

imi PharmaTrain Career Driver

MASTERING MEDICINES DEVELOPMENT

European Curricula
for Global Expertise

My View

"I am interested to widen my knowledge and know-how in integrated medicines development. It should go beyond what I have studied and applied now during two years in a pharmaceutical company. I seek to shape my profile for the next career step."

Patricia Blank from Uni Zurich just signed up at the European Center of Pharmaceutical Medicine (ECPM).



What is PharmaTrain all about?

IMI PharmaTrain is a five year project to jointly devise a new pan-European collaborative network for comprehensive postgradual training. It is designed for all key professionals involved in the process of mastering medicines development and thereby creates the "European Curricula for Global Expertise". Involved are 23 European universities, 11 learned societies, 3 regulatory agencies and 15 member companies of the European Federation of Pharmaceutical Industries and Associations (EFPIA). IMI PharmaTrain is a unique collaboration between public universities, EC and private industry (EFPIA).

In order to develop higher quality medicines in Europe the European Commission (EC) together with the EFPIA have generated the **Innovative Medicines Initiative (IMI)** and founded it with two billion Euros. One of the four training projects, **PharmaTrain**, is devoted to pharmaceutical medicine and the integration of medicines development from molecule to market place. It concerns all key professionals working in this process and helps to develop their careers. Training providers offer full **Diploma Courses** and **Master Programmes**, which can be individualized by a selection "à la carte" from the modular **Continued Professional Development (CPD)** Platform.

Trainee Programmes ready for you First Set now

DiMD Diploma in Medicines Development: Trainees understand now the integrating concepts and best practices of the medical product development and regulatory sciences. **MMD** Master of Medicines Development: Trainees' knowledge and know-how about the integrated process from molecule to market place has reached the newly assessed European standard of Master level. **MRA** Master of Regulatory Affairs: Trainees will get a comprehensive knowledge about protocols and documents, data management of clinical trial research.

Second Set in preparation

CLIC Clinical Investigator Course, **DCTP** Diploma for Clinical Trial Professionals, **DMDR** Diploma in Medical Device Regulation

CPD/Elective Platform

Since all these training courses are built in a modular way and presented on our rich PharmaTrain CPD/Elective platform they can also be used standalone and you can craft your own training.

Universities – make your choice

Brussels (Pharmed), Basel (ECPM), Budapest (CEDDC), Barcelona, Lyon (Eudipharm), Belgrade (CEDDC), Cardiff, Duisburg-Essen, Dublin (Hibernia College), Dublin (Trinity), London (King's College), Milano, Rome, Surrey

Learn more by visiting our e-Campus

Trainers' blog

Semmelweis... University (SE) is one of participating University Centres that established the first Diploma Course in Medicines Development (DiMD) using the PharmaTrain shared standards. SE now extends the Course with another six Modules, incl. two Electives, to a full Master of Medicines Development (MMD) in cooperation with other universities, the Cooperative European Drug Development Course (CEDDC).



"We are ready to provide a platform to strengthen your competence in bringing a medicine from molecule to market, what we call integrated medicines development."



Sándor Kerpel-Fronius, SE, Budapest

Industry's Voice

"The Pharmaceutical Industry recognized the need for greater competence in integrated medicines development through high quality and standardised training throughout Europe – and thereby mastering the process."

Mike Hardman, AstraZeneca

For more information visit:
www.pharmatrain.eu

