## DR. BABASAHEB AMBEDKAR TECHNOLOGICAL UNIVERSITY LONERE - RAIGAD - 402 103

Winter Semester Examination - January - 2023

Subjec Subjec Maxin	Course: B. Pharm. Subject: Biostatistics & Research Methodology Subject Code: BP801T Maximum Marks: 75 Date: 10.02.2023  Semester: Vision Seme	
Instru	ctions: i) All questions are compulsory ii) Figures to the right indicate full marks iii) Draw the diagrams or flow charts wherever necessary.	
Q. 1.	Objective Type Questions (Answer all the questions)	$(10 \times 2) = 20$
i)	Differentiate between Null and alternative hypothesis.	
i)	Enumerate Statistical Software's.	
ii)	Enlist Advantages and disadvantages Pie charts.	
iv)	Classify observational studies.	
v)	Give two methods of sample size calculation in research study	·.
vi)	Write Applications of SAS or MINITAB.	
vii)	Enlist non-parametric tests.	
viii)	Define research and enlist its types.	
ix)	Write significance of sampling in research.	
<b>x</b> )	Define bias in clinical study.	
Q. 2.	Long Answers (Answer 2 out of 3)	$(2 \times 10) = 20$
i) ii) iii) Q. 3. i)	What is hypothesis? What are different types of hypothesis? E you will formulate a hypothesis with a suitable example. Explain with suitable examples regression analysis and standaregression.  What are measures of dispersion? Explain their significance we examples.  Short Answers (Answer 7 out of 9)  What is Plagiarism? What are the needs for research?	ard error of
ii)	Explain a typical experimental study design.	
iii)	Write a short note on central composite design.	
iv) v)	What do you meant by factorial design? Write some advantage factorial design.  Discuss null hypothesis, type I and type II errors.	ges of
vi)	Explain ANOVA and its applications.	
vii) viii) viii)	Discuss the applications of EXCEL and SPSS programmes in analysis  Describe the various graphical methods of representing quant	
ix)	What are non-parametric tests? Explain chi square test-Good test.	

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		End Semester Examination - Summer 2023	
	The second secon	e: B. Pharmacy Semester: Subject Code: Aarks: 75	VIII BP802T 3 Hrs
	Q. 1. i) ii)	Enlist the names of disease that can be protected using Ervebo vaccine, Dengvaxia Vaccine, ShanChol vaccine and PCV13 vaccine  Differentiate between community medicine and preventive	(10 x 2) = 20
•	iii) iv) v) vi) vii) viii) ix)	Define prevention and control State different functions of Sodium and Fluoride as a minerals	
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	x)	A / Sydney / 05 / 97 (H3N2)    Ose in the sydney in the sy	
•	Q. 2. i) ii) iii)	Long Answers (Answer 2 out of 3) State and explain different levels of prevention and control State the objective of National Malaria Prevention Programme Explain different Malaria control strategies Explain different vitamin deficiency disorders and their prevention	(2 x 10) = 20
	Q. 3. i) ii) iii) iv) v) vi) vii) viii) ix)	Short Answers (Answer 7 out of 9) What is personal hygiene? How to maintain personal hygiene? What is SARS write its symptoms, prevention and control Explain different functions of PHC in health care system Write about National Leprosy Control Programme Discuss health promotion and education in school Explain the objectives of National Tobacco Control Programme Elaborate the role of WHO in Indian national health program Explain levels of prevention Highlight the effect of urbanization on Health	$(7 \times 5) = 35$

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## DR. BABASAHEB AMBEDKAR TECHNOLOGICAL UNIVERSITY, LONERE **End Semester Examination – Summer 2023** Date: 13/06/2023 VIII Course Sem: B. Pharmacy BP803ET Subject Name: **Pharma Marketimg Management** Subject Code: 3 Hr. Duration Max Marks 75 **Instructions:** All questions are compulsory Draw diagrams / figures wherever necessary Figures to right indicate full marks $(10 \times 2) = 20$ Objective Type Questions (Answer all the questions) Q. 1. Define needs and demands. i) Write the significance of product modification. ii) Write the importance of medical exhibition. iii) State the objectives of promotion. iv) Differentiate between marketing and selling. V) List out functions of C and F agents. vi) What do you mean by vertical and horizontal marketing? vii) Define Branding and Channel Conflict. viii) Write differences between the Consumer Buying and Industrial Buying. ix) What is the role of NPPA? x) $(2 \times 10) = 20$ Long Answers (Answer 2 out of 3) Q. 2. Define marketing. Explain its nature and scope of pharmaceutical marketing. i) What is market segmentation? Explain the different ways of segmenting the ii) pharmaceutical market with examples. Enumerate the various channels of distribution in pharmaceutical marketing. iii) How will you select the most appropriate channel for distribution of drugs? $(7 \times 5) = 35$ Short Answers (Answer 7 out of 9) Q. 3. Describe the different strategies of pricing. i) Explain importance of communication in detailing. ii) Suggest some online promotion techniques for OTP products. iii) ii) Global marketing Write short notes on i) Rural marketing iv) Write an overview on DPCO. v) Explain product branding. vi)

	vii)						
	viii) Discuss in detail role of retailers in distribution of pharmaceutical product  ix) Justify the role of professional sales representative. Highlight the process of selection and training of the sales representatives in pharma organization.						
		x) Justify the role of professional sales representative. Highlight the process of selection and training of the sales representatives in pharma organization.					
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## DR. BABASAHEB AMBEDKAR TECHNOLOGICAL UNIVERSITY, LONERE

## End Semester Examination - Summer 2023

Date: Course **B.** Pharmacy Sem: VIII Subject Code :BP804ET Subject Name: Pharmaceutical Regulatory Science : 3 Hrs Max Marks Duration Instructions: 1. All questions are compulsory 2. Draw diagrams / figures wherever necessary 3. Figures to right indicate full marks  $(10 \times 2) = 20$ Objective Type Questions (Answer all the questions) Q. 1. Differentiate innovative drug and generic drug. i) Name the regulatory authorities of India, US, EU and Australia. ii) Write down the objectives of CDSCO. iii) Enlist modules of CTD. iv) Give Constitution of IRB. v) Give examples for brand and the respective generic products. vi) Write role of regulatory affairs in pharmaceutical industry. vii) Enlist the benefits of eCTD. viii) Draw ACTD triangle. ix) Difference between NDA and ANDA. X)  $(2 \times 10) = 20$ Long Answers (Answer 2 out of 3) Q. 2. Describe in detail the stages of drug discovery process. Explain the organization and functions of CDSCO. ii) Write in detail about clinical trial protocol and explain how Pharmacovigilance as iii) safety monitoring in clinical trials?  $(7 \times 5) = 35$ Short Answers (Answer 7 out of 9) Q. 3. What is DMF? Explain the contents of DMF. i) Write the importance of 21 CFR. ii) What are the salient features of purple book? iii) Define clinical trial and explain role of Investigator. iv) What is ethics behind informed consent process? v) Explain orange book. vi) Write different modules of ACTD. vii) Explain regulatory approval process of IND. viii)

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Describe stages in development of generic formulations.

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