

ADMISSION FORM, 2023-2024

PERSONAL DETAILS				
Name				
Address				
Contact No				
Email ID				
Birth Date	DD/MM/YYYY			
Gender				
PROFESSIONAL BACKGROUND/EXPERIENCE				
Company's Name				
Company's Address				
Department		Designation		
Yr. of Exp.				
EDUCATION BACKGROUND				
Qualification Degree	Name of the College/Institute	Year of Passing	% of Marks	Special/Optional Subject

FEES PAYMENT DETAILS

<b>BANK NAME</b>	AXIS BANK
<b>ACCOUNT NAME</b>	RAAJ GPRAC PRIVATE LIMITED
<b>ACCOUNT NUMBER</b>	919020005392967
<b>IFSC</b>	UTIB0000571 <b>SWIFT CODE:</b> AXISINBB571
<b>BRANCH</b>	Hiranandani Estate, Patlipada, Ghodbundar Road, Thane West, Mumbai, India
<b>GPAY NO</b>	9821144706

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Please select the course of your choice, Mark (√) in respective column			
COURSE TITLE		DURATION	MARK (√)
<b>Sr.No</b>	<b>PHARMACEUTICALS</b>		
1	Advanced course in GLOBAL REGULATORY AFFAIRS [GRA-API, Formulations, Medical Devices, Biosimilars, OTC, USFDA, EU, ROW, Canada Drug Registration, CTD-ECTD, ACTD, NEES Submission, GMP-GCP-GLP guidance, Drug development, Product Life Cycle Management, ICH Guidelines Q1-Q14]	6 months	
2	REGULATORY AFFAIRS-Formulation	4 months	
3	REGULATORY AFFAIRS-API[DMF/ASMF/CEP]	3 months	
4	QA-QM [ Quality Assurance and Quality Management]	3 months	
5	Regulatory Submissions as per CTD-eCTD, ACTD, NeeS	3 months	
6	Advanced course in Clinical Research [CR]	3 months	
7	Advanced course in Pharmaceutical Documentation [PD]	3 months	
8	Advanced course in Pharmacovigilance [PV]	3 months	
9	Advanced course in Patents Laws and Procedures [IPR]	12 months	
10	Advanced course in Pharmaceutical Management [PM]	12 months	
11	Advanced course in Medical Writing [MW]	3 months	
12	Effective CMC (Chemistry, Manufacturing and Controls) Writing and Review Skills	7 days	
	<b>NUTRACEUTICALS</b>		
13	Advance Course in Nutraceuticals	3 months	
14	QA in Nutraceuticals/Herbal/Cosmetics	3 months	
	<b>MEDICAL DEVICE</b>		
15	GLOBAL Medical Devices (MDR) course [India, Europe, USA, ASEAN, ROW regulations, Design and Development, Clinical Evaluation, Quality Management system, Medical Device Single Audit Program (MDSAP), Risk Management System, ]	6 months	
16	Medical Device-Quality Assurance [Introduction to various medical device regulatory bodies-India, US,EU, ISO 13485-2016-Quality Management System, Risk Management & Risk Analysis]	3 months	
17	Internal Auditor/Lead Auditor- Quality Management System (QMS)- ISO 13485:2016, Auditing Management System -ISO 19011:2018	10 Days	
18	Medical Device Directive to Medical Device Regulation (MDD to MDR) Implementation	7 Days	
19	Implementation of Medical Device Single Audit Program (MDSAP), An Overview- Australia, Brazil, Canada, Japan and the United States.	7 Days	
20	Technical File Documentation for Medical Device [Technical file preparation, medical device classification, documentation design, biocompatibility, Unique design identification, clinical evaluation, general safety and performance report (GSPR)]	7 Days	
	<b>Cosmetics</b>		
21	QA/RA in Cosmetics (Introduction to Cosmetics, Technology in Cosmetic Industry, Product Development and Cosmetic Formulation, Good Manufacturing Practices for Cosmetic, Quality Assurance and Quality Control, Global Regulation for Cosmetics (USFDA, India, EU), Supply Chain Management, Cosmatovigilance)	3 months	

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Contact Details:

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For more information visit: [www.raajpharmaelearning.com](http://www.raajpharmaelearning.com), [www.raajgprac.com](http://www.raajgprac.com)