

**Interstitial
Cystitis/Bladder
Pain Syndrome
Study**



Living with Bladder Pain?

Join an interstitial
cystitis/bladder pain
syndrome study today.





About Interstitial Cystitis/Bladder Pain Syndrome (IC/BPS)

Interstitial cystitis/bladder pain syndrome (IC/BPS) is a chronic condition that can cause bladder pain and discomfort, along with urinary symptoms such as frequent voiding. This study is evaluating an investigational drug for the potential treatment of IC/BPS and its safety and effectiveness have not been established.

About Clinical Research Studies



The main objective of a clinical research study is to learn as much as possible about the safety and effectiveness of potential new drugs. These studies must be completed before a potential new drug can be made available to the public. Every day, research uncovers new information about medical conditions and possible therapies. Right now, there are more than 400,000 clinical studies in progress all over the world.¹

[1] <https://clinicaltrials.gov/ct2/resources/trends>



Study Qualifications

You may be able to participate in this Interstitial Cystitis/Bladder Pain Syndrome Study if you:

-  Are a female between 18 and 75 years old
-  Have a confirmed diagnosis of interstitial cystitis/bladder pain syndrome

If you are interested in participating, the study doctor and team will review additional study criteria with you.

Qualified participants will receive all study-related care at no cost and may be reimbursed for travel and expenses.



Study Details

This study is evaluating an investigational drug (a drug not approved for clinical use) for the potential treatment of IC/BPS in women.

In the first portion of the study, the investigational drug will be compared with a placebo (a substance that looks exactly like the investigational drug, except it does not contain any active ingredients). In the second portion of the study, should you decide to receive another treatment, you will receive the investigational drug.

Total study participation can last up to 46 weeks with a total of up to 13 visits.

Participants will also be asked to complete an electronic study diary to keep track of urination and bladder pain.



For more information:



[IC-BPSStudy.com](https://www.IC-BPSStudy.com)