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2009-J-12:

• Types of Intermolecular Forces

 $1s^2 2s^2 2p^6 3s^2 3p^5$

Cl

Marks • Complete the following table. 4 Formula Systematic name CaBr₂ calcium bromide potassium hydrogencarbonate KHCO₃ KMnO₄ potassium permanganate $Fe(NO_3)_3$ iron(III) nitrate • Consider the elements W, X, Y and Z from the same period, n, with the following 3 valence electron configurations: **Y** $ns^2 np^5$ **W** $ns^2 np^3$ $\mathbf{Z} ns^2 np^6$ **X** ns^2 Which element will conduct electricity in the solid state? X (alkaline earth) Which element will be the most electronegative? Y (halogen) Which element will possess the largest atomic radius? X (left hand side) Write the electronic configuration of lowest energy for the following species. 2 Na is given as an example. $1s^2 2s^2 2p^6 3s^1$ Na $1s^2 2s^2 2p^6$ Al^{3+}

٠	Complete the following table.

Marks	
10	

Species	Lewis structure	Arrangement of the electron pairs around the underlined atom	Geometry of species
<u>N</u> H ₃	H = N = H H H	tetrahedral	trigonal pyramidal
<u>S</u> F ₆	$ \begin{array}{c} \vdots F \\ F \\$	octahedral	octahedral
<u>B</u> F ₄ ⁻	- :F:::::::::::::::::::::::::::::::::::	tetrahedral	tetrahedral
<u>C</u> O ₂	∴o=c=o:	linear	linear
<u>I</u> Cl ₃	:ĊI: :ĊI: :ĊI:	trigonal bipyramidal	T-shaped

CI	HEM1001	2009-J-4	June 2009	22/01(a)
•	Manganese(II) chlor Manganese(II) carbo all insoluble. Descr manganese(II) chlor	ride and manganese(II) sulfate are bonate, manganese(II) hydroxide and ibe, using equations where appropriate into solid manganese(II) sulfate	ooth soluble in water. I manganese(II) phosphate are iate, how to convert solid 2.	Marks 4
	 Dissolve the manga MnCl₂(s) - 	nnese chloride in water. → Mn ²⁺ (aq) + 2CΓ(aq)		
	Add a solution of s • Mn ²⁺ (aq) +	odium carbonate. Manganese car CO₃ ^{2–} (aq) → MnCO₃(s)	bonate will precipitate.	
	 Filter off and wash MnCO₃(s) 	the precipitate and then dissolve + $2H^+(aq) \rightarrow Mn^{2+}(aq) + H_2O($	it in dilute sulfuric acid. l) + CO ₂ (g)	
	 Evaporate the solu Mn²⁺(aq) + 	tion to give manganese sulfate. $SO_4^{2-}(aq) \rightarrow MnSO_4(s)$		
•	Sodium chloride is s understanding of the	soluble in water, magnesium oxide e intermolecular forces involved, ex	is not. Using your plain why this is so.	3
	Both sodium chlor electrostatic forces required to overco	ide and magnesim oxide are ionic between the oppositely charged I me these attractions is called the I	compounds with strong particles. The energy lattice enthalpy.	
	When the ions diss the polar water mo solvation enthalpy.	olve in water strong bonds are for elecules. The energy released in th	rmed between the ions and is process is called the	
	If the solvation ent soluble.	halpy exceeds the lattice enthalpy	the compound will be	
	Sodium chloride is magnitude of ΔH (i	soluble because the magnitude of i):	ΔH (i) is less than the	
	(i) NaCl(s) \rightarrow Na	$a^+(g) + Cl^-(g)$		
	(ii) Na ⁺ (g) + Cl ⁻ ($\begin{array}{ccc} \mathbf{H}_2\mathbf{O} & & \mathbf{H}_2\mathbf{O} \\ \mathbf{g}) & \longrightarrow & \mathbf{N}\mathbf{a}^+(\mathbf{a}\mathbf{q}) + \mathbf{C}\Gamma(\mathbf{a}\mathbf{q}) \end{array}$		
	Magnesium oxide i magnitude of ΔH (i	s insoluble because the magnitud i):	e of ΔH (i) is greater than the	
	(i) MgO(s) \rightarrow M (ii) Mg ²⁺ (g) + O ²	$g^{2^+}(g) + O^{2^-}(g)$ $g^{-}(g) \xrightarrow{H_2O} Mg^{2^+}(aq) + OH^-(aq)$)	

Marks

5

• A solution is prepared by dissolving lead(II) nitrate (33.12 g) in 1.00 L of water. Write the balanced ionic equation for this dissolution reaction.

 $Pb(NO_3)_2(s) \rightarrow Pb^{2+}(aq) + 2NO_3(aq)$

When a 100.0 mL portion of this solution is mixed with a solution of potassium iodide (0.300 M, 150.0 mL), a bright yellow precipitate of lead(II) iodide forms. Write the balanced ionic equation for this precipitation reaction.

 $Pb^{2+}(aq) + 2I^{-}(aq) \rightarrow PbI_{2}(s)$

What mass of lead(II) iodide is formed?

The formula mass of Pb(NO₃)₂ is:

formula mass = $(207.2 \text{ (Pb)} + 2 \times 14.01 \text{ (N)} + 6 \times 16.00 \text{ (O)}) \text{ g mol}^{-1}$ = 331.22 g mol⁻¹

The number of moles in 33.12 g is therefore:

number of moles = $\frac{\text{mass}}{\text{formula mass}} = \frac{33.12 \text{ g}}{331.22 \text{ g mol}^{-1}} = 0.1000 \text{ mol}$

If this dissolved in 1.00 L and a 100.0 mL portion is taken, this will contain 0.01000 mol of $Pb^{2+}(aq)$.

150.0 mL of a 0.300 M solution of KI contains:

number of moles = concentration × volume = $0.300 \text{ mol } \text{L}^{-1} \times 0.1500 \text{ L} = 0.0450 \text{ mol}$

The precipitation reaction requires 2 mol of $\Gamma(aq)$ for every 1 mol of $Pb^{2+}(aq)$. The 0.01000 mol of $Pb^{2+}(aq)$ that is present requires 0.02000 mol of $\Gamma(aq)$. As there is more $\Gamma(aq)$ than this present, $\Gamma(aq)$ is in excess and $Pb^{2+}(aq)$ is the limiting reagent.

From the precipitation reaction, 1 mol of $Pb^{2+}(aq)$ will produce 1 mol of $PbI_2(s)$. Therefore 0.01000 mol of $Pb^{2+}(aq)$ will produce 0.01000 mol of $PbI_2(s)$.

The formula mass of PbI₂ is:

formula mass = $(207.2 \text{ (Pb)} + 2 \times 126.9 \text{ (I)}) \text{ g mol}^{-1}$ = 461.0 g mol⁻¹

The mass of 0.01000 mol is therefore:

mass = formula mass × number of moles = 461.0 g mol⁻¹ × 0.01000 mol = 4.61 g

Answer: **4.61 g**

What is the final concentration of $\Gamma(aq)$ ions remaining in solution after the reaction is complete?

As described above reaction of 0.01000 mol of $Pb^{2+}(aq)$ requires 0.02000 mol of Γ (aq). As 0.0450 mol are initially present, there are (0.0450 – 0.02000) mol = 0.0250 mol of $\Gamma(aq)$ after the precipitation reaction.

After mixing the two solutions, the total volume becomes (100.0 + 150.0) mL = 250.0 mL. The final concentration of $\Gamma(aq)$ is therefore:

concentration $=$	number of moles volume	$= \frac{0.0250 \text{ mol}}{0.2500 \text{ L}} = 0.100 \text{ mol } \text{L}^{-1} = 0.100 \text{ M}$
		Answer: 0.100 M

•	• Direct damage to the DNA of skin cells can be brought about by exposure to ultraviolet radiation of wavelength 300 nm. What are the frequency and energy (in kJ mol ⁻¹) of this radiation?				Marks 4
	The frequency, v, and the w Hence:	avelength, λ of t	he radia	tion are related by $c = v\lambda$.	
	$v = \frac{c}{\lambda} = \frac{2.998 \times 10^8 \text{ m}}{300 \times 10^{-9} \text{ m}}$	$\frac{s^{-1}}{n} = 1 \times 10^{15} \text{s}^{-1}$	$1 = 1 \times 1$	0 ¹⁵ Hz	
	The energy of the radiation	is given by $E = 1$	$hv=rac{hc}{\lambda}.$	Hence:	
	$E = \frac{hc}{\lambda} = \frac{\left(6.626 \times 10^{-34}\right)}{(30)}$	(z j s)(2.998 ×10 0 ×10 ^{−9} nm)	8 m s^{-1}	$) = 6.62 \times 10^{-19} \text{ J}$	
	This is the energy per photo	on. For a mole, t	he energ	y is:	
	$E = (6.62 \times 10^{-19} \text{ J}) \times (6.0)$	$22 \times 10^{23} \text{ mol}^{-1}$	= 400000) J mol ⁻¹ = 400 kJ mol ⁻¹	
Fre	equency: 1×10^{15} Hz	Ene	ergy: 400	kJ mol ⁻¹	
•	• Three different oxides of lead are known. The oxide that is red in colour is found to consist of 90.67 % lead. What is its empirical formula?			2	
	The oxide contains 90.67 %	Pb and so 9.33	% O.		
		Pb		0	
	percentage	90.67		9.33	
	divide by atomic mass	$\frac{90.67}{207.2} = 0.4$	438	$\frac{9.33}{16.00} = 0.583$	
	divide by smallest value	1		1.33	

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The ratio of Pb : O is 1 : 1.33 or 3: 4. The empirical formula is thus Pb₃O₄.

Answer: **Pb₃O₄**

•	Propane, C_3H_8 , is commonly used in barbecue gas cylinders, its complete combustion yielding water and carbon dioxide as the only products. What volume of CO_2 is produced at 0 °C and 1.0 atm from the complete combustion of 15.0 L of propane at a pressure of 4.5 atm and a temperature of 25 °C?	Marks 3
	Using the ideal gas law, $PV = nRT$, 15.0 L of propane at 25 °C (= 298 K) and 4.5 atm corresponds to:	
	$n = \frac{PV}{RT} = \frac{(4.5 \text{ atm})(15.0 \text{ L})}{(0.08206 \text{ L} \text{ atm} \text{ K}^{-1} \text{ mol}^{-1})(298 \text{ K})} = 2.76 \text{ mol}$	
	The combustion reaction is:	
	$C_3H_8(g) + 5O_2(g) \rightarrow 3CO_2(g) + 4H_2O(l)$	
	Hence, combustion of 2.76 mol of $C_3H_8(g)$ will give rise to (3 × 2.76) mol = 8.28 mol.	
	At 0 °C (= 273 K) and 1.0 atm, this amount will occupy:	
	$n = \frac{nRT}{n} = \frac{(8.28 \text{ mol})(0.08206 \text{ L atm } \text{K}^{-1} \text{ mol}^{-1})(273 \text{ K})}{1000 \text{ K}} = 190 \text{ L}$	
	<i>P</i> (1.0 atm)	
	Answer: 190 L	
•	The following cell has a potential of 0.55 V at 25 °C:	4
	$Pt(s) H_2(1.0 \text{ atm}) H^+(x M) Cl^-(1.0 M) Hg_2Cl_2(s) Hg(l)$	
	What is the concentration of H^+ in the anode compartment?	
	Data: $Hg_2Cl_2(s) + 2e^- \rightarrow 2Hg(l) + 2Cl^-(aq)$ $E^0 = 0.28 V$	
	The two half cells are:	
	$H_2(g) \rightarrow 2H^+(g) + 2e^- \qquad E^\circ = 0.00 V$ $Hg_2Cl_2(s) + 2e^- \rightarrow 2Hg(l) + 2Cl^-(aq) \qquad E^\circ = 0.28 V$	
	Hence, the overall reaction and the standard cell potential are:	
	$H_2(g) + Hg_2Cl_2(s) \rightarrow 2H^+(g) + 2Hg(l) + 2C\Gamma(aq)$ $E^0 = (0.00 + 0.28) V = 0.28 V.$	
	As $[H^+(aq)]$ is non-standard, the Nernst equation must be used to work out the cell potential:	
	$E_{\text{cell}} = E^{\circ} - 2.303 \times \frac{RT}{nF} \log Q$	
	ANSWER CONTINUES ON THE NEXT PAGE	J

The cell reaction is a 2 e⁻ process: n = 2. The reaction quotient only includes species that are gaseous or in solution:

$$Q = \frac{[H^{+}(aq)]^{2}][Cl^{-}(aq)]^{2}}{[H_{2}(g)]}$$

= [H⁺(aq)]² = x² since only [H⁺(aq)] is non-standard.
Hence, when $E_{cell} = 0.55$ V:
 $0.55 \text{ V} = (0.28 \text{ V}) - 2.303 \times \frac{(8.314 \text{ J K}^{-1} \text{ mol}^{-1})(298 \text{ K})}{(2 \times 96485 \text{ C mol}^{-1})} \log x^{2}$
 $x = [H^{+}(aq)] = 2.8 \times 10^{-5} \text{ M}$
Answer: $2.8 \times 10^{-5} \text{ M}$

Marks

3

• Adiponitrile, a key intermediate in the manufacture of nylon, is prepared by the reduction of acrylonitrile.

Anode: $2H_2O \rightarrow O_2 + 4H^+ + 4e^-$

Cathode: $2CH_2=CHCN + 2H^+ + 2e^- \rightarrow NC(CH_2)_4CN$

Write a balanced equation for the overall electrochemical reaction.

 $4CH_2=CHCN + 2H_2O \rightarrow 2NC(CH_2)_4CN + O_2$

What mass of adiponitrile (in kg) is produced in 10.0 hours in a cell that has a constant current of 3.00×10^3 A?

A current of 3.00×10^3 A passed for 10.0 hours corresponds to:

number of moles of electrons =
$$\frac{It}{F} = \frac{(3.00 \times 10^3 A)(10.0 \times 60 \times 60 s)}{(96485 C mol^{-1})}$$

= 1120 mol

From the cathode half cell reaction, 2 mol of electrons are needed to produce 1 mol of adiponitrile. Therefore, 1120 mol of electrons will produce 560. mol of this product.

The molar mass of NC(CH₂)₄CN is $(2 \times 14.01 \text{ (N)} + 6 \times 12.01 \text{ (C)} + 8 \times 1.008 \text{ (H)})$ g mol⁻¹ = 108.44 g mol⁻¹.

560 mol therefore corresponds to (560. mol) \times (108.44 g mol⁻¹) = 60500 g = 60.5 kg.

Answer: 60.5 kg

3

3

Marks • Write a balanced equation for the following reaction: $WO_3(s) + H_2(g) \rightarrow W(s) + H_2O(g)$ $WO_3(s) + 3H_2(g) \rightarrow W(s) + 3H_2O(g)$ What is the equilibrium constant expression, K_p , for the above reaction? The equilibrium constant only involves the gaseous reactants and products. The equilibrium constant in terms of partial pressures, K_p , is therefore: $K_{\rm p} = \frac{p^3({\rm H}_2{\rm O})}{n^3({\rm H}_2)}$ What is the equilibrium constant, K_c , for the above reaction, in terms of K_p ? The reaction involves no change in the moles of gas during the reaction: $\Delta n = 0$. As $K_p = K_c(RT)^{\Delta n}$, for this reaction $K_p = K_c$. • Fe₂O₃ can be reduced by carbon monoxide according to the following equation. $Fe_2O_3(s) + 3CO(g) \iff 2Fe(s) + 3CO_2(g)$ $K_{\rm p} = 19.9$ at 1000 K At 1000 K, what are the equilibrium partial pressures of CO and CO₂ if the only gas initially present is CO at a partial pressure of 0.978 atm? The reaction table is: $3CO_2(g)$ $Fe_2O_3(s)$ 3CO(g)~` 2Fe(s)0.978 0 initial / atm -change / atm -3x+3x0.978 - 3xequilibrium / atm 3xThe solids do not appear in the equilibrium constant expression and do not need to be considered. The equilibrium constant in terms of partial pressures, K_p , is given by: $K_{\rm p} = \frac{p({\rm CO}_2)^3}{p({\rm CO})^3} = \frac{(3x)^3}{(0.978 - 3x)^3} = 19.9$ Hence,

$$\frac{(3x)}{(0.978-3x)} = (19.9)^{1/3} = 2.71$$

3x = (2.71)(0.978 - 3x) = 2.65 - 8.13x or 11.1x = 2.65 or x = 0.238

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From the reaction table,	
p(CO) = (0.978 - 3x) atm = 0.264 atm	
$p(CO_2) = 3x \text{ atm} = 0.714 \text{ atm}$	
<i>p</i> (CO) = 0.264 atm	$p({\rm CO}_2) = 0.714 \text{ atm}$

Marks • Calculate the standard-free energy change for the oxidation of ammonia to nitric oxide 3 and water, according to the following equation. $4NH_3(g) + 5O_2(g) \rightarrow 4NO(g) + 6H_2O(1)$ Data: $\Delta_f G^{\circ}(NO(g)) = 87.6 \text{ kJ mol}^{-1}$ $\Delta_{\rm f} G^{\circ}(\rm NH_3(g)) = -16.5 \text{ kJ mol}^{-1}$ $\Delta_{\rm f}G^{\circ}({\rm H_2O(1)}) = -237.2 \text{ kJ mol}^{-1}$ Using $\Delta_{rxn}G^{\circ} = \Sigma m \Delta_f G^{\circ}$ (products) - $\Sigma n \Delta_f G^{\circ}$ (reactants), $\Delta_{rxn}G^{\circ} = ((4 \times 87.6 + 6 \times -237.2) - (4 \times -16.5)) \text{ kJ mol}^{-1} \text{ as } \Delta_{f}G^{\circ}(O_{2}(g)) = 0.$ $= -1007 \text{ kJ mol}^{-1}$ Answer: -1007 kJ mol⁻¹ Is the reaction spontaneous under standard conditions? Give a reason for your answer. The reaction is spontaneous as $\Delta_{rxn}G^{\circ} < 0$. This is the condition for a reaction to be spontaneous. How much heat is evolved, in kJ, when 5.00 g of Al reacts with a stoichiometric 1 amount of Fe₂O₃ according to the following equation? $2Al(s) + Fe_2O_3(s) \rightarrow 2Fe(s) + Al_2O_3(s)$ $\Lambda H^{0} = -852 \text{ kJ mol}^{-1}$ The atomic mass of Al is 26.98 g mol⁻¹ so 5.00 g corresponds to: number of moles of Al = $\frac{\text{mass}}{\text{atomic mass}} = \frac{5.00 \text{ g}}{26.98 \text{ g mol}^{-1}} = 0.185 \text{ mol}$ From the chemical equation, 852 kJ mol⁻¹ is evolved when 2 mol of Al reacts so 426 kJ is evolved when 1 mol of Al reacts. Therefore, when 0.185 mol of Al reacts, the heat evolved is (0.185 mol × 426 kJ mol^{-1}) = 78.9 kJ.

Answer: 78.9 kJ

Marks • The specific heat of Si is 0.71 J g^{-1} K⁻¹. How much heat is required to heat a Si wafer 1 weighing 0.45 g from 20.0 °C to 26.0 °C? The heat, q, required to increase a solid of mass m by ΔT is given by $q = mC_v\Delta T$ where $C_{\rm v}$ is the specific heat capacity. Hence: $q = (0.45 \text{ g}) \times (0.71 \text{ J g}^{-1} \text{ K}^{-1}) \times ((26.0 - 20.0) \text{ K}) = 1.9 \text{ J}$ Answer: 1.9 J 1 The structural formula of acetic acid is shown on the right. H = C = C'Acetic acid forms dimers (*i.e.* pairs of molecules) in the gas phase. Draw the dimer showing the H-bonding that occurs. H-bonding О-Н¹¹¹С-СН₃ 4 • Heating SbCl₅ causes it to decompose according to the following equation. $SbCl_{5}(g) \iff SbCl_{3}(g) + Cl_{2}(g)$ A sample of 0.50 mol of SbCl₅ is placed in a 1.0 L flask and heated to 450 °C. When the system reaches equilibrium there is $0.10 \text{ mol of } Cl_2 \text{ present.}$ Calculate the value of the equilibrium constant, K_c , at 450 °C. From the chemical equation, 1 mol of $Cl_2(g)$ is produced from every 1 mol of $SbCl_{5}(g)$ that reacts. As 0.10 mol of $Cl_{2}(g)$ is present at equilibrium, 0.10 mol of SbCl₅(g) has reacted. Initially, 0.50 mol of SbCl₅(g) was present so the amount left at equilibrium is (0.50 - 0.10) mol = 0.40 mol. From the chemical equation, 1 mol of SbCl₃(g) is produced from every 1 mol of $SbCl_5(g)$ that reacts. As 0.10 mol of $SbCl_5(g)$ has reacted, 0.10 mol of $SbCl_3(g)$ is present at equilibrium. $[SbCl_5(g)]_{eq} = \frac{number of moles}{volume} = \frac{0.40 \text{ mol}}{1.0 \text{ L}} = 0.40 \text{ mol } \text{L}^{-1} = 0.40 \text{ M}$ $[SbCl_3(g)]_{eq} = [Cl_2(g)]_{eq} = \frac{0.10 \text{ mol}}{1.0 \text{ L}} = 0.10 \text{ mol } \text{L}^{-1} = 0.10 \text{ M}$ The equilibrium constant in terms of concentrations, K_c is given by: $K_{\rm c} = \frac{[{\rm SbCl}_3({\rm g})][{\rm Cl}_2({\rm g})]}{[{\rm SbCl}_3({\rm g})]} = \frac{(0.1)(0.1)}{(0.40)} = 0.025$ Answer: $K_{c} = 0.025$

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•	Which of acetone, (CH ₃) ₂ CO, and water will have the greater surface tension. Why?	Marks 2
	Water will have the greater surface tension. It has much stronger intermolecular forces (H-bonds and dispersion) than acetone (dispersion and dipole-dipole forces).	
•	Melting points of the hydrogen halides increase in the order HCl < HBr < HF < HI. Explain this trend.	2
	The major intermolecular forces in HCl, HBr and HI are dispersion forces. The heavier the halogen is, the larger is its electron cloud and the more polarisable it is. Higher polarisability leads to stronger dispersion forces therefore leading to melting points which increase in the order HCl < HBr < HI.	
	F is a very small and very electronegative atom. The H–F bond is therefore highly polarised and H-bonds form. These are much stronger than dispersion forces and so HF has an anomalously high melting point. As seen by the experimental order given, this is enough to raise its melting point above that of HBr, but not above that of HI.	
•	Why is the solubility of chloroform (CHCl ₃) in water 10 times greater than that of carbon tetrachloride (CCl ₄) in water?	2
	The C–H bond in CHCl ₃ is quite polarised due to the electron-withdrawing effect of the Cl atoms. CHCl ₃ has a fairly large dipole moment with ô+ H and ô- Cl.	
	CCl ₄ has no dipole moment. Although each C-Cl bond is fairly polar, the tetrahedral shape of the molecule leads to overall cancellation of these bond dipoles.	
	The polar water molecules interact better with the polar CHCl ₃ molecules than with the non-polar CCl ₄ molecules so CHCl ₃ is more soluble.	