1960 TAIPEI MEDICAL UNIVERSITY

July 2<sup>nd</sup> – 3<sup>rd</sup>, 2024 | Taipei, Taiwan

Speaker information

## **General Information**

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Affiliation	School of Pharmacy, College of Pharmacy, Taipei Medical University	
Education Background	Ph.D., Graduate Institute of Pharmaceutical Sciences, National Taiwan University	
Professional Appointment	Distinguished Professor	
Research Interest	Structure optimization through rational drug design and intelligent modification.	
Website	https://hub.tmu.edu.tw/en/persons/jing-ping-liou	
<ol> <li>2024 Apr 1;586:216666. (I</li> <li>Sharma S, Wang SA, Yang <i>Chem.</i> 2024 Feb 22;67(4):2</li> <li>Narwanti I, Yu ZY, Sethy B (IF = 6.700; CHEMISTRY, M</li> <li>Tseng HJ, Banerjee S, Qia <i>Chem.</i> 2023 Aug 5;256:111</li> <li>Wu TY, Chen M, Chen IC, 2023 Apr;46:159-171. (IF =</li> <li>Nepali K, Wu AC, Lo WL, C = 6.700; CHEMISTRY, MED</li> <li>Mehndiratta S, Qian B, C CHEMISTRY, MEDICINAL;</li> <li>Thakur A, Faujdar C, Shar 8685. (IF = 7.300; CHEMIS</li> </ol>	n B, Lai MJ, Wu TY, Hsu TI, Lin TE, Hsu KC, Chuang KH, 5459. (IF = 6.700; CHEMISTRY, MEDICINAL; 7/60 11.6% Chen YJ, Chen CY, Wang CH, Cheng JJ, Nepali K*, Chu = 10.7; MULTIDISCIPLINARY; 10/73 13.6%) Chopra B, Lai MJ, Chuang JY*, <b>Liou JP</b> *. <i>Eur J Med Chem.</i> DICINAL; 7/60 11.6%) huang JY, <b>Liou JP</b> *, Shih JC*. <i>J Med Chem.</i> <b>2022</b> Feb 1	K*, Hsu TI*, <b>Liou JP</b> *. <i>J Med</i> %) hem. <b>2023</b> Jun 2;258:115505. <b>Liou JP</b> *, Shih JC*. <i>Eur J Med</i> ) ang KH*, <b>Liou JP</b> * <i>J Adv Res.</i> <b>2023</b> Feb 15;248:115054. (IF 0;65(3):2208-2224. (IF = 7.3; nem. <b>2022</b> Jul 14;65(13):8596-

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## Speech Topic and Abstract

Title:

New Immunosensitizer for Treatment of Colorectal cancer

## Abstract:

Utilizing rational drug design approach to synthesize a series of small molecule compounds. Lead compound exhibited not only direct cytotoxicity to cancer cells but also downregulated immune checkpoints (PD-L1 and IDO) expression in tumors via the inhibition of STAT1 pathway and degradation of oncogene proteins (Src, AKT, Rb, and FAK), leading to in vivo tumor growth inhibition. These multiple effects enabled the effector T cells to largely infiltrate into the tumor region and release granzyme B to kill cancer cells. In addition, Lead also decreased TGF- $\beta$  secretion from normal cells, resulting in the systemic reduction of immunosuppressive regulatory T cells. Delightfully, a cocktail treatment of lead compound and anti-PD-1 antibodies demonstrated synergistic efficacy to eliminate solid tumors with 83.9% of tumor growth inhibition.