## Attentional Modulation of the Nociceptive Flexion Reflex and Anxiety Sensitivity



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**NFR Facilitators** 

N

96

41/55

47/24/25

**NFR Window** 

M

30.32

SD

13.90

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Variable

Age (years)

Male/Female

NHW/NA/Other

### Introduction

Pain and the nociceptive flexion reflex (NFR) are modulated by attention. It is unknown whether psychological traits are related to attentional modulation of pain/NFR. Healthy, pain-free participants were enrolled in a study (OK-SNAP) that assessed variables associated with pain processing. Participants completed the Anxiety Sensitivity Index-Revised (ASI-R) and viewed affective pictures, during which NFR magnitudes and pain ratings were gathered in response to electric stimulations to the sural nerve.

## Participants (N = 257)

- NHW = non-HispanicWhite
- NA = Native American

#### • Exclusion criteria:

- < 18 years of age</li>BMI > 35
- Current acute illness, psychotic symptoms,

chronic pain condition, or inability to speak/read English, cardiovascular, neurological, and/or circulatory problems, and recent use of analgesic, antidepressant, anxiolytic, antihypertensive medications

**NFR** Inhibitors

27.90

N

80

41/39

43/18/16

 Participant characteristics are reported by NFR modulation groups (pain modulation group compositions differ) - refer to Methods section for an explanation of group creation

# Participant Characteristics

SD

12.47

NFR No Modulation

M

27.77

**Stimulation** 

83

40/43

40/22/21

SD

11.45

ar, neurological, and/or circulatory problems, and edications odulation group compositions differ) - refer to		
Siceps Femoris EMG	0.8- 0.7- 0.6- 0.5- 0.3- 0.2- 0.1-	

## Methods

#### Nociceptive Flexion Reflex (NFR)

- $\bullet$  The NFR is a spinally-mediated, pain-related reflex elicited by A $\delta$  fiber activation
- The size of the NFR is correlated with pain ratings, but is an independent marker of spinal nociception
- NFR magnitude is determined by measuring biceps femoris activity in the 90-150 ms post-stimulus window
- NFR magnitude = mean rectified EMG of 90 to 150 ms post-stimulation interval minus mean of rectified EMG from -60 to 0 ms prestimulation interval divided by the average standard deviation of the rectified EMG from the two intervals

#### **Group Creation**

- 3 groups were created based on modulation during distraction (no modulation of pain/NFR, inhibited pain/NFR, facilitated pain/NFR)
- NFR Inhibitors = NFR magnitude < -0.1, NFR No Modulation = NFR magnitude -0.1 to 0.1, NFR Facilitation = NFR magnitude > 0.1
- Pain Inhibitors = Pain ratings < -2, Pain No Modulation = Pain ratings -2 to 2, Pain Facilitators = Pain ratings > 2

#### **Anxiety Sensitivity Index—Revised (ASI-R)**

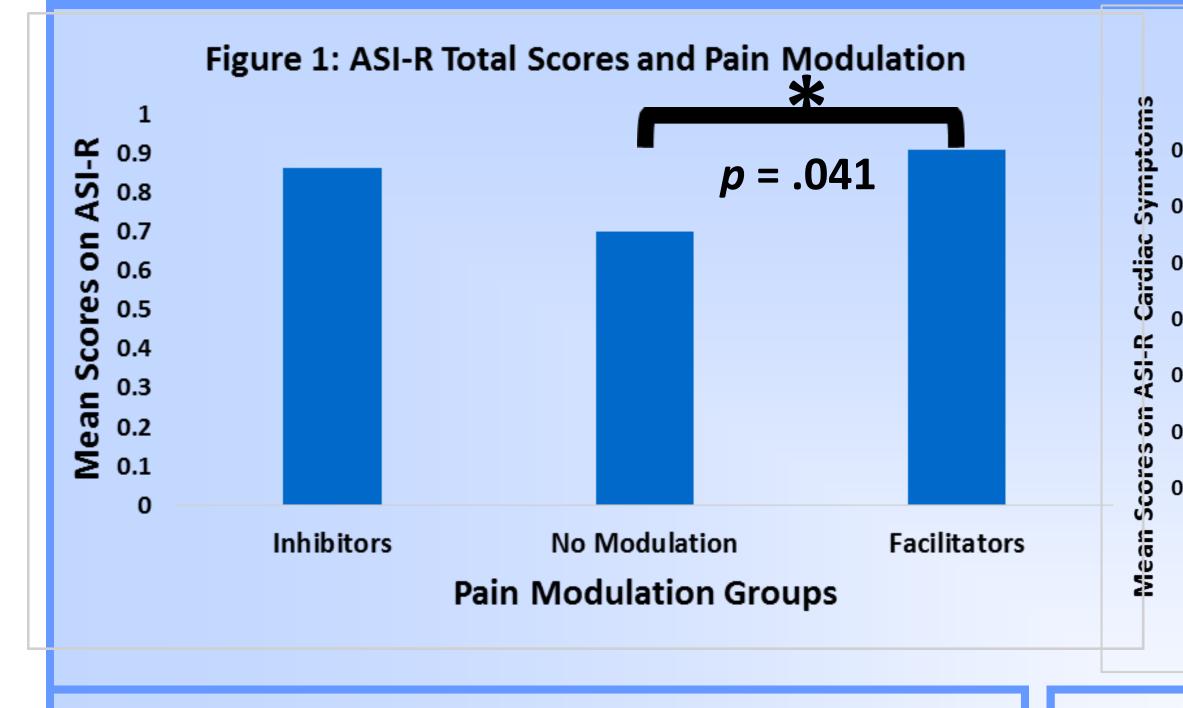
- The ASI-R is a measure of fear of anxiety symptoms with four subscales: somatic symptom interpretation, fear of cognitive dyscontrol, fear of publicly displaying anxiety, and fear of cardiac symptoms of anxiety
- Participants completed the ASI-R during a battery of questionnaires gathered before physiological data recording

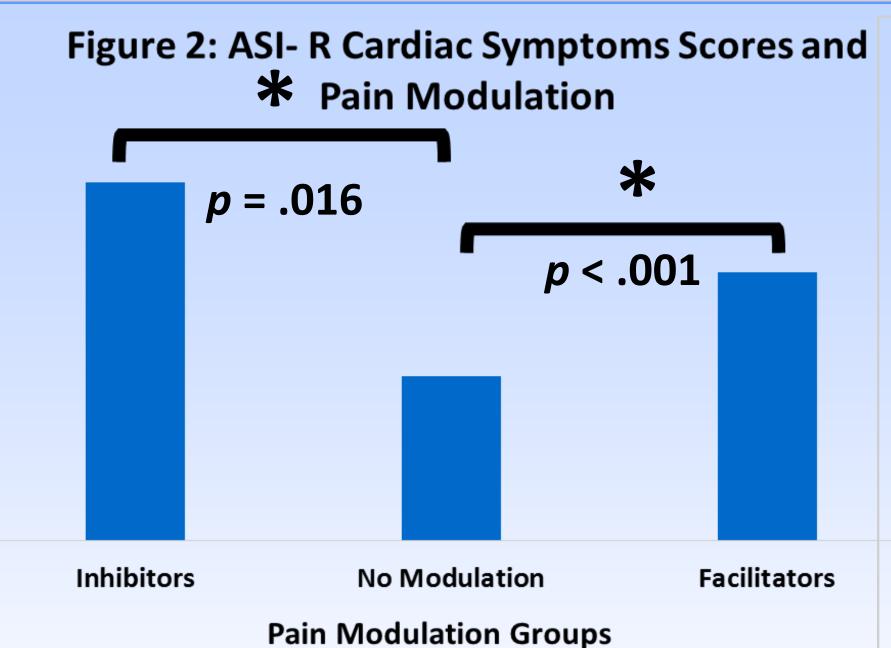
## Emotional Controls of Nociception (ECON) Paradigm

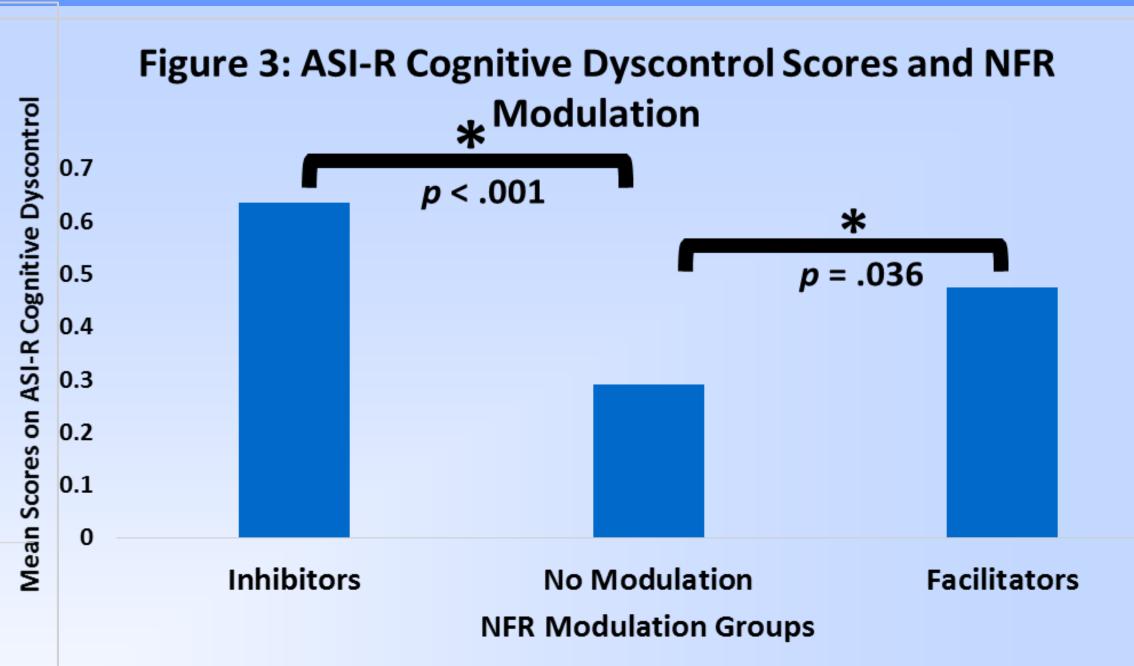
- 8 pleasant, 8 neutral, and 8 unpleasant pictures were presented in random order
- Each picture was shown for 6 seconds with a 12-22 second inter-picture interval
- Participants received painful stimulations to the ankle during 50% of the pictures (balanced across picture contents) and during 6 inter-picture intervals
- After each stimulation, participants rated their experienced pain intensity on a visual analog scale (VAS, 0-100)

#### **Attentional Modulation**

• For this study, attentional modulation was defined as the difference between pain/NFR evoked without pictures (no distractor) to those evoked during neutral pictures (distractor)







F(2, 235) = 3.49, MSE = 0.227, p = .032

F(2, 235) = 7.82, MSE = 0.186, p = .001

F(2, 232) = 10.32, MSE = 0.213, p < .001

## **Data Analysis**

- Outliers on the ASI-R were identified through Wilcox's MAD-median and replaced with the nearest non-outlier neighbor value
- One-Way Analysis of Variance
  - <u>Dependent Variables:</u> ASI-R Total, Somatic
    Symptom Interpretation, Fear of Cognitive
    Dyscontrol, Fear of Publicly Displaying Anxiety,
    Fear of Cardiac Symptoms of Anxiety
  - Independent Variables: NFR/Pain modulation groups

## Results

- In Figure 1, the "no pain modulation" group had lower anxiety sensitivity than the "facilitator" group
- In Figure 2, the "no pain modulation" group had less fear of cardiac symptoms of anxiety than the "inhibitor" or "facilitator" groups
- In Figure 3, the "no NFR modulation" group had less fear of cognitive dyscontrol related to anxiety than the "inhibitor" or "facilitator" groups

# Conclusions

- The pain facilitation group exhibited greater anxiety sensitivity and fear of cardiac symptoms than the no pain modulation group
- The no NFR modulation group exhibited greater fear of cognitive dyscontrol than the NFR facilitation and inhibition groups
- Individuals with a fear of cardiac symptoms may facilitate pain during a neutral distractor, whereas individuals with a lower fear of cognitive dyscontrol may not modulate NFR during a neutral distractor
- Future studies are needed to examine the mechanisms for this disparity

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