

Characterizing dopaminergic signaling in the nucleus accumbens core across different sign-tracking responses using fiber photometry

Sign-tracking press phenotypes and

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Introduction

- Sign-tracking (ST) ia physical manifestation of motivation in which animals attribute incentive salience—or motivational value—to a reward predictive cue (1).
- When this physical manifestation is too amplified, maladaptive behaviors like **addiction** arise (1).
- Phasic dopamine (DA) signaling in the nucleus accumbens core (NAc) is thought to encode incentive salience in ST (2), in addition to reward prediction errors when reward values change (3,4).
- It is unknown whether phasic DA may modulate the vigor and persistence of different sign-tracking responses observed across individual differences.

Methods

Subjects: 9 Long Evans rats, PN 70-90 male (n=4) and female (n=5)

Timeline:

 Sugery and Viral Expression
 Mag Training
 Sign-Tracking Acquisition Training (CS+ → pellet (noncontingent))
 Omission Testing CS+ → no press → pellet (CS+ → press → no pellet)

 21 days
 1 day
 12 days
 5 days

recording

Acq 12 ΔF/F DA peaks

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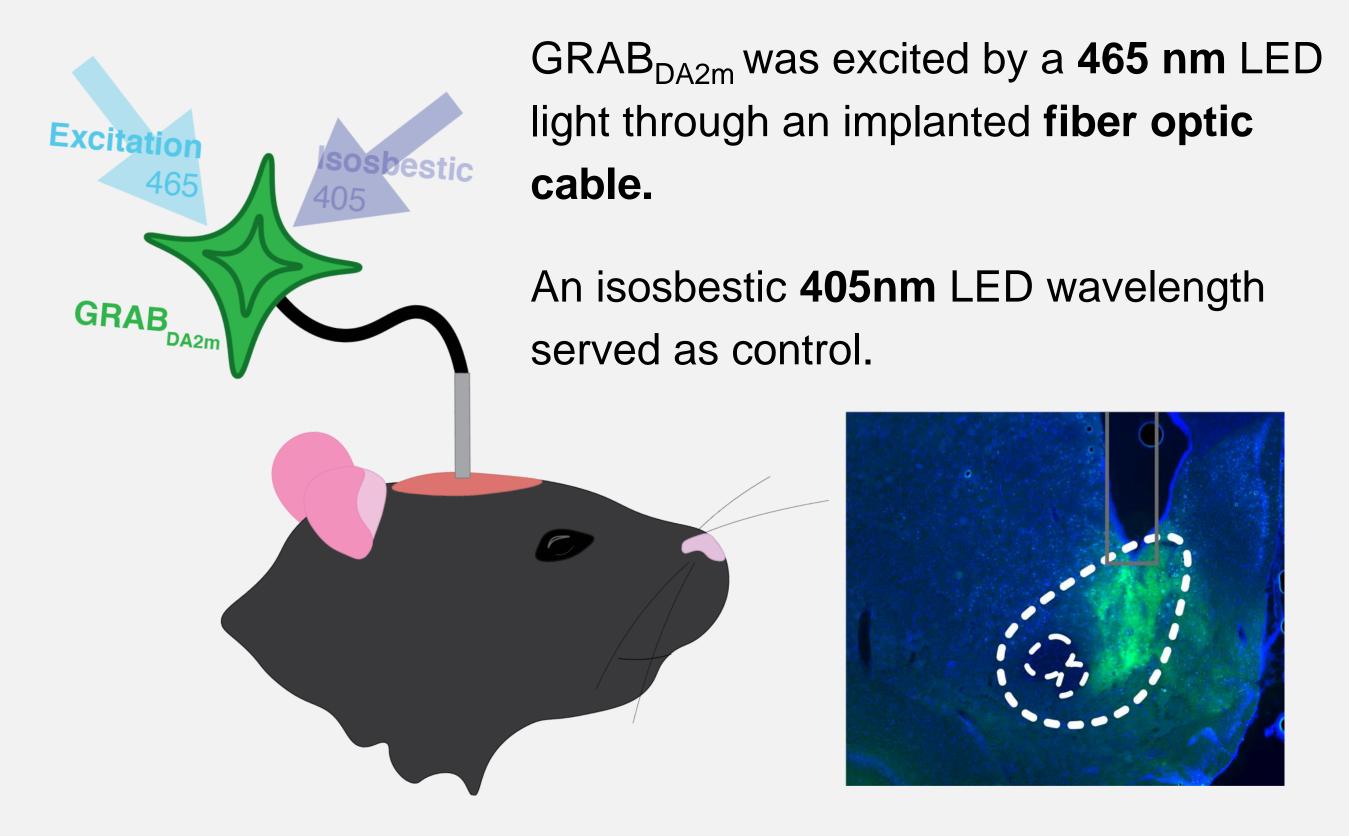
r=-0.14, p=0.86

Acq 12 ΔF/F DA peaks

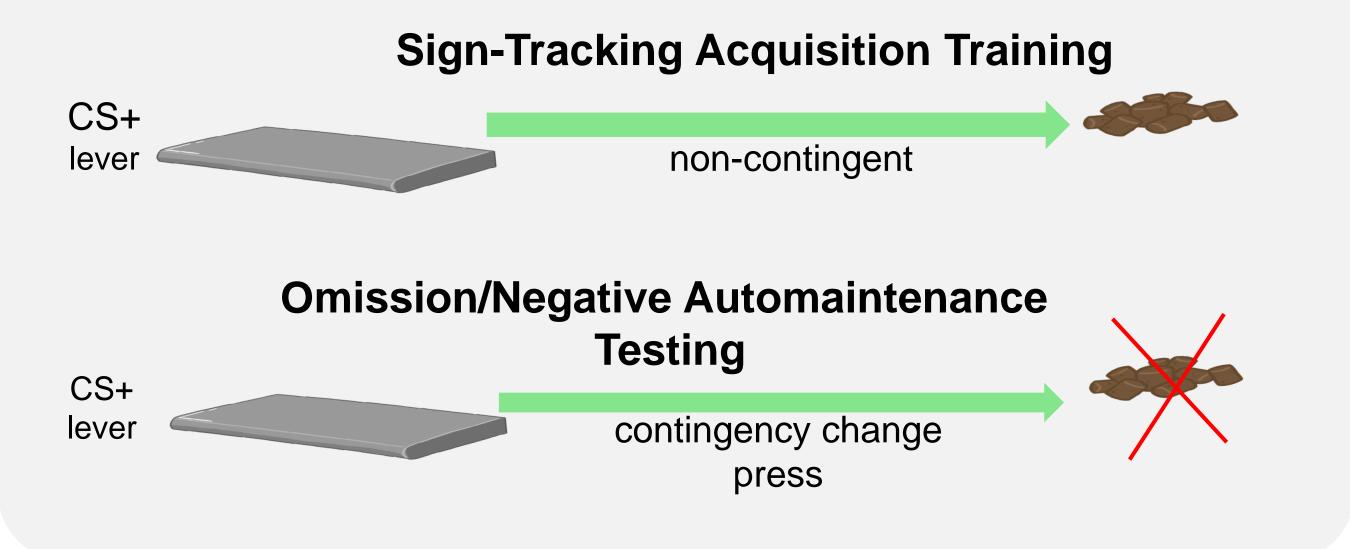
r=0.78, p=0.12

Acq 12 ΔF/F DA peaks

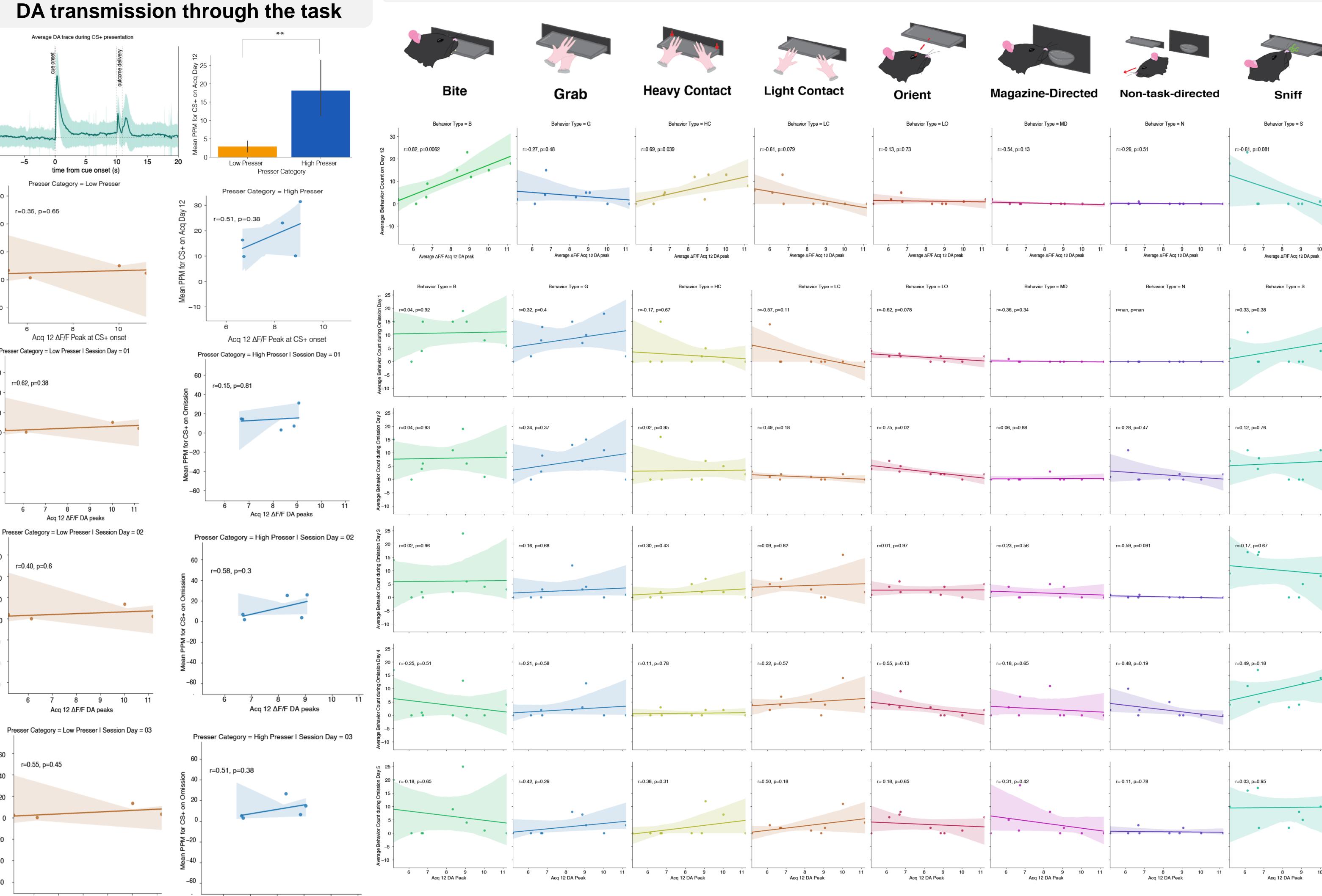
Fiber photometry: DA was recorded using GRAB_{DA2m} — a fluorescence dopamine viral sensor infused in the core of the NAc



Behavioral task:



Dynamics between behavioral microstructure and DA transmission through the task



Conclusions and Future directions

- Phasic DA in the Nac in ST animals may underlie dissociable incentive salience and prediction errors.
- The magnitude of Phasic DA signal may also explain the vigor and persistence of sign-tracking.
- ST animals with higher DA release at cue onset were more likely to engage in vigorous behavior during acquisition training and omission.
- Behavioral microstructure better characterizes the range of sign-tracking responses than pressing rates.

Future Directions:

- Using DA release during omission to predict behaviors during omitted or rewarded trials.
- Characterize cholinergic-dopaminergic dynamics in NAc across sign-tracking responses.

Funding: NSF IOS 1557987 (KSS); NIH 1R01DA044199 (KSS)

Acknowledgements: E.E Just Program Fellowship, UGAR Program, Presidential Scholar Program.

Special thanks to my advisors: Dr. Kyle Smith, Dr. Katherine Nautiyal and Erica Townsend

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