



Background

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Fixed-interval (FI) schedules of reinforcement capture patterns of anticipatory responding

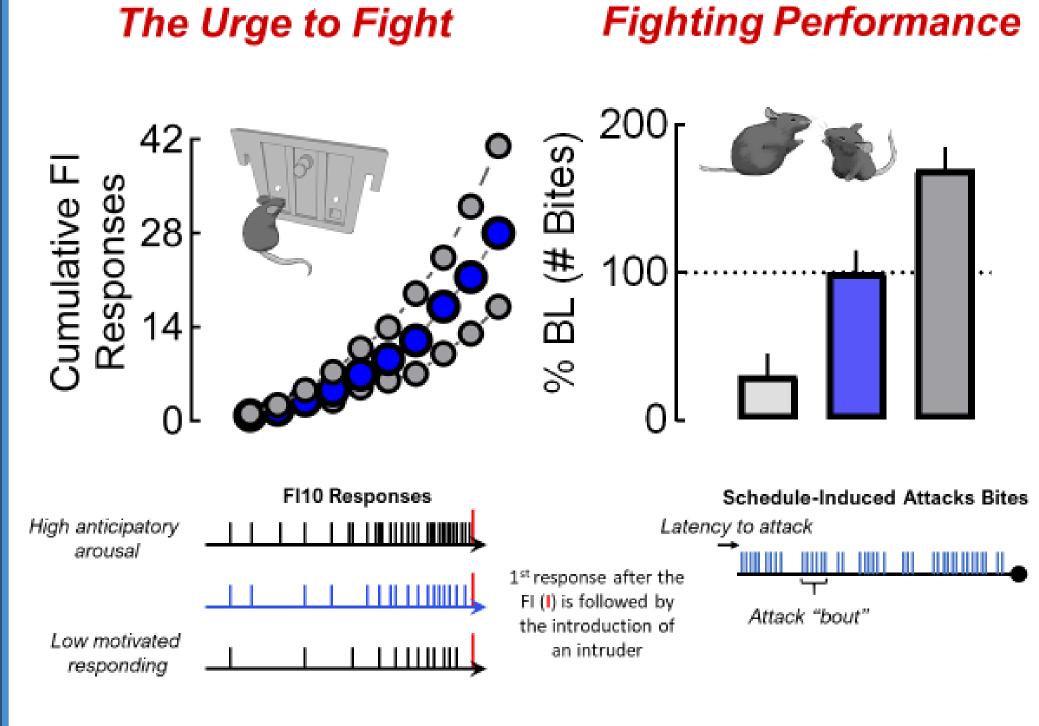
>The rate and associated scalloped "pattern" produced during FI schedules reflect the intensity of the motivation to obtain a reward The molecule CRF appears to be critical for the arousal associated with the motivation to fight

Ventral Tegmental Area (VTA) associated with reward and motivation How does the activity of CRF-containing cells that terminate in the VTA of the brain modulate the motivation for aggression?

Methods

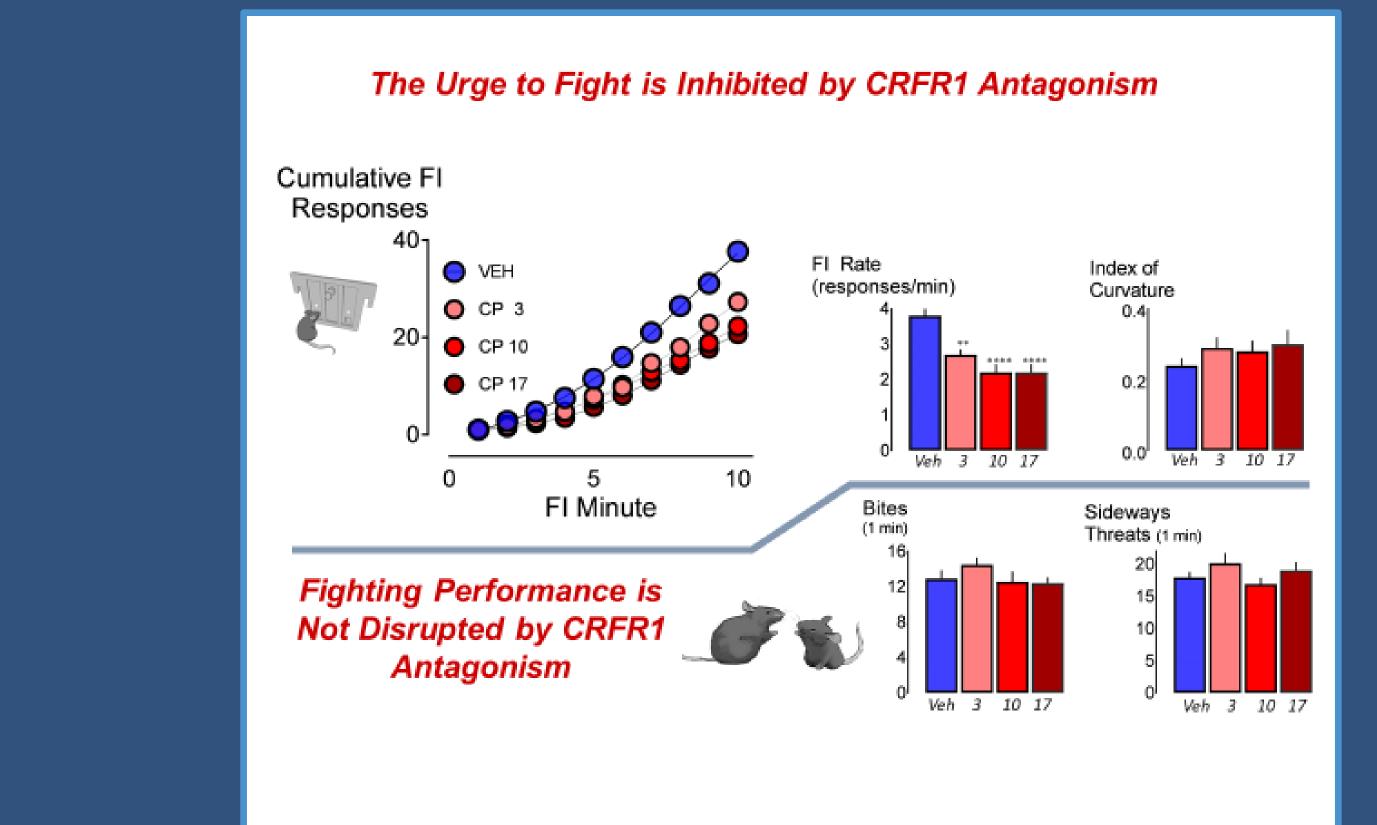
C57/BL6J mice were trained to work under the demands of a FI10 schedule for the opportunity to fight an intruder:

- > CRFR1 antagonism was evaluated during the motivation to fight and during fighting performance
- > After FI, functional activation of cells in the VTA, LH, and CeA was examined via protein markers of cellular activity
- > Using CRF-Cre mice, AAV-DIO-ChR2 or -NpHR was used to control the activity CRF neurons along the LH \rightarrow VTA pathway



Results

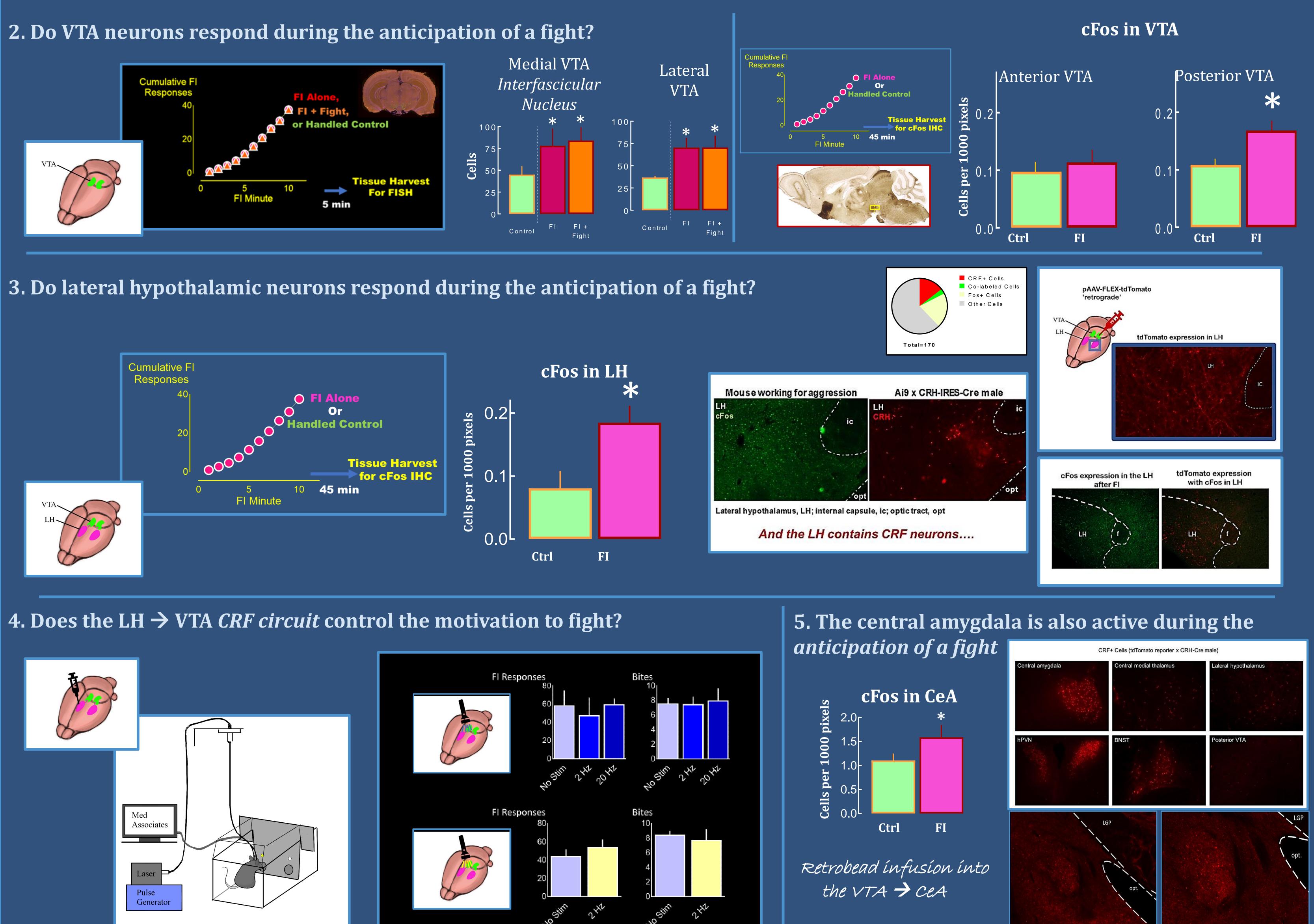
1. Effect of CRFR1 antagonism in C57/BL6J on the anticipation to fight

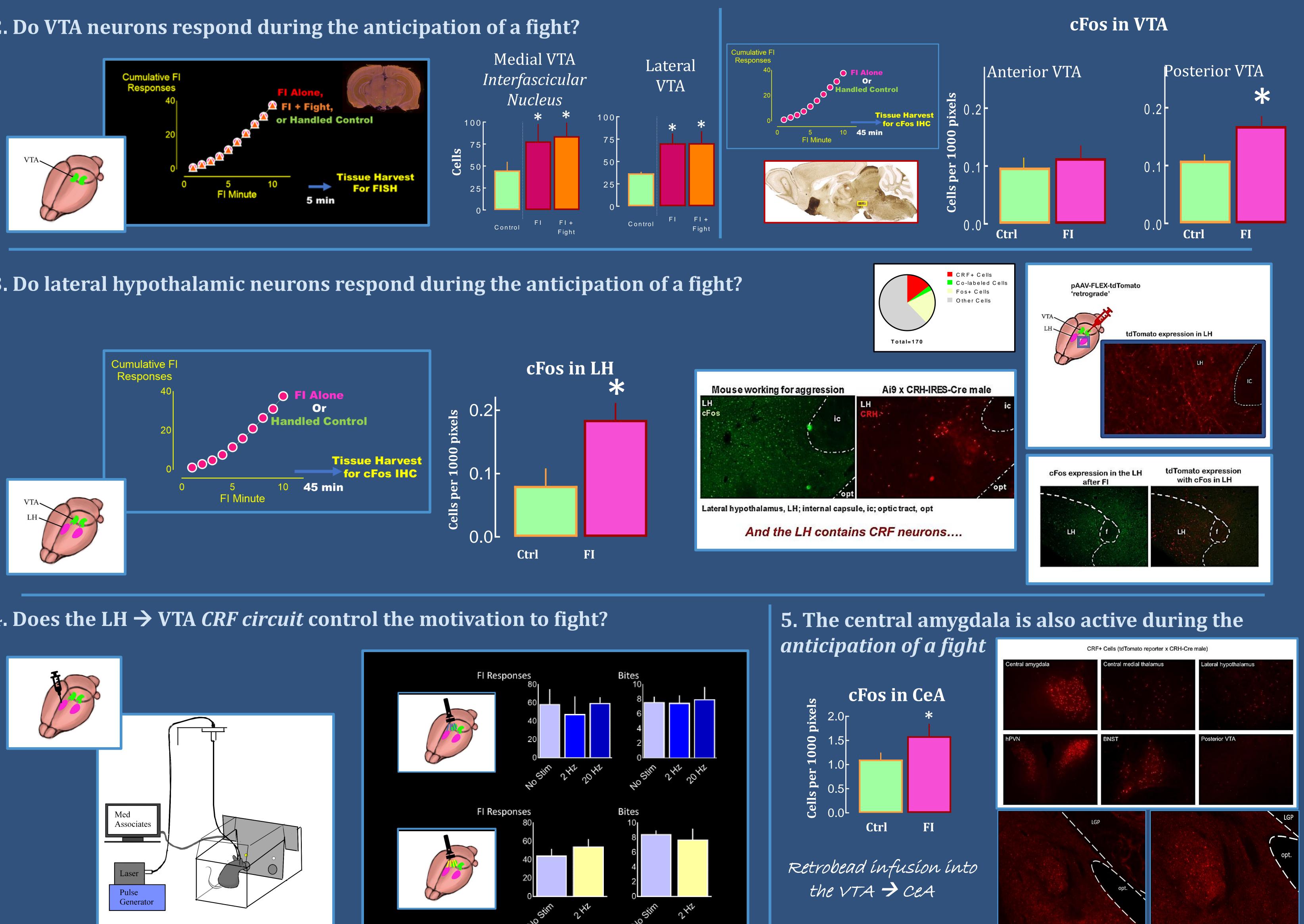


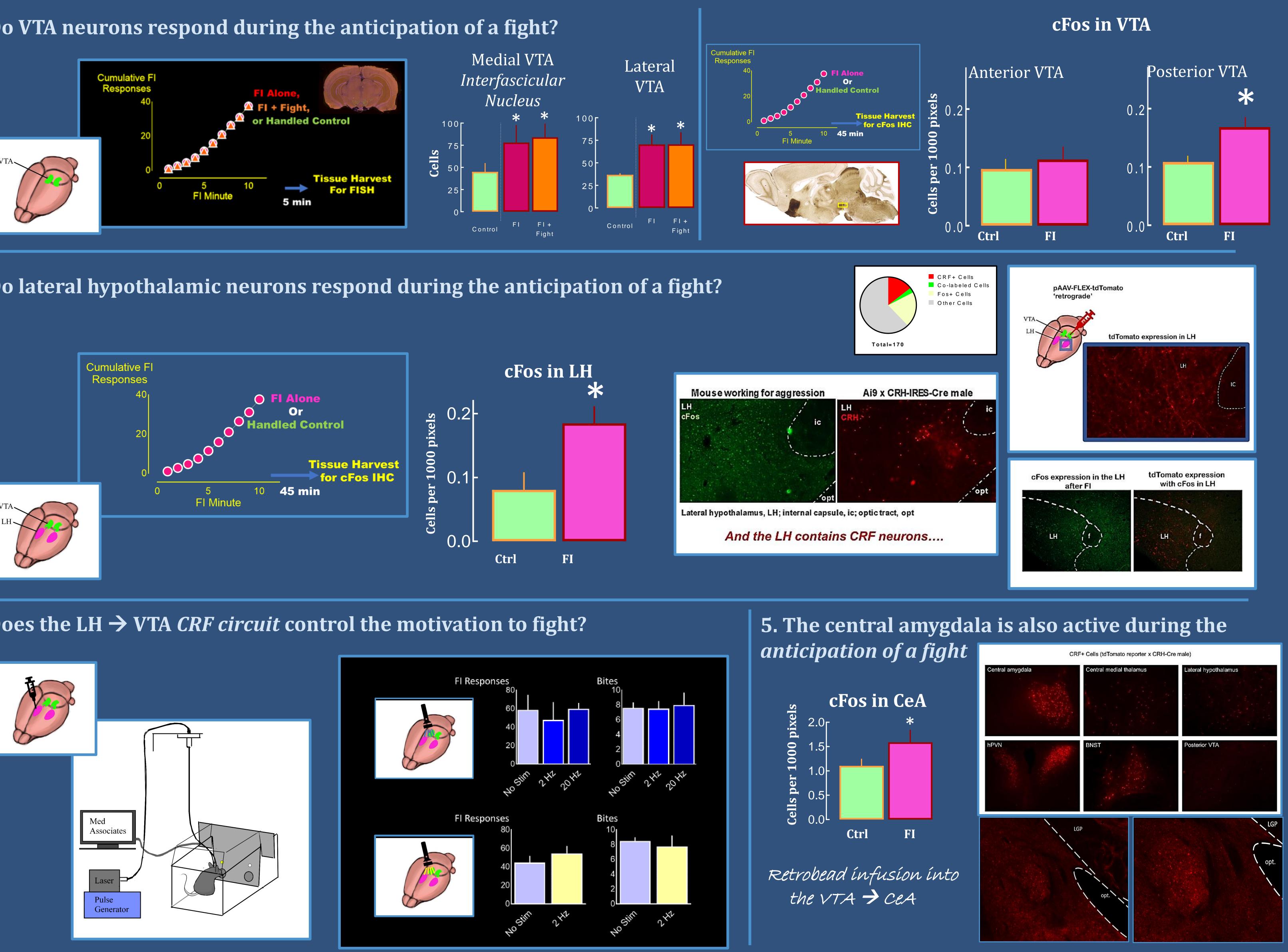
Modeling pathological aggressive motivation: VTA CRF circuits in aggression-specific arousal

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Results, cont.







Summary

>Patterns of FI responding for natural rewards are sensitive to CRF-R1 antagonism \geq Manipulation of CRF neurons along LH \rightarrow VTA pathway did not drive behavioral changes under FI responding. Potential explanations include attrition or that the $LH \rightarrow VTA$ pathway alone was not sufficient. \triangleright Next steps include manipulation of CeA and characterizing its projections, as the CeA was shown to contain many CRF neurons.

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