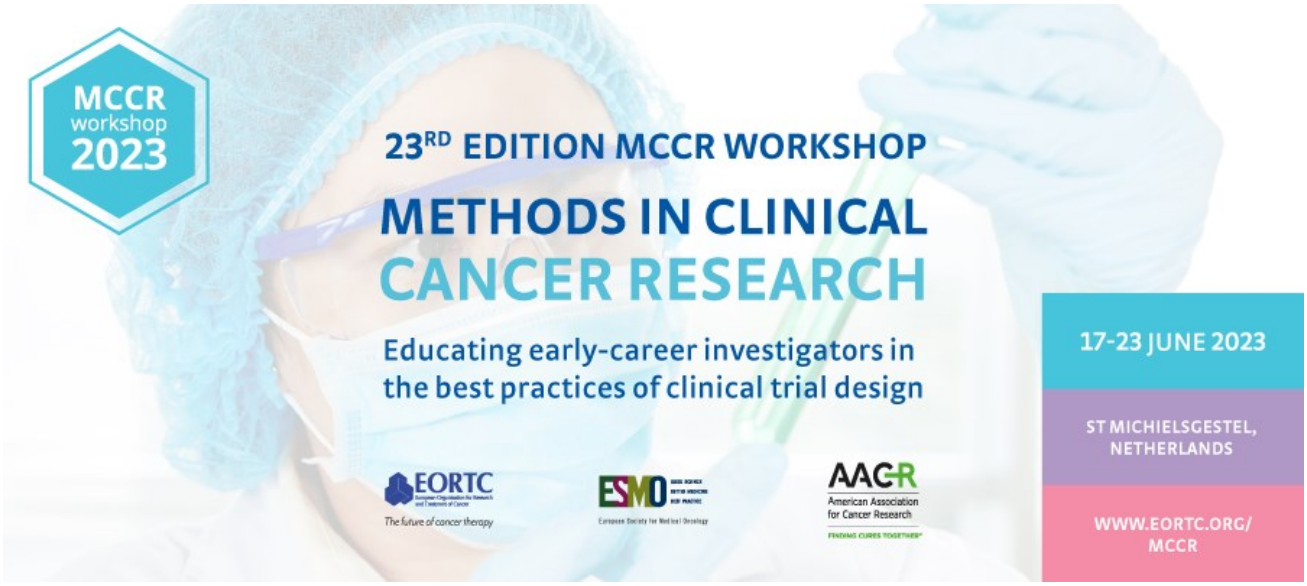
23rd MCCR WORKSHOP

METHODS IN CLINICAL CANCER RESEARCH



Corporate Support Prospectus 2023

eortc.org/MCCR

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1. INTRODUCTION

To address the need for clinical investigators in oncology, the European Organisation of Research and Treatment of Cancer (**EORTC**), European Society of Medical Oncology (**ESMO**), and American Association for Cancer Research (**AACR**) established the one-week MCCR Workshop. Since its inception in 1999 training was provided to over 1900 investigators from all over the world. This intense workshop in small groups with highly interactive discussions has had a major impact on the training of clinicians in their ability to design and implement clinical trials in cancer research.

The MCCR Workshop

The need for clinical investigators capable of designing and conducting state-of-the-art clinical trials that align innovations from basic research with early drug development is very high. As technology and its impact on clinical trial design continue to evolve at an accelerating pace, the knowledge and number of tools a clinical investigator needs to use effectively are also rapidly increasing.
The annual Methods in Clinical Cancer Research Workshop is a week-long course designed to educate and train early-career investigators in the best practices of clinical trial design and provide access to experienced clinical investigators from different institutions and countries with expertise across all areas of clinical research.

The MCCR Workshop is offered to **early-career investigators** in all disciplines working in the field of oncology who are nearing completion of their training or have recently begun their initial faculty positions and is designed to address the needs of today while maintaining flexibility to implement changes to train early-career investigators **on how to conduct effective clinical trials** in the future. In addition, it enables the development of **peer-to-peer and mentoring relationships** to enhance future career development.

Up to 80 students are selected using a competitive application process that consists of a protocol concept, personal statement, and support from an existing mentor. Those accepted to the Workshop will have access to **world-renowned faculty members** who, through the course of the Workshop, will help students **develop concept sheets into IRB-ready clinical trials**, **ready for submission at their own institutions.**

Who should attend?

The Workshop is offered to **early-career investigators in all disciplines working the field of oncology** who are nearing completion of their training or have recently begun their initial faculty positions and is designed to address the needs today, while maintaining flexibility to implement changes to train early-career investigators on how to conduct effective clinical trials in the future. In addition, it enables the development of peer-to-peer and mentoring relationships to enhance future career development.

Selection of participants

Participation to the MCCR Workshop is limited to a maximum of up to 80 participants. The Workshop Review Committee will evaluate the applications and base its decision on several factors including:

- Quality and feasibility of the proposed protocol concept and the letters of commitment submitted.
- Individual career path in medical training and competence in clinical cancer research.
- Support of relevant departments and/or institutions to help conduct the clinical trial.

The selection of applications is at the sole discretion of the Workshop Review Committee. Whilst feedback on the application process and selection is welcome, the Workshop Review Committee will not enter any discussions regarding the final decision.

Key benefits for attendees

- Exclusive access to and mentoring by up to 40 highly experienced clinical experts in the field of oncology from Europe and North America
- Exceptional opportunity to meet and network with an elite group of up to 80 junior clinical oncologists from all over the world
- Outstanding educational experience in a unique setting conducive to professional relationship building, learning and development
- Access to a variety of educational tools designed to enable participants to develop their initial protocol proposal into a complete protocol
- Establishment of a network for educational exchanges between early career cancer clinicians worldwide

Workshop Directors

Representing EORTC

Saskia Litiere

Representing ESMO

Nadia Harbeck

Representing AACR

Patricia LoRusso

2. EDUCATIONAL FORMAT & PROGRAMME

The course is comprised of a mix of academic sessions ensuring that all learning needs are catered for:

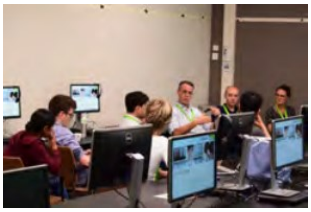
Protocol Development Group Sessions

These sessions form the core activity of this Workshop and allow students to complete the writing of their protocol by applying the knowledge acquired during the Workshop. Students will receive extensive feedback on their trial concepts from designated faculty within assigned groups comprising a maximum of ten students.

Meet the Expert Sessions

One-to-one sessions where students will have access to experts providing individual counselling on protocol related issues and advice on career development.

Small Group Discussion Sessions

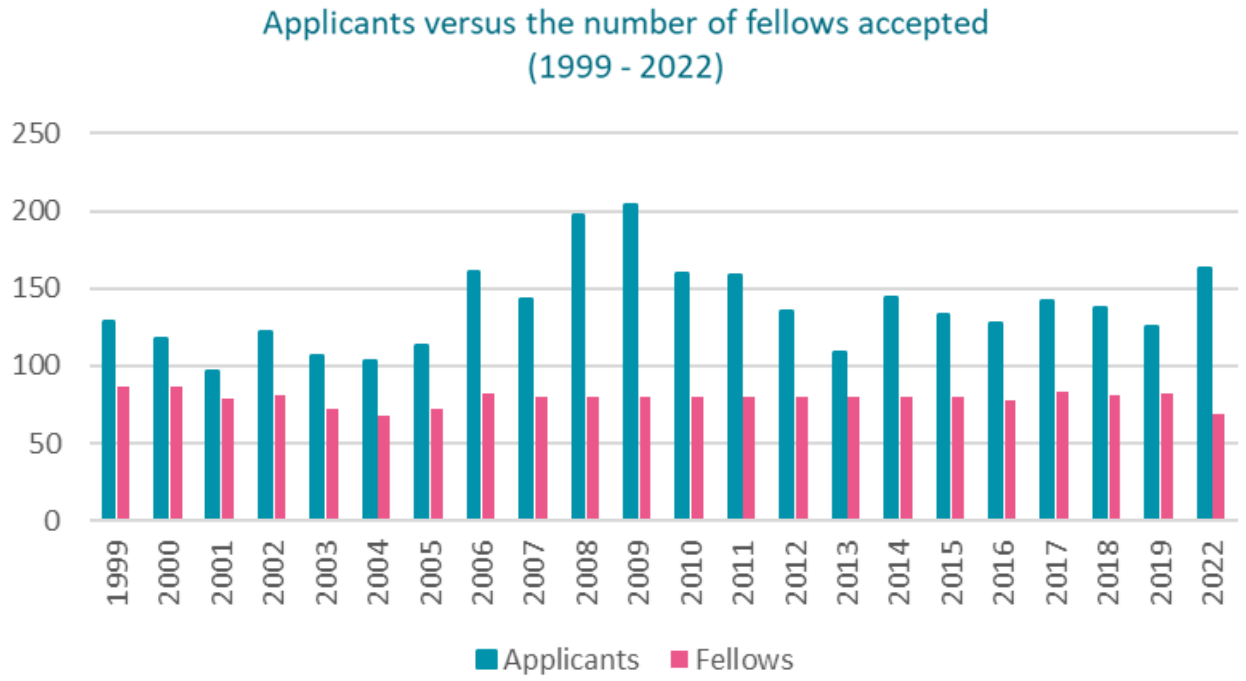


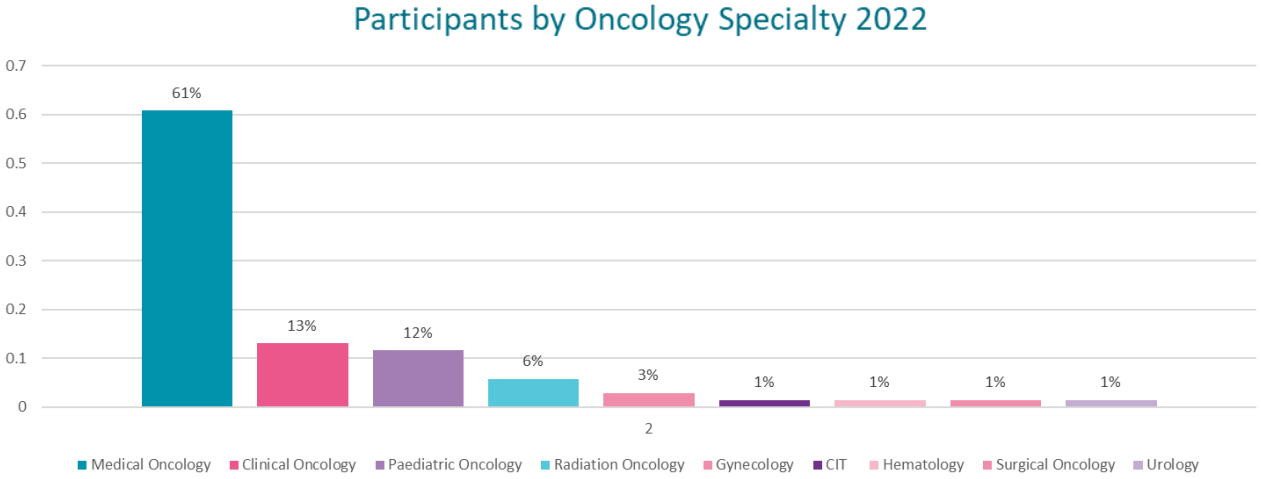
Sessions that focus on topics that are essential to the success of clinical trials and facilitating discussion on and around the difficulties and challenges of a particular type of trial. Attendance to these sessions is limited to maximise interaction and information exchanges.

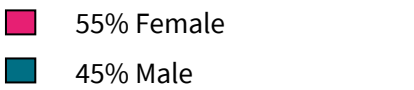
Lectures and Panel Discussions

Presentations by key experts on specific topics will provide participants with an overview of the design and implementation of high-quality clinical trials. This will be followed by a panel discussion during which Faculty and students can explore issues raised during the talks in greater depth.

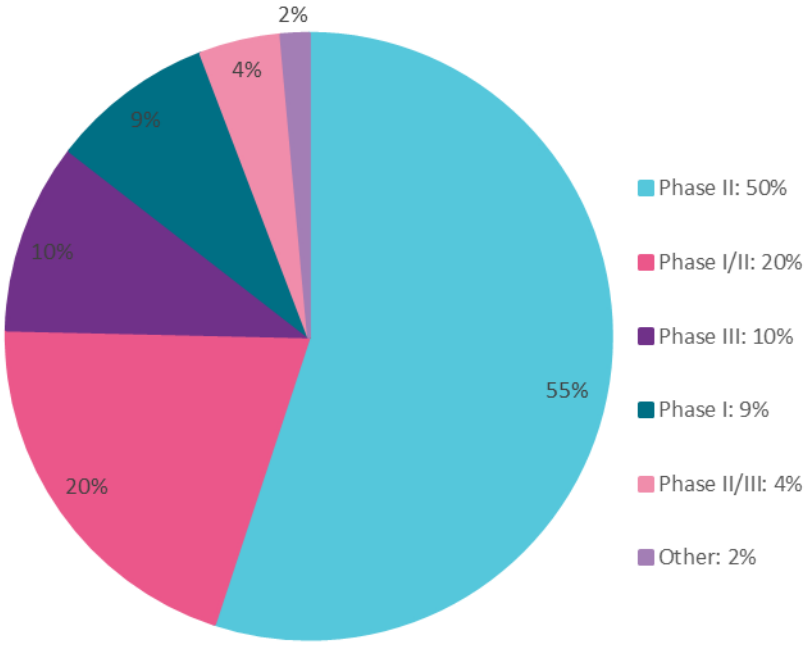
3. FACTS & FIGURES



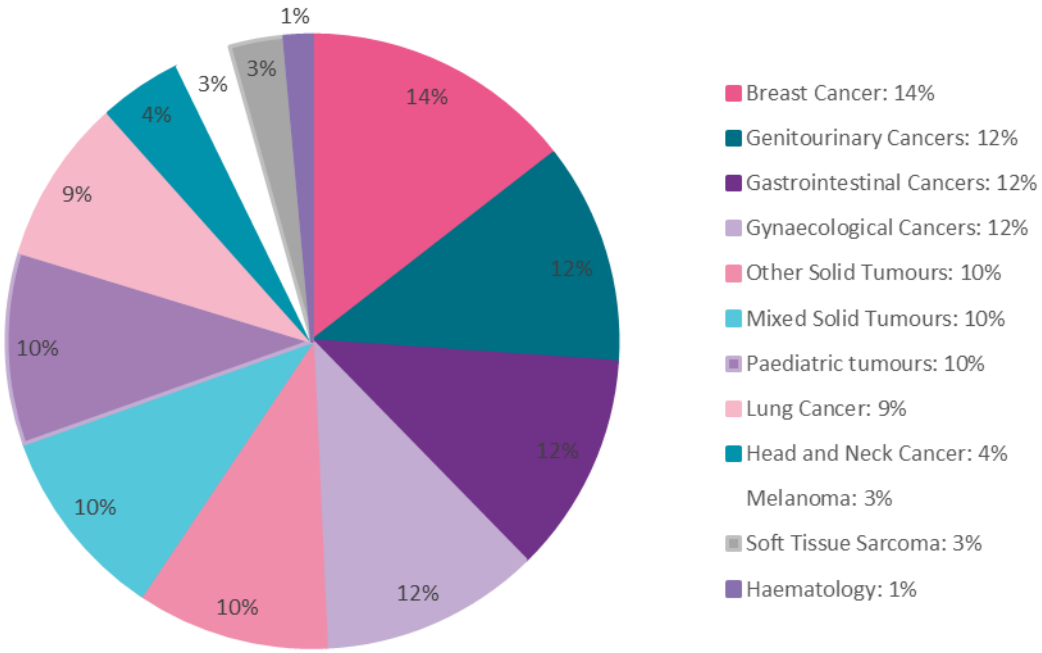




Protocols by Phase 2022 Workshop



Tumour type of protocols developed in 2022 Workshop



4. CORPORATE SUPPORT OPPORTUNITIES

As a non-profit organization we are looking to partner to support this outstanding educational event. This is a great opportunity for your company to show its involvement in an educational programme that will ultimately contribute to better research and improved public health worldwide.

	Diamond	Gold	Silver	Bronze	Contributor
	Exclusive				
	€ 95 000	€ 55 000	€ 37 000	€ 25 000	€ 10 000
Number of corporate delegates[1]	2	1	0	0	0
On-site company name recognition on message screens	✓	✓	✓	✓	✓
Acknowledgement at the Welcome Session of the Workshop	✓	✓	✓	✓	✓
Company logo on acknowledgement leaflet distributed on-site	✓	✓	✓	✓	✓
Sponsor logo on the Workshop website linked to the sponsor corporate website	✓	✓	✓	✓	
Bag insert[2]	✓	✓	✓	✓	
Observer pass for one day	✓	✓	✓		
Company name/logo included on the Workshop platform	✓	✓			
NEW Faculty member Industry (send one clinician with relevant expertise in protocol writing **or** statics from your company to join the course as faculty member) Note that CV and 400 words abstract needs to be approved by the workshop chairs.	✓				
Company name & logo printed on the delegate bag	✓				
NEW Up to 5-minute slot at the Workshop (or video message) Welcome and why this workshop is valuable	✓				

[1] To participate in the Workshop, the corporate delegate is required to fulfill the following criteria.

- The corporate delegate must be a physician specialising in oncology and submit the following items by the **28 April 2023**.
  - Personal details and contact information.
  - One-page description of the clinical therapeutic trial protocol to be written during the Workshop.
  - Statement of motivation outlining the reasons for wanting to participate in this workshop.
  - Corporate Delegates will be responsible for covering accommodation and travel expenses.

[2] A corporate flyer with the focus on research. Diamond sponsor = 2 one-page leaflets; Gold = 1 one-page leaflet

5. BOOKING FORM

Organisation:_____

Contact Person:_____

Company VAT number:_____

Purchase order number:_____

Address:_____

Zip/Postal Code:_____ City:_____

Country:_____

Telephone:_____ E-mail:_____

We would like to support the 23rd Workshop on Methods in Clinical Cancer Research as:

□	Diamond Sponsor	€ 95 000
□	Gold Sponsor	€ 55 000
□	Silver Sponsor	€ 37 000
□	Bronze Sponsor	€ 25 000
□	Contributor	€ 10 000

We agree to pay the total cost of the selected Corporate Sponsorship 30 days from the date on the invoice. We accept the sponsorship packages as described in the Corporate Support Prospectus 2023 and agree to observe and to be bound by them.

Date:_____ Signature: _____

Please complete and return to: mccr@eortc.org

Terms of Payment:

Invoices will be sent within two weeks following the confirmation. Payment is due within 30 days following the date of the invoice. Direct transfer payments should be made to the Workshop bank account: BE46 2100 5100 1036 (GEBABEBB), BNP Paribas Fortis, Rue Montagne du Parc, 3 (91616) 1000 Brussels, stating the number of the invoice. Sender's bank charges are at the expense of the sponsor. The application is legally binding on the sponsor pending its acceptance in writing by the organiser. When applicable, sponsorship packages are subject to 21% VAT

Cancellation terms:

Unexpected cancellation of the event: EORTC reserves the right to cancel MCRR without notice or compensation in the event of force majeure cases (strikes, fires, terrorist attacks, damages, pandemic, or other fatal occurrences). In such cases, EORTC is freed of all responsibility.

Contact Us

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Corporate Sponsorship Opportunities

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