amyotrophic lateral sclerosis

Amyotrophic lateral sclerosis (ALS) is a progressive disease that affects motor neurons, which are specialized nerve cells that control muscle movement. These nerve cells are found in the spinal cord and the brain. In ALS, motor neurons die (atrophy) over time, leading to muscle weakness, a loss of muscle mass, and an inability to control movement.

There are many different types of ALS; these types are distinguished by their signs and symptoms and their genetic cause or lack of clear genetic association. Most people with ALS have a form of the condition that is described as sporadic, which means it occurs in people with no apparent history of the disorder in their family. People with sporadic ALS usually first develop features of the condition in their late fifties or early sixties. A small proportion of people with ALS, estimated at 5 to 10 percent, have a family history of ALS or a related condition called frontotemporal dementia (FTD), which is a progressive brain disorder that affects personality, behavior, and language. The signs and symptoms of familial ALS typically first appear in one's late forties or early fifties. Rarely, people with familial ALS develop symptoms in childhood or their teenage years. These individuals have a rare form of the disorder known as juvenile ALS.

The first signs and symptoms of ALS may be so subtle that they are overlooked. The earliest symptoms include muscle twitching, cramping, stiffness, or weakness. Affected individuals may develop slurred speech (dysarthria) and, later, difficulty chewing or swallowing (dysphagia). Many people with ALS experience malnutrition because of reduced food intake due to dysphagia and an increase in their body's energy demands (metabolism) due to prolonged illness. Muscles become weaker as the disease progresses, and arms and legs begin to look thinner as muscle tissue atrophies. Individuals with ALS eventually lose muscle strength and the ability to walk. Affected individuals eventually become wheelchair-dependent and increasingly require help with personal care and other activities of daily living. Over time, muscle weakness causes affected individuals to lose the use of their hands and arms. Breathing becomes difficult because the muscles of the respiratory system weaken. Most people with ALS die from respiratory failure within 2 to 10 years after the signs and symptoms of ALS first appear; however, disease progression varies widely among affected individuals.

Approximately 20 percent of individuals with ALS also develop FTD. Changes in personality and behavior may make it difficult for affected individuals to interact with others in a socially appropriate manner. Communication skills worsen as the disease progresses. It is unclear how the development of ALS and FTD are related. Individuals who develop both conditions are diagnosed as having ALS-FTD.
A rare form of ALS that often runs in families is known as ALS-parkinsonism-dementia complex (ALS-PDC). This disorder is characterized by the signs and symptoms of ALS, in addition to a pattern of movement abnormalities known as parkinsonism, and a progressive loss of intellectual function (dementia). Signs of parkinsonism include unusually slow movements (bradykinesia), stiffness, and tremors. Affected members of the same family can have different combinations of signs and symptoms.

**Frequency**

About 5,000 people in the United States are diagnosed with ALS each year. Worldwide, this disorder occurs in 2 to 5 per 100,000 individuals. Only a small percentage of cases have a known genetic cause.

Among the Chamorro people of Guam and people from the Kii Peninsula of Japan, ALS-PDC can be 100 times more frequent than ALS is in other populations. ALS-PDC has not been reported outside of these populations.

**Genetic Changes**

Mutations in several genes can cause familial ALS and contribute to the development of sporadic ALS. Mutations in the \textit{C9orf72} gene account for 30 to 40 percent of familial ALS in the United States and Europe. Worldwide, \textit{SOD1} gene mutations cause 15 to 20 percent of familial ALS, and \textit{TARDBP} and \textit{FUS} gene mutations each account for about 5 percent of cases. The other genes that have been associated with familial ALS each account for a small proportion of cases. It is estimated that 60 percent of individuals with familial ALS have an identified genetic mutation. The cause of the condition in the remaining individuals is unknown.

The \textit{C9orf72}, \textit{SOD1}, \textit{TARDBP}, and \textit{FUS} genes are key to the normal functioning of motor neurons and other cells. It is unclear how mutations in these genes contribute to the death of motor neurons, but it is thought that motor neurons are more sensitive to disruptions in function because of their large size. Most motor neurons affected by ALS have a buildup of protein clumps (aggregates); however, it is unknown whether these aggregates are involved in causing ALS or are a byproduct of the dying cell.

In some cases of familial ALS due to mutations in other genes, studies have identified the mechanisms that lead to ALS. Some gene mutations lead to a disruption in the development of axons, the specialized extensions of nerve cells (such as motor neurons) that transmit nerve impulses. The altered axons may impair transmission of impulses from nerves to muscles, leading to muscle weakness and atrophy. Other mutations lead to a slowing in the transport of materials needed for the proper function of axons in motor neurons, eventually causing the motor neurons to die. Additional gene mutations prevent the breakdown of toxic substances, leading to their buildup in nerve cells. The accumulation of toxic substances can damage motor neurons and eventually cause cell death. In some cases of ALS, it is unknown how the gene mutation causes the condition.
The cause of sporadic ALS is largely unknown but probably involves a combination of genetic and environmental factors. Variations in many genes, including the previously mentioned genes involved in transmission of nerve impulses and transportation of materials within neurons, increase the risk of developing ALS. Gene mutations that are risk factors for ALS may add, delete, or change DNA building blocks (nucleotides), resulting in the production of a protein with an altered or reduced function. While genetic variations have been associated with sporadic ALS, not all genetic factors have been identified and it is unclear how most genetic changes influence the development of the disease. People with a gene variation that increases their risk of ALS likely require additional genetic and environmental triggers to develop the disorder.

Inheritance Pattern

About 90 to 95 percent of ALS cases are sporadic, which means they are not inherited. An estimated 5 to 10 percent of ALS is familial and caused by mutations in one of several genes. The pattern of inheritance varies depending on the gene involved. Most cases are inherited in an autosomal dominant pattern, which means one copy of the altered gene in each cell is sufficient to cause the disorder. In most cases, an affected person has one parent with the condition. Some people who inherit a familial genetic mutation known to cause ALS never develop features of the condition. (This situation is known as reduced penetrance.) It is unclear why some people with a mutated gene develop the disease and other people with a mutated gene do not.

Less frequently, ALS is inherited in an autosomal recessive pattern, which means both copies of the gene in each cell have mutations. The parents of an individual with an autosomal recessive condition each carry one copy of the mutated gene, but they typically do not show signs and symptoms of the condition. Because an affected person's parents are not affected, autosomal recessive ALS is often mistaken for sporadic ALS even though it is caused by a familial genetic mutation.

Very rarely, ALS is inherited in an X-linked dominant pattern. X-linked conditions occur when the gene associated with the condition is located on the X chromosome, which is one of the two sex chromosomes. In females (who have two X chromosomes), a mutation in one of the two copies of the gene in each cell is sufficient to cause the disorder. In males (who have only one X chromosome), a mutation in the only copy of the gene in each cell causes the disorder. In most cases, males tend to develop the disease earlier and have a decreased life expectancy compared with females. A characteristic of X-linked inheritance is that fathers cannot pass X-linked traits to their sons.

Other Names for This Condition

- ALS
- amyotrophic lateral sclerosis with dementia
• Charcot disease
• dementia with amyotrophic lateral sclerosis
• Lou Gehrig disease
• motor neuron disease, amyotrophic lateral sclerosis

Diagnosis & Management

Genetic Testing

• Genetic Testing Registry: Amyotrophic lateral sclerosis

• Genetic Testing Registry: Amyotrophic lateral sclerosis 14, with or without frontotemporal dementia

• Genetic Testing Registry: Amyotrophic lateral sclerosis 15, with or without frontotemporal dementia

• Genetic Testing Registry: Amyotrophic lateral sclerosis 16, juvenile

• Genetic Testing Registry: Amyotrophic lateral sclerosis 17

• Genetic Testing Registry: Amyotrophic lateral sclerosis 18

• Genetic Testing Registry: Amyotrophic lateral sclerosis 19

• Genetic Testing Registry: Amyotrophic lateral sclerosis 20

• Genetic Testing Registry: Amyotrophic lateral sclerosis and/or frontotemporal dementia 1

• Genetic Testing Registry: Amyotrophic lateral sclerosis-parkinsonism/dementia complex 1

• Genetic Testing Registry: Amyotrophic lateral sclerosis type 1

• Genetic Testing Registry: Amyotrophic lateral sclerosis type 2
• Genetic Testing Registry: Amyotrophic lateral sclerosis type 3

• Genetic Testing Registry: Amyotrophic lateral sclerosis type 4

• Genetic Testing Registry: Amyotrophic lateral sclerosis type 5

• Genetic Testing Registry: Amyotrophic lateral sclerosis type 6

• Genetic Testing Registry: Amyotrophic lateral sclerosis type 7

• Genetic Testing Registry: Amyotrophic lateral sclerosis type 8

• Genetic Testing Registry: Amyotrophic lateral sclerosis type 9

• Genetic Testing Registry: Amyotrophic lateral sclerosis type 10

• Genetic Testing Registry: Amyotrophic lateral sclerosis type 11

• Genetic Testing Registry: Amyotrophic lateral sclerosis type 12

• Genetic Testing Registry: Frontotemporal dementia and/or amyotrophic lateral sclerosis 3

• Genetic Testing Registry: Frontotemporal dementia and/or amyotrophic lateral sclerosis 4

Other Diagnosis and Management Resources

• GeneReview: ALS2-Related Disorders
  https://www.ncbi.nlm.nih.gov/books/NBK1243

• GeneReview: Amyotrophic Lateral Sclerosis Overview
  https://www.ncbi.nlm.nih.gov/books/NBK1450

• GeneReview: C9orf72-Related Amyotrophic Lateral Sclerosis and Frontotemporal Dementia
  https://www.ncbi.nlm.nih.gov/books/NBK268647

• GeneReview: TARDBP-Related Amyotrophic Lateral Sclerosis
  https://www.ncbi.nlm.nih.gov/books/NBK5942
• Massachusetts General Hospital: How is ALS Diagnosed?  
http://www.massgeneral.org/als/patienteducation/DiagnosingALS.aspx

• MedlinePlus Encyclopedia: Amyotrophic Lateral Sclerosis  
https://medlineplus.gov/ency/article/000688.htm

General Information from MedlinePlus
• Diagnostic Tests  
https://medlineplus.gov/diagnostictests.html
• Drug Therapy  
https://medlineplus.gov/drugtherapy.html
• Genetic Counseling  
https://medlineplus.gov/geneticcounseling.html
• Palliative Care  
https://medlineplus.gov/palliativecare.html
• Surgery and Rehabilitation  
https://medlineplus.gov/surgeryandrehabilitation.html

Additional Information & Resources
MedlinePlus
• Encyclopedia: Amyotrophic Lateral Sclerosis  
https://medlineplus.gov/ency/article/000688.htm
• Health Topic: Amyotrophic Lateral Sclerosis  
https://medlineplus.gov/amyotrophiclateralsclerosis.html

Genetic and Rare Diseases Information Center
• Amyotrophic lateral sclerosis  

Additional NIH Resources
• National Institute of Neurological Disorders and Stroke  
https://www.ninds.nih.gov/Disorders/All-Disorders/Amyotrophic-Lateral-Sclerosis-ALS-Information-Page

Educational Resources
• Cleveland Clinic  
https://my.clevelandclinic.org/health/articles/als
• Disease InfoSearch: Amyotrophic Lateral Sclerosis  
http://www.diseaseinfosearch.org/Amyotrophic+Lateral+Sclerosis/400
• Disease InfoSearch: Frontotemporal dementia and-or amyotrophic lateral sclerosis
  http://www.diseaseinfosearch.org/Frontotemporal+dementia+and-or+amyotrophic+
lateral+sclerosis/8445

• KidsHealth from Nemours

• MalaCards: amyotrophic lateral sclerosis 1
  http://www.malacards.org/card/amyotrophic_lateral_sclerosis_1

• MalaCards: amyotrophic lateral sclerosis-parkinsonism/dementia complex
  http://www.malacards.org/card/amyotrophic_lateral_sclerosis_parkinsonism_dementia_complex

• Merck Manual Consumer Version
  peripheral-nerve-disorders/amyotrophic-lateral-sclerosis-and-other-motor-neuron-
diseases

• Muscular Dystrophy Association: Facts About Amyotrophic Lateral Sclerosis
  https://www.mda.org/disease/amyotrophic-lateral-sclerosis

• My46 Trait Profile
  https://www.my46.org/traits-document?trait=Amyotrophic%20lateral%20sclerosis
  &type=profile

• Orphanet: Amyotrophic lateral sclerosis
  http://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=803

• Orphanet: Amyotrophic lateral sclerosis-parkinsonism-dementia complex
  http://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=90020

• Orphanet: Frontotemporal dementia with motor neuron disease
  http://www.orpha.net/consor/cgi-bin/OC_Exp.php?Lng=EN&Expert=275872

• University of Michigan Health System
  http://www.uofmhealth.org/health-library/hw179630#hw179632

• Washington University, St. Louis Neuromuscular Disease Center: Hereditary ALS
  http://neuromuscular.wustl.edu/synmot.html#Hereditaryals

• World Federation of Neurology-ALS
  http://www.wfnals.org
Patient Support and Advocacy Resources

- Centers for Disease Control and Prevention: National Amyotrophic Lateral Sclerosis Registry  
- Les Turner Amyotrophic Lateral Sclerosis Foundation  
  http://lesturnerals.org/what-is-als/
- National Organization for Rare Disorders (NORD)  
  https://rarediseases.org/rare-diseases/amyotrophic-lateral-sclerosis/
- Project ALS  
  http://www.projectals.org/
- The ALS Association  
  http://www.alsa.org/
- The ALS Society of Canada  
  https://www.als.ca/

GeneReviews

- ALS2-Related Disorders  
  https://www.ncbi.nlm.nih.gov/books/NBK1243
- Amyotrophic Lateral Sclerosis Overview  
  https://www.ncbi.nlm.nih.gov/books/NBK1450
- C9orf72-Related Amyotrophic Lateral Sclerosis and Frontotemporal Dementia  
  https://www.ncbi.nlm.nih.gov/books/NBK268647
- TARDBP-Related Amyotrophic Lateral Sclerosis  
  https://www.ncbi.nlm.nih.gov/books/NBK5942

ClinicalTrials.gov

- ClinicalTrials.gov  
  https://clinicaltrials.gov/ct2/results?cond=%22amyotrophic+lateral+sclerosis%22

Scientific Articles on PubMed

- PubMed  
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28Amyotrophic+Lateral+Sclerosis%5BMAJR%5D%29+AND+%28amyotrophic+lateral+sclerosis%5BTI%5D%29+AND+review%5Bpt%5D+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+720+days%22+AND+%22last+720+days%22+AND+%22last+720+days%22
AMYOTROPHIC LATERAL SCLEROSIS 1
http://omim.org/entry/105400

AMYOTROPHIC LATERAL SCLEROSIS 2, JUVENILE
http://omim.org/entry/205100

AMYOTROPHIC LATERAL SCLEROSIS 3
http://omim.org/entry/606640

AMYOTROPHIC LATERAL SCLEROSIS 4, JUVENILE
http://omim.org/entry/602433

AMYOTROPHIC LATERAL SCLEROSIS 5, JUVENILE
http://omim.org/entry/602099

AMYOTROPHIC LATERAL SCLEROSIS 6 WITH OR WITHOUT FRONTOTEMPORAL DEMENTIA
http://omim.org/entry/608030

AMYOTROPHIC LATERAL SCLEROSIS 7
http://omim.org/entry/608031

AMYOTROPHIC LATERAL SCLEROSIS 8
http://omim.org/entry/608627

AMYOTROPHIC LATERAL SCLEROSIS 9
http://omim.org/entry/611895

AMYOTROPHIC LATERAL SCLEROSIS 10 WITH OR WITHOUT FRONTOTEMPORAL DEMENTIA
http://omim.org/entry/612069

AMYOTROPHIC LATERAL SCLEROSIS 11
http://omim.org/entry/612577

AMYOTROPHIC LATERAL SCLEROSIS 12
http://omim.org/entry/613435

AMYOTROPHIC LATERAL SCLEROSIS 14 WITH OR WITHOUT FRONTOTEMPORAL DEMENTIA
http://omim.org/entry/613954

AMYOTROPHIC LATERAL SCLEROSIS 15 WITH OR WITHOUT FRONTOTEMPORAL DEMENTIA
http://omim.org/entry/300857

AMYOTROPHIC LATERAL SCLEROSIS 16, JUVENILE
http://omim.org/entry/614373
• AMYOTROPHIC LATERAL SCLEROSIS 17
  http://omim.org/entry/614696
• AMYOTROPHIC LATERAL SCLEROSIS 18
  http://omim.org/entry/614808
• AMYOTROPHIC LATERAL SCLEROSIS 19
  http://omim.org/entry/615515
• AMYOTROPHIC LATERAL SCLEROSIS 20
  http://omim.org/entry/615426
• AMYOTROPHIC LATERAL SCLEROSIS 22 WITH OR WITHOUT
  FRONOTEMPORAL DEMENTIA
  http://omim.org/entry/616208
• AMYOTROPHIC LATERAL SCLEROSIS-PARKINSONISM/DEMENTIA COMPLEX
  1
  http://omim.org/entry/105500
• FRONOTEMPORAL DEMENTIA AND/OR AMYOTROPHIC LATERAL
  SCLEROSIS 1
  http://omim.org/entry/105550
• FRONOTEMPORAL DEMENTIA AND/OR AMYOTROPHIC LATERAL
  SCLEROSIS 2
  http://omim.org/entry/615911
• FRONOTEMPORAL DEMENTIA AND/OR AMYOTROPHIC LATERAL
  SCLEROSIS 3
  http://omim.org/entry/616437
• FRONOTEMPORAL DEMENTIA AND/OR AMYOTROPHIC LATERAL
  SCLEROSIS 4
  http://omim.org/entry/616439

Sources for This Summary

• Andersen PM, Al-Chalabi A. Clinical genetics of amyotrophic lateral sclerosis: what do we really
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/21989245

• Andersen PM. Amyotrophic lateral sclerosis associated with mutations in the CuZn superoxide
  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/16469270


  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/21527311

• National Institute of Neurological Disorders and Stroke
  https://www.ninds.nih.gov/Disorders/All-Disorders/Amyotrophic-Lateral-Sclerosis-ALS-Information-Page

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/24981792
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4433003/

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  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4516382/

Reprinted from Genetics Home Reference:

Reviewed: March 2016
Published: August 15, 2017

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National Institutes of Health
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