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Parkinson Disease Overview

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Summary

The purpose of this overview is to increase the awareness of clinicians regarding the genetics of Parkinson disease and related genetic counseling issues.

The following are the goals of this overview.

Goal 1

Describe the clinical characteristics of Parkinson disease.

Goal 2

Review the causes of Parkinson disease.

Goal 3

Provide an evaluation strategy to identify the genetic cause of Parkinson disease in a proband.

Goal 4

Inform genetic counseling for family members of an individual with Parkinson disease.

Goal 5

Provide information on targeted therapeutic clinical trials.

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1. Clinical Characteristics of Parkinson Disease

Clinical Manifestations of Parkinson Disease

Parkinson disease, a neurodegenerative disorder, is characterized by rest tremor, muscle rigidity, slowed movement (bradykinesia), and often postural instability. Onset is typically unilateral and may include other abnormal movements such as postural or action tremor as well as limb dystonia. Common associated non-motor findings include insomnia, depression, anxiety, rapid-eye-movement (REM) sleep behavior disorder, fatigue, constipation, dysautonomia, and hyposmia. As part of a prodromal phase, non-motor features may predate formal diagnosis of Parkinson disease by years. With disease progression, bilateral manifestations, balance disturbances, and other complications such as levodopa-induced dyskinesia and motor fluctuations develop. Dementia and/or psychosis occur in 30%-40%.

The diagnosis of Parkinson disease is based on the clinical findings of bradykinesia plus rest tremor and/or rigidity. An important supportive diagnostic feature is an observable response to dopaminergic therapy. Dementia is not an exclusion criterion for diagnosis. While developed for use in a genetic research study, the diagnostic criteria summarized in the following citations are increasingly being used to guide diagnosis in a clinical setting [Postuma et al 2015, Postuma et al 2018].

In instances of diagnostic uncertainty, especially in early stages of disease, neuroimaging such as dopamine transporter single-photon emission computed tomography (DAT-SPECT) or fluoro-dopa positron emission tomography (PET) can be used to make a more definitive diagnosis of dopaminergic deficit [Ba & Martin 2015, Ibrahim et al 2016]. While these studies can document the presence of dopaminergic dysfunction, they cannot distinguish between Parkinson disease and other degenerative forms of parkinsonism such as those included in Parkinson plus syndromes (see Figure 1). DAT-SPECT is most commonly used to distinguish Parkinson disease from essential tremor; it can also be used to verify that a symptomatic individual has a non-degenerative form of parkinsonism, such as [dopa-responsive dystonia](#). Laboratory or imaging studies such as brain MRI are useful only in excluding alternative diagnoses such as infarction, tumor, or normal-pressure hydrocephalus.

The cardinal histopathologic feature of Parkinson disease is the loss of dopaminergic neurons in the substantia nigra with intracytoplasmic inclusions (alpha-synuclein-containing Lewy bodies) in the remaining intact nigral neurons.

Onset of Parkinson disease is most commonly around age 60 years but can vary. The following are generally accepted descriptions regarding age of onset:

- Juvenile-onset Parkinson disease. Age <20 years
- Early-onset adult Parkinson disease. Age 20-50 years
- Late-onset adult Parkinson disease. Age >50 years

Various terms related to Parkinson disease used in the clinical setting are diagrammed in Figure 1. Parkinsonism is the broadest term that refers to any disorder that includes bradykinesia, rigidity, tremor, and imbalance. "Parkinson plus" includes degenerative conditions in which parkinsonism is a major finding, such as multiple system atrophy, progressive supranuclear palsy, and corticobasal degeneration.

2. Causes of Parkinson Disease

Parkinson disease results from complex interplay of non-genetic and genetic factors. However, genetic factors are increasingly recognized as causative.

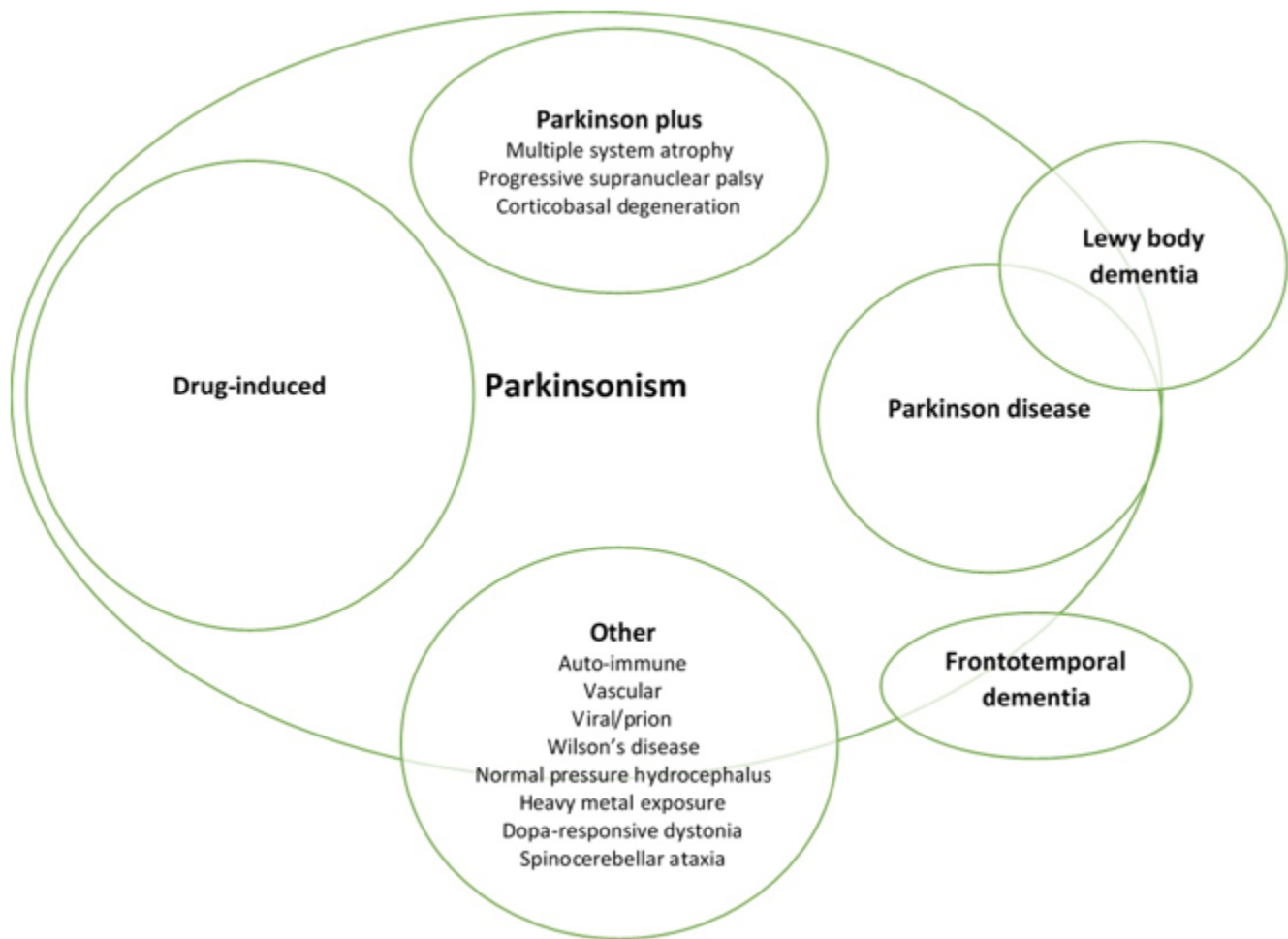


Figure 1. Parkinson disease: related terms in use in the clinical setting

Typically Parkinson disease occurs in only one family member (i.e., a simplex case); less commonly it occurs in several family members (i.e., familial Parkinson disease). Historically, about 15% of individuals with Parkinson disease have a positive family history of PD.

An estimated 5%-10% of all Parkinson disease is attributed to pathogenic variants in single genes (monogenic Parkinson disease) (Table 1, Table 2). In addition to these variants, other genetic and environmental factors – known and unknown – may contribute to overall risk [Kalinderi et al 2016, Kim & Alcalay 2017].

Monogenic Parkinson Disease

Monogenic Parkinson disease can be inherited in an autosomal dominant, autosomal recessive, or, uncommonly, X-linked manner. Age of onset in the proband can be useful in distinguishing autosomal dominant PD (typically age >50 years) from autosomal recessive PD (typically age <40 years) [Kasten et al 2018]; however, age of onset can vary greatly between individuals with the same genetic variant.

Adult-onset Parkinson disease. Table 1 lists the genes associated with early-onset adult PD (age 20-50 years) and late-onset adult PD (age >50 years). Table 1 uses the recommended International Parkinson and Movement Disorder Society Task Force for Nomenclature of Genetic Movement Disorders, in which the phenotype prefix, PARK, is followed by the italicized gene symbol (e.g., PARK-*LRRK2*) [Marras et al 2016].

Table 1. Early-Onset and Late-Onset Adult Parkinson Disease: Monogenic Causes

Gene ¹	PD Designation ²	MOI	% of Adult PD	Comments	GeneReview / Reference / OMIM Entry
<i>GBA1</i> ³ (formerly <i>GBA</i>)	PARK- <i>GBA</i>	AD	3%-7% (20% in AJ ancestry)	<ul style="list-style-type: none"> Onset age may be <50 yrs. Higher likelihood of cognitive impairment, atypical motor findings & severe progression Associated w/dementia w/ Lewy bodies Variable penetrance dependent on age, variant, & ethnicity ⁴ Consider if family history of <i>Gaucher disease</i>. 	OMIM 606463
<i>LRRK2</i>	PARK- <i>LRRK2</i>	AD	1%-2% (13%-30% in AJ ancestry; 41% in African Berber ancestry)	<ul style="list-style-type: none"> Classic manifestations w/ less non-motor involvement Variable penetrance dependent on age, variant, & ethnicity ⁵ 	LRRK2-Related PD
<i>PARK7</i> (<i>DJ1</i>)	PARK- <i>DJ1</i>	AR	Rare	<ul style="list-style-type: none"> Phenotype similar to PARK-<i>Parkin</i> ID &/or seizures occasionally Risk to heterozygotes unknown 	OMIM 606324
<i>PINK1</i>	PARK- <i>PINK1</i>	AR	Rare (3.7% of early-onset adult PD)	<ul style="list-style-type: none"> Phenotype similar to PARK-<i>Parkin</i> Non-motor manifestations incl psychiatric features more common Heterozygotes may have ↑ PD risk. 	PINK1 Type of Young-Onset PD
<i>PRKN</i>	PARK- <i>Parkin</i>	AR	1% (4.6%-10.5% of early-onset adult PD)	<ul style="list-style-type: none"> Slow progression Can have lower-limb dystonia, dyskinesias, hyperreflexia Mild non-motor manifestations Heterozygotes may have ↑ PD risk. 	Parkin Type of Early-Onset PD
<i>SNCA</i>	PARK- <i>SNCA</i>	AD	Rare	<ul style="list-style-type: none"> Onset age may be <50 yrs Cognitive & psychiatric features more likely 	OMIM 168601 , 605543
<i>VPS13C</i>	Not assigned	AR	Rare	Early-onset PD w/very rapid progression; truncating variants cause severe disease.	Puschmann [2017], Reed et al [2019]

Table 1. continued from previous page.

Gene ¹	PD Designation ²	MOI	% of Adult PD	Comments	GeneReview / Reference / OMIM Entry
VPS35	PARK-VPS35	AD	Rare	<ul style="list-style-type: none"> • Classic PD w/tremor • Fewer non-motor manifestations 	VPS35-Related PD

AD = autosomal dominant; AJ = Ashkenazi Jewish; AR = autosomal recessive; ID = intellectual disability; MOI = mode of inheritance; PD = Parkinson disease

See [Parkinson disease: Phenotypic Series](#) to view genes associated with this phenotype in OMIM.

1. Genes are listed in alphabetic order.

2. Nomenclature based on Marras et al [2016]

3. There is some disagreement among researchers as to whether *GBA1* (formerly *GBA*) should be classified as a monogenic disorder or, alternatively, a risk factor – due to its low age-related penetrance. However, of note, some *GBA1* variants (including the fairly common Leu444Pro variant) have estimated penetrance approaching that of *LRRK2* variants [Gan-Or et al 2015, Lee et al 2017].

4. The estimated penetrance of the c.1226A>G (p.Asn409Ser) *GBA1* variant is 6%-14% for all populations. Higher penetrance is reported for severe *GBA1* variants [Gan-Or et al 2015].

5. The estimated penetrance of the c.6055G>A (p.Gly2019Ser) *LRRK2* variant is 25%-43% for all populations [Marder et al 2015, Lee et al 2017]. The estimated penetrance for other *LRRK2* variants may be higher or lower, possibly related to the location of the variant within the gene [Trinh et al 2014].

Juvenile-onset Parkinson disease. Variants in several genes have been associated with juvenile-onset Parkinson disease (i.e., onset age generally <20 years but can vary) (Table 2). Inheritance of juvenile-onset PD is usually autosomal recessive; the clinical presentation often includes additional signs such as dystonia, spasticity, and dementia.

Table 2. Juvenile-Onset Parkinson Disease: Monogenic Causes

Gene ¹	PD Designation ²	MOI	% of Juvenile-Onset PD	Comments	GeneReview / Reference / OMIM Entry
<i>ATP13A2</i>	PARK- <i>ATP13A2</i>	AR	Rare	<ul style="list-style-type: none"> • Triad of spasticity, supranuclear gaze palsy, & dementia • Wide variability • Also referred to as Kufor-Rakeb syndrome or juvenile-onset atypical PD 	Park et al [2015], OMIM 606693
<i>DNAJC6</i>	PARK- <i>DNAJC6</i>	AR	Rare	<p>2 subtypes identified:</p> <ul style="list-style-type: none"> • Slowly progressing, levodopa-responsive parkinsonism; onset in 3rd-4th decade • Rare juvenile onset w/rapid disease progression; atypical features of hyperreflexia, seizures, & ID <p>Variants w/milder effect on protein function may cause adult early-onset PD w/few other symptoms. ³</p>	DNAJC6 Parkinson Disease

Table 2. continued from previous page.

Gene ¹	PD Designation ²	MOI	% of Juvenile-Onset PD	Comments	GeneReview / Reference / OMIM Entry
<i>FBXO7</i>	PARK- <i>FBXO7</i>	AR	Rare	<ul style="list-style-type: none"> Juvenile or early-onset adult, rapidly progressive, may have corticospinal signs Early-onset parkinsonism w/ bradykinesia in some families 	OMIM 260300
<i>PODXL</i>	Not assigned	AR	1 family reported	Levodopa-responsive PD w/no atypical features ⁴	Puschmann [2017]
<i>SLC6A3</i>	DYT/PARK- <i>SLC6A3</i>	AR	Rare	<ul style="list-style-type: none"> Onset in infancy Also referred to as dopamine transporter deficiency syndrome or infantile parkinsonism-dystonia-1 	SLC6A3-Related Dopamine Transporter Deficiency Syndrome
<i>SYNJ1</i> ⁵	PARK- <i>SYNJ1</i>	AR	Rare	<ul style="list-style-type: none"> Variants in SAC1-like domain cause juvenile-onset dystonia w/dyskinesia. 1 family w/early-onset adult PD reported 	OMIM 617389

AR = autosomal recessive; ID = intellectual disability; MOI = mode of inheritance; PD = Parkinson disease

1. Genes are listed in alphabetic order.

2. Nomenclature based on Marras et al [2016]

3. Variants that are deleterious or have a large effect on protein function may lead to more severe, juvenile-onset phenotype, whereas those with a milder effect on protein function may lead to "adult early-onset PD with few or no additional signs and symptoms" [Puschmann 2017].

4. Atypical features are neurologic manifestations including early cognitive decline, severe dystonia, corticospinal signs (hyperreflexia), spasticity, seizures, and oculomotor limitations – in addition to the rigidity, tremor, and bradykinesia that characterize PD.

5. Allelic disorder: AR epilepsy syndrome [Puschmann 2017]

Other suspected genetic risk factors for Parkinson disease. Evidence that variants in other genes may increase the risk for Parkinson disease is less well supported [Kalia & Lang 2015, Ruiz-Martinez et al 2015, Zhang et al 2015, Lee & Hsu 2017, Puschmann 2017, Youn et al 2019]:

- Genetic variants reported in single families include *RIC3*, *TMEM230*, and *UCHL1*.
- Evidence is conflicting for *EIF4G1* and *DNAJC13*.
- Further confirmation is needed to clarify the risk associated with *CHCHD2*, *GIGYF2*, and *HTRA2*.

Additional studies are needed to confirm and clarify the role of variants in these genes in Parkinson disease causation [Kim & Alcalay 2017]. While these genes may appear on Parkinson disease multigene testing panels, it is suggested that they not be included in diagnostic testing because of their limited clinical utility (see Evaluation Strategies).

Susceptibility Genes for Parkinson Disease

In addition to the major known monogenic causes of Parkinson disease (Table 1, Table 2), additional genes and susceptibility loci have been identified through genome-wide association studies (GWAS) and other research. Furthermore, genetic modifiers may influence lifetime risk for PD or clinical aspects of PD such as age of onset and progression of motor and non-motor features [Davis et al 2016]. While these genetic modifiers may contribute in small ways to PD risk [Reed et al 2019], most are not yet clinically applicable and more research is needed to clarify their role in clinical diagnosis and risk assessment.

Variants in such susceptibility genes differ from variants in genes known to cause Parkinson disease: no variant in any of the susceptibility genes "causes" PD. Thus, such genes should not be included in diagnostic testing (see Evaluation Strategies).

Environmental Risk Factors for Parkinson Disease

One of the most important non-genetic factors contributing to Parkinson disease risk is advancing age. Epidemiologic studies have shown possible association of Parkinson disease with environmental factors including pesticide exposure, head injury, rural living, and infectious agents.

Environmental factors that may be associated with a lower risk for PD include tobacco smoking, caffeine consumption, use of nonsteroidal anti-inflammatory drugs (NSAID), high blood urate levels, and physical activity [Kalia & Lang 2015, Ascherio & Schwarzschild 2016].

3. Evaluation Strategies to Identify the Genetic Cause of Parkinson Disease in a Proband

Establishing a specific genetic cause of Parkinson disease:

- Can aid in discussions of causation, recurrence risks, and research eligibility (see Goal 5).
- May provide some information about phenotype including prognosis of a particular monogenic cause of Parkinson disease (Table 1, Table 2).
- Usually involves evaluation of medical and family histories, and molecular genetic testing. Physical examination may be less helpful in suggesting a specific genetic cause because of the overlap of clinical features [Kasten et al 2018, Trinh et al 2019].

Medical history. Age of onset and age at diagnosis may help distinguish autosomal dominant monogenic causes of PD and autosomal recessive monogenic causes of PD (Table 1, Table 2). History of another related diagnosis or overlapping condition may aid in diagnosis and guide genetic testing. For example, if dystonia is a predominant feature of the condition, this may suggest alternative testing. (See [Dystonia Overview](#).)

Physical and mental examination. Typically, findings on a physical or mental examination will not entirely implicate a monogenic cause of Parkinson disease. However, the presence of certain clinical features can provide clues, such as significant cognitive impairment (Table 1, Table 2).

Family history. A three-generation family history should be obtained, with particular attention to any individual with a movement disorder and/or neurodegenerative disorder, with the following also noted:

- Age of onset/diagnosis of disease, which may help distinguish autosomal dominant from autosomal recessive monogenic PD if genetic (Table 1, Table 2)
- Age (or age at death), which will help distinguish unaffected individuals from individuals whose clinical status cannot be determined as they are younger than the typical age of onset for PD
- Details and results of genetic testing of relatives
- Ethnicity, as some populations are more likely to have certain causative pathogenic variants (see Table 1)
- Evaluation by a neurologist (preferably one specializing in movement disorders) of first-degree relatives who have findings concerning for PD

Molecular genetic testing approaches can include a combination of gene-targeted testing (multigene panel or single-gene testing) and less commonly, comprehensive genomic testing (exome sequencing, exome array, or genomic sequencing). Gene-targeted testing requires the clinician to hypothesize which gene(s) are likely involved, whereas genomic testing does not.

- **Single-gene or targeted variant testing.** In a family with an established molecular diagnosis of PD (i.e., a PD-related pathogenic variant has been identified in an affected family member), targeted testing for the familial variant would be indicated. Gene-targeted testing may be appropriate in other limited circumstances (e.g., a family history of [Gaucher disease](#) indicates an increased likelihood of *GBA1* (formerly *GBA*) involvement and an African Berber ancestry indicates a higher likelihood of *LRRK2* involvement). However, single-gene testing is rarely useful and typically not recommended because of the significant overlap in clinical manifestations and age of onset in PD.
- **A multigene panel** that includes some or all of the genes listed in Table 1 and Table 2 is most likely to identify a genetic cause of Parkinson disease while limiting identification of variants in genes that do not explain the underlying phenotype. Note: (1) The genes included in the panel and the diagnostic sensitivity of the testing used for each gene vary by laboratory and are likely to change over time. (2) Some multigene panels may include genes not associated with the condition discussed in this *GeneReview*. Of note, given the rarity of some of the genes associated with Parkinson disease, some panels may not include all the genes included in Table 1 and Table 2. (3) In some laboratories, panel options may include a custom laboratory-designed panel and/or custom phenotype-focused exome analysis that includes genes specified by the clinician. (4) Methods used in a panel may include sequence analysis, deletion/duplication analysis, and/or other non-sequencing-based tests.

For an introduction to multigene panels click [here](#). More detailed information for clinicians ordering genetic tests can be found [here](#).

- **Comprehensive genomic testing** (which does not require the clinician to determine which gene[s] are likely involved) may be considered; however, genomic testing in Parkinson disease is still being evaluated, and yield and utility remain to be determined [Schormair et al 2018, Trinh et al 2019].

For an introduction to comprehensive genomic testing click [here](#). More detailed information for clinicians ordering genomic testing can be found [here](#).

4. Genetic Counseling

Genetic counseling is the process of providing individuals and families with information on the nature, mode(s) of inheritance, and implications of genetic disorders to help them make informed medical and personal decisions. The following section deals with genetic risk assessment and the use of family history and genetic testing to clarify genetic status for family members; it is not meant to address all personal, cultural, or ethical issues that may arise or to substitute for consultation with a genetics professional. —ED.

Typical, Late-Onset Parkinson Disease of Unknown Cause

Typical, late-onset Parkinson disease of unknown cause is generally presumed to be multifactorial in origin (i.e., the result of combined contributions of genetic and environmental factors). Recurrence risk counseling for family members of individuals with presumed multifactorial Parkinson disease (often referred to as "idiopathic" or "sporadic" in the Parkinson disease literature; see Note) and their family members must be empiric and relatively nonspecific. PD is fairly common in the general population and the cumulative lifetime risk (i.e., to age 85 years) of developing PD approaches 4% [Tysnes & Storstein 2017, Deng et al 2018, Licher et al 2019]. Taking into account family history, the cumulative lifetime risk to first-degree relatives (parents, sibs, and offspring) of an individual with typical, late-onset Parkinson disease of unknown cause who represents a simplex case (i.e., the only affected family member) is estimated to be about twofold greater (i.e., 8%) [Gaare et al 2017, Liu et al 2018].

Note: "Idiopathic Parkinson disease" and "sporadic Parkinson disease" are terms used in the Parkinson disease medical literature to describe Parkinson disease of unknown cause diagnosed in an individual with a negative

family history. Because future advances in the understanding of genetic risk factors are likely to identify genetic causes / risk factors for some Parkinson disease currently considered "idiopathic" or "sporadic," these terms are generally not used in this *GeneReview*. Instead, the term "Parkinson disease of unknown cause representing a simplex case" is preferred because "simplex case" refers specifically to an individual with no family history of Parkinson disease without implying presence or absence of a genetic factor or a specific recurrence risk.

Monogenic (Mendelian) Parkinson Disease

Mode of Inheritance

Monogenic Parkinson disease is inherited in an autosomal dominant, autosomal recessive, or X-linked manner. (See [X-Linked Dystonia-Parkinsonism](#) for a review of X-linked inheritance.)

Autosomal Dominant Inheritance – Risk to Family Members

Parkinson disease caused by a pathogenic variant in one of four genes – *GBA1* (formerly *GBA*), *LRRK2*, *SNCA*, or *VPS35* – is inherited in an autosomal dominant manner (Table 1). This section refers only to families having a known Parkinson disease-causing pathogenic variant in one of these four genes.

Parents of a proband

- Almost all individuals diagnosed with autosomal dominant Parkinson disease inherited a pathogenic variant from a parent, who may or not be affected.
- In very rare instances, a proband diagnosed with autosomal dominant Parkinson disease has the disorder as the result of a *de novo* pathogenic variant.
- If the parents of a proband undergo molecular genetic testing for the pathogenic variant found in the proband and the pathogenic variant cannot be detected in the leukocyte DNA of either parent, possible explanations include a *de novo* pathogenic variant in the proband or the theoretic possibility of germline mosaicism in a parent.
- The family history of some individuals diagnosed with autosomal dominant Parkinson disease may appear to be negative because of failure to recognize the disorder in family members, early death of the parent before the onset of symptoms, late onset of the disease in the affected parent, variable expression, or (especially in *GBA1*- and *LRRK2*-related Parkinson disease) reduced penetrance. Therefore, an apparently negative family history can only be confirmed with parental molecular genetic testing.

Sibs of a proband. The risk to the sibs of the proband depends on the genetic status of the proband's parents:

- If a parent of the proband is known to have the pathogenic variant identified in the proband, the risk to each sib of having inherited the variant is 50%. While all sibs who inherit a familial pathogenic variant will be at increased risk for Parkinson disease, not all heterozygous sibs will manifest the disease; both reduced, age-related penetrance and intrafamilial clinical variability are observed in autosomal dominant Parkinson disease.
- If the pathogenic variant found in the proband cannot be detected in the leukocyte DNA of either parent, the recurrence risk to sibs is estimated to be 1% because of the theoretic possibility of parental germline mosaicism [Rahbari et al 2016].
- If the parents have not been tested for the pathogenic variant identified in the proband but are clinically unaffected, sibs are still presumed to be at increased risk for Parkinson disease because of the possibility of reduced penetrance in a heterozygous parent or the theoretic possibility of parental germline mosaicism.

Offspring of a proband

- Each child of an individual with an autosomal dominant Parkinson disease-causing pathogenic variant has a 50% chance of inheriting the pathogenic variant.

- The probability that offspring who inherit a pathogenic variant will manifest Parkinson disease increases with advancing age (see Table 1, penetrance footnotes).

Autosomal Recessive Inheritance – Risk to Family Members

Parkinson disease caused by biallelic pathogenic variants in *ATP13A2*, *DJ1*, *DNAJC6*, *FBXO7*, *PINK1*, *PODXL*, *PRKN*, *SLC6A3*, *SYNJ1*, or *VPS13C* is inherited in an autosomal recessive manner (Table 1, Table 2).

Parents of a proband

- The parents of an individual with an autosomal recessive Parkinson disease are obligate heterozygotes for one Parkinson disease-related pathogenic variant.
- Heterozygotes for a pathogenic variant in *DJ1*, *PINK1*, or *PRKN* (genes typically associated with autosomal recessive adult-onset Parkinson disease) may be at increased risk for adult-onset Parkinson disease (Table 1) [Kim & Alcalay 2017, Reed et al 2019].

Sibs of a proband

- At conception, each sib of an individual with autosomal recessive Parkinson disease has a 25% chance of inheriting both pathogenic variants and being affected, a 50% chance of inheriting one pathogenic variant and being heterozygous, and a 25% chance of inheriting neither pathogenic variant.
- Heterozygotes for a pathogenic variant associated with adult-onset Parkinson disease may be at increased the risk for Parkinson disease (see AR Inheritance, **Parents of a proband**).

Offspring of a proband. The offspring of an individual with an autosomal recessive Parkinson disease are obligate heterozygotes for a pathogenic variant. Heterozygotes for a pathogenic variant associated with adult-onset Parkinson disease may be at increased the risk of Parkinson disease (see AR Inheritance, **Parents of a proband**).

Related Genetic Counseling Issues

Predictive testing for *GBA1*, *LRRK2*, *SNCA*, or *VPS35*-related adult-onset autosomal dominant PD (i.e., testing of asymptomatic, at-risk individuals)

- Predictive testing for relatives at risk for *GBA1*, *LRRK2*, *SNCA*, or *VPS35*-related adult-onset autosomal dominant Parkinson disease is an option once the pathogenic variant has been identified in an affected family member. However, age of onset cannot be accurately predicted by molecular genetic testing because current data suggest that penetrance is both age-related and reduced.
- The following should be discussed in the context of formal genetic counseling prior to testing:
 - Potential consequences of predictive testing (including but not limited to socioeconomic changes and the need for long-term follow up and evaluation arrangements for individuals with a positive test result)
 - The capabilities and limitations of predictive testing.

Predictive testing in minors (i.e., testing of asymptomatic at-risk individuals age <18 years)

- For asymptomatic minors at risk for adult-onset conditions for which early treatment would have no beneficial effect on disease morbidity and mortality, predictive genetic testing is considered inappropriate, primarily because it negates the autonomy of the child with no compelling benefit. Further, concern exists regarding the potential unhealthy adverse effects that such information may have on family dynamics, the risk of discrimination and stigmatization in the future, and the anxiety that such information may cause.
- For more information, see the National Society of Genetic Counselors [position statement](#) on genetic testing of minors for adult-onset conditions and the American Academy of Pediatrics and American

College of Medical Genetics and Genomics [policy statement](#): ethical and policy issues in genetic testing and screening of children.

Note: In a family with an established diagnosis of monogenic Parkinson disease, it is appropriate to consider testing of symptomatic individuals regardless of age.

Direct-to-consumer (DTC) testing is increasingly available. Among the benefits of DTC testing:

- It provides easy, quick, and affordable genetic testing for some Parkinson disease-associated variants; and
- The results of DTC are not inserted into the medical record except when requested by the affected individual.

On the downside, DTC testing does **not** provide:

- Pretest genetic counseling;
- Personalized interpretation of results by a qualified professional; or
- Supportive counseling to ensure that after disclosure of new health information, the emotional needs of the affected individual are identified and a plan is made to address them.

For more information, see [Genetics Home Reference: What are the benefits and risks of direct-to-consumer genetic testing?](#)

Resources

GeneReviews staff has selected the following disease-specific and/or umbrella support organizations and/or registries for the benefit of individuals with this disorder and their families. GeneReviews is not responsible for the information provided by other organizations. For information on selection criteria, click [here](#).

- **American Parkinson Disease Association (APDA)**
Phone: 800-223-2732
Fax: 718-981-4399
Email: apda@apdaparkinson.org
www.apdaparkinson.org
- **Fox Trial Finder**
foxtrialfinder.michaeljfox.org
- **MedlinePlus**
[Parkinson disease](#)
- **Michael J. Fox Foundation for Parkinson's Research**
Phone: 800-708-7644 (toll-free)
Email: info@michaeljfox.org
www.michaeljfox.org
- **National Institute of Neurological Disorders and Stroke (NINDS)**
[Parkinson's Disease Information Page](#)
- **Parkinson's Disease Society (UK)**
United Kingdom
Phone: 0808 800 0303
Email: hello@parkinsons.org.uk
www.parkinsons.org.uk

- **Parkinson's Foundation**
Phone: 800-4PD-INFO (473-4636)
Email: contact@parkinson.org
www.parkinson.org

5. Participation in Targeted Therapeutic Clinical Trials

Several gene-targeted therapies for Parkinson disease are currently in development [Sardi et al 2018]. Individuals may be candidates for a given Parkinson disease clinical trial if they undergo molecular genetic testing and a pathogenic variant in the gene under study is identified.

Clinical trials for individuals with Parkinson disease and *GBA1* (formerly *GBA*) or *LRRK2* pathogenic variants are now ongoing (www.clinicaltrials.gov). Testing for common *GBA1* or *LRRK2* variants is available through direct-to-consumer (DTC) testing or, more formally, through a health care professional. See *GeneReviews Resources for Genetics Professionals* for an overview of DTC testing.

Chapter Notes

Revision History

- 25 July 2019 (bp) Comprehensive update posted live
- 27 February 2014 (me) Comprehensive update posted live
- 16 October 2006 (me) Comprehensive update posted live
- 25 May 2004 (me) Overview posted live
- 12 November 2003 (tmf) Original submission

References

Literature Cited

- Ascherio A, Schwarzschild MA. The epidemiology of Parkinson's disease: risk factors and prevention. *Lancet Neurol*. 2016;15:1257-72. PubMed PMID: 27751556.
- Ba F, Martin WRW. Dopamine transporter imaging as a diagnostic tool for parkinsonism and related disorders in clinical practice. *Parkinsonism Relat Disord* 2015;21:87-94. PubMed PMID: 25487733.
- Davis AA, Andruska KM, Benitez BA, Racette BA, Perlmutter JS, Cruchaga C. Variants in *GBA*, *SNCA*, and *MAPT* influence Parkinson disease risk, age at onset, and progression. *Neurobiol Aging*. 2016;37:209.e1-e7. PubMed PMID: 26601739.
- Deng H, Wang P, Jankovic J. The genetics of Parkinson disease. *Ageing Res Rev*. 2018;42:72-85. PubMed PMID: 29288112.
- Gaare JJ, Skeie GO, Tzoulis C, Larsen JP, Tysnes OB. Familial aggregation of Parkinson's disease may affect progression of motor symptoms and dementia. *Mov Disord*. 2017;32:241-5. PubMed PMID: 27862270.
- Gan-Or Z, Amshalom I, Kilarski LL, Bar-Shira A, Gana-Weisz M, Mirelman A, Marder K, Bressman S, Giladi N, Orr-Urtreger A. Differential effects of severe vs mild *GBA* mutations on Parkinson disease. *Neurology*. 2015;84:880-7. PubMed PMID: 25653295.
- Ibrahim N, Kusmirek J, Struck A, Floberg JM, Perlman SB, Gallagher C, Hall LT. The sensitivity and specificity of F-DOPA PET in a movement disorder clinic. *Am J Nucl Med Mol Imaging*. 2016;6:102-9. PubMed PMID: 27069770.
- Kalia LV, Lang AE. Parkinson's disease. *Lancet*. 2015;386:896-912. PubMed PMID: 25904081.

- Kalinderi K, Bostantjopoulou S, Fidani L. The genetic background of Parkinson's disease: current progress and future prospects. *Acta Neurol Scand.* 2016;134:314-26. PubMed PMID: 26869347.
- Kasten M, Hartmann C, Hampf J, Schaake S, Westenberger A, Vollstedt EJ, Balck A, Domingo A, Vulinovic F, Dulovic M, Zorn I, Madoev H, Zehnle H, Lembeck CM, Schawe L, Reginald J, Huang J, König IR, Bertram L, Marras C, Lohmann K, Lill CM, Klein C. Genotype-phenotype relations for the Parkinson's disease genes Parkin, PINK1, DJ1: MDSGene Systematic Review. *Mov Disord.* 2018;33:730-41. PubMed PMID: 29644727.
- Kim CY, Alcalay RN. Genetic forms of Parkinson's disease. *Semin Neurol.* 2017;37:135-46. PubMed PMID: 28511254.
- Lee AJ, Wang Y, Alcalay RN, Mejia-Santana H, Saunders-Pullman R, Bressman S, Corvol JC, Brice A, Lesage S, Mangone G, Tolosa E, Pont-Sunyer C, Vilas D, Schüle B, Kausar F, Foroud T, Berg D, Brockmann K, Goldwurm S, Siri C, Asselta R, Ruiz-Martinez J, Mondragón E, Marras C, Gbate T, Giladi N, Mirelman A, Marder K; Michael J. Fox LRRK2 Cohort Consortium. Penetrance estimate of LRRK2 p.G2019S mutation in individuals of non-Ashkenazi Jewish ancestry. *Mov Disord.* 2017;32:1432-8. PubMed PMID: 28639421.
- Lee YC, Hsu SD. Familial mutations and post-translational modifications of UCH-L1 in Parkinson's disease and neurodegenerative disorders. *Curr Protein Pept Sci.* 2017;18:733-45. PubMed PMID: 26899237.
- Licher S, Darweesh SKL, Wolters FJ, Fani L, Heshmatollah A, Mutlu U, Koudstaal PJ, Heeringa J, Leening MJG, Ikram MK, Ikram MA. Lifetime risk of common neurological diseases in the elderly population. *J Neurol Neurosurg Psychiatry.* 2019;90:148-56. PubMed PMID: 30279211.
- Liu FC, Lin HT, Kuo CF, Hsieh MY, See LC, Yu HP. Familial aggregation of Parkinson's disease and coaggregation with neuropsychiatric diseases: a population-based cohort study. *Clin Epidemiol.* 2018;10:631-41. PubMed PMID: 29881310.
- Marder K, Wang Y, Alcalay RN, Mejia-Santana H, Tang MX, Lee A, Raymond D, Mirelman A, Saunders-Pullman R, Clark L, Ozelius L, Orr-Urtreger A, Giladi N, Bressman S, et al. Age-specific penetrance of LRRK2 G2019S in the Michael J. Fox Ashkenazi Jewish LRRK2 Consortium. *Neurology.* 2015;85:89-95. PubMed PMID: 26062626.
- Marras C, Lang A, van de Warrenburg BP, Sue CM, Tabrizi SJ, Bertram L, Mercimek-Mahmutoglu S, Ebrahimi-Fakhari D, Warner TT, Durr A, Assmann B, Lohmann K, Kostic V, Klein C. Nomenclature of genetic movement disorders: recommendations of the International Parkinson and Movement Disorder Society task force. *Mov Disord.* 2016;31:436-57. PubMed PMID: 27079681.
- Park J-S, Blair NF, Sue CM. The role of ATP13A2 in Parkinson's disease: Clinical phenotypes and molecular mechanisms. *Mov Disord.* 2015;30:770-9. PubMed PMID: 25900096.
- Postuma RB, Berg D, Stern M, Poewe W, Olanow CW, Oertel W, Obeso J, Marek K, Litvan I, Lang AE, Halliday G, Goetz CG, Gasser T, Dubois B, Chan P, Bloem BR, Adler CH, Deuschl G. MDS clinical diagnostic criteria for Parkinson's disease. *Mov Disord.* 2015;30:1591-601. PubMed PMID: 26474316.
- Postuma RB, Poewe W, Litvan I, Lewis S, Lang AE, Halliday G, Goetz CG, Chan P, Slow E, Seppi K, Schaffer E, Rios-Romenets S, Mi T, Maetzler C, Li Y, Heim B, Bledsoe IO, Berg D. Validation of the MDS clinical diagnostic criteria for Parkinson's disease. *Mov Disord.* 2018;33:1601-8. PubMed PMID: 30145797.
- Puschmann A. New genes causing hereditary Parkinson's disease or parkinsonism. *Curr Neurol Neurosci Rep.* 2017;17:66. PubMed PMID: 28733970.
- Rahbari R, Wuster A, Lindsay SJ, Hardwick RJ, Alexandrov LB, Turki SA, Dominiczak A, Morris A, Porteous D, Smith B, Stratton MR, Hurles ME, et al. Timing, rates, and spectra of human germline mutation. *Nat Genet.* 2016;48:126-33. PubMed PMID: 26656846.
- Reed X, Bandrés-Ciga S, Blauwendraat C, Cookson MR. The role of monogenic genes in idiopathic Parkinson's disease. *Neurobiol Dis.* 2019;124:230-9 PubMed PMID: 30448284.

- Ruiz-Martinez J, Krebs CE, Makarov V, Gorostidi A, Martí-Massó JF, Paisán-Ruiz C. GIGYF2 mutation in late-onset Parkinson's disease with cognitive impairment. *J Hum Genet.* 2015;60:637-40. PubMed PMID: 26134514.
- Sardi SP, Cedarbaum JM, Brundin P. Targeted therapies for Parkinson's disease: from genetics to the clinic. *Mov Disord.* 2018;33:684-96. PubMed PMID: 29704272.
- Schormair B, Kemlink D, Mollenhauer B, Fiala O, Machetanz G, Roth J, Berutti R, Strom TM, Haslinger B, Trenkwalder C, Zahorakova D, Martasek P, Ruzicka E, Winkelmann J. Diagnostic exome sequencing in early-onset Parkinson's disease confirms VPS13C as a rare cause of autosomal-recessive Parkinson's disease. *Clin Genet.* 2018;93:603-12. PubMed PMID: 28862745.
- Trinh J, Guelle I, Farrer MJ. Disease penetrance of late-onset parkinsonism: a meta-analysis. 2014;71:1535-9.
- Trinh J, Lohmann K, Baumann H, Balck A, Borsche M, Brüggemann N, Dure L, Dean M, Volkmann J, Tunc S, Prasuhn J, Pawlack H, Imhoff S, Lill CM, Kasten M, Bauer P, Rolfs A; International Parkinson's Disease Genomics Consortium (IPDGC), Klein C. Utility and implications of exome sequencing in early-onset Parkinson's disease. *Mov Disord.* 2019;34:133-7. PubMed PMID: 30537300.
- Tysnes OB, Storstein A. Epidemiology of Parkinson's disease. *J Neural Transm (Vienna).* 2017;124:901-5. PubMed PMID: 28150045.
- Youn J, Lee C, Oh E, Park J, Kim JS, Kim HT, Cho JW, Park WY, Jang W, Ki CS. Genetic variants of PARK genes in Korean patients with early-onset Parkinson's disease. *Neurobiol Aging.* 2019; 75:224.e9-224.e15 PubMed PMID: 30502028.
- Zhang Y, Sun Q-Y, Yu R-H, Guo J-F, Tang B-S, Yan X-X. The contribution of GIGYF2 to Parkinson's disease: a meta-analysis. *Neurol Sci.* 2015;36:2073-9. PubMed PMID: 26152800.

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