

Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix Phosphoprotein or Tegument Protein UL83 or UL83) - Pipeline Review, H1 2018

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Abstracts

Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix Phosphoprotein or Tegument Protein UL83 or UL83) - Pipeline Review, H1 2018

SUMMARY

Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix Phosphoprotein or Tegument Protein UL83 or UL83) pipeline Target constitutes close to 9 molecules. Out of which approximately 8 molecules are developed by companies and remaining by the universities/institutes.

The latest report Human cytomegalovirus 65 kDa Phosphoprotein - Pipeline Review, H1 2018, outlays comprehensive information on the Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix Phosphoprotein or Tegument Protein UL83 or UL83) targeted therapeutics, complete with analysis by indications, stage of development, mechanism of action (MoA), route of administration (RoA) and molecule type.

Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix Phosphoprotein or Tegument Protein UL83 or UL83) - 65-kDa cytosolic phosphoprotein (pp65) counteracts the host antiviral immune response when activated and phosphorylated and by preventing IRF3 from entering the nucleus.

It participates in the transactivation of viral major immediate-early genes by the recruitment of host IFI16 to the promoter of these genes. The molecules developed by companies in Phase I, Phase I, IND/CTA Filed and Preclinical stages are 4, 1, 1 and 2



respectively. Similarly, the universities portfolio in Phase II stages comprises 1 molecules, respectively.

Report covers products from therapy areas Infectious Disease, Oncology and Immunology which include indications Cytomegalovirus (HHV-5) Infections, Glioblastoma Multiforme (GBM), Brain Tumor, Hematological Tumor, Human Immunodeficiency Virus (HIV) Infections (AIDS), Kidney Transplant Rejection, Liver Transplant Rejection, Medulloblastoma, Recurrent Glioblastoma Multiforme (GBM) and Transplant Rejection.

Furthermore, this report also reviews key players involved in Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix Phosphoprotein or Tegument Protein UL83 or UL83) targeted therapeutics development with respective active and dormant or discontinued projects. Driven by 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.

Note: Certain content/sections in the pipeline guide 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 Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix Phosphoprotein or Tegument Protein UL83 or UL83)

The report reviews Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix Phosphoprotein or Tegument Protein UL83 or UL83) 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 Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix Phosphoprotein or Tegument Protein UL83 or UL83) targeted therapeutics and enlists all their major and minor projects

The report assesses Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix Phosphoprotein or Tegument Protein UL83 or UL83) 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 Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix Phosphoprotein or Tegument Protein UL83 or UL83) 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 Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix Phosphoprotein or Tegument Protein UL83 or UL83)

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 it's most promising pipeline therapeutics



Devise corrective measures for pipeline projects by understanding Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix Phosphoprotein or Tegument Protein UL83 or UL83) 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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Hookipa Biotech AG

Immunomic Therapeutics Inc

Vakzine Projekt Management GmbH

Vaximm AG

VBI Vaccines Inc

Vical Inc

Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix

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Cellular Immunotherapy to Target PP65 for Cytomegalovirus Infections - Drug Profile

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Mechanism Of Action

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Triplex - Drug Profile

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Mechanism Of Action

R&D Progress

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Product Description

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R&D Progress

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Human cytomegalovirus 65 kDa Phosphoprotein (PP65 or 65 kDa Matrix

Phosphoprotein or Tegument Protein UL83 or UL83) - Product Development Milestones Featured News & Press Releases

Jan 17, 2018: VBI Vaccines Announces Dosing of First GBM Patient in Phase 1/2a Clinical Study of VBI-1901

Nov 22, 2017: Dr. Duane Mitchell Reports Findings of Study Aimed at Aggressive Brain Cancer

Nov 10, 2017: VBI Vaccines to Present New Preclinical Data for its GBM Immunotherapy, VBI-1901, at SITC 2017

Oct 11, 2017: Immunomic Therapeutics Offers Travel Fund for Clinical Trial Patients

Aug 28, 2017: VBI Vaccines to Present New Preclinical Data for its GBM

Immunotherapy, VBI-1901, at the Immuno-Oncology Summit



Aug 15, 2017: VBI Vaccines Announces FDA Acceptance of Investigational New Drug Application for VBI-1901 to Treat Glioblastoma Multiforme

May 04, 2017: Hookipa Biotech Presents Positive Data from Phase 1 First-In-Human Trial of Vaccine Against Cytomegalovirus

Apr 05, 2017: VBI Vaccines to Present Update on VBI-1901 at the World Vaccine Congress

Jan 09, 2017: Hookipa Biotech Announces Publication in Clinical and Vaccine Immunology Highlighting Vaxwave as an Effective Viral Vector for Vaccination against Congenital Cytomegalovirus Infections

Nov 10, 2016: VBI Vaccines to Present at the Society of Neuro-Oncology Annual Meeting

Oct 28, 2016: Data from Phase 1 Trial of Triplex Vaccine for Control of Cytomegalovirus Published in Blood

Oct 11, 2016: VBI Vaccines Completes Pre-IND Meeting for its Glioblastoma Immunotherapy Candidate

Oct 05, 2016: VBI Vaccines to Present at the World Vaccine Congress Europe

Aug 29, 2016: VBI Vaccines to Present at the Immuno-Oncology Summit

Jul 20, 2016: Hookipa Biotech Initiates First-in-Human Study of Cytomegalovirus

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COMPANIES MENTIONED

Hookipa Biotech AG
Immunomic Therapeutics Inc
Vakzine Projekt Management GmbH
Vaximm AG
VBI Vaccines Inc
Vical Inc



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