



ITINERARE Innovative Therapies in Rare Diseases University Research Program of the University of Zurich

Caring for the orphans: Ethical issues in the context of rare diseases

June 15-16, 2023 University of Zurich (Switzerland)

Thursday, 15 June

- 12:00 Arrival & Lunch Bag
- 12:30 Welcome & Introduction
- 12:45 **Havi Carel** (University of Bristol) Rare diseases, epistemic injustice and the good life
- 13:45 Heiner Fangerau (University of Duesseldorf) Diagnoses and historical concepts of disease – is the end of a diagnostic odyssey a useful fiction?
- 14:45 Coffee Break

Parallel sessions

15:15	Sebastian Wäscher (University of Zurich) The burden of rare diseases. Patient narratives on managing their disease	Kathryn Tabb (Bard College) Individualized genetic therapies as a treatment- research hybris	Susanne Wehrli (University of Zurich) Access to healthcare among Swiss adults with rare diseases – the influence of facilitating and inhibiting individual factors
16:00	Lainie Friedman Ross (University of Rochester) Thinking zebras not horses when the herd runs past: Atypical Diabetes Mellitus	Urte Laukaityte (UC Berkeley) Rare diseases in the context of effective altruism	Jacopo Morelli (University of Modena-Reggio) Kantian constructivism applied to rare diseases

17:00 Rachel Grob (University of Wisconsin)

Diagnostic information and uncertainty in the newborn period: qualitative dimensions of families' experiences with novel technologies

- 18:00 **Bernice Elger** (University of Basel) Genetic testing of children: Child welfare versus parental autonomy
- 19:00 End
- 19:30 Dinner

Friday, 16 June

- 9:00 **Dana Mahr** (University of Geneva) Citizen science and biomedical research: the experiential knowledge of a lived genome
- 10:00 Marcello Ienca (Technical University of Munich and Ecole Polytechnique Fédérale de Lausanne EPFL)
 Patient-engagement in rare disease research. The case study of rare neurological diseases
- 11:00 Break

Parallel sessions

11:30	Lucie Perillat (University of Toronto) Rare diseases: a universal priority	Francesca Greco (Insubria University) Understanding orphan drugs: roles and perspectives regarding orphan drugs for rare diseases
12:15	Nina Streeck (University of Zurich) The vulnerable orphans: Does vulnerability constitute a right to solidarity?	Rebeca Méndez-Veras (Universidad Mariano Gálvez de Guatemala) Ethical issues regarding clinical exome and targeted gene sequencing. A Guatemalan experience

13:00 Lunch Break

- 14:00 **Deborah Mascalzoni** (Uppsala University) Respecting the rights of children with rare disease in research within their lifespan: challenges and perspectives
- 15:00 **Eva Winkler** (University of Heidelberg) Data use for research in rare disease. Ethical considerations on the governance of data initiatives
- 16:00 Coffee Break
- 17:00 Niklas Juth (Uppsala University and Karolinska Institutet) Prioritization of orphan drugs: considerations of ethics and justice
- 18:00 End