

## A double blind, randomised, placebo-controlled trial evaluating the efficacy and safety of nerandomilast over at least 26 weeks in patients with Systemic Autoimmune Rheumatic Diseases associated Interstitial Lung Diseases (SARD-ILD)

**Status:** Recruiting

### Eligibility Criteria

**Sex:** Male or Female

**Age Group:** 18 years and over

This study is NOT accepting healthy volunteers

#### Inclusion Criteria:

- diagnosis of systemic autoimmune rheumatic diseases associated interstitial lung diseases (SARD-ILD) - lung function has not improved with immunosuppressant (IS) therapy - see link to [clinicaltrials.gov](http://clinicaltrials.gov) for complete Inclusion criteria

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#### Exclusion Criteria:

- active vasculitis - suicidal behavior in the past 2 years - see link to [clinicaltrials.gov](http://clinicaltrials.gov) for complete Exclusion criteria

### Conditions & Interventions

#### Interventions:

Drug: Nerandomilast, Drug: Placebo matching nerandomilast

#### Conditions:

Rare Diseases, Rare Diseases, Respiratory System

#### Keywords:

Clinics and Surgery Center (CSC), Autoimmune Rheumatic Diseases, Interstitial Lung Diseases

### More Information

**Description:** Nerandomilast is being developed to treat lung fibrosis. This study is to test a drug called nerandomilast in people with SARD-ILD who also take an immunosuppressant medicine. Participants are put into 2 groups randomly, which means by chance and will receive either nerandomilast or placebo tablets. Placebo tablets look like nerandomilast tablets but do not contain any study drug. Participants will be in the study for about 7.5 to 13 months depending on when they join the study.

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**Phase:** PHASE3

**IRB Number:** STUDY00024424

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