# Neurofibromatoses Type I Trends, Analysis and Review during H1 2017 To 2022

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Healthcare

Summary

Neurofibromatosis type 1 (NF1), also called von Recklinghausen's disease, is a rare genetic disorder characterized by the development of multiple noncancerous (benign) tumors of nerves and skin (neurofibromas). This is transmitted on chromosome 17 and is caused by mutation of the NF1 gene. Symptoms include liver enlargement, glioma, Lisch nodules and pheochromocytoma. Treatment includes pain medications, surgery, chemotherapy and radiation therapy.

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Report Highlights

The  Pharmaceutical and Healthcare latest pipeline guide Neurofibromatoses Type I (Von Recklinghausen’s Disease) - Pipeline Review, H1 2017, provides comprehensive information on the therapeutics under development for Neurofibromatoses Type I (Von Recklinghausen’s Disease) (Genetic Disorders), complete with analysis by stage of development, drug target, mechanism of action (MoA), route of administration (RoA) and molecule type. The guide covers the descriptive pharmacological action of the therapeutics, its complete research and development history and latest news and press releases.

The Neurofibromatoses Type I (Von Recklinghausen’s Disease) (Genetic Disorders) pipeline guide also reviews of key players involved in therapeutic development for Neurofibromatoses Type I (Von Recklinghausen’s Disease) and features dormant and discontinued projects. The guide covers therapeutics under Development by Companies /Universities /Institutes, the molecules developed by Companies in Phase II, Phase I and Preclinical stages are 1, 1 and 2 respectively. Similarly, the Universities portfolio in Discovery stages comprises 1 molecules, respectively.

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