

Photobiomodulation for Management of Temporomandibular Disorder Pain

NCT05916235

Status	RECRUITING
Phase	Not Applicable
Sponsor	University of Florida
Enrollment	130 participants

Key Eligibility Criteria

Inclusion (6)

- Male or female, aged 18 years and older
- Meets the CATI pre-screening criteria during pre-screening visit \[participant-reported facial pain for at least 3 months and an average pain intensity rating for the week preceding CATI of ≥ 30 on a numerical rating scale (NRS, 0-100)\].
- Willing to provide signed and dated informed consent
- Willing to to comply with all study procedures and to be available for the duration of the study
- Meets diagnostic criteria for TMD (Masticatory Muscle disorder, 1 A: Myalgia) during Visit 0
- ... and 1 more (see full listing online)

Exclusion (14)

- Active rheumatologic disease
- Has a medical condition, laboratory finding, or physical exam finding (e.g., renal failure or dialysis, uncontrolled diabetes mellitus, or uncontrolled seizures) that precludes participation as determined by the investigator
- Initiated occlusal appliance therapy within 30 days prior to CATI
- Initiated non-pharmacologic therapy, such as acupuncture, biofeedback, and/or TENS within 30 days prior to CATI
- Is in active orthodontic treatment
- ... and 9 more (see full listing online)

Locations (1 total)

University of Florida, Gainesville, Florida, United States

<https://clinicaltrials.gov/study/NCT05916235>

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