THE 10th EUROPEAN CONFERENCE ON
RARE DISEASES & ORPHAN PRODUCTS

ecrd

ONLINE on 14-15 May 2020

THE JOURNEY OF LIVING WITH
a RARE DISEASE in

2030

JOIN US AT THE LARGEST PATIENT-LED RARE DISEASE EVENT TAKING PLACE IN EUROPE

Leading, inspiring and engaging all rare disease community stakeholders to take action

Hundreds of attendees
including patient advocates, healthcare professionals, healthcare industry, academics, regulators, payers and policy makers

2 days
14-15 May 2020
ONLINE

Worldwide attendance

 # ECRD2020

Over 100
expert chairs, speakers and panellists

PROGRAMME COMMITTEE CO-CHAIRS

Prof. Milan Macek
Professor of Medical and Molecular Genetics, Motol University Hospital and Charles University Prague, Czech Republic

Maria Montefusco
President, Rare Diseases Sweden

Violeta Stoyanova-Beninska
Chair, Committee of Orphan Medical Products, European Medicines Agency

Organised by
EURORDIS
RARE DISEASES EUROPE

Co-organised by
orphanet
Inserm

With the support of
AFMTÉLÉTHON
INNOVER POUR GUÉRIR

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the Health Programme
of the European Union

LEARN – NETWORK – INSPIRE

- **Co-create** the policies and services of the future to support measureable and improved health outcomes for people living with a rare disease
- **Learn** about how care for rare diseases is organised across Europe, what are the trends, innovations and emerging models of care
- **Share/exchange** your knowledge and experience with other fellow clinicians from all around Europe and beyond to create collaborations
- **Meet** patients online from other countries **and learn from their** experiences
- **Showcase** and share your work with a wide, multi-stakeholder audience as a poster presenter or as a speaker in the open microphone plenary session
- **Exchange views on** technological developments that are changing the rare disease landscape and contribute to debates on the opportunities, ethical, legal and social implications in areas such as data sharing and artificial intelligence
- **Learn** about the important progress made to advocate for rare diseases as a health and human rights priority at the United Nations and international agencies
- **Learn more** about growing efforts to advocate for organised and integrated social support and patient-centred care at the national level
- **Gain** insight into European and national rare disease policies
- **Obtain** CME (Continuing Medical Education) credits by attending ECRD 2020

LANGUAGES

Simultaneous interpretation and live streaming of the Opening and Plenary sessions on 14 May 2020 will be available in:

ENGLISH | **FRENCH** | **GERMAN**

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