



Nov 02, 2021

# GCaMP6F Fluorescence in Ex Vivo Intestinal Preparation

Bryan Yoo<sup>1</sup>, Jessica Griffiths<sup>1</sup>, Sarkis Mazmanian<sup>1</sup><sup>1</sup>California Institute of Technology

1

[dx.doi.org/10.17504/protocols.io.bzqap5se](https://dx.doi.org/10.17504/protocols.io.bzqap5se)

Mazmanian Lab

Jessica Griffiths

Protocol for GCaMP6F imaging in ex vivo intestinal tissue used in Yoo et al 2021

DOI

[dx.doi.org/10.17504/protocols.io.bzqap5se](https://dx.doi.org/10.17504/protocols.io.bzqap5se)

Bryan Yoo, Jessica Griffiths, Sarkis Mazmanian 2021. GCaMP6F Fluorescence in Ex Vivo Intestinal Preparation. **protocols.io**  
<https://dx.doi.org/10.17504/protocols.io.bzqap5se>



Center for Environmental and Microbial Interactions (CEMI)

Grant ID: n/a

Emerald Foundation

Grant ID: n/a

Heritage Medical Research Institute

Grant ID: n/a

Department of Defense

Grant ID: PD160030

National Institutes of Health

Grant ID: GM007616 and DK078938

Aligning Science Across Parkinson's

Grant ID: ASAP-000375

protocol ,

Nov 02, 2021

- 1 Small intestinal tissue was quickly harvested from ChAT-Cre mice, flushed and placed in oxygenated (95% O<sub>2</sub>, 5% CO<sub>2</sub>), ice cold Krebs-Henseleit solution for 1 hour followed by 15 min at room temperature.
- 2 A segment was cut along the mesenteric attachment, and pinned flat (mucosa facing down) on a Sylgard-lined recording chamber (Warner Instruments, PH1) in oxygenated Krebs-Henseleit solution.
- 3 C21 was added at 10nM and GCaMP6F fluorescence was detected on an upright microscope (Zeiss, Oberkochen, Germany-Examiner D1).