Module A	Morning	Morning	Midday	Afternoon	Evening
Day 1: 23 September 2026	Welcome & introduction to case studies	Scientific evolution of medicines' regulation	Plenary session: "Building the European medicines regulatory system: history, governance and strategic evolution	From regulatory approval to patienst access: How the EU HTA regulation is reshaping drug development	Welcome diner
Speaker(s)	**Jens van Wijngaarden**	**Bert Leufkens**	**Bert Leufkens, Hans van Bronswijk & Thoman Lönngren**	**Eline Hendriksen**	
Time	09.30 – 10.00	10.15-12.00	13.30-14.45	15.30-17.00	19.00 – 21.00
Day 2: 24 September 2026	Other European registration procedures, why do they exist and how to deal with them	The new Pharmaceutical Legislation; an overview of the reforms	The European centralised procedure and the role of the EMA	Drug development and route to market: academic, startup and SME perspective	Case studies
Speaker(s)	**Kora Doorduyn**	**Anna Wawrzyiak**	**Maud Dirken**	**Edwin Spaans**	**Jens van Wijngaarden & Hans van Bronswijk**
Time	9.15-10.45	11.00-12.30	13.30-15.15	15.30-17.00	19.30 – 21:30
Day 3: 25 September 2026		Insights in regulatory strategy: the why, the what and the how.	The international context of medicines' regulation, convergence, reliance and collaboration	Wrap-up Module A and case studies (looking forward to Module B)	
Speaker(s)	**To be decided**	**Carolina de la Portilla**	**Roelie Marinus**	**Jens van Wijngaarden & Hans van Bronswijk**	
Time	9.15-10.30	10.45-12.15	13.30-15.00	15.15 – 15.45	