



10TH ANNUAL MEETING
Stockholm • 2026



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

Friday, August 28 to Sunday, August 30, 2026
Stockholm, Sweden

Preliminary 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

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:
 We will apply for certification by EACCME

Content

Invitation	2
Organisation	3
Meeting Overview Friday	6
Meeting Overview Saturday	8-10
Meeting Overview Sunday	11
Day 1 – Friday, September 28	
Site Visit Karolinska University Hospital	12
Registration	12
Welcome to EHTG 2026	12
New Developments in Preventive Measures and Treatment Options	12-13
Novel Approaches to Genetic Counselling	13-14
A paradigm shift in rectal cancer management	14-15
Scientific Speed-Dating	15-16
Welcome Reception	16
Young EHTG – Meet Great	16
Day 2 - Saturday 29	
ILSD Business Meeting	18
Secondary Use of Cancer Health Data in Europe	18
New Hereditary Gastrointestinal Cancer Syndromes / Genetics Session	19
Hereditary Colorectal Cancer: Mucosal Immunology and Vaccines	20
Rare cancers in Hereditary Gastrointestinal Syndromes. Clinical and Molecular Session-	21
Chemoprevention	21-22
Dynamic EHTG guidelines – Controversial and Ethical Issues in Development of Guidelines in Hereditary Syndromes	22
Interactive Discussion & Thematic Debate: “What Do We Lack, What Do We Need, and Who Should Lead?”	23
Social Event: Junibacken, Djurgården	23

Content

Parallel Working Groups

Research Collaborative Workshop: Polyposis Syndromes	24
Endoscopy Session** FAP and Lynch	25
Day 3 - Sunday 30	
EHTG Business Meeting	26
Changes in Population Screening and the Consequences for Patients with an Increased Risk	26
Ideas for the Future and New Developments	26-27
Heading Forward Together. Summary, Conclusion and Final Remarks	27
Faculty	28-33
Sponsors	34
Imprint	35

Meeting Overview

Day 2 – Saturday, August 29

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

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/Norrbj

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 spaces)

8:30-10:00 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
Aletha Thang, United Kingdom

11:45-12:15 **Precision Medicine in Hereditary Gastrointestinal Tumors**
Richard Rosenquist, 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' **The Genetic Landscape of Colon Cancer Susceptibility Genes among the Israeli Arab Population**
Yael Goldberg, Israel

8'+2' **Cell-Free DNA in Hereditary and High-Risk Malignancies 2 (CHARM2): A Randomized Controlled Trial for Early Cancer Detection**
Sarah Ridd, Canada

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 Lagercrantz, Sweden

14:30-14:40 **Artificial Intelligence and Genetic Counseling in Hereditary Cancer Syndromes**
Elen Siglen, Norway

14:40-15:30 **ABSTRACTS**

6'+2' **Genetic Counselling Regarding Recessive Syndromes Associated with Cancer Predisposing Genes: A Consensus on Shared Decision-Making**
Judit Sanz Buxo, Spain

6'+2' **Exploring Quality of Life, Illness Representations, Coping Strategies, and Needs in Patients with APC-Related Desmoid-Type Fibromatosis**
Bianca Scacciati, Italy

6'+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

6'+2' **Implementing a Canadian Population Genomic Screening Study for Hereditary Cancers**

Laura Redondo, Canada

6'+2' **Diagnostic Value of Long-Read Sequencing in Unexplained Mismatch Repair Deficient MMR) Cancer Cases**

Sanne Baiwa-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 **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

6'+2' **Upper Gastrointestinal Neoplasia Detection in Lynch Syndrome: High Yield of Esophagogas Troduodenoscopy Surveillance**

Elizabeth Half, Israel

Scientific Programme

Day 1 – Friday, August 28

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

Kalle Hokkanen, Sweden

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

5'+2' **Haplotype GWAS identifies Sex-Specific Differences in Modifying Risk Loci in Colorectal Cancer**

Litika Vermani, India

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: NN
Johannes Blom, Sweden
Asaf Maoz, USA

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 Baiwa-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' **The Interplay Between Aging, Physical Activity and Immune Cells in Cancer Risk Regulation in Lynch syndrome**
Tiina Jokela, 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

Scientific Programme

Day 2 – Saturday, August 29

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

10:00-11:15 **New Hereditary Gastrointestinal Cancer Syndromes / Genetics Session**
Chairs: Ingrid Winship, Australia
Yael Goldberg, Israel
Romy Walker, Australia

ABSTRACTS

8'+2' **A Founder ATM Pathogenic Variant Among Ashkenazi Jews: Cancer Risk and Clinical Implications**
Elizabeth Half, Israel

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 Kingdom

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
NN
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**

8'+2' **Higher-than-expected Prevalence of Germline PTEN Variants Reveals Attenuated PTEN Hamartoma Tumour Syndrome Phenotypes in Cancer Patients**
Henriett Butz, Hungary

8'+2' **Integrative Analysis of Plasma cfDNA Features in Endometrial Cancer and Benign Endometrial Pathology**
Jakub Styk, Slovakia

8'+2' **Risk Factors for Gastric Cancer in Patients with Lynch syndrome: A Systematic Review and Meta-analysis**
Daniela Silva, Portugal

8'+2' **Overall Survival of Individuals with Pancreatic Adenocarcinoma Harboring Pathogenic Germline Variants in BRCA1, BRCA2, PALB2 or ATM**
Asaf Maoz, USA

8'+2' **Non-Invasive Urine Based Detection of Mismatch Repair Deficiency in Endometrial Cancer**
Rachel Phelps, United Kingdom

8'+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
Naim Abu-Freha, Israel*

Scientific Programme

Day 2 – Saturday, August 29

15:00-15:30 **Aspirin in Lynch syndrome – Standard of Care or Premature?**

*Tim Bishop, United Kingdom
Elizabeth Half, Israel
Ann-Sofie Backman, Sweden
Maria Rasmussen, Denmark*

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

8'+2' **Feasibility and Preliminary Results of an Electronic Dietary Assessment eHealth Intervention in Patients at Increased Hereditary/Genetic Cancer Risk**

Michael Hall, USA

8'+2' **How are Patients with Lynch syndrome Informed about Aspirin Chemoprevention by Genetic Counselors, and what Factors Influence Communication Behaviors?**

Michael Hall, USA

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**

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
- 09:30-09:50 **Patients’ Perspective on what is Important to Measure when Having a Polyposis Syndrome?**
Emil Willman, Sweden
- 09:50-10:00 **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
NN
- 10:00-10:30 **Management of Duodenal Polyposis in FAP Including Surveillance and Management of Gastric Lesions**
John Gasdal Karstensen, Denmark
- 10:30-11:00 **Endoscopic Management of Polyposis in FAP, Lower GI**
John Gasdal Karstensen, Denmark
- 11:00-11:05 **Short Break**
- 11:05-11:40 **Endoscopic Surveillance in Lynch syndrome**
Lars Joachim Lindberg, Denmark
- 11:40-12:00 **Discussion**
- 12:00-13:00 **Lunch**

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, Sweden

09:00-09:30 **The Swedish Cancer Strategy, Screening and Surveillance**
Mef Nilbert, Sweden
Timo Nykopp, Finland

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' **Classification of MSH6 Variants of Uncertain Significance (VUS) Using High-throughput CRISPR Functional Analysis for Improved Precision Genomic Medicine**
Signe Grue Andreassen, Denmark

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

10:30-11:45 **Ideas for the Future and New Developments**
Chairs: Sanne Baiwa-ten Broeke, The Netherlands
Patrick Benusiglio, France
Rachel Phelps, United Kingdom

Scientific Programme

Day 3 – Sunday, August 30

10:45-11:30 ABSTRACTS

7'+2' **Novel Methodologies for Lynch syndrome Registry Research in Finland**
Piia Takabe, Finland

7'+2' **Correlation of the Mismatch Repair-deficient Crypt foci Density with Clinical Patient Characteristics**
Hugo Montemont, Germany

7'+2' **Epigenetic Treatment of Lynch syndrome-Associated Colorectal Carcinoma and BRAF-Mutant Colorectal Carcinoma**
Anna Scheßl, Germany

7'+2' **Histopathological, Epidemiological, and Molecular Profiling of PMS2 Lynch Syndrome Adenomas**
Katarina Andini, Netherlands

7'+2' **Pap Smear Liquid Biopsy Enables Non-invasive Detection of Microsatellite Instability in Endometrial Cancer**
Alexandre Perrier, France

11:30-11:45 **Discussion**

11:45-12:15 **Heading Forward Together. Summary, Conclusion and Final Remarks**
Chairs: Annika Lindblom, Sweden
NN

Panel Discussion, with:
NN

12:30 **Farewell / Adjourn**

Faculty

Abu-Freha, Naim	Israel
Ahadova, Aysel	Heidelberg University Hospital Heidelberg, Germany
Andini, Katarina	University Medical Centre Groningen Groningen, Netherlands
Andreassen, Signe Grue	Rigshospitalet Copenhagen, Denmark
Backman, Ann-Sofie	Karolinska Institute Stockholm, Sweden
Baiwa-ten Broeke, Sanne	University Medical Center Groningen Groningen, The Netherlands
Benusiglio, Patrick	Sorbonne, Hôpitaux Pitié-Salpêtrière Paris, France
Bishop, Tim	University of Leeds Leeds, United Kingdom
Björk, Jan	Karolinska Institute Stockholm, Sweden
Blom, Johannes	Karolinska Institute Stockholm, Sweden
Brinch, Heidi	Copenhagen University Hospital Copenhagen, Denmark
Butz, Henriett	Semmelweis University Budapest, Hungary
Capella, Gabriel	Catalan Institute of Oncology IDIBELL Hospitalet de Llobregat, Spain
de Vries	Jolanda RadboudUMC Nijmegen, The Netherlands
Frank, Alexander	Karolinska Institute Stockholm, Sweden
Fritzell, Kaisa	Karolinska Institute Huddinge, Sweden
Goldberg, Yael	Rabin Medical Center Princess Margaret Hospital Toronto, Canada

Faculty

Gorini, Alberto	Fondazione IRCCS Istituto Nazionale dei Tumori Milan, Italy
Grohs, Laura	University of Bonn, Medical Faculty Bonn, Germany
Grolmusz, Vince Kornél	National Institute of Oncology Budapest, Hungary
Hadar, Tal	Hadassah Medical Center Jerusalem, Israel
Half, Elizabeth	Rambam Health Care Campus Haifa, Israel
Hall, Michael	Fox Chase Cancer Center Philadelphia, USA
Hoffmann, Georgina	SemiColon Schorndorf, Germany
Hokkanen, Kalle	Tampere University and Tays Cancer Centre Tampere, Sweden
Jokela, Tiina	University of Jyväskylä Jyväskylä, Finland
Karstensen, John Gásdál	Amager og Hvidovre Hospital Copenhagen, Denmark
Kloor, Matthias	Heidelberg University Hospital Heidelberg, Germany
Kohut, Kelly	University of Exeter Medical School Exeter, United Kingdom
Lagercrantz, Svetlana	Karolinska Institute Stockholm, Sweden
Lagerstedt Robinson, Kristina	Sweden
Lesand, Lovisa	Lund University Lund, Sweden
Lindberg, Lars Joachim	Hvidovre Hospital Copenhagen, Denmark
Lindblom, Annika	Karolinska Institute Stockholm, Sweden

Faculty

Lindén, Ann	Stockholm, Sweden
Ljungman, David	Sahlgrenska University Hospital Göteborg, Sweden
Löhr, Matthias	Karolinska Institute Stockholm, Sweden
Maoz, Asaf	Harvard Medical School, Dana-Farber Cancer Institute Boston M.A., USA
Marin Grez, Leon	University Hospital Heidelberg, Institute of Pathology Heidelberg, Germany
McVeigh, Terri	Royal Marsden Hospital NHS Trust Lonen, United Kingdom
Mecklin, Jukka Pekka	University of Jyväskylä Jyväskylä, Finland
Möslein, Gabriela	Evangelisches Bethesda Krankenhaus Duisburg, Germany
Montemont, Hugo	University Hospital Heidelberg, Institute of Pathology Heidelberg Germany
Nielsen, Maartje	Leiden University Medical Center Leiden, The Netherlands
Nieminen, Taina	University of Helsinki Helsinki, Finland
Nilbert, Mef	Lund University Lund, Sweden
Nilsson, Per	Karolinska Institute Stockholm, Sweden
Nykopp, Timo	University of Eastern Finland Kuopio, Finland
Pappas, Christoforos	Karolinska Institutet, Department of Molecular Medicine and Surgery Stockholm, Sweden
Perrier, Alexandre	Sorbonne Université, INSERM Paris, France

Faculty

Pestoff, Rebecka	Linköping, Sweden
Phelps, Rachel	Newcastle University Newcastle upon Tyne, United Kingdom
Phelps, Rachel	Newcastle University Newcastle upon Tyne United Kingdom
Rasmussen, Maria	Copenhagen University Hospital, The Danish HNPCC Register Copenhagen, Denmark
Rausa, Emanuele	Italy
Redondo, Laura	Princess Margaret Cancer Centre, University Health Network Toronto, Canada
Reents, Nicola	SemiColon Moenchengladbach, Germany
Ridd, Sarah	Princess Margaret Cancer Centre, University Health Network Toronto, Canada
Rosen, Anna	Umea University Umea, Sweden
Rosenquist, Richard	Karolinska Institute Stockholm, Sweden
Sanz Buxo, Judit	Althaia Xarxa Assistencial Universit�ria Manresa Manresa, Spain
Sche�l, Anna	University Hospital Heidelberg, Institute of Pathology Heidelberg, Germany
Sepp�l�, Toni	Tampere University Tampere, Finland
Siglen, Elen	Western Norway Familial Cancer Center Bergen, Norway
Silander, Gustav	Ume� University Ume�, Sweden

Faculty

Silva, Sofia	University Hospital A Coruña A Coruña, Spain
Silva, Daniela	University of Porto, Faculty of Medicine Porto, Portugal
Styk, Jakub	Comenius University Science Park Bratislava, Slovakia
Takabe, Piia	Tampere University Tampere, Finland
Tang, Aletha	United Kingdom
Tham, Emma	Karolinska Institute Stockholm, Sweden
Therkildsen, Christina	Amager og Hvidovre Hospital Copenhagen, Denmark
Usman Babagari, Musa	Modibbo Adama University Yola Nigeria, Nigeria
van Beekum, Katrin	University Hospital Bonn, National Center for Hereditary Tumor Syndromes Bonn, Germany
Vermani, Litika	Cachar Cancer Hospital & Research Centre Silchar, India
Vermani, Litika	Cachar Cancer Hospital & Research Centre Silchar, India
Vermani, Litika	Cachar Cancer Hospital & Research Centre Silchar, India
Walker, Romy	University of Melbourne Melbourne, Australia
Walton Bernstedt, Sophie	Sweden
Wethe, Vibeke	Oslo University Hospital, Department of Medical Genetics Oslo, Norway

Faculty

White, Helen	University of Manchester Manchester, United Kingdom
Willman, Emil	Sweden
Winship, Ingrid	Royal Melbourne Hospital Melbourne, Australia
Wollborn, Laura	Harvard Medical School, Dana-Farber Cancer Institute Boston, MA; USA

Sponsors



**Karolinska
Institutet**

110.800 SEK
(10.000 €)

**RADIUMHEMMETS
FORSKNINGSFONDER**

The Cancer Research Foundations of Radiumhemmet

100.000 SEK
(9.200 €)



CANCERFONDEN

100.000 SEK
(9.200 €)

SPRINGER NATURE

2 Book-Vouchers
å 250 €

Imprint

Responsible in accordance with press law:

European Hereditary Tumour Group (EHTG) e.V.
Toni Seppälä (Chair)
Weissdornstr. 6
40474 Düsseldorf
toni.seppala@ehtg.org

Organisation:

EHTG secretariat: Gabriele Sponholz, gs007@ehtg.org

Image Credits

Cover: PantherMedia (now Smartpix)
iStock
Page 9 ERSTA Conference Centre
Page 10 Junibacken



**Thank you for attending
EHTG 2026**



**We look forward to
seeing you next year
in Edinburgh
September 17-19, 2027**

