



Psychiatry and Neuroscience Seminar Series 2019

Pr Francesco Papaleo

(Host R Piskorowski)



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**Perinatal antinflammatory
oxytocin effects ameliorates
developmental trajectories in
22q11.2 deletion**

Friday March 22nd, 2019, noon

Amphitheater, 102-108 rue de la santé - 75014 Paris

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22q11.2 deletion syndrome (DS) is a chromosomal anomaly which causes a congenital malformation disorder whose common features include cardiac defects, palatal anomalies, facial dysmorphism, developmental delay and immune deficiency. Approximately 1% of patients with schizophrenia have 22qDS. Using genetically modified mice for genes relevant to schizophrenia, Papaleo's lab will then employ a combined approach beginning at the behavioral level and culminating at the cellular and molecular levels. Cognitive abnormalities are core enduring symptoms in schizophrenia, dramatically contribute to poor functional outcomes in patients and currently represent a great "unmet therapeutic need". Papaleo's lab mainly focus its work on behavioral cognitive analyses and relative neuronal correlates

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- Cognitive Neuroscience
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