CAPRA ASOCIATION of Professionals in Regulatory Affairs Association canadienne des professionnels en réglementation

MEMBERSHIP FORM 2017/2018

(Membership Year: Aug.1/17 to July 31/18)

NOTE: Where possible, Membership Applications and Renewals should be done by way of the website at www.capra.ca. If this is not possible, then please use this form. If you change companies or wish to change any other information in your file during the membership year, be sure to update your contact information in your file at www.capra.ca, to continue uninterrupted service.

Please fill out both pages completely and print clearly.

How many years of regulatory affairs experience do you have?

Dr Mr Ms N	Miss Mrs	NEW Member?	Renewal?	_ Honorary* Mem	nber?
Name:					
Job Title:					
Organization:					
Mailing Address:					
Street/ Suite					
Town/City					
Province/Country					
Postal Code					
Telephone:					
Fax:					
E-Mail Address:					
	(Required, as most co	orrespondence is ser	nt via e-mail)		
Date:	Signature:				
*Honorary Members are appointed	I d by the Board of Director	s and pay no fee.			
Please answer the following q	uestions so we may kr	now something abou	t you:		
Do you wish to have your membership information accessible for other members to see?			Yes	No	
Do you wish to volunteer to y	work on a Committee?			Yes	No

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More than 20

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NAME:				
What type of <u>company</u> are you <u>currently</u> em Trade / Professional Organization / Law Industry – Biotech / small pharma / Cal Industry - Large Pharma / biotech / sub Contract Organization (laboratory, mar Academic Institution (College / University Other (Please specify)	□ Government / legal □ Industry - Cosmetics □ Industry - Medical Device □ Industry - NHP □ Consultancy			
What types of companies where previously Trade / Professional Organization / Law Industry – Biotech / small pharma / Cal Industry - Large Pharma / biotech / sub Contract Organization (laboratory, mar Academic Institution (College / University Other (Please specify)	Firm nadian Sponsor company sidiary of global organization sufacturing, clinical)	□ Government / legal □ Industry - Cosmetics □ Industry - Medical Device □ Industry - NHP □ Consultancy		
What types of <u>products</u> are you responsible ☐ Biologics/Biotechnology ☐ Veterinary Medicines ☐ Natural Health Products (NHPs) / Cons ☐ Innovative Pharmaceuticals / Drugs	 ☐ Medical Devices ☐ Combination Products ☐ Cosmetics ☐ Generic Pharmaceuticals / Drugs 			
What types of <u>activities</u> are you responsible □ Business/Strategy/Management □ Submissions/Registrations - Pre-market □ Advertising and Promotion □ Teaching / Training □ Research and Development, including development (Chemistry, Manufacturing)	□ Qualit st □ Subm □ Gover Preclinical (pharmacology / toxicology)	uality Assurance /Quality Control ubmissions/Registrations - Post-market overnment Affairs / Policy / Trade ogy); Clinical trials; pharmaceutical		
What jurisdictions are you responsible for (o ☐ Canada ☐ USA ☐ Europe (EU) ☐ Non-EU Europe ☐ Worldwide	check all that apply; if worldwide, check Asia Australia/New Zealand Africa Latin America	conly worldwide)		
Are you a member of any of these other Or ☐ RAPS ☐ TOPRA ☐ DIA	ganizations: CSPS SQA Other (Please specify)			
Memberships are subject to approval by the Board of Directors. Memberships will only be processed with payment.				

Please allow TWO WEEKS for processing.

Mail this form with payment of \$55.00 CDA (or equivalent) to:

CAPRA/ACPR Membership 2425 Matheson Blvd E., Suite 795 Mississauga ON L4W 5K4 Canada

Make payment payable to CAPRA/ACPR. HST is already included in the \$55.00 (HST # 85475 8349).

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