

## NEUROFIBROMATOSIS TYPE 1

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[Neurofibromatosis type 1 - Genetics Home Reference - NIH](#)

Neurofibromatosis type 1 is a condition characterized by changes in skin coloring (pigmentation) and the growth of tumors along nerves in the skin, brain, and other

[Neurofibromatosis 1 - NORD \(National Organization for Rare ...](#)

Eleftherios F, Kolanczyk M, Schindeler A, et al. Skeletal abnormalities in neurofibromatosis type 1: approaches to therapeutic options. Am J Med Genet A. 2009 Oct;

[Neurofibromatosis type 1 - Wikipedia](#)

Neurofibromatosis type 1 (NF-1) is a complex, multi-system human disorder caused by the mutation of a gene on chromosome 17 that is responsible for production of a

[Neurofibromatosis 1 and 2: Symptoms, Treatments, Causes](#)

The disorders are known as neurofibromatosis type 1 (NF1) and neurofibromatosis type 2 (NF2). NF1 is the more common type of neurofibromatosis.

[Neurofibromatosis type 1 \(NF1\) - Tumour Foundation of BC](#)

NF1 - Neurofibromatosis type 1, formerly known as von Recklinghausen NF or Peripheral NF, is the most common of the three types.

[Neurofibromatosis type 1 - NHS](#)

Neurofibromatosis type 1 (NF1) is a genetic condition that causes tumours to grow along your nerves. The tumours are usually non-cancerous (benign) but may cause a

[Neurofibromatosis Type 1 | Children's Hospital of Philadelphia](#)

Neurofibromatosis type 1 (NF1) is a genetic disorder characterized by the presence of skin differences. Ten percent of people with NF1 develop cancerous neurofibromas.

[Neurofibromatosis Type 1 \(NF1\) Fact Sheet](#)

Visit us on the web at [benf.bc.ca/ca](http://benf.bc.ca/ca) Fact Sheet.

[Neurofibromatosis Type 1](#) Neurofibromatosis Type 1 (NF1) is the most common single gene disorder to affect the human

[Neurofibromatosis type 2 | Genetic and Rare Diseases ...](#)

A collection of disease information resources and questions answered by our Genetic and Rare Diseases Information Specialists for Neurofibromatosis type 2

[Neurofibromatosis type II - Wikipedia](#)

Neurofibromatosis type II (also known as MISME syndrome - multiple inherited schwannomas, meningiomas, and ependymomas) is a genetic condition

[Harnuntersuchungsmethoden Und Ihre Klinische Bedeutung Stabe Mit Undeformierbaren Querschnitten Die Metallhüttenkunde Lehrbuch Der Organischchemischen Methodik](#)

which may be inherited.

[Neurofibromatosis Type 1: Practice Essentials, Background ...](#)

Neurofibromatosis type 1 (NF1) is a multisystem genetic disorder that commonly is associated with cutaneous, neurologic, and orthopedic manifestations. It

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(c.1354 C>T) in exon 10a of the NF1 gene. Translation of this mutation would result in the replacement

[Neurofibromatosis type 1 - Symptoms - NHS](#)

The severity of neurofibromatosis type 1 (NF1) can vary considerably. The condition is often mild and causes no serious health problems but some people experience

[Neurofibromatosis-1: MedlinePlus Medical Encyclopedia](#)

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