



Composur, A Patient-centric, Phase IV, Open-label, Prospective, Real World US Study to Evaluate Vibegron on Patient Treatment Satisfaction, Quality of Life, and Healthcare Resource Utilization in Patients with Overactive Bladder

Status: Recruiting

# Eligibility Criteria

Sex: Male or Female

**Age Group:** 18 years and over This study is NOT accepting healthy

volunteers

#### **Inclusion Criteria:**

- diagnosis of overactive bladder (OAB) with or without urgency urinary incontinence - symptoms of OAB for at least 3 months

### **Exclusion Criteria:**

- specific previous treatments for OAB (study staff will review) - neurologic conditions associated with OAB symptoms, e.g., multiple sclerosis - women who are pregnant or breast feeding or planning to become pregnant

## Conditions & Interventions

Interventions:

Drug: Vibegron

Conditions:

Kidney, Prostate & Urinary, Women's Health

Keywords:

Clinics and Surgery Center (CSC), OAB, Overactive Bladder

### More Information

Description: This study will evaluate treatment satisfaction, discontinuation, reasons for discontinuation, quality of life, healthcare resource utilization, and safety with vibegron for the treatment of OAB in the context of real-world clinical practice.

Study Contact: Maressa Twedt - twedt050@umn.edu

Principal Investigator: Nissrine Nakib

IRB

Number: SITE00001699

Thank you for choosing StudyFinder. Please visit http://studyfinder.umn.edu to find a Study which is right for you and contact sfinder@umn.edu if you have questions or need assistance