



**PAMIBIA UNIVERSITY
OF SCIENCE AND TECHNOLOGY**

FACULTY OF HEALTH AND APPLIED SCIENCES

DEPARTMENT OF NATURAL AND APPLIED SCIENCES

QUALIFICATION: BACHELOR OF SCIENCE	
QUALIFICATION CODE: 07BOSC	LEVEL: 7
COURSE CODE: ACS701S	COURSE NAME: APPLIED COLLOID AND SURFACE CHEMISTRY
SESSION: JUNE 2019	PAPER: THEORY
DURATION: 3 HOURS	MARKS: 100

FIRST OPPORTUNITY EXAMINATION QUESTION PAPER	
EXAMINER(S)	Prof Habauka M. Kwaambwa
MODERATOR:	Prof Edet F. Archibong

INSTRUCTIONS
1. Answer ALL the questions. 2. Write clearly and neatly. 3. Number the answers clearly.

PERMISSIBLE MATERIALS

Non-programmable Calculators

ATTACHMENT

List of Useful Constants

THIS QUESTION PAPER CONSISTS OF 6 PAGES (Including this front page and List of Useful Constants)

QUESTION 1**[23]**

- (a) Colloids can be described in terms **size, dispersed phase/dispersion medium**, and **lyophilic or lyophobic** colloids. Write briefly about this statement. (5)
- (b) Surfactants are classified according to the type of the hydrophilic group. In many respects, the aqueous behaviour of dodecylbetaine is closer to octyl poly(oxyethylene) glycol (Triton X-100) than to hexadecyltrimethylammonium bromide. Explain or discuss this statement. (4)
- (c) Water treatment Moringa seed proteins are cationic. Arrange in increasing order of interaction of the following surfactants with Moringa seed proteins and explain briefly your answer:
Cetylpyridinium bromide; Polyoxyethylene alkyl ether, and; Sodium dodecyl sulphate.
Explain your answer. (4)
- (d) There is a variety of physical properties that can be used to determine the critical micelle concentration (CMC) of a surfactant such as sodium dodecyl sulphate (SDS). On the same diagram, show the variation of the following physical quantities with SDS concentration, showing clearly the position of the CMC: (4)
- (i) Osmotic pressure
 - (ii) Turbidity
 - (iii) Surface tension
- (e) Explain briefly the observed behaviours in (d) above. (6)

QUESTION 2**[27]**

- (a) Define the terms **solubilisation, Krafft temperature, T_K , and cloud point** as used in colloid chemistry. (6)
- (b) The cloud point for TX-100 was studied as function temperature. What would you observe if the same experiment was done using sodium dodecyl sulphate (SDS)? (2)
- (c) Using well-labelled schematic diagrams illustrate how (i) solubilisation varies with surfactant concentration, and; (ii) how solubility of surfactants with temperature indicating clearly the position T_K and critical micelle concentration. (7)
- (d) State whether the critical micelle concentration (CMC) would **increase, decrease or not change** after the following changes: (5)
- (i) Changing the surfactant from $\text{CH}_3(\text{CH}_2)_9(\text{OCH}_2\text{CH}_2)_5\text{OH}$ to $\text{CH}_3(\text{CH}_2)_7(\text{OCH}_2\text{CH}_2)_5\text{OH}$
 - (ii) Increasing the temperature
 - (iii) Addition of electrolyte to an ionic surfactant
 - (iv) Presence of impurity when CMC is determined by surface tension
 - (v) Branching of the hydrophobic part of the surfactant

(e) A Moringa shower gel formula has the following components:

Identification	Component
A	Water
B	SDS 30%
C	Coconut diethanolamide
D	Alkyl amido propyl betaine 30%
E	Cocoamine oxide
F	NaCl
G	Perfume, colour, preservative
H	Lactic acid
J	Moringa seed oil

Match each of the following functions to corresponding letter (A-J) of the components in the table above: (7)

- (i) Increases thickness by causing the surfactant to restructure into the high viscosity cylindrical micelle structures
- (ii) Anionic surfactant
- (iii) Thought to be a component of the skin and used to the pH compatible with that of the skin, i.e. adjust pH to 6.5
- (iv) Antiaging and antifungal component
- (v) Amphoteric surfactant to generate foam or cold water detergent
- (vi) Nonionic surfactant that imparts excellent viscosity enhancing and foam stabilisation in anionic based systems
- (vii) Foam booster surfactant

QUESTION 3

[12]

(a) Outline any **three** main assumptions involved in the derivation of the BET adsorption isotherm equation for molecules at the gas/liquid interface. (3)

(b) The linear BET equation is of the form:

$$\frac{p}{V(p_0 - p)} = \frac{1}{V_m c} + \frac{(c - 1) p}{V_m c p_0}$$

(i) State what each of the quantities in this equation represents. (4)

(ii) A graphical plot of $\frac{p}{V(p_0 - p)}$ against $\frac{p}{p_0}$ data for the adsorption of nitrogen gas on 1 g of sample of alumina at 77 K gave a slope of $2.88 \times 10^{-2} \text{ cm}^{-3} \text{ (s.t.p.)}$ and an intercept of $9.87 \times 10^{-4} \text{ cm}^{-3} \text{ (s.t.p.)}$. Calculate the specific surface area (m^2g^{-1}) for the alumina sample, taking the molecular area of nitrogen as $16.2 \times 10^{-20} \text{ m}^2$. (5)

QUESTION 4**[20]**

- (a) Compare and contrast the following terms as used in colloid stability: (6)
- (i) Sedimentation and Creaming
 - (ii) Depletion flocculation and Bridging flocculation
- (b) Explain how the following factors would affect the stability of colloidal dispersions. (6)
- (i) Brownian motion
 - (ii) Increase in particle size of colloidal particles
 - (iii) Decrease in medium viscosity
- (c) Define the terms **point of zero charge** and **potential determining ions**. As a Colloid Scientist, use/apply these concepts to explain why AgI particles are negatively charged and how you can manipulate them so that you have a dispersion with zero charged AgI particles and another with positively charged AgI particles. (6)
- (d) Apart from the above mechanism in (c), **isomorphous substitution** is another mechanism particles acquire charge. Deduce the resulting charge of clay particles if metal X (valency = 4⁺) replaces metal M (valency = 3⁺)? (2)

QUESTION 5**[18]**

- (a) One form of the van Waals interaction potential between two particles is given by:
- $$V_A(h) = -\frac{Aa}{12h},$$
- (i) State **any** two conditions under which this equation is valid. (2)
 - (ii) Briefly state the effect on V_A if the particles are immersed in medium instead particles instead of particles in a vacuo? (1)
 - (iii) What effect on V_A is observed if the Hamaker constant of the medium approaches that of the particles? (1)
- (b) Using combining relations based on the Hamaker constants of pure materials (A_i), calculate the composite Hamaker constants for the following interacting systems: (6)
- (i) Polystyrene in water; (ii) SiO₂ in water, and; (iii) Polystyrene-Water-SiO₂ (6)
- Comment on the results with respect colloid stability of the systems. (3)

Given:

Material	$A_i \times 10^{-20} \text{ J}$
Polystyrene	7.2
SiO ₂	0.8
Water	4.1

- (c) On the same well-labelled diagram, show schematically the variation of the van der Waals attraction potential (V_A), electrostatic potential (V_R) and total pair potential ($V_T = V_A + V_R$) with the interparticle separation, h , for a **marginally stable** dispersion of nanoparticles indicating clearly the positions, if any, of primary minimum, primary maximum, secondary minimum and Born repulsion potential (V_B). (5)

END OF EXAM QUESTIONS

USEFUL CONSTANTS:

Universal Gas constant	R	=	8.314 J K ⁻¹ mol ⁻¹
Boltzmann's constant,	k	=	1.381 x 10 ⁻²³ J K ⁻¹
Planck's constant	h	=	6.626 x 10 ⁻³⁴ J s
Debye-Huckel's constant,	A	=	0.509 (mol dm ⁻³) ^{1/2} or mol ^{-0.5} kg ^{0.5}
Faraday's constant	F	=	96485 C mol ⁻¹
Mass of electron	m _e	=	9.109 x 10 ⁻³¹ kg
Velocity of light	c	=	2.998 x 10 ⁸ m s ⁻¹
Avogadro's constant	N _A	=	6.022 x 10 ²³
1 electron volt (eV)		=	1.602 x 10 ⁻¹⁹ J
