

2018 China/EU Pharmaceutical Industry Forum

AGENDA

16 April 2018, Basel, Switzerland

Venue: Auditorium WSJ 510, Novartis Campus, Fabrikstrasse 2, CH-4056 Basel, Switzerland

Organisers: CPIA and EFPIA

Presenters: CPIA, EFPIA, EU Commission, EMVO, scienceindustries

Focus areas: Pharmaceuticals in the Environment, Coding & Serialisation, Compliance and CFDA in the ICH

Moderators: Ms Lei Ying, Vice President & Secretary General, CPIA (morning), Mr Faraz Kermani, Medical writer (afternoon)

	Topic	Title of presentation	Whom
09:00-09:10		Welcome words	Ms Couzette Kleynhans, Head International Public Affairs, Novartis Mr Pan Guangcheng, Executive President, CPIA Ms Kang Wei, Managing Director, RDPAC
09:10-10:10	CFDA in the ICH	1. The pharmaceutical market in China	Mr Tu Yongrui, Vice President, CPIA, Chairman & General Manager, Chang Zhou Si Yao Pharmaceuticals Co., Ltd
		2. Overview of ICH and industry support to implementation of ICH guidelines by CFDA	Mr Pär Tellner, Director, Team Leader, EFPIA
		3. New pharmaceutical policies of China in the ICH	Mr Liu Xiaohan, Special Guest Vice President, CPIA, Director-General, Hebei Pharmaceutical Association, Secretary of the Party Committee, Hebei Pharmaceutical Profession Association
10:10-11:20	Pharmaceuticals in the Environment (PIE)	4. European pharm industry view PIE consultation	Mr Bengt Mattson, Chair, PIE Task Force, EFPIA
		5. Green pharmaceuticals in China	Mr Hong Hao, Special Guest Vice President, CPIA, Chairman & CEO, Asymchem Laboratories (Tianjin) Co., Ltd.
		6. Comprehensive environmental protection in the pharmaceutical industry in China	Mr Liu Wenfu, Special Guest Vice President, CPIA, General Manager, North China Pharmaceutical Group Corporation
11:20-12:20	Compliance (Marketing ethics)	7. Insights in the Swiss Pharma Self-regulation and its benefits for the local industry	Mr Jürg Granwehr, Attorney at Law, Head Pharma, scienceindustries
		8. Innovation of the pharmaceutical industry in China	Mr Zhang Yaohua, Vice President, Shanghai Pharmaceuticals Holding Co., Ltd.
		9. Chinese Consensus Framework for the Ethical Collaboration in the Pharmaceutical and Medical Device Sectors	Mr Pan Guangcheng, Executive President, CPIA
12:20-13:15	LUNCH		
13:15-14:00	CFDA in the ICH cont.	10. CFDA in the ICH	Mr Qin Xiaolin, CFDA
14:00-15:20	Coding & Serialisation	11. Identification and authentication of medicines in Europe: Less than a year to the deadline of implementation 9 February 2019	Ms Patrizia Tosetti, Policy officer, Medical products: quality, safety, innovation, European Commission,
		12. Identification and authentication of medicines in Europe: Opportunities and challenges to comply with EU legislation	Mr Andreas Walter, General Manager, European Medicines Verification organization (EMVO)
15:20-15:40	Coffee/tea break		
15:40-16:30	Panel discussion re PIE, Compliance, Coding and serialization and CFDA in the ICH		Mr Faraz Kermani (moderator), Mr T Yongrui, CPIA, Mr Pär Tellner, EFPIA, Mr Liu Xiaohan, CPIA, Mr Bengt Mattson, EFPIA, Mr Hong Hao, CPIA, Mr Liu Wenfu, Mr Jürg Granwehr, Mr Zhang Yaohua, CPIA, Mr Pan Guangcheng, Ms Lei Ying, CPIA, Ms Patrizia Tosetti, European Commission and Mr Andreas Walter, EMVO
16:30-16:40	Conclusions	Conclusions	Mr Pär Tellner, Director, Team Leader, EFPIA

