10TH ANNUAL MEETING
Stockholm • 2026

10th Meeting of the European Hereditary Tumour Group (EHTG)

Friday, August 28 to Sunday, August 30, 2026

Stockholm, Sweden

Final Programme

Invitation 10th Meeting – Stockholm

Dear Members, Friends, Affiliates and Sponsors,

We are delighted to welcome you to the 10th Annual EHTG Meeting, taking place from August 28-30, 2026 in Stockholm.

Join us for the full program focusing on new technologies, emerging treatments and other developments in the fields of genetics, gastroenterology, oncology, endoscopy and surgery in hereditary gastrointestinal tumours.

The program will feature cutting-edge and controversial topics, expert discussions and debates, and valuable insights from both clinical and patient perspectives. Together, we will explore advances in precision medicine, prevention, surveillance, guideline development, and collaborative European research initiatives.

Beyond the scientific sessions, this year's meeting offers a unique opportunity to strengthen collaboration within our growing international community. We look forward to inspiring discussions, new connections, and shaping the future of hereditary tumour research and care together.

Please visit **www.ehtg.org** for more information and contact **gs007@ehtg.org** for organizational queries.

We are excited to welcome you to Stockholm at the end of August!

Sincerely yours,
EHTG Programme Committee and Board Members

Organisation

Board Members:

Chair: Toni Seppälä
Vice-Chair: Gabriela Möslein
General Secretary: Kelly Kohut
Treasurer: Elizabeth Half
Y-EHTG: Maria Rasmussen
Communications Chair: Patrick Benusiglio
Programme Chair: Maartje Nielsen
Education: Neil Ryan
Nurses/Genetic Counseling: Mechelle Loughrey

Programme Committee:

Maartje Nielsen, The Netherlands
Ann-Sofie Backman, Sweden
(Programme Directors)
Patrick Benusiglio, France
Kaisa Fritzell, Sweden
Elizabeth Half, Israel
Kelly Kohut, United Kingdom
Florian Kühn, Germany
Annica Lindblom, Sweden
David Ljungman, Sweden
Matthias Löhr, Sweden
Maria Rasmussen, Denmark

Congress Organisation/ Secretariat:

Gabriele Sponholz
gs007@ehtg.org
+49 160 8459502
65207 Wiesbaden, Germany

MCE Medical Congresses & Events

Location:

ERSTA Conference Centre
Erstagatan 1K
Södermalm, Stockholm
+46 (0)8-714 63 41

Registration and Registration Fee:

See online via www.ehtg.org

Certification:

The event has been certified by UEMS/ EACCME. Credit points to follow soon, see website

10TH ANNUAL MEETING Stockholm • 2026

Content

Content

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Meeting Overview

Day 1 – Friday, August 28

Time	Event – Room Bring/Norrby
08:30-10:00	Site Visit Karolinska University Hospital, Solna (limited spaces)
09:30-11:00	Y-EHTG Come together at ERSTA Conference center
11:00-11:30	Registration / Coffee
11:30-11:45	Welcome to EHTG 2026
11:45-12:15	New Developments in Preventive Measures and Treatment Options
12:15-13:00	New Developments in Preventive Measures and Treatment Options in Hereditary Forms of (GI) Tumors
13:00-14:00	Lunch Break
14:00-15:30	Novel Approaches to Genetic Counselling
15:30-16:00	Coffee Break
16:00-17:30	A Paradigm Shift in Rectal Cancer Management
17:30-18:30	Scientific Speed-Dating and Poster Presentation
18:30-20:00	Welcome Reception with a View – ERSTA Conference Centre
20:00	Young EHTG – Meet & Great

Notes

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Meeting Overview

Day 2 – Saturday, August 29

Time	Event – Room Bring/Norrby
08:00-08:30	**ILSD Business meeting**
08:30-09:30	**Secondary use of cancer health data in Europe** Managing hereditary cancer registries Study proposals /abstract ILSD/other EU collaborations
09:30-10:00	**Coffee Break**
10:00-11:15	**New hereditary gastrointestinal cancer syndromes** Genetics session
11:15-12:00	**Hereditary Colorectal Cancer: Mucosal Immunology and Vaccines** Vaccination in Lynch Syndrome: Clinical Feasibility and Promising Early Outcomes
12:00-13:00	**Lunch**
13:00-14:30	**Rare Cancers in Hereditary Gastrointestinal Syndromes Clinical and molecular session** Infrequent cancers in Lynch syndrome, with a focus on glioma
14:30-15:00	**Coffee Break**

Meeting Overview

Day 2 – Saturday, August 29

Parallel Working Groups

Time	Event – Cederschiöldsalen
08:30-10:00	**Research Collaborative Workshop: Polyposis Syndromes** Patient Perspectives: Do we really know how patients are feeling, and can we acquire that knowledge with patient reported data?
10:00-12:00	**Endoscopy Session** FAP and Lynch** Management of Duodenal Polyposis in FAP Including Surveillance and Management of Gastric Lesions Endoscopic Management of Polyposis in FAP, Lower GI. Endoscopic Surveillance in Lynch syndrome
12:00-13:00	**Lunch**

Meeting Overview

Day 2 – Saturday, August 29

Time	Event – Room Bring/Norrby
15:00-15:50	**Chemoprevention** Aspirin in Lynch Syndrome — Standard of Care or Premature?
15:50-16:00	**Short Break**
16:00-16:30	**Dynamic EHTG guidelines – Controversial and Ethical Issues in Development of Guidelines in Hereditary Syndromes**
16:30-17:30	**Interactive Discussion & Thematic Debate: "What Do We Lack, What Do We Need, and Who Should Lead?"**
18:15	**Transfer to Dinner at Djurgården**
19:30	**Congress Dinner at Junibacken, Djurgården**

Meeting Overview

Day 3 – Sunday, August 30

Time	Event – Room Bring/Norrby
08:30-09:00	**EHTG Business meeting**
09:00-10:00	**Changes in Population Screening and the Consequences for Patients with an Increased Risk** The Swedish Cancer Strategy, Screening and Surveillance
10:00-10:30	**Coffee Break**
10:30-11:45	**Ideas for the Future and New Developments**
11:45-12:15	**Heading forward together. Summary, conclusion and final remarks**

Scientific Programme

Day 1 – Friday, August 28

08:30-10:00	**Site Visit Karolinska University Hospital, Solna (limited space)**
9:30-11:00	**Y-EHTG Come together at ERSTA Conference center**
11:00-11:30	**Registration / Coffee**
11:30-11:45	**Welcome to EHTG 2026** *Ann-Sofie Backman, Sweden* *Maartje Nielsen, The Netherlands* *Toni Seppälä, Finland*
11:45-13:00	**New Developments in Preventive Measures and Treatment Options** *Chairs: Matthias Löhr, Sweden* *Emma Tham, Sweden* *Alethea Tang, United Kingdom*
11:45-12:15	**Precision Medicine in Hereditary Gastrointestinal Tumors** *Richard Rosenquist Brandell, Sweden*
12:15-13:00	**New Developments in Preventive Measures and Treatment Options in Hereditary Forms of (GI) Tumors**
	ABSTRACTS
8'+2'	**Engineering the Two-in-one Hit Mechanism of Colorectal Carcinogenesis in Normal Colon Organoids Derived from Lynch syndrome Carriers** *Aysel Ahadova, Germany*
8'+2'	**Somatic CTNNB1 Mutations as a Positive Predictor of Lynch syndrome** *Aysel Ahadova, Germany*

Scientific Programme

Day 1 – Friday, August 28

8'+2'	**Cell-Free DNA in Hereditary and High-Risk Malignancies 2 (CHARM2): A Randomized Controlled Trial for Early Cancer Detection** *Sarah Ridd, Canada*
8'+2'	**Epigenetic Treatment of Lynch Syndrome-associated Colorectal Carcinoma and BRAF-mutant Colorectal Carcinoma** *Anna Scheßl, Germany*
13:00-14:00	**Lunch Break**
14:00-15:30	**Novel Approaches to Genetic Counselling** *Chairs Anna Rosen, Sweden* *Kaisa Fritzell, Sweden* *Sanne Baiwa ten Broeke,The Netherlands*
14:00-14:20	**Implementing Mainstream Germline Genetic Testing in Breast Cancer** *Svetlana Bajalica Lagercrantz, Sweden*
14:20-14:40	**Artificial Intelligence and Genetic Counseling in Hereditary Cancer Syndromes** *Elen Siglen, Norway*
14:40-15:30	ABSTRACTS
8'+2'	**Genetic Counselling Regarding Recessive Syndromes Associated with Cancer Predisposing Genes: A Consensus on Shared Decision-Making** *Judit Sanz Buxo, Spain*
8'+2'	**Exploring Quality of Life, Illness Representations, Coping Strategies, and Needs in Patients with APC-Related Desmoid-Type Fibromatosis** *Bianca Scacciati, Italy*
8'+2'	**Frequency of Second Hits in Normal Lynch syndrome Colonic Mucosa: a Paired Spatial Mapping Pilot Comparing MMR-deficient Crypts With a Clonal Reference Marker** *Leon Marin Grez, Germany*

Scientific Programme

Day 1 – Friday, August 28

8'+2' **Implementing a Canadian Population Genomic Screening Study for Hereditary Cancers**
Laura Redondo, Canada

8'+2' **Diagnostic Value of Long-Read Sequencing in Unexplained Mismatch Repair Deficient MMR) Cancer Cases**
Sanne Bajwa-ten Broeke, The Netherlands

15.30-16.00 **Coffee Break**

16:00-17:30 **A Paradigm Shift in Rectal Cancer Management**
Chairs: David Ljungman, Sweden
Toni Seppälä, Finland
Gustav Silander, Sweden

16:00-16:20 **The Surgeon's Perspective: Organ Preservation and Watch and Wait in Rectal Cancer**
Per Nilsson, Sweden

16:20-16:40 **The Oncologist's Perspective Management of dMMR Rectal Cancer (Consequences for Patient with Hereditary Cancer Lynch syndrome)**
Mef Nilbert, Sweden

16:40.17:30 **Greetings and welcome from Ersta hospital CEO, an organizational perspective**
Sara Lindholm Larsson, Sweden

ABSTRACTS

6'+2' **Incidental Colorectal Cancer in APC-Mutated Familial Adenomatous Polyposis: Implications for Surgical Strategy**
Emanuele Rausa, Italy

6'+2' **Surgery for MUTYH-Associated Polyposis: Surgical Management and Outcomes in a Single-Centre Cohort of 171 Patients**
Alberto Gorini, Italy

Scientific Programme

Day 1 – Friday, August 28

6'+2' **Upper Gastrointestinal Neoplasia Detection in Lynch Syndrome: High Yield of Esophagogas troduodenoscopy Surveillance**
Elizabeth Half, Israel

6'+2' **Association of Prior Cancer Burden with Cancer Incidence in MLH1 and MSH2 Lynch syndrome Carriers Undergoing Colonoscopy Surveillance**
Kalle Hokkanen, Finland

6'+2' **High Prevalence of Lynch syndrome Identified via Clinical Whole-Genome Sequencing in a Large Swedish Colon Cancer Cohort**
Kristina Lagerstedt Robinson, Sweden

17:30-18:30 **Scientific Speed-Dating and Poster Presentation (BIBLIOTEK)**
Y-EHTG

5'+2' **A Truncating NLRC4 Gene Variant Predisposing to Familial Colorectal Carcinoma**
Taina Nieminen, Finland

5'+2' **Not All Deficiencies Are Created Equal: Clinical Landscapes and Immunotherapy Outcomes in Mismatch Repair Deficient Colon Cancer**
Sofia Silva, Spain

5'+2' **Endometrial Cancer in Lynch Syndrome: Age at Diagnosis and Family History in a US Genetic Testing Registry**
Laura Wollborn, USA

5'+2' **Personalised Risk Stratification in Colorectal Cancer Screening: Integrating Germline Genetic Predisposition and Clinical Data in a Danish Population-Based Screening Cohort**
Heidi Brinch, Denmark

5'+2' **Attenuated, Variable Phenotype in a Family with Peutz-Jeghers Syndrome due to a Splice Site Variant with Reduced Splicing Efficiency**
Laura Grohs, Germany

Scientific Programme

Day 1 – Friday, August 28

5'+2'	**Endoscopic Communication and Risk Interpretation in Hereditary Colorectal Cancer: the ENCOUNTER Study – A Qualitative Interview Study of Endoscopists** *Alexander Frank, Sweden*
5'+2'	**Barriers to Implementation of Hereditary Gastrointestinal Cancer Screening in Low-Resource Health Systems: a Narrative Review** *Musa Usman Babagari, Nigeria*
18:30-20:00	**Welcome Reception with a View – ERSTA Conference Centre**
20:00	**Young EHTG – Meet & Great**
	EVENING for free disposition

Notes

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Scientific Programme

Day 2 – Saturday, August 29

08:00-08:30 **ILSD Business Meeting**
Chairs: Christina Therkildsen, Denmark
Gabriel Capella, Spain

08:30-09:30 **Secondary Use of Cancer Health Data in Europe**
Chairs: Johannes Blom , Sweden
Asaf Maoz, USA
Rakefet Chen-Shtoyerman, Israel

08:30-08:50 **Managing hereditary cancer registries**
Christina Therkildsen, Denmark
John Gásdal Karstensen, Denmark

08:50-09:30 **Study Proposals / Abstracts – ILSD / other EU Collaborations**

8'+2' **Extrapolating the CanRisk Framework to Hereditary Colorectal Cancer (CRC) and Polyposis: A Collaborative Proposal**
Sanne Bajwa-ten Broeke, The Netherlands

8'+2' **Possible Period Bias in Cancer Incidence Rates in the Prospective Lynch syndrome Database Following the Introduction of Amsterdam Criteria**
Kalle Hokkanen, Finland

8'+2' **Familial Colorectal Cancer History and Participation in Organised Stool-Based Colorectal Cancer Screening in Sweden: The FENCE Register-Linked Cohort Study**
Alexander Frank, Sweden

8'+2' **Novel methodologies for Lynch syndrome registry research in Finland**
Piia Takabe, Finland

09:30-10:00 **Coffee Break**

Scientific Programme

Day 2 – Saturday, August 29

10:00-11:15 **New Hereditary Gastrointestinal Cancer Syndromes / Genetics Session**
Chairs: Ingrid Winship, Australia
Kristina Lagerstedt-Robinson, Sweden
Romy Walker, Australia

ABSTRACTS

8'+2' **A Founder ATM Pathogenic Variant Among Ashkenazi Jews: Cancer Risk and Clinical Implications**
Yael Goldberg, Canada

8'+2' **Early-Onset Cancers Without Familial Aggregation Reveal Significant Germline Pathogenic Variants**
Litika Vermani, India

8'+2' **Germline KIT Mutation in a Norwegian Family with GIST: Predictive Testing and Follow up**
Vibeke Wethe, Norway

8'+2' **Breast Cancer Phenotypes in Carriers of Pathogenic POT1 Variants**
Tal Hadar, Israel

8'+2' **Cumulative Incidence of First Cancer Severely Underestimates Lifetime Cancer Burden in MLH1 and MSH2, but not in MSH6 or PMS2, Lynch syndrome Carriers**
Kalle Hokkanen, Finland

8'+2' **Timing of Germline Testing and Stage of Colorectal and Endometrial Cancer in Lynch syndrome Carriers: a Retrospective Cohort Study Using Linked Electronic Health Records**
Helen White, United Kongdom

8'+2' **The Genetic Landscape of Colon Cancer Susceptibility Genes Among the Israeli Arab Population**
Yael Goldberg, Israel

Scientific Programme

Day 2 – Saturday, August 29

11:15-12:00 **Hereditary Colorectal Cancer: Mucosal Immunology and Vaccines**
Chairs: Aysel Ahadova, Germany
Matthias Kloor, Germany
Hugo Montemont, Germany

11:15-11:35 **Vaccination in Lynch syndrome: Clinical Feasibility and Promising Early Outcomes**
Jolanda de Vries, The Netherlands

11:35-12:00 ABSTRACTS

6'+2' **Clinicopathological and Molecular Characteristics of Double Somatic DNA Mismatch Repair Gene Mutation Cases: a Systematic Review and Meta-Analysis**
Romy Walker, Australia

6'+2' **The Immunological Environment and Tumor Evasion in Lynch syndrome-associated Urothelial Tumors**
Maria Rasmussen, Denmark

6'+2' **Comprehensive Peripheral Immunophenotyping Analyses Reveal Systemic Immune Activation in Hereditary Cancer Predisposition Syndromes**
Vince Kornél Grolmusz; Hungary

12:00-13:00 **Lunch Break**

13:00-14:30 **Rare Cancers in Hereditary Gastrointestinal Syndromes. Clinical and Molecular Session**
Chairs: Elizabeth Half, Israel
Yael Goldberg, Israel
Henriett Butz, Hungary

13:00-13:20 **Infrequent Cancers in Lynch syndrome, with a Focus on Glioma**
Patrick Benusiglio, France

Scientific Programme

Day 2 – Saturday, August 29

13:20-14:30 ABSTRACTS

6'+2' **Higher-than-expected Prevalence of Germline PTEN Variants Reveals Attenuated PTEN Hamartoma Tumour Syndrome Phenotypes in Cancer Patients**
Henriett Butz, Hungary

6'+2' **Integrative Analysis of Plasma cfDNA Features in Endometrial Cancer and Benign Endometrial Pathology**
Jakub Styk, Slovakia

6'+2' **Risk Factors for Gastric Cancer in Patients with Lynch syndrome: A Systematic Review and Meta-analysis**
Daniela Silva, Portugal

6'+2' **Overall Survival of Individuals with Pancreatic Adenocarcinoma Harboring Pathogenic Germline Variants in BRCA1, BRCA2, PALB2 or ATM**
Asaf Maoz, USA

6'+2' **Non-Invasive Urine Based Detection of Mismatch Repair Deficiency in Endometrial Cancer**
Rachel Phelps, United Kingdom

6'+2' **Mismatched Lynch Syndrome Diagnostics; Suboptimal Attrition Challenges Benefit from Universal Screening for Defective Mismatch Repair in Colorectal Cancer**
Lovisa Lesand, Sweden

14:30-15:00 **Coffee Break**

15:00-15:50 **Chemoprevention**
Chairs: Ann-Sofie Backman, Sweden
Maria Rasmussen, Denmark
Kelly Kohut, United Kingdom

Scientific Programme

Day 2 – Saturday, August 29

15:00-15:30 **Aspirin in Lynch syndrome — Standard of Care or Premature?**
Tim Bishop, United Kingdom
Peter Bauerfeind, Switzerland

15:30-15:50 **Abstract/Proposal/Ongoing Studies – Preventive Medicine**

6'+2' **Feasibility and Preliminary Results of an Electronic Dietary Assessment eHealth Intervention in Patients at Increased Hereditary/Genetic Gancer Risk**
Michael Hall, USA

6'+2' **How are Patients with Lynch syndrome Informed about Aspirin Chemoprevention by Genetic Counselors, and what Factors Influence Communication Behaviors?**
Michael Hall, USA

5' **Space for study proposal/Discussion**

15:50-16:00 **Short Break**

16:00-16:30 **Dynamic EHTG Guidelines – Controversial and Ethical Issues in Development of Guidelines in Hereditary Syndromes**
Chairs: Maartje Nielsen, The Netherlands
Kaisa Fritzell, Sweden
Litika Vermani, India

16:00-16:15 **Overview of Existing Guidelines – Which Guidelines currently exist? – Where do they overlap or conflict?**
Maartje Nielsen, The Netherlands

16:15-16:30 **How Guidelines Are Made – The UK Approach (example MUTYH testing guideline)**
Terri McVeigh, United Kingdom

Scientific Programme

Day 2 – Saturday, August 29

16:30-17:30 **Interactive Discussion & Thematic Debate: "What Do We Lack, What Do We Need, and Who Should Lead?"**
Chairs: Maartje Nielsen,The Netherlands
Kaisa Fritzell, Sweden

Maartje Nielsen, The Netherlands
Kaisa, Fritzell, Sweden
Gabriela Möslein Germany
Anna Rosen, Sweden
Kelly Kohut, United Kingdom
Terri McVeigh, United Kingdom
Gustav Silander, Sweden

18:15 **Transfer to Dinner at Djurgården**

19:30 **Social Event: Junibacken, Djurgården**

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Scientific Programme

Day 2 – Saturday, August 29

Parallel Working Groups – Cederschiöldsalen

08:30-10:00	**Research Collaborative Workshop: Polyposis Syndromes** *Chair: Kaisa Fritzell*
08:30-08:40	**Brief Description on PROM and PREM – How to use and what to do with the results.** *Kaisa Fritzell*
08:40-08:55	**Presentation of the Swedish aAdaptation of the Genetic Counseling Outcome Scale-24 (GCOS-24)** *Rebecka Pestoff, Sweden*
08:55-09:10	**Experience of using GCOS-24 in the Clinic** *Lisa Åkerman, Sweden* *Ann Lindén, Sweden*
09:10-09:20	**Results from the Study: "Mental- and Physical Health, and General Well-being in Patients with Polyposis Syndromes: a Scoping Review** *Sophie Walton Bernstedt, Sweden*
09:20-09:30	**Presentation of a Research Project Aiming to Develop and Implementing Polyposis Syndrome Specific PROMs in Sweden – Looking for Collaborators** *Kaisa Fritzell, Sweden*
09:30-09:40	**Patients' Perspective on what is Important to Measure when Having a Polyposis Syndrome?** *Emil Willman, Sweden* *Samantha Hermannsson, Sweden*
09:40-10:00	**General Discussion: Do we Need to Collect Data in a Structured Way with PROM/PREM?** *Georgina Hoffmann, Germany* *Nicola Reents, Germany*

Scientific Programme

Day 2 – Saturday, August 29

Parallel Working Groups – Cederschiöldsalen

10:00-12:00	**Endoscopy Session** FAP and Lynch** *Chair: Maria Elmberg, Sweden* *Elizabeth Half, Israel* *Peter Thelin-Schmidt, Sweden*
10:00-10:15	**Endoscopic Management of duodenal polyposis and gastric lesions in FAP. What to remove, when to remove and how should it be done?** *John Gásdal Karstensen, Denmark*
10:15-10:25	**When the endoscopist had enough, aspects of surgical management in duodenal polyposis and gastric lesions** *NN*
10:30-10:40	**Endoscopic management of polyposis in FAP, lower GI** *John Gasdal Karstensen, Denmark*
10:40-11.00	**Surgical aspects in FAP, lower GI, where are we now?** *David Ljungman, Sweden*
11:05-11:15	**Upper endoscopic Surveillance in Lynch syndrome, time to individualize?** *Lars Joachim Lindberg, Denmark*
11:15-11:25	**Lower endoscopic surveillance in Lynch syndrome, aspects of quality, frequency and organization** *Lars Joachim Lindberg, Denmark*
10:25:11:40	**Fullthickness resection eFTR in hereditary cancer** *Peter Bauerfeind, Switzerland*
11:40-11:50	**The surgeons role in Lynch syndrome** *David Ljungman, Sweden*
11:50-12:00	**Discussion** *ALL ABOVE*

Scientific Programme

Day 3 – Sunday, August 30

08:30-09:00	**EHTG Business Meeting** *Chairs: Toni Seppälä, Finland* *Gabriela Moeslein, Germany*
09:00-10:00	**Changes in Population Screening and the Consequences for Patients with an Increased Risk** *Chairs: Johannes Blom, Sweden* *Kalle Hokkanen, Finland*
09:00-09:10	**The Swedish Cancer Strategy, Screening and Surveillance** *Mef Nilbert, Sweden*
09:10-09:20	**The Finnish cancer strategy, screening and surveillance** *Timo Nykopp, Finland*
09:20-09:30	**Discussion**
09:30-10:00	ABSTRACTS
8'+2'	**Evaluation of Upper Gastrointestinal Tract Surveillance in Individuals with Lynch syndrome (EARLY): An International, Multicenter Registry** *Katrin van Beekum, Germany*
8'+2'	**Reflex CRC Initiative - Advancing Colorectal Cancer Care through Comprehensive Genetic Testing** *Christoforos Pappas, Sweden*
8'+2'	**Pap Smear Liquid Biopsy Enables Non-invasive Detection of Microsatellite Instability in Endometrial Cancer** *Alexandre Perrier, France*
10:00-10:30	**Coffee Break**
10:30-11:45	**Ideas for the Future and New Developments** *Chairs: Sanne Bajwa-ten Broeke, The Netherlands* *Patrick Benusiglio, France* *Rachel Phelps, United Kingdom*

Scientific Programme

Day 3 – Sunday, August 30

10:45-11:30	ABSTRACTS
8'+2'	**Correlation of the Mismatch Repair-deficient Crypt foci Density with Clinical Patient Characteristics** *Hugo Montemont, Germany*
8'+2'	**Classification of MSH6 variants of uncertain significance (VUS) using high-throughput CRISPR functional analysis for improved precision genomic medicine** *Signe Grue Andreassen, Denmark*
8'+2'	**Histopathological, Epidemiological, and Molecular Profiling of PMS2 Lynch Syndrome Adenomas** *Katarina Andini, Netherlands*
8'+2'	**The Interplay Between Aging, Physical Activity and Immune Cells in Cancer Risk Regulation in Lynch syndrome** **Tiina Jokela, Finland**
5'	**Time for study proposal/late breaking abstract**
11:30-11:45	**Discussion**
11:45-12:15	**Heading Forward Together. Summary, Conclusion and Final Remarks** *Chairs: Toni Seppälä, Finland* *Ann-Sofie Backman, Sweden* *Maartje Nielsen, The Netherlands*
	Panel Discussion, with: *Annika Lindblom, Sweden* *Aysel Ahadova, Germany* *Yael Goldberg, Canada* *David Ljungman, Sweden* *Patrick Benusiglio, France* *Maria Rasmussen, Denmark* *Kaisa Fritzell, Sweden*
12:30	**Farewell / Adjourn** *Toni Seppälä, Finland* *Gabriela Möslein, Germany*

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Faculty

Ahadova, Aysel p. 12, 20, 21, 27	Heidelberg University Hospital Heidelberg, Germany
Åkerman, Lisa p. 24	Sweden
Andini, Katarina p. 27	University Medical Centre Groningen Groningen, Netherlands
Andreassen, Signe Grue p. 13, 26	Rigshospitalet Copenhagen, Denmark
Backman, Ann-Sofie p. 12, 21, 27	Karolinska Institute Stockholm, Sweden
Bajalica Lagercrantz Svetlana p. 13	Karolinska Institute Stockholm, Sweden
Bajwa-ten Broeke, Sanne p. 13, 14, 18, 27	University Medical Center Groningen Groningen, The Netherlands
Bauerfeind, Peter p. 22	Hirslanden Hospital Luzern/Zürich, Switzerland
Benusiglio, Patrick p. 20, 27	Sorbonne, Hôpitaux Pitié-Salpêtrière Paris, France
Bishop, Tim p. 21	University of Leeds Leeds, United Kingdom
Blom, Johannes p. 18, 26	Karolinska Institute Stockholm, Sweden
Brinch, Heidi p. 15	Copenhagen University Hospital Copenhagen, Denmark
Butz, Henriett p. 20, 21	National Institute of Oncology Budapest, Hungary
Capella, Gabriel p. 18	Catalan Institute of Oncology IDIBELL Hospitalet de Llobregat, Spain
Chen-Shtoyerman, Rakefet p. 18	Kaplan Medical Center Rehovot, Israel
de Vries, Jolanda p. 20	RadboudUMC Nijmegen, The Netherlands

Faculty

Elmberg, Maria p. 25	Sweden
Frank, Alexander p. 16, 18	Karolinska Institute Stockholm, Sweden
Fritzell, Kaisa p. 13, 22, 23, 24, 27	Karolinska Institute Huddinge, Sweden
Goldberg, Yael p. 19, 20, 27	Rabin Medical Center Princess Margaret Hospital Toronto, Canada
Gorini, Alberto p. 14	Fondazione IRCCS Istituto Nazionale dei Tumori Milan, Italy
Grohs, Laura p. 15	University of Bonn, Medical Faculty Bonn, Germany
Grolmusz, Vince Kornél p. 20	National Institute of Oncology Budapest, Hungary
Hadar, Tal p. 19	Hadassah Medical Center Jerusalem, Israel
Half, Elizabeth p. 15, 20, 21, 25	Rambam Health Care Campus Haifa, Israel
Hall, Michael p. 22	Fox Chase Cancer Center Philadelphia, USA
Hermannsson, Samantha p. 24	Sweden
Hoffmann, Georgina p. 24	SemiColon Schorndorf, Germany
Hokkanen, Kalle p. 15, 18, 19, 26	Tampere University and Tays Cancer Centre Tampere, Finland
Jokela, Tiina p. 18	University of Jyväskylä Jyväskylä, Finland
Karstensen, John Gásdal p. 18, 25	Amager og Hvidovre Hospital Copenhagen, Denmark

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Faculty

Kloor, Matthias p. 20	Heidelberg University Hospital Heidelberg, Germany
Kohut, Kelly p. 23	University of Exeter Medical School Exeter, United Kingdom
Lagerstedt Robinson, Kristina p. 15, 19	Karolinska Institute Sweden Stockholm, Sweden
Lesand, Lovisa p. 21	Lund University Lund, Sweden
Lindberg, Lars Joachim p. 24, 25	Hvidovre Hospital Copenhagen, Denmark
Lindblom, Annika p. 27	Karolinska Institute Stockholm, Sweden
Lindén, Ann p. 24	Stockholm, Sweden
Lindholm Larsson, Sara p. 14	Karolinska Institute Stockholm, Sweden
Ljungman, David p. 14, 27	Sahlgrenska University Hospital Göteborg, Sweden
Löhr, Matthias p. 12	Karolinska Institute Stockholm, Sweden
Maoz, Asaf p. 18, 21	Harvard Medical School, Dana-Farber Cancer Institute Boston M.A., USA
Marin Grez, Leon p. 14	University Hospital Heidelberg, Institute of Pathology Heidelberg, Germany
McVeigh, Terri p. 22, 23	Royal Marsden Hospital NHS Trust Lonen, United Kingdom
Möslein, Gabriela p. 23, 26	Evangelisches Bethesda Krankenhaus Duisburg, Germany
Montemont, Hugo p. 20, 27	University Hospital Heidelberg, Institute of Pathology Heidelberg, Germany

Faculty

Nielsen, Maartje p. 12, 22, 23, 27	Leiden University Medical Center Leiden, The Netherlands
Nieminen, Taina p. 15	University of Helsinki Helsinki, Finland
Nilbert, Mef p. 14, 26	Lund University Lund, Sweden
Nilsson, Per p. 14	Karolinska Institute Stockholm, Sweden
Nykopp, Timo p. 26	University of Eastern Finland Kuopio, Finland
Pappas, Christoforos p. 26	Karolinska Institutet, Department of Molecular Medicine and Surgery Stockholm, Sweden
Perrier, Alexandre p. 27	Sorbonne Université, INSERM Paris, France
Pestoff, Rebecka p. 24	Linkoping, Sweden
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We look forward to seeing you next year in Edinburgh

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