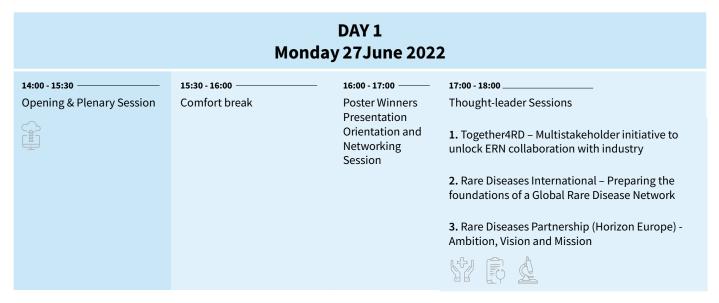
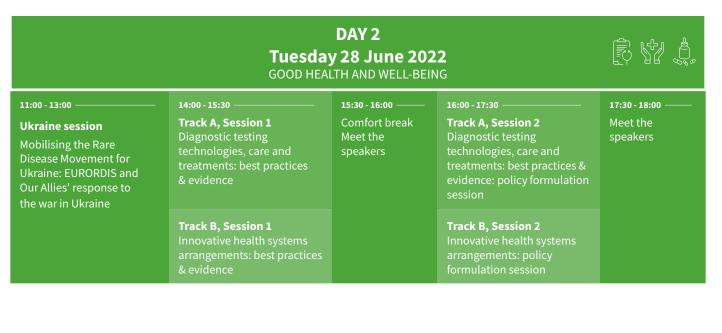
ECRD 2022 is taking place fully online and is spread over 5 days.

On the first day of ECRD 2022, the Opening Plenary Session sets the scene for this year's online conference and will be followed by an orientation and networking session along with several parallel Thought Leader sessions. The following three days are dedicated to discussions on how to reach our 3 visionary goals for people living with a rare disease, inspired by the Rare 2030 project and implemented through a new policy framework. A Closing Plenary on Day 5 reminds the audience that rare diseases must be addressed across all of Europe (beyond the EU) and on a global scale. This closing session will also leave participants with a clear call to action in the immediate, medium and long-term future.

All times listed are Central European Summer Time (CEST)









Reducing inequalities

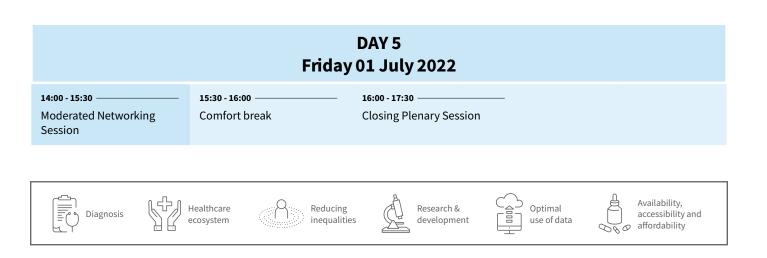




Availability, accessibility and affordability

PROGRAMME AT A GLANCE

DAY 3 Wednesday 29 June 2022 **REDUCED INEQUALITIES** 11:00 - 12:00 14:00 - 16:00 16:00 - 16:30 16:30 - 18:00 18:00 - 18:30 Comfort break Track C/D, Session 1 Track C, Session 2 Meet the Language Networking Hub Invisibility as a roadblock Meet the Inequalities in accessing speakers Connect with fellow health and social care towards reducing speakers participants in your inequalities native language Track D, Session 2 Inequalities in accessing and 11:00 - 13:00 remaining in education & Swedish Satellite Meeting employment 12:00 - 13:30 **Danish Satellite Meeting** (Forberedelsesmøde for patientrepræsentanter 11:00 - 11:45) DAY 4 Thursday 30 June 2022 INDUSTRY, INNOVATION AND INFRASTRUCTURE 14:00 - 15:30 -15:30 - 16:00 -16:00 - 17:30 09:30 - 11:30 17:30 - 18:00 **Dutch Satellite Meeting** Track E, Session 1 Track E, Session 2 Meet the speakers data ecosystem 11:00 - 12:00 17:00 - 19:30 Language Networking Hub Romanian Connect with fellow Satellite Meeting participants in your native language



LEARNING PATHWAYS

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Diagnosis: improving access to effective diagnostic tools for rare diseases and accelerating diagnosis through the optimised sharing of data. Found in both sessions of Tracks A, B and E Thought leader session. RD Partnership



A highly specialised <u>healthcare ecosystem</u>, with political, financial and technical support at European and national levels, that leaves no person living with a rare disease in uncertainty regarding their diagnosis, care or treatment. Found in both sessions of Tracks A, B and E



<u>Reducing inequalities</u> through the identification and removal of systemic barriers through targeted policy interventions to ensure people living with a rare disease are given the same opportunities as others to reach their maximum level of health and well-being. Found in both sessions of Tracks C and D



Rare disease <u>research and development</u> of orphan medicinal products maintained as a priority - across basic, clinical, translational and social research. Found in both sessions of Track F Thought leader session. RD Partnerships. Opening plenary session



Data used optimally to improve the health and well-being of people living with a rare disease. Found in Session 1 of Track C/D and both sessions of Track E



Improve the availability, <u>accessibility and affordability</u> of rare disease treatments, by attracting investments, fostering innovation and collaboration across countries, to address inequalities. Found in both sessions of Tracks A, B and F