Date: August 19, 2017

Time: 16:15-18:45

Venue: Jiangmen Hall, 3rd Floor, Block 2

Session: Clinical Investigation I (Cancer,

Liver Disease and Inflammation)

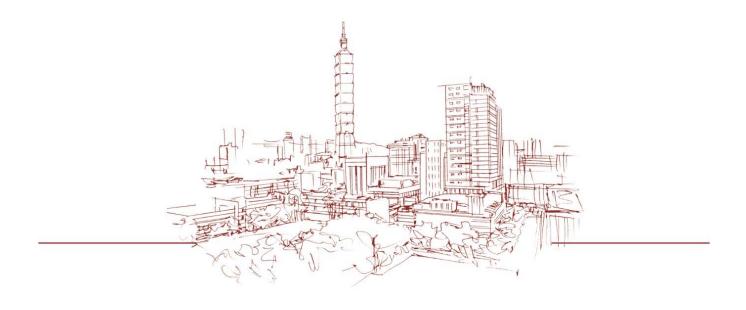
Chairman: Jeffrey D. White (NCI)

Co-chairman: Yun Yen (TMU)



Clinical Investigation I (Cancer, Liver Disease and Inflammation)

- ➤ Chairman: Jeffrey D. White (NCI)
- **≻**Co-chairman:
 - **Yun Yen**, Taipei Medical University
 - ➤ **Kequn Chai**, Tongde Hospital of Zhejiang Province
- **≻**Panelists:
 - **➢ Jian-Ping Chen**, The University of Hong Kong
 - ➤ Shwu-Huey Liu, President of PhytoCeutica Inc



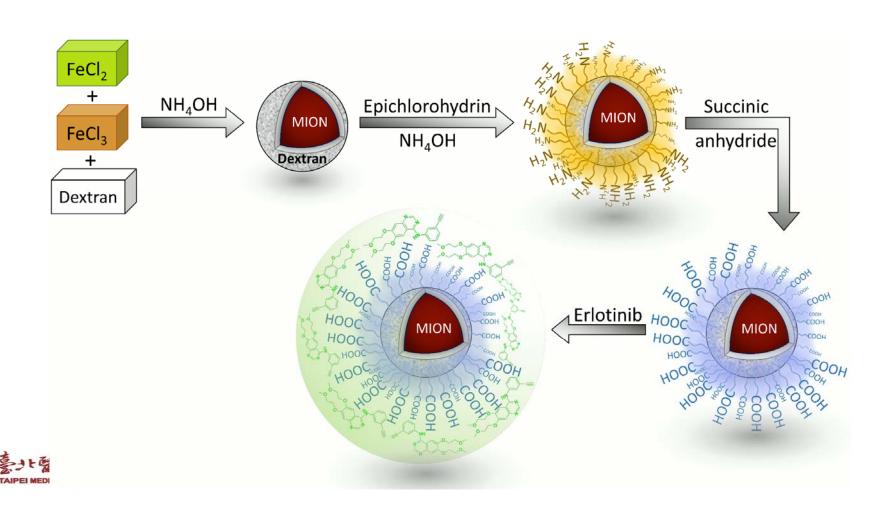
Novel theranostic biomarker probes: Can we predict and monitor the treatment of cancer?

Prof. Dr. Hsu-Shan Huang



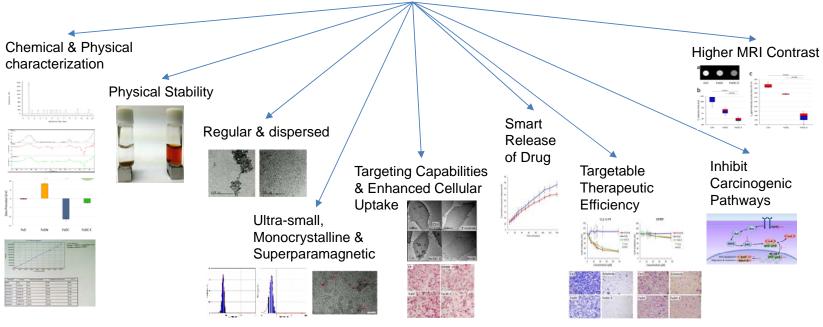


Design and synthesis of the erlotinib-loaded NPs



Conclusions







Identification of predictive metabolites for the time to tumor progression of phase II trail of Irinotecan plus PHY906 (YIV906) in patient with metastatic colorectal cancer

Jiehuan Sun, Wing Lam, Shwu-Huey Liu, Guoxiang Xie, Wei Jia, Howard S Hochster, Edward Chu, Hongyu Zhao, YungChi Cheng



Summary for phase IIB clinical of PHY906+/- CPT-11 for patient with metastasis colorectal cancer

- Patients of CPT11/PHY906 group appears to have longer time to progression than CPT11 alone group
- When metabolites data and cytokine/chemokines from patient plasma was analyzed by using L1 penalized Cox regression model, the dichotomized score derived from these metabolites and cytokine can separate the patients well into non-responders and responders to CPT11, regardless of PHY906.
- PHY906 appears to improve quality of patient receiving CPT11.



Thank you for your attention!

See you in Malaysia