Who is eligible?

Inclusion Criteria

- Age 30 to 60 years
- Systolic BP ≥ 130 mm Hg based on two screenings
- Insomnia, either currently diagnosed or confirmed at screening

Exclusion Criteria

- Uncontrolled hypertension $(BP \ge 160/100 \text{ mm Hg})$
- Antihypertensive medications
- Cardiovascular medications
- Obstructive sleep apnea
- $BMI > 40 \text{ kg/m}^2$
- Pacemaker or atrial fibrillation
- Current pregnancy
- Alcohol or drug abuse in past year







Research studies are voluntary.

We encourage you to talk with your healthcare provider, family, and friends before you decide to take part in this research study. All participants will receive Cognitive Behavioral Therapy for Insomnia (CBT-I), which has been shown to improve sleep quality and promote consolidated sleep.

Over 9 months, participants will have assessments of sleep quality, blood pressure (BP), blood vessel elasticity, and lab work before and after CBT-I.

What does participation involve?

Screening assessments: 2 visits

• BPs and sleep surveys

Pre- CBT-I assessments: 2 visits*

- Questionnaires
- Sleep quality and BP monitors
- Ultrasound imaging
- Blood and urine collection

CBT-I: 6 sessions

• Weekly one-hour sessions

Post- CBT-I assessments: 2 visits*

- Questionnaires
- Sleep quality and BP monitors
- Ultrasound imaging
- Blood and urine collection

What are the benefits?

- CBT-I to improve sleep quality
- Assessments of sleep quality
- 24-hour blood pressure monitoring

