

**Katherine Amberg-Johnson** | Cell & Chemical Biologist  
katieambergjohnson@gmail.com • Cell: 808 428 9703 • www.linkedin.com/in/katieambergjohnson

## SUMMARY

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I am a highly motivated cell/chemical biologist with extensive experience in drug target discovery. As one of the first students in a newly formed lab, I thrive in a fast-paced and collaborative team environment. I am seeking a research science position within a cutting-edge biotechnology company where my intellectual curiosity, technical training, and hard work will have a positive impact on human health.

## RESEARCH EXPERIENCE

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### STANFORD UNIVERSITY

JULY 2014-PRESENT

#### Stanford Bio-X Fellow Ph.D. Student, Advisor: Ellen Yeh

- Performed phenotypic drug screening and identified an antimalarial compound with a novel inhibition phenotype.
- Analyzed whole-genome sequencing data of drug-resistant *T. gondii* strains to determine potential drug targets.
- Utilized CRISPR/Cas9 genome editing to validate drug targets in both *P. falciparum* and *T. gondii*.
- Coordinated collaboration with enzymologists at MIT to establish high-throughput *in vitro* drug assays for inhibition of enzymatic activity.
- Pioneered and optimized live video microscopy experiments of *T. gondii* to study defects in division in collaboration with *T. gondii* geneticists.

### UNIVERSITY OF

### CALIFORNIA-BERKELEY

JULY 2010-MAY 2013

#### Undergraduate Research Assistant, Advisor: David Wemmer

- Performed NMR spectroscopy and optical tweezers experiments to understand the structure and function of the bacterial transcriptional regulatory factor,  $\sigma^{54}$ .

## EDUCATION

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### STANFORD UNIVERSITY

Ph.D. in Microbiology and Immunology  
Anticipated Fall/Winter 2017

### UNIVERSITY OF CALIFORNIA, BERKELEY

B.S. in Microbial Biology, with honors  
August 2009-May 2013

## LEADERSHIP AND TEACHING

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### HUME CENTER FOR WRITING & SPEAKING

MARCH 2017-PRESENT

#### Oral Communications Tutor

Fostered supportive scientific communication through one-on-one mentoring at all stages of the oral presentation process

### BIOCHEMISTRY DEPT.

FEB 2015-PRESENT

#### Flow Cytometry Manager

Provided both instrument training and technical support for 70+ flow cytometry users. Aided users with experimental design and data analysis.

#### Scientific Mentor

Directly mentored 3 first year Ph.D. students and 1 undergraduate researcher on diverse projects including synthetic biology, super-resolution microscopy, protein immunoprecipitation, and gene expression profiling.

**MICROBIOLOGY &  
IMMUNOLOGY DEPT.**  
JAN 2015-MARCH 2016

**Teaching Assistant**  
Techniques in Biotechnology (Jan 2015-March 2015)  
Innate Immunology (Jan 2016-March 2016)

## RELEVANT TECHNICAL AND TRANSFERABLE SKILLS

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Drug screening	Microscopy-live, IFA	Western blotting, ELISA
Drug characterization	NGS analysis	Protein expression & purification
SAR analysis	Data analysis- python	Protein structure modeling-
Tissue culture-primary cells	CRISPR/Cas9 genome engineering	PyMOL
Transfection-stable	Molecular cloning	
FACs cell sorting and analysis	PCR, qPCR, dPCR	
Teamwork	Problem solving	Collaboration
Organizational skills	Publication, grant writing	Communication
Project management	Time management	Market knowledge

## AWARDS AND HONORS

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2016-2019	Bio-X Stanford Interdisciplinary Graduate Fellowship (Stanford University)
2016	Two-Photon and Super-Resolution Microscopy Pilot Grant (Stanford University)
2015, 2016	Biosciences Office of Graduate Education Travel Grant (Stanford University)
2013-2016	Cellular and Molecular Biology Training Grant (Stanford University)
2012	Amgen Scholars Program (UC Berkeley)
2012-2013	Barry Goldwater Scholarship (UC Berkeley)
2011	Science Undergraduate Laboratory Internship (LBNL)
2009-2010	Leadership Award (UC Berkeley)

## PUBLICATIONS

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- **Amberg-Johnson, K.**, Hari, S.B., Ganesan, S.M., Lorenzi, H.A., Sauer, R.T., Niles, J.C., Yeh, E. Small molecule inhibition of apicomplexan FtsH1 disrupts plastid biogenesis in human pathogens. *eLife*. (2017).
  - **Amberg-Johnson, K.** Yeh, E. Kinetics of *Toxoplasma gondii* Apicoplast Loss Upon Treatment with Apicoplast Inhibitors. (in preparation).
  - Foe, I. **Amberg-Johnson, K.** Onguka, O., Bogyo, M. The *Toxoplasma gondii* protein Active Serine Hydrolase 4 (Ash4) is important for parasite growth and the formation of ordered vacuoles *in vitro*. (in preparation).

## PRESENTATIONS

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2017	Toxo-14 Meeting (Oral Presentation, Portugal)
2017	Bay Area Microbial Pathogenesis (Oral Presentation, UCSF)
2016	Biochemistry Postdoc Seminar (Oral Presentation, Stanford)
2016	Microbiology and Immunology Retreat Seminar (Oral Presentation, Stanford)
2015, 2016	Molecular Parasitology Meeting (Poster Presentation, Woods Hole)
2015	Bay Area Meeting on Organelle Biology (Oral Presentation, UCSF)
2013	Undergraduate Honors Thesis Research Symposium. (Oral Presentation, UC Berkeley)
2012	Amgen Symposium (Oral Presentation, UC Berkeley)