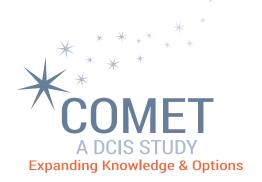
COMET Study (AFT-25): A Clinical Trial for Low-Risk Ductal Carcinoma in Situ (DCIS)



#### What is COMET?

The Comparison of Operative to Monitoring and Endocrine Therapy (COMET) Study is a clinical trial that looks at different treatment choices for ductal carcinoma in situ (also called DCIS).

DCIS is a non-invasive breast condition where cells that do not appear to be normal are found in the milk ducts. These cells are often harmless and may not need treatment.

### Why is the COMET Study being done?

The COMET Study will help researchers learn more about low-risk DCIS. The goal is to help many women avoid unnecessary treatments and their physical and/or emotional side effects. COMET Study (AFT-25): A Clinical Trial for Low-Risk Ductal Carcinoma in Situ (DCIS)



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## Why is the COMET Study being done?

The COMET Study will help researchers learn more about low-risk DCIS. The goal is to help many women avoid unnecessary treatments and their physical and/or emotional side effects. The study will compare two groups of women who are 40 years of age or older and have been diagnosed with low-risk DCIS. Participants will be randomly assigned to one of these two groups:

- Breast surgery, with the option of radiation and/or endocrine (hormone-blocking) therapy.
- 2. Active surveillance (sometimes called "close monitoring" or "watchful waiting"). This includes more follow-up exams and tests as well as the option of endocrine (hormone-blocking) therapy.

## Why would women join the COMET Study?

- → Many types of DCIS pose no threat to a woman's life.
- → Women may want to avoid or delay surgery and/or radiation while being closely watched by their medical team.
- → Close monitoring of low-risk DCIS may result in the same excellent outcomes as the standard treatments of surgery and radiation.

#### To learn more about this study or see if you may be eligible, talk to your doctor or go to: www.DClSoptions.org.

This work is supported through a Patient-Centered Outcomes Research Institute (PCORI) Award (PCS-1505-30497). The study will compare two groups of women who are 40 years of age or older and have been diagnosed with low-risk DCIS. Participants will be randomly assigned to one of these two groups:

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