

APR 05, 2024

OPEN ACCESS



DOI:

dx.doi.org/10.17504/protocols.io.k xygxprqkl8j/v1

Protocol Citation: Scott Vermilyea 2024. Brain Homogenization and MSD Protocol for Mouse Brain and Serum. protocols.io https://dx.doi.org/10.17504/protoc ols.io.kxygxprqkl8j/v1

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Protocol status: Working We use this protocol and it's working

Created: Nov 12, 2021

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ABSTRACT

This protocol details about the homogenization and MSD protocol of the mouse brain using various panel kits.

ATTACHMENTS

320-681.pdf



Last Modified: Apr 05, 2024

MATERIALS

PROTOCOL integer ID: 55080

Reagent, Kits and Equipment List

Keywords: Homogenization,

1.

MSD Protocol, Mouse Brain and Serum, ASAPCRN

Equipment

gentleMACS™ C Tubes

NAME

C Tubes

TYPE

gentleMACS™

BRAND

130-093-237

SKU

 $https://www.miltenyibiotec.com/US-en/products/gentlemacs-c-tubes.html\#gref^{LINK}$

- 2. RIPA Lysis and Extraction Buffer Thermo Fisher Catalog #89901
- 3. Halt PIC (ThermoFisher 1861279)

4.

Equipment

Direct Detect® Assay-free Cards

NAME

TYPE

Assay-free Cards

Direct Detect®

BRAND

DDAC00010-GR

SKU

https://www.emdmillipore.com/US/en/product/Direct-Detect-Assay-free-Cards,MM_NF-DDAC00010-GR?bd=1

SPECIFICATI

Membrane cards used for protein quantitation in the Direct Detect®

ONS

Infrared Spectrometer

- 5. HCl (Millipore Sigma H1758-500ML)
- 6. NaOH (Millipore Sigma 415413-500ML)
- 7. X HEPES Sigma Aldrich Catalog #83264-500ML-F

8.

Proinflammatory Panel 1 mouse MESO SCALE DIAGNOSTICS, LLC. Catalog #MSD K15048D

9.

Cytokine Panel 1 Mouse Kit MESO SCALE DIAGNOSTICS, LLC. Catalog #K15245D

10.

U-PLEX TGF-β Combo mouse **MESO SCALE DIAGNOSTICS**, **LLC. Catalog #**MSD K15242K-2

11.

Equipment

Octo Dissociator with Heaters

NAME

Octo Dissociator

TYPE

gentleMACS™

BRAND

130-096-427

SKU

https://www.miltenyibiotec.com/US-en/products/gentlemacs-octo-dissociator-with-heaters.html

tor- K

12.

13. Meso Sector S 600 (MSD 1201)

Brain Homogenization Using Miltenyi gentleMACS Octodissociator

43m

- 1 Weight out brain using pre-chilled C-tube.
- 2 Use 🚨 2 mL RIPA (1x Halt PIC) per whole brain, 🚨 1 mL RIPA (1x Halt PIC) per half brain.
- 3 Homogenize brain using "protein_01_01" routine on Miltenyi gentle MACS Octodissociator.
 - **3.1** Repeat homogenization routine if not fully homogenized.

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- 4 30m Vortex briefly and incubate 00:30:00 00:30:00 00:30:00
- - 5 3m Centrifuge at 3500 rpm, 4°C, 00:03:00 .
 - - Transfer homogenate to two 4 1.5 mL eppendorf tubes.

 - 7 10m Vortex briefly and centrifuge at 14000 x g, 4°C, 00:10:00. •
 - Transfer supernatant to new 4 1.5 mL eppendorf tubes
 - Perform 1:10 serial dilution.

9.1

9.2 Dilute using \perp 10 μ L sample to \perp 90 μ L RIPA (1x Halt PIC) to achieve 1:100 dilution. (2/2)

Dilute using \perp 10 μ L sample to \perp 90 μ L RIPA (1x Halt PIC) to achieve 1:100 dilution. (1/2)



- 10 Blot Δ 2 μL diluted sample onto Direct Detect assay card (run duplicates).
- 11 Use Δ 2 μL RIPA (1x Halt PIC) as blank.
- 12 Read assay card on Direct Detect Spectrometer.
- Detectable range is \triangle 0.2 mg/mL to \triangle 5.0 mg/mL , dilute or concentrate as needed.
- 14 Average all readings for each sample to determine concentration.

Brain MSD Protocol Proinflammatory Panel 1 mouse and Cytokine Panel 1...

- 15 Using concentration determined, calculate volume needed for 🚨 200 µg protein.

- 17 Combine $\[\[\] \]$ sample and $\[\] \[\] \]$ MSD diluent in U-bottom plate.





Load 4 50 µL diluted sample to MSD plate per duplicate.



19 Proceed with MSD assay protocol from manufacturer.

Brain MSD Protocol U-PLEX TGF-β Combo mouse

10m

20 Using concentration determined, calculate volume needed for Δ 200 μg protein.

Use RIPA(1x PIC) to dilute \perp 200 µg protein to \perp 100 µL (\perp 2 undetermined).



22 Load Δ 100 μL sample into U-bottom plate.



Add Δ 20 μL [M] 1 Molarity (M) HCl into U-bottom plate and shake for 00:10:00 at 25 °C.



Neutralize sample with Δ 14 μL [M] 1.2 Molarity (M) NaOH in [M] 0.5 Molarity (M) HEPES.



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25 Combine \perp 100 µL treated sample and \perp 100 µL MSD diluent in new U-bottom plate.



26 Load 4 50 µL diluted sample to MSD plate per duplicate.



27 Proceed with MSD assay protocol from manufacturer.

Serum MSD Protocol Proinflammatory Panel 1 mouse

28 Add 🚨 25 µL MSD Diluent to MSD Plate per duplicate.



Add 🚨 25 µL serum sample to MSD Plate per duplicate.



29

30 Proceed with MSD assay protocol from manufacturer.

Serum MSD Protocol Cytokine Panel 1 mouse

31 Add 🚨 37.5 µL MSD Diluent to MSD Plate per duplicate.





32 Add 🚨 12.5 µL serum sample to MSD Plate per duplicate.



33 Proceed with MSD assay protocol from manufacturer.

Serum MSD Protocol U-PLEX TGF-β Combo mouse Add Δ 50 μL serum sample to U-bottom plate. Add Δ 10 μL [M] 1 Molarity (M) HCl into U-bottom plate and shake for ৩0:10:00 at 25 °C. 10m

- Neutralize sample with Δ 7 μL [M] 1.2 Molarity (M) NaOH in [M] 0.5 Molarity (M) HEPES.
- 37 Add Δ 25 μL MSD Diluent to MSD Plate per duplicate.
- Add 25 pc Wisb blidefit to Wisb Flate per duplicate
- Add \triangle 25 μ L treated serum sample to MSD Plate per duplicate.
- **39** Proceed with MSD assay protocol from manufacturer.

