

Akronym	Bezeichnung	Phase	Indikation	Beschreibung	Rekrutierung
<b>IPF STUDIEN</b>					
<b>ISABELA 1 &amp;2</b>	GLPG1690-CL-304	III	Idiopathische Lungenfibrose	A Phase 3, randomized, double-blind, parallel-group, placebo-controlled, multi-center study to evaluate the efficacy and safety of two doses of GLPG1690 in addition to local standard of care for minimum 52 weeks in subjects with idiopathic pulmonary fibrosis.	<b>Aktiv</b>
<b>Juniper</b>	ND-L02s-0201-005	II	Idiopathische Lungenfibrose	A Phase 2, Randomized, Double-Blind, Placebo-Controlled Study to Evaluate the Safety, Tolerability, Biological Activity, and PK of ND-L02-s0201 in Subjects with Idiopathic Pulmonary Fibrosis (IPF)	<b>Aktiv</b>
<b>VAY736 Novartis</b>	CVAY736X2207	II	Idiopathische Lungenfibrose	A subject-, investigator-, and sponsor-blinded, randomized, placebo-controlled, multicenter study to investigate efficacy, safety, and tolerability of VAY736 in patients with idiopathic pulmonary fibrosis	<b>Aktiv</b>
<b>ZEPHYRUS</b>	FibroGen FGCL-3019-091	III	Idiopathische Lungenfibrose	A Phase 3, Randomized, Double-Blind, Placebo-Controlled Efficacy and Safety Study of Pamrevlumab in Subjects with Idiopathic Pulmonary Fibrosis (IPF)	<b>In Vorbereitung</b>
<b>Celgene</b>	CC-90001-IPF-001	II	Idiopathische Lungenfibrose	A Phase 2, 24-Week, Randomized, Double-blind, Placebo-controlled, multicenter Study, with an 80 week active Treatment Extension, to evaluate the efficacy and safety of CC-90001 in Subjects with IPF.	<b>Aktiv</b>
<b>BI 1305-0013</b>	1305-0013	II	Idiopathische Lungenfibrose	A randomized, double-blind, placebo-controlled parallel group study in IPF patients over 12 weeks evaluating efficacy, safety and tolerability of BI 1015550 18 mg taken orally b.i.d.	<b>In Vorbereitung</b>

<b>Pliant</b>	PLN-74809-IPF-202	IIa	Idiopathische Lungenfibrose	A randomized, double-blind dose-ranging, placebo-controlled Phase 2a evaluation of the safety, tolerability and pharmacokinetics of PLN-74809 in participants with idiopathic pulmonary fibrosis (IPF).	<b>Aktiv</b>
<b>BMS</b>	IM027-040	II	Idiopathische Lungenfibrose	A Multicenter, randomized, double-blind, placebo-controlled, phase 2 study of the Efficacy and Safety and Tolerability of BMS-986278 in Participants with pulmonary Fibrosis.	<b>In Vorbereitung</b>
<b>Galactic</b>	Galactic-1	IIb	Idiopathische Lungenfibrose	A randomized, double-blind, multicentre, parallel, placebo-controlled Phase 2b study in subjects with idiopathic pulmonary fibrosis (IPF) investigating the efficacy and safety of TD139, an inhaled galectin-3 inhibitor administered via a dry powder inhaler over 52 weeks.	<b>Aktiv</b>
<b>Roche</b>	WA-42293	III	Idiopathische Lungenfibrose	A Phase III Randomized, Double –Blind, placebo-controlled trial to evaluate the efficacy and Safety of PRM-151 in patients with idiopathic pulmonary fibrosis.	<b>In Vorbereitung</b>
<b>Nicht-IPF STUDIEN</b>					
<b>INBUILD-ON</b>	1199.248	III	PF-ILD	An open-label extension trial of the long term safety of nintedanib in patients with Progressive Fibrosing Interstitial Lung Disease (PF-ILD)	<b>Rekrutierung beendet</b>

**Alveolar Proteinose STUDIEN**

<b>IMPALA</b>	IMPALA MOL PAP 202 Extension	III	Alveolar Proteinose	An open-label,non-controlled, multicentre trial of inhaled Molgramotism in autoimmune pulmonary alveolar proteinosis patients	<b>Rekrutierung beendet</b>
<b>IMPALA 2</b>	IMPALA MOL PAP	III	Alveolar Proteinose	An open-label,non-controlled, multicentre trial of inhaled Molgramotism in autoimmune pulmonary alveolar proteinosis patients	<b>In Vorbereitung</b>

**Sarkoidose STUDIEN**

<b>Novartis</b>	CCMK389X2201	II	Sarkoidose	A subject and investigator blinded, randomized, placebocontrolled, repeat-dose, multicenter study to investigate efficacy, safety, and tolerability of CMK389 in patients with chronic pulmonary sarcoidosis.	<b>Aktiv</b>
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