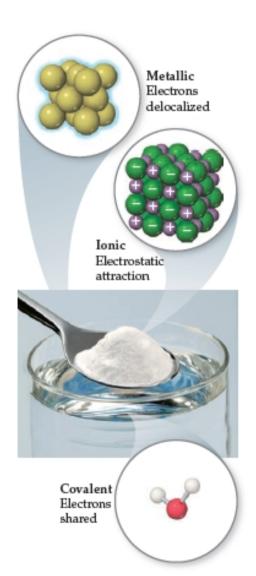
Ligações Químicas



Por que as propriedades das substâncias são tão diferentes?

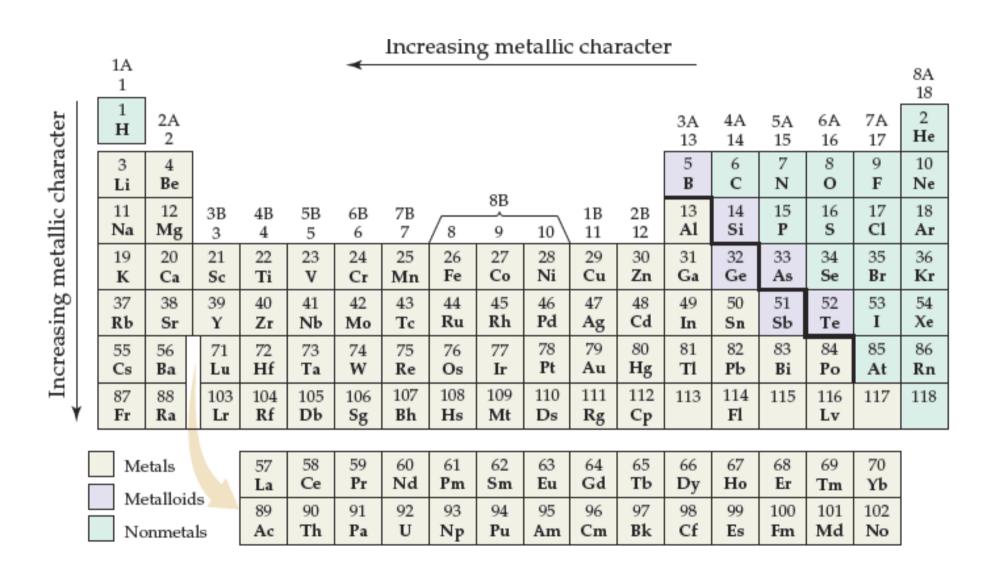
Tipos de Ligações Químicas

- Ligação iônica
- Ligação covalente
- Ligação metálica

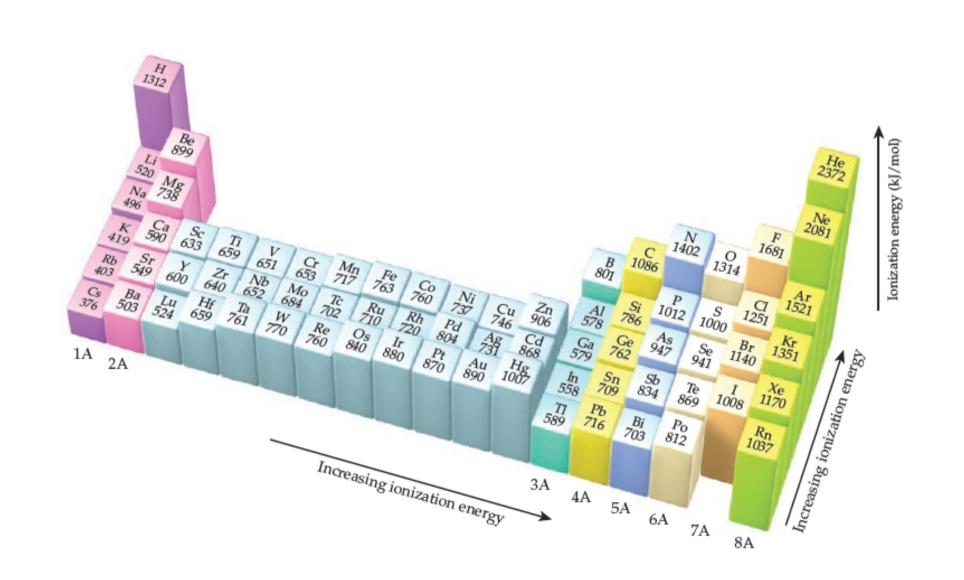
Por que os átomos se ligam?

Table 8.1 Lewis Symbols

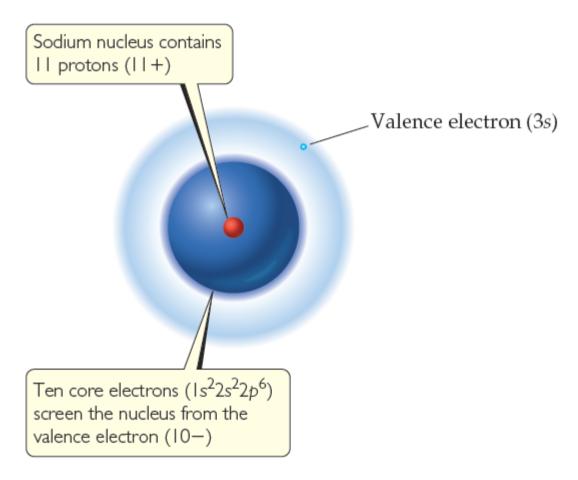
Group	Element	Electron Configuration	Lewis Symbol	Element	Electron Configuration	Lewis Symbol
1A	Li	[He]2s ¹	Li·	Na	[Ne]3s1	Na∙
2A	Be	$[He]2s^2$	·Be·	Mg	[Ne]3s ²	⋅Mg⋅
3A	В	$[He]2s^22p^1$	٠Ġ٠	Al	$[Ne]3s^23p^1$	٠Ål٠
4A	С	$[He]2s^22p^2$	٠Ċ٠	Si	[Ne] $3s^23p^2$	·Śi·
5A	N	$[He]2s^22p^3$	·Ņ:	P	[Ne] $3s^23p^3$	·ṛ:
6A	О	[He] $2s^22p^4$:ọ:	S	[Ne] $3s^23p^4$:\$:
7A	F	[He] $2s^22p^5$	·F:	Cl	[Ne]3s ² 3p ⁵	٠ċj:
8A	Ne	$[He]2s^22p^6$:Ne:	Ar	[Ne]3s ² 3p ⁶	:Àr:



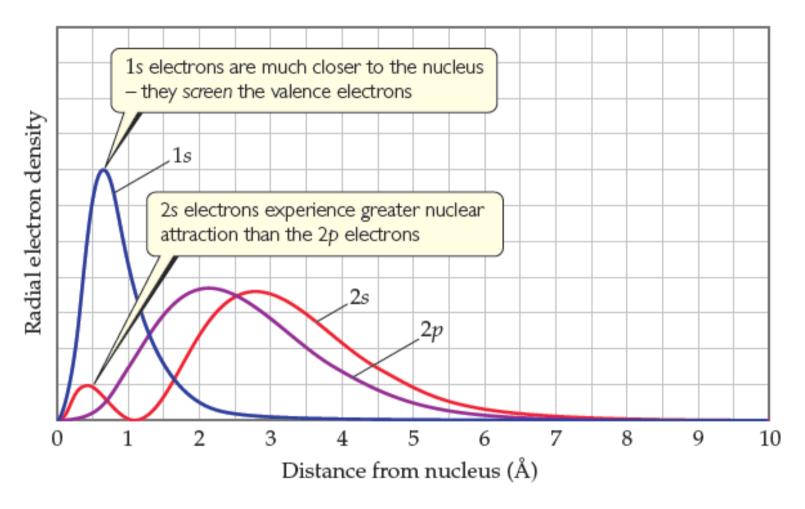
Energia de Ionização



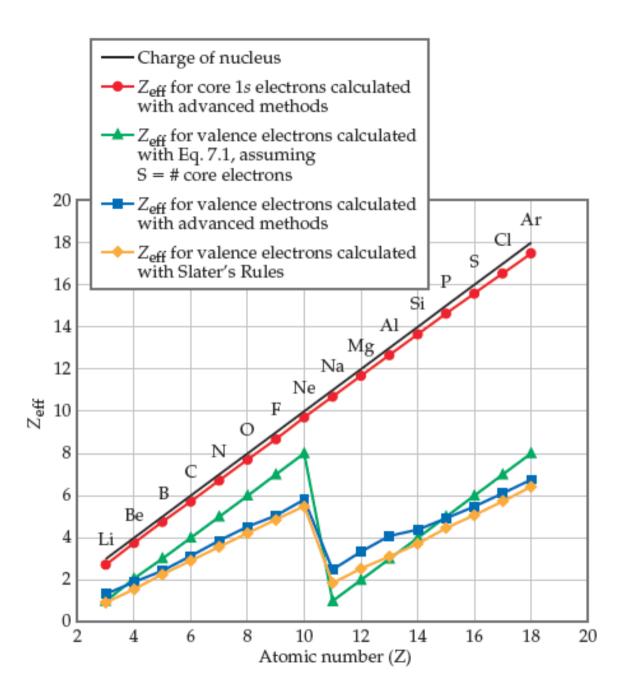
Raio Atômico



▲ Figure 7.3 Effective nuclear charge. The effective nuclear charge experienced by the 3s electron in a sodium atom depends on the 11+ charge of the nucleus and the 10− charge of the core electrons.



▲ Figure 7.4 Comparison of 1s, 2s, and 2p radial probability functions.



Formação de Redes Cristalinas

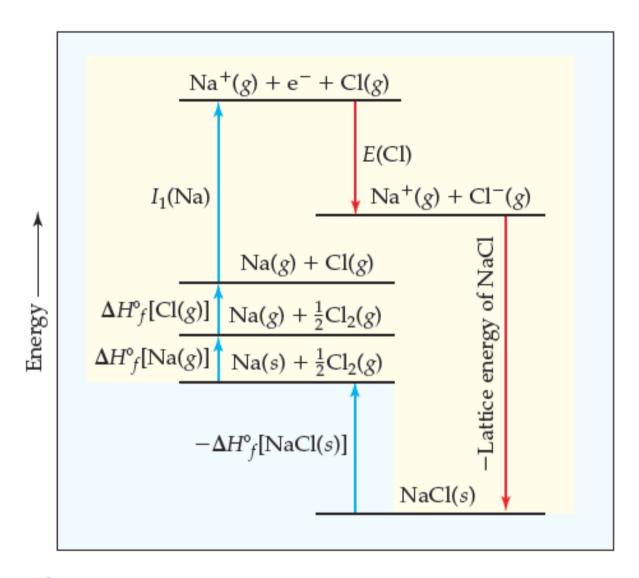
Table 8.2 Lattice Energies for Some Ionic Compounds

Compound	Lattice Energy (kJ/mol)	Compound	Lattice Energy (kJ/mol)
LiF	1030	MgCl_2	2326
LiCl	834	SrCl ₂	2127
LiI	730		
NaF	910	MgO	3795
NaCl	788	CaO	3414
NaBr	732	SrO	3217
NaI	682		
KF	808	ScN	7547
KCl	701		
KBr	671		
CsCl	657		
CsI	600		

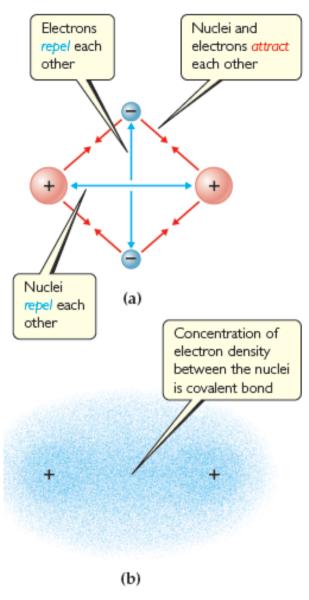
$$E_{lattice} = K \frac{Q_1 Q_2}{d}$$

Qual é a ordem crescente de energia de retículo cristalino?

NaF, CsI e CaO



▲ Figure 8.5 Born–Haber cycle for formation of NaCl. This Hess's law representation shows the energetic relationships in the formation of the ionic solid from its elements.



▲ Figure 8.6 The covalent bond in H₂.

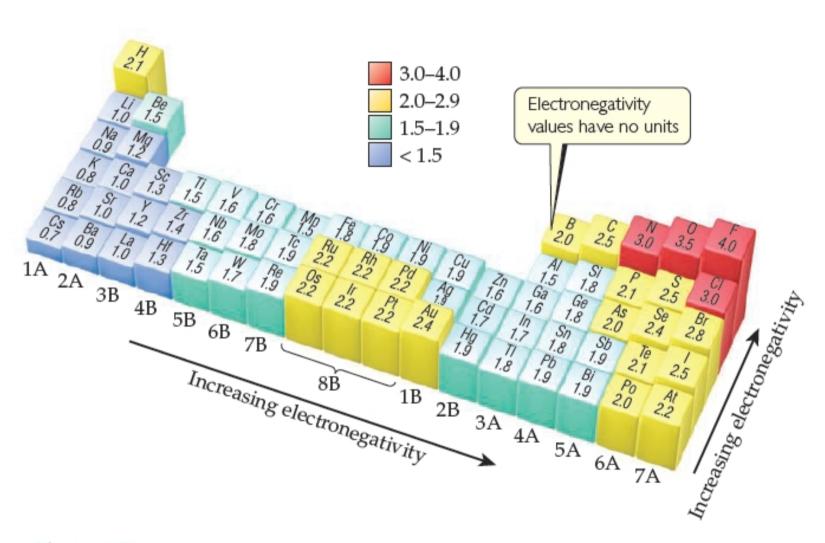
(a) The attractions and repulsions among electrons and nuclei in the hydrogen molecule. (b) Electron distribution in the H₂ molecule.

Ligação Covalente

$$H \cdot + \cdot H \longrightarrow H \cdot H$$

$$: \dot{\bigcirc}: + \cdot \dot{\bigcirc}: + : \dot{\bigcirc}: \longrightarrow \ddot{\bigcirc}:: C:: \ddot{\bigcirc} \qquad (or \ \ddot{\bigcirc} = C = \ddot{\bigcirc})$$

Polaridade de Ligações Químicas



▲ Figure 8.7 Electronegativity values based on Pauling's thermochemical data.

Electronegativity difference

 F_2

HF

LiF

4.0 - 4.0 = 0

4.0 - 2.1 = 1.9

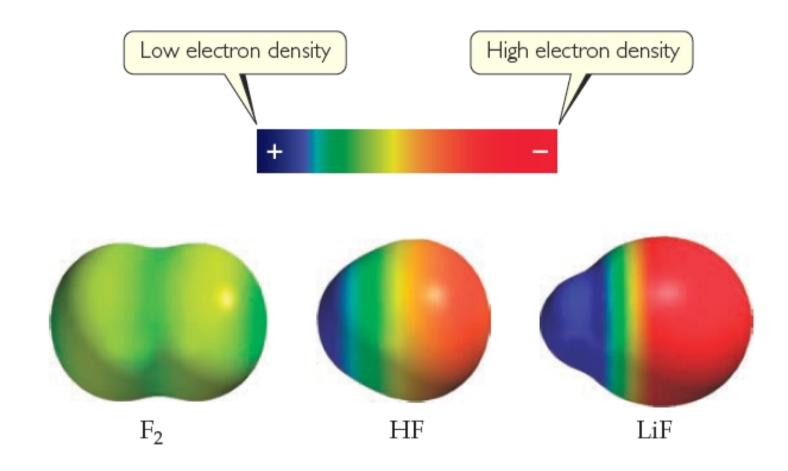
4.0 - 1.0 = 3.0

Type of bond

Nonpolar covalent

Polar covalent

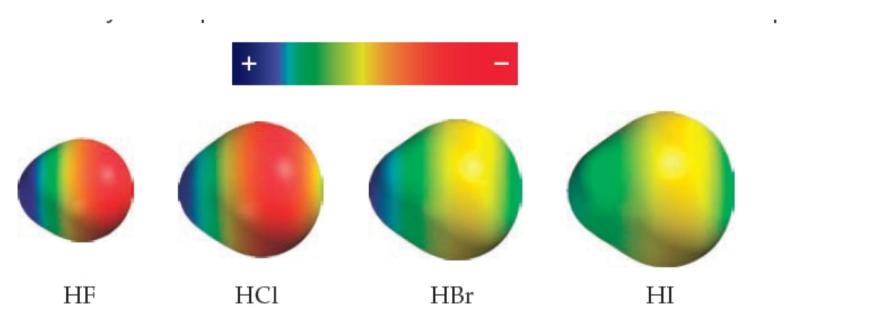
Ionic



Qual ligação é mais polar?

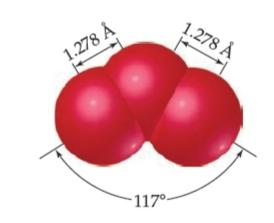
(a) B - Cl or C - Cl,

(b) P — F or P — Cl.

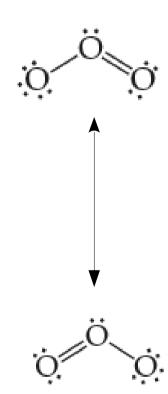


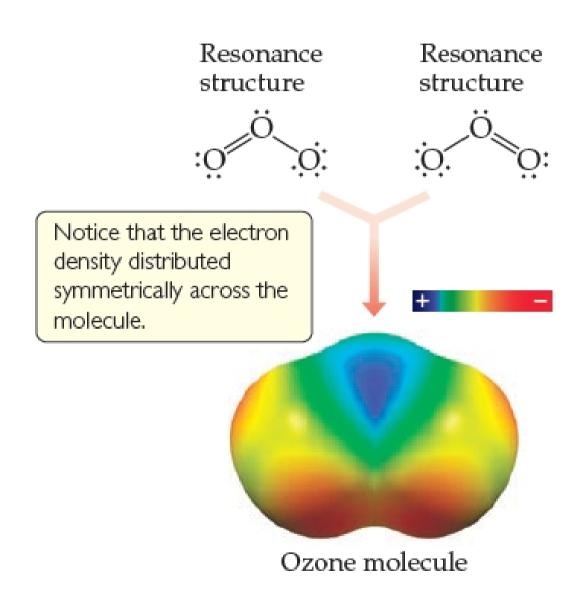
▲ Figure 8.10 Charge separation in the hydrogen halides. In HF, the strongly electronegative F pulls much of the electron density away from H. In HI, the I, being much less electronegative than F, does not attract the shared electrons as strongly, and, consequently, there is far less polarization of the bond.

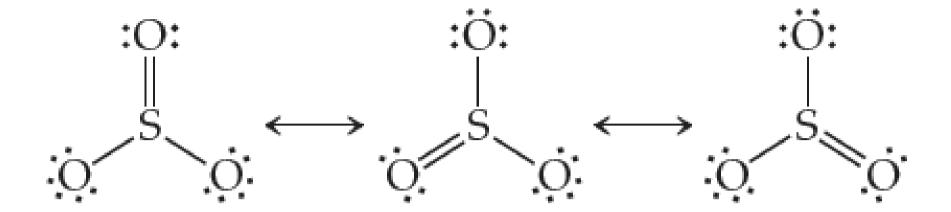
Estruturas de Ressonância

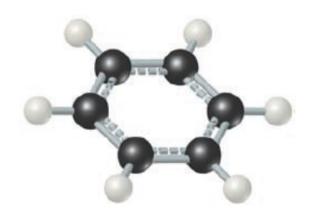


▲ Figure 8.12 Molecular structure of ozone.



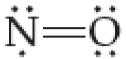




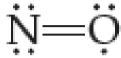


▲ Figure 8.14 Benzene, an "aromatic" organic compound. The benzene molecule is a regular hexagon of carbon atoms with a hydrogen atom bonded to each one. The dashed lines represent the blending of two equivalent resonance structures, leading to C—C bonds that are intermediate between single and double bonds.

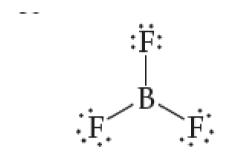
Exceções à Regra do Octeto



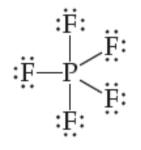
and

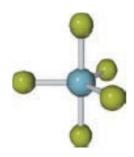


Número ímpar de elétrons



Número de elétrons menor que 8





Número de elétrons maior que 8 (átomos hipervalentes)