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Generation of knockout and rescue cell lines using CRISPR-Cas9 genome editing

William Hancock-Cerutti[1,2,3,4,5], Jun Hyun Park[1,2,3,6,5], Pietro De Camilli[1,2,3,5]

[1]Departments of Neuroscience and of Cell Biology, Yale University School of Medicine, New Haven, Connecticut 06510, USA;

[2]Howard Hughes Medical Institute;

[3]Program in Cellular Neuroscience, Neurodegeneration and Repair, Yale University School of Medicine, New Haven, Connecticut 06510, USA;

[4]Interdisciplinary Neuroscience Program and MD-PhD Program, Yale University School of Medicine, New Haven, Connecticut 06510, USA;

[5]Aligning Science Across Parkinson's (ASAP) Collaborative Research Network, Chevy Chase, MD, 20815;

[6]Interdisciplinary Neuroscience Program, Yale University School of Medicine, New Haven, Connecticut 06510, USA

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William Hancock-Cerutti

This protocol describes the genetic modification of cultured cells using CRISPR-Cas9, including synthesis of reagents, transfection, selection and screening of single clones, and sequencing of genomic DNA to confirm mutations. In addition, this protocol describes CRISPR-Cas9 mediated repair of single base deletions or insertions by CRISPR-Cas9.

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