



UPPSALA  
UNIVERSITET



# MIND THE RISK FINAL CONFERENCE

29 November 2019 • Clarion Hotel Gillet, Uppsala



RIKSBANKENS  
JUBILEUMSFOND  
STIFTELSEN FÖR HUMANISTISK OCH  
SAMHÄLLSVETENSKAPLIG FORSKNING

Register at [crb.uu.se/mind-the-risk/conference](http://crb.uu.se/mind-the-risk/conference)

# PRELIMINARY PROGRAMME

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08.30 Registration, coffee

08.50 **Introduction**  
Mats G. Hansson

## PANEL I

09.00 **GMS: From Mendel to Gene sequencing in the clinic**  
Anna Lindstrand

09.25 **Perceptions of risk and predictive testing amongst people with, or at risk of, rheumatoid arthritis**  
Marie Falahee

09.50 **Biologic calculators for targeting the use of biologics in rheumatoid arthritis**  
Katherine Payne, Caroline Vass

10.15 Questions and comments

10.25 Bio break

## PANEL II

10.30 **Non-invasive prenatal testing and the delicate balancing of risk information**  
Susanne Georgsson

10.55 **Peoples' conceptions and preferences regarding genetic risk information**  
Jennifer Viberg Johansson

11.20 **How do family aspects affect expectations towards predictive genetic tests?**  
Sabine Wöhlke

11.45 Questions and comments

12.00 Buffét lunch and poster presentations

## PANEL III

13.00 **Genetic risk information across cultures**  
Silke Schickanz

13.25 **Psychological implications of Direct to Consumer Genetic testing: consumers' profiling and psycho-decisional follow up**  
Serena Oliveri, Ilaria Cutica, Alessandra Gorini, Gabriella Pravettoni

13.50 **Serious Games to improve Genetic Literacy and Genetic Risk Awareness in the General Public**  
Serena Oliveri, Renato Mainetti, Ilaria Cutica, Alessandra Gorini, Gabriella Pravettoni

14.15 **Genetic risk policy in the Big Data age**  
Frederic Bouder, Sanja Mrksic Kovacevic

14.40 **Ethical implications of providing genetic risk information**  
Ulrik Kihlbom

15.00 Questions, comments and closure of English part

15.20 Coffee and poster presentations

## PANEL IV: Swedish section

16.00 Genetisk risk i mötet med patienter

- **Rutinmässig cancerdiagnostik** - Maria Johansson-Soller
- **Gensekvensering för sällsynta diagnoser** - Anna Lindstrand
- **Rapport från projektet** - Anna Holm, Pär Segerdahl
- **Vad tycker patienterna?** - Karin Schölin Bywall

17.30 Closure of Swedish section