

Wednesday 20 March

1330 - 1430	PLSD Business Meeting: All interested are welcomed
1600 - 1900	Meeting registration desk open - wine and cheese served
	Public lecture: Familial Bowel Cancer Syndromes - an update and international perspective
1930 - 1935	Welcome
1935 - 1950	Update on diagnosis and management <i>Patrick Lynch</i>
1950 - 2020	Panel: What is happening in other countries? <i>Ian Frayling, Nicoline Hoogerbrugge, Finlay Macrae</i>
2020 - 2100	Discussion

Thursday 21 March

0700 - 1830	Registration desk open
0830 - 0845	Official opening <i>Ashley Bloomfield, Director General of Health</i>
0845 - 0900	Gastrointestinal hereditary tumours: what we know and what we need to know? <i>Sue Clark</i>
0900 - 0920	Cancer risks associated with MMR gene mutations - how can we define further? <i>Mark Jenkins</i>
0920 - 0950	Free papers - 5 minutes each followed by 10 minutes discussion Cancer risks by age and gender in carriers of pathogenic MMR variants: Findings from the Prospective Lynch Syndrome Database (PLSD) report <i>Mev Dominguez Valentin</i> Impact from compliance with surveillance on risk of colorectal cancer in Lynch Syndrome <i>Lars J. Lindberg</i> Breast cancer risk not increased in women with Lynch Syndrome identified by multi-gene panel testing <i>Sonia Kupfer</i> Occurrence of polyps and incident colorectal carcinomas in patients with PMS2-associated Lynch Syndrome: a prospective cohort analysis <i>Sanne Ten Broeke</i>
0950 - 1030	Free papers - 5 minutes each followed by 15 minutes discussion Molecular tumour testing in Lynch-like patients reveals de novo mosaic DNA mismatch repair gene pathogenic variants transmitted to offspring

Chrystelle Colas

Predictors of class: Using protein structure and function information to predict and understand mismatch repair variant pathogenicity

Bernard Pope

cDNA analyses of the MMR genes MLH1, MSH2, MSH6 and PMS2 investigate the effect of VUS upon splicing, detect unexpected splicing defects, and find allelic losses indicating a germline defect

Elke Holinski-Feder

Highly sensitive MLH1 methylation analysis in blood identifies a cancer patient with low-level mosaic MLH1 epimutation

Gabriel Capella

Comprehensive constitutional genetic and epigenetic characterization of Lynch-like individuals

Gabriel Capella

1030 - 1100 Morning Tea

1100 - 1125 Finding Lynch Syndrome and beyond

John Burn

1125 - 1200 **Free papers - 5 minutes each followed by 15 minutes discussion**

Reflex Mis-Match-Deficiency testing of colorectal cancer below age 70 to detect Lynch Syndrome: a prospective, multicenter, multidisciplinary evaluation of uptake, yield and appreciation

Nicoline Hoogerbrugge

UK National External Quality Assessment Scheme for Immunocytochemistry and In Situ Hybridisation: 10 years of international experience with mismatch repair proteins shows that participation improves performance

Ian Frayling

Evaluating tumour mutational signatures for classification of mismatch repair deficiency and identification of Lynch syndrome and MLH1 methylated subtypes

Peter Georgeson

RAID-LS: a non-invasive tool based on faecal bacterial signature for Lynch Syndrome surveillance

Joan Brunet

1200 - 1215 **Free papers - 5 minutes each followed by 5 minutes discussion**

Multiple Genetic Tumor Syndromes: When to suspect them?

Maurizio Genuardi

Germline pathogenic variants of hereditary cancer genes in 12,347 colorectal cancer patients and 27,706 controls in Japanese population

Hidewaki Nakagawa

1215 - 1230	Somatic mutations in colorectal cancer and implications for genetic testing <i>Gabriel Capella</i>
1230 - 1325	Lunch Break
1325 - 1350	The immune system in Lynch Syndrome/hereditary colorectal cancer <i>Magnus von Knebel Doeberitz</i>
1350 - 1430	Free papers - 5 minutes each followed by 15 minutes discussion The shared mutation and neoantigen landscape of MMR-deficient cancers suggests immunoediting during tumor evolution <i>Matthias Kloor</i> High endothelial venules are associated with immune evasion and hereditary background in microsatellite-unstable colorectal cancers <i>Aysel Ahadova</i> Germline variants associated with immune infiltration in solid tumors <i>Sahar Shahamatdar</i> Intratumoural assessment of colorectal cancer diagnostic and prognostic markers using RNA in situ hybridization <i>Tim Eglinton</i> A mouse model for a vaccine against Lynch Syndrome-associated cancers <i>Matthias Kloor</i>
1430 - 1445	Free papers - 5 minutes each followed by 5 minutes discussion Genetic Cancer Susceptibility in Adolescents and Young Adults with Colorectal Cancer <i>Richarda de Voer</i> Therapy-associated polyposis in childhood and young adulthood cancer survivors <i>Matthew Yurgelun</i>
1445 - 1510	Update on targeted therapies in Gastrointestinal Oncology <i>Michael Hall</i>
1510 - 1540	Afternoon Tea
1540 - 1605	Hereditary Diffuse Gastric Cancer (HDGC) - past, present, future <i>Parry Guilford</i>
1605 - 1615	Genetic counselling in HDGC - the bi-cultural context <i>Kim Gamet</i>
1615 - 1630	Challenges in genetic counselling - case discussions <i>Kim Gamet</i>
1630 - 1655	Free papers - 5 minutes each followed by 10 minutes discussion Early Genetic Counseling and Detection of CDH1 Mutation in Asymptomatic Carriers Improves Survival in Hereditary Diffuse Gastro Cancer <i>R Matthew Walsh</i>

CDH1 Gastric Cancer: Does Family History Change Your Risk?
Margaret O'Malley

Multiple-gene panel analysis in an Italian cohort of patients with familial gastric cancer

Gianluca Tedaldi

1655 - 1710 **Free papers - 5 minutes each followed by 5 minutes discussion**

Technical and Endoscopic Factors in CDH1 gastric cancer surveillance

Carol Burke

Gastroscopic outcomes compared with histology in CDH1 mutation carriers; 9 years experience with the International Gastric Cancer Linkage Consortium Consensus Guideline

Jolanda van Dieren

1710 - 1725 Gastroscopy in HDGC - an update

Massimiliano Di Pietro

1725 - 1740 Guideline update - hot news from Wanaka

Nicoline Hoogerbrugge

1745 Leave venue to walk to Welcome Reception

1800 - 2000 Welcome Reception - Maritime Museum

2000 Council Dinner

Friday 22 March

0730 - 0830 CaPP3 Collaborators meeting (invitation only)

0830 - 0850 New colorectal cancer genes: one big happy family?

Ian Frayling

0850 - 0905 Genetics of Serrated Polyposis Syndrome (SPS)

Dan Buchanan

0905 - 0930 **Free papers - 5 minutes each followed by 10 minutes discussion**

Colorectal cancer risk in NTHL1 heterozygous mutation carriers

Abi Raganathan

Germline POLE and POLD1 variation in persons with colorectal cancer from the Colon Cancer Family Registry Cohort

Khalid Mahmood

Variant profiling of colorectal adenomas from patients with MSH3-related adenomatous polyposis

Claudia Perne

0930 - 0955 **Free papers - 5 minutes each followed by 10 minutes discussion**

Patient derived intestinal mucosal organoids: a new technology to study

	pathogenesis in familial adenomatous polyposis <i>Roshani Patel</i>
	Pathogenic variants in new colorectal cancer/polyposis genes rarely identified among patients with colorectal, breast, prostate, and pancreatic cancer <i>Brandie Heald Leach</i>
	Exome sequencing identified potential causative candidate genes for serrated polyposis syndrome <i>Sophia Peters</i>
0955 - 1015	Pathogenesis of Colorectal Cancer in SPS <i>Christophe Rosty</i>
1015 - 1045	Morning Tea
1045 - 1100	Clinical interpretation of genetic variants in hereditary GI Cancer: Where we are and where do we go? <i>Maurizio Genuardi</i>
1100 - 1115	The InSiGHT Database – continuing the mission of centralising variants of the GI Cancer genes <i>John Paul Plazzer</i>
1115 - 1155	Free papers - 5 minutes each followed by 15 minutes discussion
	Interpretation of inheritable DNA variation: How much room for error across genetic services? <i>Matthew Daly</i>
	Variant analyses of PMS2 by Single-Molecule Long-Read Sequencing <i>Richarda de Voer</i>
	Curation and classification of Adenomatous Polyposis Coli (APC) gene variants responsible for familial adenomatous polyposis (FAP) in ClinVar and the International Society for Gastrointestinal Hereditary Tumours (InSiGHT) locus-specific database <i>Xiaoyu Yin</i>
	Splicing Effects and In Silico Pathogenicity Predictions For APC Missense Variants Reported in ClinVar <i>Marc Greenblatt</i>
	The detection of hybrid mosaic mutations during analysis for APC mosaicism <i>Manon Suerink</i>
1155 - 1205	Genetic testing - which genes on which panel? <i>Ian Frayling, Gabriel Capella</i>
1205 - 1230	Panel Discussion <i>Heather Hampel, Ian Frayling (Panel Co-ordinator)</i>
1230 - 1330	Lunch and Asia Pacific Meeting
1330 - 1350	What constitutes good colonoscopy and gastroscopy in Lynch Syndrome and FAP <i>Andrew Latchford</i>

1350 - 1425	<p>Free papers - 5 minutes each followed by 15 minutes discussion</p> <p>Risk of interval colorectal cancer in patients with Lynch Syndrome undergoing surveillance in New Zealand – results from the New Zealand Familial Gastrointestinal Cancer Service <i>Mehul Lamba</i></p> <p>Quality of and compliance with colonoscopy in Lynch Syndrome surveillance: Are we getting it right? <i>Karen Hartery</i></p> <p>Stage of CRC is not associated with time since last colonoscopy in Lynch Syndrome: A Prospective Lynch Syndrome Database (PLSD) report <i>Toni Seppala</i></p> <p>The impact of a risk management clinic model on surveillance and colorectal cancer incidence in patients with Lynch Syndrome <i>Andrew Buckle</i></p>
1425 - 1440	<p>Free papers - 5 minutes each followed by 5 minutes discussion</p> <p>Identifying clinical features associated with advanced gastric pathology in familial adenomatous polyposis <i>Gautam Mankaney</i></p> <p>Individualized surveillance for serrated polyposis syndrome: Results from a prospective 5-year international cohort study <i>Arne Bleikenberg</i></p>
1440 - 1500	<p>Management of SPS <i>Evelien Dekker</i></p>
1500 - 1530	Afternoon tea
1530 - 1550	<p>Management of duodenal adenomas in FAP <i>Evelien Dekker</i></p>
1550 - 1625	<p>Free papers - 5 minutes each followed by 15 minutes discussion</p> <p>The effect of endoscopic duodenal interventions in patients with familial adenomatous polyposis <i>Victorine Roos</i></p> <p>Duodenal Adenomas and Cancer in Familial Adenomatous Polyposis <i>Isabel Martin</i></p>
1625 - 1645	<p>Pancreatic screening for high risk familial syndromes <i>John Windsor</i></p>
1645 - 1730	InSiGHT Annual General Meeting
1830	Buses depart SkyCity Auckland Convention Centre for Auckland Museum
1900 - Midnight	<p>Meeting Gala Dinner- Auckland Museum</p> <p><i>Shuttle buses will depart from 2230 for the hotels</i></p>

Saturday 23 March 2019

0730 - 0830	InSiGHT Database Governance Committee Meeting
0830 - 0845	Genetic counselling - implications of panel and tumour testing <i>Heather Hampel</i>
0845 - 0915	Free papers - 5 minutes each followed by 10 minutes discussion The clinical utility and impact on risk categorisation of a lifestyle and genomic risk prediction model for colorectal cancer <i>Sibel Saya</i> Shared Medical Appointments for Lynch Syndrome: An Effective and Efficient Model for Patient Management <i>Lisa LaGuardia</i> "When do I tell my family, what do I tell them?"; the importance of psychological adaptation to a genetic diagnosis before patients are able to share information about their diagnosis – findings from the Family Web study. <i>Selina Goodman</i> Directly approaching individuals at risk of inherited colorectal cancer syndromes: The New Zealand experience <i>Julie Arnold</i>
0915 - 0930	Timing of surgery in FAP <i>Sue Clark</i>
0930 - 0945	Oligopolyposis/SPS - when is colectomy indicated <i>John Keating</i>
0945 - 1000	Proctectomy and advanced pouch adenomas in FAP <i>Matthew Kalady</i>
1000 - 1030	Free papers - 5 minutes each followed by 15 minutes discussion Indications and outcomes for pouch excision in patients with familial adenomatous polyposis (FAP) <i>Roshani Patel</i> ATZ Neoplasia: A comprehensive examination of a dangerous phenomenon <i>James Church</i> Safety and efficacy of laparoscopic near-total colectomy and ileo-distal sigmoid anastomosis as a modification of total colectomy and ileorectal anastomosis for prophylactic surgery in patients with adenomatous polyposis syndromes– a comparative study <i>Chukwuemeka Anele</i>
1030 - 1100	Morning tea
1100 - 1115	Free papers - 5 minutes each followed by 5 minutes discussion The Impact of Desmoid Tumors on Quality of Life and Pouch Survival, in

	patients with Familial Adenomatous Polyposis who have undergone Ileal Pouch-Anal Anastomosis <i>James Church</i>
	Laparotomy results in more desmoid tumour when compared to laparoscopy in a preclinical model of desmoid tumour in familial adenomatous polyposis <i>Timothy Chittleborough</i>
1115 - 1145	Case discussion - Panel <i>Christopher Wakeman (Panel Coordinator)</i>
1145 - 1155	New considerations in surgery for Lynch Syndrome? <i>Ian Bissett</i>
1155 - 1210	Extensive Surgery in LS - factoring in genes and gender <i>Gabriela Moslein</i>
1210 - 1220	Lynch Syndrome - segmental colectomy and aspirin <i>John Burn</i>
1220 - 1230	Discussion
1230 - 1250	Hereditary Colorectal Cancer 1989 - 2019: Perspectives from the past and predictions for the future <i>James Church</i>
1250 - 1305	InSiGHT 2021 and close of meeting
1305 - 1310	Poroporoaki <i>All delegates to remain seated</i>
1310 - 1345	Lunch
1345 - 1400	Walk to Waiheke Island Ferry
1430 - 1930	Waiheke Island Trip (optional)