

UNIVERSITY OF SWAZILAND

Faculty of Health Science

Department of Environmental Health Sciences

Final Examination 2009

Title of paper: Instrumental Methods for Environmental Analysis 1

Course code: EHS 573

Time allowed: 2 hours

Marks allocation: 100 Marks

Instructions:

- 1) Read the questions and instructions carefully
- 2) Answer ANY FOUR (4) questions
- 3) Each question is weighted 25 marks
- 4) Write neatly and clearly
- 5) Begin each question on a separate sheet of paper

This paper is not to be opened until the invigilator has granted permission

Question 1(25 marks)

- (a) State the factors that would guide you in selecting an appropriate method for the analysis of an environmental sample. [6]
- (b) Explain the term 'sampling' of an environmental sample for analysis. What steps should be taken to ensure that an appropriate sampling has been carried out?

 [4]
- (c) Why is sample pretreatment usually a necessary step prior to the analysis of the sample? Give four of the usual pre-treatment steps often employed during analysis of environmental samples. [5]
- (d) Using a labeled diagram, identify the basic components of a typical instrument for chemical analysis. State the functions of any two of the components and give an example in a named instrument. [10]

Question 2 (25 marks)

- (a) Explain the term 'deviation from Beer's law'. Using a graphical illustration, distinguish between positive and negative deviation from Beer's law. [4]
- (b) Briefly discuss the causes and the possible corrections of true (real), deviation from Bee's law. [5]
- (c) The combined absorbance, A_c , when a beam of radiation made of two wavelengths λ and λ^1 , with molar absorbances of ϵ and ϵ^1 respectively, pass through an absorbing solution is given by:

$$A_c = log(P_o + P_o^1) - log(P_o 10^{-\epsilon bc} + P_o^1 10^{-\epsilon' bc})$$

- (i) Assuming Beer's law holds, obtain this expression. [4]
- (ii) What type of deviation from Beer's law (if any), occurs when:

$$\epsilon = \epsilon^1, \quad \epsilon > \epsilon^1 \text{ and } \epsilon < \epsilon^1 ?$$

- (d) Stray radiations have been identified as one of the instrumental causes of deviation from Beer's law during spectroscopic analysis:
 - (i) What are the characteristics of these radiations? [4]
 - (ii) Give the expression for the measured absorbance, A_m , due to them and state the type of deviation they cause i.e. positive or negative. [2]

Que	estion 3 (25 marks)						
For	the Electrothermal Atomic Absorption Spectrophotometer, EAAS, (Graphite	furnace)					
a)	Draw and label its schematic diagram.	[3]					
b)	Give four of its advantages and three if its disadvantages over the flame AAS. [7]						
c)	Discuss the stages involved in the atomization process of a sample.						
d)	Account for the uses of 'the flowing inert gas' and the Matrix modifier' du analysis involving the use of this method.	ıring [6]					
Que	estion 4 (25 marks)						
a)	For the following spectrophotometric methods: FAAS, FAES, AFS, ICP, ICP-AFS i) Classify them into two groups as 'emission' or 'absorption' method. ii) State the quantity measured in each case.	[5]					
b)	Using the 'Spectronic 20' as a typical example of a single beam spectrophotometer i) Draw and label the schematic diagram of its optical train. ii) State the material used for the source of radiation, the wavelength disp medium, and the detector iii) Why is it referred to as 'a single beam instrument'?	[6] ersing [3] [1]					
c)	Give a brief description of the working principles of the FAES (flame aton emission) spectrometry.	nic [10]					
Qu	estion 5(25 marks)						
a)	Give at least one difference in the instrumental design for the following a spectroscopic methods of analysis. i) AAS and FES; ii) AAS and AFS; iii) AFS and FES.	atomic [5]					
b)	Which is more sensitive to flame temperature stability, AAS or FES and	•					
c)	Give five advantages of ICP (Inductively Coupled Plasma) spectroscopic of analysis over the other conventional spectroscopic methods.						
d)	Briefly describe the working principles of ICP.	[5] [10]					

Question 6	25	marks)	۱
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- a) What is meant by "interference" with regards to flames and furnaces? [2]
- b) For the 'chemic4', 'ionization' and 'spectral' types of interferences.
 - (i) Explain their causes
 - (ii) Discuss the steps normally taken to correct or eliminate each of them.
 - (iii) Discuss the steps usually taken to correct or eliminate 'ionization interference.

[15]

- c) (i) Give two examples each of a 'cool flame' and a hot flame'
 - (ii) Offer an explanation for the following observation. "Although chemical interferences are more prevailent in 'cool' flames, yet this flame is preferred for the analysis of alkali metals"

[8]

PERIODIC TABLE OF ELEMENTS

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() indicates the mass number of the isotope with the longest half-life.

Quantity	Symbol	Value	General data and							
Speed of light	c	2.997 924 58 × 10 ⁸ m s ⁻¹	fundamental							
Elementary charge		7.602.177 x 10-19 C	constants-							
Faraday constant	$F = eN_{\lambda}$	9.6485 x 10° C mol ⁻¹								
Boltzmann constant	k	1.380 66 × 10 ⁻²³ J K ⁻¹	· •							
Gas constant	$R = kN_A$	8.31451 J K ⁻¹ mol ⁻¹	. •							
		8.205 78 × 10 ⁻² dm³ atm K ⁻¹	mol ⁻¹							
		62.364 L Torr K ⁻¹ mol ⁻¹								
Planck constant	t h	6.626 08 × 10 ⁻³⁴ J s								
:	$h = h/2\pi$	$1.054^{\circ}57 \times 10^{-34} \text{ J s}$								
Avogadro .	N;	6.022 14 × 10 ²³ mol ⁻¹								
constant										
Atomic mass unit	u ·	1.660 54 × 10 ⁻²⁷ kg								
Mass of	*		•							
electron	m•	$9.10939 \times 10^{-31} \text{ kg}$	•							
proton	m ₂ .	$1.672-62 \times 10^{-27} \text{ kg}$								
neutron	m,	1.674 93 \times 10 ⁻²⁷ kg $^{\circ}$								
Vacuum ∵: permeability		$4\pi \times 10^{-7} \text{ J s}^2 \text{ C}^{-2} \text{ m}^{-1}$								
	, .	$4\pi \times 10^{-7} \mathrm{T}^2 \mathrm{J}^{-1} \mathrm{m}^3$: - 							
Yacuum :. permittivity	$\varepsilon_0 = 1/c^2 \mu_0$	$8.854 19 \times 10^{-12} \text{ J}^{-1} \text{ C}^2 \text{ r}$								
• • • •	4πε ₀	1.112 65 × 10 ⁻¹⁰ J ⁻¹ C ² i	n−1							
Bohr magneto	$n \mu_2 = e \hbar/2m_*$	9.274 02 × 10 ⁻²⁴ J T ⁻³	• • •							
Nuclear magneton	$\mu_{N} = e fi/2m_{p}$	$5.05079 \times 10^{-27} \text{J} \text{T}^{-1}$								
Electron g	5.	2.002 32.								
value		,								
Bohr radius	$a_0 = 4\pi \varepsilon_0 \hat{n}^2/\pi$	1, £ 5.291 77 × 10 ⁻¹¹ m	-							
Rydberg constant	R_ = m,e*/8h	3 c: 1.097 37 × 10 5 cm $^{-1}$								
Fine structure constant	$c = \mu_0 e^2 c/2h$	7.29735×10^{-3}								
Gravitational constant	G	$6.67259 \times 10^{-11} \text{ N m}^2 \text{ k}$	$6.67259 \times 10^{-11} \text{ N m}^2 \text{ kg}^{-2}$							
Standard (. g	2.806.65_n_s_ ⁷								
of free fall†			. † Exact (defined) values							
f P	η μ	nedk M	g Prefixes							
femto pico	nano micro m	illi centi deci kilo meg	a giga							
10-15 10-1)-3 10 ⁻² 10 ⁻¹ 10 ³ 10 ⁶								