

**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 <i>Prof. Bruno Le Bizec, NRL Nantes, LABERCA</i>
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 <i>Mr Frans Verstraete, EU DG SANTE Bruxelles</i>
09:45 – 10:15	PRESENTATION	25 min presentation + 5 min questions Commission Implementing Regulation (EU) 2021/808 – Overview of changes <i>Dr Joachim Polzer, EURL Berlin, BVL</i>
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) <i>Dr Eric Verdon and Dr Valerie Gaudin, EURL Fougères, Anses</i>
10:50 – 11:15	PRESENTATION	20 min presentation + 5 min questions EURL Guidance Document on Confirmation Method Validation (conventional approach) <i>Drs Saskia Sterk, EURL Wageningen, WFSR</i>
11:15 – 11:40	PRESENTATION	20 min presentation + 5 min questions EURL Guidance Document on Confirmation Method Validation (alternative approach) <i>Dr Joachim Polzer, EURL Berlin, BVL</i>
11:40 – 12:00	CONCLUSION	1-2 cross-cutting questions answered by each expert in his/her field