Autism spectrum disorder

Autism spectrum disorder (ASD) is a condition that appears very early in childhood development, varies in severity, and is characterized by impaired social skills, communication problems, and repetitive behaviors. These difficulties can interfere with affected individuals' ability to function in social, academic, and employment settings. People with ASD also have an increased risk of psychiatric problems such as anxiety, depression, obsessive-compulsive disorder, and eating disorders.

From as early as 1 to 2 years of age, people with ASD have an impaired ability to interact with other people; they are often more comfortable dealing with objects. Affected individuals have difficulty understanding and using non-verbal social cues such as eye contact, facial expressions, gestures, and body language. Inability to recognize and use these cues makes it hard for affected individuals to understand the feelings of others or communicate their own feelings appropriately. Behavioral signs of ASD, such as reduced eye contact and social interaction, can sometimes be detected before age 2. However, the condition is usually diagnosed between ages 2 and 4, when more advanced communication and social skills, such as learning to play with others, typically begin to develop.

Repetitive behaviors in ASD can include simple actions such as rocking, hand-flapping, or repetition of words or noises (echolalia). Affected individuals often dwell on or repeatedly express particular thoughts; this behavior is called perseveration. People with ASD tend to be rigid about their established routines and may strongly resist disruptions such as changes in schedule. They may also have difficulty tolerating sensory stimuli such as loud noises or bright lights.

While social and communication difficulties and unusual behaviors define ASD, affected individuals can have a wide range of intellectual abilities and language skills. A majority of people with ASD have mild to moderate intellectual disability, while others have average to above-average intelligence. Some have particular cognitive abilities that greatly surpass their overall level of functioning, often in areas such as music, mathematics, or memory.

Some people with ASD do not speak at all, while others use language fluently. However, fluent speakers with ASD often have problems associated with verbal communication. They might speak in a monotone voice, have unusual vocal mannerisms, or choose unusual topics of conversation.

Several diagnoses that used to be classified as separate conditions are now grouped together under the diagnosis of ASD. For example, autistic disorder was a term that was used when affected individuals had limited or absent verbal communication, often in combination with intellectual disability. By contrast, Asperger syndrome was
a diagnosis formerly applied to affected individuals of average or above-average intelligence who were not delayed in their language development. The broader diagnosis of ASD was established because many affected individuals fall outside of the strict definitions of the narrower diagnoses, and their intellectual and communication abilities may change over time. However, some individuals who were previously diagnosed with one of the subtypes now do not meet all the criteria of the new umbrella diagnosis.

Frequency

ASD is a common condition, and affects almost five times as many males as females. The number of children diagnosed with ASD has been increasing rapidly in the past few decades. The prevalence of the disorder in the United States was estimated as 1 in 68 children in 2014, up from 1 in 88 only two years earlier. In the 1980s, before the term ASD was used, the prevalence of autism was reported to be about 1 in 2,000. However, it is unclear whether this represents a true increase in the prevalence of ASD or reflects changes in the way behaviors characteristic of the disorder have been diagnosed and categorized.

Causes

Changes in over 1,000 genes have been reported to be associated with ASD, but a large number of these associations have not been confirmed. Many common gene variations, most of which have not been identified, are thought to affect the risk of developing ASD, but not all people with the gene variation will be affected. Most of the gene variations have only a small effect, and variations in many genes can combine with environmental risk factors, such as parental age, birth complications, and others that have not been identified, to determine an individual’s risk of developing this complex condition. Non-genetic factors may contribute up to about 40 percent of ASD risk.

By contrast, in about 2 to 4 percent of people with ASD, rare gene mutations or chromosome abnormalities are thought to be the cause of the condition, often as a feature of syndromes that also involve additional signs and symptoms affecting various parts of the body. For example, mutations in the ADNP gene cause a disorder called ADNP syndrome. In addition to ASD and intellectual disability, this condition involves distinctive facial features and a wide variety of other signs and symptoms. Some of the other genes in which rare mutations are associated with ASD, often with other signs and symptoms, are ARID1B, ASH1L, CHD2, CHD8, DYRK1A, POGZ, SHANK3, and SYNGAP1. In most individuals with ASD caused by rare gene mutations, the mutations occur in only a single gene.

Many of the genes associated with ASD are involved in the development of the brain. The proteins produced from these genes affect multiple aspects of brain development, including production, growth, and organization of nerve cells (neurons). Some affect the number of neurons that are produced, while others are involved in
the development or function of the connections between neurons (synapses) where cell-to-cell communication takes place, or of the cell projections (dendrites) that carry signals received at the synapses to the body of the neuron. Many affect development by controlling (regulating) the activity of other genes or proteins.

The specific ways that changes in these and other genes relate to the development of ASD are unknown. However, studies indicate that during brain development, some people with ASD have more neurons than normal and overgrowth in parts of the outer surface of the brain (the cortex). In addition, there are often patchy areas where the normal structure of the layers of the cortex is disturbed. Normally the cortex has six layers, which are established during development before birth, and each layer has specialized neurons and different patterns of neural connection. The neuron and brain abnormalities occur in the frontal and temporal lobes of the cortex, which are involved in emotions, social behavior, and language. These abnormalities are thought to underlie the differences in socialization, communication, and cognitive functioning characteristic of ASD.

Inheritance Pattern
ASD has a tendency to run in families, but the inheritance pattern is usually unknown. People with gene changes associated with ASD generally inherit an increased risk of developing the condition, rather than the condition itself. When ASD is a feature of another genetic syndrome, it can be passed on according to the inheritance pattern of that syndrome.

Other Names for This Condition
- ASD
- autistic continuum
- pervasive developmental disorder

Diagnosis & Management
Genetic Testing Information
- What is genetic testing?
  /primer/testing/genetictesting

Research Studies from ClinicalTrials.gov
- ClinicalTrials.gov https://clinicaltrials.gov/ct2/results?cond=%22autism+spectrum+disorder%22
Other Diagnosis and Management Resources

- Academy of Nutrition and Dietetics: Autism Spectrum Disorders and Diet

- Agency for Healthcare Research and Quality: Therapies for Children With Autism Spectrum Disorder
  https://effectivehealthcare.ahrq.gov/topics/autism-update/consumer/

- American Occupational Therapy Association: Occupational Therapy’s Role with Autism

- Job Accommodation Network: Accommodation and Compliance Series: Employees with Autism Spectrum
  https://askjan.org/publications/Disability-Downloads.cfm?pubid=206344

- Simons Foundation Autism Research Initiative (SFARI)
  https://www.sfari.org/

Additional Information & Resources

Health Information from MedlinePlus

- Encyclopedia: Autism Spectrum Disorder
  https://medlineplus.gov/ency/article/001526.htm

- Health Topic: Autism Spectrum Disorder
  https://medlineplus.gov/autismspectrumdisorder.html

Genetic and Rare Diseases Information Center

- Autism spectrum disorder

Additional NIH Resources

- Eunice Kennedy Shriver National Institute of Child Health and Human Development: Autism Spectrum Disorder
  https://www.nichd.nih.gov/health/topics/autism/conditioninfo

- National Human Genome Research Institute: Learning About Autism
  https://www.genome.gov/Genetic-Diseases/Autism

- National Institute of Mental Health: Autism Spectrum Disorder
• National Institute of Neurological Disorders and Stroke: Autism Spectrum Disorder Fact Sheet
  https://www.ninds.nih.gov/Disorders/Patient-Caregiver-Education/Fact-Sheets/Autism-Spectrum-Disorder-Fact-Sheet

• National Institute on Deafness and Other Communication Disorders: Autism Spectrum Disorder Communication Problems in Children

Educational Resources

• American Psychiatric Association: What is Autism Spectrum Disorder?
  https://www.psychiatry.org/patients-families/autism/what-is-autism-spectrum-disorder

• American Psychological Association: Autism
  https://www.apa.org/topics/autism/

• Centers for Disease Control and Prevention: Autism Spectrum Disorder (ASD)
  https://www.cdc.gov/ncbddd/autism/index.html

• HelpGuide.org: Autism Spectrum Disorders

• MalaCards: autism spectrum disorder
  https://www.malacards.org/card/autism_spectrum_disorder

• Merck Manual Consumer Version: Autism Spectrum Disorders

• National Professional Development Center on Autism Spectrum Disorder
  https://autismpdc.fpg.unc.edu/national-professional-development-center-autism-spectrum-disorder

• Spectrum: Autism Research News
  https://www.spectrumnews.org/

• TeensHealth: Autism

• World Health Organization: Autism Spectrum Disorders
Patient Support and Advocacy Resources

- American Speech-Language-Hearing Association: Autism
  https://www.asha.org/content.aspx?id=14091
- Autism Research Institute
  https://www.autism.org/
- Autism Science Foundation
  https://autismsciencefoundation.org/
- Autism Society
  https://www.autism-society.org/
- Autism Speaks
  https://www.autismspeaks.org/
- Interactive Autism Network
  https://iancommunity.org/
- University of Kansas Support Resources: Autism
  http://www.kumc.edu/gec/support/autism.html

Scientific Articles on PubMed

- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28Autism+Spectrum+Disorder%5BMAJR%5D%29+AND+%28autism+spectrum+disorder%5BTDI%5D%29+AND+review%5Bpt%5D+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+720+days%22%5Bdp%5D

Catalog of Genes and Diseases from OMIM

- AUTISM
  http://omim.org/entry/209850

Sources for This Summary

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/27911742
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  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/22068992

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  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/24670167
  Free article on PubMed Central: https://www.ncbi.nlm.nih.gov/pmc/articles/PMC4499461/

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/28153685

  Citation on PubMed: https://www.ncbi.nlm.nih.gov/pubmed/28260791

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