2016 China/EU Pharmaceutical Industry Forum

AGENDA

5 April 2016, Hamburg, Germany

Venue: InterCityHotel Hamburg Dammtor Messe, St Petersburger Strasse 1, Hamburg
Organisers: CPIA and EFPIA
Presenters: CPIA, EFPIA, European Commission, European Medicines Agency and IMS.

Focus areas: Overview of pharmaceutical market in China/EU, Pharmaceuticals in the Environment, the Drug Administration Law (DAL) revision and implementation of new EU legislation

Moderators: Ms Zhou Yan, Vice Chairman and Secretary General, CPIA (morning session)
Mr Par Tellner, Director, Team Leader, Regulatory Affairs, EFPIA (afternoon session)

	Topic	Title of presentation	Whom
09:00-09:10		Welcome words	CPIA Mr Richard Bergström, Director General, EFPIA and Mr Zhang Ming Yu, Senior Vice Chairman, CPIA
09:10-10:10	**Overview of pharmaceutical market in China/EU**	Overview of pharmaceutical market in China	Mr Zhang Zhen En, CPIA Vice Chairman and Chairman North China Pharmaceutical Group
		Overview of pharmaceutical market in EU	Mr Per Troein, Vice President, Strategic Partners, IMS
10:10-11:10	**Pharmaceuticals in the Environment (PIE)**	European Commission strategy for Pharmaceuticals in the Environment (PIE)	Ms Patrizia Tosetti, Medical products: quality, safety, innovation, European Commission
		Proposals from European pharm industry for PIE	Mr Bengt Mattson, PIE Task Force,EFPIA
11:10-11:30	**Coffee/tea break**		
11:30–13:00	**Drug Administration Law (DAL) revision**	EFPIA's view on regulatory reform in China	Mr Jin Shun, Vice chair, EFPIA China regulatory network
		Progress of the revision of drug registration and approval in China	Mr Zhang Ping, Chief Engineer, Tianjin Pharmaceutical Group
13:00-14:00	**LUNCH**		
14:00–15:30	**Implementation of new EU legislation**	Identification and authentication of medicines in Europe: What will happen in 2019	Ms Patrizia Tosetti, Medical products: quality, safety, innovation, European Commission
		Implementation of Clinical trial regulation	Mr Fergus Sweeney, Head of Inspections and Human Medicines Pharmacovigilance, European Medicines Agency (EMA)

15:30-15:50	**Coffee/tea break**		
15:50-16:20	**EU System for marketing authorisation**	EU system for marketing authorisations	Ms Bente Jessen, chair EFPIA China regulatory network
16:20–16:50	**Panel discussion re DAL revision and implementation of EU legislation**		Jin Shun, Zhang Ping, Patrizia Tosetti, Fergus Sweeney and Bente Jessen
16:50-17:00	**Conclusions**	Conclusions	Mr Zhang Ming Yu, Senior Vice Chairman CPIA
17:00		End of the meeting	