

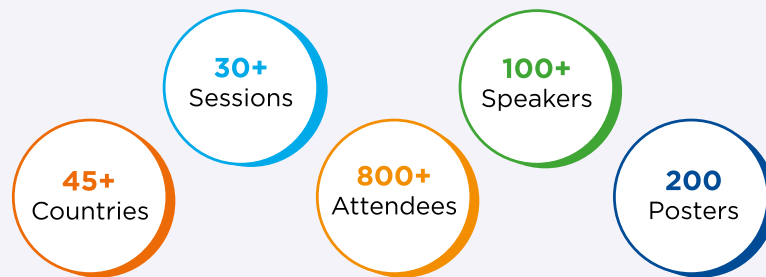


# Rare Diseases 360°

Collaborative Strategies  
to leave no-one behind

9th European Conference  
on Rare Diseases & Orphan Products  
**10-12 May 2018 Vienna**  
Messe Wien Exhibition & Congress Center

## Looking for information and community networking opportunities?



### ECRD PROGRAMME COMMITTEE CO-CHAIRS



**Rainer Riedl**  
President  
Pro Rare Austria  
Austria



**Justina Januševičenė**  
Executive for the development of health care  
technologies and innovations  
Lithuanian University of Health Sciences  
Former Director of Healthcare resources and  
innovation management department at the  
Health Ministry of Lithuania



**Vinciane Pirard**  
Co-Chair Joint Task Force on Orphan Drugs  
& Rare Diseases of EFPIA – EuropaBio  
Sanofi Genzyme  
Netherlands

### 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:



Co-organised by:



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## LEARN – DIALOGUE – CONNECT

- › **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](http://www.rare-diseases.eu)

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