Commission Implementing Regulation (EU) 2021/808 of 22 March 2021 on the performance of analytical methods for residues of pharmacologically active substances used in food-producing animals and on the interpretation of results

1 March 2022
(09:00 – 12:00 CET)

09:00 – 09:15	INTRODUCTION	Presentation of the speakers, welcome to the participants *Prof. Bruno Le Bizec, NRL Nantes, LABERCA*
09:15 – 09:45	PRESENTATION	25 min presentation + 5 min questions Current activities of the European Commission in the area of residues of veterinary medicines and the implementation of Reg. (EU) 2017/625 *Mr Frans Verstraete, EU DG SANTE Bruxelles*
09:45 – 10:15	PRESENTATION	25 min presentation + 5 min questions Commission Implementing Regulation (EU) 2021/808 – Overview of changes *Dr Joachim Polzer, EURL Berlin, BVL*
10:15 – 10:25	BREAK	
10:25 – 10:50	PRESENTATION	20 min presentation + 5 min questions EURL Guidance Document on Screening Method Validation (focus on microbiological, immunological and physico-chemical screenings) *Dr Eric Verdon and Dr Valerie Gaudin, EURL Fougères, Anses*
10:50 – 11:15	PRESENTATION	20 min presentation + 5 min questions EURL Guidance Document on Confirmation Method Validation (conventional approach) *Drs Saskia Sterk , EURL Wageningen, WFSR*
11:15 – 11:40	PRESENTATION	20 min presentation + 5 min questions EURL Guidance Document on Confirmation Method Validation (alternative approach) *Dr Joachim Polzer, EURL Berlin, BVL*
11:40 – 12:00	CONCLUSION	1-2 cross-cutting questions answered by each expert in his/her field