CRMO SOP Training Workshop + Post CFDA Inspection Sharing

- Training Workshop on the following 5 updated CRMO SOPs (which will be effective on 1 June 2016)
- Sharing after China-FDA inspection conducted in early March

Date	Session 1: 17 May 2016 (Tuesday), 1 to 2pm Session 2: 25 May 2016 (Wednesday), 1 to 2pm *(Light lunch will be provided)*
Venue	Seminar Room, 2/F, Lui Che Woo Clinical Sciences Building, PWH
5 updated CRMO SOPs	
1. CRMO-SOP-003	-SOP for application for clinical trial certificate from Department of Health
2. CRMO-SOP-006	-SOP for Study Document Filing, Retention and Storage
3. CRMO-SOP-007	-SOP for Investigational Medicinal Product (IMP) management
4. CRMO-SOP-019	-SOP for Inspection of Research Unit
5. CRMO-SOP-020	-SOP for biological sample management
Remarks	*The training record of these 5 SOPs for all attendees will be registered on CRMO website.* *Online training on these SOPs will also be launched in late May*

Registration
(by 10 May 2016):

For Enquiry: 2632 4276/ crmo@cuhk.edu.hk

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