Advances in the understanding of hereditary ataxia
– implications for future patients

Abstract

Introduction: Hereditary ataxias are caused by mutations in a plethora of different genes. Advances in sequencing technologies have led to an exponential increase in novel gene discoveries, highlighted the genetic overlap with other neurological diseases and improved our understanding of genotype-phenotype relationships. Together, these developments allowed the identification of new therapeutic targets that are subsequently making their way into clinical trials.

Areas covered: This review focuses on the shared genetic characteristics and the latest insights into the molecular cause of the most prevalent hereditary ataxias. Furthermore, conventional genetic diagnosis and the gradual implementation of next-generation sequencing (NGS) approaches in clinical practice is discussed. Finally, the latest investigated disease-modifying therapeutic agents are reviewed. A literature search was performed in PubMed and the Cochrane Library. Additional information on previous and on-going trials was obtained from the ClinicalTrials.gov website.

Expert opinion: The implementation of NGS in clinical practice has led to an increase in detected sequence variants of unknown clinical significance. Determining their pathogenicity is an expensive and time-consuming process. However, misinterpretation of these variants can have far-reaching consequences for the patient and their relatives. In accordance with the progresses in genetics, there is a need for the simultaneous definition of novel biomarkers and functional assays that can assist in the interpretation of genetic tests. Moreover, the identification of biomarkers that are relevant to specific diseases has the potential to improve clinical trial design.
**Article highlights**

- Hereditary ataxias are mono-genetic diseases that show genetic and clinical heterogeneity with marked phenotypical overlap.
- The most prevalent dominant, recessive and X-linked ataxias are due to a trinucleotide repeat expansion in their respective genes.
- Depending on the disease-causing gene, the length and configuration of these trinucleotide repeat expansions variably influences age of onset, intergenerational instability and phenotypic presentation.
- Advances in sequencing technologies, such as next-generation sequencing, have revolutionised the genetic landscape of ataxias and are increasingly implemented in clinical practice.
- Disease-modifying treatments target the genetic cause of the disease directly or focus on shared pathological downstream mechanisms. Biomarkers could prove useful in monitoring therapeutic response.
1. Introduction

The term hereditary ataxia encompasses a clinically and genetically heterogeneous group of disorders. They share a progressive incoordination of motor activity affecting gait, extraocular movements and speech.\textsuperscript{1} Underlying these conditions is a well-described genetic association with autosomal dominant, autosomal recessive, X-linked or mitochondrial transmission.\textsuperscript{2} Even though marked variability in intra- and interfamilial phenotypical presentation were identified early in their description, and different genetic backgrounds suspected, the number of distinct genetic causes has only recently begun to be unravelled.\textsuperscript{3} The first milestones of gene discovery in hereditary ataxia were made in the early 1990s with the discovery of pathogenic repetitive trinucleotide repeat (TR) expansions within the \textit{ATXN1} gene in Spinocerebellar ataxia type 1, and the \textit{FXN} gene in Friedreich’s ataxia (FRDA).\textsuperscript{4,5} Both causative genes were discovered by linkage analysis and subsequent Sanger sequencing.

In the last two decades, conventional sequencing techniques have gradually been replaced by next-generation sequencing (NGS). These new approaches are capable of reading huge amounts of genetic sequences in parallel.\textsuperscript{6} NGS has undoubtedly revolutionised the field of ataxia by broadening our knowledge about the phenotypic spectrum of known ataxia associated genes and exponentially increasing the number of novel gene discoveries (Figure 1).\textsuperscript{7,8} To date, over 40 different disease-causing gene loci have been mapped in autosomal dominant cerebellar ataxias (ADCA) (Table 1). These genetic loci have been labelled spinocerebellar ataxia (SCA) and numbered in chronological order of their discovery. The group of eight known episodic ataxias and Dentatorubral-pallidoluysian atrophy (DRPLA) are an exception to this nomenclature. In addition, an equally high number of causative genes has been identified in recessive ataxias (Table 2). However, their classification remains a challenge due to a lack of uniform nomenclature and the vast number of recessive neurological diseases presenting
with symptoms of ataxia. This is reflected by the marked variability in disorders that are included to the group of hereditary recessive ataxias in literature.⁹ Although numerous, both dominant and recessive ataxias belong to the group of rare diseases. The average prevalence is 2.7 in 100,000 and 3.3 in 100,000, respectively.¹⁰ An awareness of the great variation in prevalence of distinctive ataxic disorders that exists in different ethnical groups and geographic regions may facilitate the choice of genetic tests in clinical practice.

The observed phenotypical overlap between hereditary ataxias is surprising considering the broad structural and functional differences in the disease-causing proteins. Physiological functions of disease-causing genes in SCAs include ion transport, deubiquitination, dephosphorylation and phosphorylation, transcriptional regulation and translational elongation.¹¹ The nuclear gene \( \text{FXN} \) that is mutated in FRDA encodes for a well-studied, ubiquitously expressed mitochondrial protein that is crucial for the biogenesis of iron-sulphur clusters and haem.⁵,¹² The literature on pathological downstream mechanisms is rapidly growing, however their individual contribution to neurodegeneration remains unclear. The fact that this genetic heterogeneity results in such a similar clinical picture drives the search for shared molecular pathways that could potentially have an implication beyond the field of progressive ataxias.¹³ A cross-disciplinary cooperation between clinicians, cell biologist and physiologist seems crucial in the definition of new therapeutic targets. Based on the postulated toxic gain-of-function hypothesis in the polyglutamine ataxias and animated by the breakthrough clinical success of antisense oligonucleotide (ASO) treatments in spinal muscular atrophy (SMA) and positive preliminary results in a Phase II clinical trial in Huntington disease (HD), different genetic approaches with the common goal of decreasing toxic protein levels are currently under investigation for SCAs.¹⁴⁻¹⁶ On the other hand, to combat the pathogenic loss-of-function, restoration of physiological frataxin protein levels appears to be a viable
approach for FRDA patients.\textsuperscript{17} In addition, new agents aimed at improving the mitochondrial defects in FRDA are currently being assessed in clinical trials.\textsuperscript{18}

This review will focus on the most prevalent dominant, recessive and X-linked ataxias in adults that are intriguingly all associated with pathological repetitive DNA sequences. We aim to outline the progress made in the molecular understanding of disease pathogenesis and the techniques that have made these advances possible. Lastly, we comment on the most recent developments in preclinical and clinical trials.

2. Repeat expansions as a shared theme in the most prevalent ataxias

2.1 Repeat expansions in autosomal dominant ataxia

ADCA are clinically characterized by a typically late-onset, progressive ataxia, but can present at any age. This incoordination is caused primarily by the degeneration of the cerebellum, which is variably associated with the involvement of other central and peripheral nervous system regions.\textsuperscript{19–21} In the individual patient, cerebellar ataxia is often accompanied by a variety of additional neurological symptoms.\textsuperscript{22}

A plethora of causative mutations have been described in dominant inherited ataxias, including: conventional mutations (SCA5, SCA11, SCA13, SCA14, SCA19/22, SCA23, SCA26, SCA27, SCA28, SCA19, SCA35); rearrangements (SCA15, SCA16, SCA20); as well as expansions of variable length in intronic (SCA8, SCA10, SCA12, SCA31, SCA36) and exonic regions (SCA1, SCA2, SCA3, SAC6, SCA7, SCA17, DRPLA).\textsuperscript{23} The latter encompass the most common and best studied dominantly inherited ataxias. Together with other late-onset neurodegenerative diseases, namely HD and spinal-bulbar muscular atrophy, they form the group of nine known polyglutamine (polyQ) diseases. These disorders share an exonic (CAG)\textsubscript{n}
TR expansion in their respective disease genes. Simple repetitive elements are considered pathological if the number of triplets are greater than the number found in wild-type alleles. Once above a critical threshold, the excessive polyQ stretches in the translated proteins promote cell specific degeneration associated with a toxic gain-of-function at the protein and mRNA level, which leads to the pathological hallmark of these disorders, cellular aggregation.

In addition to pathogenic exonic TR expansions, repetitive DNA elements have more recently been discovered in untranslated regions of ataxia associated genes. SCA8, for example, belongs to the group of pure adult-onset cerebellar ataxias. It is caused by a unique overlapping intronic CAG-CTG repeat in two complementary genes in the SCA8 locus. The CTG expansion in the ATXN8OS gene is transcribed into mRNA containing an expanded CUG repeat, while the CAG expansion in the ATXN8 gene encodes a polyQ protein. This polyalanine was one of the initial proteins described in repeat associated non-ATG (RAN) translation. RAN translation occurs in different frames and causes accumulation of homopolymeric toxic SCA8-polyalanine peptides in vitro. This mechanism has since been described in other trinucleotide repeat diseases such SCA2, SCA7, HD, myotonic dystrophy, c9orf72-mediated amyotrophic lateral sclerosis (ALS) and frontotemporal dementia.

2.2 Repeat expansions in autosomal recessive ataxia
FRDA is the most prevalent recessive ataxia and shares the causative agent of a TR expansion. At the genetic level, FRDA is caused by a homozygous transcribed, but not translated (GAA)n expansion located in the first intron of the FXN gene in about 95% of patients. The residual individuals are compound heterozygote for a pathological expansion and a point mutation, insertion or deletion. In contrast to ADCA, onset of symptoms is typically around puberty,
family history is often negative except for affected siblings and atrophy is observed mainly in
the dorsal root ganglia, spinocerebellar and pyramidal tracts; with less predominant
involvement of the cerebellum. Two recent studies suggest an underlying developmental
component adding to degeneration of the dorsal columns. Together with axonal peripheral
neuropathy, these neurodegenerative patterns contribute to hallmark clinical features, such as
progressive unsteadiness of gait, bilateral Babinski sign and loss of deep tendon reflexes. Multisystem involvement is reflected by the high incidence of cardiomyopathy and insulin
resistance that require close surveillance and active management. Late-onset presentations
associated with smaller expansions on the shorter FXN allele (GAA1) are increasingly well
characterized, broadening the phenotypic spectrum of the disease.

2.3 Repeat expansions in X-linked ataxia

The most common and only recently recognized adult-onset X-linked ataxia is fragile X-
associated tremor/ataxia syndrome (FXTAS). The typical clinical presentation is characterized
by intention tremor, gait ataxia, and parkinsonism in combination with variable cognitive
decline. FXTAS is caused by what was previously considered as a premutation of 55 to 200
CGG·CCG repeats in the 5’ untranslated region of the FMR1 gene. Similarly to SCA8, RAN
translation results in the transcription of toxic proteins (FXTAS-polyglycine, FXTAS-
polyproline, FXTAS-polyalanine) which have been detected in different brain regions. In
addition, the RNA-mediated sequestering of proteins has been described. These pathological
mechanisms are in contrast to the hypermethylation associated silencing of the FMR1 gene in
fragile X-syndrome.

3. Advances in genetics and genomics
Advances in the field of genome sequencing have led to the identification of novel genes, broadened the phenotypic spectrum of known ataxia associated genes, ameliorated genotype-phenotype relationships and increased accuracy in prognosis. Recently, research in polyQ SCAs and FRDA has focused the influence of expansion configuration and genetic modifiers on age of disease onset (AOO), intergenerational instability and phenotypical presentation.

3.1 Modifiers of AOO

As mentioned above, the first genes identified by linkage analysis in ADCA share a dynamic CAG expansion that is translated into polyQ stretches. The threshold for pathogenicity is specific for each disease. The expansion of these structurally unstable TRs in the germline is referred to as genetic anticipation and is associated with higher levels of mutant proteins.27 This, in turn, translates into clinic in the form of increased disease severity, earlier manifestation of symptoms and reduced life expectancy in successive generations.50

Anticipation accounts for 50 – 80% of variance in AOO; however, it does not explain variances in AOO observed in individuals with expansions of the same size. Once the length of the CAG repeat has been accounted for, the residual AOO variance can be considered as a heritable trait, implying the existence of genetic modifiers.51,52 A recent study identified non-pathological repeat lengths in other CAG-containing disease loci, that act in trans with wildtype alleles (SCA1, SCA6, SCA7), as modifiers of disease onset in 1255 patients from the EUROSCA cohort.53 These findings were partly replicated by other studies in smaller cohorts.54,55 However, overall conflicting evidence exists from the analysis of allelic associations across distinct populations.56–58 Inconclusive reports may reflect ethnical differences, small sample size or differences in methods. Moreover, correlation does not imply causation and functional relationship between genes necessitate conformation in in vitro or in vivo models.
CAG repeat length and the presence of genetic modifiers are still insufficient to fully account for AOO variance. There is an increasing body of literature illustrating the influence of polyQ expansion configuration in the form of silent (CAA) and missense (CAT) insertions on disease onset.\(^5\) In the polyQ tract of SCA1, 98\% of the normal \textit{ATXN1} alleles (19-36 repeats) are interrupted by at least one histidine (CAT) trinucleotide when the tract exceeds 21 repeats.\(^6\) In addition to stabilizing (CAG)n expansions, histidine interruptions appear to prevent or delay phenotypic presentation in individuals with borderline, and even clearly pathogenic repeat lengths.\(^5\) However, up to 11\% of the patients in a SCA1 cohort harboured CAT interruptions in expanded alleles.\(^6\) Together with the observance of the loss of an interruption in maternal transmission, this implies an incomplete, more complex protective effect. In that cohort, the longest uninterrupted or pure repeat tract correlated best with disease onset and severity, highlighting the importance of repeat interruption analysis in SCA1 clinical practice.\(^6\) Interestingly, in two other TR diseases, repeat interruptions were not found to account for the variable age at onset.\(^6,6\)

Most recently, a genome-wide association study found a significant association between AOO and genetic variants in DNA repair pathways.\(^6\) The modifying effects of these variants were examined in a cohort of subjects with HD and polyQ SCAs. The results yielded the most significant association with AOO when grouping all of the polyQ diseases, with rs3512 in \textit{FAN1} and rs1805323 in \textit{PMS2} being the top variants for HD and SCAs.\(^6\)

FRDA is the only recessive TR disease, and thus is generally restricted to one generation. Even though anticipation is therefore not evident, there is a correlation between the repeat size of the shorter allele (GAA1) and AOO.\(^4\) Disease onset is inversely correlated with GAA1, with a
prediction of a 2.6 years earlier onset for every 100 GAA repeats.\textsuperscript{65} The GAA1 repeat size correlates with frataxin protein levels, cardiac complications, AAO and disease progression, implicating that the shorter allele correlates best with the genotype-phenotype relationship.\textsuperscript{66--68} Similarly to SCAs, GAA1 length explains approximately 50\% of variance in AOO.\textsuperscript{67} In compound heterozygotes patients with loss-of-functions mutations, AOO is decreased.\textsuperscript{40}

### 3.3.2 Modifiers of intergenerational instability

The complex molecular mechanisms and genetic factors influencing intergenerational instability likewise remain insufficiently understood. The tendency of polyQ repeats to expand is disease dependent (low in SCA17; high in SCA7 and DRPLA). With the exception of SCA8, a paternal expansion bias is observed in polyQ diseases that is mainly attributed to the greater number of mitotic divisions in spermatogenesis.\textsuperscript{69} In SCA2 and SCA7, cases of extreme anticipation result in infantile onset of a severe, multisystemic variant of the disease. This phenomenon has recently, and for the first time within the group of SCAs, been described through maternal transmission in SCA7.\textsuperscript{70} Standard laboratory methods may not detect the very high number of repeats seen in infantile presentation, raising awareness of potentially false negative results in pre- and antenatal testing.\textsuperscript{70}

Similarly, GAA-expansions in the $FXN$ gene in FRDA show instability in intergenerational transmission. In contrast to polyQ SCAs, paternally transmitted alleles tend to contract, whereas maternal alleles are equally likely to increase or decrease in repeat-size.\textsuperscript{71} A strong paternal contraction bias is likewise observed in Fragile-X syndrome, which is considered to be related to CpG methylation.\textsuperscript{69}
In summary, extensive research performed on modifiers of AOO and intergenerational in trinucleotide associated ataxias points to repeat number, single-nucleotide polymorphism in DNA repair genes, repeat length in trans alleles and insertions. However, to date, CAT interruptions in SCA1 remain the only discovery that ameliorates prognosis for patients in clinical practice.\textsuperscript{61}

3.3 New insights in genotype-phenotypic presentation – implications beyond ataxia

SCA2 serves as an excellent example of how the full phenotypic spectrum of distinct disorders and the influence of the composition of repeats are only beginning to be revealed. Patients with SCA2 harbour between 33 and 200 repeats in the $ATXN2$ gene and typically present with ataxia accompanied by peripheral neuropathy and slowed ocular saccades.\textsuperscript{72} However, the presence of glutamine coding CAA interruptions within intermediate length repeat expansions in the $ATXN2$ gene has been shown to predispose to a parkinsonian phenotype in pedigrees of both Chinese and European heritage.\textsuperscript{73,74} Interrupted $ATXN2$ tracts appear to have a far-reaching effect on other neurodegenerative diseases as well. Recent investigations focused on the modifying effect of $ATXN2$ on mutations in two ALS associated RNA regulation genes, $c9orf72$ and $TARDBP$. It has been demonstrated that long normal-length expansions of glutamines in the $ATXN2$ gene with CAA-interruptions serve as one of the most important risk factors for ALS, although the exact threshold for this effect remains controversial.\textsuperscript{75–77} Furthermore, experimental evidence demonstrating that $ATXN2$ increases the cytotoxicity of TDP-43\textsuperscript{75}, a protein that is found aggregated in 95% of ALS patients, might serve as an explanation for this.\textsuperscript{78} More recently, the coexistent of an uninterrupted pathogenic $ATXN2$ expansion with 37 repeats and $C9orf71$ mutation was described in two individuals with parkinsonism, ataxia and dementia, thus complicating the contribution of CAA-interruptions in ATXN2 to phenotypic presentation of other neurodegenerative diseases.\textsuperscript{79}
Contrarily to these recent findings, it has been long recognized that SCA3 patients occasionally present with prominent extrapyramidal features, in particular parkinsonism. Genetic variations have been implicated in these cases, and it has been suggested that polymorphism in Parkinson’s related genes and the presence of an APOE ε2 genotype may influence phenotypical presentation, although this has not been confirmed in larger cohorts. The role of ATXN3 as a deubiquitinating enzyme, thus involved in the same pathway as several Parkinson disease associated proteins, supports a probable interaction between these genes.

In SCA17, the length of the CAG/CAA tract, rather than the configuration of the repeats, appears to be correlated with the phenotype.

4. First-line genetic work-up

Considering the marked phenotypic overlap between the different forms of ataxia and the variability of presentation even among patients with the same affected gene, genetic testing is the only way of establishing a conclusive diagnosis in the majority of patients. The choice of genetic work should be guided by a detailed family history, ethinical background, physical examination, biochemical analysis and cerebral MRI studies. Once acquired causes have been excluded, it is generally recommended to test for the most prevalent forms of autosomal dominant and recessive ataxias first, regardless of any known family history. These include: SCA1, SCA2, SCA3, SCA6, SCA7, DRPLA in the Asian population and FRDA. Reduced penetrance and repeat lengths of unknown pathogenicity, as described in several disorders including SCA6 and SCA8, necessitate careful interpretation of results.

In the UK, the cost of first-line SCA genetic test for the polyQ test lies between £160 and £452 with a turnover between 14 and 28 working days. The targeted FXN gene testing costs between £80 and £420 and has a turnover between 3 and 28 working days. Together, these
disorders represent between 50 and 60% of hereditary ataxias worldwide.\textsuperscript{90,91} Thus, approximately half of the patients remain without a confirmed diagnosis after conventional first-line genetic testing.\textsuperscript{10,19} Mutations in rarer genes are often only investigated in cases with guiding clinical or biochemical findings. Moreover, in addition to the group of hereditary cerebellar ataxias, there are nearly 300 genetic conditions in which cerebellar ataxia can be an associated clinical feature\textsuperscript{92}.

5. Next-generation sequencing in clinical practice

The vast number of involved genes and the molecular complexity of ataxias has recently encouraged the implementation of NGS in clinical genetic diagnosis. Indeed, neurological diseases in which a variety of mono-genetic mutations can result in a very uniform phenotype, such as ataxia, have profited enormously from the development of the massively paralleled sequencing methods.\textsuperscript{93} They have provided researchers and clinicians with an unprecedented opportunity to gain genetic information in base pair resolution across the genome in a single experiment at an increasingly affordable cost. NGS approaches range from targeted gene panels to whole exome (WES) and whole genome sequencing (WGS).

Németh \textit{et al.} were the first to demonstrate the utility of targeted gene panels in familial and sporadic ataxia patients.\textsuperscript{94} The reported mean diagnostic delay of 18 years in patients who had tested negative for SCA1, SCA2, SCA3, SCA6, SCA7 and FRDA highlights the diagnostic journey of many individuals. An NGS panel capturing 117 known and putative ataxia genes confirmed a molecular diagnosis in 18% of patients, with the highest detection rate of 75% in familial cases with adolescent onset.\textsuperscript{94} Similar diagnostic rates have been reported in heterogeneous cohorts of ataxia patients who remained without diagnosis after standardised genetic testing.\textsuperscript{91,95,96} WES represents an unbiased NGS approach, thus enabling the detection
of novel genes. Using WES, positive results were obtained in 21 and 41% in heterogeneous cohorts of sporadic and familial ataxia patients in whom previous screening for common mutations had not yield any results. Across all NGS studies, a history of affected family members and an early AOO appear to be the most consistent factors associated with a higher diagnostic success rate.

WES has played a crucial role in the identification of rare ataxia-associated genes that appear to be more prevalent than hitherto expected, such as SYNE1, ANO10 and SPG7. Homozygous mutations in the mitochondrial AAA protease encoded by the SPG7 gene are known to cause autosomal recessive hereditary spastic paraplegia, but were also found by WES in ataxia patients from four independent cohorts in the UK and USA. Encouraged by these findings, it has since been shown that SPG7 mutations are responsible for a significant percentage of unexplained ataxia cases, who may initially present without spasticity.

The greatest limitations that currently prevent a widespread application of NGS in clinical practice in hereditary ataxia are the avalanche of generated data, the risk of incidental findings and the poor ability to sequence repetitive DNA stretches and regions with a high Guanine-Cytosine content. Firstly, debate remains around the critical evaluation of pathogenicity of variants of unknown significance through bioinformatics tools and functional analysis. This currently also represents the main cost factor and time-limiting step. Secondly, ethical challenges may arise from unexpected discoveries of potential medical relevance that are unrelated to the initial diagnostic indication. Several commercial and academic laboratories offer a combination of NGS implementation strategies by exome sequencing with a greater coverage of up to 1000 ataxia associated genes, thus reducing the risk for incidentals findings. Thirdly, it must be kept in mind that copy number variation, such trinucleotide repeat expansions, larger duplications and deletions, cannot reliably be detected by NGS.
sequencing techniques. Large repetitive regions fail to be aligned and mapped to a single position on the reference genome efficiently. Thus, prior exclusion of prevalent ataxias caused by trinucleotide repeat expansions remains paramount to date.

5.1 Newly discovered genes through next-generation sequencing

The success of NGS in a research setting is outlined by its role in the discovery of several new ataxia associated genes, including TGM6 (SCA36), CACNA1G (SCA42), ATP2B3 (X-linked congenital cerebellar ataxia), PNKP (ataxia with oculomotor apraxia type 4), AB1CB7 (X-linked congenital cerebellar ataxia), KCND3 (SCA19/22), and TPP1 (SCAR7). Some of these genes have previously been described in the context of other neurological and non-neurological diseases, such as ceroid lipofuscinosis (TPP1) and Brugada syndrome type 9 (KCND3). Mechanism of genetic pleiotropy include different downstream effects of mutations within the same gene, modifier genes, and oligogenic inheritance. The best known example of genetic pleiotropy within the group of ataxias are CACNA1A mutations that can present as SCA6, episodic ataxia type II and familial hemiplegic migraine due to different functional downstream mechanisms.

6. From bench to bedside

While new genes are being mapped, functional consequences and the reason behind the selective vulnerability of certain neurons to mutations in abundant transcribed ataxia-causing proteins remain largely uncharted. New insights into recurrent pathophysiological mechanism are expected to facilitate the discovery of therapeutic targets that may prove useful in several disorders in the future. Gene co-expression networks recently revealed two SCA gene enriched modules that included genes involved in the ubiquitin-proteasome pathway in granule cells and calcium homeostasis in Purkinje cells. Dysfunction in DNA repair genes, disturbance of
protein expression at the transcriptional and post-transcriptional level, and perturbed glutaminergic signaling represent additional emerging mechanisms of dominant and recessive ataxia associated genes.\textsuperscript{111,116–119} Intervening at the level of the mutant gene can bypass the obstacle of multiple downstream pathogenic pathways. These efforts have already advanced into clinical trials in other neurodegenerative diseases.

6.1 Spinocerebellar ataxias

The two main therapeutic pipelines in SCAs encompass pharmacological agents targeting disrupted downstream pathways and genetic therapy aiming to reduce toxic polyQ gene products.\textsuperscript{120} Given their monogenetic inheritance, the number of involved pathways and the insufficient knowledge of their individual contribution to neurodegeneration, intervening at the source of dysfunction by decreasing the expression level of mutant proteins appears to be a promising approach towards developing a disease-modifying therapy.\textsuperscript{11}

6.1.1 Gene-based approaches

To date, preclinical research focuses on the modulation of protein expression through antisense oligonucleotides (ASOs) and RNA interference (RNAi). ASOs are short, single stranded DNA sequences that bind complementary mRNA transcripts through Watson-Crick hybridisation. The DNA-RNA complex recruits ubiquitously expressed RNase H enzymes, resulting in decreased expression of the targeted protein.\textsuperscript{121} Advances in delivery methods, allele-specificity and intracellular stability pave the way for safe and successful application in humans. ASO-based altering of SMN2 pre-mRNA splicing in children with infantile-onset SMA exemplifies a remarkably successful translation of genetic therapy from bench to bedside.\textsuperscript{14} The broad therapeutically potential of ASO-based therapy is further underlined by completed and ongoing clinical trials in SOD1-associated ALS\textsuperscript{122} and HD (NCT02519036)\textsuperscript{123}. 
In relation to SCA, ASO-mediated removal of the toxic polyQ tract in the mutant ATXN3 gene via exon skipping has been successfully demonstrated in SCA3 fibroblasts\textsuperscript{124,125} and transgenic mice harbouring full-length human ATXN3\textsuperscript{126,127}, however amelioration of motor phenotype was not observed or not assessed. Strikingly, ATXN2-targeting ASOs significantly improved motor performance and extended the average survival not only in SCA2\textsuperscript{128}, but also in TDP43-transgenic mice models of ALS.\textsuperscript{129} This could be advantageous over current strategies directly targeting ALS-associated mutated proteins such as SOD1 and TDP-43 directly, as they only account for 2-5% of ALS cases\textsuperscript{130} and are vital for development and cellular function respectively.\textsuperscript{131}

RNAi is a naturally occurring post-transcriptional gene suppression process, which functions through non-coding double-stranded RNA sequences. RNAi effectors can be introduced into the cell in the form of short interfering RNAs, short hairpin RNAs or artificial miRNAs.\textsuperscript{132} Both non-allele specific and allele-specific RNAi approaches have demonstrated improvement on disease and molecular phenotype in SCA7\textsuperscript{133,134} and SCA3\textsuperscript{135,136} rodent models. Most recently, combined gene-knockdown-replacement therapy using mirtrons has been explored in fibroblast cell lines from SCA7 patients.\textsuperscript{137}

6.1.2. Completed and on-going clinical trials

Five randomized, placebo-controlled clinical trials investigating the safety and efficacy of lithium\textsuperscript{138,139}, varenicline\textsuperscript{140}, riluzole\textsuperscript{141} and tiriluzole\textsuperscript{142} have been completed in the last years. The former three drugs exemplify the increasing use of drug repurposing and are already licenced for other indications, namely bipolar disorders, nicotine addiction and ALS. In 2014, two separate groups reported no significant difference in the Scale for the Rating and Assessment of Ataxia (SARA) and Neurological Examination Score for the Assessment of
Spinocerebellar Ataxia (NESSCA) after lithium treatment for 48 weeks in patients with SCA2 and SCA3. Varenicline, a partial agonist at $\alpha_4\beta_2$ neuronal nicotinic acetylcholine receptors, improved several SARA subscores in a small cohort of SCA3 patients compared to the placebo group. However, owing to the high dropout rate of 40% and the side effect profile, further studies are needed to assess clinical efficacy. Romano and colleagues reported that the glutamate modulator Riluzole improved SARA scores in a heterogeneous group of patients with SCA and FRDA compared to a placebo group at 3 and 12 months. A phase III multicentric trial (NCT03347344) assessing its effect in SCA2 patients will start recruiting patients soon. However, in a phase II/III multicentre clinical trial, Trigriluzole, a prodrug formulation of riluzole, failed to differentiate from placebo on the primary and secondary endpoints after eight weeks of treatment. An open-label extension phase is currently on-going and results are expected at the end of 2018.

Preliminary results published from an open-label pilot trial administrating allogeneic adipose tissue-derived mesenchymal stem cells (MSCs) in six SCA3 patients reported safety and unaltered SARA scores at 12 months. In light of a reported annual decline of 3.00 ± 1.52 in the SARA score reported from a natural history study of a similar cohort, this was cautiously interpreted as a stabilizing effect. The postulated neuroprotective mechanism of MSCs in SCAs include secretion of neurotrophic factors, immune modulation and neuronal replacement. A phase II randomized controlled trial assessing safety and efficacy of MSCs in patients with SCA2 and SCA3 is currently recruiting patients in Taiwan (NCT02540655).

6.2. Friedreich’s ataxia

Based on the current understanding of the functional consequences of decreased FXN expression on iron-sulphur-cluster biogenesis and mitochondria function, the majority of studies in the past have focused on reducing reactive oxygen species (ROS). In brief, perturbed
iron-sulphur-cluster assembly results in mitochondria dysfunction and iron accumulation in the membrane. Consequently, generated ROS triggers an avalanche of toxic downstream mechanisms such as lipid peroxidation. The failure to demonstrate consistent clinical benefit of the most extensively studied antioxidants in FRDA, coenzyme Q\textsubscript{10} (CoQ\textsubscript{10}) and idebenone, has been, among other factors, attributed to their insufficient concentration both within cells and the central nervous system. Several ongoing clinical trials investigate new, tailored antioxidants with increased potency and improved bioavailability. However, as in SCAs, targeting only one factor of the incompletely understood downstream cascade may not be sufficient to improve symptoms. Again, pre-clinical research focuses on targeting the source of dysfunction, hence restoring FXN levels.

6.2.1 Clinical trials

Similar to coenzyme Q10 and idebenone, anti-oxidative properties of the currently investigated agent named EPI-743 (alpha-tocotrienol quinone) rely on a redox active para-benzoquinone ring that undergoes a two-electron cycling reaction. Promising results were obtained in an open-label trial in a small, heterogeneous cohort of patients with mitochondrial disease that included one patient with FRDA. A double-blind placebo-controlled trial including 61 patients with FRDA demonstrated the safety of EPI-743, however its primary endpoint of improvement of visual acuity was not met after 6 months. After completing an 18 month open-label extension phase, treated patients demonstrated improvement in the Friedreich’s Ataxia Rating Scale (FARS)-NEURO, when compared to natural history data.

RTA-408 (Omaveloxolone), an agent with anti-inflammatory and antioxidant properties, slows the degradation rate of nuclear factor erythroid-derived 2-related factor2 (Nrf2). Nrf2 functions as a transcription factor that targets genes, including antioxidant enzymes, that subsequently
impact on mitochondrial function by reducing ROS production.\textsuperscript{156} RTA-408 clinical trials are currently recruiting for a phase II randomized controlled trial named MOXie (NCT02255435).\textsuperscript{157} The part I results of the MOXie trial demonstrated a dose-dependent improvement in a modulated FARS and in Nrf2 associated markers, such as CK and AST. This was also associated with improvements in mitochondrial and neurological function.

At present, there are many innovating clinical trials in the pipeline for therapeutic intervention in FRDA (Table 3). To tackle the low levels of FXN, a small cell-penetrant fusion protein, trans-activator transcription (TAT), has been engineered to shuttle synthetic FXN directly into the mitochondria. In frataxin knockout mouse models, an injection of TAT-frataxin resulted in a prolonged life span of up to 53\% longer with improved cardiac function, growth velocity and cardiac output\textsuperscript{158}. An additional method to overcome the transcriptional deficit in FRDA is to directly provide encapsulated mRNA to cells, as naked mRNA is rapidly degraded, in order to raise mRNA levels.\textsuperscript{159} The application of FXN mRNA to cultured cells or animal models was successfully translated into the FXN protein.\textsuperscript{159} In principle, these techniques provide a novel method to replace the pathogenic depleted protein stores.

\subsection*{6.2.2 Gene-based approaches}

Another valid approach to increase FXN is to reactivate the gene by inhibition of histone deacetylases (HDACs). Several studies of FRDA cell and animal models have shown that specific HDAC inhibitors reverse the epigenetic silencing of the frataxin gene resulting in downstream protein upregulation.\textsuperscript{160} RG2833, a synthetic HDAC inhibitor did not surpass Phase I clinical trials due to the adverse formation of metabolites in the body.\textsuperscript{161} Currently, a new generation of HDAC inhibitors are being developed to prevent harmful metabolites forming.
ASOs that activate, rather than inhibit, gene expression have been developed for the use in FRDA. In FRDA patient-derived fibroblasts, the addition of synthetic duplex RNA, complementary to the GAA repeat region, increased expression of the frataxin gene mRNA by 3 - 4 fold and protein levels by 4 - 6 fold. This increase is consistent with wild-type frataxin levels. Similarly, single-stranded locked nucleic acid (LNA) oligonucleotides increased frataxin gene expression and protein levels in patient-derived fibroblasts. An alternative approach is to use oligonucleotides to eliminate the long noncoding RNA, which suppresses the frataxin gene expression. This method of frataxin upregulation couples the use of oligonucleotides for site recognition with the above mentioned RNase-H. Applying this technique to FRDA patient-derived fibroblasts has shown a significant upregulation of frataxin. In theory, oligonucleotide-based techniques can be used to modulate frataxin expression, but also possibly the downstream events. Therefore, this indicates the widespread use of antisense oligonucleotides as potential therapies in FRDA.

7. Discussion

Even though individually, hereditary ataxias belong to the group of rare disorders, taken together, they represent a prevalent group of disabling neurodegenerative diseases with significant economic burden. NGS has provided insight into the molecular cause of hereditary ataxias, the relationship with other neurodegenerative diseases and is currently making its way into clinical practice. The identification of novel ataxia associated genes in the future is expected, which has the potential to increase molecular diagnoses as a significant proportion of patients with hereditary ataxia still remain undiagnosed. There is a broad consensus that the transition of NGS from a research to a clinical setting possesses a great potential to increase the molecular diagnostic success rates and to improve the clinical management of patients who
remain without diagnosis after standard genetic testing. The continuous decrease of WES costs, the reported higher diagnostic yield and its independence from an a priori hypothesis renders this technique especially suitable for the investigation of these genetically extremely heterogeneous disorders.

The theory of toxic gain-of-function caused by the transcription and accumulation of polyQ proteins certainly represents part of the pathological jigsaw, but increasingly emerges as an oversimplified portrayal of a far more complex process that needs further elaboration.163 For the first time, genetic modulation allows the direct targeting of the prima causa of these disorders. Important advances have been made in allele-specific targeting, as functional consequences of long-term downregulation of wild-type alleles of ataxia associated genes in humans are unknown.124,126 Moreover, the vast number of involved genes certainly complicates the generation of an agent that can be extrapolate to a larger patient population. Therefore, specific pathway-based approaches are still being pursued and many have advanced into clinical trials.

Regardless of the therapeutic approach, the development of disease-modifying agents create the need for robust, objective and easily accessible markers to monitor disease progression and assess treatment response.164 The successful quantification of the mutant huntingtin protein in the cerebrospinal fluid of HD patients and its relationship with disease progression has recently been described.165 Similar quantification of ataxia disease proteins are currently underway and could potentially serve as biomarkers for experimental gene modulation therapies.166

8. Expert Opinion

Over the last two decades, the progress in genetic sequencing has provided us with a better
understanding of hereditary ataxia. Yet, at a closer look, these advances seem to have raised as many questions as they have answered. With the increased implementation of massively paralleled sequencing techniques in routine clinical practice, the number of patients with variants of unknown clinical significance has risen exponentially. To determine the pathogenicity of previously undescribed sequencing variants, time consuming and expensive functional analysis is frequently required. Allele segregation can be helpful in obtaining a conclusive decision, however family members are not always available for molecular testing. Systematic documentation of sequencing variants along with precise clinical information will enable faster segregation between causative and non-causative variants in the future. Misinterpretation of variants of unknown significance can have a major impact on the clinical management of a patient and their families. The affected individual may receive the wrong therapy, prognosis and crucial information about recurrence risk. In patients with inconclusive genetic tests, biomarkers and functional assays could help to support the genetic results. The combined approach of laboratory markers and genetics has the potential to greatly increase the sensitivity of genetic tests and decrease expenses as well as time to diagnosis. Several helpful non-genetic tests are already established in diagnosing ataxic disorders, such as screening for elevated oxysterol markers in Niemann-Pick Type C.167 The need for blood tests to improve diagnosis has also been established in other disorders with underlying genetic mutations, such as cancer syndromes.168

Beyond ameliorating the interpretation of genetic testing, research in the field of biomarkers is urgently needed to optimize the design of clinical trials. As potential disease-modifying drugs are extensively investigated in preclinical and clinical studies, it will become crucial to identify easily accessible biomarkers to monitor the activity and therapeutic response of these agents.
In summary, concomitant with the progress in genetic techniques, the field of ataxia could profit tremendously from the integration of robust biomarkers in both clinical diagnosis and therapeutic studies. Preliminary results of plasma biomarkers, such as neurofilaments, have shown promising potential in other neurodegenerative diseases and subsequently encourages the investigation of biomarkers in hereditary ataxia.\textsuperscript{169}
References

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*Breakthrough Phase 2 trial investigating antisense-oligonucleotide therapy in spinal muscular atrophy.*


*Positive preliminary results of a Phase II clinical trial assessing a huntingtin-lowering drug in the treatment of Huntington's disease.*


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Figure 1. Timeline of gene discoveries in autosomal dominant inherited ataxia. Grey background indicates genes that were discovered by positional cloning and subsequent Sanger sequencing. Blue background indicates genes that were identified by next-generation sequencing approaches. The most prevalent ataxias were discovered with conventional sequencing techniques, whereas novel genes generally underlie a small proportion of ataxia patients. * Disorders that are part of first-line genetic testing. SCA Spinocerebellar ataxia. EA Episodic ataxia. DRPLA Dentatorubral-pallidoluysian atrophy. SPAX1 Spastic ataxia type 1.
Figure 2. Timeline of gene discoveries in autosomal recessive inherited ataxias. Grey background indicates genes that were discovered by positional cloning and subsequent Sanger sequencing. Blue background indicates genes that were identified by next-generation sequencing approaches. The most prevalent ataxias were discovered with conventional sequencing techniques, whereas novel genes generally underlie a small proportion of ataxia patients. * Disorders that are part of first-line genetic testing. # Complex disorders that present with ataxia as a prominent clinical feature. ** X-chromosomal inheritance.
**Figure 2.** Timeline of gene discoveries in autosomal recessive inherited ataxias. Grey background indicates genes that were discovered by positional cloning and subsequent Sanger sequencing. Blue background indicates genes that were identified by next-generation sequencing approaches. The most prevalent ataxias were discovered with conventional sequencing techniques, whereas novel genes generally underlie a small proportion of ataxia patients. * Disorders that are part of first-line genetic testing. # Complex disorders that present with ataxia as a prominent clinical feature. ** X-chromosomal inheritance. LO-GM2 late-onset Tay-Sachs. CTX Cerebrotendinous xanthomatosis. ABLP Abetaliproteinemia. AVED ataxia with Vitamin E deficiency. AT Ataxia telangiectasia. Cockayne S. Cockayne syndrome. FRDA Friedreich’s ataxia. NPC1 Nieman Pick type C1. Refsum Refsum disorder. ATLD Ataxia-telangiectasia-like disorder. ARSACS Autosomal recessive spastic ataxia of Charlevoix-Saguenay. AOA Ataxia with oculomotor apraxia. FXTAS Fragile – tremor/ataxia syndrome. SCAN1 Spinocerebellar ataxia with axonal neuropathy. Cayman Cayman ataxia. SANDO Sensory ataxic neuropathy with dysarthria/dysphagia. MSS Marineseo–Sjögren syndrome. IOSCA Infantile-onset spinocerebellar ataxia. DCMA Dilated cardiomyopathy with ataxia. ARCA Autosomal recessive cerebellar ataxia. SeSAME Seizures, Sensorineural deafness, Ataxia, Mental retardation and Electrolyte imbalance. CAMRQ Cerebellar ataxia, mental retardation, and dysequilibrium syndrome. PHARC Polyneuropathy, hearing loss, ataxia, retinitis pigmentosa, and cataract. SCAR Autosomal recessive spinocerebellar ataxia. SPAX Spastic ataxia.
<table>
<thead>
<tr>
<th>Disease</th>
<th>Gene</th>
<th>Loci</th>
<th>Mutation</th>
<th>Function</th>
<th>Comments</th>
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<tr>
<td>ADCA I – Cerebellar Ataxias with additional features</td>
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<tr>
<td>DRPLA</td>
<td>ATN1</td>
<td>12q13.31</td>
<td>CAG exp.</td>
<td>Transcriptional activator</td>
<td>Myoclonus, epilepsy, choreoathetosis, intellectual decline; DD Huntington disease</td>
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<tr>
<td>SCA1</td>
<td>ATXN1</td>
<td>6p22.3</td>
<td>CAG exp.</td>
<td>Transcriptional repression, involved in developmental processes</td>
<td>Fast progression with early bulbar involvement, pyramidal involvement</td>
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<tr>
<td>SCA2</td>
<td>ATXN2</td>
<td>12q24.13</td>
<td>CAG exp.</td>
<td>Translational modification</td>
<td>Saccade slowing, peripheral neuropathy</td>
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<tr>
<td>SCA3</td>
<td>ATXN3</td>
<td>14p32.12</td>
<td>CAG exp.</td>
<td>Ubiquitin-protease</td>
<td>Also known as Machado-Jacob disease; parkinsonian phenotype in a subgroup of patients</td>
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<tr>
<td>SCA4</td>
<td>Unknown</td>
<td>16q22.1</td>
<td>Unknown</td>
<td>Unknown</td>
<td>Sensory axonal neuropathy</td>
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<tr>
<td>SCA5</td>
<td>SPTBN2</td>
<td>11q13.2</td>
<td>Deletion, MM</td>
<td>Forming of neuronal membrane skeleton</td>
<td>Early-onset severe phenotype described in de novo missense mutations</td>
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<tr>
<td>SCA10</td>
<td>ATXN10</td>
<td>22q13.31</td>
<td>ATTCT exp.</td>
<td>Activation of the mitogen-activated protein kinase cascade</td>
<td>Seizures; reports restricted to Latin American population</td>
</tr>
<tr>
<td>SCA12</td>
<td>PPP2R2B</td>
<td>5q32</td>
<td>CAG exp. (non-coding)</td>
<td>Protein phosphatase</td>
<td>Action tremor; common in Indian ancestry</td>
</tr>
<tr>
<td>SCA13</td>
<td>KCNC3</td>
<td>19q13.33</td>
<td>MM</td>
<td>Membrane potential regulation</td>
<td>Occasionally intellectual disability</td>
</tr>
<tr>
<td>SCA17</td>
<td>TBP</td>
<td>6q27</td>
<td>CAG exp.</td>
<td>DNA-binding subunit of RNA-polymerase II transcription factor</td>
<td>Dementia, chorea, psychiatric symptoms</td>
</tr>
<tr>
<td>SCA18</td>
<td>Unknown</td>
<td>7q22.23</td>
<td>Unknown</td>
<td>Unknown</td>
<td>Sensory-motor neuropathy, atrophy, nystagmus</td>
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<tr>
<td>SCA19/22</td>
<td>KCND3</td>
<td>1p13.2</td>
<td>MM</td>
<td>Voltage-gated potassium channel</td>
<td>Slow progression, rare cognitive impairment, myoclonus, pyramidal signs</td>
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<tr>
<td>SCA20</td>
<td>Unknown</td>
<td>11q12</td>
<td>Duplication</td>
<td>Unknown</td>
<td>Dysphonia, bradykinesia</td>
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<td>SCA21</td>
<td>TMEM240</td>
<td>1p36.33</td>
<td>MM</td>
<td>Transmembrane protein</td>
<td>Intellectual impairment</td>
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<td>SCA23</td>
<td>PDYN</td>
<td>20p13</td>
<td>MM, FS</td>
<td>Synaptic transmission</td>
<td>Dysarthria, myoclonus, peripheral neuropathy</td>
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<td>SCA25</td>
<td>Unknown</td>
<td>2p15-21</td>
<td>Unknown</td>
<td>Unknown</td>
<td>Sensory neuropathy</td>
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<td>SCA26</td>
<td>EEF2</td>
<td>19p13.3</td>
<td>MM</td>
<td>Translation</td>
<td>Sensory neuropathy, dysarthria</td>
</tr>
<tr>
<td>Disease</td>
<td>Gene</td>
<td>Loci</td>
<td>Mutation</td>
<td>Function</td>
<td>Clinic</td>
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<tr>
<td><strong>SCA27</strong></td>
<td>FGF14</td>
<td>13q13.1</td>
<td>MM</td>
<td>Cell growth and survival</td>
<td>Cognitive deficits, dyskinesia, tremor</td>
</tr>
<tr>
<td><strong>SCA28</strong></td>
<td>AFG3L2</td>
<td>18p11.21</td>
<td>MM</td>
<td>ATP-dependent protease</td>
<td>Ophthalmoparesis, ptosis; Allelic to SPAX5</td>
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<tr>
<td><strong>SCA29</strong></td>
<td>ITPR1</td>
<td>3p26.1</td>
<td>MM</td>
<td>Ca(^{2+}) signalling</td>
<td>Slow progressive, learning deficits; Allelic to SCA15</td>
</tr>
<tr>
<td><strong>SCA34</strong></td>
<td>ELOVL4</td>
<td>5q14</td>
<td>MM</td>
<td>Lipid metabolism</td>
<td>Erythrokeratodermia variabilis described</td>
</tr>
<tr>
<td><strong>SCA35</strong></td>
<td>TGM6</td>
<td>20p13</td>
<td>MM</td>
<td>Protein crosslinking</td>
<td>Hyperreflexia, dystonia</td>
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<tr>
<td><strong>SCA36</strong></td>
<td>NOP56</td>
<td>20p13</td>
<td>GGGCCTG exp. (non-coding)</td>
<td>RNA-procession</td>
<td>Fasciculations, tongue atrophy</td>
</tr>
<tr>
<td><strong>SCA42</strong></td>
<td>CACNA1G</td>
<td>17q21.33</td>
<td>MM</td>
<td>Ca(^{2+}) signalling</td>
<td>Mild pyramidal signs</td>
</tr>
<tr>
<td><strong>SCA43</strong></td>
<td>MME</td>
<td>3q25.2</td>
<td>MM</td>
<td>Zinc-dependent metalloprotease</td>
<td>Reported in one family</td>
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</table>

**ADCA II – Cerebellar ataxia with pigmental retinal degeneration**

<table>
<thead>
<tr>
<th>Disease</th>
<th>Gene</th>
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<th>Mutation</th>
<th>Function</th>
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<td>SCA7</td>
<td>ATXN7</td>
<td>3p14.1</td>
<td>CAG exp.</td>
<td>Transcription factor</td>
<td>Visual loss caused by retinopathy</td>
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**ADCA III – ‘Pure’ cerebellar ataxias**

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<tr>
<th>Disease</th>
<th>Gene</th>
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<th>Mutation</th>
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<tr>
<td>SCA6</td>
<td>CACNA1A</td>
<td>19p13.13</td>
<td>CAG exp.</td>
<td>Voltage-gated calcium channel</td>
<td>Allelic to EA2 and familial hemiplegic migraine</td>
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<tr>
<td>SCA8</td>
<td>ATXN8OS-ATXN8</td>
<td>13q21</td>
<td>CTA.CTG exp. (non-coding)</td>
<td>Non-protein coding; Unknown</td>
<td>Standardized genetic test not established.</td>
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<td>SCA11</td>
<td>TTBK2</td>
<td>15q15.2</td>
<td>Deletion</td>
<td>Tau phosphorylation</td>
<td>Benign course</td>
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<tr>
<td>SCA14</td>
<td>PRKCG</td>
<td>19q13.42</td>
<td>MM</td>
<td>Protein phosphorylation</td>
<td>Myoclonus</td>
</tr>
<tr>
<td>SCA15/16</td>
<td>ITPR1</td>
<td>3p26.1</td>
<td>MM, Deletion</td>
<td>Ca(^{2+}) signalling</td>
<td>Slow progression, occasionally intellectual disability</td>
</tr>
<tr>
<td>SCA31</td>
<td>BEAN1</td>
<td>16q22</td>
<td>TGGAA exp. (non-coding)</td>
<td>Ubiquitin-pathway</td>
<td>Sensorineural hearing loss</td>
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<tr>
<td>SCA38</td>
<td>ELOVL5</td>
<td>6p12</td>
<td>MM</td>
<td>Lipid metabolism</td>
<td>Slow progression</td>
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<tr>
<td>SCA40</td>
<td>CCDC88C</td>
<td>14q32.12</td>
<td>MM</td>
<td>WNT signaling</td>
<td>Reported in one patient</td>
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<tr>
<td>SCA41</td>
<td>TRPC3</td>
<td>4q27</td>
<td>MM</td>
<td>Regulations MP, Ca signalling</td>
<td>Reported in one family</td>
</tr>
</tbody>
</table>

**Table 1. Genes, loci, gene function and distinguishing features of autosomal dominant hereditary ataxias.** Classified according to the Harding classification (1982). Exp Expansion. FS Frameshift. MM Missense mutation.
<table>
<thead>
<tr>
<th>Disorder</th>
<th>Gene</th>
<th>Loci</th>
<th>Mutation</th>
<th>Function</th>
<th>Comments</th>
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<tr>
<td>Friedreich's ataxia</td>
<td>FXN</td>
<td>9q13</td>
<td>GAA repeat, Deletion</td>
<td>Mitochondria iron transport and respiration</td>
<td>Neuropathy, insulin resistance, cardiomyopathy, scoliosis, visual and hearing impairment</td>
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<tr>
<td>Cayman ataxia</td>
<td>ATCAY</td>
<td>19p13.3</td>
<td>MM</td>
<td>Neural tissue development</td>
<td>Psychomotor retardation; frequent in Cayman population</td>
</tr>
<tr>
<td>Refsum's disease</td>
<td>Pex7, PHYH</td>
<td>6q23.3, 10p13</td>
<td>MM, FS</td>
<td>Fatty acid oxidation</td>
<td>Neuropathy, ichthyosis, retinopathy</td>
</tr>
<tr>
<td>Abetalipoproteinaemia</td>
<td>MTTP</td>
<td>4q23</td>
<td>NM, MS, FS</td>
<td>Lipoprotein assembly</td>
<td>Fat malabsorption, acanthocytosis, neuropathy, spasticity</td>
</tr>
<tr>
<td>Autosomal recessive spastic ataxia of Charlevoix-Saguenay</td>
<td>SACS</td>
<td>13w12</td>
<td>MM, FS</td>
<td>Regulates ataxia proteins</td>
<td>Spastic paraparesis; OCT: hypertrophy of myelinated fibres</td>
</tr>
<tr>
<td>Ataxia telangiectasia</td>
<td>ATM</td>
<td>11q22.3</td>
<td>MM, NM, deletion</td>
<td>Phosphorylation</td>
<td>Telangiectasias, cancer, immunodeficiency; raised α-fetoprotein</td>
</tr>
<tr>
<td>Ataxia telangiectasia-like disorder</td>
<td>MRE11</td>
<td>11q21</td>
<td>MM, NM</td>
<td>DNA double-strand break repair</td>
<td>Mimics ataxia telangiectasia; no ocular telangiectasias</td>
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<td>Ataxia with oculomotor apraxia type 1</td>
<td>APTX</td>
<td>9p21.1</td>
<td>MM, LOF</td>
<td>Single-stranded DNA repair</td>
<td>Oculomotor apraxia, chorea, dystonia, neuropathy</td>
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<tr>
<td>Ataxia with oculomotor apraxia type 2</td>
<td>SETX</td>
<td>9q34.13</td>
<td>MM</td>
<td>DNA and RNA processing</td>
<td>Oculomotor apraxia, chorea, dystonia, neuropathy</td>
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<tr>
<td>Ataxia with oculomotor apraxia type 3</td>
<td>PIK3R5</td>
<td>17p13.1</td>
<td>MM</td>
<td>Phosphorylation</td>
<td>Oculomotor apraxia, neuropathy</td>
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<td>Cerebrotendinous xanthomatosis</td>
<td>CYP27A1</td>
<td>2q35</td>
<td>MM</td>
<td>Oxidation</td>
<td>Xanthomas, spasticity, neuropathy, intellectual disability, cataracts</td>
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<td>Marinesco-Sjogren syndrome</td>
<td>SIL1</td>
<td>5q31.2</td>
<td>LOF</td>
<td>Nucleotide exchange factor</td>
<td>Hypotonia, intellectual disability, early-onset cataract</td>
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<td>SANDO/MIRAS</td>
<td>POLG1</td>
<td>15q26.1</td>
<td>MM</td>
<td>Mitochondrial DNA polymerase</td>
<td>Myoclonus, dysarthria, ophthalmalgin</td>
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<td>Autosomal recessive ataxia type 1</td>
<td>SYNE1</td>
<td>6q25.2</td>
<td>Splice site</td>
<td>Nuclear membrane localisation</td>
<td>Cerebellar atrophy, lack of neuropathy</td>
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<td>ADCK3</td>
<td>1q42.13</td>
<td>Splice site</td>
<td>Electron transfer, respiratory chain</td>
<td>Mental disability, epilepsy myoclonus, exercise intolerance</td>
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<td>Autosomal recessive ataxia type 3</td>
<td>ANO10</td>
<td>3p22.1-p21.33</td>
<td>LOF</td>
<td>Calcium-activated chloride channels</td>
<td>Pure ataxia</td>
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<td>Condition</td>
<td>Gene</td>
<td>Chromosome</td>
<td>Mutations</td>
<td>Function</td>
<td>Phenotype</td>
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<tr>
<td>Late onset Tay-Sachs</td>
<td>HEXA</td>
<td>15q23</td>
<td>MM, LOF</td>
<td>Ganglioside degradation</td>
<td>Motor neuron involvement, psychiatric features</td>
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<tr>
<td>Ataxia with Vitamin E deficiency</td>
<td>TTPA</td>
<td>8q12.3</td>
<td>FS</td>
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<td>Mimics Friedreich’s ataxia, decreased Vitamin E, head tremor</td>
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<td>Cockayne Syndrome</td>
<td>ERCC8</td>
<td>5q12.2</td>
<td>NM, MM</td>
<td>Signal transduction</td>
<td>Microcephaly, growth retardation, photosensitivity, progeria</td>
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<td>Niemann-Pick type C1</td>
<td>NPC1</td>
<td>18q11.2</td>
<td>MM, FS</td>
<td>Cholesterol trafficking</td>
<td>Vertical supranuclear palsy, splenomegaly, dystonia, cognitive disability</td>
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<td>Fragile X-associated tremor/ataxia syndrome</td>
<td>FMR1</td>
<td>Xq27.3</td>
<td>CGG repeat (5’ UTR)</td>
<td>RNA binding</td>
<td>Late-onset, intention tremor, cognitive problems</td>
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<tr>
<td>Spinocerebellar ataxia with axonal neuropathy</td>
<td>TDP1</td>
<td>14q32.11</td>
<td>MM</td>
<td>Repairing stalled topoisomerase I-DNA complexes</td>
<td>Peripheral neuropathy</td>
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<td>Infantile-onset spinocerebellar ataxia</td>
<td>C10orf2</td>
<td>10q24.31</td>
<td>MM</td>
<td>mtDNA-specific helicase</td>
<td>Atheosis, muscle hypotonia, optic atrophy, primary hypogonadism in females</td>
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<td>DNAJC19</td>
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<td>Splice site</td>
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<td>Early-onset cardiomyopathy</td>
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<td>SeSAME syndrome</td>
<td>KCNJ10</td>
<td>1q23.2</td>
<td>LOF, MM</td>
<td>Potassium buffering</td>
<td>Seizures, sensorineural deafness, ataxia, mental retardation and electrolyte imbalance</td>
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<td>CA8</td>
<td>8q12.1</td>
<td>MM</td>
<td>Gene transcription</td>
<td>Cognitive impairment</td>
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<td>PHARC</td>
<td>ABHD12</td>
<td>20p11.21</td>
<td>LOF</td>
<td>Hydrolysis of lipid transmitters</td>
<td>Polyneuropathy, hearing loss, ataxia, retinitis pigmentosa and cataract</td>
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<td>KIAA0226</td>
<td>3q29</td>
<td>FS</td>
<td>Vesicular trafficking, endosome maturation</td>
<td>Epilepsy</td>
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<td>Spastic ataxia type 4</td>
<td>MTPAP</td>
<td>10p11.23</td>
<td>MM</td>
<td>mRNA degradation</td>
<td>Spasticity</td>
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<td>Posterior column ataxia with retinitis pigmentosa</td>
<td>FLVCR1</td>
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<td>Heme transporter</td>
<td>Impairment of vision and proprioception</td>
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<td>TTBK2</td>
<td>15q15.2</td>
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<td>Phosphorylation</td>
<td>Occasionally pyramidal involvement and peripheral neuropathy</td>
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<td>Cerebellar ataxia, mental retardation</td>
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<td>17p13.3</td>
<td>MM</td>
<td>Endolysosomal trafficking</td>
<td>Cognitive impairment</td>
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<td>Chromosome</td>
<td>Mode of Inheritance</td>
<td>Pathway/Function</td>
<td>Phenotype</td>
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<td>11q13.2</td>
<td>MM</td>
<td>Glutamate signaling regulation</td>
<td>Early-onset ataxia, psychomotor developmental delay</td>
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<td>GRM1</td>
<td>6q24.3</td>
<td>MM</td>
<td>Glutamatergic neurotransmission</td>
<td>Delayed psychomotor development, mental retardation</td>
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<td>Spastic ataxia type 3</td>
<td>MARS2</td>
<td>2q33.1</td>
<td>DR, Duplication-deletion</td>
<td>Mitochondrial</td>
<td>Spasticity, Leukoencephalopathy</td>
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<td>Autosomal recessive spinocerebellar ataxia 18</td>
<td>GRID2</td>
<td>4q22.1-q22.2</td>
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<td>Synapse organisation</td>
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<td>STUB1</td>
<td>16p13.3</td>
<td>MM</td>
<td>E3 ubiquitin ligase</td>
<td>Occasionally hypogonadism</td>
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<td>8q12.3</td>
<td>Splice site, MM</td>
<td>Regulating vitamin E levels</td>
<td>Developmental delay</td>
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<td>Cerebellar ataxia, mental retardation and disequilibrium syndrome 4</td>
<td>ATP8A2</td>
<td>13q12.13</td>
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<td>Epilepsy, delayed psychomotor development, mental retardation</td>
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<td>6p22.3</td>
<td>Splice site</td>
<td>DNA repair</td>
<td>Delayed psychomotor development, seizures</td>
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<td>mRNA splicing</td>
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<td>19q13.33</td>
<td>LOF</td>
<td>DNA repair</td>
<td>Oculomotor apraxia, microcephaly, seizures</td>
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<td>9q34.3</td>
<td>MM</td>
<td>Cleavage</td>
<td>Cognitive impairment, dystonia, spasticity</td>
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<tr>
<td>Condition</td>
<td>Gene</td>
<td>Locus</td>
<td>Mutation Type (MM)</td>
<td>Function</td>
<td>Feature</td>
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<td>MM</td>
<td>Activates ubiquitin-fold modifier 1</td>
<td>Early-onset cataract</td>
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<td>VWA3B</td>
<td>2q11.2</td>
<td>MM</td>
<td>Transcription/DNA repair</td>
<td>Spasticity, intellectual disability</td>
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</table>

**Table 1. Genes, loci, gene function and distinguishing features of autosomal recessive hereditary ataxias.** DR duplication rearrangement. FS frame shift. NM nonsense mutation. MM missense mutation. LOF loss of function. TM truncation mutation.