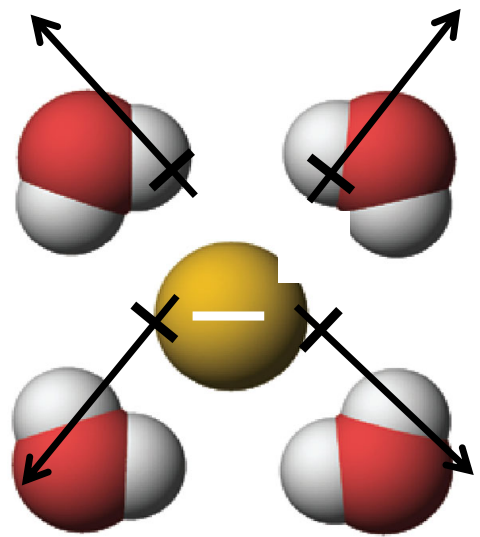
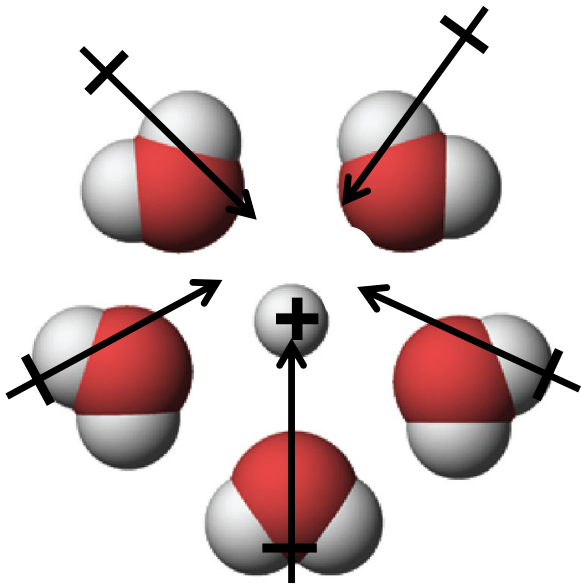
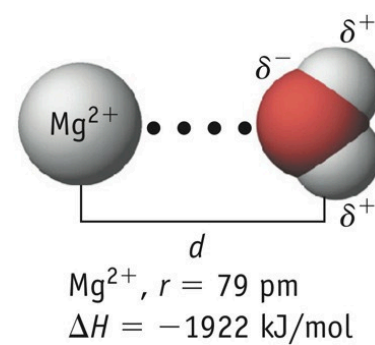
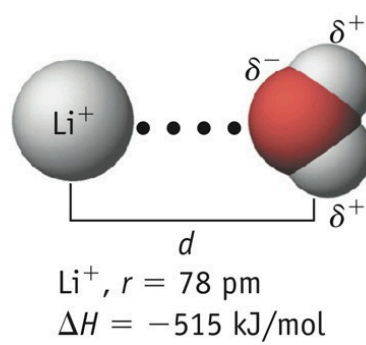
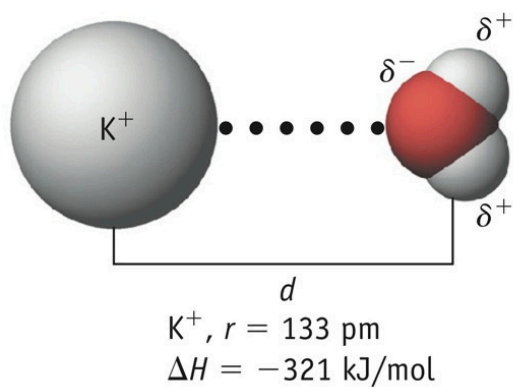


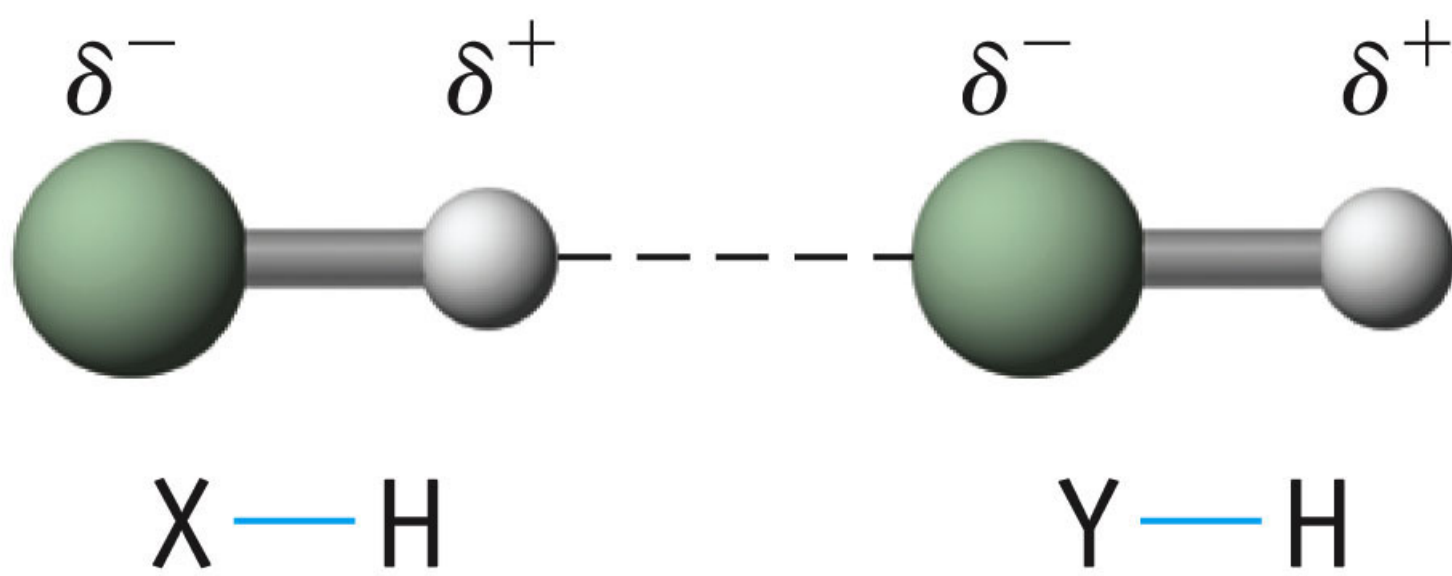
Solvation of Ions



When a *cation* exists in solution, it is surrounded by the *negative* dipole ends of water molecules.
When as *anion* exists in solution, it is surrounded by the *positive* dipole ends of water molecules.



Increasing force of attraction; more exothermic enthalpy of hydration

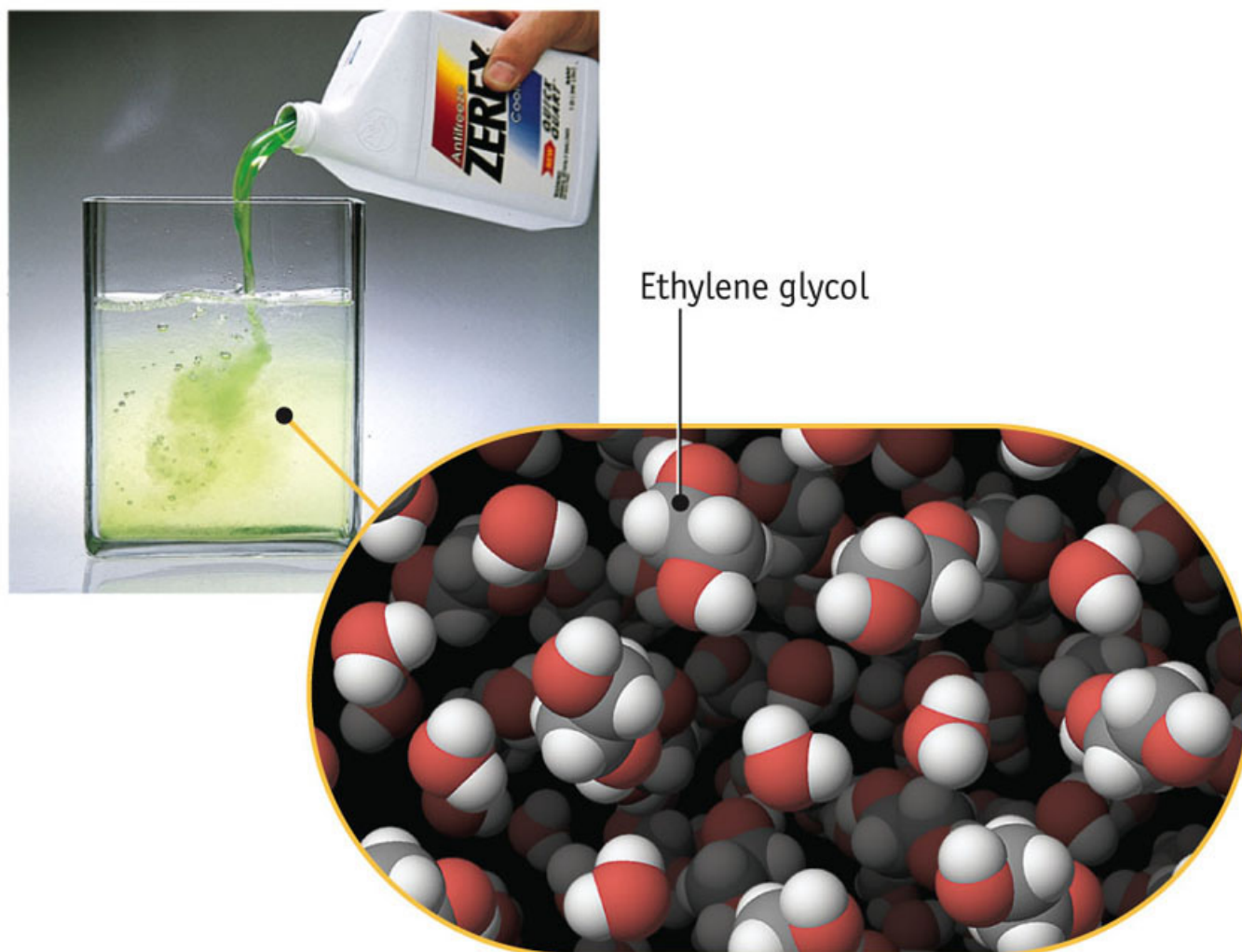


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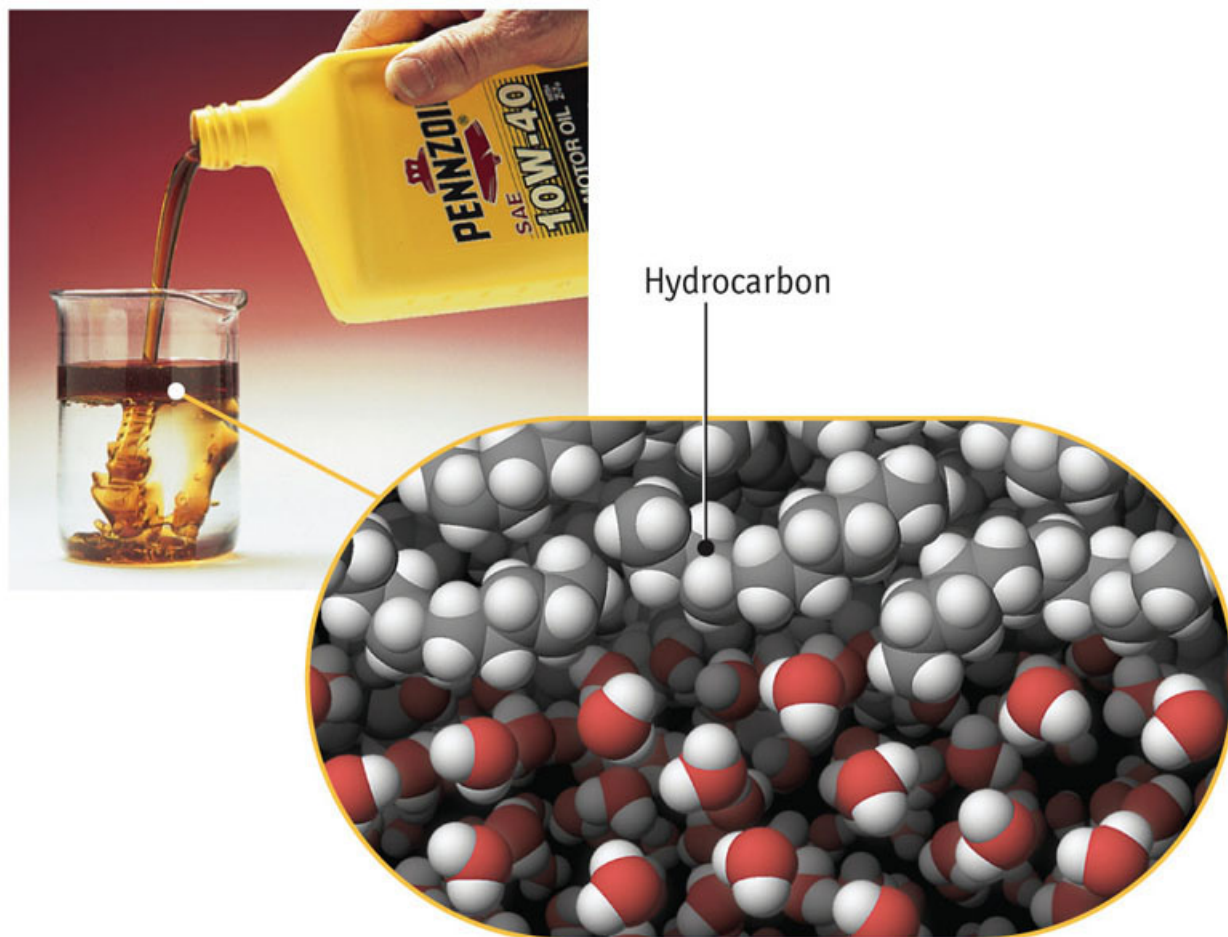
Dipole – Dipole Forces

Table 11.2 Molar Masses, Boiling Points, and $\Delta_{\text{vap}}H^\circ$ of Nonpolar and Polar Substances

NONPOLAR				POLAR			
	<i>M</i> (g/mol)	BP (°C)	$\Delta_{\text{vap}}H^\circ$ (kJ/mol)		<i>M</i> (g/mol)	BP (°C)	$\Delta_{\text{vap}}H^\circ$ (kJ/mol)
N ₂	28	−196	5.57	CO	28	−192	6.04
SiH ₄	32	−112	12.10	PH ₃	34	−88	14.06
GeH ₄	77	−90	14.06	AsH ₃	78	−62	16.69
Br ₂	160	59	29.96	ICl	162	97	—



(a) Ethylene glycol ($\text{HOCH}_2\text{CH}_2\text{OH}$), a polar compound used as antifreeze in automobiles, dissolves in water.

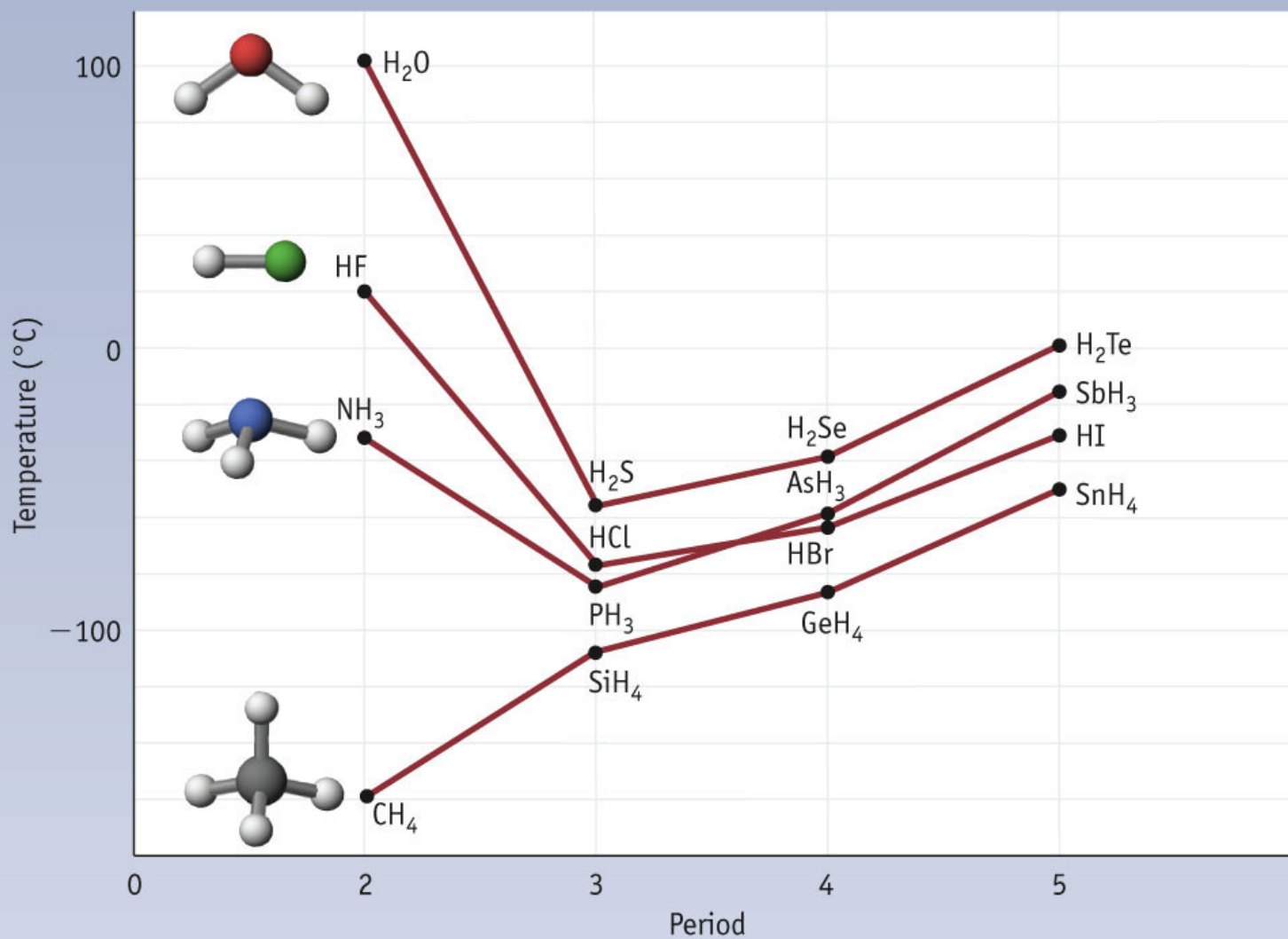


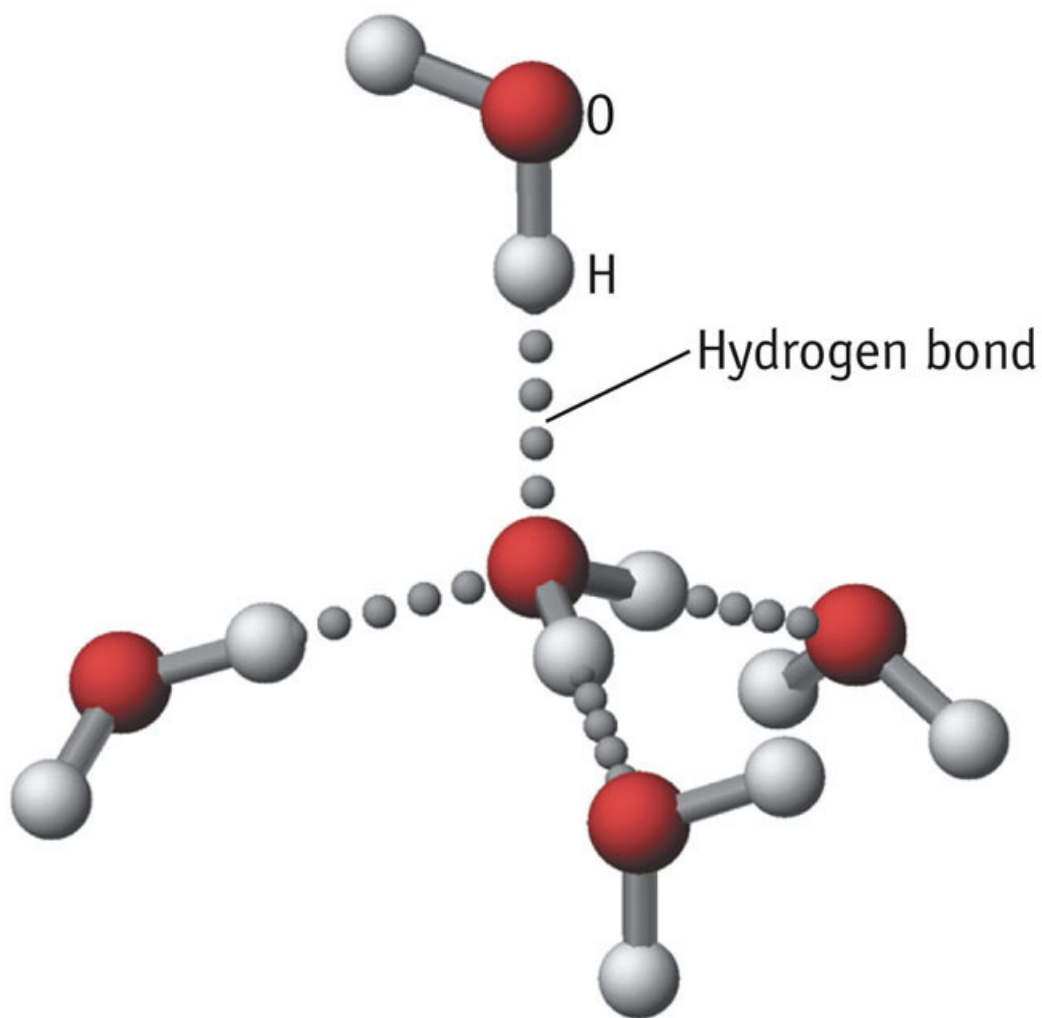
(b) Nonpolar motor oil (a hydrocarbon) dissolves in nonpolar solvents such as gasoline or CCl_4 . It will not dissolve in a polar solvent such as water, however. Commercial spot removers use nonpolar solvents to dissolve oil and grease from fabrics.

Dipole – Dipole Forces

Table 11.2 Molar Masses, Boiling Points, and $\Delta_{\text{vap}}H^\circ$ of Nonpolar and Polar Substances

NONPOLAR				POLAR			
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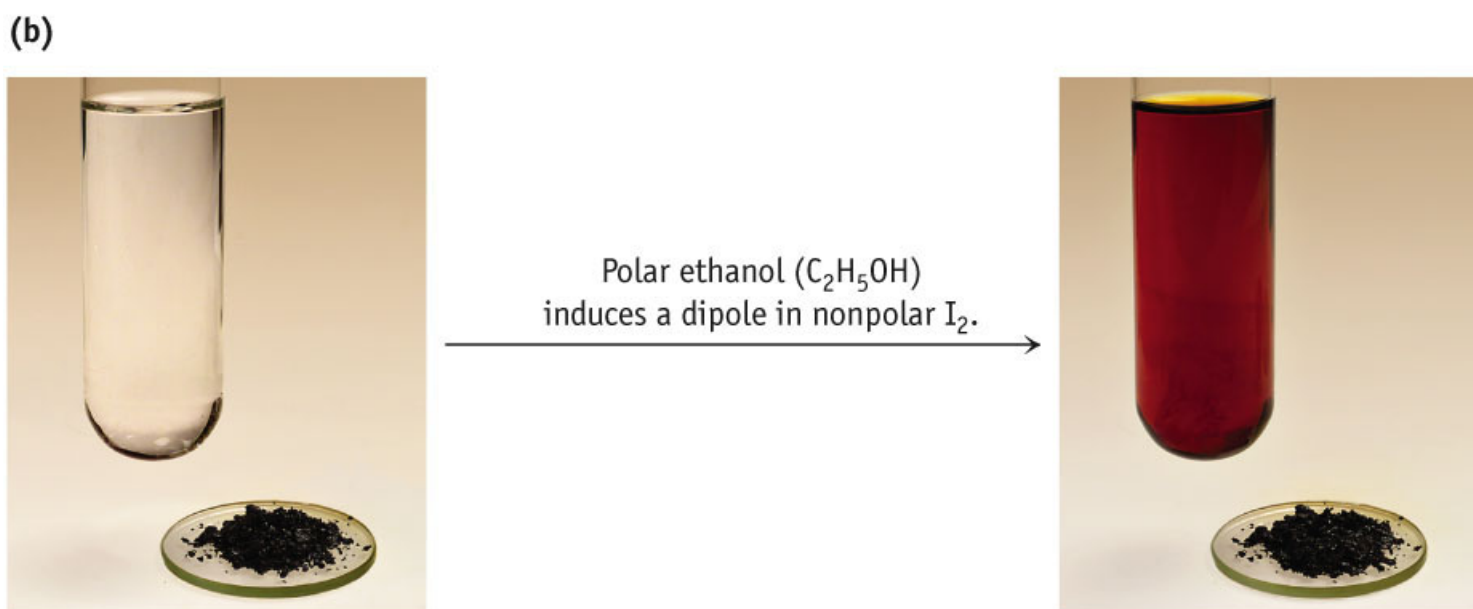
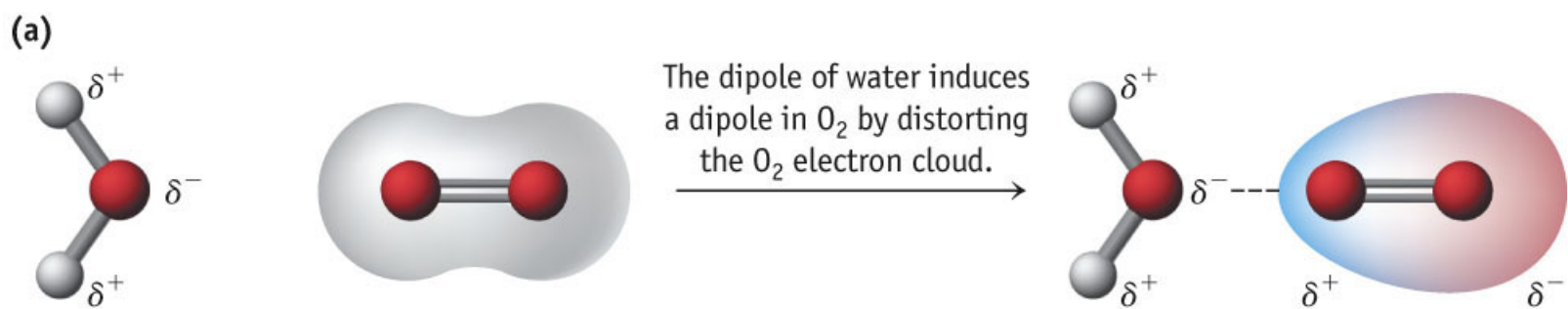




(a)

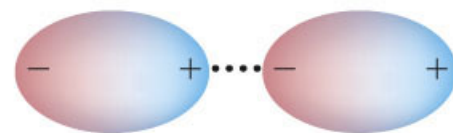
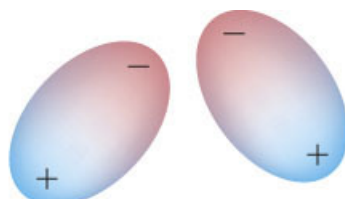
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Fig. 12-8, p. 564



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Fig. 12-10, p. 566



Two nonpolar atoms or molecules (depicted as having an electron cloud that has a time-averaged spherical shape).

Momentary attractions and repulsions between nuclei and electrons in neighboring molecules lead to induced dipoles.

Correlation of the electron motions between the two atoms or molecules (which are now dipolar) leads to a lower energy and stabilizes the system.

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Fig. 12-11, p. 567



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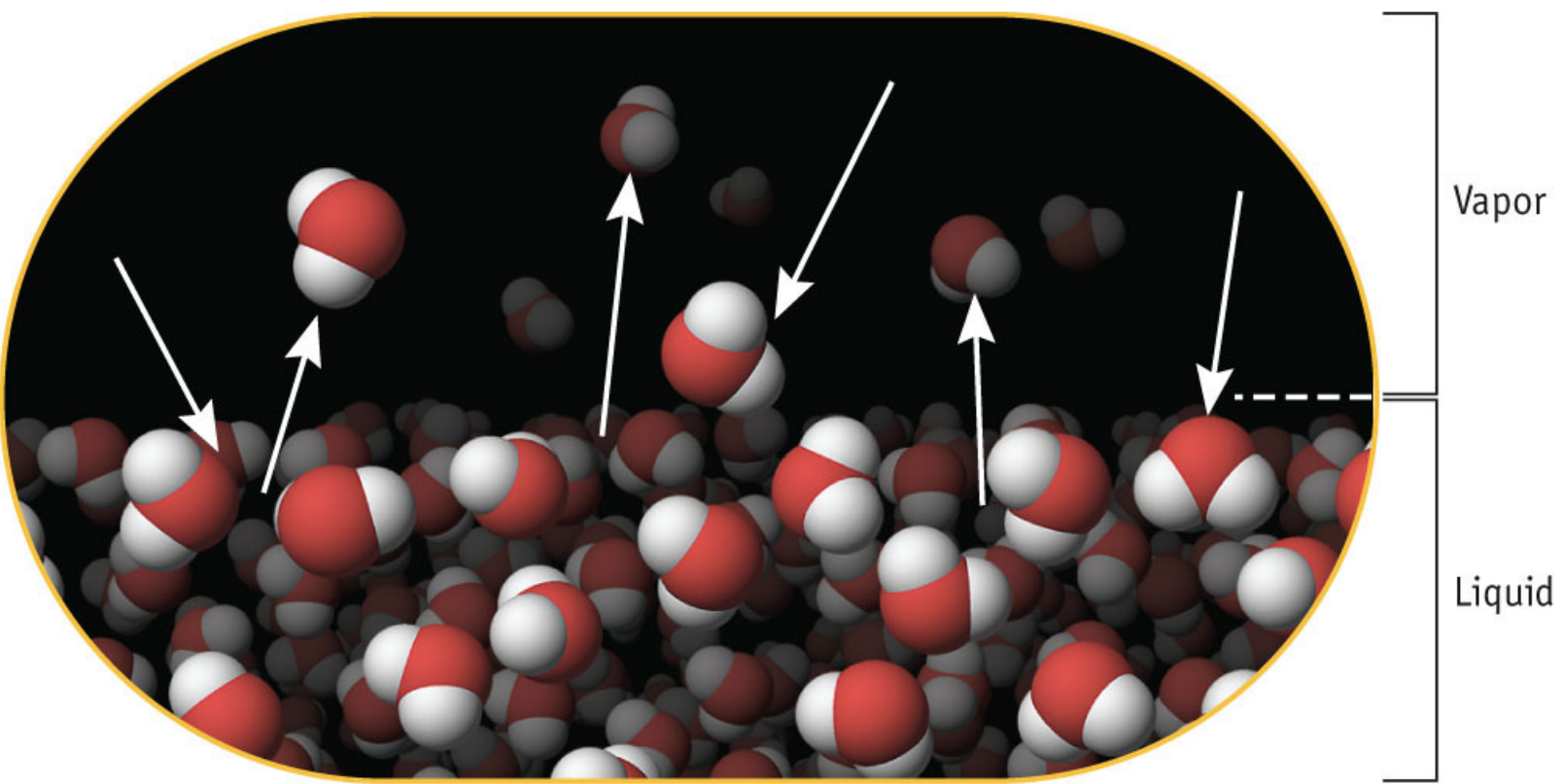


TABLE 12.6 Molar Enthalpies of Vaporization and Boiling Points for Common Substances*

Compound	Molar Mass (g/mol)	$\Delta_{\text{vap}}H^\circ$ (kJ/mol)†	Boiling Point (°C) (Vapor pressure = 760 mm Hg)
<i>Polar Compounds</i>			
HF	20.0	25.2	19.7
HCl	36.5	16.2	−84.8
HBr	80.9	19.3	−66.4
HI	127.9	19.8	−35.6
NH ₃	17.0	23.3	−33.3
H ₂ O	18.0	40.7	100.0
SO ₂	64.1	24.9	−10.0
<i>Nonpolar Compounds</i>			
CH ₄ (methane)	16.0	8.2	−161.5
C ₂ H ₆ (ethane)	30.1	14.7	−88.6
C ₃ H ₈ (propane)	44.1	19.0	−42.1
C ₄ H ₁₀ (butane)	58.1	22.4	−0.5
<i>Monatomic Elements</i>			
He	4.0	0.08	−268.9
Ne	20.2	1.7	−246.1
Ar	39.9	6.4	−185.9
Xe	131.3	12.6	−108.0
<i>Diatomic Elements</i>			
H ₂	2.0	0.90	−252.9
N ₂	28.0	5.6	−195.8
O ₂	32.0	6.8	−183.0
F ₂	38.0	6.6	−188.1
Cl ₂	70.9	20.4	−34.0
Br ₂	159.8	30.0	58.8

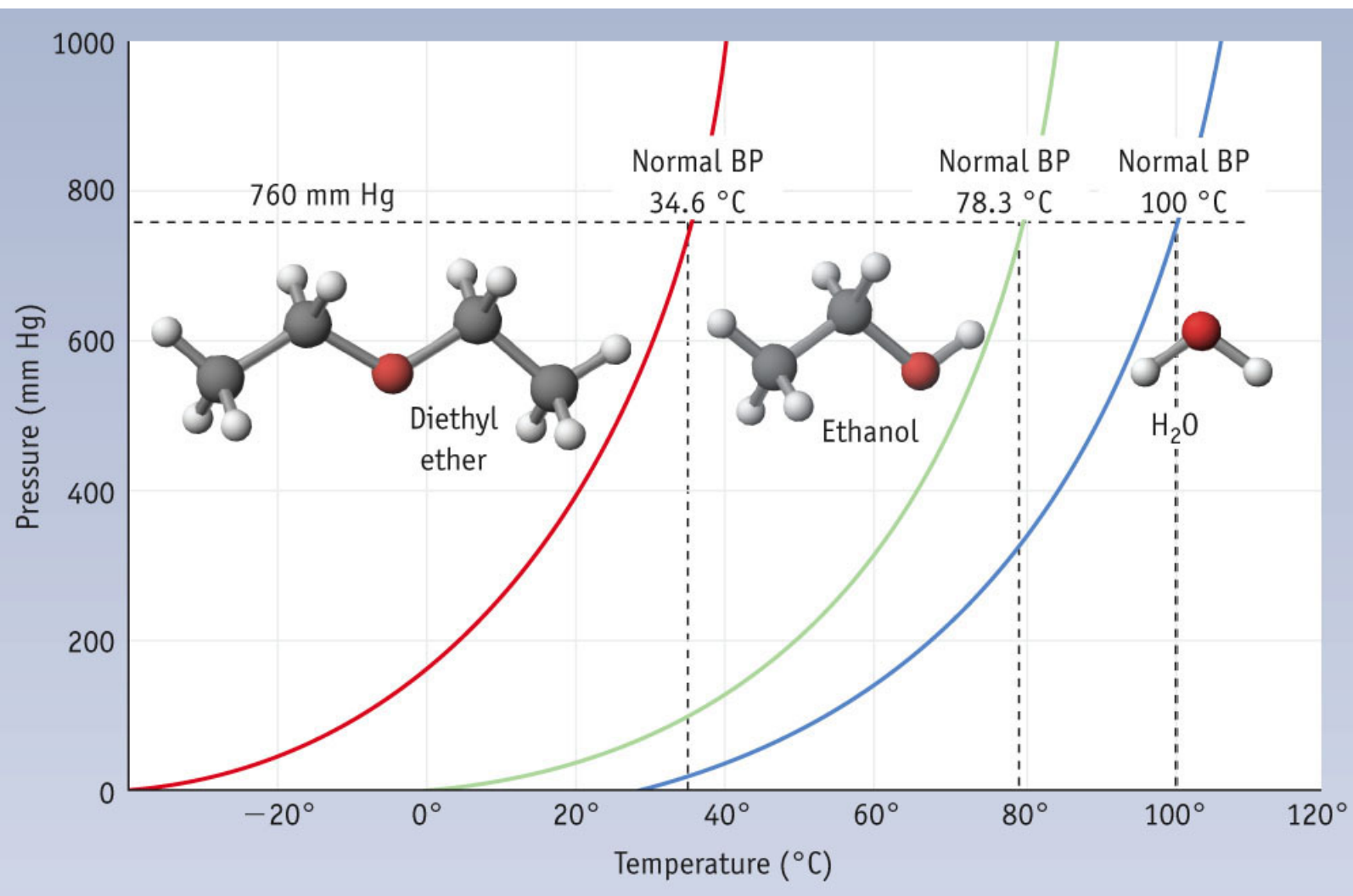
*Data taken from D. R. Lide: *Basic Laboratory and Industrial Chemicals*, Boca Raton, FL, CRC Press, 1993.

† $\Delta_{\text{vap}}H^\circ$ is measured at the normal boiling point of the liquid.



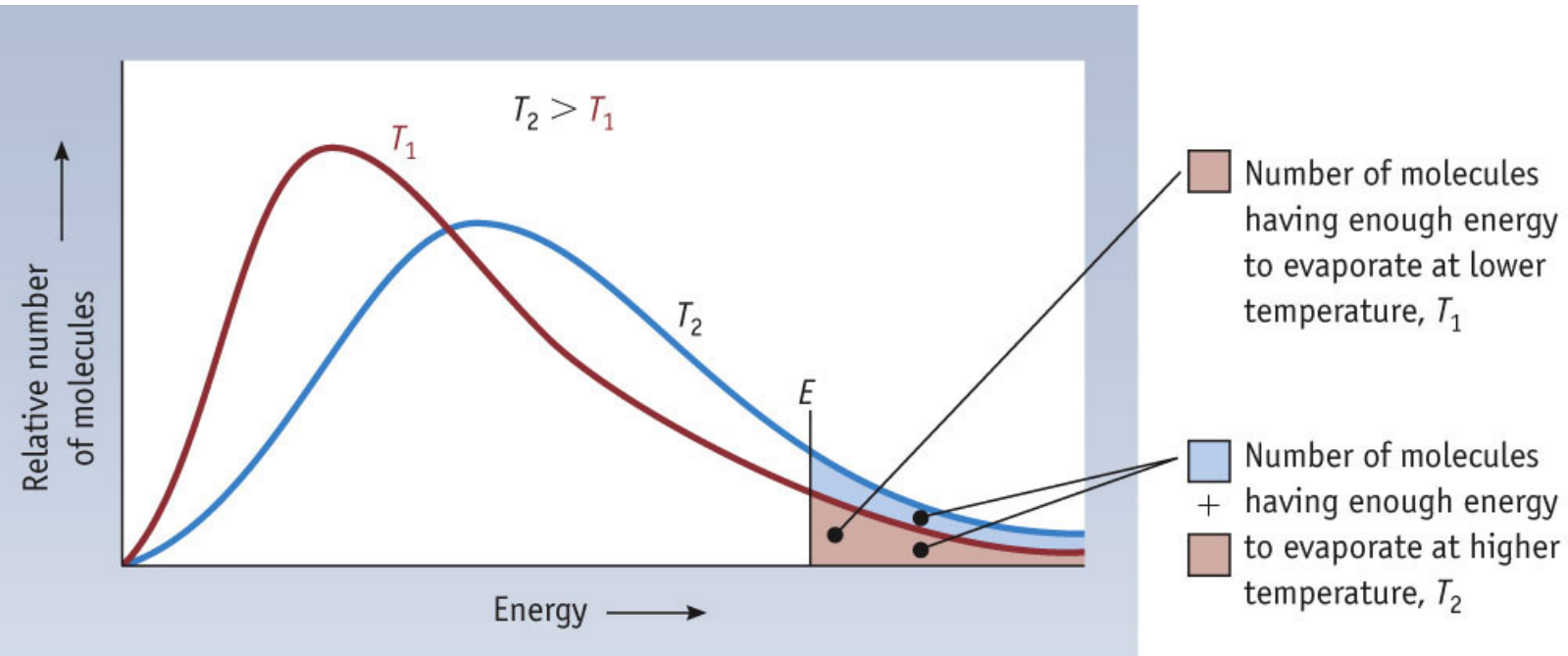
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Fig. 12-14, p. 571



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Fig. 12-17, p. 574



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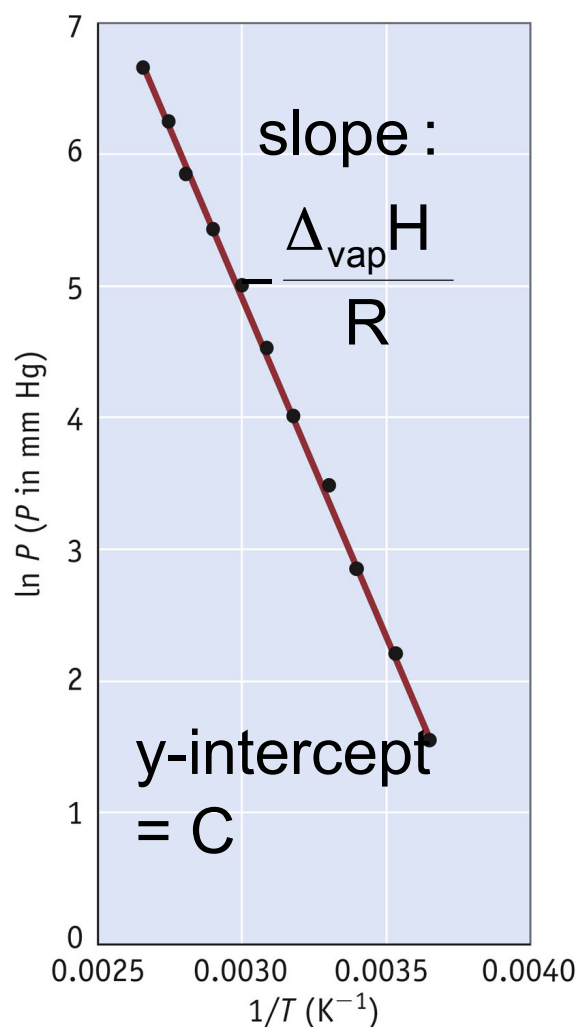
Fig. 12-13, p. 571

The Temperature Dependence of Vapor Pressure Goes As:

$$\ln P_{\text{vap}} = -\frac{\Delta_{\text{vap}} H^{\circ}}{RT} + C$$

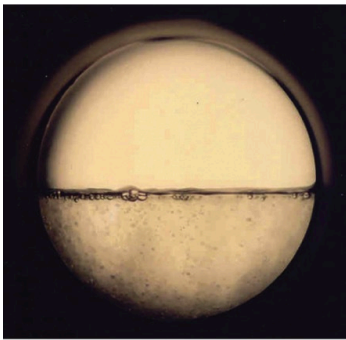
A plot of $\ln P_{\text{vap}}$ vs. $\frac{1}{T}$ yields a slope of:

$\Delta_{\text{vap}} H^{\circ}$ is related to T and P by the *Clausius-Clapeyron* equation

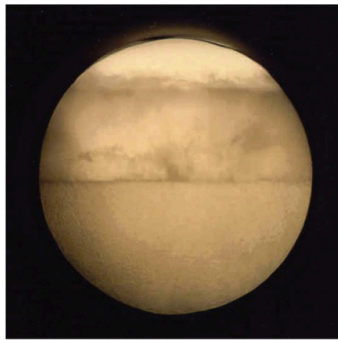


Critical Temperature and Pressure for CO₂

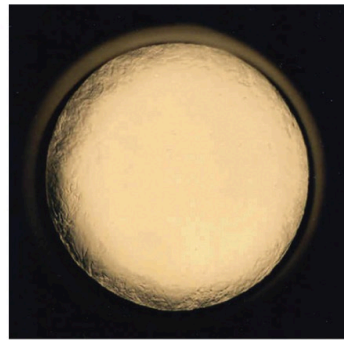
The separate phases of CO₂ are seen through the window in a high-pressure vessel.



As the sample warms and the pressure increases, the meniscus becomes less distinct.



Once the critical T and P are reached, distinct liquid and vapor phases are no longer in evidence. This homogeneous phase is "supercritical CO₂."



Photos: Dr. Christopher M. Rayner/
University of Leeds

