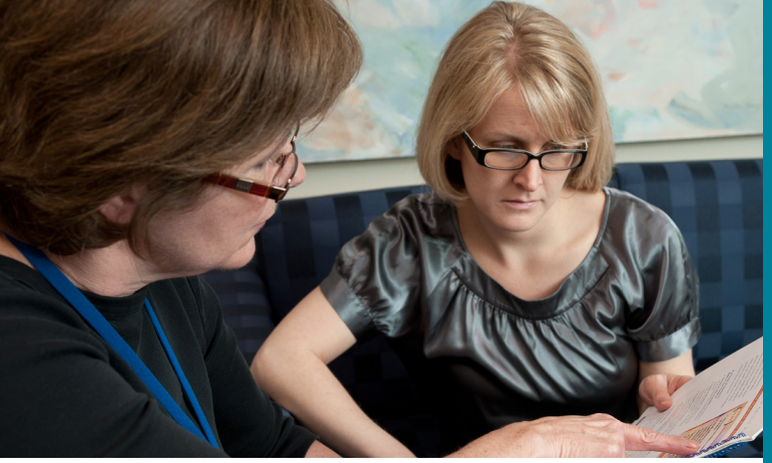


How can I learn more?

If you would like more information about the BP25543 Clinical Research Study, please contact the study doctor or staff. They will be able to answer any questions that you have and determine if you may be eligible to participate.

BP25543

Clinical Research Study

If you have Down Syndrome, you may wish to participate in a clinical research study.

What is a clinical research study?

- Clinical research studies are a way of developing new medicines.
- As a possible volunteer in a clinical trial, you may help researchers test the safety and other effects of study drugs.
- We do not know if volunteers will benefit from the study drug.
- There are possible risks if you choose to participate in this study.

What is the purpose of the BP25543 Study?

- We are studying a drug that is being developed to improve attention and memory in people with Down Syndrome.
- We want to know more about the study drug's safety, and about how your body handles the study drug.
- This study drug is a pill.
- If you participate, you will be instructed to take a certain number of pills twice a day.
- We are enrolling 33-44 volunteers with Down Syndrome, between the ages of 18 and 30 years for this study.
- The study will last for about 16 weeks.
- If you participate, you will receive the study drug and the medical care necessary for the study at no charge.

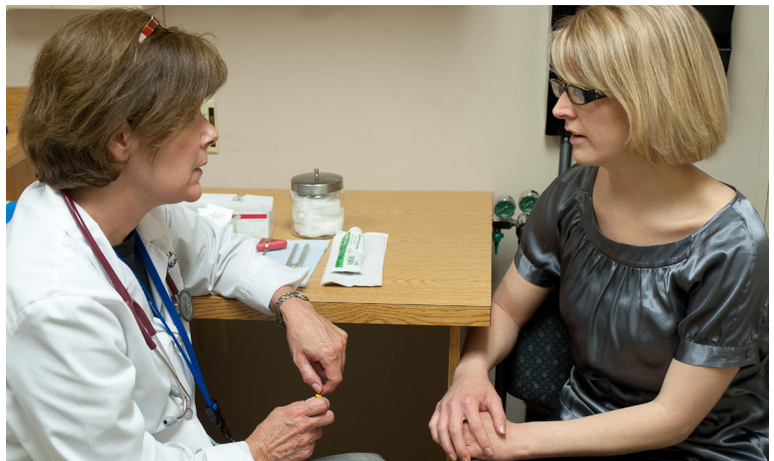
Who can participate?

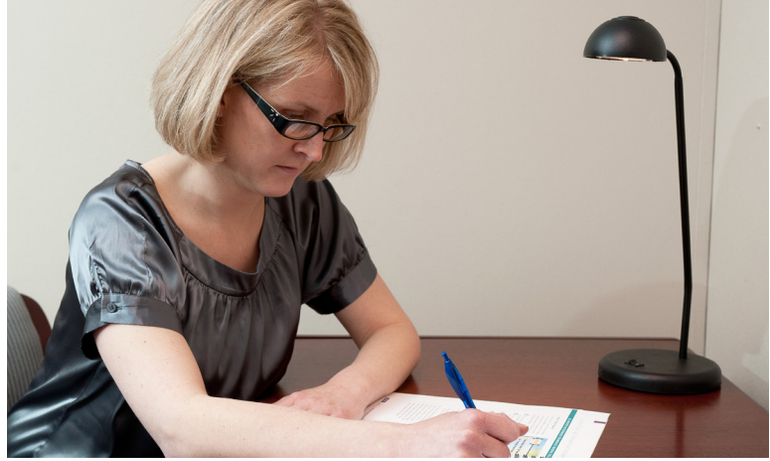
In order to be able to participate in this study:

- You must have Down Syndrome.
- To participate, you will need to first sign a consent/assent form.
- You must sign the consent/assent form and agree to certain restrictions. For example, certain medications are not allowed during the study.)
- You must use contraception (a form of birth control).

Participation is voluntary

- If you are eligible for the study, you will be informed of study details and any possible risks of participation.
- You can then decide if you would like to participate.
- You can change your mind about participating at any time during the study.





Responsibilities of study volunteers

If you choose to participate, you will be expected to:

- Take the study drug as instructed
- Complete a study drug diary
- Come to the clinic for study appointments
- Tell study staff about any side effects, other doctor visit, or hospital visits
- Tell study staff if you believe you might be pregnant
- Have medical tests and answer questions during study visits.