

2020 Spinal Muscular Atrophy (SMA) Drug Pipeline Report- Current Status, Phase, Mechanism, Route of Administration, and Companies, of Pre-Clinical And Clinical Drugs

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Abstracts

The 2020 Spinal Muscular Atrophy (SMA) pipeline report presents a comprehensive overview of the research and development of Spinal Muscular Atrophy (SMA) drug candidates. It presents drugs in development that could potentially reach the market in the next 5 to 10 years: Four drugs in Research Phase, 17 drugs in Pre-clinical phase, three drugs in Phase 1, four drugs in Phase 2 and two drugs in Phase 3

As of February 2020, the Spinal Muscular Atrophy (SMA) pipeline remains robust with 30 therapeutic candidates under development. An increasing number of companies are actively participating in the development of Spinal Muscular Atrophy (SMA) treatment. Diverse types of targeted therapies are being explored through clinical trials including Acetylcholine stimulant; activin II receptor antagonist; GABA transaminase inhibitor; MAS receptor agonists; myostatin inhibitor; Nerve tissue protein stimulants; oxygen species (ROS) inhibitor; Potassium channel antagonist; Ryanodine receptor calcium release channel modulators; SMN protein activator; survival motor neuron 2 (SMN2) protein modulator; Troponin stimulants.

The report provides complete details of pipeline drugs including the development phase, mechanism of action, companies involved, clinical trial developments, molecule type, and other details. Further, the report also provides Spinal Muscular Atrophy (SMA) drug development history, latest news, and other developments.

It assists companies, governments, investors and research organizations to understand the current status in 2020 and possible development in the next 10 years. Further, it



enables readers to track new companies in the market and their developments. The product portfolio of different companies and their growth strategies are also detailed in the report.

PUBLISHER EXPERTISE

VPAResearch online databases analyze pipeline drugs and developments for over 2,000 diseases worldwide. All our reports and databases are developed through intensive primary and secondary research methods. The insights and data presented in the databases are validated through industry experts and represent completely unbiased opinions.

SCOPE:

The report scope comprises of both pre-clinical phase and clinical phase development drugs for Spinal Muscular Atrophy (SMA) development

Spinal Muscular Atrophy (SMA) pipeline compounds and molecules under study by both large scale and small companies are included in the report

Spinal Muscular Atrophy (SMA) pipeline across different phases including discovery, research, and pre-clinical stage, phase 1, phase 2, phase 3 and pre-registration phases are covered

Drug profile comprising of current development status, regulatory progress, companies, sponsors, mechanism of action, route of administration, molecule, and discovery details are covered

Further, orphan drug status, fast track designation, different grants awarded and special status for Spinal Muscular Atrophy (SMA) pipeline candidates included

Business overview and snapshot of all companies involved in Spinal Muscular Atrophy (SMA) pipeline are included

Latest market and pipeline developments are provided in the report

Spinal Muscular Atrophy (SMA) pipeline companies included in the report are- Allianz Pharmascience Ltd, Armgo Pharma Inc, AurimMed Pharma Inc, Biogen Inc, BioMarin



Pharmaceutical Inc, Biophytis SA, Cytokinetics Inc, Exicure Inc, Genea Biocells, Genethon SA, GNT Pharma Co Ltd, Kowa Co Ltd, Neurodyn Life Sciences Inc, Neurotune AG, Novartis AG, Ono Pharmaceutical Co Ltd, PTC Therapeutics Inc, Ractigen Therapeutics Inc, Reborna Biosciences Inc, Recursion Pharmaceuticals Inc, Sarepta Therapeutics Inc, Scholar Rock Inc, Skyhawk Therapeutics Inc, Spotlight Innovation Inc, Vybion Inc, Xcelthera INC

Spinal Muscular Atrophy (SMA) pipeline drugs profiled in the report include- ALZ002, ARM210, SMN2 target for Spinal Muscular Atrophy, ALG-802, BIIB110, amifampridine, BIO-103, reldesemtiv (CK-2127107), SMN2 mRNA for Spinal Muscular Atrophy, Small Molecules for Spinal Muscular Atrophy, SMN1 gene therapy for spinal muscular atrophy, AAD-2004 (Crisdesalazine), Sodium valproate (K-828-SP), ND602, NT-1654, branaplam (LMI070), maresin 1, risdiplam, RAG-06, TEC-1, REC-0000716, REC-0001202, PMO25, SRK-105, Small Molecule for Spinal Muscular Atrophy, LDN-5178, STL-182, INT41, Xcel-hNu, Xcel-hNuP



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