Receive case studies from Merck Serono, Teva, Cinfa Biotech GmbH, MHRA and more at Biosimilars and Biobetters

This year's conference will address regulatory updates, pharmacovigilance of biosimilars, patent litigation, market access and global market developments.

LONDON, ENGLAND, UNITED KINGDOM, August 31, 2015 /EINPresswire.com/ -- SMi's 6th annual Biosimilars and Biobetters conference, taking place on 30th September - 1st October 2015 in London, will provide the ideal platform to engage in scientific discussions and debate the best practices and solutions to improve industry performance through technical case studies and practical advice from leading biosimilar and regulatory players from organisations including Harvest Moon Pharmaceuticals USA Inc, GfK, MHRA, Selecta Biosciences, Cinfa Biotech GmbH, Norwegian Medicines Agency and more. Unique perspectives will be shared on:

• Evolving biosimilar regulatory landscape
• Market access and product commercialisation barriers
• Global marketing developments, including a focus on emerging market trends
• In-depth protein characterisation and analytical comparability to efficiently and effectively collect data

Speaker Panel includes:

• Richard DiCicco, Chairman, Harvest Moon Pharmaceuticals USA, Inc.
• Shahin Kauser, Senior Scientific Assessor, MHRA
• Chris Teale, Vice President Europe, GfK NOP Ltd
• Bracha Timan, Director, Israel Site Head, Global Bioassays & Technology, Global R&D, Teva
• Takashi Kei Kishimoto, Chief Scientific Officer, Selecta Biosciences
• Karsten Roth, Director Clinical Operations, Cinfa Biotech GmbH
• Alan Sheppard, Principal, Global Generics and Biosimilars, IMS Health
• Steinar Madsen, Medical Director, Norwegian Medicines Agency

To view the full speaker line-up and conference programme, visit http://www.biosimilars-biobetters.co.uk/einpresswire

Plus, don't miss two interactive conference workshops on:

A: Assessing the Regulatory Framework for Europe and the US – Developing Future Biologics
Led by: Lincoln Tsung, Daniel Kracov, Jennifer Sklenar, Partners, Arnold & Porter (UK) LLP
29th September 2015, 13.30 - 17.30

B: Biosimilars – Understanding the Regulatory Processes and the Commercial Realities
Led by: Peter Wittner, Senior Consultant, Interpharm Consultancy

2nd October 2015, 08.30 - 12.30

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