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Faculty Research Profile: Dr. Shang Jia

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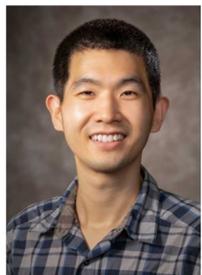
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**Research Assistant
Professor
Dr. Shang Jia**



Emphasis

- Fluorophore
- Bioimaging
- Biosensing

- Protein Chemistry
- Chemoproteomics

Professional Experience

- 2022- Research Assistant Professor, University of Arkansas

- 2019-2022 - Postdoctoral Fellow, Department of Chemistry & Biochemistry, University of California Los Angeles
Research Area: near-infrared imaging in animals, molecular fluorophore design and synthesis

- 2014-2019-Graduate Research Assistant, Department of Chemistry, University of California Berkeley
Research Area: fluorescent sensors, live cell imaging, protein bioconjugation, proteomics

- 2010-2014- Undergraduate Researcher, College of Chemistry, Peking University, China.
Research Area: unnatural amino acid incorporation, biorthogonal and biocompatible chemistry

Education

- B.S- Chemistry, Peking University, Beijing, China
- PhD- Chemistry, University of California- Berkeley

Research Interest

The Jia Group performs studies in bioorganic chemistry. One direction is the development of molecular dyes for purposes in imaging, sensing and bioengineering. We get inspired on existing about photochemistry to create sparkles at the interface of different realms. Another topic is new avenues for chemoproteomics study by developing new tools for more efficient proteomics analysis as well as reactive molecules for probing protein function.

AWARDS

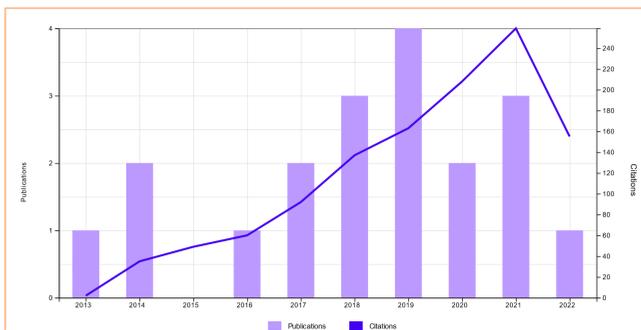
- 2020- Chinese Government Award for Outstanding Self-Financed Students Abroad
- 2016-2017- The Dauben Memorial Endowment Fund Fellowship
- 2016- Gerald K. Branch Fellowship
- 2012-2013- First Class Scholarship of Education Abroad Program
- 2012- 2013- Kwang-Hua Scholarship
- 2010-2011- Zhang Wenjin Scholarship
- 2010-2011- Excellent Students Awards at Peking University

PATENTS

- Sletten, E. M., Jia, S., Garcia, C. A., Pengshung, M. H. & van de Wouw, H. L. Water Soluble Heterocyclic Polymethine Chromophores. Provisional patent application filed.
- Chang, C. J. & Jia, S. Thiophosphorodichloridate Reagents for Chemoselective Histidine Bioconjugation. PCT/US2019/056018

Profile of the Newest CHBC Research Faculty - Welcome Dr. Shang Jia !

CITATION SNAPSHOT



Publications 19 Total	Citing Articles 957 Total 946 Without self-citations	Times Cited 1,160 Total 1,143 Without self-citations	61.05 Average per item	14 H-Index
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TOP CO-AUTHORS

Authors	Count	% of 19
Chang CJ	12	63.158
Ackerman CM	3	15.789
Chan J	3	15.789
Chen PR	3	15.789
Li J	3	15.789
Lin SX	3	15.789
Cao W	2	10.526
Lee J	2	10.526
Toste FD	2	10.526
Wang J	2	10.526
Yang MY	2	10.526
Zhang XY	2	10.526
Alshimiri SA	1	5.263
Aron AT	1	5.263
Atallah TL	1	5.263
Bao SD	1	5.263
Bulek K	1	5.263

TOP SUBJECT AREAS

Research Areas	Count	% of 19
Chemistry	9	47.368
Biochemistry Molecular Biology	6	31.579
Science Technology Other Topics	3	15.789
Life Sciences Biomedicine Other Topics	1	5.263
Pharmacology Pharmacy	1	5.263

TOP CITING INSTITUTIONS/ORGANIZATIONS

Institution/ Organization- USA	No of Times Cited	Institution/ Organization- Other Countries	No of Times Cited
UNIVERSITY OF CALIFORNIA SYSTEM	76	CHINESE ACADEMY OF SCIENCES	70
UNIVERSITY OF CALIFORNIA BERKELEY	39	PEKING UNIVERSITY	62
HOWARD HUGHES MEDICAL INSTITUTE	26	CENTRE NATIONAL DE LA RECHERCHE SCIENTIFIQUE CNRS	26
HARVARD UNIVERSITY	25	UDICE FRENCH RESEARCH UNIVERSITIES	26
UNIVERSITY OF CAMBRIDGE	24	UNIVERSITY OF EDINBURGH	26
HARVARD MEDICAL SCHOOL	18	UNIVERSITY OF CHINESE ACADEMY OF SCIENCES CAS	20
SCRIPPS RESEARCH INSTITUTE	17	UNIVERSIDADE DE LISBOA	18
PENNSYLVANIA COMMONWEALTH SYSTEM OF HIGHER EDUCATION PCSHE	14	CIQUS	13
UNIVERSITY OF CALIFORNIA LOS ANGELES	14	SICHUAN UNIVERSITY	13
CORNELL UNIVERSITY	11	UNIVERSIDADE DE SANTIAGO DE COMPOSTELA	13
JOHNS HOPKINS UNIVERSITY	11	UNIVERSITY OF PITTSBURGH	13
UNIVERSITY OF ILLINOIS SYSTEM	11	CNRS INSTITUTE OF CHEMISTRY INC	12
STATE UNIVERSITY OF NEW YORK SUNY SYSTEM	10	JAPAN SCIENCE TECHNOLOGY AGENCY JST	12
		KYOTO UNIVERSITY	12
		PEKING TSINGHUA CTR LIFE SCI	11
		TECHNICAL UNIVERSITY OF MUNICH	11

TOP CITED PUBLICATIONS

- (1) Li, J.; Yu, J. T.; Zhao, J. Y.; Wang, J.; Zheng, S. Q.; Lin, S. X.; Chen, L.; Yang, M. Y.; Jia, S.; Zhang, X. Y.; et al. Palladium-triggered deprotection chemistry for protein activation in living cells. *Nature Chemistry* **2014**, *6* (4), 352-361, Article. DOI: 10.1038/nchem.1887. **Times Cited- 257**
- (2) Lin, S. X.; Yang, X. Y.; Jia, S.; Weeks, A. M.; Hornsby, M.; Lee, P. S.; Nichiporuk, R. V.; Iavarone, A. T.; Wells, J. A.; Toste, F. D.; et al. Redox-based reagents for chemoselective methionine bioconjugation. *Science* **2017**, *355* (6325), 597-602, Article. DOI: 10.1126/science.aal3316. **Times Cited- 221**
- (3) Li, J.; Jia, S.; Chen, P. R. Diels-Alder reaction-triggered bioorthogonal protein decaying in living cells. *Nature Chemical Biology* **2014**, *10* (12), 1003-+, Article. DOI: 10.1038/nchembio.1656. **Times Cited- 165**
- (4) Li, J.; Lin, S. X.; Wang, J.; Jia, S.; Yang, M. Y.; Hao, Z. Y.; Zhang, X. Y.; Chen, P. R. Ligand-Free Palladium-Mediated Site-Specific Protein Labeling Inside Gram-Negative Bacterial Pathogens. *Journal of the American Chemical Society* **2013**, *135* (19), 7330-7338, Article. DOI: 10.1021/ja402424j. **Times Cited- 112**
- (5) Krishnamoorthy, L.; Cotruvo, J. A.; Chan, J.; Kaluarachchi, H.; Muchenditsi, A.; Pendyala, V. S.; Jia, S.; Aron, A. T.; Ackerman, C. M.; Vander Wal, M. N.; et al. Copper regulates cyclic-AMP-dependent lipolysis. *Nature Chemical Biology* **2016**, *12* (8), 586-+, Article. DOI: 10.1038/nchembio.2098. **Times Cited- 87**
- (6) Xiao, T.; Ackerman, C. M.; Carroll, E. C.; Jia, S.; Hoagland, A.; Chan, J.; Thai, B.; Liu, C. S.; Isacoff, E. Y.; Chang, C. J. Copper regulates rest-activity cycles through the locus coeruleus-norepinephrine system. *Nature Chemical Biology* **2018**, *14* (7), 655-+, Article. DOI: 10.1038/s41589-018-0062-z. **Times Cited- 55**
- (7) Jia, S.; He, D.; Chang, C. J. Bioinspired Thiophosphorodichloridate Reagents for Chemoselective Histidine Bioconjugation. *Journal of the American Chemical Society* **2019**, *141* (18), 7294-7301, Article. DOI: 10.1021/jacs.8b11912. **Times Cited- 52**
- (8) Liao, Y.; Zhao, J. J.; Bulek, K.; Tang, F. Q.; Chen, X.; Cai, G.; Jia, S.; Fox, P. L.; Huang, E.; Pizarro, T. T.; et al. Inflammation mobilizes copper metabolism to promote colon tumorigenesis via an IL-17-STEAP4-XIAP axis. *Nature Communications* **2020**, *11* (1), 15, Article. DOI: 10.1038/s41467-020-14698-y. **Times Cited- 36**
- (9) Guo, L.; Jia, S.; Diercks, C. S.; Yang, X. J.; Alshimiri, S. A.; Yaghi, O. M. Amidation, Esterification, and Thioesterification of a Carboxyl-Functionalized Covalent Organic Framework. *Angewandte Chemie-International Edition* **2020**, *59* (5), 2023-2027, Article. DOI: 10.1002/anie.201912579. **Times Cited- 36**
- (10) Christian, A. H.; Jia, S.; Cao, W.; Zhang, P.; Meza, A. T.; Sigman, M. S.; Chang, C. J.; Toste, F. D. A Physical Organic Approach to Tuning Reagents for Selective and Stable Methionine Bioconjugation. **Times Cited- 26**

THREE MOST RECENT PUBLICATIONS

Spatiotemporal Control of Biology: Synthetic Photochemistry Toolbox with Far-Red and Near-Infrared Light
Published as part of the *ACS Chemical Biology Young Investigators virtual special issue*
Shang Jia and Ellen M. Sletten*

Cite This: <https://doi.org/10.1021/acscchembio.1c00518> | [Read Online](#)

ABSTRACT: The complex network of naturally occurring biological pathways motivates the development of new synthetic molecules to perturb and/or detect these processes for fundamental research and clinical applications. In this context, photochemical tools have emerged as an approach to control the activity of drug or probe molecules at high temporal and spatial resolutions. Traditional photochemical tools, particularly photolabile protecting groups (photocages) and photoswitches, rely on high energy UV light that is only applicable to cells or transparent model animals. More recently, such designs have evolved into the visible and near-infrared regions with deeper tissue penetration, enabling photochemical to study biology in tissue and model animal contexts. This review highlights recent developments in synthetic far red and near infrared photocages and photoswitches and their current and potential applications at the interface of chemistry and biology.

Source: ACS Chem Bio, 2021.

Establishing design principles for emissive organic SWIR chromophores from energy gap laws

Hannah C. Friedman,¹ Emily D. Cosco,^{1,2} Timothy L. Atallah,^{1,3} Shang Jia,^{1,3} Ellen M. Sletten,¹ and Justin R. Caram^{1,3,4*}

SUMMARY
The rational design of bright near and shortwave infrared (NIR-700–1,000; SWIR: 1,000–2,000 nm) emitters remains an open question with applications spanning imaging and photonics. Combining experiment and theory, we derive an energy gap quantum yield master equation (EQME), describing the fundamental limits in SWIR quantum yields (ϕ_F) for organic chromophores. Evaluating the photophysics of 21 polymethine NIR/SWIR chromophores to parameterize the EQME, we explain the precipitous decline of ϕ_F past 900 nm through decreasing radiative rates and increasing nonradiative losses via high-frequency vibrations relating to the energy gap. Using the EQME, we develop an energy-gap-independent ϕ_F NIR/SWIR chromophore comparison metric. We show electron-donating character on polymethine heterocycles results in relative increases in radiative efficiency obscured by a simultaneous redshift. Finally, the EQME yields rational chromophore design insights shown by how deuteration (backed by our experimental results) or molecular aggregation increases SWIR ϕ_F .

Source: Chem, 2021, vol 7(12), pp. 3359-3376

A microtubule-localizing activity-based sensing fluorescent probe for imaging hydrogen peroxide in living cells

Shang Jia¹, Christopher J. Chang^{1,2,3*}

¹Department of Chemistry, University of California, Berkeley, Berkeley, CA 94720, USA
²Department of Molecular and Cell Biology, University of California, Berkeley, Berkeley, CA 94720, USA

ABSTRACT
Hydrogen peroxide (H₂O₂) is a major reactive oxygen species (ROS) in living systems with broad roles spanning both oxidative stress and redox signaling. Indeed, owing to its diverse redox activity, regulating levels (and thus of H₂O₂ generation and trafficking) is critical to determining downstream physiological and/or pathological consequences. The new report the design, synthesis, and biological evaluation of microtubule-targeting fluorescent probe (MTP-F), an activity-based sensing fluorescent probe bearing a microtubule-targeting moiety for detection of H₂O₂ in living cells. MTP-F exhibits a biphasic response to show a collective and robust response to H₂O₂ in aqueous solution and in living cells. Live-cell microscopy experiments establish that the probe co-localizes with microtubules and enables to visualize other changes in levels of H₂O₂, including detection of endogenous H₂O₂ during neuronal open growth factor stimulation. This work adds to the arsenal of activity-based sensing probes for biological analysis that enable selective molecular imaging with subcellular resolution.

Source: Bioorganic & Medical Chemistry Letters, 2021, vol 48, Article #128252

HIGHLY CITED PAPERS

Redox-based reagents for chemoselective methionine bioconjugation

Shixian Lin,^{1,3} Xiaoyu Yang,^{1,3,6} Shang Jia,¹ Amy M. Weeks,⁶ Michael Hornsby,⁶ Peter S. Lee,⁶ Rita V. Nichiporuk,⁴ Anthony T. Iavarone,⁴ James A. Wells,^{6,7} F. Dean Toste,^{1,2,3} Christopher J. Chang^{1,2,3,4,5*}

Cysteine can be specifically functionalized by a myriad of acid-base conjugation strategies for applications ranging from probing protein function to antibody-drug conjugates and proteomics. In contrast, selective ligation to the other sulfur-containing amino acid, methionine, has been precluded by its intrinsically weaker nucleophilicity. Here, we report a strategy for chemoselective methionine bioconjugation through redox reactivity, using oxaziridine-based reagents to achieve highly selective, rapid, and robust methionine labeling under a range of biocompatible reaction conditions. We highlight the broad utility of this conjugation method to enable precise addition of payloads to proteins, synthesis of antibody-drug conjugates, and identification of hyperactive methionine residues in whole proteomes.

Source: Science, 2017, vol 355(6325), pp 597-602. Times Cited-221

Diels-Alder reaction-triggered bioorthogonal protein decaying in living cells

Jie Li^{1,3}, Shang Jia^{1,3} & Peng R Chen^{1,2*}

Abstract
Small molecules that specifically activate an intracellular protein of interest are highly desirable. A generally applicable strategy, however, remains elusive. Herein we describe a small molecule-triggered bioorthogonal protein decaying technique that relies on the inverse electron-demand Diels-Alder reaction for eliminating a chemically caged protein side chain within living cells. This method permits the efficient activation of a given protein (for example, an enzyme) in its native cellular context within minutes.

Source: Nature Chemical Biology, 2014, vol 10(12), pp. 1003-+, Times Cited- 165

PUBLICATIONS - BY FREQUENCY (FROM ISI JOURNAL CITATION REPORTS)

Publication Titles	Count	Category Name	Journal Ranking	Quartile in Category
JOURNAL OF THE AMERICAN CHEMICAL SOCIETY	4	Chemistry, Multidisciplinary	16/179	Q1
NATURE CHEMICAL BIOLOGY	3	Biochemistry & Molecular Biology	13/296	Q1
NATURE COMMUNICATIONS	2	Multidisciplinary Sciences	6/73	Q1
ACS CHEMICAL BIOLOGY	1	Biochemistry & Molecular Biology	130/296	Q2
ANGEWANDTE CHEMIE INTERNATIONAL EDITION	1	Chemistry, Multidisciplinary	15/179	Q1
BIOORGANIC MEDICINAL CHEMISTRY LETTERS	1	Chemistry, Organic-Chemistry, Medicinal	42/63	Q3
CHEM	1	Chemistry, Multidisciplinary	6/179	Q1
ELIFE	1	Biology	8/94	Q1
INORGANIC CHEMISTRY	1	Chemistry, Inorganic & Nuclear	5/46	Q1
JOURNAL OF BIOLOGICAL CHEMISTRY	1	Biochemistry & Molecular Biology	94/296	Q2
METALLOMICS	1	Biochemistry & Molecular Biology	129/296	Q2
NATURE CHEMISTRY	1	Chemistry, Multidisciplinary	8/179	Q1
SCIENCE	1	Multidisciplinary Sciences	2/73	Q1

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