

ARCS Professional Development Pathways for Regulatory Affairs Professionals



Professional Development for the Therapeutics Industry™

	Introductory/Entry Level	Intermediate (upto 3 years)	Advanced/ Extra Curricular
Workshops			
Introduction to Australian Regulatory Affairs (3 days)	✓		
Overview of Drug Development (1 day)	✓		
Requirements of Quality – Module 3 and Module 2.3 (2 days)	✓	✓*	
Bioavailability An Bioequivalence – Understanding and Applying Bioavailability and Bioequivalence (BABE) Guidelines (2 days)		✓	
EU Regulatory Affairs – Pharmaceutical (1day)			✓
USA Regulatory Affairs – Pharmaceutical (1day)			✓
Foundations in Biostatistics for the Biotechnology and Pharmaceutical Industry (2 days)		✓	✓
An Introduction to Project Management for the Therapeutic Industry (1 day)	✓	✓	
Project Management Fundamentals for Therapeutic Goods Development (3 days)		✓	✓
An Introduction to Pharmacovigilance Requirements (0.5 day)		✓	
Business Writing Skills (1day)	✓	✓	✓
Advanced Business Writing (1 day)			✓
The Effective Manager (2 days)			✓
Increasing your Effectiveness at Work (1 day)		✓	✓
Managing Conflict Situations (1 day)		✓	✓
Assertive Negotiation (1 day)		✓	✓
TGA Overview Day (1 day)	✓		
TGA Industry Update Day (1 day)		✓	✓
E-learning			
Some Simple Techniques to Make your Next Presentation More Effective		✓	
Preparing Submission in the Common Technical Document (CTD) Format	✓	✓	
Electronic Common Technical Document (eCTD)		✓	✓
Other ARCS Professional Development Opportunities			
Evening Update Series ^a	✓	✓	✓
Volunteer pathway	✓	✓	✓
ARCS / Industry Events ^a	✓	✓	✓

^a There are a number of key forums applicable to all levels of experience where ARCS members can remain current on new/changing industry wide initiatives (go to the ARCS website for further information).

* Refer course description