

2018 Primary biliary cholangitis Drug Development-Pipeline Analysis Report

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Abstracts

Primary biliary cholangitis (PBC) is a long term chronic and autoimmune disease in which the bile ducts are inflamed and gradually destroyed. This results in built of bile in the liver, resulting in damage to the liver. The rare disease has limited treatment options with FDA approving obeticholic acid plus ursodeoxycholic acid as a single therapy.

Over 25 companies and universities are focusing on developing treatment options for Primary biliary cholangitis.

To assist investigators and funding and regulatory organizations, VPA Research has come up with a comprehensive report on Primary biliary cholangitis pipeline. The report provides insights into different therapeutic candidates in preclinical, research, discovery, NDA/IND, pre registration, phase 1, phase 2, and phase 3 trials). Drugs under development directly and through combination with other drugs are also included.

Current status, developmental phase, participating companies and entities, recent developments, orphan drug/fast track/other designations, drug class are provided for each Primary biliary cholangitis pipeline product. Mechanism of Action and the target area of the pipeline product are also provided. Further, clinical and preclinical trials along with results of the trials are also included in the report.

In addition to complete details of each product, the report provides key trends in Primary biliary cholangitis pipeline studies. The products under development are categorized according to their development phase, mechanism and company to provide detailed insights into the type of drugs being developed and the stages of development.

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Arena Pharmaceuticals

CymaBay Therapeutics

Enanta Pharmaceuticals

FF Pharma

Genfit

Genkyotex

GlaxoSmithKline

Intercept Pharmaceuticals

KAN Research Institute

NGM Biopharmaceuticals

Novartis

Phenex Pharmaceuticals

Retrophin

Suzhou Zelgen Biopharmaceutical Co Ltd

Tiziana Life Sciences

Virobay Inc

Zydus Cadila

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Originator
Co-Developer/License Partner
Orphan Drug Designation
Development Phase
Mechanism of Action
Current Status
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