

Yi Li

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Education

China Pharmaceutical University Nanjing, P. R. China
B.S. in Pharmaceutical Science (Top-notch Innovation Program) 09/2016-07/2020
GPA: 3.6/4.0

Honors:

- Outstanding Student Scholarship, Top-notch Innovation Program (2018 and 2019)
- Overseas Outstanding Student Scholarship (2018)
- Third Prize Scholarship (top 10%) (2017)

University of Strathclyde Glasgow, United Kingdom
Summer Exchange Program 07/2018-08/2018

Research

Key Laboratory for Druggability of Biopharmaceuticals, China Pharmaceutical University, Nanjing, P. R. China
Advisor: Dr. Lifang Yin and Dr. Wei He

- **Project Leader:** Study on a Smart Paclitaxel-Dichloroacetic Acid Nanocrystals for Efficient MDR Reversal and Enhanced Apoptosis. 01/2019-present
 - 1) Constructed a multifunctional drug-delivery-drug self-assembled nanoplateform to reverse MDR effects and enhance apoptosis via a non-lysosomal pathway.
 - 2) Explored methods of preparing a two-drug platform and optimized the prescription via simulation.
 - 3) Characterized particles *in vitro* and improved experimental conditions based on their effects on the nanocrystals' behavior and properties.
 - 4) Studied cytotoxicity and apoptosis in normal and drug-resistant A549 strains.
 - 5) Studied intracellular distribution and non-lysosomal inoculation pathways.
- **Research Associate:** Using Marimastat-loaded Thermosensitive Liposomes (MATT-LTSLs) and Paclitaxel Nanocrystals (PTX-Ns) as Dual-nanomedicines to Treat Metastatic Cancer. 11/2018-03/2019
 - 1) Designed MATT-LTSLs to lock cancer cells into their microenvironment via hyperthermia as well as PTX-Ns to enhance apoptosis via local treatment.
 - 2) Analyzed the thermosensitivity of LTSLs by measuring fluorescence anisotropy in relation to membrane fluidity.
 - 3) Studied cellular uptake of PTX using flow cytometry; examined the antimetastatic ability of MATT-LTSLs through transwell assays; and assessed the targeting ability and penetration by CLSM.
 - 4) Examined apoptosis and proliferation of cancer cells via TUNEL and Ki67 assays.
 - 5) Evaluated biodistribution and antitumor efficacy *in vivo* in a 4T1 tumor-bearing mouse model.
- **Research Associate:** Study on a Drug-delivering-drug Platform-mediated Potent Protein Therapeutics via a Non-lysosomal Route. 03/2018-06/2018
 - 1) Studied the apoptosis mechanisms of caspase3 and performed quantitative analyses via western blot.
 - 2) Compared endocytosis and intracellular distribution of the lysosomal and non-lysosomal routes of CLSM.
 - 3) Studied the influence of nanoparticle properties on their endocytosis and performed pharmacodynamics assays in cells.
- **Project Leader:** Baicalein-mediated Delivery of p53 in Therapeutics of Pulmonary Hypertension (Student Entrepreneurship Competition Program). 10/2017-04/2018

- 1) Constructed a gene-drug co-delivery platform for a novel chemotherapy for PAH and confirmed its efficacy in animals.
- 2) Designed synthetic routes to assemble the BCL-p53-βlg complex, in order to reduce pulmonary vascular resistance and apoptosis.
- 3) Performed electrophoresis for a quantitative component analysis of p53, tsp1, and Bax; performed RIP assays to study the regulation mechanism of p53 in relation to these two proteins.
- 4) Established a pulmonary embolism in a rat model and studied the efficacy of anti-PAH.

■ **Research Assistant:** Assembling Nanoplatforms from a CD44-targeted Drug and Liposomes for Dual-targeting of TME and Cancer Cells. 05/2017-01/2018

- 1) Designed self-assembled smart thermosensitive liposomes to function as a dual-targeting platform for co-delivery of MATT and HA-PTX, with the goal of inhibiting metastasis and angiogenesis.
- 2) Designed the synthetic route of a HA-PTX prodrug and compared its effects with those of normal liposomes and free drugs.
- 3) Performed pharmacokinetic experiments in a mouse 4T1 tumor model.

State Key Laboratory of Natural and Biomimetic Drugs, Peking University

Beijing, P. R. China

Advisor: Dr. Wangliang Lu

■ **Research Assistant:** Constructing Nanosized Functional miRNA Liposomes to treat TNBC by Silencing the Slug Gene. 06/2019-08/2019

- 1) Constructed miRNA liposomes using CRISPR/Cas9 to silence the slug gene, in order to inhibit the TGF-β1/Smad pathway, thereby inhibiting the invasiveness and growth of TNBC cells.
- 2) Utilized TargetScan, miRanda/mirSVR, and PicTar to investigate the region of miRNA used to target slug and confirmed that miRNA-203 was the regulator of slug.
- 3) Used solid phase synthesis to prepare miRNA-203 and a negative control miRNA.
- 4) Designed primers, performed PCR and gene sequencing, and analyzed cellular localization of liposomes via qPCR.

Publications

Lyu, Y., Xiao, Q., Li, Y., Wu, Y., He, W., & Yin, L. (2019). "Locked" cancer cells are more sensitive to chemotherapy. *Bioengineering & Translational Medicine*, 4(2), e10130. doi:10.1002/btm2.10130

Affiliations

Student Representative: The 4th Chinese American Society of Nanomedicine and Nanobiotechnology 08/2019

Student Representative: The 12th China Pharmaceutical Conference 11/2018

Student Representative: Generic Drug Consistency Evaluation Training 08/2018

Volunteer

Physician's Assistant: National Hospice Service Program, Fujian, China 09/2016-Present

Teaching Assistant: Science College of CPU 09/2016-09/2018

Pharmacist's Assistant: Outpatient Pharmacy of Nanjing First Hospital 10/2016-06/2017

Pharmacist's Assistant: Department of Pharmacy of Fujian Provincial Hospital 12/2017-02/2018

Skills

Instruments: CLSM, TEM/SEM, PXRD, FCM, CD, MS, NMR, WB, cryo-EM, DLS particle analyzer, Biacore.

Cell Experiments: MTT assay, transwell assay, ChIP/RIP assays, TUNEL assay, intracellular distribution of drugs.

Animal Experiments: Constructing tumor models, drug administration, inducing hyperthermia, bio-sample analyzing.

Gene Engineering: CRISPR, PCR, RT-qPCR, primer design, DNA sequencing.

Software: Origin, SPSS, MATLAB, Chemdraw, Primerbank, ProtParam, Design Expert, Endnote, Mathematica.