## Cell Centered Database University of California, San Diego maryann@ncmir.ucsd.edu

#### Microscopy Product #:4045 112006aaa

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Image2D	Reconstruction	Segmentation

## **Project Information:**

PROJECT_ID	P1723
PROJECT_NAME	Localization of Metabotropic Glutamate Receptors in Alpha Synuclein Overexpressing Mouse
PROJECT_DESCRIPTION	Characterization of staining for mGluR5 glutamate receptor in animal model of Parkinsonian disorders
LEADER	Diana Price
FUNDING_AGENCY	Branfman Family Foundation
PROJECT_START_DATE	
PROJECT_END_DATE	
COLLABORATORS	Edward Rockenstein, Eliezer Masliah, Mark Ellisman
PUBLICATION1	
PUBLICATION2	
PUBLICATION3	

Experiment Information -	
PURPOSE	To determine the relationship between mGluR5 and alpha synuclein
	staining in different lines of alpha synuclein overexpressing mouse
TITLE	Comparison of mGluR5 and synuclein staining
EXPERIMENTER	Diana Price
EXPERIMENT_NAME	
EXPERIMENT_DATE	

Subject Information -	
GROUP_BY	genetic modification
SUBJECT_NAME	A53T15+
FIXATION_METHOD_ID	
SCIENTIFIC_NAME	mus musculus
SPECIES	mouse
STRAIN	C57BL/6-DBA/2
AGE	days
AGECLASS	adult
ANIMAL_NAME	
LITTER_ID	
SEX	unspecified
VENDOR	Eliezer Masliah
WEIGHT	grams

Tissue -	
ANATOMIC_LOCATION	unknown
MICROTOME	Vibratome
ORIENTATION	coronal
THICKNESS	80 um
TISSUE_PROD_STORAGE	
EXTERNAL_FILE_NAME	
TISSUE_GROUP_TYPE	triple label

Microscopy Product Information -		
MICROSCOPY_PRODUCT_ID	4045	
IMAGE_BASENAME	112006aaa	
CREATE_DATE		
INSTRUMENT	Olympus Fluoview 1000	
MICROSCOPE_TYPE	LASER SCANNING CONFOCAL	
PLANE_COUNT	1	
PRODUCT_TYPE	SURVEY	
PURL		
SESSION_NAME		
TELESCIENCE_SRB	P1723/Experiment_3482/Subject_252/Tissue_365/Microscopy_4045	
X_RESOLUTION	.207 um/pixels	
Y_RESOLUTION	.207 um/pixels	
XSIZE	1024	
YSIZE	1024	

## **Protocol:**

Specimen processing: Tissue section acquisition from transgenic animals

Animals were deeply anesthetized with Nembutal¿ (pentobarbital) and perfused via intracardiac catheterization. Perfusion with oxygenated Ringer's solution containing 250U/ml heparin, 0.2 mg/ml xylocaine and 1% dextrose was followed 4% paraformadehyde in 0.1 M phosphate buffer solution (PBS) (both at 37 degrees Celsius). The brains were carefully removed from the skull and postfixed for 1 hour in the same fixative used in the perfusion. The brain was blocked and cut into 2 mm thick sections using an acrylic brain matrix (David Kopf; Tujunga, CA) to facilitate reproducibility of sections. These thick sections were then sectioned into 80 micron thick coronal sections using a Vibratome (VT1000E, Leica Microsystems, Wetzlar , Germany).

#### Specimen processing: Immunocytochemistry

Tissue sections were incubated with monoclonal anti- a-syn (1:250; BD Transduction Laboratories, San Diego, CA, Catalog #AB610787) and rabbit anti-mGluR5 (1:250; Chemicon, Temecula, CA, Catalog #AB5675) followed by incubation with donkey anti-mouse Alexa Fluor 488 (1:100, Molecular Probes, Carlsbad, CA) and donkey anti-rabbit RRX (1:100, Jackson Immunoresearch Laboratories, Inc., West Grove, PA, USA) overnight at 4C. The immunolabeling procedure consisted of the following steps: (1) 6x5 min rinses in 0.1 M PBS; (2) 1 hr blocking step in PBS containing 3% normal donkey (NDS), 0.1% Triton X-100, 1% fish gelatin, and 1% BSA; (3) 48 hr incubation in primary antibodies diluted in working buffer (PBS, 1% NDS) at 20 degrees C; (4) 6 x 5 minute rinsed in working buffer; (5) 24 hr incubation in working buffer containing donkey anti-mouse Alexa Fluor 488 (Molecular Probes, Carlsbad, CA) and donkey anti-rabbit RRX (Jackson Immunoresearch Laboratories, Inc., West Grove, PA, USA). (6) 6 x 10 min rinses in working buffer; (7) 3 x 10 min rinses in PBS; (8) the sections were free floated onto slides and coverslipped using ProLong mounting media (Invitrogen Molecular Probes, Carlsbad, CA) with DAPI nuclear stain. Controls for the mGluR5 antibody experiments included both preabsorption with the control peptide (Chemicon, Catalog #AG374), as well as primary omission studies, which both revealed a lack of non-specific staining. Controls for other antibodies used were performed via omission of primary antibodies, and revealed no non-specific staining. All steps were conducted at 4 degrees C, on wet ice and with ice-cold solutions.

# **Specimen Preparation Information:**

Specimen Preparation Information -	
PROTOCOL_ID	15692
PROTOCOL_NAME	Immunolabeling P1723
PROTOCOL_DESCRIPTION	Double labeling immunolabeling of alpha synuclein and mGIR5
Protocol Steps:	1)Molecular Localization(15740)
	2)Molecular Localization(15749)
	3)Stain(15765)
	4)Chemical(15690)
	5)Microtomy(15691)

Molecular Localizat	tion (15740)	
Molecular Target	MOLECULAR TARGET ID: 15741 MOLECULAR LOCALIZATION ID: 15740 MOLECULE: synuclein ISO FROM: alpha MOLECULAR CLASS: protein ABBREVIATION: Snca ENTREZ_ID: 20617	
Probe used	PROBE ID: 15742   CONTROLS: omitted primary antibody   Antibody ID: 15743   Clonality: monoclonalRaised in animal: mouse   Antibody type: IgG   Reagent (15696)	
	Reagent name	anti alpha synuclein antibody
	Temperature	
	Chemical	Chemical ID: 15695 Chemical name: anti alpha synuclein antibody Vendor: BD Transduction Laboratories Concentration: .25 % Catalog number: AB610787
		Chemical ID: 15704 Chemical name: normal donkey serum Concentration: 1 %
		Chemical ID: 24 Chemical name: phosphate buffer Concentration: .1 M pH: 7.4
		Chemical ID: 31 Chemical name: saline Concentration: .9 % Chemical notes: normal saline
Detection method	Molecule reagent ID: 15709 Molecular type: antibody Chromagen :Alexa 488	

Molecular Localization (15749)		
Molecular Target	MOLECULAR TARGET ID: 15750 MOLECULAR LOCALIZATION ID: 15749 MOLECULE: metabotropic glutamate receptor ISO FROM: 5 MOLECULAR CLASS: protein ABBREVIATION: GRM5 ENTREZ_ID: 108071	
Probe used	PROBE ID: 15751 CONTROLS: omitted pr	imary antibody
	Antibody ID: 15752 Clonality: polyclonalRaised in animal: rabbit Antibody type: IgG	
	Reagent (15714)	
	Reagent name	anti mGluR5 antibody
	Temperature	
	Chemical	Chemical ID: 15719 Chemical name: anti mGluR5 antibody Vendor: Chemicon Concentration: .25 % Catalog number: AB5675
Detection method	Molecule reagent ID: 15721 Molecular type: antibody Chromagen :Rhodamine Red X	

Stain (15765)		
Stain ID	15765	
Prepared by	Diana Price	
Temperature		
Stain notes	DAPI is dissolved in Pro coverslipping	Long Mounting medium and applied at time of
Reagent	Reagent (15760)	
	Reagent name	DAPI in ProLong
	Temperature	
	Chemical	Chemical ID: 15758 Chemical name: DAPI Concentration: Chemical ID: 15759 Chemical name: ProLong mounting medium Vendor: Molecular Probes Concentration:

Chemical Fixation (15690)		
Time of fixation		
Temperature	37 C	
Fixative volume		
Fixation method	perfusion	

Microtomy (15691)	
Microtome	0
Thickness	80 um
Temperature	
Embedding agent	0
Microtomy notes	Vibratome

Specimen Description -		
ANATOMICAL_DETAIL	15777	
ATLAS_COORD	, ,	
ORGAN	brain	
REGION	hippocampus	
SYSTEM	central nervous system	

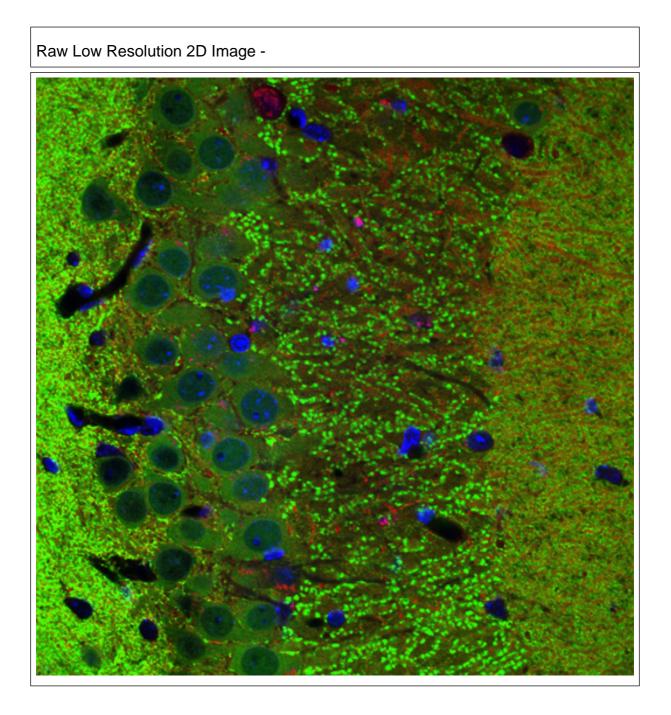
# Imaging Parameters:

Image Type -		
OPTICAL_SECTION_SERIES	15776	
OPTICAL_SECTION_SERIES_D	Only a single optical section was obtained for each section.	
ESC		

Light Microscopy Product -		
LMPRODUCT_ID	15778	
IMMERSION_MEDIUM	oil	
LENS	Olympus PlanApo 60X oil	
LENS_MAGNIFICATION	60 X	
MOUNTING_MEDIUM	Prolong (Molecular Probes)	
NUMERICAL_APERTURE	1.4	
LM_NOTES	DAPI was added to the mounting medium	

Confocal channel (15781)		
Confocal image ID	15781	
Fluorophor	Alexa 488	
Color	Green	
Exitation wavelength	488 nm	
Emission wavelength	520 nm	
Molecular Localization (15740)		
Molecular Target	MOLECULAR TARGET ID: 15741 MOLECULAR LOCALIZATION ID: 15740 MOLECULE: synuclein ISO FROM: alpha MOLECULAR CLASS: protein ABBREVIATION: Snca ENTREZ_ID: 20617	

## Raw 2D Image



Raw 2D Image -	
IMAGE2D_ID	15805
IMAGE_DATE	2006-11-20 00:00:00.0
IMAGE_DESC	Zip archive containing the 3 channel image file in tiff format (112006aaa_RGB.tiff). Also included are the .oif header file generated by the Olympus Fluoview, which give additional detail on microscope settings.
IMAGE_FILE_FORMAT	tiff
IMAGE_FILE_NAME	/usr/local/tomcat/webapps/FileUploadTool/temp_file_upload/112006 aaa_img.jpg
RAW_DATA_FILE	/telescience/home/CCDB_DATA_USER.portal/P1723/Experiment_3 482/Subject_252/Tissue_365/Microscopy_4045/112006aaa_img.zip
THUMBNAIL_DESC	Triple labeled confocal image of hippocampal area CA3 of a transgenic mouse engineered to overexpress a mutant form of human alpha synuclein (A53T), immunolabeled for mGluR5 (red), alpha synuclein (green) and counterstained with DAPI (blue) to reveal cell nuclei.
THUMBNAIL_FILE	/usr/local/tomcat/webapps/FileUploadTool/temp_file_upload/112006 aaa_img_thmb.jpg
X_RESOLUTION	.207 um/pixel
Y_RESOLUTION	.207 um/pixel
X_SIZE	1024 pixels
Y_SIZE	1024 pixels
NOTES	Pixel resolution was read from the .oif file submitted with the tiff image but has not been verified.

## USER AGREEMENT

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#### **USER NOTIFICATION**

For large size image data, it will take several minutes to download, please be patient. Thanks!

### ACKNOWLEDGEMENT

Data used from the CCDB should be appropriately referenced, including both the author of the data and the CCDB. If the data were from a published study, the reference is included in the database record. The following reference should be cited for the CCDB:

Martone, M. E., Gupta, A., Wong, M., Qian, X., Sosinsky, G., Ludaescher, B., and Ellisman, M. H. A cell centered database for electron tomographic data. J. Struct. Biology 138: 145-155, 2002.

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Maryann Martone