Why join the Butterfly™ Medical BPH study?

Your involvement will help the medical field learn about a potential new BPH treatment–The Butterfly. This study will provide data for the FDA review and weigh, in advance of clearing Butterfly for further use in the United States.

Butterfly Medical

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CAUTION–investigational device: Limited by Federal (or United States) law to investigational use

MK-RA-USA-EN-PRB-22(001)

Butterfly Medical

INNOVATIVE INVESTIGATIONAL TECHNOLOGIC ADVANCEMENT FOR MEN WITH BPH!

What is The Butterfly™ and how does it work?

The Butterfly™ represents an innovate minimally invasive medical device technology. The Butterfly is a small, nitinol "Retractor implant" system designed relief BPH symptoms. The Butterfly procedure can be conducted in a Urologist's office or an Ambulatory Surgical Center (ASC)–BPH patients is no longer required to be treated in a hospital. The entire procedure takes approximately six-minutes!

The Butterfly™

Early clinical data with The Butterfly show that this treatment option helps reduce common LowerUrinary Tract (LUTS) symptoms. The data show improvement in IPSS indicating improvement in BPH symptoms, and in urinary flow (as measured in Qmax).

What should I expect, if treated by The Butterfly™ implant?

According to the available clinical data most men feel relief from prior urinary symptoms and resume common lifestyle activities within three-four weeks. The tiny, Butterfly implant will allow for continued sexual function and is placed without any burning or cutting of the tissue.

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BPH Study– am I eligible for The Butterfly study?

- **Men 50 – 80 years old**
- **Currently suffering from symptomatic BPH**
- **No previous BPH (device-based) treatment (other than a desire to eliminate BPH pills)**
- **Willingness to participate and return for appropriate follow-up visits**

To ask questions about the study or associated compensation, please contact: 512-410-3773 or email: UA-research@urologyaustin.com