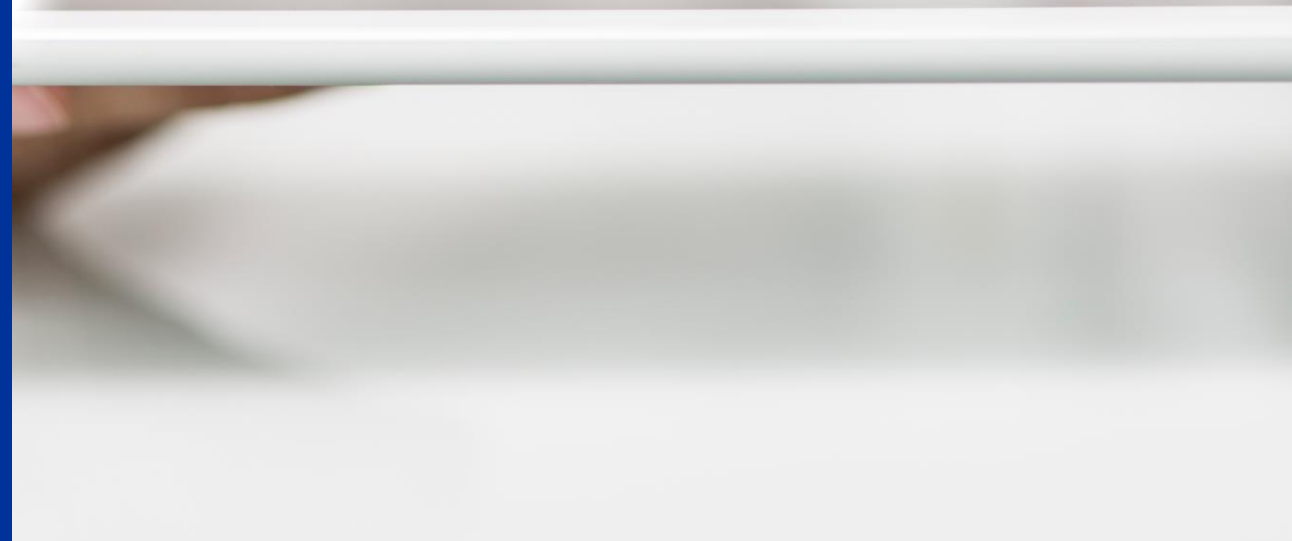


New Fee Regulation

Human Industry Q&A clinic

10 February 2025



Session's scope



Questions on:

- **Fees**
- **Incentives**
- **The prepayment of Scientific Advice, Certificates and Parallel Distribution**
- **The new Certificates Processing System (including access management)**
- **Regulatory matters**



System (e.g., IRIS) technical queries, malfunctions etc.

Available regulatory guidance and support

Updated documents are available on EMA's website

- [Regulation \(EU\) 2024/568](#)
- [Working arrangements](#)
- [Fee Q&As](#)
- [Guidance on how to pay fees](#)

Support channels

Standard support channels (currently available and ongoing)



- **Technical queries:** [EMA Service Desk](#)
- **Fee-related inquiries:** form on [Fees payable to the European Medicines Agency | European Medicines Agency \(EMA\)](#)
- **Invoicing queries:** [How to pay | European Medicines Agency \(EMA\)](#)

Until 31st March 2025



(in addition to the above) nfr@ema.europa.eu