

INNOVATIONACCESS AVAILABLE TECHNOLOGIES CONTACT US

Request Information

Permalink

Peptide Inhibitors of Idiopathic Pulmonary Fibrosis

Tech ID: 31787 / UC Case 2019-089-0

ABSTRACT

Researchers at the University of California, Davis have developed a peptide that targets fibrogenic pathways in order to treat idiopathic pulmonary fibrosis.

FULL DESCRIPTION

Fibrosis is the development of excess fibrous, connective tissue. The development of some fibrous tissue is a normal step in many tissue or organ repair processes. In the lungs, however, repeated injury and repair can lead to life-threatening diseases such as idiopathic pulmonary fibrosis (IPF). IPF has a poor prognosis, and current therapeutics are ineffective - as they tend to focus on the inflammation aspects of the disease (not the fibrosis stage). These therapeutic have limited success due to their non-specific suppression of the inflammatory response. They can also act as powerful immunosuppressants. So there is a strong need for improved therapeutics to treat IPF.

Researchers at the University of California, Davis have developed a novel peptides targeting fibrogenic pathways.

Phospho-MARCKS can act as a specific marker for activated fibroblasts. This marker can be targeted by a MARCKS PSD sequence (MPS) peptide to inhibit the marker's activity. The novel peptides will destroy activated fibroblasts/myofibroblasts without affecting dormant fibroblasts. It is also effective in inhibiting fibroblast cell movement, proliferation and differentiation into myofibroblasts without exhibiting any toxicity to normal cells.

APPLICATIONS

Therapeutic treatment for pulmonary fibrosis through fibrosis suppression

FEATURES/BENEFITS

- More effective than IPF therapeutics that focus on inflammation
- · Effective in inhibition of fibroblast cell migration, proliferation, and differentiation
- Does not affect dormant fibroblasts
- · Peptide is soluble, stable and easy to manipulate

PATENT STATUS

Country	Туре	Number	Dated	Case
United States Of America	Published Application	20220267390	08/25/2022	2019-089

CONTACT

Raj Gururajan rgururajan@ucdavis.edu tel: 530-754-7637.



INVENTORS

- ▶ Chen, Ching-Hsien
- ▶ Wu, Reen
- ► Yang, David

OTHER INFORMATION

KEYWORDS

Idiopathic Pulmonary
Fibrosis, Fibroblast, MPS
peptide, MARCKS

CATEGORIZED AS

- Medical
 - ▶ Disease:

Respiratory and

Pulmonary System

New Chemical

Entities, Drug Leads

▶ Therapeutics

RELATED CASES

2019-089-0

- ► Controlling Tumor Growth And Malignancy
- ▶ Suppression of Allergic Lung Inflammation and Hyperactivity
- ► Mucin-Specific Monoclonal Antibodies

University of California, Davis
InnovationAccess
1850 Research Park Drive, Suite 100, ,
Davis,CA 95618

Tel: 530.754.8649
innovationAccess@ucdavis.edu
research.ucdavis.edu/u/s/ia
Fax: 530.754.7620

@ 2019 - 2024, The Regents of the University of California $\frac{\text{Terms of use}}{\text{Privacy Notice}}$