Master's degree programme "Drug Regulatory Affairs"
Study plan: MDRA-27
Academic year 2025/2026

(subject to change, 13 Sept 2025)

Module 1 Definition and Scope of Drug Regulatory Affairs and Good Regulatory Practice

Wednesday	17 September 2025 from 12:00 to 18:30
Thursday	18 September 2025 from 8:00 to 18:30
Friday	19 September 2025 from 8:00 to 18:30
Saturday	20 September 2025 from 8:00 to 16:00

Pharmaceutical Institute, Bonn-Endenich Submission of study paper: 20/Oct/2025

Module 2 Pharmaceutical Law

Part 1:	Friday	10 October 2025 from 8:30 to 18:30
	Saturday	11 October 2025 from 8:00 to 16:00
Part 2:	Friday	24 October 2025 from 8:30 to 18:30
	Saturday	25 October 2025 from 8:00 to 16:00

Part 1 Online, Part 2 Online

Module 3 International Registration Procedures

Part 1:	Friday	07 November 2025 from 8:30 to 18:30
	Saturday	08 November 2025 from 8:00 to 16:00
Part 2:	Friday	21 November 2025 from 8:30 to 18:30
	Saturday	22 November 2025 from 8:00 to 16:00
Part 3:	Friday	12 December 2025 from 8:30 to 18:30
	Saturday	13 December 2025 from 8:00 to 16:00

Part 1 Online, Part 2 Online, Part 3 Online

Module 4 General Aspects of Module 1 (CTD), Registration of Special Medicinal Products

Part 1:	Friday	09 January 2026 from 8:30 to 18:30
	Saturday	10 January 2026 from 8:00 to 16:00
Part 2:	Friday	23 January 2026 from 8:30 to 18:30
	Saturday	24 January 2026 from 8:00 to 16:00

Part 1 Online, Part 2 Online Submission of study paper: 23/Feb/2026

Module 5 Maintenance of Marketing Authorisations / Pharmacovigilance

Part 1:	Friday _PV	06 February 2026 from 8:30 to 18:30
	Saturday _PV	07 February 2026 from 8:00 to 16:00
Part 2:	Friday _MMA	20 February 2026 from 8:30 to 18:30
	Saturday _MMA	21 February 2026 from 8:00 to 16:00

Part 1 Online, Part 2 Online

Module 6 Information Management, e-CTD

Part 1:	Friday	06 March 2026 from 8:30 to 18:00
	Saturday	07 March 2026 from 8:30 to 18:00
Part 2:	Friday	20 March 2026 from 8:30 to 18:30
	Saturday	21 March 2026 from 8:30 to 14:00 (project work)

Part 1 Online, Part 2 Pharmaceutical Institute, Bonn-Endenich

Written examination (Module 2 / 3 / 5), Thursday 16 April 2026, 13 o'clock
lecture hall X / University of Bonn

Module 7 Quality Management / Medical Devices

Part 1: Friday _QM 17 April 2026 from 13:00 to 18:30
Saturday _MD 18 April 2026 from 8:00 to 16:00

Part 2: Friday _MD 24 April 2026 from 8:30 to 18:30
Saturday _QM 25 April 2026 from 8:00 to 16:00

Part 1 Pharmaceutical Institute, Bonn-Endenich
Part 2 Online Submission of study paper: 26/May/2026 (Tuesday!)

Module 8 Chemical Pharmaceutical Documentation

Part 1: Friday 08 May 2026 from 8:30 to 18:30
Saturday 09 May 2026 from 8:00 to 16:00

Part 2: Friday 15 May 2026 from 8:30 to 18:30
Saturday 16 May 2026 from 8:00 to 16:00

Part 1 Online, Part 2 Online

Module 9 Pharmacology and Toxicology Documentation

Part 1: Friday 29 May 2026 from 8:30 to 18:30
Saturday 30 May 2026 from 8:00 to 16:00

Part 2: Friday 12 June 2026 from 8:30 to 18:30
Saturday 13 June 2026 from 8:00 to 16:00

Part 1 Online, Part 2 Online

Module 10 Clinical Documentation

Part 1: Friday 26 June 2026 from 8:30 to 18:00
Saturday 27 June 2026 from 8:30 to 18:00

Part 2: Friday 10 July 2026 from 8:30 to 18:00
Saturday 11 July 2026 from 8:30 to 16:15

Part 1 Online, Part 2 Online

Module 11 Benefit, Efficiency, Reimbursement

Thursday 23 July 2026 from 14:00 to 18:45
Friday 24 July 2026 from 08:30 to 18:30
Saturday 25 July 2026 from 08:00 to 16:00

Online Submission of study paper: 24/Aug/2026

Module 12 Regulatory Management / Decision Making

Friday 07 August 2026 from 8:30 to 18:30
Saturday 08 August 2026 from 8:00 to 16:00

Pharmaceutical Institute, Bonn-Endenich Submission of study paper: 07/Sept/2026

Oral examination (Module 8 / 9 / 10): Time period: September – November 2026