2913/106
APPLIED SCIENCE AND MICROBIOLOGY
PRACTICAL
Oct./Nov. 2019
Time: 4 hours



THE KENYA NATIONAL EXAMINATIONS COUNCIL

DIPLOMA IN FOOD SCIENCE AND PROCESSING TECHNOLOGY

MODULE I

APPLIED SCIENCE AND MICROBIOLOGY PRACTICAL

4 hours

INSTRUCTIONS TO CANDIDATES

You should have the following for this examination:
Answer booklet;

Non-programmable scientific calculator.

This paper consists of THREE practical tests.

Carry out ALL the practical tests and answer ALL the questions in the answer booklet provided.

Maximum marks for each part of the practical test are as indicated.

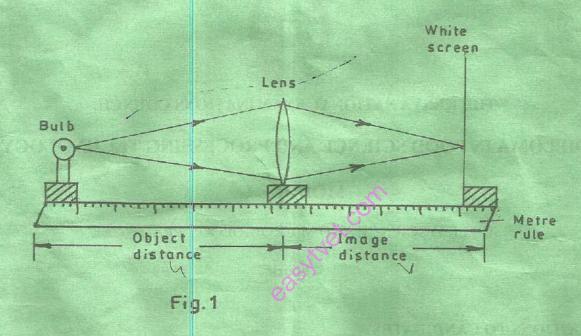
This paper consists of 4 printed pages.

Candidates should check the question paper to ascertain that all the pages are printed as indicated and that no questions are missing.

1. You are required to determine the focal length of a lens using magnification (M) method.

Apparatus

- A 3 Volt bulb and a bulb holder.
- A convex lens with lens holder.
- A meter rule.
- A white screen with a support.
- A pair of dry cells (size D) with a cell holder.
- Connecting wires.



Procedure

- (a) Set up the apparatus as shown in Figure 1.
- (b) With the bulb on, set the object distance U = 25 cm and move the screen to and fro until a sharp image of the object is formed on the screen. Measure the image distance (V cm) and record in table 1.
- (c) Repeat step (b) for the other values of U by adjusting lens and complete table 1.

(14 marks)

Object distance U cm	25	30	40	50	60	70	80
Image distance V cm	5c浇	lucm.	Bin	13cm	Born	lacon	pem
$m = \frac{V}{U}$	or Gum	6.46mi	0:3cm	0,26,	012162	0.171	0.15
M		4-611	73:3	2.6	2:16	1.41	1.5

2913/106 Oct./Nov. 2019

(d) Plot a graph of m against V in cm on the graph paper provided. Determine the gradient of the graph. 3- 51 (e) (4 marks, Determine the value of V in cm on the graph when m = 0.00. (f) (2 weeks) Calculate the value of $f = \frac{1}{gradient}$. (g) (2 marks) (h) Calculate the difference between value of f and V obtained in (g) and (f) above. (2 marks) (a) You are provided with the following. (i) Analar grade sodium carbonate. Dilute hydrochloric acid solution. (ii) (iii) Sodium hydroxide solution. (iv) Methyl orange indicator. (v) Titration apparatus. Naot Naco3 (b) You are required to: prepare a standard solution of 0.2 M sodium carbonate; (i) standardise dilute hydrochloric acid solution; (ii) (iii) determine the molarity of sodium hydroxide solution. Proceed as follows: (c) calculate the mass of sodium carbonate required to prepare 250 cm³ of (i) 0.2 M sodium carbonate (Na = 23, C = 12, O = 16). weigh accurately the mass of sodium carbonate calculated in (i) above and (ii) record the exact mass weighed. (1 mark) Transfer completely the mass weighed into a 250 cm³ volumetric flask, (iv) add 100 cm3 of distilled water, shake until all sodium carbonate dissolves and top up with distilled water to the 250 cm³ mark. Calculate the exact molarity of the solution you have prepared. (v) (4 marks) (d) (i) Fill a burette with HCl solution.

Pipette 25 cm³ of sodium carbonate solution prepared into a clean conical flask (ii) and add three drops of methyl orange indicator.

Applied s in a microproced

Turn over

2.

LIW = 1000M Titrate the sodium carbonate solution with HCl until a permanent colour change occurs. Repeat the titration two more times recording the volume of the titre. (4 marks) (iv) Calculate the molarity of the dilute HCl solution. (6 marks) Measure 200 cm³ of sodium hydroxide provided and dilute to the mark in a (i) 1 litre volumetric flask with distilled water, and set aside. 1000cm (ii) Fill a burette with HCl solution. Transfer 25 cm3 of sodium hydroxide in (i) above into a conical flask, add (iii) three drops of methyl orange indicator. (iv) Titrate the sodium hydroxide solution with HCl to a permanent colour change. Repeat the fitration two more times recording the titre volume. (4 marks) (v) Calculate the molarity of the sodium hydroxide solution. (9 marks) 3. You are provided with the following. (a) - Two micro-organism labelled X and Y growing on nutrient agar (NA) plates. - 2 microscope slides. - 2 cover slips. - 1 wire loop. - Distilled water in a wash bottle. - Crystal violet solution. - Gram's iodine. - Ethanol/Acetone - Safranin. 1/-- Blotting paper. - Bunsen burner. - Immersion oil. - Staining tray. - Microscope with X10, X40 and X100 objectives. Examine the cultures and describe the colony characteristics of the two microorganisms (b) provided. (12 marks) Make a smear for each micro-organism and stain using the Gram stain method. Write (c) the procedure for Gram staining. (15 marks) Describe the appearance of the micro organisms upon staining. (d) (6 marks) Applied surner & micro princt THIS IS THE LAST PRINTED PAGE 2913/106 Oct./Nov. 2019