Acid bases Practice worksheet

| Α | hydrogen | sulfate/sulfate | system is re | enresented by | / the | following | equation |
|---------------|----------|-----------------|--------------|---------------|-------|------------|-----------|
| $\overline{}$ | Hydrogen | Sullate/Sullate | 373101111311 | spiesenieu bi | y uio | IOIIOWIIIQ | equation. |

$$\mathsf{HSO_4^-}(\mathsf{aq}) \quad + \quad \mathsf{H_2O}(\ell) \quad \leftrightarrows \quad \mathsf{SO_4^{2-}}(\mathsf{aq}) \quad + \quad \mathsf{H_3O^+}(\mathsf{aq})$$

- (a) Predict how
 - the forward reaction rate and
 - the pH

will differ from their original values after the following changes are imposed on the system and equilibrium has been re-established. Use the terms **increase**, **decrease**, **no change**. (6 marks)

| Change imposed by the addition of | Effect on forward reaction rate when equilibrium is re-established | Effect on pH when equilibrium is re-established |
|---|--|---|
| a few drops of concentrated hydrochloric acid | | |
| a few drops of concentrated lead(II) nitrate solution | | |
| distilled water | | |

| The reaction in part (a) is endothermic in the forward direction as written. Predict what will |
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| happen to the pH when the temperature is increased. Justify this prediction. (4 marks) |
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| ues | stion 29 (10 marks) | |
|------------|---|------------|
| a b 028 | peaker 12.00 mL of 0.0334 mol L ⁻¹ sulfuric acid solution, H ₂ SO ₄ (aq), is added to 3 mol L ⁻¹ potassium hydroxide solution, KOH(aq). | 2.50 mL of |
| 1) | Identify the limiting reagent in this reaction. Show all workings. | (5 marks) |
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|) | Calculate the final concentration of the excess reagent. Show all workings. | (3 marks |
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|) | Calculate the pH of the final solution. Show all workings. | (2 marks |
| | | |
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| vvaic | er is capable of s | seit-ionisation. | | |
|-------|---|--|---|--|
| (a) | Write an equa | ation for the self-ionisation of | water. | (2 ma |
| (b) | Write the equ | uilibrium constant expression | for the self-ionisation of \ | water. (1 m |
| (c) | | um constant for the self-ionisa ue indicate about this reactior | | x 10 ⁻¹⁴ at 25 °C. Wh (1 m |
| | | | | |
| | \(\text{values for the eratures.} \) | self-ionisation of water at 100 | | or a number of diffe |
| | | Temperature (°C) | K value | or a number of differ |
| | | Temperature (°C) | K value 0.114 x 10 ⁻¹⁴ | or a number of differ |
| | | Temperature (°C) | K value | or a number of differ |
| | | Temperature (°C) 0 25 | K value 0.114 x 10 ⁻¹⁴ 1.00 x 10 ⁻¹⁴ | or a number of differ |
| | | Temperature (°C) 0 25 50 | K value 0.114 x 10 ⁻¹⁴ 1.00 x 10 ⁻¹⁴ 5.48 x 10 ⁻¹⁴ | or a number of differ |
| | eratures. | Temperature (°C) 0 25 50 75 | K value 0.114 x 10 ⁻¹⁴ 1.00 x 10 ⁻¹⁴ 5.48 x 10 ⁻¹⁴ 19.9 x 10 ⁻¹⁴ | or a number of differ |

| (e) | Is water acidic, basic or neutral at 50 °C? State a reason for your answer. | (2 marks) |
|-------|---|----------------------------|
| | | |
| Que | estion 26 (1 | 0 marks) |
| Solid | d copper(II) hydroxide is added to excess 0.100 mol L ⁻¹ carbonic acid solution. | |
| (a) | Write the balanced equation, with appropriate state symbols, for the reaction that place between the copper(II) hydroxide and carbonic acid. | t takes (3 marks) |
| (b) | Predict all visible changes that would be observed, if any, while the reactants are together and afterwards. | e mixed (3 marks) |
| (c) | Predict two observations that would be different if excess 0.100 mol L ⁻¹ hydrowas used instead of the 0.100 mol L ⁻¹ carbonic acid. One: | ochloric acid (2 marks) |
| | Two: | |
| d) | State two personal safety measures the experimenter should take when conthese experiments. | ducting (2 marks) |
| | One: | |
| | Two: | |

| Phos | phoric acid, H ₃ PO ₄ (aq), is a weak, triprotic acid. | |
|------|--|------------------|
| (a) | Write the ionisation equation for phosphoric acid in water which shows the second proton of the acid being released into solution. | ond (2 marks) |
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| Mag | nesium carbonate, MgCO ₃ (s), is an ingredient of a commonly-used antacid. | |
| (b) | Other than water, list three species (elements, compounds, ions) that would be the reacting vessel open to the atmosphere at the completion of the reaction be excess solid magnesium carbonate and an aqueous solution of phosphoric acid | tween |
| | One: | |
| | Two: | |
| | Three: | |
| | ium hydroxide solution, NaOH(aq), was used in a titration to determine the concent sphoric acid. | tration of |
| (c) | Other than it having too low a molar mass, state two reasons why the concentrathe sodium hydroxide solution cannot be reliably determined by weighing out art of solid sodium hydroxide and dissolving it in a known volume of distilled water. | n amount |
| | One: | |
| | | |
| | Two: | |
| | | |

(12 marks)

Question 27

Question 29 (15 marks)

Wines and other alcoholic drinks can spoil when the alcohol (ethanol) they contain oxidises to acetic acid (ethanoic acid). An acidity regulator, monosodium citrate, is often added to drinks to prevent the formation of acetic acid. The monosodium citrate does this by acting as a buffer.

A citric acid/dihydrogen citrate ion buffer can be prepared from citric acid, $\rm H_3C_6H_5O_7$ and monosodium citrate, $\rm NaH_2C_6H_5O_7$.

| (a) | Write an equation for the buffer system $(H_3C_6H_5O_7/H_2C_6H_5O_7^-)$ containing cit $H_3C_6H_5O_7$ and monosodium citrate, $NaH_2C_6H_5O_7$. | citric acid, (2 marks) | |
|---------|---|---------------------------|--|
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| | | | |
| solutio | rs that contain equal concentrations of both components are most effective. The on is prepared by mixing 100.0 mL of citric acid solution with 100.0 mL of mone solution. The citric acid solution, $H_3C_6H_5O_7(aq)$, has a concentration of 0.200 | osodium | |
| (b) | Calculate the mass of sodium citrate, $NaH_2C_6H_5O_7$, that would need to be di 100.0 mL of distilled water to make the most effective buffer solution. | ssolved in (3 marks) | |
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| (c) | If a citric acid buffer was prepared to a pH of 3.5, what would b hydroxide ion at 25.0 °C? | | tration of the (3 marks) | |
|-----|---|------------------------|-----------------------------|--|
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| (d) | Explain why only a small change in pH is observed in this buffer solution when a small amount of sodium hydroxide solution is added, compared to adding a similar amount of sodium hydroxide solution to a system that is not a buffer solution. Your answer should refer to the buffer equilibrium in part (a). (4 marks) | | | |
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| 1. | . Consider the following data: | | | |
| | Barbituric acid $HC_4H_3N_2O_3$ Ka = 9.8 x 10^{-4} | ⁴ at 25°C | | |
| | If you were to put 2 .0 M of $HC_4H_3N_2O_3$ into pure distilled water | what would: | | |
| a) |) The pH of the solution be to 2 significant figures [3 marks , -1 for | r incorrect sig figs] | | |

| ۲) | Calculate the percentage ionication of Davbituvia acid to 2 significant figures [2 months 4 fee |
|-----|---|
| IJ) | Calculate the percentage ionisation of Barbituric acid, to 2 significant figures [2 marks , -1 for incorrect sig figs] |
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