

The Effects of Maternal Autoimmune Dysregulation on ASD Associated Behaviors in Offspring

Team Rugrats

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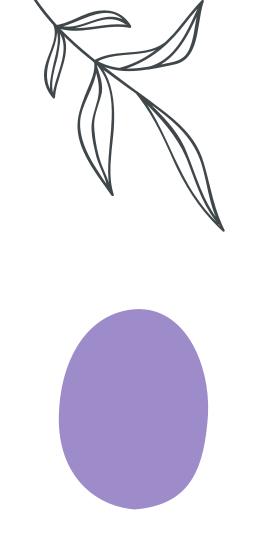
Overview

01 Goals & Hypothesis

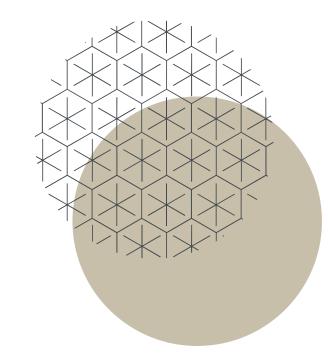
02 Specific Aims 03 Research Design

04 Methods & Results

Interpretations & Future Directions



01 Goals & Hypothesis



Goal

To investigate whether ASD - like behaviors arise from exposure to maternal autoantibodies in lupus mice (NZB).

Hypothesis

The lupus mice will display ASD like behaviors.



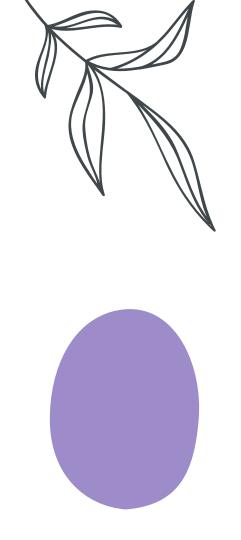
Specific Aims

Specific Aim:

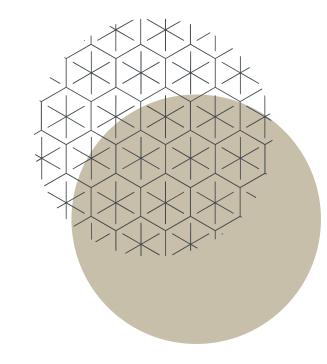
To evaluate Lupus mice models for repetitive, social, and motor behaviors at 3 and 12 weeks old and compare the behaviors to those associated with ASD.

Working Hypothesis:

The Lupus mice will demonstrate ASD - like behaviors at a higher rate than the control mice, with the 3 week old lupus mice displaying the most deficits.



Research Design



Research Design

Repetitive Behavior (Marble Test)



Tests the mice' repetitive behavior patterns by surveying how many marbles are buried.

Motor Function Test (Beam Test)



Tests the motor function of the mice by surveying their ability to walk across the beam; coordination.

Social Behavior Test (Partition)



Tests their sociability (amount of time they spend with another mouse).

Timeline

3 and 12 week old Control & Lupus Mice

1 day

2 days

1 day



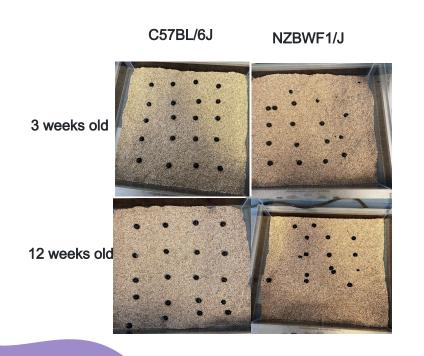
1 trial of marble test; 30 minutes each Day 1: Training Day 3 trials; 10 minute intervals

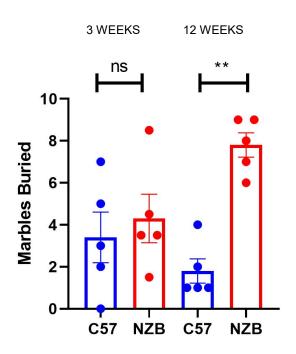
Day 2: Test Day, 2 successful trials

1 trial of partition test; Habituation: 10 minutes of familiarity 15 minutes of testing



NZB Present Repetitive Behavior at 12 Weeks But Not 3 Weeks



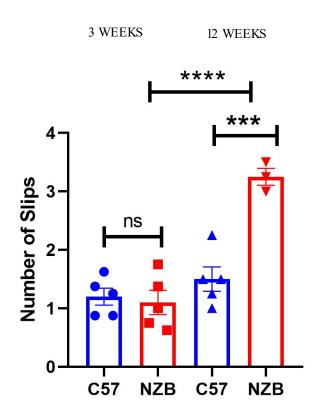


NZB Present Motor Deficits at 12 Weeks But Not 3 Weeks

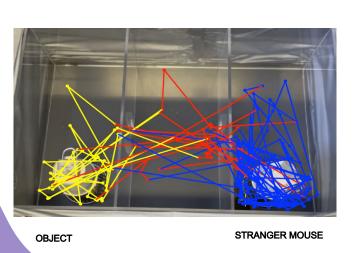
NZBWF1J

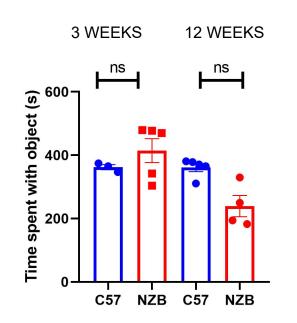


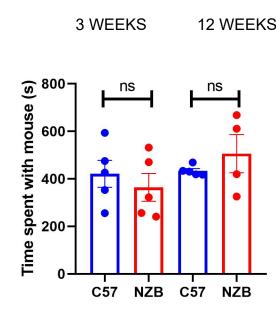
12 WEEK OLD



NZB Mice Do Not Present Social Deficits









Interpretations

- + At 3 weeks, the lupus mice show no deficits while the 12 week old lupus mice demonstrated repetitive behavior and impaired motor function, however no social deficits.
- + Our hypothesis was partially supported by our data since one of the age groups presented ASD like behavior

Future Directions:

- Larger sample size
- + Analysis of brain morphology

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THANK YOU!







Questions?

