ECFS Patient Registry (ECFSPR) FREE webinar

Highlights and insights from the ECFS Patient Registry (ECFSPR)

Tuesday 9th July 2024 13:00-14:00 CEST / 12:00 – 13:00 UK (GMT+1)

If you are able to attend, please click here to register. You will receive your personal log in after registration

Time	Presentation	Speaker
13:00 - 13:05	Welcome	Chris Smith (UK)
13:05 - 13:15	Introduction to the ECFSPR	Alice Fox (UK)
13:15 - 13:35	ECFSPR Highlights Report 2022	Domenique Zomer-van Ommen (NL)
13:35 - 13:45	Collecting and interpreting data from ECFSPR	Anna Zolin (IT)
13:45 - 13:55	Q&A moderated by Chris Smith (UK)	All speakers
13:55 - 14:00	Closing	Chris Smith (UK)

Objectives

At the end of the session, the participant will be able to:

- Describe how the ECFSPR is organised, which countries are involved and the goals of the registry
- 2. Discuss the key information about how cystic fibrosis (CF) affects people with CF and their families throughout Europe
- Explain how the ECFSPR collects, measures, compares and interprets data of people with CF living in Europe and neighbouring countries

Target audience

All members of the CF multi-disciplinary team, patient organisations, and people with CF and their families.

The webinar will be recorded and made available on the <u>ECFS Education Platform</u> and <u>ECFS website</u> afterwards.

