

ICGCW 2025 7th INDIAN CANCER GENETICS CONFERENCE & WORKSHOP Tata Memorial Hospital & ACTREC, Mumbai, India



ICGC2025 CONFERENCE TMH, PAREL: 22-23 Feb '25 (Saturday & Sunday)

THREE PRECONFERENCE HANDS ON WORKSHOPS 20-21st Feb 2025 ❑Genetic Counselling of Common Hereditary Cancers: TMH, Mumbai ❑NGS Wetlab, DryLab & Reporting: ACTREC, Navi Mumbai ❑Functional Genomics & Tissue Culture, ACTREC, Navi Mumbai

Deep Dive To know More about Less Known Genes Navigating The Rising Tide of VUS



- Should Germline or Somatic report include these genes / variants ??
- How should they be described ?
- How to act (or not) on these ?

Registration Link for a Deep Dive to learn together & reach consensus

ALL ABOUT LESS KNOWN or MODERATE RISK GENES, HYPOMORPHIC ALLELES & VUS

TMC Cancer Genetics Unit has CURATED germline genotype & phenotype data of 10,000+ cases - not only well characterised genes like BRCA1/2, MLH1, MSH2, TP53, PALB2) but also Less known genes (CHEK2, BRIP1, RAD51, CDH1, MSH6, PMS2, SDH, & many more We have reclassified hundreds of ClinVar VUS as Likely Benign & a few as Likely Pathogenic

Registration fee (with 18% GST) *Will be Half for Trainees & GCs & *Double for Industry delegates	Early Bird 31-12.2024	Regular 31-1-2025	SPOT REG Feb 2025
Conference Only	6000	7500	9000
Conference + GC Workshop	8000	9500	12000
Conference + 1 Lab Workshop NGS or Functional Genomics	10000	12000	If available

DATES TO REMEMBER → EARLY BIRD: 31/12/2024

→ ABSTRACT: 05/02/2025 Mail 1 page PDF abstract to icgcw.tmc@gmail.com Best Paper Award for One abstract each in

- Clinical Genetics
- Genetic Counselling
- Molecular Genetics

**Abstracts eligible for fee waiver

*May Request Reg Fee Refund for abstract with Genotype-Phenotype description of >10 cases of less known genes or VUS

<< <u>CLICK TO REGISTER</u> >>

Email icgcw.tmc@gmail.com

ail.com Phone +91-22-27405000 ext. 5318 M Registration Link: https://forms.gle/5H6csUvmQ1NvkYUx5

Mobile: +91-8433733212

PATRON	CHAIRPERSON	ORGANIZING SECRETARY	TREASURER		
Prof. Sudeep Gupta	Prof. Rajiv Sarin	Dr Poonam Gera	Dr Somya Srivastava		

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Less Known or Moderate Risk Genes, Hypomorphic Alleles & VUS

- Why are they tested in the first place?
- Should they be in the report & how should they be described?
- Genotype-Phenotype uncertainty of these genes or variants
- Family & functional studies to characterise these variants
- Pitfalls in ignoring or acting upon these genes / variants
- My RARE GENE DIARY of Low to Moderate Cancer Risk Genes

Three Pre Conference Workshops

1)CANCER GENETIC COUNSELLING

• Genetic counselling overview

20-21 FEB. 2025

PRE CONFERENCE HANDS ON WORKSHOPS

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Only 40 participants

- Drawing informative complex pedigrees
- Psychosocial issues in genetic counselling
- Counselling for VUS & Moderate penetrance genes
- Counselling for HBOC, Lynch, Li-Fraumeni, NF, FAP, MEN2, Cowden, RB1, Dicer, WT1 & other syndromes
- Prenatal & Pre-Implantation genetic testing
- Case based discussions & role plays

2) NGS WET LAB, DRY LAB & REPORTING

- Library prep Targeted Panel & Exome
- Setting up NGS run on MiSeq & NextSeq
- Bioinformatics fastq to vcf, genomic databases
- Variant annotation, clinical report drafting on ACMG guidelines. Case based discussions on imp. aspects
- Sanger sequencing, Long Range PCR & Digital MLPA

3) FUNCTIONAL GENOMICS & TISSUE CULTURE

- Functional Genomics Overview
- Mammalian Cell Culture Techniques
- Gene Editing Technologies CRISPR/Cas9
- Functional Assays in Tissue Culture
- DNA Damage Repair and Genomic Integrity
- Apoptosis and Cell Survival Assays

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AI TMH, PAREL MUMBA

Only 24 participants

Only 14 participants

Pre Conference Workshop 1 Cancer Genetic Counselling in Practice

(TMH)

Thursday & Friday: 20th & 21st February,2025 Venue: Choksi Auditorium, Tata Memorial Hospital, Parel, Mumbai

Who should attend: Genetic Counsellors, Oncologists, Gynecologists, Preventive Medicine Faculty: Rajiv Sarin, Somya Srivastava, Neena Bhatnagar, Deepali Kapoor, Aparna Dhar, Deepanjana Dutta, Yoon Sook Yee

Instructors: TMC Clinical Genetic & Counselling Team

Format: Faculty Talks, Case Based Interactive Discussions & Role Plays

Day 1 (10 am - 6:00 pm)

- 1. Overview of Cancer Epidemiology, risk factors & Cancer Genetics: Risk factors for cancers, Global burden of Cancer & Hereditary Cancers
- 2. Cancer Genetic Counselling in Indian Practice Setting & Barriers
- 3. Taking Personal & Family history, Phenotyping, Syndromic Diagnosis & drawing complex pedigrees
- 4. Navigating Medical Records of relevance and Verbal Autopsy techniques for cancer
- 5. Knudson's model for suspecting inherited predisposition to cancer: Concept and case scenarios
- 6. Choosing the right test and panel- Germline, Somatic or Both and their Sequence
- 7. Do's & Don'ts in Pretest and Post Test counselling: Theoretical framework and real life scenarios
- 8. Minimizing Pre-analytical errors, Consents & disclaimers: Systematic approach to spot the red flags.
- 9. Navigating through Genomic Databases to improve reporting and improve public health
- **10.** Interpreting Genetic Test Reports
- 11. Dealing with VUS in the Clinic

Day 2 (9 am - 5:30 pm)

- 1. HBOC syndrome, BRCA1/2 & other HRR Genes
- 2. Lynch Syndrome, MMR Genes
- 3. Li-Fraumeni Syndrome, TP53
- 4. MEN1, MEN2 & PPGL syndromes
- 5. NF1 & NF2 Syndrome
- 6. Cowden & Peutz Jegher Syndrome
- 7. DICER, WAGR and other syndromes, genes
- 8. Screening / Preventive Surgery for Ovary & Endometrial Cancer Risk
- 9. Screening / Preventive Surgery for Breast Cancer Risk
- **10.** Chemoprevention, Endoscopic management & Surgical Prevention of colorectal cancer risk
- 11. Reporting & counseling for moderate penetrance genes
- 12. Ethical Legal and Psychosocial Issues in Genetic Counselling
- 13. Feedback, Post Workshop Evaluation & Group Photo

Pre Conference Workshop 2 NGS & Digital MLPA - Wet Lab, Dry Lab & Reporting (ACTREC)

Thursday & Friday: 20th & 21st February,2025 Venue: Genomics Lab, CRI, ACTREC, Kharghar, Navi Mumbai

Who should attend: Genome analysts, Molecular Biologists, Molecular Pathologists, Clinicians, Genetic Counsellors, PhD students

Faculty: Rajiv Sarin, Poonam Gera, Somya Srivastava, Nikhil Gadewal

Instructors: TMC Genomics Lab & Reporting Team

Format: Faculty Talks, NGS Wet Lab, Sanger / MLPA Overview & Brief Demonstration,

Interrogating Database & Drafting Reports, Case Based Interactive Discussions & Troubleshooting [BRING YOUR LAPTOPS, IF POSSIBLE]

Day 1 (10 am - 6:30 pm)

- 1. Overview of somatic & Germline testing & common hereditary cancer syndrome
- 2. DNA extraction on Semi-Automated platforms & QC
- 3. NGS Hands-on workshop (in 2 batches)
- 4. Primer Designing & Troubleshooting for PCR
- 5. MLPA (including Digital MLPA): Principles, steps & report analysis
- 6. Sanger Sequencing: Principle, steps, run set up demo and report analysis
- 7. Bioinformatic pipelines
- 8. Population & disease database & resources used for variant classification

Day 2 (9 am - 5:30 pm)

- 1. Sequencing Outputs, QA & QC
- 2. ACMG Classification: Applying various criteria for variant classification & reporting
- 3. Variant interpretation & Reporting with Sophia DDM (Demo including CNV analysis)
- 4. NGS Run Set up, Sequencing Methods & Illumina platforms
- 5. Interesting & informative cases: Variant Analysis, Interpretation, Clinical correlation
- 6. Report Analysis in clinics: Tips & tricks on fingertips
- 7. Feedback, Post Workshop Evaluation & Group Photo

Pre Conference Workshop 3 Functional Genomics, Tissue Culture & CRISPR (ACTREC)

Thursday & Friday: 20th & 21st February,2025 Venue: Sarin Lab & Hasan Lab, CRI, ACTREC, Kharghar, Navi Mumbai

Who should attend: Molecular Biologists, PhD students and Post Docs & Scientists Faculty: Rajiv Sarin, Syed Hasan, Ashwin Kotnis, Poonam Gera, Jasoda Chaudhari Instructors: Amartya Chakraborty, Om Yadav & Neha Gupta & Anup Jha Format: Faculty Talks, Wet Lab techniques and Demonstration,

Day 1 (9:30 am - 6:00 pm)

1. Introduction to Functional Genomics: (Talk - Dr Ashwin Kotnis)

Overview of functional genomics, its role in understanding gene function, and how cell culture models like LCLs contribute to this research.

2. Mammalian Cell Culture Techniques:

Key techniques in culturing and maintaining mammalian cells (including LCLs, PBMCs)

3. Functional Assays in Mammalian Cell Culture:

a. Cell Proliferation Assay:

Measuring cell growth and division in response to gene modifications.

b. Cell Migration Assay:

Investigating how gene modifications impact cellular motility.

4. DNA Damage Repair and Genomic Integrity:

Molecular techniques for studying DNA damage response in cell culture models.

Day 2 (9:30 am - 6:00 pm)

1. CRISPR in Gene Editing: An Introduction to Its Applications: (Talk - Dr Syed Hasan)

In-depth discussion on CRISPR/Cas9 applications in LCLs & other tissue culture systems for functional genomics. Exploring genetic modifications, gene knockouts, and gene overexpression techniques to study gene function

2. Gene Editing Technologies - CRISPR/Cas9:

- a. Designing the Guide RNA (sgRNA) b. Vector digestion and gel extraction
- c. Annealing of sgRNA Oligonucleotides d. Ligation
- e. Plasmid Isolation

f. Cell Transfection

g. Targeted DNA Cleavage

h. Validation of edits.

Measuring apoptosis in response to genetic modifications using functional assays

3. Apoptosis Assay and Cell Survival:

4. Troubleshooting and Discussion.

5. Feedback, Post Workshop Evaluation & Group Photo

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Topics to be covered in Main Conference (full programme will be shared soon)

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