



PACT – Patient New Clinical Trial and Research

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| Trial title | A 52-Week, Randomized, Double-Blind, Placebo-Controlled, Multi-center Study to Evaluate the Efficacy, Safety, and Tolerability of Axatilimab in Subjects with Idiopathic Pulmonary Fibrosis (IPF) |
| Trial synopsis | This is a study to evaluate the effectiveness and safety of axatilimab given for 52 weeks to patients with IPF. Patients will randomly (like flipping a coin) receive axatilimab or placebo (containing no medication) infusion (an injection into the vein) every two weeks. The effect of axatilimab on IPF will be measured through pulmonary function tests (PFTs), 6-minute walk tests and by assessing patients' IPF symptoms (including breathing and cough) using standard questionnaires. |
| Investigational medicinal product | Axatilimab |
| Disease target | Idiopathic Pulmonary Fibrosis (IPF) |
| Sponsor | Syndax Pharmaceuticals, Inc. |
| Duration | Patients will participate in the study for up to 70 weeks, including an 8 week Screening Period, a 52-week Treatment Period, and a 10-week Follow-up Period that includes a Follow-up telephone call 28 days post final dose and an ADA blood test Follow-up Visit 12 weeks post final dose. |
| Trial Status | Not yet recruiting |
| Lead site(s) in Australia | Lung Research Queensland |
| Lead site(s) in New Zealand | |
| Additional sites | |
| Contact | pactcoordinator@cre-pf.org.au |