

CONCEPT: DIPROTIC ACID TITRATIONS PT.1

Equivalence Volume (V_e)

The titration of 100.0 mL of 0.100 M H_2SO_3 with 0.050 M KOH

Before any Strong Base is added

The titration of 100.0 mL of 0.100 M H_2SO_3 with 0.00 mL of 0.050 M KOH

	H_2SO_3	+	H_2O	\rightleftharpoons	HSO_3^-	+	H_3O^+
Initial	0.100 M				0.00 M		0.00 M
Change	- x				+ x		+ x
Equilibrium	0.100 - x				+ x		+ x

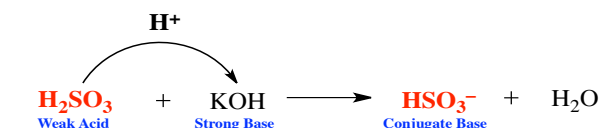
$$K_a = \frac{x^2}{[]_0 - x} \therefore x = [\text{H}^+] \therefore \text{pH} = -\log[\text{H}^+]$$

$$\frac{[]_0}{K_a} > 500 \quad \text{5\% Approximation Method}$$

Before 1st Equivalence Point

The titration of 100.0 mL of 0.100 M H_2SO_3 with 75.00 mL of 0.050 M KOH

	H_2SO_3 Weak Acid	+	KOH Strong Base	\longrightarrow	HSO_3^- Conjugate Base	+	H_2O
Initial	0.01000 moles		0.00375 moles		0.00000 moles		
Change	- 0.00375 moles		- 0.00375 moles		+ 0.00375 moles		
Final	0.00625 moles		0		0.00375 moles		



- **Weak acid** and **Conjugate base** will be present at the end.
- Use the Henderson Hasselbalch Equation.

$$\text{pH} = \text{pK}_a + \log\left(\frac{\text{Conjugate Base}}{\text{Weak Acid}}\right)$$

At 1st Equivalence Point

The titration of 100.0 mL of 0.100 M H_2SO_3 with 200.00 mL of 0.050 M KOH

	H_2SO_3 Weak Acid	+	KOH Strong Base	\longrightarrow	HSO_3^- Conjugate Base	+	H_2O
Initial	0.01000 moles		0.00100 moles		0.00000 moles		
Change	- 0.00100 moles		- 0.00100 moles		+ 0.00100 moles		
Final	0		0		0.00100 moles		

$$F = [\text{HA}]_0 \left(\frac{V_{\text{Initial Acid}}}{V_{\text{Solution}}} \right)$$

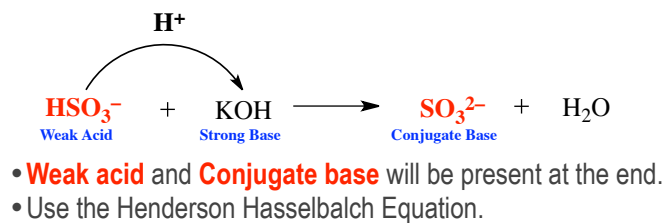
$$[\text{H}^+] = \sqrt{\frac{K_1 K_2 F + K_1 K_w}{K_1 + F}}$$

CONCEPT: DIPROTIC ACID TITRATIONS PT.2

After 1st Equivalence Point

The titration of 100.0 mL of 0.100 M H₂SO₃ with 250.00 mL of 0.050 M KOH

	HSO₃⁻ Weak Acid	+	KOH Strong Base	→	SO₃²⁻ Conjugate Base	+	H ₂ O
Initial	0.01000 moles		0.00250 moles		0.00000 moles		
Change	- 0.00250 moles		- 0.00250 moles		+ 0.00250 moles		
Final	0.00750 moles		0		0.00250 moles		



$$\text{pH} = \text{pK}_{a2} + \log\left(\frac{\text{Conjugate Base}}{\text{Weak Acid}}\right)$$

At 2nd Equivalence Point

The titration of 100.0 mL of 0.100 M H₂SO₃ with 400.00 mL of 0.050 M KOH

$$F = [\text{HA}]_0 \left(\frac{V_{\text{Initial Acid}}}{V_{\text{Solution}}} \right) \quad \left| \quad K_{b1} = \frac{x^2}{F - x} \therefore x = [\text{OH}^-] \therefore \text{pOH} = -\log[\text{OH}^-] \right.$$

$$\text{pH} = -\log[\text{H}^+]$$

After 2nd Equivalence Point

The titration of 100.0 mL of 0.100 M H₂SO₃ with 420.00 mL of 0.050 M KOH

$$[\text{OH}^-] = [\text{OH}^-]_0 \left(\frac{V_{\text{Excess Base}}}{V_{\text{Base}} + V_{\text{Acid}}} \right) \quad \left| \quad \text{pOH} = -\log[\text{OH}^-] \therefore \text{pH} = 14 - \text{pOH} \right.$$

PRACTICE: DIPROTIC ACID TITRATIONS CALCULATIONS 1

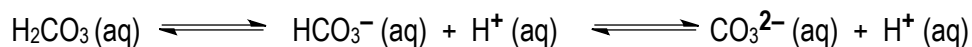
EXAMPLE 1: Calculate the pH of 100 mL of a 0.25 M H_2CO_3 when 70.0 mL of 0.25 M NaOH are added. $K_{a1} = 4.3 \times 10^{-7}$ and $K_{a2} = 5.6 \times 10^{-11}$.

EXAMPLE 2: Calculate the pH of 75.0 mL of a 0.10 M of phosphorous acid, H_3PO_3 , when 80.0 mL of 0.15 M NaOH are added. $K_{a1} = 5.0 \times 10^{-2}$, $K_{a2} = 2.0 \times 10^{-7}$.

PRACTICE: DIPROTIC TITRATIONS CALCULATIONS 2

EXAMPLE 1: Find the pH when 100.0 mL of a 0.1 M dibasic compound B ($pK_{b1} = 4.00$; $pK_{b2} = 8.00$) was titrated with 11 mL of a 1.00 M HCl.

EXAMPLE 2: Carbonic acid, H_2CO_3 , is a diprotic acid that dissociates, losing its two protons, to create bicarbonate, HCO_3^- , and carbonate, CO_3^{2-} , according to the following reactions given below:



As a diprotic acid system, it has two dissociation constants that $pK_{a1} = 6.30$ and $pK_{a2} = 10.30$ for the two steps. In the reaction you titrate 50.0 mL solution of 0.50 M H_2CO_3 with 1.00 M solution of NaOH. What would be the expected pH after the addition of 35 mL of the NaOH titrant?