Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13) - Pipeline Review, H1 2016

Description:

Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13) - Pipeline Review, H1 2016

Summary

‘Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13) - Pipeline Review, H1 2016’, provides in depth analysis on Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13) targeted pipeline therapeutics.

The report provides comprehensive information on the Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13), targeted therapeutics, complete with analysis by indications, stage of development, mechanism of action (MoA), route of administration (RoA) and molecule type. The report also covers the descriptive pharmacological action of the therapeutics, its complete research and development history and latest news and press releases. Additionally, the report provides an overview of key players involved in Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13) targeted therapeutics development and features dormant and discontinued projects.

Our report features investigational drugs from across globe covering over 20 therapy areas and nearly 3,000 indications. The report is built using data and information sourced from proprietary databases, company/university websites, clinical trial registries, conferences, SEC filings, investor presentations and featured press releases from company/university sites and industry-specific third party sources. Drug profiles featured in the report undergoes periodic review following a stringent set of processes to ensure that all the profiles are updated with the latest set of information. Additionally, various dynamic tracking processes ensure that the most recent developments are captured on a real time basis.

The report helps in identifying and tracking emerging players in the market and their portfolios, enhances decision making capabilities and helps to create effective counter strategies to gain competitive advantage.

Note*: Certain sections in the report may be removed or altered based on the availability and relevance of data.

Scope

- The report provides a snapshot of the global therapeutic landscape for Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13)
- The report reviews Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13) targeted therapeutics under development by companies and universities/research institutes based on information derived from company and industry-specific sources
- The report covers pipeline products based on various stages of development ranging from pre-registration till discovery and undisclosed stages
- The report features descriptive drug profiles for the pipeline products which includes, product description, descriptive MoA, R&D brief, licensing and collaboration details & other developmental activities
- The report reviews key players involved in Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13) targeted therapeutics and enlists all their major and minor projects
- The report assesses Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13) targeted therapeutics based on mechanism of action (MoA), route of administration (RoA) and molecule type
- The report summarizes all the dormant and discontinued pipeline projects
- The report reviews latest news and deals related to Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13) targeted therapeutics

Reasons to buy

- Gain strategically significant competitor information, analysis, and insights to formulate effective R&D strategies
- Identify emerging players with potentially strong product portfolio and create effective counter-strategies
to gain competitive advantage
- Identify and understand the targeted therapy areas and indications for Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13)
- Identify the use of drugs for target identification and drug repurposing
- Identify potential new clients or partners in the target demographic
- Develop strategic initiatives by understanding the focus areas of leading companies
- Plan mergers and acquisitions effectively by identifying key players and its most promising pipeline therapeutics
- Devise corrective measures for pipeline projects by understanding Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13) development landscape
- Develop and design in-licensing and out-licensing strategies by identifying prospective partners with the most attractive projects to enhance and expand business potential and scope

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Aprea AB
Cellceutix Corporation
Critical Outcome Technologies Inc.
Eleos Inc.
ORCA Therapeutics B.V.
OSE Pharma SA
PCI Biotech Holding ASA
Quark Pharmaceuticals, Inc.
Stemline Therapeutics, Inc.
Shenzen SiBiono GeneTech Co., Ltd.
SK Biopharmaceuticals Co., Ltd.
Tara Immuno-Oncology Therapeutics LLC
Z53 Therapeutics, LLC

Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13) - Drug Profiles

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Cellular Tumor Antigen P53 (Tumor Suppressor P53 or Antigen NY-CO-13) - Featured News & Press Releases

Apr 19, 2016: Critical Outcome Commences Treatment of Second Cohort in Phase 1 Study of COTI-2 in Gynecologic Cancers

Mar 31, 2016: Cellceutix Successfully Completes In Vitro Study in Support of Planned Phase 2 Trial of Kevetrin for Ovarian Cancer

Mar 14, 2016: Quark Pharmaceuticals Doses First Patient In Two Pivotal Clinical Studies of RNAi-Based Therapeutic QPI-1002

Feb 22, 2016: Assay Results From Cellceutix Phase 1 Clinical Trial of Kevetrin for Cancer Show Increased p21 Expression in 67.5% of Evaluable Patients

Feb 22, 2016: First patients enrolled and dosed in the pivotal trial of Phase 3 of the immunotherapy Tedopi for advanced non-small cell lung cancer

Feb 16, 2016: Critical Outcome Treats First Patient in Phase 1 Study of COTI-2 in Gynecologic Cancers

Feb 16, 2016: Cellceutix Completes Clinical Trial of Kevetrin for Advanced Solid Tumors

Feb 10, 2016: Cellceutix Meets With FDA for Phase 2 Clinical Trial of Kevetrin for Ovarian Cancer

Feb 04, 2016: OSE Pharma Announces U.S. Initiation of Atalante 1, the Company’s Global, Pivotal Phase 3 Trial for Tedopi Immunotherapy in Non-Small Cell Lung Cancer

Jan 22, 2016: Cellceutix Releases Data Used in Receiving FDA Orphan Drug Designation of Kevetrin for Pancreatic Cancer

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Disclaimer

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