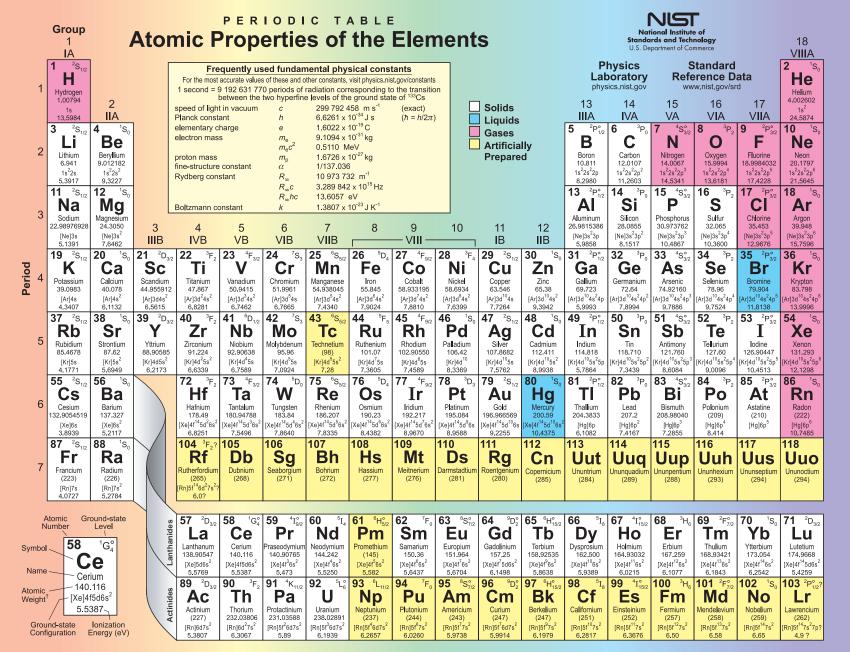
The Heroes of the Story



]	Peri	od 4	P	od 5	Pe	d 6	Period 7					
Element	Z	Electron Configuration	Element	Z	Electron Configuration	Element	z	Electron Configuration	Element	Z	Electron Configuration	
								$[Xe] 6s^2 \frac{5d^1}{5d^1}$	Actinium	89	[Rn] 7s ² 6d ¹	
						Cerium	58	[Xe] 6s2 4f1 5d ¹	Thorium	90	[Rn] 7s ² 6d ²	
						Praseodymium	59	$[Xe] 6s^2 4f^3$	Protactinium	91	[Rn] 7s ² 5f ² 6d ¹	
						Neodymium	60	[Xe] 6s ² 4f ⁴	Uranium	92	[Rn] 7s ² 5f ³ 6d ¹	
						Promethium	61	[Xe] 6s ² 4f ⁵	Neptunium	93	[Rn] 7s ² 5f ⁴ 6d ¹	
						Samarium	62	[Xe] 6s ² 4f ⁶	Plutonium	94	[Rn] 7s ² 5f ⁶	
						Europium		[Xe] 6s ² 4f ⁷	Americium	95	[Rn] 7s ² 5f ⁷	
						Gadolinium	64	[Xe] $6s^2 4f^7$ $5d^1$	Curium	96	[Rn] 7s ² 5f ⁷ 6d ¹	
						Terbium	65	[Xe] $6s^2 4f^9$	Berkelium	97	[Rn] 7s ² 5f ⁹	
											-	
Scandium	21	[Ar] 4s ² 3d ¹	Yttrium	39	[Kr] 5s ² 4d ¹	Lutetium	71	[Xe] 6s ² 4f ¹⁴ 5d ¹	Lawrencium	103	[Rn] 7s ² 5f ¹⁴ 7p ¹	
Titanium	22	[Ar] 4s ² 3d ²	Zirconium	40	[Kr] 5s ² 4d ²	Hafnium		5d ²	Rutherfordium	104	[Rn] 7s ² 5f ¹⁴ 6d ²	
Vanadium	23	$[Ar] 4s^2 3d^3$	Niobium	41	[Kr] 5s ¹ 4d ⁴			[Xe] 6s ² 4f ¹⁴ 5d ³			•	
Chromium	24	[Ar] 4s ¹ 3d ⁵	Molybdenum	42	[Kr] 5s ¹ 4d ⁵	Tungsten	74	[Xe] 6s ² 4f ¹⁴ 5d ⁴				
Manganese	25	[Ar] 4s ² 3d ⁵	Technetium	43	[Kr] 5s ² 4d ⁵	Rhenium	75	[Xe] 6s ² 4f ¹⁴ 5d ⁵				
Iron	26	[Ar] 4s ² 3d ⁶	Ruthenium	44	[Kr] 5s ¹ 4d ⁷	Osmium	76	[Xe] 6s ² 4f ¹⁴ 5d ⁶				
Cobalt	27	[Ar] 4s ² 3d ⁷	Rhodium	45	[Kr] 5s ¹ 4d ⁸	Iridium	77	[Xe] 6s ² 4f ¹⁴ 5d ⁷				
Nickel	28	[Ar] $4s^2 3d^8$ or [Ar] $4s^1 3d^9$ (disputed) ^[22]	Palladium	46	[Kr] 4d ¹⁰	Platinum		5d ⁹				
Copper	29	[Ar] 4s ¹ 3d ¹⁰	Silver	47	[Kr] 5s ¹ 4d ¹⁰			[Xe] 6s ¹ 4f ¹⁴ 5d ¹⁰				
Zinc	30	$[Ar] 4s^2 3d^{10}$	Cadmium	48	[Kr] 5s ² 4d ¹⁰	Mercury	80	[Xe] 6s ² 4f ¹⁴ 5d ¹⁰				

Electron shells filled in violation of Madelung's rule^[21] (red)

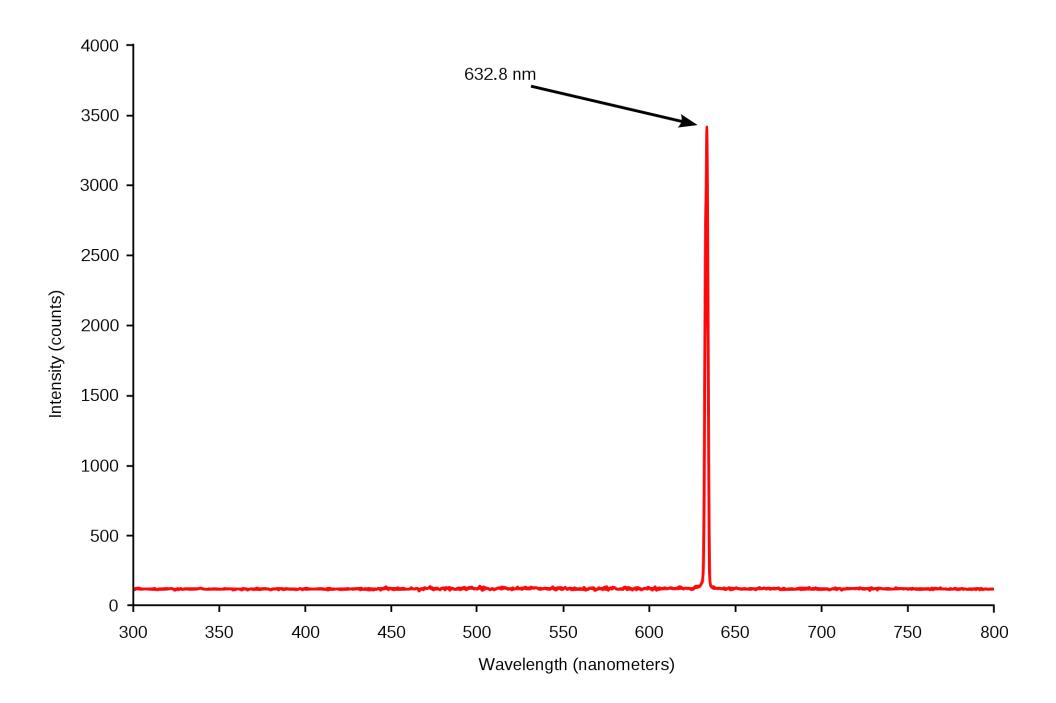
The electron-shell configuration of elements beyond rutherfordium is not yet known.

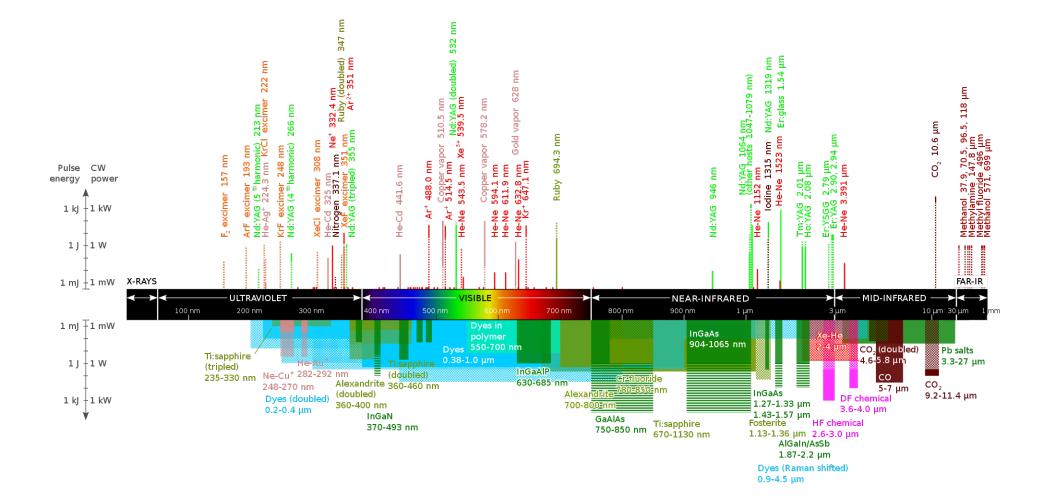


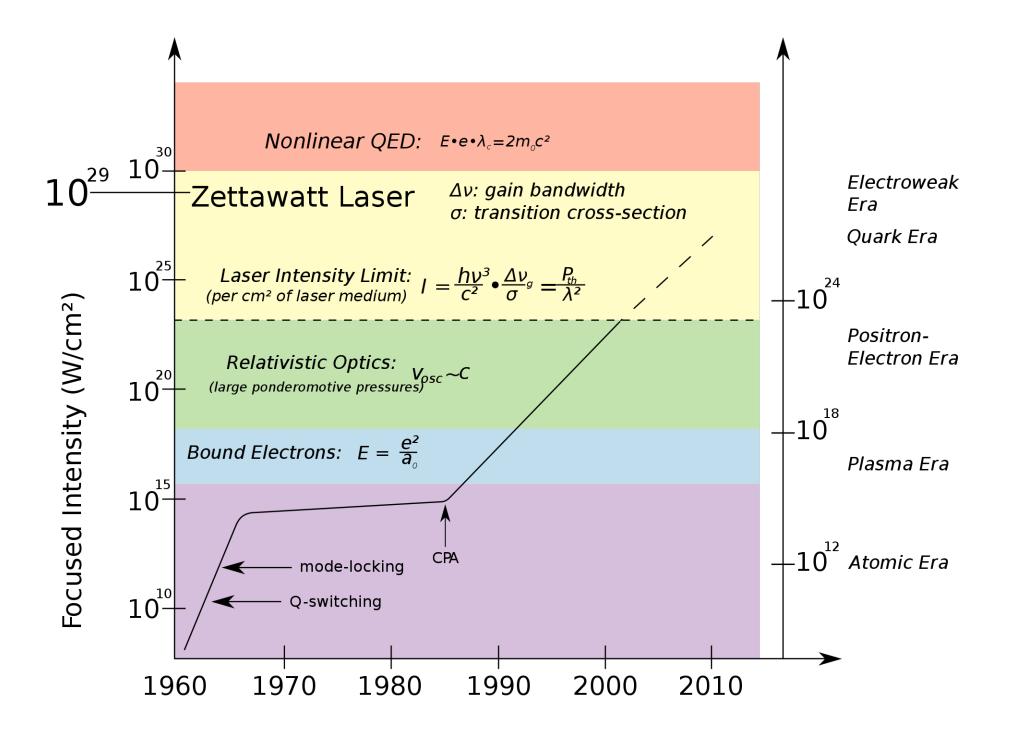
[†]Based upon ¹²C. () indicates the mass number of the longest-lived isotope.

For a description of the data, visit physics.nist.gov/data

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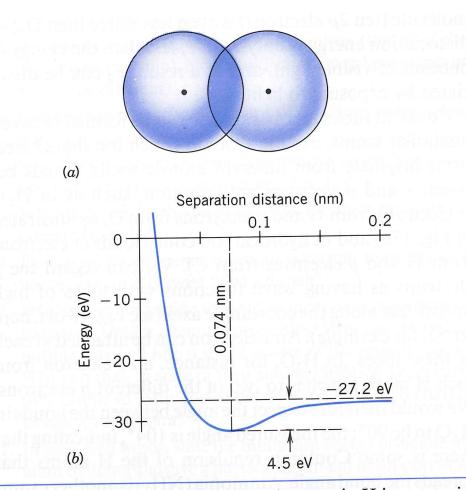


Figure 16 (a) The overlap of the s electrons in H is responsible for the formation of the H₂ molecule. (b) The total energy of the two electrons in the bound state of the H₂ molecule, as a function of the atomic separation distance. When the separation is large, the energy is -27.2 eV (twice the energy of the single electron in atomic hydrogen, -13.6 eV). The minimum energy of the bound molecule is -31.7 eV when the separation is 0.074 nm.

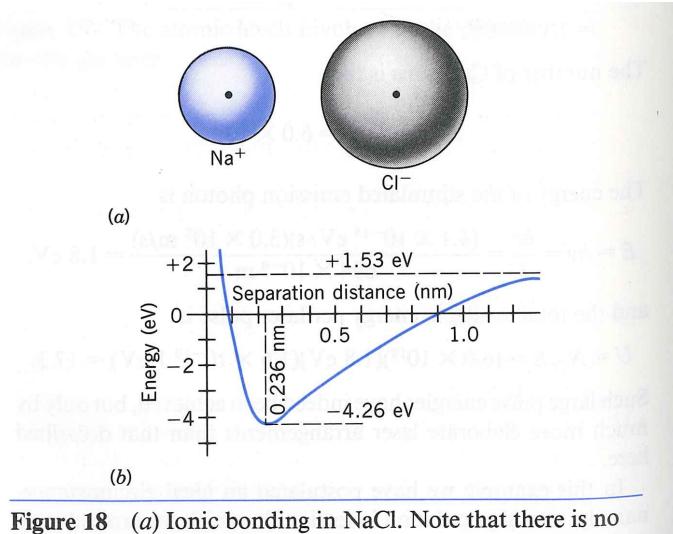
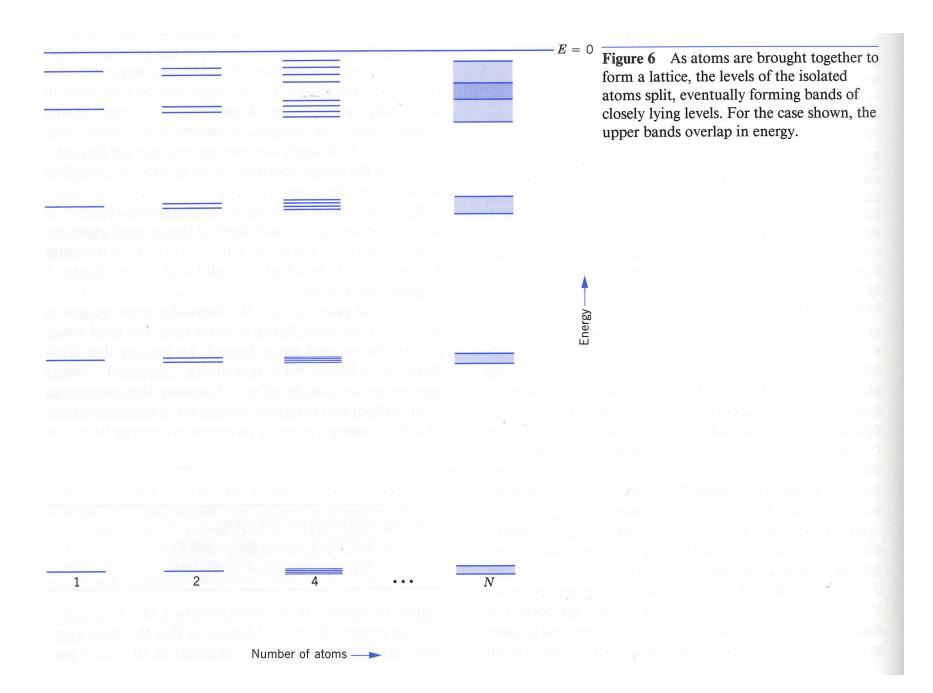
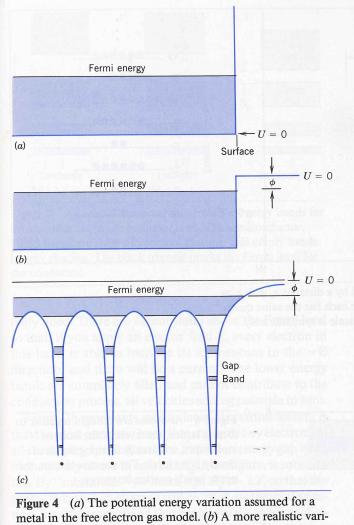
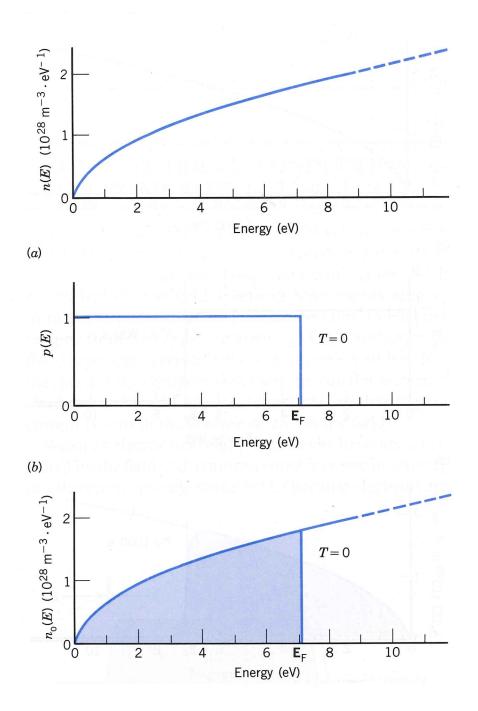


Figure 18 (a) Ionic bonding in NaCl. Note that there is no appreciable overlap of the electron distributions. (b) Binding energy in NaCl. The zero of energy corresponds to Na and Cl *atoms* separated by a large distance. The dashed line represents the energy of Na⁺ and Cl⁻ *ions* separated by a large dis-





right 4^{-1} (a) The potential energy valuation assumed for a metal in the free electron gas model. (b) A more realistic variation, showing a finite change in potential energy at the surface of the sample. (c) A still more realistic variation, taking the lattice of ion cores into account. This curve is a one-dimensional cut along a line of ion cores (shown as dots at the bottom of the figure). The shaded regions are the energy bands permitted for the electrons. Electrons are not permitted to have energies corresponding to the gaps.



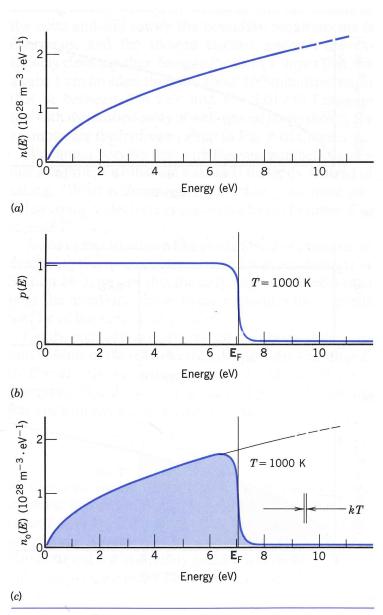
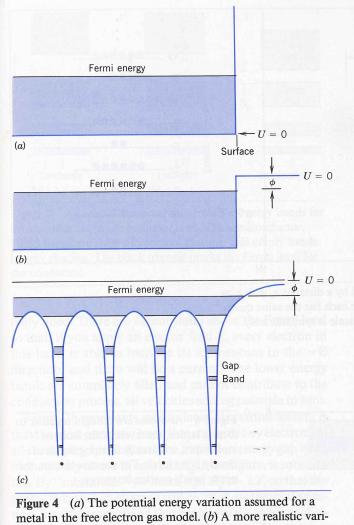


Figure 2 Same as Fig. 1, but for T = 1000 K. Note how little the plots differ from those of Fig. 1. (These plots are somewhat idealized in that they assume the electrons move in a region of uniform potential. Measured density of states plots in real metals do not have this simple shape.)



right 4^{-1} (a) The potential energy valuation assumed for a metal in the free electron gas model. (b) A more realistic variation, showing a finite change in potential energy at the surface of the sample. (c) A still more realistic variation, taking the lattice of ion cores into account. This curve is a one-dimensional cut along a line of ion cores (shown as dots at the bottom of the figure). The shaded regions are the energy bands permitted for the electrons. Electrons are not permitted to have energies corresponding to the gaps.

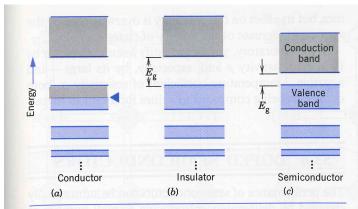


Figure 7 An idealized representation of the energy bands for (a) a conductor, (b) an insulator, and (c) a semiconductor. Filled bands are shown in colored shading, and empty bands in gray shading. The black triangle marks the Fermi level for the conductor.