

Spinocerebellar Ataxia

SSA - POMS: DI 23022.500 - Spinocerebellar Ataxia -

SPINOCEREBELLAR ATAXIA. ALTERNATE NAMES. SCA; Infantile-onset Spinocerebellar Ataxia; Autosomal Dominant Spinocerebellar Ataxia (ADSCA) DESCRIPTION

<https://secure.ssa.gov/apps10/poms.nsf/lnx/0423022500>

Spinocerebellar Ataxia | Specialist -

Spinocerebellar Ataxia (SCA) is the name given to a group of hereditary conditions where the cerebellum (a part of the

<http://spinocerebellarataxia.org/>

Spinocerebellar Ataxia - PatientsLikeMe -

What is Spinocerebellar Ataxia? Spinocerebellar ataxia is a progressive, degenerative, genetic disease with multiple types which are characterized by slowly

<https://www.patientslikeme.com/conditions/529-spinocerebellar-ataxia>

Spinocerebellar ataxia | definition of -

Looking for online definition of spinocerebellar ataxia in the Medical Dictionary? spinocerebellar ataxia explanation free. spinocellular; spinocerebellar;

<http://medical-dictionary.thefreedictionary.com/Spinocerebellar+ataxia>

Spinocerebellar ataxia | Radiology Reference -

Spinocerebellar ataxias comprise a large (and expanding) group of diseases characterised by degeneration of the spinal cord and cerebellum. There are well over 25

<http://radiopaedia.org/articles/spinocerebellar-ataxia>

OMIM Entry - # 183090 - SPINOCEREBELLAR ATAXIA 2; -

183090 - spinocerebellar ataxia 2; sca2 - spinocerebellar atrophy ii;; olivopontocerebellar atrophy, holguin type;; olivopontocerebellar atrophy ii

<http://omim.org/entry/183090>

Spinocerebellar ataxia 2 | Disease | Overview | -

Jun 22, 2014 A collection of disease information resources and questions answered by our Genetic and Rare Diseases Information Specialists for Spinocerebellar ataxia 2

<https://rarediseases.info.nih.gov/gard/4072/spinocerebellar-ataxia-2/resources/1>

Spinocerebellar ataxia - Wikipedia, the free -

Spinocerebellar ataxia (SCA) or also known as Spinocerebellar atrophy or Spinocerebellar degeneration, is a progressive, degenerative, [1] genetic disease with

https://en.m.wikipedia.org/wiki/Spinocerebellar_ataxia

Spinocerebellar Ataxia -- Medical Definition -

spinocerebellar ataxia Type: Term. Definitions: 1. a generic term now increasingly used to describe autosomal dominant-inherited ataxias that have a progressive course.

<http://www.medilexicon.com/medicaldictionary.php?t=8211>

Ataxia - NHS Choices -

Ataxia is the term for a group of disorders that affect co-ordination, balance and speech. Find out about the main types, what causes them, and how they're treated.

<http://www.nhs.uk/Conditions/Ataxia/Pages/Introduction.aspx>

Ataxia - Mayo Clinic -

Ataxia describes a lack of muscle control during voluntary movements, NINDS ataxias and cerebellar or spinocerebellar degeneration information page.

<http://www.mayoclinic.org/diseases-conditions/ataxia/basics/definition/CON-20030428>

Spinocerebellar ataxia - Psychology Wiki -

Spinocerebellar ataxia (SCA) is a genetic disease with multiple types, each of which could be considered a disease in its own right.

http://psychology.wikia.com/wiki/Spinocerebellar_ataxia

Spinocerebellar Ataxia with Axonal Neuropathy - -

Jan 23, 2012 Spinocerebellar ataxia with axonal neuropathy (SCAN1) is a neurodegenerative disorder that is inherited in an autosomal recessive pattern. SCAN1 is

<http://www.webmd.com/cancer/brain-cancer/spinocerebellar-ataxia-with-axonal-neuropathy>

Ataxia - Wikipedia, the free encyclopedia -

Ataxia (from Greek - [a negative prefix] + - [order] = "lack of order") is a neurological sign consisting of lack of voluntary coordination of muscle

<http://en.wikipedia.org/wiki/Ataxia>

Spinocerebellar Ataxia : Dr David Steenblock -

Spinocerebellar Ataxia. Spinocerebellar ataxia (SCA) is a slowly progressive genetic disorder characterized by slowly poor coordination of hands, speech, and eye

<http://sironatechnology.com/steenblock/spinocerebellar-ataxia/>

National Ataxia Foundation - Fact Sheets -

Fact Sheets. The National Ataxia Foundation has an extensive catalog of free and for purchase publications to help you stay informed. Free Fact Sheets

<http://www.ataxia.org/resources/publications.aspx>

Spinocerebellar Degeneration | Specialist -

Spinocerebellar Degeneration, also known as Spinocerebellar Ataxia or SCA, is a progressive disease that can affect the spine, the cerebellum,

<http://spinocerebellarataxia.org/spinocerebellar-degeneration/>

Spinocerebellar ataxia, dominantly inherited -

Spinocerebellar ataxia, dominantly inherited forms is a rare disorder described in the database for rare diseases of the Swedish National Board of Health and Welfare.

<http://www.socialstyrelsen.se/rarediseases/spinocerebellarataxia-dominant>

Spinocerebellar ataxia - Living With Ataxia -

Spinocerebellar ataxia. Spinocerebellar ataxia (SCA) is a genetic disease with multiple types, each of which could be considered a disease in its own right.

<http://www.livingwithcerebralpalsy.com/spinocerebellar-ataxia.php>

Spinocerebellar Ataxia -

There are several different types of Spinocerebellar Ataxia, the characteristics of each of which are such that they could be categorized as a separate diseases.

<http://spinocerebellarataxia.net/>

Spinocerebellar Ataxia Medical Definition | -

Medical Definition of SPINOCEREBELLAR ATAXIA: any of a group of inherited neurodegenerative disorders that are characterized by cerebellar dysfunction manifested

<http://www.merriam-webster.com/medical/spinocerebellar%20ataxia>

Spinocerebellar Ataxia Case Study | HHMI's -

Dr. Huda Zoghbi interviews Milan Cloud, a patient who has inherited the neurological disorder spinocerebellar ataxia 1, or SCA1.

<http://www.hhmi.org/biointeractive/spinocerebellar-ataxia-case-study>

Spinocerebellar Ataxia Australia Inc -

Spinocerebellar Ataxia Australia Support Group This site is hosted by. Javascript Menu by Deluxe-Menu.com: Press F5 to refresh page Latest news updated

<http://www.scars.org.au/>

Spinocerebellar ataxia - Genes and Disease - NCBI -

Persons with spinocerebellar ataxia experience a degeneration of the spinal cord and the cerebellum, the small fissured mass at the base of the brain, behind the

<http://www.ncbi.nlm.nih.gov/books/NBK22234/>

The Spinocerebellar Ataxias - PubMed Central (PMC) -

The definition of ataxia is loss of coordination, particularly of gait. Thus, when a clinician diagnoses someone with ataxia, it typically is someone with gait

<http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2739122/>

Spinocerebellar ataxia | Disease | Overview | -

Dec 01, 2014 Spinocerebellar ataxia (SCA) is a group of inherited conditions that are characterized by degenerative changes of the nervous system (brain and spinal cord

<https://rarediseases.info.nih.gov/gard/10748/spinocerebellar-ataxia/resources/1>

Ataxia Symptoms - Mayo Clinic -

Ataxia, a sign of a number of neurological disorders, may cause: Poor coordination; NINDS ataxias and cerebellar or spinocerebellar degeneration information page.

<http://www.mayoclinic.org/diseases-conditions/ataxia/basics/symptoms/CON-20030428>

Ataxia - Types - NHS Choices -

Some types of ataxia affect children from an early age, while other types may not develop until much later in adulthood. Depending on the type of ataxia, the symptoms

<http://www.nhs.uk/Conditions/Ataxia/Pages/Symptoms.aspx>

Ataxia Spinocerebellar (SCA) - Facts and -

Spinocerebellar ataxia (SCA) is a genetically inherited disorder characterized by abnormalities in brain functioning

<http://www.disabled-world.com/disability/types/sca.php>

OMIM Entry - # 164500 - SPINOCEREBELLAR ATAXIA 7; -

164500 - spinocerebellar ataxia 7; sca7 - olivopontocerebellar atrophy iii; opca3;; opca iii;; opca with retinal degeneration;; opca with macular

<http://www.omim.org/entry/164500>