

12 and 13 November 2020

Thursday N	loveniber 12 2020		
10:00-10:15	Welcome		
	<u>P. Devilee</u> : Chair Hebon Steering Group		
Session 1 – RF	RSO		
10:15-10:35	<u>M.P. Steenbeek</u> : Early salpingectomy (TUbectomy) with delayed oophorectomy to improve quality of life as alternative for risk-reducing salpingo-oophorectomy in women with a BRCA1/2 pathogenic variant (TUBA study): a prospective multicentre preference trial		
10:35-10:55	<u>I.A.S. Stroot</u> : Inter- and intra-observer variability in the assessment of muscle mass with the whole body DEXA scan		
10:55-11:15	M.H.D. van Bommel: Evaluation of a Patient Decision Aid for Risk Reducing Surgery in premenopausal BRCA1/2 mutation carriers		
Session 2 – Ge	enetics		
11:15-11:35	A. Hollestelle: The genomic landscape of breast cancers from CHEK2 c.1100delC mutation carriers		
11:35-11:55	<u>M. Vreeswijk</u> : CRAFT: Cancer Ri <mark>sk Assessment throug</mark> h Functional Testing of variants in BRCA1, BRCA2 and PALB2		
11:55-12:00	Closing session 1 and 2 - P. Devil <mark>ee</mark>		
Session 3 – Pe	ersonalised risk prediction		
15:00-15:20	R. de Groot: Routine recruitment and data acquisition: the Hebon Infra project		
15:20-15:40	Xin Yang: Development of a Comprehensive Risk Prediction Model for BRCA1 and BRCA2 mutation carriers – TRANsIBCCS		
15:40-16:05	<u>G. Evans</u> : What are the benefits <mark>and harms of risk stratified scre</mark> ening as part of the NHS breast screening Programme?		
16:05-16:30	L.J. van 't Veer: The WISDOM Study: breaking the deadlock in the breast cancer screening debate		
16:30-16:55	Future of personalised risk prediction: a discussion		
16:55-17:00	Closing session 3 - P. Devilee		
Friday Nov	ember 13 2020		
Session 4 – M	lainstreaming genetic services		
10:00-10:20	N. Hoogerbrugge: Tumor-First-workflow: Nationwide Implementation of Ovarian Cancer		

Heredity Prescreening to stratify both Genetic Testing and Treatment Options K. Bokkers: First experiences of oncologic surgeons, medical oncologists and nurse 10:20-10:40

specialists with a mainstreaming approach for germline genetic testing for patients with breast cancer

Session 5 – Risk communication and management

10:40-11:00	N.M. Medendorp: Discussing ur	ncertainty about multigene panel	testing: different manners,
	different outcomes?		

- 11:00-11:20 <u>M.L. Haadsma:</u> Informing family members about predisposition for hereditary cancer: results of a proactive and personalized approach
- 11:20-11:40 <u>M. Hooning</u>: Prognosis of T1 breast cancer in BRCA1/2 mutation carriers: is screening beneficial?
- 11:40-12:00 <u>M.M.A. Tilanus-Linthorst</u>: MRI breast screening for women with $\geq 20\%$ lifetime risk is costeffective in Europe and preferred by most women the Famrisc trial shows

12:00-12:05	Closing
	P. Devilee: Chair Hebon Steering Group