CA8 Mutations Cause a Novel Syndrome Characterized by Ataxia and Mild Mental Retardation with Predisposition to Quadrupedal Gait

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Abstract

We describe a consanguineous Iraqi family in which affected siblings had mild mental retardation and congenital ataxia characterized by quadrupedal gait. Genome-wide linkage analysis identified a 5.8 Mb interval on chromosome 8q with shared homozygosity among the affected persons. Sequencing of genes contained in the interval revealed a homozygous mutation, S100P, in carbonic anhydrase related protein 8 (CA8), which is highly expressed in cerebellar Purkinje cells and influences inositol triphosphate (IP(T)) binding to its receptor IP3R1 on the endoplasmatic reticulum and thereby modulates calcium signaling. We demonstrate that the mutation S100P is associated with proteasome-mediated degradation, and thus presumably represents a null mutation comparable to the CA8 mutation underlying the previously described waddles mouse, which exhibits ataxia and appendicular dystonia. CA8 thus represents the third locus that has been associated with quadrupedal gait in humans, in addition to the VLDLR locus and a locus at chromosome 17p. Our findings underline the importance of ITP-mediated signaling in cerebellar function and provide suggestive evidence that congenital ataxia paired with cerebral dysfunction may, together with unknown contextual factors during development, predispose to quadrupedal gait in humans.

Introduction

The hereditary ataxias comprise a diverse groups of disorders characterized by loss of balance and coordination. They are classified as autosomal recessive, autosomal dominant, X-linked, and mitochondrial, and are clinically and pathogenetically diverse [1,2]. The identification of genes associated with individual forms of ataxia has gone a long way towards the identification of factors responsible for development, homeostasis, and function of the cerebellum, which is the organ primarily affected in most forms of ataxia.

Recently, a form of ataxia characterized by quadrupedal gait has been described in several families. A syndrome of nonprogressive cerebellar ataxia and mental retardation associated with inferior cerebellar hypoplasia and mild cerebral gyral simplification was initially identified in patients with a disorder termed “dysequilibrium syndrome” (MIM 224050). This disorder was found to be due to a 199 kb deletion on chromosome 9p24 encompassing all of the very low-density lipoprotein receptor gene (VLDLR) and part of the poorly characterized gene LOC401491 [3]. Homozygous premature-truncation-codon (PTC) mutations in VLDLR were subsequently found to be associated with severe ataxia, cerebellar hypoplasia, dystarthisia, and severe mental retardation. Affected persons walked on all four extremities. Although the affected persons could stand upright and even walk bipedally, they preferred quadrupedal walking [4,5]. An unrelated Iranian family with a PTC mutation in VLDLR was subsequently reported in which affected persons had mental retardation, strabismus, short stature, disturbed equilibrium, and walking disability, but no tendency towards quadrupedal gait [6]. A further unrelated family with cerebellar hypoplasia, mental retardation, and quadrupedal gait demonstrated linkage to a locus on chromosome 17p [7].

The observation that some, but not all, VLDLR mutation carriers walk on all four extremities raised the question of whether quadrupedal gait is a functional adaptation that can be seen in congenital ataxia syndromes depending on unknown internal or external influences. Although many mammals and other animals can stand or walk bipedally for shorter or longer periods of time, the manner in which humans do so is unique to our species. Our upright posture is the result of constant skeletalmuscular adjustments of posture. In contrast, other primates make frequent use of “holds” during locomotion. In effect, we are always a second or two away from falling down. Human bipedalism and higher cognition are closely integrated. Especially running places high requirements on cognition, such as the need to incorporate...
Author Summary

We identified a homozygous missense mutation (S100P) in the gene encoding carbonic anhydrase VIII in a consanguineous Iraqi family in which affected siblings had mild mental retardation and congenital ataxia characterized by quadrupedal gait. The affected persons walk on their hands and feet with their legs held straight with a “bear-like” gait. Our results show that the mutation S100P induces proteasome-mediated degradation with a severe reduction of the level of Ca8 protein. The waddles (wdl) mouse, a spontaneous animal model with ataxia, was previously shown to harbor a 19-bp deletion in Ca8 that leads to an almost complete lack of detectable Ca8 protein, resulting in abnormalities in cerebellar synaptic transmission. Therefore, we speculate that the reduction in Ca8 protein concentration associated with the S100P mutation could result in similar pathophysiological effects. With the current report, alterations at three gene loci (CA8, VLDLR, and a yet-to-be discovered gene on chromosome 17p) have been reported to be associated with quadrupedal gait. It is unknown whether quadrupedal gait is related to specific molecular abnormalities or is an adaptive response to ataxia in some circumstances. However, we note that ataxia associated with mutations at all three loci is congenital and also associated with mental retardation, which is not generally a feature of other hereditary ataxias.

information about uneven parts of the ground ahead. Nonhuman primates such as gorillas can only run for short distances, and human children do not develop adult competence in walking and running until about the age of seven years [8].

In the current work, we report on mutations in the Ca8 gene in a consanguineous family from Iraq. Affected family members displayed a syndrome of ataxia and mild mental retardation (AMMR) and ambulate on all four extremities (quadrupedal gait). In contrast to the previously reported cases associated with VLDLR and a yet-to-be discovered gene on chromosome 17p, affected persons have only mild mental retardation. Including Ca8 as described in the present report, there are now three gene loci, mutations in which are associated with ataxia and cerebral defects, whereby some, but not all affected persons display quadrupedal gait. We therefore suggest that congenital ataxia together with cerebral deficits may, together with other, currently unknown external or internal factors, predispose to quadrupedal gait in humans.

Results

We identified an Iraqi family with a syndrome characterized by ataxia and mild mental retardation (AMMR). The healthy parents are first cousins, and four of eight sibs are affected. The parents claimed that the affected persons never learned to crawl on their hands and feet with their legs held straight with a “bear-like” gait. They also claimed that attempts to teach the children to walk on two legs with crutches or with other supports failed. Our own observations confirm that four of the affected children as adults are predominantly quadrupedal (see Video S1). They walk with straight legs, placing weight on the palms of their hands. Although the affected are able to walk on two legs for several steps, they tend to tumble into a quadrupedal position quickly, complain of lack of balance and occasionally fall from a sitting position. Affected persons were noted to have mild mental retardation and dysarthric, slurred speech, but there were no other symptoms such as retinopathy or pyramidal signs. Table 1 provides an overview of the clinical features observed in the four affected siblings encoded using terms of the Human Phenotype Ontology [9].

We used a genome-wide linkage approach to identify the genetic basis of AMMR in this family. We established significant genetic linkage to a 6-centiMorgan interval on chromosome 8q12 with a lod score of 3.01 (Figure 1). This interval spans nucleotides 50,801,724–64,607,419 on chromosome 8 (March 2006, UCSC hg18 assembly) and contains 17 protein-coding genes, including an obvious candidate, carbonic anhydrase-related protein VIII (Ca8), which is abundantly expressed in cerebellar Purkinje cells [10]. Carbonic anhydrases are a family of monomeric zinc metalloenzymes that catalyze the reversible hydration of CO2. However, Ca8 lacks one of the three histidine residues required for binding to the zinc ion and thus has no catalytic carbonic anhydrase activity [11]. The waddles (wdl) mouse is a spontaneous animal model with ataxia and appendicular dystonia without morphological abnormalities in the central or peripheral nervous system. The wdl mouse was shown to harbor a 19-bp deletion in Ca8 that leads to rapid degradation of mutant Ca8 mRNA and to the almost complete lack of detectable Ca8 protein [12]. Although the morphology of the cerebellum appeared normal by confocal microscopy [12], subsequent investigations revealed abnormalities of parallel fiber-Purkinje cell synapses in the cerebellum together with defects in excitatory transmission [13].

We therefore sequenced all 8 exons of Ca8 and identified the homozygous transition c.298T>C, predicted to lead to the substitution of a serine by a proline residue (S100P, in all affected individuals (Figure 2), c.298T>C segregated correctly with the disease in the family and was not found in 200 population-matched controls, thus making a previously undescribed polymorphism unlikely. In addition, we found no other nonsynonymous sequence changes in any of the other 16 genes in the mapping interval, including the TTPA gene, mutations in which are found in ataxia with vitamin E deficiency [14] (AVED). The sole known biochemical function of Ca8 is to inhibit inositol 1,4,5-triphosphate (IP3) binding to IP3 receptor 1 (ITPR1) [15]. ITPR1 plays a critical role in the modulation of intracellular calcium (Ca2+) signaling [16]. Like Ca8, ITPR1 is also highly expressed in Purkinje cells. Mutation of ITPR1 underlies ataxia in mice and mutations in ITPR1 have been identified in spinocerebellar ataxia 15 in humans [17]. Interestingly, mutant ataxin-3, which is the protein product of the gene mutated in spinocerebellar ataxia type 3 (SCA3), was shown to bind to ITPR1 and thereby cause a destabilization of neuronal calcium signaling [18]. These observations suggest that disturbances of ITPR1-mediated calcium signaling may be an important and common phenomenon in hereditary ataxias. We therefore investigated whether S100P affects Ca8-ITPR1 binding using recombinant full-length Ca8 and a region of ITPR1 containing the known Ca8-binding domain [15]. No detectable difference was observed between wildtype and mutant Ca8 constructs using blot-overlay assays (Figure 3). Therefore, since the wdl mouse displays a hypomorphic Ca8 mutation [12], we hypothesized that S100P might affect protein stability.

HEK cells were stably transfected with Flag-tagged wildtype and S100P mutant Ca8 constructs using a system that allows single-copy targeted integration of a tetracyclin-inducible construct. This allowed us to compare the stability of mutant and wildtype Ca8 mRNA and protein at several different levels of expression. Following selection of stable clones, the expression of wildtype and mutant Ca8 was examined using quantitative realtime PCR, and the quantity of wildtype and mutant protein was examined by Western blots. Whereas mRNA levels for the mutant Ca8 were not different from those of the wildtype Ca8 gene product, using the Western blot detection system we observed that in HEK cells, the amount of Ca8 protein was reduced by about one-fifth in the presence of mutant S100P Ca8 compared with wildtype Ca8. A more sensitive way of comparing the stability of wildtype and mutant Ca8 constructs was to use a system that allows measurement of real-time Ca8 protein synthesis. Expression of wildtype Ca8 and mutant Ca8 was studied in HEK cells in which a Ca8 construct was expressed under the control of a inducible expression system. We confirmed that the amount of Ca8 protein produced in HEK cells is independent of the level of protein synthesis, thereby allowing us to perform quantitative analysis of Ca8 synthesis in the presence of wildtype and mutant Ca8 from transcription to translation. Consistent with our observation that there is a reduction of wildtype Ca8 protein in the presence of mutant S100P Ca8, we observed a reduction of wildtype Ca8 protein synthesis in HEK cells expressing wildtype Ca8 and mutant S100P Ca8 compared with wildtype Ca8 alone.
Table 1. Clinical features of the affected persons of the family presented in this work and summary of features found related to mutations in VLDLR and at a locus on chromosome 17p.

| HPO term                                          | CA8   | VLDLR | 17p   |
|--------------------------------------------------|-------|-------|-------|
|                                                   | S100P | ns/fs | unknown |
| Cerebellar ataxia (HPO:0001253),                  | Q     | Q/B   | Q/B   |
| Cerebellar ataxia associated with quadrupedal gait (HP:0009878) |       |       |       |
| Dysarthria (HP:0001260)                           | +     | + (or no speech) | +     |
| Mental retardation, mild (HP:0001256)            | Mild  | Moderate/Severe | Severe |
| Mental retardation, moderate (HP:0002342)        |       |       |       |
| Mental retardation, severe (HP:0001261)          |       |       |       |
| Tremor (HP:0001337)                              | +     | +/-   | +     |
| Seizures (HP:0001250)                            | –     | very rare | +/-  |
| Strabismus (HP:0000486)                          | +/-   | +     | +     |
| Cerebellar hypoplasia (HP:0001321)               | n/a   | +     | +     |
| Cortical gyral simplification (HP:0009879)       | n/a   | +     | +     |
| Corpus callosum hypoplasia (HP:0002079)          | n/a   | –     | +     |

The features are coded using terms from the Human Phenotype Ontology [9]. Abbreviations: Q: quadrupedal gait (HP:0001253); B: cerebellar ataxia with bipedal gait (annotated as HP:0001253); Q/B: some affected show quadrupedal gait, some bipedal; Mild: mild mental retardation (HP:0002342), Moderate: moderate mental retardation (HP:0002342); severe: severe mental retardation (HP:0001261). +: present; --: absent; +/-: present in some, absent in other affected persons. ns/fs: information summarized for nonsense/frameshift mutations in VLDLR R257X [4], c2339delT [4,5] and R448X [6]. Information for the 17p locus summarized from ref. [7]. n/a: information not available.

doi:10.1371/journal.pgen.1000487.t001

Discussion

The cerebellum is a complex neurological structure, containing more than half of the brain’s total number of neurons. Cerebellar networks show long-term synaptic plasticity, which indicates that experience-dependent adaptive and learning processes are a salient feature of cerebellar function. Most afferent information enters the cerebellum via climbing fibers (CF) and mossy fibers, which excite the Purkinje cells indirectly through the parallel fiber (PF) pathway. Binding of CA8 to ITTPR1 inhibits IP3 binding to ITTPR1 by reducing the affinity of the receptor for IP3 [15]. This is one of several factors that modulate the ability of ITTPR1 to rapidly release calcium stores from the endoplasmatic reticulum [19]. Mice with mutations in either Itp3r1 or Ca8 do not display cerebellar atrophy, but rather both show neurophysiological defects [20,21]. Modulation of intracellular calcium is important for a number of cerebellar functions such as long-term depression [22]. Therefore, we speculate that the consequences of a CA8 mutation may involve improper modulation of the ITTPR1 with resultant functional and/or developmental defects in the cerebellum.

S100P leads to a proteasome-mediated reduction in protein stability in our in vitro assay. Loss of protein stability is a common mechanism of missense mutations associated with human disease [23]. We suggest that it is plausible that the mutation leads to a reduction in the amount of CA8 in the cerebellum of the affected individuals, which might lead to a similar defect as that observed in the waddles mouse, in which Ca8 is nearly undetectable in cerebellum owing to a 19 bp deletion in the gene [12].

Mutations at three loci (CA8, VLDLR and the yet-to-be-discovered gene at 17p) have now been found to be associated with quadrupedal locomotion in humans, although not in all affected individuals. The clinical picture of the disorders associated with mutations at the three loci is similar but shows important differences. The affected persons of the family described in this work showed a relatively mild degree of mental retardation. Seizures have not been observed in the family described in this study, but did represent a characteristic clinical feature of the family in whom linkage to a locus on chromosome 17p was reported [7]. Table 1 gives an overview of the clinical features encountered in the three forms of ataxia which have been observed with quadrupedal gait. Given the variable incidence of quadrupedalism in individuals with mutations in the same gene, we think it probable that contextual factors during development – either internal or external – contribute to this particular phenotypic outcome [24]. As one possibility, we note that ataxia associated with mutations at all three loci is congenital and also associated with mental retardation, which is not generally a feature of other hereditary ataxias, such as Joubert syndrome [25] or AVED [14]. Thus, perhaps it is only when congenital ataxia is coupled to a certain kind of malfunction of the cerebral cortex that individuals are likely to remain walking on all fours.

Materials and Methods

Written informed consent was obtained from the subjects to publish details of their case and family history including the photograph and video documentation.

Genome-Wide Linkage Analysis

The family was genotyped using the Affymetrix GeneChip® Human Mapping 10 K Array. The maximum expected lod score for a first cousin marriage with 4 affected children is approximately 3.01. The two-point and multipoint linkage analyses (Genehunter [26], Allegro [27] and Merlin [28]) were performed assuming a fully penetrant autosomal recessive trait with a disease
frequency of 0.001 and no phenocopies. Analyses were performed using the easyLINKAGE and easyLINKAGE-Plus tools [29,30].

Recombinant Constructs
All PCRs were performed using High-Fidelity Taq Polymerase (Invitrogen). Two large amplicons comprising the full-length human \textit{CA8} transcript and a region of \textit{ITPR1} containing the known \textit{CA8}-binding domain [15] (amino acids 1387–1647) of \textit{ITPR1} were amplified by PCR from human fibroblast cDNA. A \textit{CA8} amplicon corresponding to positions +91 to +1190 (numbering based on GenBank entry AY075022) was amplified using the primers 5'-tgcactcacactgcggttca-3' (f) and 5'-aagggcattataggacactcact-3' (r). An \textit{ITPR1} amplicon corresponding to positions +4361 to +5270 (numbering based on GenBank entry L38019) was amplified using the primers 5'-ctcatgtaccacatccactt-3' (f) and 5'-ccttaatgcagagcttctct-3' (r).

Figure 1. Abbreviated pedigree showing the inferred haplotypes of the parents and the four affected children. The SNPs A-1511204 and A-1509289 are homozygous in the mother and therefore noninformative for the maternally derived recombination in this region. Therefore, these markers were not fully informative with respect to the critical recombination on the maternally inherited haplotype in affected individual II-1.

doi:10.1371/journal.pgen.1000487.g001

CA8 Mutations and AMMR Syndrome
We first produced CA8 and ITPR1 constructs for investigation of protein-protein binding. A recombinant CA8 construct corresponding to the complete coding region of human CA8 was then amplified by nested PCR with the primers 5'-gtatGGATCCgatggcggacctgagcttca-3' (f) and 5'-cttgCTCGAGtattcctactctacggtggtgct-3' (r). The resulting fragments were subcloned into the plasmid pFastBac HT A (Invitrogen) using the BamH1 and XhoI restriction sites (capitalized in the primer sequences). pFastBac HT A adds a hexa-histidine tag and an rTEV protease cleavage site to the N-terminus of the expressed protein.

The c.298T>C (S100P) mutant CA8 construct was prepared by using the GeneTailor Site-Directed Mutagenesis kit (Invitrogen) using the primers 5'-tcaaaatcagttcttCcgggaggaccattgc-3' (f) and 5'-aagaactgattttgacttcaggataacctga-3' (r) according to the manufacturer's protocol.

Additional Flag-tagged CA8 constructs were produced for investigation of the influence of S100P on protein stability. PCR amplification of the full-length wild-type or S100P mutant CA8-pFastBac HT A constructs was used to subclone to the BamH1 and XhoI sites of pcDNA3 (Invitrogen) using the primers 5'-gtatGGATCCgatggcggacctgagcttca-3' (f) and 5'-cttgCTCGAGtattcctactctacggtggtgct-3' (r). The reverse primer introduces the Flag tag sequence DYKDDDDK to the C-terminus of the CA8 protein. pcDNA5/FRT/TO Flp-In constructs were then obtained by subcloning Flag-CA8(WT) and Flag-CA8(S100P) from pcDNA3.1 into the pcDNA5/FRT vector using the BamH1 and XhoI restriction sites. All resulting recombinant vectors were verified by sequencing.

Figures 2 and 3. CA8 Mutations in two families with ataxia and mild mental retardation. (A) Affected members of an Iraqi family displayed quadrupedal gait. (B) Sequence traces of the mutation c.298T>C (S100P) in CA8 and a wildtype control (wt). (C) Sequence logo of an alignment of CA8 orthologs from 12 species from human to sea urchin showing the sequence affected by S100P. Position 100 does not show a high degree of conservation.

Similarly, a recombinant ITPR1 was amplified with the primers 5'-gtatGGATCCgaagaatgtctacacagaga-3' (f) and 5'-cttgCTCGAGctactgaaatgcagctctaa-3' (r). The resulting fragments were subcloned into the plasmid pFastBac HT A (Invitrogen) using the BamH1 and XhoI restriction sites (capitalized in the primer sequences). pFastBac HT A adds a hexa-histidine tag and an rTEV protease cleavage site to the N-terminus of the expressed protein.

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Expression of Recombinant Human CA8 Protein and the CA8 Binding Domain of ITPR1

Spodoptera frugiperda (Sf9) cells were maintained in serum-free Insect-Xpress medium (Biowhittaker) supplemented with 5% fetal bovine serum, 10 U/ml penicillin, and 10 μg/ml streptomycin, and cultured at 27°C. The production of the protein was performed according to the protocol of the Bac-to-Bac Baculovirus expression system (Invitrogen). Briefly, the recombinant pFastBac HT A vectors were transformed into E. coli DH10Bac cells (Invitrogen). The isolated positive recombinant bacmids were verified by PCR and used to transfect Sf9 insect cells for viral particle formation. Following three rounds of amplification of baculovirus, the virus titer was determined by plaque assay.

Small scale time-course expression experiments were conducted in 6 well plates to optimize the multiplicity of infection (MOI) and expression time. Bands corresponding to the size of recombinant protein were visualized on Western blots by detection with monoclonal anti-His6 antibody (Novagen). For large-scale production of protein, 5 × 10^8 Sf9 cells were infected at an MOI of 1 plaque-forming unit (pfu)/cell with recombinant baculovirus. After about 48 hours cells were harvested and resuspended in 10 ml lysis buffer (10 mM Tris-HCl, 10 mM Imidazol, 25 mM EDTA free protease inhibitor cocktail (Sigma-Aldrich), 0.1 mM PMSF), sonication was performed on ice for 2 × 15 s at 50% maximum energy output and was cleared by centrifugation at 12,000 rpm for 30 min. Purification of the recombinant protein was performed as described before [31]. Concentration of the recombinant polypeptide was determined with the BCA protein assay (Pierce) according to the manufacturer’s instructions.

Blot Overlay Assay

For the blot overlay assay, recombinant wildtype and S100P mutant CA8 were separated by sodium dodecyl sulfate-12% polyacrylamide gel electrophoresis (SDS-PAGE) and then transferred onto a polyvinylidene difluoride (PVDF) membrane (Immobilon-P transfer membrane, Millipore). A recombinant fibrillin-1 fragment [32] was used as a control. Non-specific binding sites were blocked with dilution buffer (ProFound Far-Western Biotinylated Protein-Protein Interaction Kit, Pierce), according to the manufacturer’s instructions, and incubation was performed for 2 h at room temperature with 2 μg/ml of the biotinylated ITPR1 fragment in dilution buffer and was followed by washing the PVDF membrane three times with PBS including 0.025% Tween 20. For detection, incubation was performed with streptavidin-horseradish peroxidase conjugate for 3 h at room temperature. Membranes were washed six times in 1× PBS including 0.025% Tween 20. Membranes were then incubated in the UnBlot substrate working solution (ProFound Far-Western Biotinylated Protein-Protein Interaction Kit, Pierce). Hyperfilm ECL chemiluminescence films (Amersham Pharmacia) were exposed according to the manufacturer’s instructions.

Generation of Stable Flip-In T-REx 293 Cells Expressing Flag-CA8(WT) and Flag-CA8(S100P)

To generate stable tetracycline-inducible Flip-In T-REx cells, 0.5 μg pcDNA/FRT/TO encoding Flag-CA8(WT) or Flag-CA8(S100P) or empty pcDNA/FRT/TO and 1 μg pOG44 were cotransfected into Flp-In T-REx 293 cells (1 × 10^6) using Lipofectamine 2000 (Invitrogen) according to the manufacturer’s directions. 48 hours after transfection cells were reseeded at less than 25% confluence, after 4 hours, the medium was changed to medium supplemented with 100 μg/ml hygromycin B (Invitrogen) and 15 μg/ml blasticidin. After two weeks, hygromycin-resistant colonies were picked and sub-cultured. Selection of positive colonies was performed by immunoblotting as described below.

Detection of mRNA by Quantitative RT-PCR

Tetracycline inducible Flip-In T-REx 293 cells were grown in the absence or presence of 1 μg/ml tetracycline for 24 hours, total
RNA of those cells were extracted with NucleoSpin RNA II (Macherey-Nagel, Duren, Germany) according to the manufacturer’s protocol. 1 mg RNA from cells was reverse-transcribed using the Superscript First Strand synthesis kit (Invitrogen). Quantitative RT-PCR was performed on an ABI Prism 7500 Sequence Detection System using SYBR-Green PCR Master Mix at 50°C for 2 min, 95°C for 10 min, then 40 cycles at 95°C for 15 s, and at 60°C for 1 min. The forward primer for CA8 is complementary to a sequence at the C-terminal region of CA8, 5’-cttgtggaaggctgtgatgg-3’, and the reverse primer is complementary to a sequence of the Flag tag, 5’-ttgtcatcgtcgtccttgtag-3’. The primer pair for amplification of GAPDH was: 5’-ctcaacgaccacatgggtcagctca-3’, 5’-ggctcattctctggaggccatgtg-3’.

Immunoblotting to Detect Flag-CA8 Proteins
To detect the expression of CA8-Flag WT and S100P protein, tetracycline-inducible Flip-in T-REx cells seeded in 24 well plate (2 x 10⁶) were induced by 0.1 μg/ml or 1 μg/ml tetracycline (Sigma-Aldrich). To inhibit the proteasome, cells were incubated in proteasome inhibitor MG132 (Sigma-Aldrich) at different concentrations (0.4 μM, 2.0 μM) in the presence of 1 μg/ml tetracycline. After 24 hours cells were washed twice with PBS and lysed with 100 μl of 1X SDS-sample buffer (50 mM Tris-HCL pH 6.8, 10% glycerol, 2% SDS, 1% 2-mercaptoethanol, 0.02% Bromphenol blue). After brief sonication on ice, 10 μl aliquots were separated on 12% SDS-gel and electrophoretically transferred to a PVDF membrane. Immunodetection was performed by using mouse monoclonal anti-Flag (1:2000, Sigma-Aldrich) as primary antibody, followed by subsequent incubation with HRP (horseradish peroxidase)-conjugated rabbit anti-mouse secondary antibody (1:2000, Calbiochem). Parallel Western blots were probed with a mouse anti-GAPDH antibody (1:5000, Abcam) as a loading control. Protein expression was quantified using the ImageJ program [33].

CA8 Sequence Logos
A CA8 protein multiple alignment from Human (Homo sapiens; NP_004047), dog (Canis familiaris; XP_544094), cow (Bos taurus; NP_001077159), horse (Equus caballus; XP_001496523), mouse (Mus musculus; NP_031618), rat (Rattus norvegicus; NP_001009662), opossum (Monodelphis domestica; XP_001368351), chicken (Gallus gallus; XP_419221), frog (Xenopus (Silurana) tropicalis; NP_001011213), trout (Oncorhynchus mykiss; NP_001118116), zebrafish (Danio rerio; NP_001017571), and sea urchin (Strongylocentrotus purpuratus; XP_795365) was prepared with ClustalX [34] and sequence logos were prepared using texshade [35].

Supporting Information
Video S1 Film demonstrating quadrupedal gait of one of the families described in this manuscript. Found at: doi:10.1371/journal.pgen.1000487.s001 (1.94 MB WMV)
Acknowledgments

The authors wish to thank Uwe Kornak for help with the protein stability assays. The authors acknowledge additional support by the Deutsche Forschungsgemeinschaft (RO 2005/4-1).

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Author Contributions

Conceived and designed the experiments: ST GG NH SM PNR. Performed the experiments: GG MG CM. Analyzed the data: ST GG MG KH AK PNR. Contributed reagents/materials/analysis tools: AJA NH. Wrote the paper: ST GG PNR.