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国际原子能机构
International Atomic Energy Agency
Agence internationale de l'énergie atomique
Международное агентство по атомной энергии
Organismo Internacional de Energía Atómica

Vienna International Centre, PO Box 100, 1400 Vienna, Austria
Phone: (+43 1) 2600 • Fax: (+43 1) 26007
Email: Official.Mail@iaea.org • Internet: https://www.iaea.org

In reply, please refer to: SP-TC-RER6041-2502428
Dial directly to extension: (+43 1) 2600-25981

National Liaison Officers/National Coordinators

2025-04-14

Subject: **TC Sponsored Participation on GMP for Radiopharmaceuticals from 05 to 06 June in Vienna, Austria.**

Dear National Liaison Officer / National Coordinator,

I am pleased to inform you that the International Atomic Energy Agency (IAEA) is organizing the above event under the IAEA technical cooperation project RER6041**,** "Enhancing and Harmonizing Nuclear Medicine and Diagnostic Imaging Capabilities".

The purpose of the workshop is to provide guidance through EU-cGMP regulation for manufacturing, testing, and quality assurance of radiopharmaceuticals to ensure that a manufactured product is safe for human use.

Selection of participants will be in accordance with IAEA procedures. Member States are strongly encouraged to identify women participants.

The IAEA will provide non-local participants with a round-trip air ticket based on the most direct and economical route between the airport nearest the participant's residence and Vienna or a travel allowance to purchase an air ticket. Travel details will be agreed with the participants upon receipt of their official nomination. Participants will also receive an allowance from the IAEA sufficient to cover their costs of lodging, daily subsistence and miscellaneous expenses for the duration of the event in line with IAEA rules and procedures.

We would appreciate receiving your country's nominations by 05 May 2025 through the IAEA's InTouch+ platform (https://Intouchplus.iaea.org). Should this not be possible, applicants may download the Nomination Form for the course from the IAEA's webpage. Completed forms must be endorsed by the relevant government authority and may be sent to the IAEA, preferably by email to Official Mail - IAEA Mail address Official.Mail@iaea.org, with copy to Ms Sibel Unlu s.unlu@iaea.org and Ms Angie Mieses A.Mieses-Concepcion@iaea.org. Please be advised that late nominations or replacements of participants after the closing date for nominations will not be accepted.

We look forward to receiving your early response.

Yours sincerely,

Sibel Unlu
Programme Management Officer
Division for Europe
Department of Technical Cooperation

Enclosure: Information Sheet

The 4th International Congress Imaging infections and inflammation

under the Regional TC Project RER/6/041

Event Number	SP-TC-RER6041- EVT2502428
Event Title	TC Sponsored Participation on GMP for Radiopharmaceuticals
Location	Vienna, Austria
Date	05 to 06 June 2025
Nomination Deadline	05 May 2025
Event Information	The GMP guidelines provide minimum requirements that a manufacturer of radiopharmaceuticals must meet to assure that their products are consistently high in quality, from batch to batch, for clinical use. The **Basics of cGMP** is 2 days on-site ESMIT course, providing guidance through EU-cGMP regulation for manufacturing, testing, and quality assurance of radiopharmaceuticals to ensure that a manufactured product is safe for human use. Principles regarding facilities, equipment qualification, controlled environmental conditions, processes and methods validation, standard operation procedures and user specific requirement, risk assessment, and specifications will be detailed. Also, the main components of investigational new drug and marketing authorisation dossiers will be discussed. LEARNING OBJECTIVES General knowledge regarding GMP and cGMP for Radiopharmaceuticals principles and guidelines
Organizer	European School of Multimodality Imaging & Therapy Link: GMP for Radiopharmaceuticals - The European School of Multimodality Imaging & Therapy (ESMIT)
Selection Criteria	Radiopharmacists and radiochemists in translational research, hospital radiopharmacy and radiopharmaceutical production centres