

Introduction

- Impaired reward sensitivity has been shown to be a promising marker in major depressive disorder in that baseline and early treatment related changes in reward sensitivity can predict ultimate treatment response¹.
- Different ERPs components contribute to the anticipatory (cue-P3 and SPN) and consummatory (feedback-P3) stages of reward processing, which also correspond to different forms of anhedonia in clinical population ².
- Limited studies have used high temporal resolution EEG/ERP to study anticipatory and consummatory in the same study design.

Objective: Validate current task performance aiming to be used in future clinical studies Aim 1: Examine whether ERPs amplitudes change as a function of reward magnitudes and valence at anticipatory and consummatory stages.

Aim 2: Exploratory examination of the association of severities in depressive symptoms and reward sensitivity.

Methods		
	Participant	Data Analys
Ν	33	EEG data were pre-process
Age, M (SD)	22.6 (6.0)	epoched to extract different components. Peak-to-peak were extracted and company one-way analysis of varian
% Female	21 (63.6%)	
BDI Score	10.4375 (9.0)	
GAD Score	4.75 (4.6)	
33 participants were recruited through the HSP system and received course for participating in the study.		and ERP amplitudes were a Linear regression analysis.



Developing An EEG Assay of Anticipatory and Consummatory Reward Sensitivity in Anhedonia

Yuexiaoxi (Cecilia) Yu^{*1}, Rebecca M. Todd¹ & Trisha Chakrabarty¹ ¹University of British Columbia

*Contact: cecilia.yu@ubc.ca



Analysis

-processed and different ERP -to-peak amplitudes

d compared with a

of variance.

ores and task ratings es were analyzed with





Higher reward rating > Low reward rating (p < .001) Anticipatory rating < Consummatory rating (p <.001)

BDI, GAD, HCL scores did not show significant correlation in anticipatory or consummatory ratings

- ERP amplitudes were not different in win and not win condition.
- In general population, reward sensitivity did not show significant correlation with selfreported BDI, GAD and HCL scores. Limitations
- Further research with a larger is needed.
- Possible confounders caused by visual stimuli are observed in earlier visual ERPs (N170).
- **Further direction**
- Analysis in other channel locations need to be conducted.
- Adjusted paradigm without the confounding stimuli were implanted and testing. • Analysis on other channel locations and exploratory analysis of self-reported scores and ERP amplitudes need to be conducted.

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