



SwAPP

Swiss Association of Pharmaceutical Professionals

CH-3000 Bern

Summary of Theoretical Training for *Specialty Diploma Regulatory Affairs*

Last name/First name/Title: _____ Date of Birth (D/M/Y): _____ Sheet No.: _____

No	Event Titel of Event Place/Organiser/Date	Duration (hrs) in each Specialty Area - Please fill in the specialty areas in the grey boxes below. Use one column per specialty area.														
		Clinic aspects drug development	CH regulatory submissions	CH regulations and law	EU regulatory submissions of drugs & biologics	US regulatory submissions of drugs & biologics	Variations & change control	Reimbursement & pricing CH	Pharmacovigilance	Good manufacturing practice	Good distribution practice	Promotion pharmaceutical products in CH	Special regulations: generics, biologics etc.	Relevant literature and publications	Trainings in any field of regulatory affairs	Other fields pharmaceutical medicine
Match with attachments		8	24	16	8	8	8	8	8	4	4	4	4	8	30	18
Total hours per specialty area:																

Add sheets as needed, add up total hours on last sheet. Attach a certificate for each training.

Place and Date

Signature of Applicant: