

# YI-PIN CHANG

*A meticulous Ph.D. enjoys working at the interface between Chemistry and Biology*

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## SUMMARY & SKILLS

### Protein and Peptide Chemistry

- Automated and manual solid-phase synthesis of peptides, peptidomimetics, cyclic peptides and combinatorial peptide libraries through Fmoc or Boc chemistry (*vide infra* publications #2-6, 9-10, 12, 13 and 15).
- Protein purification (publications #9 and 12)
- Molecular modeling (publications #1, 2, 5, 9 and 13)

### Assay Development

- Surface plasmon resonance (publications #8, 9, 11, 13 and 15)
- Gel electrophoresis (publications #2, 5, 9 and 12)
- High-performance liquid chromatography (publications #1)
- Cell-based assays (publications #1, 3 and 4)
- Histochemical analysis (publications #1)
- Liquid chromatography–mass spectrometry

### Highlights

- A mannosidase inhibitor was identified from an iminosugar library by HPLC, cell-based and histochemical assays that capable of reproduce a human disease in guinea pigs (publication #1).
- The most selective and potent antagonist of a nicotinic acetylcholine receptor was discovered from a positional scanning library (> 100 million compounds) by cell-based assays (publication #3).
- The most potent inhibitor of antitrypsin aggregation was identified from a split-and-mix library by a unique conformation-sensitive gel electrophoresis, which has been used to validate the disease mechanism (publication #5 and 9).
- The binding kinetics and thermodynamics between VanX and peptidemimetics (the molecular weight was the detection limit) were revealed by surface plasmon resonance (publication #13).
- The binding affinity of a streptavidin-binding peptides was enhanced 1000-fold by cyclization and their equilibrium dissociation constants were determined by surface plasmon resonance (publication #15).

## EDUCATION & PROFESSIONAL

### Postdoctoral Research Fellow (Proteomics)

2014-present

The Forsyth Institute (Harvard Medical School affiliate), Cambridge, USA

### Consultant (Peptide Drug Development)

2014-present

TaiPhar Management & Consulting Corp, Hsinchu, Taiwan

### Postdoctoral Research Fellow (Peptide Drug Discovery)

2012-2013

Torrey Pines Institute for Molecular Studies, Port St. Lucie, USA

### Postdoctoral Research Assistant (Glycobiology)

2008-2011

University of Oxford, Oxford, UK

### Ph.D. in Chemistry and Biochemistry

2001-2007

National Chung Cheng University, Chiayi, Taiwan

### Ph.D. Graduate Research Assistant (Bioorganic Chemistry)

2003-2007

National Chung Cheng University, Chiayi, Taiwan

### M.S. in Applied Chemistry

1999-2001

Chinese Culture University, Taipei, Taiwan

### M.S. Graduate Research Assistant (Natural Product Chemistry)

1999-2001

National Research Institute of Chinese Medicine, Taipei, Taiwan

### B.Eng. in Chemical Engineering

1994-1999

Feng Chia University, Taichung, Taiwan

## EDITORIAL & SOCIETIES

### Editorial Board

Journal of Liver and Clinical Research

Hereditary Genetics: Current Research

Journal of Research and Development

2013-present

### Member

American Peptide Society

European Peptide Society

American Chemical Society

2012-present

## UNIVERSITY SERVICES & EXTRACURRICULAR ACTIVITIES

### Safety Committee

2014-present

The Forsyth Institute (Harvard Medical School affiliate), Cambridge, USA

<b>Team Manager and Coach for Eastern Collegiate Taekwondo Conference</b> Harvard University Taekwondo Club, Cambridge, USA	2014–present
<b>Oxford Blue (Varsity Team)</b> Oxford University Taekwondo Club, Oxford, UK	2010-2011
<b>Teaching Assistant for Undergraduate (Organic Chemistry LAB I &amp; II)</b> National Chung Cheng University, Chiayi, Taiwan	2004-2005
<b>Teaching Assistant for Undergraduate (Instrumental Analysis)</b> Chinese Culture University, Taipei, Taiwan	2001-2002
<b>President</b> Feng Chia University Taekwondo Club, Taichung, Taiwan	1995-1997
<b>Publicity for the Student Association</b> Feng Chia University, Taichung, Taiwan	1995-1996

## HONORS & AWARDS

<b>Young Peptide Scientist Award</b> 12 <sup>th</sup> International Chinese Peptide Symposium, Shenyang, China	2012/07/04
<b>Travel Grant Award</b> 5 <sup>th</sup> International Peptide Symposium, Kyoto, Japan	2010/12/04
<b>Finalist of European Young Chemist Award</b> 3 <sup>rd</sup> EuCheMS Chemistry Congress, Nuremberg, Germany	2010/08/29
<b>Service Award</b> Ministry of National Defense, Kaohsiung, Taiwan	2008/08/01
<b>Excellent Dissertation Award in Analytical Chemistry</b> Annual Meeting of Chinese Chemical Society Located in Taipei, Hsinchu, Taiwan	2007/12/16
<b>Best Poster Award &amp; Travel Grant Award</b> 29 <sup>th</sup> European Peptide Symposium, Gdańsk, Poland	2006/09/08
<b>Best Poster Award</b> Annual Meeting of Chinese Chemical Society Located in Taipei, Kaohsiung, Taiwan	2005/11/20
<b>Valedictorian</b> Chinese Culture University, Taipei, Taiwan	2001/06/08
<b>Outstanding Student Award</b> Chinese Culture University, Taipei, Taiwan	2001/06/08

## CONFERENCE TALKS

### Discovery of a selective lysosomal $\alpha$ -mannosidase inhibitor from an imino-sugar library

2014/09/02

33<sup>rd</sup> European Peptide Symposium, Sofia, Bulgaria

### Blocking formation of large protein aggregates by small peptides

2012/07/04

12<sup>th</sup> International Chinese Peptide Symposium, Shenyang, China

### Combinatorially selected 4-mer peptide probes inhibit the pathogenic polymerization of $\alpha_1$ -antitrypsin

2010/12/04

5<sup>th</sup> International Peptide Symposium, Kyoto, Japan

### Inhibiting the pathogenic polymerization of protein: Combinatorial chemistry in combating antitrypsin-related liver and lung diseases

2010/08/31

3<sup>rd</sup> EuCheMS Chemistry Congress, Nürnberg, Germany

### Combinatorial search for anti-protein aggregation ligands, streptavidin-binding cyclopeptides, and cell cycle inhibitors

2005/11/20

Annual Meeting of Chinese Chemical Society Located in Taipei, Kaohsiung, Taiwan

## PUBLICATIONS

1. **Chang, Y. -P.**; Muckeen, M.; Alonzi, D. S.; Sagar, R.; Dickens, A. M.; Chapman, T. M.; Butters, T. D.; Anthony, D. C.; Davis B. G. A selective lysosomal carbohydrate mimetic induces a chemical model of congenital disease. *Nat. Chem. Biol.* **2015**, to be submitted.
2. **Chang, Y. -P.**; Chu, Y. -H. Mixture-based combinatorial libraries from small individual peptide libraries: A paradigm for a conformational disease. *Molecules* **2014**, *19*, 6330-6348. (IF: 2.095; 70/172 in Organic Chemistry)
3. **Chang, Y. -P.**; Banerjee, J.; Christensen, S.; Wu, J.; Gyanda, R.; Houghten, R. A.; Toll, L.; McIntosh, J. M.; Armishaw, C. J. Discovery of potent and selective  $\alpha 3\beta 4$  nicotinic acetylcholine receptor antagonists from a  $\alpha 4/4$ -conotoxin mixture based synthetic combinatorial library. *J. Med. Chem.* **2014**, *57*, 3511-3521. (IF: 5.480; 7/169 in Drug Discovery)
4. Banerjee, J.; Yongye, A. B.; **Chang, Y. -P.**; Gyanda, R.; Medina-Franco, J. L.; Armishaw, C. J. Design and synthesis of  $\alpha$ -conotoxin GID analogues as selective  $\alpha 4\beta 2$  nicotinic acetylcholine receptor antagonists. *Biopolymers* **2013**, *102*, 78-87. (IF: 2.288; 40/172 in Organic Chemistry)
5. **Chang, Y. -P.**; Chu, Y. -H. Blocking formation of large protein aggregates by small peptides. *Chem. Commun.* **2013**, *49*, 4591-4600. (IF: 6.718; 17/417 in Chemistry)
6. Gyanda, R.; Banerjee, J.; **Chang, Y. -P.**; Phillips, A. M.; Toll, M.; Armishaw, C. J. Oxidative folding and preparation of  $\alpha$ -conotoxins for use in high-throughput structure-activity relationship studies. *J. Pept. Sci.* **2013**, *19*, 16-24. (IF: 1.862; 159/346 in Pharmacology)
7. Cheng, C. I.; **Chang, Y. -P.**; Chu, Y. -H. Biomolecular interactions and tools for their recognition: focus on the quartz crystal microbalance and its diverse surface chemistries and applications. *Chem. Soc. Rev.* **2012**, *41*, 1947-1971. (IF: 30.425; 3/417 in Chemistry)
8. **Chang, Y. -P.**; Mahadeva, R.; Patschull, A. O. M.; Nobeli, I.; Ekeowa, U. I.; McKay, A. R.; Thalassinou, K.; Irving, J. A.; Haque, I. U.; Nyon, M. P.; Christodoulou, J.; Gonzalez, A.; Miranda, E.; Gooptu, B. Targeting serpins in high-throughput and structure-based drug design. *Method. Enzymol.* **2011**, *501*, 139-175. (IF: 1.626; 85/390 in Biochemistry)
9. **Chang, Y. -P.**; Mahadeva, R.; Chang, W. -S. W.; Lin, S. -C.; Chu, Y. -H. Small-molecule peptides inhibit Z  $\alpha_1$ -antitrypsin polymerization. *J. Cell. Mol. Med.* **2009**, *13*, 2304-2316. (IF: 3.698; 33/166 in Molecular Medicine)

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10. Tseng, M. -C.; **Chang, Y. -P.**; Chu, Y. -H. Quantitative measurements of vancomycin binding to self-assembled peptide monolayers on chips by quartz crystal microbalance D-Ala-D-Ala. *Anal. Biochem.* **2007**, 371, 1-9. (IF: 2.749; 182/390 in Biochemistry)
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11. **Chang, Y. -P.**; Tseng, M. -C.; Chu, Y. -H. Emerging biosensor-based technology in the field of drug discovery. *Instruments Today* **2007**, 156, 56-67.
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12. **Chang, Y. -P.**; Mahadeva. R.; Chang, W. -S. W.; Shukla, A.; Dafforn, T.; Chu, Y. -H. Identification of a 4-mer peptide inhibitor that effectively blocks the polymerization of pathogenic Z  $\alpha_1$ -antitrypsin. *Am. J. Respir. Cell Mol. Biol.* **2006**, 35, 540-548. (IF: 2.749; 9/130 in Pulmonary and Respiratory Medicine)
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13. **Chang, Y. -P.**; Tseng, M. -J.; Chu, Y. -H. Using surface plasmon resonance to directly measure slow binding of low-molecular mass inhibitors to a VanX chip. *Anal. Biochem.* **2006**, 359, 63-71. (IF: 2.749; 182/390 in Biochemistry)
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14. **Chang, Y. -P.**; Chang, W. -S. W.; Chu, Y. -H.  $\alpha_1$ -Antitrypsin deficiency and the conformational diseases. *Chemistry.* **2005**, 63, 419-430.
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15. **Chang, Y. -P.**; Chu, Y. -H. Using surface plasmon resonance to directly determine binding affinities of combinatorially selected cyclopeptides and their linear analogs to a streptavidin chip. *Anal. Biochem.* **2005**, 340, 74-79. (IF: 2.749; 182/390 in Biochemistry)

## BOOK CHAPTERS

Banerjee, J.; Gyanda, R.; **Chang, Y. -P.**; Armishaw, C. J. In *The chemical synthesis of  $\alpha$ -conotoxins and structurally modified analogs with enhanced biological stability*; Cudic, P., Ed.; Peptide modifications to increase metabolic stability and activity; Humana Press Inc.: New York, USA, **2013**, pp 13-34.

**Chang, Y. -P.** In *Inhibiting pathogenic protein aggregation: Combinatorial chemistry in combating alpha-1 antitrypsin deficiency*; Pignataro, B., Ed.; New strategies in chemical synthesis and catalysis; Wiley-VCH Verlag GmbH: Weinheim, Germany, **2012**, pp 299-323.

## REFERENCES

**Prof. Terry D. Butters** (e-mail: [terry.butters@bioch.ox.ac.uk](mailto:terry.butters@bioch.ox.ac.uk))

Oxford Glycobiology Institute, University of Oxford, Oxford, UK

**Dr. Ravi Mahadeva** (e-mail: [rm232@cam.ac.uk](mailto:rm232@cam.ac.uk))

Department of Medicine, University of Cambridge, Cambridge, UK

**Prof. Yen-Ho Chu** (e-mail: [cheyhc@ccu.edu.tw](mailto:cheyhc@ccu.edu.tw))

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