

Gizelle N. K. Fauss

512-992-3704 • gizelle.nicole.kayleen@gmail.com • gnkleal@tamu.edu

Resident Address:

3780 Copperfield Dr. Apt. 218
Bryan, TX 77802

Office Address:

Texas A&M University
Dept. of Psychological and Brain Sciences
ILSB BLDG Rm: 3149
301 Old Main Dr.
College Station, TX 77843

Education

Texas A&M University , College Station, TX Bachelors of Science in Psychology Minor in Neuroscience Graduated	August 2012 – December 2015 December 2015
Texas A&M University , College Station, TX Doctor of Philosophy in Neuroscience Department of Psychology Expected Graduation	August 2016 – Present May 2021

Research Experience

Undergraduate Research Assistant Texas A&M University <i>Laboratory of Mark G. Packard, Ph.D.</i> <ul style="list-style-type: none">- Lab exploring the behavioral and neurobiological mechanisms of memory- Use of peripheral and intra-cerebral drug injections to anatomically dissociate multiple memory systems	August 2014 – December 2015
Undergraduate Research Project Texas A&M University <i>Laboratory of Mark G. Packard, Ph.D.</i> <ul style="list-style-type: none">- Trained Long-Evans rats to find food in a hippocampus-dependent place-learning version of the elevated plus-maze task and were then given extinction training using two distinct protocols- Rats given “response-extinction” were provided the opportunity to make the previously reinforced response- Rats given “latent extinction” were simply confined to the previously rewarded spatial location without the reinforcement- Findings indicate that, contrary to classical learning, animals do not need to make the previously reinforced response for extinction to occur- The findings were published as a part of a larger project that is cited below	January 2015 – May 2015
Graduate Research Assistant/Student Texas A&M University <i>Laboratory of James W. Grau, Ph.D.</i> <ul style="list-style-type: none">- Examined the role of brain circuits in the exacerbation of hemorrhage at the injury site of spinally contused rats after administration of pain caudal to the injury- Produced a proposal of experiments that determines the role of descending brain circuits and local cellular mechanisms that underlie and mediate secondary injury after nociceptive stimulation in spinally injured rats- Daily duties include behavioral assessment of rodents and cellular assays in the lab- Currently completing tasks required of a Ph.D. student	June 2016 – Present

Training

Laboratory

- Histology, including:
 - o Cresyl violet tissue staining
 - o Cryostat slicing
 - o Tissue collection and preparation
 - o Western blots
 - o Protein extraction
 - o Bradford Assay
- Small animal surgery, including:
 - o Stereotaxic surgery, prep, and assistance in rats
 - o Use of gas anesthesia in rats (isoflurane)
 - o Use of intraperitoneal anesthesia in rats (ketamine and xylocine)
 - o Intra-cardial perfusion of rats, including brain harvesting
 - o Cervical spinal cord transection and sham operations
 - o Thoracic and cervical contusion and sham operations
 - o Peroneal nerve dissection
 - o Sciatic nerve transection
- Behavioral testing, including:
 - o Morris water maze
 - o Elevated plus-maze task
 - o BBB Locomotor scoring
 - o Beam and Ladder
 - o Instrumental Learning
 - Electrophysiological nerve stimulation
- Pain testing, including:
 - o Von Frey (plantar and girdle)
 - o Non-invasive blood pressure and heart-rate assessment
- Microscopy, including:
 - o Confocal microscopy
 - o Brightfield microscopy
- Care and handling of rats, including:
 - o Care of spinally injured animals

Certifications

- Ohio State University Spinal Cord Injury Training Program (SCITP) 2018
 - o May 6th – 18th, 2018
 - o Learned surgical and behavioral techniques for SCI research
 - o One-on-one faculty interaction and formal lectures regarding my own research or research in the field
 - o Only 12 out of 60 applicants accepted

Organizations

- BRAINS (Building Researchers and Innovators in Neuroscience and Society)
 - o Previously known as SANDI (until May 2018)
 - o Historian for 2017 – 2018 Year
 - o Historian for 2018 – 2019 Year
- SfN (Society for Neuroscience)
 - o Attended conference in November 2016
 - San Diego, California, United States
 - o Attending conference in November 2018
 - San Diego, California, United States

- NNS (National Neurotrauma Society)
 - o Attending conference in August 2018
 - Toronto, Ontario, Canada

Awards

- TAMIN Travel Award 2016: \$600
- ISNR Travel Award 2017: \$500
- TAMIN Travel Award 2017: \$820
- SANDI Travel Award 2017: \$149

Abstracts and Presentations

Undergraduate:

1. SfN Local Chapter Symposium, December 2015
 - a. College Station, Texas, United States
 - b. The memory systems engaged during acquisition determines the effectiveness of different extinction protocols

Graduate:

2. Mission Connect Conference, December 2016
 - a. Houston, Texas, United States
 - b. Brain circuits mediate increases in blood pressure and hemorrhage caused by nociceptive stimulation after spinal cord injury
3. SfN Local Chapter Symposium, December 2016
 - a. College Station, Texas, United States
 - b. Brain circuits mediate increases in blood pressure and hemorrhage caused by nociceptive stimulation after spinal cord injury
4. Annual TAMIN Symposium, April 2017
 - a. College Station, Texas, United States
 - b. Shock-induced hemorrhage and hypertension after spinal cord injury depend on brain systems
5. International Symposium of Neural Regeneration (ISNR) Conference, November 2017
 - a. Monterey, California, United States
 - b. Nociception-induced hemorrhage and hypertension after spinal cord injury depend on brain systems
6. SfN Local Chapter Symposium, December 2017
 - a. College Station, Texas, United States
 - b. Does nociceptive input rostral to a spinal cord injury affect recovery?
7. Mission Connect Conference, December 2017
 - a. Houston, Texas, United States
 - b. Does nociceptive input rostral to a spinal cord injury affect recovery?
8. Annual TAMIN Symposium, April 2018
 - a. College Station, Texas, United States
 - b. Pain-induced hypertension is neither necessary nor sufficient to drive hemorrhage after spinal cord injury
9. Neurotrauma Conference, August 2018
 - a. Toronto, Ontario, Canada
 - b. Noxious stimulation after spinal cord injury (SCI) induces a brain-dependent increase in hemorrhage
10. SfN Conference, November 2018
 - a. San Diego, California, United States
 - b. Noxious stimulation after spinal cord injury (SCI) induces a brain-dependent increase in hemorrhage

Publications

Acknowledgements:

1. Goodman J and Packard M (2015). THE MEMORY SYSTEM ENGAGED DURING ACQUISITION DETERMINES THE EFFECTIVENESS OF DIFFERENT EXTINCTION PROTOCOLS. *Front. Behav. Neurosci.* 9:314.

Publications: