WISPer

From Breath to Breakthrough

Are You Living with Idiopathic Pulmonary Fibrosis (IPF)?

The WISPer Trial is Enrolling Participants

What is the purpose of this study?

The purpose of this study is to evaluate the efficacy and safety of an investigational medicine in people living with IPF.

Am I eligible to participate in this study?

You may be eligible to participate if you are:

- Age 40 years or older
- Diagnosed with IPF within the past 7 years
- Have a Forced Vital Capacity (FVC) of 45% predicted or greater

You may be eligible for this trial while using approved treatments. Additional eligibility criteria will be assessed by the study doctor or staff during the screening process prior to being enrolled in the study. Not all individuals may qualify for this study.

What's involved with participating in the WISPer Trial?

The time commitment for the study includes 8 visits over the course of up to 32 weeks.

Does participating in this study cost anything?

Participants in the clinical trial may be reimbursed for study required travel and receive study required medical care at no cost. The investigational medication will be provided at no cost to you. This investigational medication has not been approved for use by the FDA outside of this study.

What can I expect as a participant in this study?

As a participant in the WISPer Trial, you will be randomly assigned to receive an infusion; this can be either the active investigational medication or placebo (the randomization ratio is 1:1). Placebo does not contain any active medication.

What if I change my mind?

You can withdraw from the study at any time, for any reason. Even if you begin the study, you can change your mind at any point. The study doctor will discuss your care options with you if you decide to withdraw from the study.

What if my condition worsens?

Each clinical trial has a well-documented plan, or protocol, about what you will need to know and what is expected. Your study team will keep you fully informed about the plan, including all known benefits and risks of the research. If there is a significant change in your health while you are participating, the study doctor will inform you and discuss your options, as your physical and mental health are of paramount importance. You are encouraged to ask any questions at any time. If you decide to participate, you will be asked to sign an informed consent form.

Where do I find more information?

To learn more about this clinical research study, please visit **wispertrial.com**

Mediar Therapeutics