The Summer School on Pharmaceutical Analysis (SSPA) is planned under the auspices of the Division of Medicinal Chemistry of the Italian Chemical Society (SCI) and the European Federation for Medicinal Chemistry (EFMC).

The school is open to a limited number of students and the official language is English.

International scientists coming to Rimini for the SSPA 2010 may take the opportunity to enlarge their knowledge into the field of pharmaceutical analysis in the frame of the following aspects: physical properties characterization during pharmaceutical product development, preformulation and formulation, analytical quality by design and method validation.

Tutorials on specific subjects will be organized in the afternoons.

All inclusive registration is 400 euro and includes access to all scientific sessions, all meals, coffee, the final program, the abstract book, and 3-night accommodation in a double/twin room at a three-star hotel located at a walking distance from the SSPA 2010 venue.

Deadline for on line registration is 30 May 2010.

Visit www.scpaweb.org or contact SSPA 2010 secretariat at sspa2010@scpaweb.org for further information and registration.

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13-16 June 2010

Summer School on Pharmaceutical Analysis

SS PA

ADVANCED
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IN DRUG
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PROGRAM PROGRAM PROGRAM PROGRAM PROGRAM PROGRAM PROGRAM PROGRAM PROGRAM

| Sunday, June 13, 2010 | | Tuesday lung 15, 2010 | | 16:00 -16:15 | Coffee break | |
|---|--|---|--|--------------------------|---|--|
| 16:00 - 18:00 18:00 - 18:30 | Registration Welcome and opening ceremony | Tuesday, June 15, 2010 II SESSION – PREFORMULATION AND FORMULATION | | 16:15 – 18:15 | Tutorial on multivariate analysis. M. Machin – Novartis Pharma AG (Switzerland) | |
| 18:30 - 19:30 | Opening lecture. G. Gaviraghi , Siena Biotech (Italy) | 9:00 - 10:00 | Drug delivery: an overview. S. Nicoli - | Wednesday, June 16, 2010 | | |
| 19:30 - 20:30 | Welcome buffet | 10:00 - 11:00 | University of Parma (Italy) 10:00 - 11:00 Formulation in preclinical studies. G. | | IV SESSION – METHOD VALIDATION | |
| Monday, June 14, 2010 | | | Verreck - Johnson & Johnson (Belgium) | 9:00 - 10:30 | Objective of validation, | |
| 9:00 - 9:15 | Opening of SSPA 2010 | 11:00 - 11:15 | Coffee break | | documentation and regulatory bodies, validation and quality | |
| I SESSION – PHYSICAL PROPERTIES CHARACTERIZATION DURING | | 11:15 - 12:15 | Active pharmaceutical ingredients and choice of formulation principle: biopharmaceutical considerations. W. Weitschies - University of | | assurance , traceability. S. Rudaz - University of Geneva (Switzerland) and E. Rozet - University of Liège (Belgium) | |
| PHARMACEUTICAL PRODUCT DEVELOPMENT | | | Greifswald (Germany) | 10:30 - 10:45 | Coffee break | |
| 9.30 - 10.30 | DS and DP characterization across product development. B. Allieri - GSK (Italy) Polymorphism assessment. D. Braga | 12:15 - 13:15 13:15 - 14.00 | Formulation development: drug/excipient interaction and formulation design. M. Bresciani - GSK (Italy) | 10:45 - 12:30 | Terminology and definition. Practical guidelines for method validation. S. Rudaz - University of Geneva (Switzerland) and E. Rozet - University of Liège (Belgium) | |
| 10.00 11.00 | - University of Bologna (Italy) | | Lunch | | | |
| 11:30 -11:45 | Coffee break | | | 12:30 - 13:30 | Lunch | |
| 11:45 - 12:45 | Spectroscopical methods toward physical properties | (QbD) | ANALYTICAL QUALITY BY DESIGN | 13:30 - 15:00 | Tutorial on validation and data treatment, examples, software demonstration. S. Rudaz - University of Geneva (Switzerland) and E. Rozet - University of Liège (Belgium) | |
| | characterization. H. Lucas - Kaiser Optical System (France) | 14:00 - 15:00 | Combining QbD and Process Analytical Technology: understand and improve pharmaceutical processes. D. Lochmann - Merck- Serono (Germany) | | | |
| 12:45 - 14:00 | Lunch | | | 15:00 - 16:30 | Tutorial on experimental design, examples, software demonstration. S. Furlanetto - University of Florence (Italy) | |
| 14:00 - 15.45 | Ph.D. thesis dissertations | | | | | |
| 15:45 - 16:00 | Coffee break | 15:00 -16:00 | QbD: Regulatory requirements and expectations. F. Bird – GSK Ware (UK) | | | |
| 16:00 - 18.00 | Tutorial on forced degradation and stability studies. M. Galvani - GSK (Italy) | | | 16:30 -16:45 | Closing remarks | |

SSPA 2010 Secretariat

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