DIR/003 (E)

CODE: SEM3-02-2017-SMTA

TITLE: ISO 13485 medical devices quality management system & international standards for pharmaceutical supply chain

OBJECTIVE: This seminar will give you an overview about international standards such as ISO 13485, GMP, GDP and how they can support your pharmaceutical quality control.

CONTENT:

- Introduction to ISO 13485:2016 Medical Devices Quality Management System
- Good Manufacturing Practices (GMP) for Pharmaceutical industry (Western & Chinese Pharmaceutical Products)
- Good Manufacturing Practices (GMP) for Secondary Packaging Manufacture in Pharmaceutical industry
- Good Distribution Practices (GDP) for Pharmaceutical industry
- Risk Management for Medical Devices and Pharmaceutical industry

DESIGNED FOR:

- Manufacturers or Traders for Medical Devices
- Manufacturers or Traders for Pharmaceuticals
- Logistics companies for Medical Devices and/or Pharmaceuticals
- Govetnment departments
- Hospitals, public health centers, medicine research teams in universities.

SPEAKER(S): Mr. Kelvin Sze

ORGANIZED BY: Macau Productivity and Technology Transfer Center (CPTTM)

NO. OF SEATS: 50 LANGUAGE: Chinese

DATE & TIME: February 15, 2017 (Wednesday), 15:00-17:30

FEE: Free of charge REGISTRATION DEADLINE: 14/02/2012 (Tuesday)

VENUE: CPTTM Head Office – Auditorium
Rua de Xangai 175, Ed. ACM., 7 andar, Macau

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	House of Apparel Technology	Rua dos Pescadores, Ed. Ind. Ocean, Fase II, 10 andar Tel: (853) 8898 0701 Fax: (853) 2831 2079
	Cyber-Lab	Rua Comandante Mata Oliveira, Ed. Associacao Ind. 3 andar Tel: (853) 8898 0601 Fax: (853) 2837 3085

~ The organizer reserves the right to cancel the Workshop or modify its topic, content and/or speaker without prior notice. ~