

## NEUTRALIZATION CURVE DF WEAK ACID VS STRONG BASE

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## VEREIUYTRALIZATION CURVE OF WEAK ACID VS STRONG BASE

- The neutralization of 100mL of O.IM acetic acid (ethanoic acid) with O.IM sodium hydroxide solution will be considered here.
- **The pH** of the solution at the equivalence point is given by,

enturion

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 $(pH = \frac{1}{2} PKw + \frac{1}{2} PKa - \frac{1}{2} pc = 7 + 2.37 - \frac{1}{2} (1.3) = 8.72$ 

For other concentrations, we may employ the approximate Mass Action expression: [H+] x [CH3COO-]/[CH3COOH] = Kw

or, [H+] = [CH3COOH] x K a/[CH3COO-]

or pH = log[Salt]/[Acid] + <u>pKa</u>



concentration of the salt (and of the acid) at any point is calculated from the me of alkali added, due allowance being made for the total volume of the

- The initial pH of O.I M acetic acid is computed, the dissociation of the acid is relatively so small that it may be neglected in expressing the concentration of acetic acid.
- Hence from the above equation,

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[H+] x [CH3COO-]/[CH3COOH] = 1.82 x 10-5
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or [H+]2/0.1 = 1.82 x 10-5
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or [H+] = J1.82 x 10 6 = 1.35 X 10-3
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or pH = 2.87
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 $[Salt] = 50 \ge 0.1/150 = 3.33 \ge 10-2$ 

and [Acid] =  $50 \ge 0.1/150 = 3.33 \ge 10-2$ 

 $pH = log(3.33 \times 10-2/3.33 \times 10-2)+4.74 = 4.74$ 

- The pH values at other points on the titration curve are similarly calculated.
- After the equivalence point has been passed, the solution contains excess of OHions which will repress the hydrolysis of the salt; the pH may be assumed, due to the excess of base present, so that in this region the titration curve will almost coincide with thatfor 0.1M hydrochloric acid.





## **THANK YOU**