PTPRR Protein Tyrosine Phosphatase Isoforms and Locomotion of Vesicles and Mice

Wiljan J. A. J. Hendriks · Gönül Dilaver · Yvet E. Noordman · Berry Kremer · Jack A. M. Fransen

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Abstract Protein tyrosine phosphatases (PTPs) are central players in many different cellular processes and their aberrant activity is associated with multiple human pathologies. In this review, we present current knowledge on the PTPRR subfamily of classical PTPs that is expressed in neuronal cells and comprises receptor-type (PTPBR7, PTP-SL) as well as cytosolic (PTPPBSγ-37, PTPPBSγ-42) isoforms. The two receptor-type isoforms PTPBR7 and PTP-SL both localize in late endosomes and the Golgi area. PTPBR7, however, is additionally localized at the cell surface and on early endosomes. During cerebellar maturation, PTPBR7 expression in developing Purkinje cells ceases and is replaced by PTP-SL expression in the mature Purkinje cells. All PTPRR isoforms contain a kinase interacting motif that makes them mitogen-activated protein kinase phosphatases. The distinct subcellular localization of the different PTPRR isoforms may reflect differential roles in growth-factor-induced MAPK-mediated retrograde signaling cascades. Studies in PTPRR-deficient mice established that PTPRR isoforms are physiological regulators of MAPK phosphorylation levels. Surprisingly, PTPRR-deficient mice display defects in motor coordination and balancing skills, while cerebellar morphological abnormalities, which are often encountered in ataxic mouse models, are absent. This is reminiscent of the phenotype observed in a handful of mouse mutants that have alterations in cerebellar calcium ion homeostasis. Elucidation of the molecular mechanisms by which PTPRR deficiency imposes impairment of cerebellar neurons and motor coordination may provide candidate molecules for hereditary cerebellar ataxias that still await identification of the corresponding disease genes.

Keywords Calcium signaling · Endosomes · Ultrastructural localization · Live cell imaging · Motor coordination · Signal transduction

Introduction

Reversible tyrosine phosphorylation is a powerful mechanism to regulate protein-mediated processes that steer the proliferation, differentiation and functioning of cells and the development, homeostasis and functioning of multicellular organisms. Protein tyrosine phosphatases (PTPs) and protein tyrosine kinases are the enzyme classes that are instrumental in controlling the spatial and temporal ratio of tyrosine-phosphorylated and non-phosphorylated targets and thus coordinately regulate cellular responses to intracellular and extracellular cues. Tyrosine phosphatases can be roughly divided into two distinct groups: the truly phosphorylosine-specific PTPs on the one hand and the dual-specificity phosphatases (DSPs) on the other [1, 2]. DSPs can dephosphorylate phosphoserine and phosphothreonine in addition to phosphorylosine residues, and some are also active against phospholipids [3]. The classical, phosphorylosine-specific PTPs consist of a subgroup that possesses a
transmembrane domain, a receptor-like extracellular region and one or two catalytic PTP domains and a second subgroup that is non-transmembrane and mostly cytoplasmic in location [4]. The non-transmembrane PTPs and DSPs all contain a single catalytic PTP domain. This ~250-residue-long PTP domain is highly conserved and contains an essential cysteine residue within its active site that is used in a cysteinyl-phosphate enzyme intermediate during dephosphorylation [5].

How PTP activity is regulated is still subject of intense investigation, but mechanisms include alternative mRNA splicing, modulation of steady state levels, posttranslational modification (including phosphorylation), dimerization and/or subcellular confinement [6]. With regard to the physiological functions of individual PTPs, also much more work needs to be done. Many studies at the cellular level are hampered by the very low endogenous expression levels of PTPs and their firm catalytic activity that is some log orders of magnitude over that of kinases. In combination, this may cause overexpression artifacts in transfected cell-based readouts, and thus, it should come as no surprise that most information on PTP functioning resulted from studies of mammalian pathologies or via exploitation of genetically modified animal models [4]. In this paper, we will review the current knowledge on the neuronal, phosphotyrosine-specific PTP isoforms that are encoded by the gene Piprr in mouse and PTPRR in man. This gene and its products may be of particular relevance to cerebellar function, as rodent postnatal cerebellar Purkinje cells express a unique PTPRR isoform (PTP-SL) and Piprr knockout mice display an ataxia phenotype. We will focus on the ways that confine and regulate PTPRR enzymatic activity, most notably the distinct subcellular locations displayed by the various isoforms, and will elaborate on the motor coordination defect displayed by PTPRR-deficient mice.

### Protein Tyrosine Phosphatase Receptor-Type R Isoforms

PTPRR gene orthologues are present in many vertebrates and homologous genes can still be found in distant species like Drosophila (Table 1) [7–10]. Interestingly, like in human and rat, the mouse Piprr gene gives rise to multiple neuronal PTPRR protein isoforms [11, 12]. Four different transcripts (represented by PTPBR7, PTP-SL, PTPPBSγ+, and PTPPBSγ− cDNAs) have been identified [7, 13, 14] that result from the differential use of three distinct transcription start sites [12, 15] and the alternative inclusion of a 117-nucleotide sequence stretch in the unique 5′ UTR of the shortest mRNA (PTPPBSγ) [7]. In situ hybridization studies, Northern blot experiments, and reverse transcription polymerase chain reaction (RT-PCR) analyses have gathered RNA expression data for these individual transcripts. PTPBR7 messengers are expressed during early embryogenesis in spinal ganglia and Purkinje cells (PCs). PTP-SL transcripts are not present during prenatal development, but postnatally, PTP-SL transcripts replace the decreasing PTPBR7 levels in PCs [15]. PTPBR7 expression persists in all other brain regions after birth. Trace amounts of PTP-SL messengers are present in midbrain, brainstem, and cortex, but firm expression is exclusively in adult cerebellar PCs (Fig. 1a). Intriguingly, this cell-specific expression pattern is fully conserved in rat [8, 9]. Mouse PTPPBSγ variants are expressed at extremely low levels throughout the brain. In fact, they represent the only Piprr transcripts that have been detected in non-neuronal tissues; trace amounts have been detected in the gastrointestinal tract and in developing cartilage [7, 12]. Unfortunately, most cell lines in culture do not express PTPRR-encoding transcripts. Thus far, only in the rat neuroendocrine PC12 cells has a Piprr-derived transcript, the one that encodes the receptor-type protein isoform PCPTP1, been detected [9, 16].

| Species          | Homo sapiens | Mus musculus | Rattus norvegicus | Drosophila melanogaster |
|------------------|--------------|--------------|-------------------|------------------------|
| Gene (position)  | PTPRR        | Piprr        | Piprr             | Piprr                  |
| Protein isoforms | PTPPBSα      | PTPBR7       | PCPTP1            | –                      |
|                  | PTPBSβ       | PTP-SL       | PCPTP1-Ce         | –                      |
|                  | PTPPBSγ−     | PTPPBSγ−42   | –                 | PTP-ER                 |
|                  | PTPPBSγ+     | –            | –                 | –                      |

The different Piprr transcripts give rise to multiple PTPRR protein isoforms (Fig. 1b). Mutagenesis studies revealed exactly which AUG start codons are being used in the various Piprr mRNAs [12]. Intriguingly, the second as well as the third start site in the two types of PTPPBSγ mRNAs are preferred by the ribosomes, generating isoforms PTPPBSγ−42 and PTPPBSγ−37 with sizes of 42 and 37 kDa, respectively. PTPBR7 and PTP-SL are 72 and 60 kDa in size, respectively. PTPBR7 is a classical type I transmembrane protein: its functional signal-peptide is cleaved of in the endoplasmic reticulum (ER) and a hydrophobic segment further on in the protein functions as transmembrane anchor. The PTP-SL isoform contains
two hydrophobic regions in the N-terminal part, which led originally to the prediction that it also would be a type I transmembrane protein [13]. However, subsequent studies revealed that PTP-SL is lacking a functional signal peptide and thus may be a membrane-associated protein [15]. Only recently, the use of a selective cell permeabilization method in combination with dedicated antisera directed against PTPRR domains allowed the determination of PTP-SL’s exact membrane topology to be a type III transmembrane protein [17]. The two smallest isoforms, PTPPBSγ-42 and PTPPBSγ-37, are located in the cytosol. Interestingly, the two receptor-type isoforms, PTPBR7 and PTP-SL, form homo- and heteromeric complexes, whereas the cytosolic species PTPPBSγ-42 and PTPPBSγ-37 cannot. Furthermore, in line with data for a number of other RPTPs, the enzymatic activity of PTPRR monomers towards an artificial substrate decreased significantly upon multimerization [17]. This opens up the possibility that the activity of the receptor-type PTPRR isoforms can be modulated through ligand binding. Ligand hunting strategies using tagged PTPBR7 ectodomain fusion proteins [18] may serve in the identification of such molecules.

PTPRR Can Bind and Inactivate Mitogen-Activated Protein Kinases

Growth factor binding induces dimerization of their receptor tyrosine kinase (RTK), which results in increased kinase activity and the generation of phosphotyrosine sites on the RTK cytosolic portion that serve as docking sites for interacting proteins [19, 20]. The consequent recruitment, assembly, and phosphorylation of these downstream signaling complexes, in turn, switches on many different signaling pathways, including the intensively studied mitogen-activated protein kinase (MAPK) cascades [20]. MAPK pathways are organized into three layers of consecutive acting kinases that through the dual kinase specificity of MAPK-kinases lead to the phosphorylation of two regulatory residues (pT-X-pY) in MAPKs, serine/
threonine kinases that continuously shuttle between the nucleus and the cytoplasm (Fig. 1c). Once activated by dual phosphorylation, MAPKs can phosphorylate a wide range of cytosolic and nuclear substrates [21, 22]. Termination of MAPK signaling thus can be achieved, in principle, by phosphatases that are either serine/threonine-specific (PP phosphatases), tyrosine-specific (classical PTPs), or of the dual-specificity type (DSPs) [23–25]. The subcellular localization and activity of such MAPK phosphatases is highly relevant for proper spatiotemporal control of MAPK activity and thus a key determinant of the final cellular response triggered by ligand-induced activation of RTKs [26, 27].

In the human genome, a subset of 11 DSP genes encode phosphatases that are especially suited to bind, dephosphorylate, and thus inactivate MAPKs, and these are generally referred to as MKPs (acronym for MAP kinase phosphatases) [28]. An additional three genes encode classical, tyrosine-specific PTPs capable of binding and inactivating MAPK family members, among which PTPRR [29]. Both the dual-specific MKPs and the tyrosine-specific MAPK phosphatases (MAPK-PTPs) have in common the presence of a 16-amino acid kinase interaction motif (KIM), located just N-terminal of the phosphatase domain, which is required for the association with MAPKs. All the PTPRR isoforms contain such a KIM (Fig. 1b) which in part determines the binding specificity towards the various MAPKs [30, 31]. Association specificity is also influenced by a small sequence that is located in between the KIM and the PTP domain [32]. As a substrate, the MAPK-PTPs prefer ERK1/2/5 and p38 over JNK [31–33]. Intriguingly, overexpression of MAPK-PTPs demonstrated that they not only are able to inactivate MAPKs but that their binding also prevents the translocation of the MAPKs into the nucleus [31, 34, 35]. Furthermore, upon binding to activated MAPKs, the MAPK-PTP itself first becomes phosphorylated by the MAPK (on a threonine residue just outside the KIM domain) before it can dephosphorylate the protein [30]. Importantly, the interaction between the two proteins is regulated by protein kinase A (PKA). Phosphorylation of a conserved serine residue within the KIM domain, which can be reversed by PP1 [36, 37], abolishes the binding and subsequent dephosphorylation of MAPKs and results in the release of MAPKs to the nucleus [34, 38]. This PKA-mediated regulatory principles also hold for most of the dual-specificity MKPs that contain a KIM-like domain [28].

**Distinct Subcellular Localization for Receptor-Type PTPRR Isoforms**

The PTPRR receptor-type isoforms PTPBR7 and PTP-SL share a common membrane topology [17] and both localize at the trans-Golgi network and on endocytic vesicles [15, 39]. However, PTPBR7 is found at the cell surface, whereas PTP-SL is not. Since detailed knowledge on their subcellular localization bears relevance for retrograde signaling cascades by endocytosed growth factor receptors, PTPBR7 and PTP-SL decorated vesicles were classified to different stages of the endocytic pathway (see Electronic Supplementary Material). Endogenous expression of PTPRR proteins in either primary neurons (Electronic Supplementary Material Fig. S1) or PC12 cells [40] is very low but reflects the localization pattern observed in transfected cells [39]. Fluorescent Dextran uptake experiments with mouse Neuro-2a cells transiently expressing EGFP-tagged PTPBR7 (Electronic Supplementary Material Fig. S2) or PTP-SL (Electronic Supplementary Material Fig. S3) corroborated that PTPBR7 and PTP-SL are present on endosomes [39]. Electron microscopic monitoring of timed 5-nm bovine serum albumin gold uptake established that PTP-SL locates predominantly to late endosomes, whereas PTPBR7 is also present at the plasma membrane and endosomes from earlier stages (Electronic Supplementary Material Fig. S4). Co-transfection of Neuro-2a cells with PTPBR7 and PTP-SL expression plasmids confirmed that PTPBR7 and PTP-SL both are present on late endosomal vesicles (Fig. 2). It should be noted, however, that it is unlikely that co-localization of the two isoforms will actually occur in vivo since PTPBR7 transcripts are detectable throughout the developing and adult brain, but in PCs, in fact, PTP-SL transcripts take over during maturation [15]. Single-positive PTPBR7 vesicles are particularly found in the vicinity of the plasma membrane, reminiscent of the early endocytic compartment in which PTP-SL is sparse (Fig. 2a–c). Analysis of the dynamics of the vesicles harboring receptor-type PTPRR isoforms, by time-lapse fluorescence microscopy of PTPBR7-EGFP and PTP-SL-EGFP proteins in living cells, revealed both anterograde and retrograde movement (Electronic Supplementary Material Fig. S5) with an average speed that did not differ significantly between PTPBR7 and PTP-SL positive vesicles. The PTPRR positive vesicle trafficking patterns are reminiscent of movement along microtubule tracks (Electronic Supplementary Material Fig. S6). Together, these findings demonstrate that PTPBR7 is present on vesicles throughout the endocytic pathway from the plasma membrane to the Golgi apparatus. In contrast, PTP-SL is only present on vesicles participating in the later stages of endocytosis, close to the trans-Golgi network, indicative of differential functions of these PTPRR isoforms within the endocytic pathway.

**Towards the Function of PTPRR Protein Isoforms**

It could be that the PTPRR isoforms PTPBR7 and PTP-SL are merely cargo proteins that for example need to be...
delivered to lysosomes for degradation. In line with this, all PTPRR isoforms are short-lived proteins with half-lives varying from 3 to 5 h [11]. However, PTPRR proteins do not co-localize with lysosomal marker proteins [39] and both anterograde and retrograde movements for PTPRR-positive vesicles are observed (Electronic Supplementary Material Fig. S5), whereas cargo proteins are expected to travel unidirectional and to accumulate at locations within the cell. This leaves two other options, namely that these phosphatases make part of the endocytic transport machinery itself and/or are involved in localized signaling events. Accurate packaging of cargo and (coated) vesicle formation and membrane fusion in vivo requires tight regulation for which phosphorylation is indeed one of the mechanisms [41]. For example, EGF-induced tyrosine phosphorylation of the β2-subunit of the adapter protein complex AP-2 turned out to be important in regulating EGFR protein turnover [42]. Likewise, the activity of PTP-MEG2 (PTPN9) on secretory vesicle membranes is required to locally bind and dephosphorylate NSF and promote secretory vesicle fusion [43]. PTPRR interacted with the β4-subunit of the AP-4 complex in a yeast two-hybrid system and co-localized with AP-4 at the Golgi apparatus and on endosomes in neuronal cells [39], suggesting a role in AP-4-mediated vesicle sorting. In line with this, co-expression of PTP-SL and β4-adaptin in Neuro-2a cells led to a displacement of PTP-SL from vesicles and the Golgi apparatus and a redistribution throughout the cytoplasm [39].

The alternative option would be a role for PTPRR in spatiotemporal control of growth factor signaling involving receptor internalization [44]. For example, after ligand binding and autophosphorylation at the plasma membrane, the EGF tyrosine kinase receptor is internalized and subsequently dephosphorylated by PTP1B that resides at the ER membrane, eventually resulting in receptor silencing (signal termination) and RTK breakdown in lysosomes [45]. Localized signaling has also been demonstrated for the MAPK scaffold Sef that resides on the Golgi apparatus; it binds ERK—thus blocking nuclear translocation—without inhibiting its activity in the cytoplasm [46]. In view of their KIM domain-mediated association with MAPKs, resulting in MAPK inactivation and prevention of nuclear translocation [30, 31, 33], a comparable role for PTPRR isoforms may be expected. Most of these studies, however, involved (ectopic) overexpression of PTPRR isoforms. The endogenous expression of PTPRR in PC12 cells [16], a paradigm model for the study of spatial and temporal control of MAPK signaling [26, 27], allowed assessment of the impact of PTPRR on MAPK activity under more physiological conditions. Modulation of PTPRR activity in these cells, however, had no effect on the temporal profile of EGF- and NGF-induced ERK1/2 activity [40]. Since in PC12 cells the two other MAPK-PTPs are not expressed, this finding suggests that rather the dual-specificity MKPs serve as key MAPK regulators.

Biological Activities of Human PTPRR Proteins

Information on putative roles of PTPRR isoforms in health and disease is limited. The human PTPRR gene has been
PTPPR-Deficient Mice Display Motor Coordination Defects

As an alternative approach to gain insights in the physiological role of PTPRR isoforms, PTPRR-deficient mice have been generated [55]. Knockout animals appeared healthy and were fertile, and inheritance of the mutant allele followed Mendelian rules. The high expression levels of PTPRR isoforms in the hippocampal area and especially in the cerebellar Purkinje cells led to the assessment of mouse basic behavior and locomotion. Nocturnal activity of PTPRR-deficient mice was significantly reduced as compared to wild-type control mice. In addition, PTPRR-deficient mice displayed significant defects in their fine motor coordination and balance skills that are reminiscent of a mild ataxia [55]. Since MAPK signaling pathways are implicated in various neuropathological states [56] and PTPRR can interact with several MAPKs, it should perhaps not come as a surprise that ERK1/2 phosphorylation levels were indeed found to be significantly elevated in PTPRR-deficient brains. However, brain morphology, Purkinje cell number, or dendritic branching was not affected [55].

The majority of ataxia mouse models display histological phenomena (i.e., loss of cerebellar granule cells or PCs) that explain cerebellar dysfunction [57]. Only a few mouse models exist that display ataxic behavior in combination with normal cerebellar morphology. These include junctophilin 3 [58], carbonic anhydrase-related protein VIII (Car8) [59], calbindin [60], and calretinin [61] knockout animals. Collectively, these models point to alterations in Ca2+ homeostasis related to Purkinje cell physiology as causative for the ataxia. Junctophilins contribute to the formation of junctional membrane complexes, by spanning the endoplasmic/sarcoplasmic reticulum membrane and interacting with the plasma membrane, that function as structural platforms for Ca2+-mediated crosstalk between cell surface and intracellular channels. Car8 has no carbonic anhydrase enzymatic activity but was identified as a binding protein for the IP₃ receptor expressed in cerebellar PCs [62]. Thus, it is conceivable that Car8 plays a role in fine-tuning of the intracellular calcium level in the PC dendrites to optimize parallel fiber innervation and proper synaptic formation and/or maintenance in the cerebellum. Calbindin and calretinin both are calcium-binding proteins that turned out to be crucial in the shaping of intraneuronal Ca2+ fluxes like PC firings. In the cerebellar cortex of alert mice, calbindin or calretinin deficiency resulted in high-
frequency oscillations that emerged from interplay between synchronized PC assemblies and coupled interneurons in the cerebellar molecular layer [61].

Calcium, Phosphorylation, and Motor Coordination

Also the work on human disorders demonstrates that aberrant cerebellar Ca\(^{2+}\) signaling may result in cerebellar ataxia. For example episodic ataxia 2 (EA2) and spinocerebellar ataxia type 6 (SCA6) result from mutations in the pore-forming subunit of P/Q-type voltage-gated Ca\(^{2+}\) channels [57, 63, 64]. EA2 manifests as an episodic, fully reversible ataxia that only after decades may lead to persistent ataxia and mild cerebellar degeneration. Several other ataxia syndromes are directly linked to alterations in the neuronal phosphoproteome [63, 64] that ultimately may have bearing for cell membrane receptors and ion channels [65, 66]. In SCA12, for example, a repeat expansion in the regulatory subunit PR55/β alters phosphatase PP2A expression and, consequently, the phosphorylation status of target proteins (including MAPKs). And SCA14 mutations create a PKCγ superkinase that, in response to Ca\(^{2+}\) influx, readily phosphorylates brain glutamate receptors [65]. Thus, some SCA mutations apparently alter the phospho-status of neuronal Ca\(^{2+}\) channels and receptors and change their membrane localization, activity, and hence functioning in synaptic connectivity and plasticity [67]. The notion that MAPK signaling is a key process in synaptic plasticity [68] and that aberrant ERK1/2 activity has been noted in several neuropathologies [56, 69] nicely merges with the above.

Continued excessive stimulation of Ca\(^{2+}\)-permeable glutamate receptors or voltage-dependent Ca\(^{2+}\) channels (VDCCs) is expected to result in neuronal excitotoxicity. The chronic Ca\(^{2+}\) overshoot and concomitant inappropriate activation of Ca\(^{2+}\)-dependent processes in cells may then result in production of free radicals, altered mitochondrial function, and ultimately cell death. Apparently, Purkinje cells and cerebellar granule cells are quite vulnerable to such cell stress conditions, but it is believed that excitotoxicity may also contribute to other neurodegenerative illnesses like Alzheimer’s, Parkinson’s, and Huntington’s disease [70].

Concluding remarks

The picture emerges that aberrancies in cellular phosphorylation-dependent processes can deregulate, at the transcriptional and/or the posttranslational level, the calcium-ion-dependent communication circuitry of cerebellar neurons, thereby ultimately causing ataxias. In Fig. 3, we provide a simplified model in which the various lines of evidence are integrated. Activated growth factor receptors recruit distinct adaptor proteins and start up various signaling cascades, such as the PI3K, PLC\(^{\gamma}\), and MAPK pathways. Transient ERK activation via Ras will affect cytosolic targets that may well include transmembrane proteins regulating cellular Ca\(^{2+}\) concentration (e.g., glutamate-gated ion

![Fig. 3](image-url). Schematic overview of growth factor-induced and Ca\(^{2+}\)-releasing signaling pathways in neuronal cells. Growth factor (GF) binding to receptors (RTK) may activate PI3K, PLC\(^{\gamma}\), and MAPK signaling cascades. Transient ERK activation via Ras will affect cytosolic targets that may extend to transmembrane proteins involved in Ca\(^{2+}\) or glutamate signaling. RTK endocytosis may lead to Rap1-mediated sustained activation of ERK and altered transcription. PTPRR targets are depicted in blue. Proteins mutated in SCA6, SCA12, and SCA14 patients are in reddish purple. Proteins that are down-regulated in SCA1 mouse models and connect to IP\(_3\) receptor-regulated Ca\(^{2+}\) stores are in orange. Ca\(^{2+}\)-modulating proteins (CaBP\(_{\alpha}\)) and PTPRR, for which deficiency in mice led to ataxic behavior but not neurodegeneration, are in green.
channels and VDCCs) and membrane potential (e.g., Kv4.2). Following receptor endocytosis, Rap1-mediated sustained activation of ERK will lead to alterations in gene expression. PTPRR may indirectly influence such transcriptional events by inactivating ERK, but could also, much like the proteins mutated in SCA12 and SCA14 patients, directly dephosphorylate and thus regulate relevant cytosolic proteins or even ion channels and receptors. The net effect on Ca²⁺ levels thus may resemble that caused by EA2 mutations in VDCCs.

Many dominant ataxias (i.e., SCA1) are caused by an expanding polyglutamine sequence in the responsible protein [63, 64]. In a mouse model for SCA1, the metabotropic glutamate receptor and its auxiliary proteins that connect to IP₃ receptor-regulated intracellular Ca²⁺ stores were all found to be down-regulated [71], providing yet another link to aberrant calcium handling in ataxias. Further support comes from the mouse models that are deficient in Ca²⁺ binding or Ca²⁺ modulatory proteins [58–61], which all display ataxic behavior without neurodegeneration. Since PTPRR-deficient mice combine a lack of apparent brain morphological abnormalities with mild ataxic symptoms and hyperactivity of ERK1/2 MAP kinases, further studies towards the underlying mechanism will have bearing for the field of human cerebellar ataxia-type locomotive disorders. This may yield novel candidate SCA genes for the many hereditary cerebellar ataxias that still await disease gene identification and may provide clues to symptomatic or neuroprotective strategies that can be applied in early stages of degenerative ataxias.

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