Looking for information and community networking opportunities?

ECRD PROGRAMME
COMMITTEE CO-CHAIRS

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WHAT IS ECRD?

› The European Conference on Rare Diseases & Orphan Products is the unique forum across all rare diseases, across all European countries, bringing together all stakeholders - patients' representatives, academics, researchers, health care professionals, industry, payers, regulators and policy makers.

› ECRD provides the state-of-the-art of the rare disease environment, monitoring and benchmarking initiatives.

› ECRD covers research, development of new treatments, health care, social care, information, public health and support at European, national, regional and international levels.

Organised by: EURORDIS
Co-organised by: DIA
With the support of: orphanet, AMFTELETHON, the European Union
› Get the latest information related to the rare disease field
› Share inspiring and innovative strategies, services or projects that could be useful to others during the plenary “soap box” session or as a poster presenter
› Network with fellow rare disease patients, EURORDIS staff and all other stakeholders from across Europe and beyond
› Find out how you can benefit from and/or be involved in European Reference Networks
› Learn about initiatives fostering the development of orphan medicinal products and getting treatments available to patients
› Discover important breakthroughs for rare diseases
› Discover the potential brought by digital technologies
› Contribute to addressing the challenges of healthcare pathways
› Hear about innovative solutions and programmes that address quality of life issues
› Feel part of the rare disease community
› Take advantage of the special registration prices and fellowship programme for patient advocates

LANGUAGES

Simultaneous interpretation and live streaming of the Opening and Plenary sessions on 11 May 2018 will be available in:

ENGLISH  FRENCH  GERMAN

For more information, please visit www.rare-diseases.eu

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