JPAG
JOINT PHARMACEUTICAL ANALYSIS GROUP

ensuring knowledge, enhancing quality

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*run **for** pharmaceutical analysts **by** pharmaceutical analysts*

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JPAG future events

Thursday 3rd December 2026
Regulatory Hot Topics XI
Royal Society of Chemistry, London

APS GB join JPAG at the 11th Regulatory Hot Topics symposium which will include an update on BP activities and specifically look at the BPs new radiopharmaceuticals guidance. The programme will cover a discussion of international reliance procedures by MHRA, various ICH updates from ongoing guideline revisions. We will catch up with CERSIs and AI, and the changing landscape for combination products due to changes in the medical device regulations.

As usual, the meeting will provide an excellent opportunity for dialogue, discussion and debate in an open forum and allows learning from expert speakers involved in the development of quality regulatory guidance and best practice.

The meeting is open to the submission of abstracts for posters in line with the above proposed key themes.

Speakers:
Dr Paul Dickinson - Founder and Chief Scientist, Seda Pharmaceutical Development Services
Dr Matt Heath - Innovation Specialist, MHRA
Steve Hoare - Standards & Regulatory Governance, MHRA
Marine Joly Battaglini - PharmDev Consulting
Tasneem Keshavji - Head of International Recognition, MHRA
Dr Abha Raveau-violette - Combination Products and Medical Devices Ltd
James Rickard - Chief Scientific Officer, Richmond Research Institute
Prof Walkiria Schlindwein - De Montfort University, Leicester
Dr Stuart Williams - Dove Analytics
Dr Christian Zeine - Scientific Affairs EMEA, USP

Register now at www.jpag.org/cp196

Thursday 21st January 2027
Advanced Dissolution
Royal Society of Chemistry, London

This event brings together scientists, analysts, and industry colleagues for a focused, one day exploration of advanced dissolution and its growing role in modern pharmaceutical development. It is a follow on from the Fundamentals event held in October 2025. As formulations become more complex and regulatory expectations evolve, dissolution has shifted from a simple quality test to a powerful, mechanistic tool for understanding drug performance.

This seminar creates space for sharing experiences, practical insight alongside open discussions. During the day, we aim to highlight new approaches, identify emerging challenges, and discuss the analytical strategies that will shape the next generation of oral medicines. The challenge for us all is: can a single test be sensitive enough to predict changes that may be of clinical relevance, whilst being robust enough to satisfy the needs of a quality control environment?

We're open for abstracts for both oral presentations and posters - deadline: 31-Oct-26 (oral) and 1-Jan-27 (posters).

Register now at www.jpag.org/cp199

Interested in any of these meetings?
Please visit the JPAG website for full details at www.jpag.org

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