

## 14.5: The pH of Solutions of Weak Bases

The pH of a solution of a weak base can be calculated in a way which is very similar to that used for a weak acid. Instead of an acid constant  $K_a$ , a base constant  $K_b$  must be used. If a weak base B accepts protons from water according to the equation

$$B + H_2O \rightleftharpoons BH^+ + OH^- \tag{14.5.1}$$

then the base constant is defined by the expression

$$K_b = \frac{{
m BH^+~OH^-}}{{
m B}}$$
 (14.5.2)

A list of  $K_b$  values for selected bases arranged in order of strength is given in the table below. This table is part of our larger collection of acid-base resources.

Table 14.5.1: The Base Constants for Some Bases at 25°C. Taken from Hogfelt, E. Perrin, D. D. Stability Constants of Metal Ion Complexes, 1<sup>st</sup> ed. Oxford; New Pergamon, 1979-1982. International Union of Pure and Applied Chemistry, Commission on Equilibrium. ISBN:

0080209580				
Base	Formula and Ionization Equation	$K_b$	Molecular Shape	
Ammonia	$NH_3 + H_2O  ightleftharpoons NH_4^+ + OH^-$	1.77 × 10 <sup>-5</sup>		
Aniline	$C_6H_5NH_2+H_2O ightleftharpoons C_6H_5NH_3^+$	$^{-} + OH^{-}$ 3.9 × 10 <sup>-10</sup>		
Carbonate ion	$CO_3^{2-} + H_2O  ightharpoonup HCO_3^- + OH^-$	2.1 × 10 <sup>-4</sup>		
Hydrazine	$N_2H_4 + H_2O  ightharpoonup N_2H_5^+ + OH^- \ N_2H_5^+ + H_2O  ightharpoonup N_2H_6^{2+} + OH^-$	$K_1 = 1.2 \times 10^{-6}$ $K_2 = 1.3 \times 10^{-15}$	330	
Hydride ion	$H^- + H_2O  ightarrow H_2 + OH^-$	1.0		
Phosphate ion	$PO_4^{3-} + H_2O \rightleftharpoons HPO_4^{2-} + OH^-$	$5.9 \times 10^{-3}$		



Base	Formula and Ionization Equation	$K_b$	Molecular Shape
Pyridine	$C_5H_5N+H_2O ightleftharpoons C_5H_5NH^++$	$OH^{-}$ 1.6 × 10 <sup>-9</sup>	

To find the pH we follow the same general procedure as in the case of a weak acid. If the stoichiometric concentration of the base is indicated by  $c_b$ , the result is entirely analogous to equation 4 in the section on the pH of weak acids; namely,

$$K_b = rac{[ ext{OH}^-]^2}{c_b - [ ext{OH}^-]}$$
 (14.5.3)

Under most circumstances we can make the approximation

$$c_b - [OH^-] pprox c_b$$

in which case Equation 14.5.3 reduces to the approximation

$$[OH^{-}] \approx \sqrt{K_b c_b} \tag{14.5.4}$$

which is identical to the expression obtained in the acid case (approximation shown in equation 6 in the section on the pH of weak acids) except that  $OH^-$  replaces  $H_3O^+$  and b replaces a. Once we have found the hydroxide-ion concentration from this approximation, we can then easily find the pOH, and from it the pH.

## Example 14.5.1: pH using K<sub>b</sub>

Using the value for  $K_b$  listed in the table, find the pH of 0.100 M NH<sub>3</sub>.

## **Solution**

It is not a bad idea to guess an approximate pH before embarking on the calculation. Since we have a dilute solution of a weak base, we expect the solution to be only mildly basic. A pH of 13 or 14 would be too basic, while a pH of 8 or 9 is too close to neutral. A pH of 10 or 11 seems reasonable. Using Equation 14.5.4 we have

$$egin{aligned} [ ext{ OH}^-] &= \sqrt{K_b c_b} \ &= \sqrt{1.8 \, imes 10^{-5} \, ext{mol L}^{-1} imes 0.100 \, ext{mol L}^{-1}} \ &= \sqrt{1.8 \, imes 10^{-6} \, ext{mol}^2 \, ext{L}^{-2}} \ &= 1.34 \, imes 10^{-3} \, ext{mol L}^{-1} \end{aligned}$$

Checking the accuracy of the approximation, we find

$$rac{[ ext{ OH}^-]}{c_{
m b}} = rac{1.34\, imes\,10^{-3}}{0.1} pprox 1 ext{ percent}$$

The approximation is valid, and we thus proceed to find the pOH.

$$\rm pOH = -log\frac{[~OH^-]}{mol~L^{-1}} = -log(1.34~\times~10^{-3}) = 2.87$$

From which

$$pH = 14.00 - pOH = 14.00 - 2.87 = 11.13$$

This calculated value checks well with our initial guess.

Occasionally we will find that the approximation

$$c_b - [OH^-] pprox c_b$$



is not valid, in which case we must use a series of successive approximations similar to that outlined above for acids. The appropriate formula can be derived from Equation 14.5.3 and reads

$$[OH^-]pprox \sqrt{K_b(c_b-[OH^-])}$$

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