**Title**: Informed Consent Document for a Blood Pressure Control Study

**Introduction**  
You have been invited to participate in this research study that investigates a new medication aimed at helping people with high blood pressure. Whether you join is entirely your decision.

**Study Purpose**  
High blood pressure increases the risk of heart disease and stroke. The goal of this study is to measure how well the experimental drug, HL-105, can lower blood pressure in adults over 18.

**Procedures**

1. Screening: A blood pressure check and blood tests will determine if you qualify.
2. Treatment: You will be randomly assigned to receive HL-105 or a placebo once a day.
3. Clinic Visits: You must attend clinic sessions every six weeks for blood pressure readings and to answer health questions.
4. Duration: The study lasts 28 weeks, including a final follow-up assessment.

**Potential Risks and Benefits**

* Typical side effects may include fatigue or mild dizziness.
* There is a small chance of more serious events, such as heart rhythm changes.
* You could experience better control of your blood pressure and help advance medical research.

**Confidentiality**  
Your identity will be protected. Information collected will only be shared with study team members and authorized regulatory bodies.

**Voluntary Participation**  
You can decide not to participate or withdraw from the study at any time. Refusal to participate will not impact your standard medical treatment.