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(SEM VIII) THEORY EXAMINATION 2021-22 Visit - pharmacyindia.co.in PHARMACOVIGILANCE

Time: 3 Hours Total Marks: 75

Note: Attempt all Sections. If require any missing data; then choose suitably.

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SECTION A

1.	Attempt all questions in brief.	$10 \times 2 = 20$
a.	What is drug event monitoring?	
b.	List four drugs contraindicated in pregnant and lactat	ing women.
c.	Suggest the requirements for CIOMS form	
d.	What is eudravigilance?	https://pharmacyindia.co.in/
e.	What is teratogenicity? Give examples.	
f.	What is post marketing safety?	
g.	Narrate the minimum criteria required for a valid repo	ort?
h.	What is phase IV of clinical trials?	
i.	What are the basic objectives of pharmacovigilance p	olanning?
i.	Mention few examples of predictable adverse drug re	eactions.

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4.	Attempt any two parts of the following: \sqrt{f}	,
a.	Classify adverse drug reaction. How will you deject and report ADR along with	ŀ
	causality assessment scales?	
b.	Discuss in detail basic and trectalized drug information resources in	l
	pharmacovigilance	l
c.	Discuss in detail of Cohort and Cocontrol study. Explain the applications of	l
	MedDRA and standard Med Pragueries.	l

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SECTION C

3.	Attempt any five parts of the following: $7 \times 5 = 35$	
a.	Explain the establishing pharmacovigilance program in the bospital.	
b.	What is vaccine safety surveillance? Explain detail different types of	
	pharmacovigilance methods used for passive and active surveillance.	
c.	Discuss in detail establishment and operation of drug safety department in	
	pharmaceutical industry. https://pharmacyindia.co.in/	
d.	Mention the importance aspects of ICH guidennes for expedited reporting.	
e.	Write short note on pharmacogenomics and adverse drug reaction	
f.	How will you carry out drug safety evaluation in geriatric and pediatric populations?	
g.	What is the role of CDSCO in pharmacovigilance? Write a note on ATC classification	
	of drugs. https://pharmacyindia.co.in/	

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