

# **Retinoic Acid Receptor Gamma (RAR Gamma or Nuclear Receptor Subfamily 1 Group B Member 3 or NR1B3 or RARG) - Pipeline Review, H2 2018**

<https://marketpublishers.com/r/RFEFF89DE91EN.html>

Date: August 2018

Pages: 75

Price: US\$ 3,500.00 (Single User License)

ID: RFEFF89DE91EN

## **Abstracts**

Retinoic Acid Receptor Gamma (RAR Gamma or Nuclear Receptor Subfamily 1 Group B Member 3 or NR1B3 or RARG) - Pipeline Review, H2 2018

### **SUMMARY**

According to the recently published report 'Retinoic Acid Receptor Gamma - Pipeline Review, H2 2018'; Retinoic Acid Receptor Gamma (RAR Gamma or Nuclear Receptor Subfamily 1 Group B Member 3 or NR1B3 or RARG) pipeline Target constitutes close to 21 molecules.

Retinoic Acid Receptor Gamma (RAR Gamma or Nuclear Receptor Subfamily 1 Group B Member 3 or NR1B3 or RARG) - Retinoic acid receptor gamma (RARG) is a nuclear receptor that is encoded by the RARG gene. RARG bind as heterodimers to the target response elements in response to their ligands (all-trans or 9-cis retinoic acid) and regulate gene expression in various biological processes. In the absence or presence of hormone ligand, acts mainly as an activator of gene expression due to weak binding to co-repressors. It is required for limb bud development.

The report 'Retinoic Acid Receptor Gamma - Pipeline Review, H2 2018' outlays comprehensive information on the Retinoic Acid Receptor Gamma (RAR Gamma or Nuclear Receptor Subfamily 1 Group B Member 3 or NR1B3 or RARG) targeted therapeutics, complete with analysis by indications, stage of development, mechanism of action (MoA), route of administration (RoA) and molecule type; that are being developed by Companies/Universities.

It also reviews key players involved in Retinoic Acid Receptor Gamma (RAR Gamma or Nuclear Receptor Subfamily 1 Group B Member 3 or NR1B3 or RARG) targeted therapeutics development with respective active and dormant or discontinued projects. Currently, The molecules developed by companies in Pre-Registration, Filing rejected/Withdrawn, Phase III, Phase II, Preclinical and Discovery stages are 1, 1, 5, 6, 7 and 1 respectively.

Report covers products from therapy areas Dermatology, Musculoskeletal Disorders, Immunology, Oncology, Ophthalmology, Respiratory, Gastrointestinal and Genetic Disorders which include indications Acne Vulgaris, Bronchopulmonary Dysplasia, Allergies, Autoimmune Disorders, Congenital Ichthyosis, Fibrodysplasia Ossificans Progressiva (Myositis Ossificans Progressiva), Fibrosis, Gorlin Syndrome (Basal Cell Nevus Syndrome/Nevoid Basal Cell Carcinoma Syndrome), Head And Neck Cancer, Inflammatory Bowel Disease, Keratoconjunctivitis Sicca (Dry Eye), Multiple Hereditary Exostoses, Plaque Psoriasis (Psoriasis Vulgaris) and Retinopathy Of Prematurity.

**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 Retinoic Acid Receptor Gamma (RAR Gamma or Nuclear Receptor Subfamily 1 Group B Member 3 or NR1B3 or RARG)

The report reviews Retinoic Acid Receptor Gamma (RAR Gamma or Nuclear Receptor Subfamily 1 Group B Member 3 or NR1B3 or RARG) 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 Retinoic Acid Receptor Gamma (RAR

Gamma or Nuclear Receptor Subfamily 1 Group B Member 3 or NR1B3 or RARG) targeted therapeutics and enlists all their major and minor projects

The report assesses Retinoic Acid Receptor Gamma (RAR Gamma or Nuclear Receptor Subfamily 1 Group B Member 3 or NR1B3 or RARG) 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 Retinoic Acid Receptor Gamma (RAR Gamma or Nuclear Receptor Subfamily 1 Group B Member 3 or NR1B3 or RARG) 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 Retinoic Acid Receptor Gamma (RAR Gamma or Nuclear Receptor Subfamily 1 Group B Member 3 or NR1B3 or RARG)

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 Retinoic Acid Receptor Gamma (RAR Gamma or Nuclear Receptor Subfamily 1 Group B

Member 3 or NR1B3 or RARG) 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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Galderma SA

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Phosphagenics Ltd

Promius Pharma LLC

Sol-Gel Technologies Ltd

Retinoic Acid Receptor Gamma (RAR Gamma or Nuclear Receptor Subfamily 1 Group B Member 3 or NR1B3 or RARG) - Drug Profiles

(adapalene + benzoyl peroxide + clindamycin phosphate) - Drug Profile

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Mechanism Of Action

R&D Progress

(adapalene + clindamycin hydrochloride) - Drug Profile

Product Description

Mechanism Of Action

R&D Progress

(benzoyl peroxide + tretinoin) - Drug Profile

Product Description

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(benzoyl peroxide + tretinoin) - Drug Profile

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Featured News & Press Releases

Jun 18, 2018: FDA declines to approve Valeant's plaque psoriasis lotion

Jun 18, 2018: FDA Issues Complete Response Letter For DUOBRII (Halobetasol Propionate and Tazarotene) Lotion

May 23, 2018: Clementia Reports Positive Phase 2 Part B Data Showing Treatment with Palovarotene Significantly Reduces New Bone Growth in Patients with FOP

Apr 20, 2018: Clementia Initiates Phase 2 MO-Ped Trial for Palovarotene in Patients with Multiple Osteochondromas

Apr 09, 2018: Ortho Dermatologics Announces Publication of Pivotal Efficacy And Safety Data For Psoriasis Treatment DUOBRII In The Journal of the American Academy of Dermatology

Feb 15, 2018: Sol-Gel's Phase 2 Data on TWIN to be Presented at the 2018 American Academy of Dermatology Annual Meeting

Jan 12, 2018: Ortho Dermatologics Announces U.S. FDA Filing Acceptance For IDP-121 Acne Treatment In Lotion Form

Dec 12, 2017: Clementia Initiates Pivotal Phase 3 MOVE Trial for Palovarotene in Patients with Fibrodysplasia Ossificans Progressiva

Nov 20, 2017: Preclinical study demonstrates promising treatment for rare bone disease

Nov 09, 2017: Promius Pharma Initiates Phase 3 Clinical Trials of DFD-03 for The Treatment of Acne Vulgaris

Nov 02, 2017: Ortho Dermatologics Announces U.S. FDA Filing Acceptance For IDP-118, Novel Plaque Psoriasis Treatment

Sep 05, 2017: Ortho Dermatologics Submits New Drug Application To The U.S. Food And Drug Administration For Psoriasis Treatment IDP-118

Aug 31, 2017: Clementia Announces Data Presentations at Upcoming Medical Conferences in September

Jul 20, 2017: Sol-Gel Announces Positive Phase 2 Clinical Trial Results for TWIN in Patients with Acne Vulgaris

Jun 05, 2017: Clementia Pharmaceuticals to Host Symposium at the 8th International



Conference on Children's Bone Health

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### COMPANIES MENTIONED

3SBio Inc

Bausch Health Companies Inc

Boehringer Ingelheim GmbH

Clementia Pharmaceuticals Inc

Galderma SA

Lee's Pharmaceutical Holdings Ltd

Phosphagenics Ltd

Promius Pharma LLC

Sol-Gel Technologies Ltd

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