

報名須知

一、報名費用

每人 3,600 元。

二、報名資格

歡迎所有對臨床試驗工作領域有興趣之醫藥相關人士報名。

三、報名方式

採網路線上報名：請至 <http://www.mpat.org.tw> 報名

四、付款方式

受理匯款、劃撥、ATM 轉帳、信用卡方式付款，詳情請參考報名網站說明。

五、注意事項

1. 即日起受理報名。因場地座位有限，敬請儘早完成報名。
2. 全程參與研討會者將核發完成訓練證書(6 小時)
3. 本次研討會將申請醫事人員繼續教育學分(醫師、藥師、護理人員類)及公務人員終身學習時數。
4. 聯絡方式：
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5. 凡因故取消報名或替換與會者(替換與會者，限更改一次)，須於活動前五天完成相關手續，取消報名將扣除手續費 600 元/每場；逾期(含活動當天)取消恕不退費。詳細退費說明參考報名網站。
6. 備註：
 - (1). 本次研討會部分講題(含外籍講師)將以英文進行演講授課；
 - (2). 本次研討會不提供同步口譯服務；
 - (3). 本會保留報名名額資格審核權，以及修改議程、額滿截止報名及未達最低人數時取消研討會辦理之權利。

7. 詳細議程內容如下

**Conference on Recent Advances in Clinical Trials
Agenda (June 12, 2018)**

| Time | Topic | Speaker |
|--------------------|---|---|
| 09:00-09:20 | Opening Remarks | FMPAT, NHRI |
| Session I | Chair: Chin-Fu Hsiao Investigator/Deputy Director, Institute of Population Health Sciences, National Health Research Institutes | |
| 09:20-09:50 | CDE's experiences to review MRCT results and expectations for E17 guideline | I-Chun Lai (賴怡君) Team Leader, Center for Drug Evaluation |
| 09:50-10:20 | Important literature in MRCT | Chieh Chiang (姜杰) Post-doctor, Institute of Population Health Sciences, National Health Research Institutes |
| 10:20-10:50 | Statistical issues in design and analysis of clinical trials including real-world data | Toshimitsu Hamasaki (濱崎俊光) Director, Department of Data Science, National Cerebral and Cardiovascular Center |
| 10:50-11:10 | Break | |
| 11:10-11:40 | Clinical innovation network | Haruko Yamamoto (山本晴子) Director, Center for Advancing Clinical and Translational Sciences, National Cerebral and Cardiovascular Center, Japan |
| 11:40-12:10 | Introduction to pragmatic trials | I-Shou Chang (張憶壽) Investigator, National Institute of Cancer Research, National Health Research Institutes |
| 12:10-12:20 | Floor Discussion | |
| 12:20-13:30 | Lunch Break | |
| Session II | Chair: Hsiao-Hui Tsou Associate Investigator, Institute of Population Health Sciences, National Health Research Institutes | |
| 13:30-14:00 | Adaptive dose-finding methods in early-phase clinical trials | Yuh-Ing Chen (陳玉英) Distinguished Professor, Institute of Statistics, National Central University |
| 14:00-14:30 | Early phase designs | Chin-Fu Hsiao (蕭金福) Investigator/Deputy Director, Institute of Population Health Sciences, National Health Research Institutes |
| 14:30-15:00 | Break | |
| 15:00-15:30 | How the ICH E9 addendum around estimands may impact our clinical trials (I) | Frank Bretz Novartis |
| 15:30-16:00 | How the ICH E9 addendum around estimands may impact our clinical trials (II) | Mey Wang (王玫) Senior Reviewer, Center for Drug Evaluation |
| 16:00-16:30 | TBD | TBD |
| 16:30-17:00 | Discussions | |
| 17:00-17:10 | Closing Remark | Li-Tzong Chen (陳立宗) Director / Distinguished Investigator / Attending Physician, National Institute of Cancer Research, National Health Research Institutes |