



# 17<sup>th</sup> OPEN SCIENTIFIC EIP SYMPOSIUM ON IMMUNOGENICITY OF BIOPHARMACEUTICALS

**Program** 

March 16<sup>th</sup> – 19<sup>th</sup> 2026

# Monday, March 16th 2026

# Trainings

09:00	Welcome and Introduction by the EIP Chairman Daniel Kramer, Sanofi	
09:15	Immunogenicity Training (one beginner and one advanced immunogenicity training in parallel sessions) – topics to be announced EIP Team	
10:15	Coffee Break	
10:45	Bring Your Own Problems – Break Out Sessions EIP Team	
12:15	Lunch Break	
Main Program		
13:15	Keynote Presentation: Innate Immunity Bruno Lemaître, EPFL	
Session 1: Immunogenicity of New Modalities		
Session Chair: Lydia Michaut, Novartis		
14:15	Immunogenicity Risk Assessment for Nucleic Acid Therapeutics: A Comprehensive Evaluation for ASOs, siRNAs, and non-vaccine mRNA/LNPs Joanna Grudzinska-Goebel, Bayer	
14:45	Unraveling xRNA Immunogenicity: Insights and Future Horizons Alessio Maiolica, Novartis	
15:15	Minimizing immunogenicity testing for development of oligonucleotide therapeutics: A risk-based strategy An Zhao, Regeneron Pharmaceuticals	
15:45	Coffee Break	
16:15	Transient blockade of the CD28/B7 costimulatory pathway with abatacept inhibits immune responses to AAV vector and transgene product Rebecca Xicluna, Roche	
16:45	Advancing Immunogenicity Testing: Overcoming Drug Tolerance in a Neutralising Antibody Assay for an mRNA-Based Oncology Therapeutic Juliane Ober-Blobaum, Moderna	
17:15	Immunogenicity Risk Assessment and Mitigation Strategy for AAV Gene Therapy Hélène Haegel, Roche	
17:45	End of Day 1	



# Tuesday, March 17th 2026

#### Session 2: Regulatory

Chair: Daniel Kramer, Sanofi

9:00 New Version of the FDA Immunogenicity Guidance (title to be confirmed)

Joao Pedras-Vasconcelos, FDA

9:30 Future Direction of Immunogenicity at FDA (title to be confirmed)

Susan Kirshner, FDA

10:00 Harmonizing Adalimumab Anti-drug Antibody Assays through WHO International

Standards

Meenu Wadhwa, Medicines and Healthcare Products Regulatory Agency

10:30 Coffee Break

## **Session 3: Immunogenicity of Therapeutic Peptides**

Chair: Arno Kromminga, BioNTech

11:00 Immunogenicity Risk in Peptide Therapeutics: Navigating Complexity, Prediction

tools, and Bioanalytical Strategies
Montserrat Puig, Merck Sharp & Dohme

11:30 Clinical Pharmacology Perspectives on Immunogenicity of Peptide Drugs

Rajabi Mohsen, Novartis

12:00 Aligning Regulatory Expectations on Immunogenicity to Support Peptide Drug

**Product Approvals** 

Laura Salazar-Fontana, Immunogenicity Integrated

12:30 Lunch Break

# **Session 4: Immunogenicity Assays**

Chair: Christopher Tiedje, BioAgilytix

13:30 EIP Immunogenicity Assays and EIP Immunogenicity Risk Assessment Working

**Group Update** 

Linlin Luo, Merck Sharp & Dohme on behalf of EIP Assay Working Group Veerle Snoeck, UCB & Joanna Grudzinska-Goebel, Bayer on behalf of the EIP

Immunogenicity Risk Assessment (IRA) Working Group

13:45 Overcoming Unexpected Interference in a Highly Drug-Tolerant and Ultrasensitive

ADA-SPEAD Assay

Florian Anlauff, BioAgilytix

14:15 PABAD in Practice: Achieving High Drug Tolerance in NAb Assays

Lysie Champion, Celerion

14:45 When Generic Assays Succeed (and Fail): Case Studies and a Fit-for-Purpose

**Strategy for Non-Clinical ADA Support** 

Noor Bodrul, Roche

15:15 Coffee Break



15:45	The Immunogenicity Nexus: Investigating the Predictive Relationship Between Early-Onset ADA Kinetics in NHP and Human Immune Responses Griet Conickx, Sanofi
16:15	Positive Controls in Immunogenicity Assays: Case Studies and Best Practices Ester Biemans, Ardena
16:45	A Drug and Target-Tolerant Hybrid LBA-LC-MS Method for (Semi-)Quantification of ADA Isotopes, Using a Single-Tier Approach Hendrik Folkerts, ICON Bioanalytical Laboratory
17:15	End of Day 2



# Wednesday, March 18th 2026

#### **Session 5: Biosimilars**

Chair: Denise Sickert

09:00 EIP Biosimilar Working Group Update

Martin Ullmann, Fesenius Kabi on behalf of EIP

09:15 Establishing in-vitro assays for non-clinical immunogenicity assessment of bio-

similars

Marc Rosenbaum, Hexal AG

09:45 Title to be confirmed

Speaker to be confirmed

## Session 6: Prediction of immunogenicity

Chair: Sofie Pattyn

10:15 EIP NCIRA Working Group Update

Sebastian Spindeldreher, Integrated Biologix

10:30 Immunogenicity Risk Assessment: Enabling Quality by Design in Therapeutic

**Protein Discovery and Development** 

Tim Hickling, Quasor & Sophie Tourdot, Pfizer

11:00 Coffee Break

11:30 How to translate pre-clinical in silico immunogenicity profiling results to clinical

immunogenicity rates? Michael Gutknecht, Novartis

12:00 Unmasking Hidden Epitopes: Bridging In Silico Prediction and MAPPs Detection in

**Immunogenicity Assessment** Andreas Hollenstein, Roche

12:30 Lunch Break

13:30 Advancing Preclinical Immunogenicity Prediction: Machine Learning on Clinical

**Data and Pathogen Cross-Reactivity Integration** 

Olga Obrezanova, Astra Zeneca

14:00 An Innovative Reductionist Antigen Presenting System to Rapidly Screen Protein

**Immunogenicity in Human Lysosomes** 

Sergio Gabriele Colangelo, RBM S.p.A., an affiliate of Merck KGaA

14:30 Coffee Break

#### Session 7: Clinical Relevance of Immunogenicity

Chair: Noel Smith, Lonza

15:00 Short Talk: Immunogenicity Assessment in Acromegaly Patients Treated with

Ready-to-Use, Sustained-Release, Octreotide Subcutaneous Depot

K. Järås, Camurus



15:15 Assessment of Long-term Immunogenicity of Lonapegsomatropin in Children With

**Growth Hormone Deficiency**Per Holse Mygind, Ascendis

15:45 Short Talk: Immunogenicity Publication Bias and Its Consequences for Predictive

**Models: A Call for Transparent Reporting** 

Sophie Tascedda, RC2NB

16:00 End of Day 3

Session 8: Social Event (reconvene at 17:00 in the Altis lobby)

Chair: Barbara Vercruyssen



# Thursday, March 19th 2026

## **Workshop – Parallel Sessions**

09:00 Deep Dive:

Recent Publication "Immunogenicity Risk Assessment for Tailored Mitigation and

Monitoring of Biotherapeutics and Biosimilars"

Veerle Snoeck, UCB and Joanna Grudzinska-Goebel, Bayer on behalf of the EIP

Immunogenicity Risk Assessment (IRA) Working Group

9:00 Deep Dive:

Three Recent Publications of the NCIRA Working Group (EIP Recommendations for

Dendritic Cell Maturation Assays, MAPPs and T-Cell Assays)

Sebastian Spindeldreher, Integrated Biologix on behalf of the EIP NCIRA Working

Group

10:30 Coffee Break

11:00 Bring Your Own Problems – Break Out Sessions

**EIP Team** 

12:30 Summary and Outlook of the EIP Chairman

Daniel Kramer, Sanofi

12:45 End of the conference

