

# **Dementia with Lewy bodies**

# **Description**

Dementia with Lewy bodies is a nervous system disorder characterized by a decline in intellectual function (dementia), a group of movement problems known as parkinsonism, visual hallucinations, sudden changes (fluctuations) in behavior and intellectual ability, and acting out dreams while asleep (REM sleep behavior disorder). This condition typically affects older adults, most often developing between ages 50 and 85. The life expectancy of individuals with dementia with Lewy bodies varies; people typically survive about 5 to 7 years after they are diagnosed.

REM sleep behavior disorder may be the first sign of dementia with Lewy bodies. It can occur years before other symptoms appear. Individuals with REM sleep behavior disorder act out their dreams, talking and moving in their sleep when they should be still. This behavior becomes less pronounced as dementia with Lewy bodies worsens and additional features develop.

Dementia is often the second major feature to develop in dementia with Lewy bodies. Initially, this intellectual decline may be mild or seem to come and go. In this condition, dementia often leads to impaired ability to perform visual-spatial tasks such as assembling puzzles. Affected individuals may also have poor problem-solving skills (executive functioning), speech difficulties, and reduced inhibitions. Problems with memory typically do not occur until later.

Most people with dementia with Lewy bodies experience visual hallucinations, which often involve people or animals. Fluctuations in behavior and thought processes (cognition) include sudden changes in attention, unintelligible speech, and brief episodes of altered consciousness that may appear as staring spells.

Parkinsonism is usually the last major feature to develop in people with dementia with Lewy bodies, although it can appear earlier in some individuals. The movement problems typically include tremors, rigidity, unusually slow movement (bradykinesia), and impaired balance and coordination (postural instability). Affected individuals may require walking aids or wheelchair assistance over time.

Individuals with dementia with Lewy bodies may also experience a sharp drop in blood pressure upon standing (orthostatic hypotension), fainting episodes (syncope), reduced sense of smell, increased saliva production and drooling, difficulty controlling the flow of urine (incontinence), or constipation.

## Frequency

Dementia with Lewy bodies is estimated to affect 1.4 million people in the United States. It accounts for about 5 percent of all dementia cases in older individuals and is the second most common dementia after Alzheimer's disease.

#### Causes

Variants (also called mutations) in genes known as *SNCA* and *SNCB* can cause dementia with Lewy bodies. The *SNCA* and *SNCB* genes provide instructions for making proteins, called alpha-synuclein and beta-synuclein, respectively, that are found primarily in the brain. Alpha-synuclein plays a role in communication between nerve cells (neurons), helping to regulate the release of chemical messengers (neurotransmitters). Beta-synuclein is likely involved in a process that allows neurons to change and adapt over time, which is necessary for learning and memory. Beta-synuclein may also prevent harmful accumulation of alpha-synuclein in neurons.

Variants in another gene called *GBA1* or a certain version of a gene called *APOE* increase the risk of developing the condition, but are not a direct cause. The enzyme produced from the *GBA1* gene is found throughout the body in cell structures called lysosomes that digest and recycle proteins and other materials that are no longer needed. The *APOE* gene provides instructions for making a protein, called apolipoprotein E, which packages cholesterol and other fats and carries them through the bloodstream.

Changes in these four genes can lead to the formation of Lewy bodies, which are clusters of alpha-synuclein protein. *SNCA* gene variants result in misshapen alpha-synuclein proteins that cluster together (aggregate). *SNCB* gene variants lead to the production of an altered beta-synuclein protein that allows accumulation of alpha-synuclein. *GBA1* gene variants are thought to disrupt the normal function of lysosomes. Research suggests that malfunctioning lysosomes impair the breakdown of alpha-synuclein, increasing the risk of its accumulation and the formation of Lewy bodies. There are multiple versions of the *APOE* gene, one version, called the e4 allele, seems to increase an individual's risk for developing dementia with Lewy bodies, although the mechanism is unclear. It is thought that the apolipoprotein E produced from the e4 allele of the *APOE* gene may disrupt the transport of alpha-synuclein into and out of cells, leading to its accumulation.

In dementia with Lewy bodies, alpha-synuclein clusters accumulate inside and outside of neurons throughout the brain where they impair cell function and ultimately cause cell death. Neurons that produce the neurotransmitter dopamine seem to be particularly vulnerable to Lewy bodies. Dopamine has many important functions, including playing complex roles in cognition, motivation, behavior, and control of movement. Over time, the loss of dopamine-producing neurons can increasingly impair intellectual and motor function and the regulation of emotions, resulting in the signs and symptoms of dementia with Lewy bodies.

Learn more about the genes associated with Dementia with Lewy bodies

- APOE
- GBA1
- SNCA
- SNCB

## Inheritance

When dementia with Lewy bodies is caused by *SNCA* or *SNCB* gene variants, it is inherited in an autosomal dominant pattern, which means one copy of the altered gene in each cell is sufficient to cause the disorder. In these cases, an affected person usually has one parent with the condition.

People with a variant in one copy of the *GBA1* gene or one copy of the *APOE* e4 allele inherit an increased risk of developing dementia with Lewy bodies, not the condition itself. Some people with these genetic variants never develop dementia with Lewy bodies. This increased risk is inherited in an autosomal dominant pattern.

#### Other Names for This Condition

- Dementia of the Lewy body type
- Dementia, Lewy body
- Diffuse Lewy body disease
- DLB
- LBD
- Lewy body dementia
- Lewy body disease

#### Additional Information & Resources

## **Genetic Testing Information**

Genetic Testing Registry: Lewy body dementia (https://www.ncbi.nlm.nih.gov/gtr/conditions/C0752347/)

## Patient Support and Advocacy Resources

National Organization for Rare Disorders (NORD) (https://rarediseases.org/)

## Clinical Trials

ClinicalTrials.gov (https://clinicaltrials.gov/search?cond=%22Dementia with Lewy bo

dies%22)

# Catalog of Genes and Diseases from OMIM

DEMENTIA, LEWY BODY; DLB (https://omim.org/entry/127750)

## Scientific Articles on PubMed

PubMed (https://pubmed.ncbi.nlm.nih.gov/?term=%28dementia+with+Lewy+bodies %5BTI%5D%29+AND+review%5Bpt%5D+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1800+days%22%5Bdp%5D)

## References

- Dickson DW, Heckman MG, Murray ME, Soto AI, Walton RL, Diehl NN, van GerpenJA, Uitti RJ, Wszolek ZK, Ertekin-Taner N, Knopman DS, Petersen RC, Graff-RadfordNR, Boeve BF, Bu G, Ferman TJ, Ross OA. APOE epsilon4 is associated withseverity of Lewy body pathology independent of Alzheimer pathology. Neurology. 2018 Sep 18;91(12):e1182-e1195. doi: 10.1212/WNL.0000000000006212. Epub 2018 Aug24. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/30143564) or Free article on PubMed Central (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC616 1556/)
- Ferman TJ, Boeve BF, Smith GE, Lin SC, Silber MH, Pedraza O, Wszolek Z, Graff-Radford NR, Uitti R, Van Gerpen J, Pao W, Knopman D, Pankratz VS, KantarciK, Boot B, Parisi JE, Dugger BN, Fujishiro H, Petersen RC, Dickson DW. Inclusionof RBD improves the diagnostic classification of dementia with Lewy bodies. Neurology. 2011 Aug 30;77(9):875-82. doi: 10.1212/WNL.0b013e31822c9148. Epub 2011Aug 17. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/21849645) or Free article on PubMed Central (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3162640/)
- Hogan DB, Fiest KM, Roberts JI, Maxwell CJ, Dykeman J, Pringsheim T, SteevesT, Smith EE, Pearson D, Jette N. The Prevalence and Incidence of Dementia withLewy Bodies: a Systematic Review. Can J Neurol Sci. 2016 Apr;43 Suppl 1:S83-95.doi: 10. 1017/cjn.2016.2. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/27307129)
- McKeith IG, Boeve BF, Dickson DW, Halliday G, Taylor JP, Weintraub D, AarslandD, Galvin J, Attems J, Ballard CG, Bayston A, Beach TG, Blanc F, Bohnen N,Bonanni L, Bras J, Brundin P, Burn D, Chen-Plotkin A, Duda JE, El-Agnaf O, Feldman H, Ferman TJ, Ffytche D, Fujishiro H, Galasko D, Goldman JG, Gomperts SN,Graff-Radford NR, Honig LS, Iranzo A, Kantarci K, Kaufer D, Kukull W, Lee VMY, Leverenz JB, Lewis S, Lippa C, Lunde A, Masellis M, Masliah E, McLean P, Mollenhauer B, Montine TJ, Moreno E, Mori E, Murray M, O'Brien JT, Orimo S, Postuma RB, Ramaswamy S, Ross OA, Salmon DP, Singleton A, Taylor A, Thomas A,Tiraboschi P, Toledo JB, Trojanowski JQ, Tsuang D, Walker Z, Yamada M, Kosaka K.Diagnosis and management of dementia with Lewy bodies: Fourth consensus report of the DLB Consortium. Neurology. 2017 Jul 4;89(1):88-100. doi:10.

- 1212/WNL.0000000000004058. Epub 2017 Jun 7. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/28592453) or Free article on PubMed Central (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5496518/)
- Morra LF, Donovick PJ. Clinical presentation and differential diagnosis ofdementia with Lewy bodies: a review. Int J Geriatr Psychiatry. 2014Jun;29(6):569-76. doi: 10. 1002/gps.4039. Epub 2013 Oct 21. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/24150834)
- Nalls MA, Duran R, Lopez G, Kurzawa-Akanbi M, McKeith IG, Chinnery PF, MorrisCM, Theuns J, Crosiers D, Cras P, Engelborghs S, De Deyn PP, Van Broeckhoven C,Mann DM, Snowden J, Pickering-Brown S, Halliwell N, Davidson Y, Gibbons L, HarrisJ, Sheerin UM, Bras J, Hardy J, Clark L, Marder K, Honig LS, Berg D, Maetzler W,Brockmann K, Gasser T, Novellino F, Quattrone A, Annesi G, De Marco EV, RogaevaE, Masellis M, Black SE, Bilbao JM, Foroud T, Ghetti B, Nichols WC, Pankratz N,Halliday G, Lesage S, Klebe S, Durr A, Duyckaerts C, Brice A, Giasson BI,Trojanowski JQ, Hurtig HI, Tayebi N, Landazabal C, Knight MA, Keller M, SingletonAB, Wolfsberg TG, Sidransky E. A multicenter study of glucocerebrosidasemutations in dementia with Lewy bodies. JAMA Neurol. 2013 Jun; 70(6):727-35. doi:10.1001/jamaneurol.2013.1925. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/23588557) or Free article on PubMed Central (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3841974/)
- Pietrzak M, Papp A, Curtis A, Handelman SK, Kataki M, Scharre DW, Rempala G, Sadee W. Gene expression profiling of brain samples from patients with Lewy bodydementia. Biochem Biophys Res Commun. 2016 Oct 28;479(4):875-880. doi:10. 1016/j.bbrc.2016.09.114. Epub 2016 Sep 22. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/27666482) or Free article on PubMed Central (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC5079284/)
- Tsuang D, Leverenz JB, Lopez OL, Hamilton RL, Bennett DA, Schneider JA, Buchman AS, Larson EB, Crane PK, Kaye JA, Kramer P, Woltjer R, Kukull W, NelsonPT, Jicha GA, Neltner JH, Galasko D, Masliah E, Trojanowski JQ, Schellenberg GD, Yearout D, Huston H, Fritts-Penniman A, Mata IF, Wan JY, Edwards KL, Montine TJ, Zabetian CP. GBA mutations increase risk for Lewy body disease with and withoutAlzheimer disease pathology. Neurology. 2012 Nov 6;79(19):1944-50. doi:10.1212/WNL.0b013e3182735e9a. Epub 2012 Oct 3. Citation on PubMed (https://pubmed.ncbi.nlm.nih.gov/23035075) or Free article on PubMed Central (https://www.ncbi.nlm.nih.gov/pmc/articles/PMC3484986/)
- Vergouw LJM, van Steenoven I, van de Berg WDJ, Teunissen CE, van Swieten JC, Bonifati V, Lemstra AW, de Jong FJ. An update on the genetics of dementia withLewy bodies. Parkinsonism Relat Disord. 2017 Oct;43:1-8. doi:10.1016/j. parkreldis.2017.07.009. Epub 2017 Jul 13. Citation on PubMed (https://pubmed.ncbi. nlm.nih.gov/28734699)

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