

Partition coefficient K – Worksheet

Partition coefficients describe how a solute is distributed between two

immiscible solvents. Solubility of Solute in Organic Layer Partition coefficient $K = \frac{\text{Solubility of Solute in Organic Layer}}{\text{Solubility of Solute in Aqueous Layer}}$

- The partition coefficient (K_{pc}) is the ratio of the concentrations of a solute in two different immiscible solvents in contact with each other when equilibrium has been established (at a particular temperature)
- For example, methylamine (CH_3NH_2) is dissolved in two immiscible solvents: o Water o An organic solvent
- A separating funnel is shaken with the organic solvent and aqueous methylamine
- The methylamine is soluble in both solvents, so when the mixture is left to settle an equilibrium is established o The rate of methylamine molecules moving from the organic layer into the aqueous layer is equal to the rate of molecules moving from the aqueous layer to the organic layer

$\text{CH}_3\text{NH}_2(\text{aq}) \rightleftharpoons \text{CH}_3\text{NH}_2(\text{organic solvent})$

- The value of its equilibrium constant is also called the partition coefficient K_{pc}

Calculating Partition Coefficients

- The partition coefficient (K_{pc}) for a system in which the solute is in the same physical state in the two solvents can be calculated using the equilibrium expression

Worked example Calculating the partition coefficient

100 cm³ of a 0.150 mol dm⁻³ solution of aqueous methylamine (CH_3NH_2) was shaken with 75.0 cm³ of an organic solvent and left in the separating funnel to allow an equilibrium to be established. Only 50.0 cm³ of the aqueous layer was run off and titrated against 0.225 mol dm⁻³ hydrochloric acid (HCl) with an end-point of 14.1 cm³ of HCl. Calculate the partition coefficient of methylamine the organic solvent and water.

Answer

- Step 1: Write down the equilibrium equation: $\text{CH}_3\text{NH}_2(\text{aq}) \rightleftharpoons \text{CH}_3\text{NH}_2(\text{organic solvent})$
- Step 2: Write down the equilibrium expression: $K_{pc} = \frac{\text{CH}_3\text{NH}_2(\text{Organic solvent})}{\text{CH}_3\text{NH}_2(\text{aq})}$
- Step 3: Determine how many moles of CH_3NH_2 has reacted with HCl at the end-point: o At the end-point, all $\text{CH}_3\text{NH}_2(\text{aq})$ has been neutralised by HCl (aq) $\text{CH}_3\text{NH}_2(\text{aq}) + \text{HCl}(\text{aq}) \rightarrow \text{CH}_3\text{NH}_3\text{Cl}(\text{aq})$ o CH_3NH_2 and HCl react in a ratio of 1:1 o Mol (HCl) = mol (CH_3NH_2) = $0.225 \times 0.0141 = 0.00318 = 3.18 \times 10^{-3} \text{ mol}$
- Step 4: Determine the number of moles of CH_3NH_2 present in the aqueous layer o Only 50.0 cm³ of the aqueous layer was used to titrate against HCl o Thus, $3.18 \times 10^{-3} \text{ mol}$ of CH_3NH_2 was present in only 50.0 cm³ of the aqueous layer o The number of moles of CH_3NH_2 in 100 cm³ aqueous layer is, therefore: o Mol (CH_3NH_2 aqueous layer) = $3.18 \times 10^{-3} \times 2 = 6.34 \times 10^{-3} \text{ mol}$
- Step 5: Determine the number of moles of CH_3NH_2 in the organic layer: o Mol $\text{CH}_3\text{NH}_2(\text{organic layer}) = \text{mol } \text{CH}_3\text{NH}_2(\text{total}) - \text{mol } \text{CH}_3\text{NH}_2(\text{aqueous layer})$ o Mol $\text{CH}_3\text{NH}_2(\text{total}) = 0.100 \times 0.150 = 0.015 \text{ mol}$ o Mol $\text{CH}_3\text{NH}_2(\text{organic layer}) = 0.015 - 6.34 \times 10^{-3} = 8.67 \times 10^{-3} \text{ mol}$
- Step 6: Change the number of moles into concentrations: o Concentration (CH_3NH_2 in aqueous layer) = $6.34 \times 10^{-3} / 0.100 = 0.063 \text{ mol dm}^{-3}$ o Concentration (CH_3NH_2 in organic layer) = $8.67 \times 10^{-3} / 0.075 = 0.116 \text{ mol dm}^{-3}$
- Step 7: Substitute the values into the K_{pc} expression: o $K_{pc} = 0.116 / 0.063 = 1.83$ o Since the value of K_{pc} is larger than 1, methylamine is more soluble in the organic solvent than in water

Problem 1 1.0g of a compound dissolved in 100ml of water. If the Partition coefficient K is 5, how much of the compound will you extract by doing one extraction with 25ml of dichloromethane?

Set the compound you will extract as X g. If you start with 1g of compound, the amount of compound left in the aqueous layer at the end of one extraction will be (1-X)g.

Volume of aqueous = 100ml H₂O, Volume of organic = 25ml CH₂Cl₂ K = 5 = $\frac{X/25}{(1-X)/100}$

$X/25 \times 100 / (1-X) = 5$ $X = 0.55$ 100 1-X This means you extract 0.55g of the compound using 25ml of CH₂Cl₂ by doing only one extraction

Problem 2 1.0g of a compound dissolved in 100ml of water. If

the Partition coefficient K is 5, how much of the compound will you extract by doing one extraction with 50ml of dichloromethane? Set the compound you will extract as X g. If you start with 1g of compound, the amount of compound left in the aqueous layer at the end of one extraction will be $(1-X)$ g. Volume of aqueous = 100ml H_2O , Volume of organic = 50ml CH_2Cl_2 $K = 5 = \frac{X/50}{(1-X)/100}$ $X \cdot 50 \cdot K = 5 = X = 0.71$ $1 - X \cdot 100$ This means you extract 0.71g of the compound using 50ml of CH_2Cl_2 by doing only one large extraction

Factors Affecting the Partition Coefficient

- The partition coefficient (K_{pc}) depends on the solubilities of the solute in the two solvents
- The degree of solubility of a solute is determined by how strong the intermolecular bonds between solute and solvent are
- The strength of these intermolecular bonds, in turn, depends on the polarity of the solute and solvent molecules
- For example, ammonia is more soluble in water than in an organic solvent such as carbon tetrachloride (CCl_4)
- o Ammonia and water are both polar molecules that form hydrogen bonds with each other
- o Ammonia forms permanent dipole-induced dipole forces with the non-polar CCl_4 molecules
- o Since these forces are much weaker than hydrogen bonding, ammonia is less soluble in CCl_4
- When K_{pc} is 1 the solute is more soluble in water than the organic solvent
- When K_{pc} is 1 the solute is more soluble in the organic solvent than the water