INDICATORS
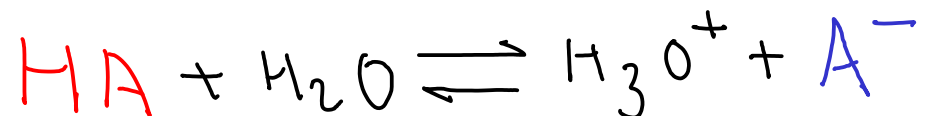
-Instead of using a pH meter to monitor acidity, we may choose to use an acid-base INDICATOR.

- Acid-base indicators are weak acids or weak bases which are highly colored.

- The color of the undissociated indicator MUST BE DIFFERENT than the color of the dissociated form!

$$HA + H_2O \rightleftharpoons H_3O^+ + A^-$$

The indicator must be present in very low concentrations - so that the indicator's equilibrium DOES NOT CONTROL the pH of the solution!

$$HA + H_2O \rightleftharpoons H_3O^+ + A^-$$
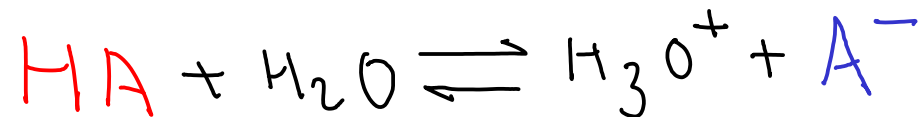
Look at the Henderson-Hasselbalch equation - we want to know how much of the red form and how much of the blue form are present!

$$pH = pK_{a,ind} + \log\left(\frac{[A^-]}{[HA]}\right)$$

When does the color of the indicator change?

IF the pH is << pKa, then the log term above must be both large AND negative!

 - What color is the solution?

$$[HA] >> [A^-]$$ ... so the SOLUTION IS RED!

If the pH is >> pKa, then the log term above must be both large AND positive!

 - What color is the solution?

$$[A^-] >> [HA]$$ ... so the SOLUTION IS BLUE!

- So, the color changes when the pH of the solution is near the pKa of the indicator, BUT we can only DETECT the change when enough of the other form is present.

Titration

- also called volumetric analysis. See the end of Ebbing chapter 4 for more details.

- frequently used to determine concentration of unknown acids or bases.

- typically react a basic sample with a STRONG ACID, or an acidic
sample with a STRONG BASE

Example:
Titrate 20 mL of vinegar (acetic acid) with 0.35 M NaOH. Let's study this titration.
What happens to the pH of the solution during the titration? How does an
indicator work?

0.35 M NaOH

Reaction:

$$NaOH + HC_2H_3O_2 \rightarrow NaC_2H_3O_2 + H_2O$$

20 mL
0.88 M $HC_2H_3O_2$
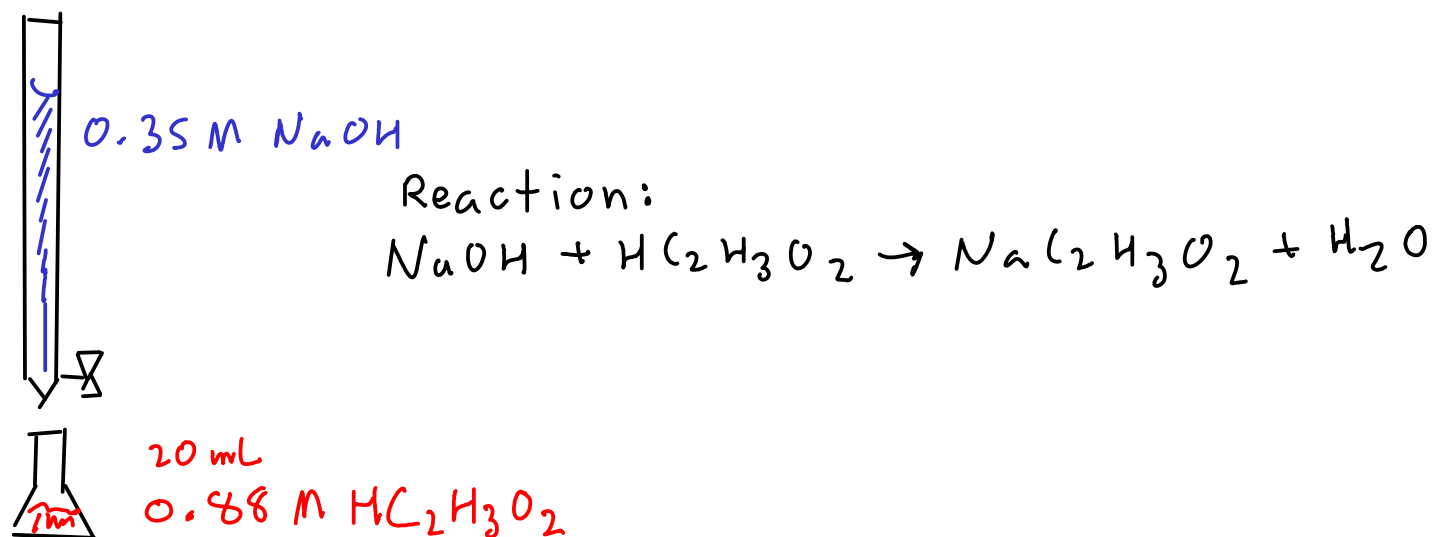
Vinegar is typically about 0.88M acetic acid. What would the EQUIVALENCE POINT (the point where we react away all of the acetic acid) be?

$NaOH + HC_2H_3O_2 \rightarrow Na C_2H_3O_2 + H_2O$

20.0 mL of 0.88M HC_2H_3O w/ 0.35 M NaOH

$20.0 \, mL \times \dfrac{0.88 \, mol}{L} = 17.6 \, mmol \; HC_2H_3O_2$

$17.6 \, mmol \; HC_2H_3O_2 \times \dfrac{mol \; NaOH}{mol \; HC_2H_3O_2} \times \dfrac{L}{0.35 \, mol \, NaOH} = \boxed{\begin{array}{l} 50.3 \, mL \\ of \; 0.35M \\ NaOH \end{array}}$

But how do we tell the titration is over if we don't already know the concentration of the acid?

In the lab, we have used phenolphthalein indicator for vinegar titrations. Phenolphthalein changes from colorless to pink over the range of about pH 9 to pH 10. How does this indicator show where the endpoint is?

Let's look at the pH of the solution during the titration- that may show us what's going on!

Titration curve for the titration of 20 mL of 0.88 M acetic acid with 0.35 M sodium hydroxide

pH = 9.08

Volume base added (mL)

buffer region: With a moderate amount of NaOH added, we have a solution that contains significant amounts of both acetic acid and its conjugate base (acetate ion). We have a buffer.

$$NaOH + HC_2H_3O_2 \rightarrow NaC_2H_3O_2 + H_2O$$

weak
acid

conjugate
base

The equivalence point:
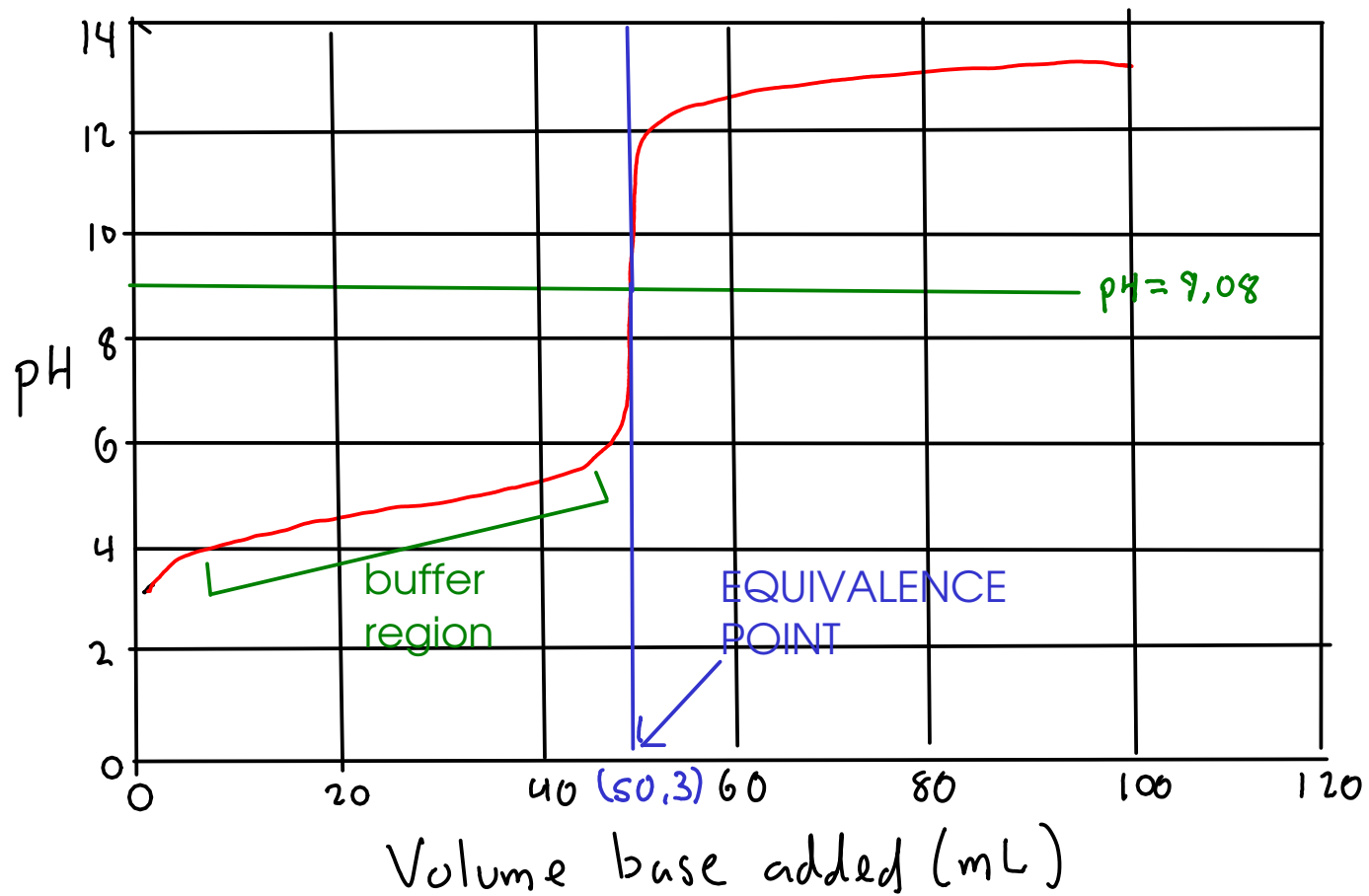
Equivalence point: We're reacting away more and more of the original acetic acid and converting it to acetate ion. At the equivalence point, all of the acetic acid has been converted, and we have only a solution of acetate ion.

Let's calculate the pH at the equivalence point.

$$NaOH + HC_2H_3O_2 \rightarrow NaC_2H_3O_2 + H_2O$$

20.0 mL of 0.88M HC_2H_3O w/ 0.35 M NaOH

$$20.0 \, mL \times \frac{0.88 \, mol}{L} = 17.6 \, mmol \; HC_2H_3O_2$$

$$17.6 \, mmol \; HC_2H_3O_2 \times \frac{mol \; NaOH}{mol \; HC_2H_3O_2} \times \frac{L}{0.35 \, mol \; NaOH} = \boxed{50.3 \, mL \; of \; 0.35M \; NaOH}$$

At the equivalence point, we have 17.6 mmol of ACETATE ION in 70.3 (20+50.3) mL of solution.

$$[C_2H_3O_2^-] = \frac{17.6}{70.3} = 0.250 \, M \; C_2H_3O_2^-$$

$$C_2H_3O_2^- + H_2O \rightleftharpoons OH^- + HC_2H_3O_2$$

	init	Δ	eqvil
$[C_2H_3O_2^-]$	0.250	$-x$	$0.250-x$
$[OH^-]$	0	$+x$	x
$[HC_2H_3O_2]$	0	$+x$	x

$$\frac{x^2}{0.250-x} = K_b$$

$$\frac{x^2}{0.250} = 5.88 \times 10^{-10}$$

$$Ka_{, HC_2H_3O_2} = 1.7 \times 10^{-5}$$

$$Kb_{, C_2H_3O_2^-} = 5.88 \times 10^{-10} \quad (K_a \times K_b = K_w)$$

$$x = 1.21 \times 10^{-5}, \quad pOH = 4.92, \quad \boxed{pH = 9.08}$$

# What about that phenolpthalein indicator?

Near the equivalence point, a very small volume of base added (a drop!) will change the pH from slightly over 6 to near 12. Since phenolphthalein changes colors at about pH 9-10, we can stop the titration within a drop of the equivalence point.

Another interesting point: The halfway point

pH (y-axis), Volume base added (mL) (x-axis)

phenolphthalein color change

pH = 9.08

buffer region

EQUIVALENCE POINT

(50,3)
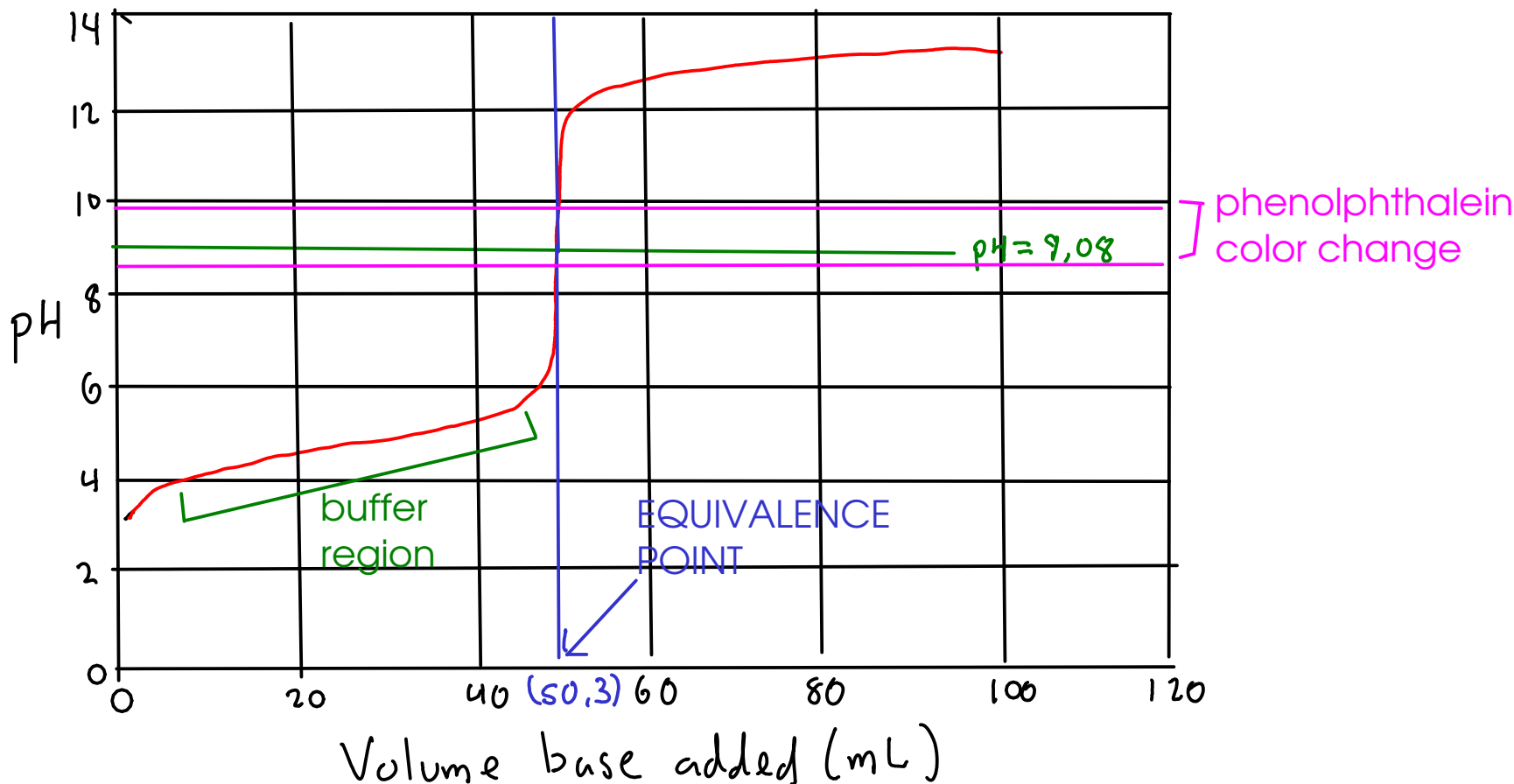
What's special about it? It's the point where we have added half the required acid to reach the equivalence point

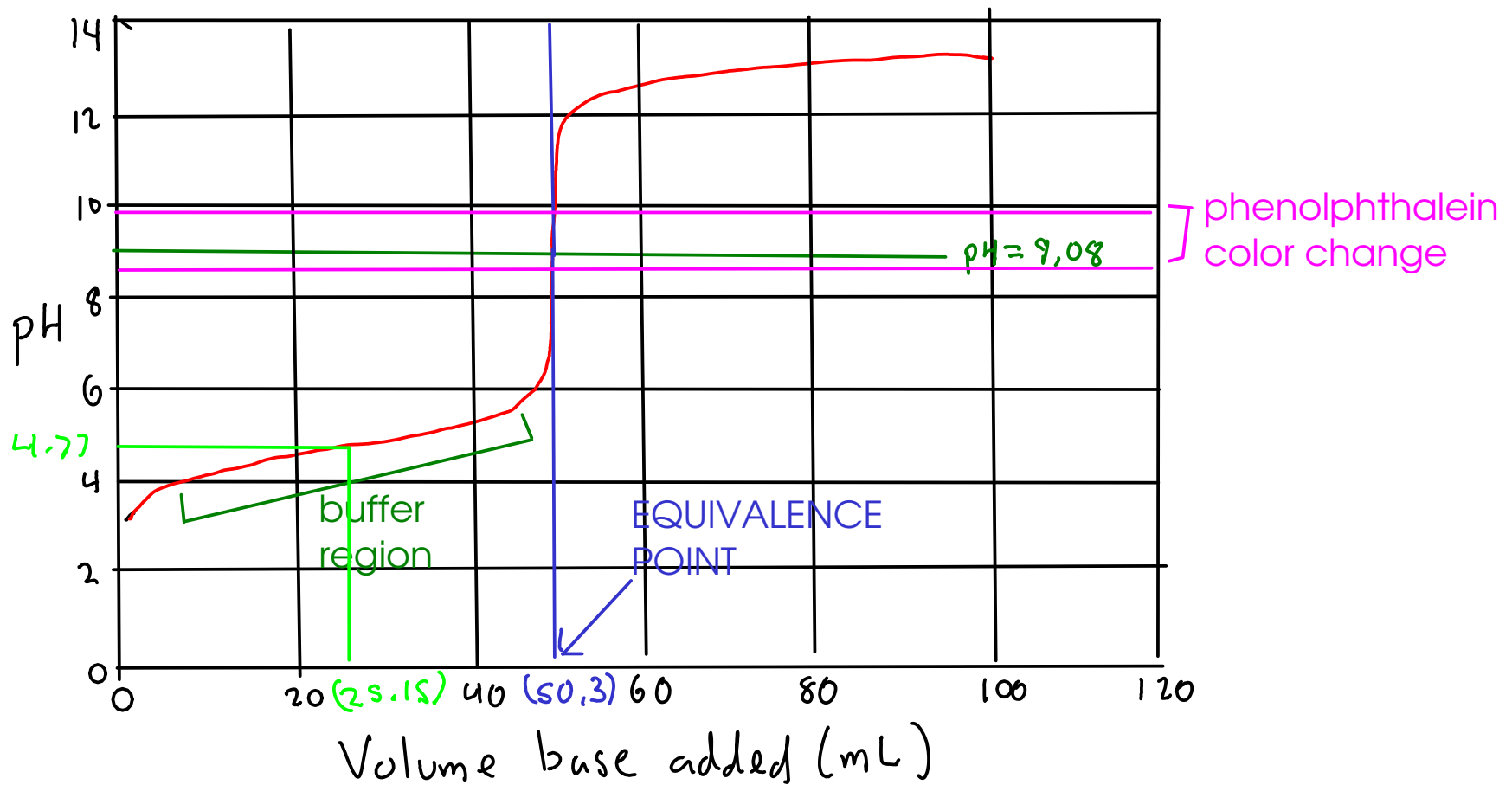$$NaOH + HC_2H_3O_2 \rightarrow NaC_2H_3O_2 + H_2O$$

17.6 mmol $HC_2H_3O_2$ initially

Add 25.15 mL base

$$25.15\ mL \times \frac{.350mol}{L} = 8.8025\ mmol\ NaOH$$

8.8 millimoles is also the amount of acid left, and the added base gets converted to acetate ion!

14
12
10
8
6
4.77
4
2
0

pH

O   20 (25.15) 40 (50,3) 60   80   100   120

pH = 9.08

phenolphthalein color change

buffer region

EQUIVALENCE POINT

Volume base added (mL)

The total volume is 45.15 mL, and both the acid and base are present at the same concentration. We have a BUFFER.

Find the pH of this buffer using the Henderson-Hasselbalch equation.

$$pH = pKa_{,HC_2H_3O_2} + \log\left(\frac{[C_2H_3O_2^-]}{[HC_2H_3O_2]}\right)$$

$= 0$, since the ratio $= 1$

At the halfway point, the pH = pKa of the acid!

Useful for finding acid ionization constants!