

Chp. 14 Acids/Bases

Characteristics:

Acids

litmus ^{re} acid

pH < 7

sour taste - citrus fruit

corrosive

caustic - burns

electrolyte - strong

formed from acid

anhydride + H₂O

↑
nonmetallic oxide SO₂
CO₂



Acid/Base Theories: 3 of these

Bases

pH > 7

bitter taste

corrosive

caustic

electrolyte

formed from

basic anhydride

+ H₂O

soap
baking
soda

metallic
oxide

MgO
CaO

NH_3^-
base-
not a
hydroxide
compd

1. **Arrhenius**

Acids produce $[\text{H}^+]$ in solution
Bases produce $[\text{OH}^-]$ in solution

Limiting--must have solutions and
bases must be hydroxides

2. **Bronsted-Lowry**

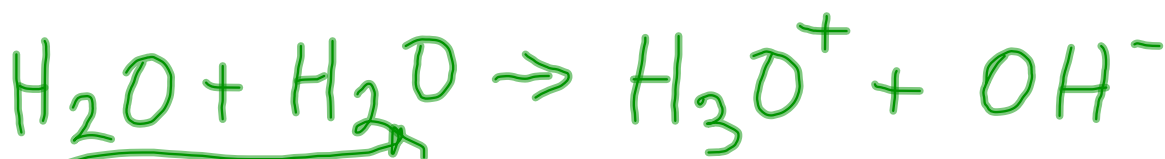
Acid is proton (H^+) donor
Base is proton acceptor

Broader—can include gases

3. **Lewis**

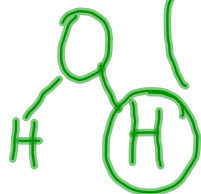
Acid accepts lone electron pair.
Base donates lone electron pair.

Broadest of the 3



general equation for dissolution of acid:

hydronium ion

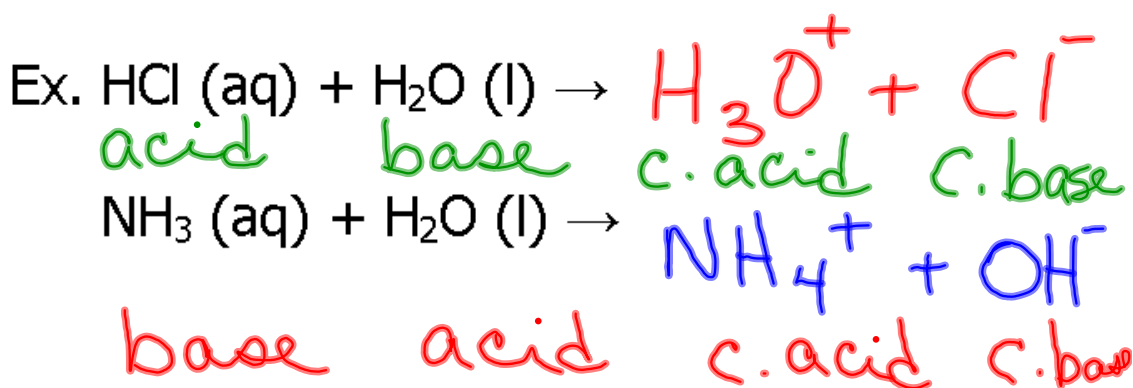


(acid) (base) (c. acid) (c. base)

Conjugate acid—substance formed when a base gains a proton (*base + H⁺*)

Conjugate base—ion that's left after the acid has donated a proton (*acid left over*)

Note: water can act as a base because of the lone prs. of electrons on the oxygen atom.



amphoteric subst -
has properties of acids
and bases
ex H_2O

Practice:

1. Give the conjugate base for each of the following Bronsted-Lowry acids:

what's left of acid

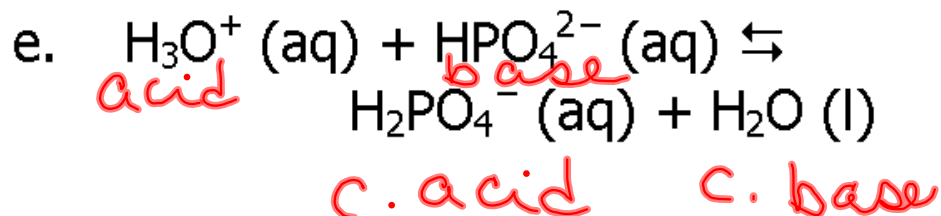
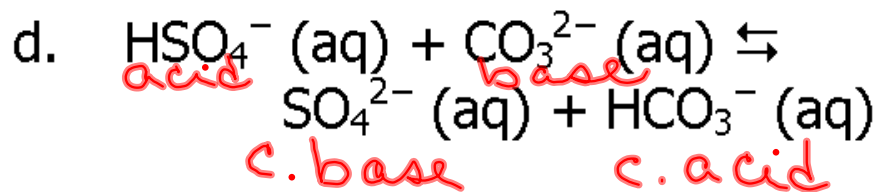
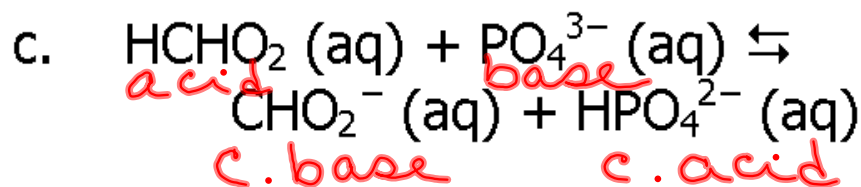
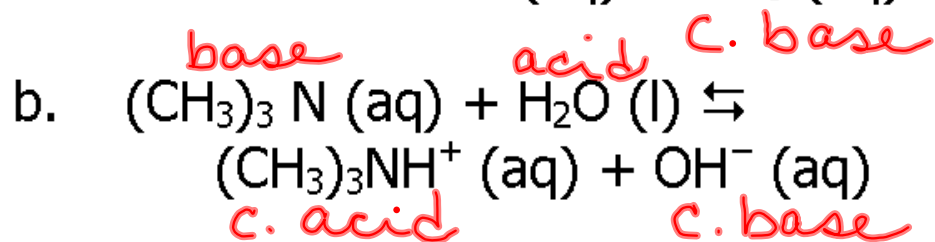
- a. HClO_2 ClO_2^-
- b. H_2S HS^-
- c. HSO_4^- SO_4^{2-}
- d. NH_4^+ NH_3

2. Give the conjugate acid for the following Bronsted-Lowry bases:

add proton

- a. IO^- HIO
- b. CH_3COO^- CH_3COOH
- c. HASO_4^- H_2AsO_4
- d. NH_3 NH_4^+

3. For each of the following equations label the Bronsted-Lowry acid, Bronsted-Lowry base, conjugate acid and conjugate base.



Acid dissociation constant (K_a):

Equilibrium expression for a general acid dissolved in water reaction

$$K_a = \frac{[\text{H}_3\text{O}^+][\text{A}^-]}{[\text{HA}]}$$

Water is not included because in dilute soln. the conc. of water is high and changes so little it is assumed to be constant.

If H_2O is a stronger base than A^- (water is more attracted to the H^+ than the A^-) the equilibrium position will lie to the far right (products favored) because most of the acid has dissolved (ionized). K has a large value.

If the A^- is a stronger base than the water the equilibrium position will lie to the far left (reactants favored). Most of the acid (HA) will be present in the equilibrium mixture (acid has partially ionized). K has a small value.

~~Do the examples on p. 656 (black book p. 639-40).~~

Strong acid:

Almost all is ionized at equilibrium

Has weak conjugate base (one that's not attracted to a p^+ and is weaker than water)

Equilibrium lies to the right

ex. H_2SO_4 , HCl , $HClO_4$, HNO_3 , HBr

Weak Acid:

Most of original acid is still present at equilibrium – dissociates a little

Has strong conjugate base (conjugate base is more attracted to proton than water)

Equilibrium lies to left

Ex. H_3PO_4 , HNO_2 , HClO

Most acids are oxyacids—have acidic H^+ attached to oxygen atom

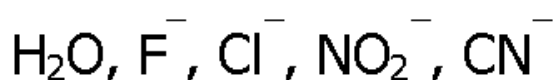
Organic acids contain C backbone and a carboxyl group

equilibrium lies to left

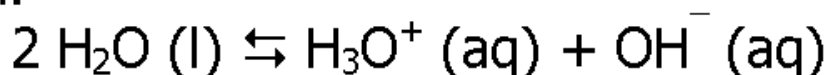
are weak acids

ex. CH_3COOH , $\text{C}_6\text{H}_5\text{COOH}$

Arrange the following bases in order of increasing strength: (remember H₂O is stronger base than the conjugate base of a strong acid but weaker than the conjugate base of a weak acid)



Water is amphoteric. It can behave as an acid or a base. It autoionizes—transfers proton from water molecule to another to produce a hydroxide ion and a hydronium ion.



NH₃ is also amphoteric.