

# Emsella Chair vs Sham for Male Sexual Dysfunction

NCT05370651

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<b>Status</b>	RECRUITING
<b>Phase</b>	Not Applicable
<b>Sponsor</b>	Corewell Health East
<b>Enrollment</b>	117 participants

## Key Eligibility Criteria

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### Inclusion (5)

- Able to read, understand, and provide written, dated, informed consent prior to screening, and be likely to comply with study protocol, including independently complete study questionnaires and communicate with study personnel about adverse events and other clinically important information.
- Men e 18 years of age.
- Sexually active within the past 12 weeks and plan to be sexually active during the next 12 weeks.
- \. Self-reported ejaculatory dysfunction symptoms present \>3 months 6. Self-reported failed conservative care of behavioral modifications and/or oral medications.
- \. Subject agrees not to start any new treatment for erectile dysfunction or ejaculatory dysfunction during the treatment and follow-up periods.

### Exclusion (17)

- Botox® use in bladder or pelvic floor muscles in the past year
- Subject weighs greater than 330 pounds, due to weight limits of the Emsella Chair.
- Pulmonary insufficiency, defined as difficulty breathing and fatigue, especially during exercise; chest pain, such as squeezing, pressure of tightness; the sensation of rapid or irregular heartbeat (palpitations); swelling of the legs or feet; dizziness or fainting; and/or bluish discoloration of the nails and/or lips (cyanosis)
- Any condition that causes a lack of normal skin sensation to the pelvis, thigh, or buttocks.
- Major metal implants such as: metal plates, screws, joint replacements, implanted cardiac pacemakers, drug pumps, neurostimulators, electronic implants, defibrillators, and metal implants in the pelvic area. Patients with other metal implants will be evaluated by the investigator for inclusion in the study.

... and 12 more (see full listing online)

## Locations (1 total)

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Corewell Health William Beaumont University Hospital, Royal Oak, Michigan, United States

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<https://clinicaltrials.gov/study/NCT05370651>

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