

# **Technical University of Mombasa**

## **Faculty of Applied and Health Sciences**

### DEPARTMENT OF PURE AND APPLIED SCIENCES UNIVERSITY EXAMINATION FOR THE DEGREE OF BACHELOR OF TECHNOLOGY IN INDUSTRIAL MICROBIOLOGY AND BIOTECHNOLOGY BIMBT 10M

# SBT 2442: GENETIC ENGINEERING I

SPECIAL/SUPPLEMENTARY EXAMINATION

FEBRUARY 2013 SERIES	2 HOURS
Instructions to candidates:	

This paper consist of **FIVE** guestions Answer guestion **ONE** (compulsory) and any other **TWO** guestions

#### **Question ONE**

	)		
		i) Sieving	(2marks)
		ii) Chimera DNA	(2marks)
		iii) Palmdromic sequence	(2marks)
1	b)	Differentiate between southern and northern blotting	(4marks)
(	c)	List the FOUR stages of DNA separation from cellular components	(4marks)
	d)	Name TWO reasons for the limited range of applications for crude lysates	as DNA source ( <b>4marks</b> )
(	e)	Name TWO types of cohesive ends	(2marks)
]	f)	State how an enzymes activity is estimated	(2marks)
:	g)	List any FOUR types of cloning vectors in order of the DNA size each	can pack (star

a) Define the following terms :

with the smallest)

- h) State any TWO reasons for amplifying DNA by polymerase chain reaction (PCR) (2marks)
- i) Name TWO uses that polyacrylamide gel electrophoresis (PAGE) is suitable for (2marks)

#### Question TWO

a)	Describe the classical procedure of gene cloning	(12marks)
b)	Describe DNA isolation by salting-out method	(8marks)

#### **Question THREE**

a)	Highlight the main distinctions between restriction enzymes I, II and III	(7marks)
b)	Describe the principle of conventional and real time PCR technology	(13marks)

#### **Question FOUR**

a)	Describe restriction mapping procedure	(10marks)
b)	Describe DNA spacing by addition of homopolymers	(10marks)

#### **Question FIVE**

a)	Plasmids are suitable as cloning vectors. Explain	(10marks)
b)	Describe the principle of nucleic acid hybridization	(10marks)