

CHARGE Syndrome (or Association)

First Description

First described as associated features independently by Hall (1979) and Hittner, Hirsch, Kreh, & Rudolph (1979). Called CHARGE in 1981 (Pagon, Graham, Zonana, & Yong).

Genetics/aetiology

In 2004 mutations in the CHD7 gene (chromodomain helicase DNA-binding protein 7), locus 8q12.1, were identified as a primary cause of CHARGE (Vissers, et al.). Its genomic structure spans 188 kb and encompasses 38 exons, the first of which is noncoding. The encoded CHD7 protein is 2,997 amino acids in length and contains several domains that are characteristic for its protein family, the chromodomain helicase DNA binding proteins, an ATP-dependent chromatin remodeling protein family. Subsequent research has found a mutation in this gene in 65-75% of cases, but in >90% of "typical" CHARGE patients based on clinical diagnosis.

Incidence/prevalence

While most sources estimate incidence at 1/10,000 births, a comprehensive study of individuals in the Netherlands found between 1:15,000 and 1:17,000 (Janssen et al., 2012).

Physical phenotype

The acronym was suggested by Pagon and colleagues (1981) based on common features: C - coloboma of the eye (missing part of iris and/or retina); H – heart defects; A – atresia of the choanae (bony or membranous blocking of nasal passage); R – restrictions of growth and/or development; G – genitourinary anomalies; E – ear anomalies and/or deafness. While all six features may appear, there is wide variation in both occurrence and severity. This has led to a more refined diagnostic system (Blake et al, 1998) with four major features (coloboma, choanal atresia, cranial nerve dysfunction, characteristic CHARGE ear) and a number of minor anomalies. Diagnosis is based on 3 major or 2 major and 2 minor. Other diagnostic systems have since been proposed (e.g., Hale, 2016). Diagnosis is difficult because there is great variability in presence and severity of the features.

CHARGE has become the most common cause of congenital deaf-blindness (after "other" and "unknown"). These difficulties are present due to missing or malformed semi-circular canals and otoliths. Swallowing difficulties are common due to cranial nerve dysfunction. Other problems include gastroesophageal reflux, airway difficulties, chronic otitis media, sinusitis, detached retina, scoliosis, chronic constipation with megacolon, and sleep disturbances.

Cranial nerve anomalies are common in CHARGE with as many as 92% having symptoms of at least one anomaly. It is speculated that in some cases all 12 cranial nerves might be affected, but the most common are I, V, VII, VIII, and IX/X.

Behavioural and psychiatric characteristics

There is variability in the presence of behavioural difficulties. Behaviour has been described as very goal directed and persistent, socially interested but socially immature, demonstrating high rates of repetitive behaviour and sensation seeking, including problems with self-regulation and transitioning between activities. The behaviours have led to diagnoses of autism, attention deficit/hyperactivity disorder, obsessive-compulsive disorder, and tic disorder. It has been proposed that pain, sensory issues, and anxiety, which produce problems with self-regulation, are major sources of the behavior (Hartshorne, Stratton, Brown, Madavan-Brown, & Schmittel, 2017).

Neuropsychological characteristics

There is considerable variability, with some children very developmentally delayed, and others graduating college. Adaptive behaviour tends to be in the low normal range, with little change over four years. Deficits in executive functions have been identified, particularly difficulties with inhibition, shifting focus, and self-monitoring. Due at least in part to sensory impairments, language development and communication may be severely delayed.

Useful websites/associations for more information

- www.chargesyndrome.org US CHARGE foundation
- www.chargesyndrome.org.uk UK support group
- www.chargesyndrome.org.nz Australasian support group
- www.cmich.edu/colleges/class/Psychology/charge CHARGE research lab focused on behaviour

References

1. Blake K.D., Davenport S.L., Hall B.D., Hefner M.A., Pagon R.A., Williams M.S., et al. (1998) CHARGE association: an update and review for the primary pediatrician. Clin Pediatr 37(3),159-73.

2. Hale, C. L., Niederriter, A. N., Green, G. E., Martin, D. M. (2016). Atypical phenotypes associated with pathogenic CHD7 variants and a proposal for broadening CHARGE syndrome clinical diagnostic criteria. Am J Med Genet A, 170, 344–354.

3. Hall B. (1979) Choanal atresia and associated multiple anomalies. J Pediatrics 95, 395-98.

4. Hartshorne T.S., Hefner M.S., Davenport S.L.H., Thelin J.W. (Eds.). CHARGE Syndrome. In press. San Diego: Plural Publishing.

5. Hartshorne, T. S., Stratton, K. K., Brown, D. M., Madhavan-Brown, S. A., Schmittel, M. C. (2017). Behavior in CHARGE syndrome. Am J Med Genet C, 175, 431–438.

6. Hittner H.M., Hirsch N.J., Kreh G.M., Rudolph A.J. (1979) Colobomatous microphthalmia, heart disease, hearing loss and mental retardation: a syndrome. J Pediatr Ophthalmol Strabismus 16,122-128.

7. Janssen, N.; Bergman, J. E. H.; Swertz, M. A.; Tranebjaerg, L.; Lodahl, M.; Schoots, J.; Hofstra, R. M. W.; van Ravenswaaij-Arts, C. M. A.; Hoefsloot, L. H. (2012). Mutation update on the CHD7 gene involved in CHARGE syndrome. Human Mutation, 33(8), 1149-1160.

8. Pagon R.A., Zonana J., Graham J.M. (1982). CHARGE association. Pediatrics 70(5), 827-8.

9. Vissers L.E., van Ravenswaaij C.M., Admiraal R., Hurst J.A., de Vries B.B., Janssen I.M., et al. (2004) Mutations in a new member of the chromodomain gene family cause CHARGE syndrome. Nat Genet 36(9), 955-7.

9. Wachtel L.E., Hartshorne T.S., & Dailor A.N. (2007) Psychiatric diagnoses and psychotropic medications in CHARGE syndrome: A pediatric survey. J Dev Phys Disabil 19, 471-483.

Timothy S. Hartshorne, May, 2019

Copyright © 2019 T. Hartshorne

The SSBP hopes that readers will find the syndrome information sheets useful. They represent the views of the authors who kindly prepared them, and not necessarily those of the SSBP.