

Smartphones for Opiate Addiction Recovery

NCT05033028

Status	RECRUITING
Phase	Phase 3
Sponsor	NYU Langone Health
Enrollment	225 participants

Key Eligibility Criteria

Inclusion (17)

- Patients:
- Meet DSM-5 criteria for opioid-use disorder (heroin and/or prescription opioids);
- Have entered, or expressed a clear intention to enter, an OUD treatment program that prescribes either methadone or buprenorphine
- Any gender;
- years of age and older;
- ... and 12 more (see full listing online)

Exclusion (15)

- Patients:
- Serious medical, psychiatric or non-opioid substance use disorder that, in the opinion of a study physician, would make study participation hazardous to the participant or compromise study findings or would prevent the participant from completing the study. Examples include: (a) Disabling or terminal medical illness (e.g., uncompensated heart failure, cirrhosis or end-stage liver disease) as assessed by medical history, review of systems, physical exam and/or laboratory assessments;
- Severe, untreated or inadequately treated mental disorder (e.g., active psychosis, uncontrolled manic-depressive illness) as assessed by history and/or clinical interview;
- Current severe alcohol, benzodiazepine, or other depressant or sedative hypnotic use likely to require a complicated medical detoxification (routine alcohol and sedative detoxifications may be included);
- Suicidal or homicidal ideation that requires immediate attention;
- ... and 10 more (see full listing online)

Locations (2 total)

Rutgers University, Piscataway, New Jersey, United States
NYU Langone, New York, New York, United States

<https://clinicaltrials.gov/study/NCT05033028>

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