Patient - PACT New Clinical Trial and Research



Trial title	Genetic research in Idiopathic Pulmonary Fibrosis (GRIPF)
Trial synopsis	We are seeking suitable participants for a research study that examines how genetic factors may contribute to Idiopathic Pulmonary Fibrosis (IPF), a rare but severe lung disease. We are interested in involving families in this study as this will allow us to look at genetic similarities and differences in family members with and without disease. This national research project is being carried out by a team of researchers from the Menzies Institute for Medical
	Research, University of Tasmania in collaboration with the Australian IPF Registry. Both the Registry and this study are facilitated by Lung Foundation Australia.
	This study plans to learn more about pulmonary fibrosis and determine if genetic factors contribute to the development of diseases which result in scarring in the lung. What we learn may be helpful in the future to people diagnosed with these diseases, known as pulmonary fibrosis, and their family members.
Investigational medicinal product, comparator and randomisation	Not applicable
Disease target	IPF
Sponsor	Philanthropic support
Duration	5 years
Trial Status	Recruiting
Lead site(s) in Australia	Royal Prince Alfred Hospital, Camperdown, NSW
Lead site(s) in New Zealand	Not applicable
Additional sites	The Alfred Hospital, VIC The Prince Charles Hospital QLD Fiona Stanley Hospital WA Royal Adelaide Hospital, SA John Hunter Hospital NSW Calvary Surgical Suites TAS Launceston General Hospital TAS Clinical A/Prof Peter Kendal WA (private) Equinox Specialist Centre, ACT

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