Compound heterozygous \textit{PLA2G6} loss-of-function variants in Swaledale sheep with neuroaxonal dystrophy

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\textbf{Abstract}
Sporadic occurrences of neurodegenerative disorders including neuroaxonal dystrophy (NAD) have been previously reported in sheep. However, so far no causative genetic variant has been found for ovine NAD. The aim of this study was to characterize the phenotype and the genetic aetiology of an early-onset neurodegenerative disorder observed in several lambs of purebred Swaledale sheep, a native English breed. Affected lambs showed progressive ataxia and stiff gait and subsequent histopathological analysis revealed the widespread presence of axonal spheroid indicating neuronal degeneration. Thus, the observed clinical phenotype could be explained by a novel form of NAD. After SNP genotyping and subsequent linkage mapping within a paternal half-sib pedigree with a total of five NAD-affected lambs, we identified two loss-of-function variants by whole-genome sequencing in the ovine \textit{PLA2G6} gene situated in a NAD-linked genome region on chromosome 3. All cases were carriers of a compound heterozygous splice site variant in intron 2 and a nonsense variant in exon 8. Herein we present evidence for the occurrence of a familial novel form of recessively inherited NAD in sheep due to allelic heterogeneity at \textit{PLA2G6}. This study reports two pathogenic variants in \textit{PLA2G6} causing a novel form of NAD in Swaledale sheep which enables selection against this fatal disorder.

\textbf{Keywords} \textit{Ovis aries} · Neurogenetic disorder · Rare disease · Compound heterozygosity · Precision medicine

\section*{Introduction}
Neuroaxonal dystrophy (NAD) comprises a clinically and genetically heterogeneous group of neurodegenerative diseases of central nervous system (Hayflick et al. 2018). It is characterized by progressive signs of neurological dysfunction including ataxia, hypermetria, proprioceptive deficits, head incoordination and tremors. The characteristic
neuropathological changes involve formation of axonal swellings (spheroids) in specific regions, usually relay nuclei, of the brainstem and spinal cord (Sisó et al. 2006). In humans, this condition is usually described as neurodegeneration with brain iron accumulation (NBIA) or infantile neuroaxonal dystrophy (INAD; OMIM PS234200) and is most frequently monogenically recessively inherited. The genetic basis is described for ~ 85% of patients diagnosed with NBIA and involves eight genes (Hayflick et al. 2018). Human NBIA2A (INAD1) and NBIA2B are associated with autosomal recessive variants in the PLA2G6 gene (OMIM 256600). Phospholipase A2 group VI gene (PLA2G6) encodes 85/88 kDa calcium-independent phospholipase A2, an enzyme that catalyzes the hydrolysis of phospholipids to produce free fatty acids, and has a critical role in cell membrane homeostasis (Baburina and Jackowski 1999).

In veterinary medicine, sporadic cases of NAD have been previously described in sheep (OMIA 000715-9940), as well as cattle (Hanshaw et al. 2015), dogs (OMIA 000715-9615), cats (OMIA 000715-9685), rabbits (OMIA 000715-9986), and horses (OMIA 000715-9796). Although, there is a variation in the age of affected lambs in the previous reports of ovine NAD cases, the clinicopathological features are comparable in all the different sheep breeds including Merino (Harper and Morton 1991; Kessell et al. 2012), Suffolk (Cordy et al. 1967), Romney (Nuttall 1988), Perendale (Nuttall 1988), and Coopworth (Nuttall 1988), as well as multiple crossbred Merino-Border Leicester×Polled Dorset lambs (Finnie et al. 2014; Hawes et al. 2017). However, the underlying genetic variants to date have been characterized only in dogs, representing four breed-specific autosomal recessive variants in MFN2 (Fyfe et al. 2011), TECPR2 (Hahn et al. 2015), PLA2G6 (Tsuboi et al. 2017), and VPS11 (Lucot et al. 2018) genes.

The aim of this study was to characterize the phenotype and the genetic etiology of an early-onset neurodegenerative disorder observed in several lambs of purebred Swaledale sheep, a native English breed. Herein we present evidence for the occurrence of a familial novel form of recessively inherited NAD due to two compound heterozygous loss-of-function variants in ovine PLA2G6, which enables selection against this fatal disorder.

Methods

Animals

This study did not require official or institutional ethical approval as it was not experimental. A total of 71 purebred Swaledale sheep from one flock were used in this study, including 52 ewes, 1 ram, 5 NAD-affected lambs, and 13 apparently normal offspring. All lambs were sired by the same ram and also shared the same maternal grandsire. Blood samples were taken from all available animals. Additionally, tissue samples (brain and spinal cord) were collected postmortem from 2 representative cases for further analyses. Genomic DNA was isolated from blood using the Maxwell RSC whole blood DNA kit (Promega).

Neuropathology

Samples of brain and spinal cord from affected lambs were fixed in 10% neutral-buffered formalin for histopathological analysis. Representative samples of brain including cerebral cortex (frontal, parietal, temporal and occipital lobes), striatum, hippocampus, rostral and caudal thalamus, rostral and caudal midbrain, cerebellum (vermis and hemisphere), rostral and caudal medulla and spinal cord including cervical segments 3 and 7, thoracic segment 7 and lumbosacral intumescence were routinely processed to 5 µm haematoxylin-and-eosin (HE)-stained sections for histopathological examination. Selected affected regions were immunolabelled for amyloid precursor protein (APP) as described previously (Garcia et al. 2015).

SNP array genotyping and analyses

Genomic DNA of 14 sheep (5 ewes, 1 ram, 5 NAD-affected, and 3 unaffected lambs) was genotyped on the Illumina OvineSNP50 BeadChip array. All genome positions refer to the ovine reference genome assembly Oar_rambouillet_v1.0. PLINK v1.9 software (Chang et al. 2015) was used for quality control of the genotyping data, parentage confirmation, as well as autozygosity mapping. All samples had good quality as indicated by > 90% call rate per individual. The final dataset included 44 066 SNP markers after pruning based on missing genotype calls per marker (> 10%). Non-parametric linkage analysis was carried out using the Merlin software (Abecasis et al. 2002) to test for co-segregation of any chromosomal regions and the NAD phenotype.

Whole-genome sequencing and variant calling

Whole-genome sequencing data of two parent–offspring trios (the sire, two dams, two NAD-affected lambs) were obtained after the preparation of a PCR-free fragment library at an average 21.5 × coverage. Fastq-files were mapped to the ovine reference genome assembly Oar_rambouillet_v1.0. Variant calling of single nucleotide and small indel variants was performed using NCBI annotation release 103 to predict their functional effects as described before (Paris et al. 2019). Private protein-changing variants present in the two NAD-affected animals were identified by comparison with 60 publically available control genomes, designated...
as local control cohort, which were produced during other ongoing projects of our group (Online Resource 3). In addition, the Sheep genomes project variant database of further 453 samples (Naval-Sanchez et al. 2018) available from the European Nucleotide Archive (ENA), herein designated as a global control cohort, was searched for the presence of identified variants. The Integrative Genome Viewer (Thorvaldsdóttir et al. 2013) was used for confirmation of the identified sequence variants and for visual inspection to exclude any structural variants in the critical region.

**RNA isolation and RT-PCR**

RNA was extracted from samples of the spinal cord, sciatic nerve, and cerebellum of three NAD-affected lambs using the RNeasy Fibrous Tissue Mini Kit (Qiagen). The tissue was first finely crushed by mechanically using TissueLyser (Qiagen), and RNA was extracted by centrifugation following the instructions of the manufacturer. Total mRNA was reverse-transcribed into cDNA using the SuperScript IV Reverse Transcriptase kit (Thermo Fisher Scientific) with oligo d(T) primers. RT-PCR was carried out using primers spanning the different exons’ boundaries (Additional File 5) of the PLA2G6 gene, and the sequences of the RT-PCR products were obtained by Sanger sequencing as described below. ACTB was included as a reference gene control (Additional File 5).

**PCR and targeted genotyping**

Sanger sequencing was used to confirm the WGS results and to perform targeted genotyping for the identified PLA2G6 variants as well as to sequence the obtained RT-PCR products. Primers were designed using the Primer-BLAST tool (Ye et al. 2012). After amplification with AmpliTaqGold-360Mastermix (Thermo Fisher Scientific), the purified PCR products were directly sequenced on an ABI3730 capillary sequencer (Thermo Fisher Scientific). The sequence data were analyzed using Sequencher 5.1 software (GeneCodes). All primer sequences are available in Additional File 5.

**Results**

**Phenotype**

In May 2017, five 6-week-old purebred Swaledale lambs presented with ataxia and a stiff gait, which over a period of days progressed to tremor, lateral recumbency, paddling and nystagmus (Additional File 1). The parents of all available affected animals were healthy. The lambs were part of a 300-ewe Swaledale flock of which every year 60 Swaledale shearlings (18 month old females) were all bred to one Swaledale ram, and all older sheep were bred to a Blue-faced Leicester ram to produce North Country Mules for sale as breeding sheep. The Swaledale ram was changed every 2 years so that he did not mate his own daughters. Thus, all Swaledale shearlings were also sired by the same ram (the previous one, which was the maternal grandsire of the affected lambs). Clinical signs were confined to the purebred Swaledale lambs, in which group an estimated 15 NAD-affected lambs were seen from a total of 110 lambs. Some lambs suspected to have been affected were found dead having fallen into a ditch. Lambs of both genders were affected. No clinical signs were seen at any time in crossbred lambs. After mating (in November 2016), all sheep were managed as one group until lambing, which occurred outside in April 2017. Border disease virus was not detected by PCR in any of the affected lambs.

**Neuropathology**

Neuropathological analysis detected widespread spheroid formation predominantly involving brainstem grey matter, including accessory cuneate, olives, rostral colliculus, lateral geniculate body, caudal colliculus, medial geniculate body, and oculomotor nuclei (Fig. 1a). Spheroid formation was also present in the cerebellar nuclei, cortex with proximal Purkinje axonal spheroid (torpedo) formation and variable smaller spheroid formation in the internal granule cell layer and Purkinje neuronal dendrites, accompanied by occasional Purkinje neuronal degeneration and loss with focal gliosis in the Bergmann layer. Spheroid formation was also detectable to a much lower extent in the white matter associated with grey matter lesions. In the spinal cord, spheroid formation was most prominent in the intermediate grey matter at segment T7. Immunohistochemistry for amyloid precursor protein, a known marker of axonal injury (Garcia et al. 2015), highlighted the presence of widespread nerve fiber swelling including perineuronal sites (Fig. 1b). Thus, the observed clinico-pathological phenotype could be explained by a novel form of neuroaxonal dystrophy (NAD).

**SNP data analyses**

Array genotypes of 50 k single-nucleotide polymorphism (SNP) markers were available for 14 Swaledale sheep including 5 NAD-affected and 3 unaffected lambs, 5 dams and their assumed sire. The sampled ram was confirmed as a sire of all eight lambs. Maternity status was investigated for the five selected ewes to determine the potential dams of the NAD-affected lambs. This enabled the construction of a pedigree (Fig. 2a) and a selection of five animals forming two complete parent–offspring trios for whole-genome sequencing (WGS). Furthermore, SNP genotyping data confirmed the reported common origin of the ewes sired
by a single ram, which was used in the flock in the previous years. Based on the pedigree analysis, we assumed recessive inheritance and carried out autozygosity mapping. However, this analysis revealed no intervals of extended homozygous regions with alleles shared by all five NAD-affected animals. In a further attempt to map the disease-associated locus, a non-parametric linkage analysis was performed, which resulted in 41 NAD-linked genome regions showing positive LOD scores (Online Resource 2) in the studied Swaledale pedigree.

Whole-genome sequence analysis

We obtained whole-genome sequences of two NAD-affected lambs, their two dams, and the sire. We hypothesized that a rare breed-specific deleterious variant is responsible for the described NAD phenotype. Therefore, filtering for protein-changing variants present only in these five Swaledale sheep compared to a local control cohort of 60 available genomes from 16 unrelated breeds (Online Resource 3) was performed. Thereby six variants inherited in a simple recessive way or as compound heterozygous remained (Online Resource 4). Additional filtering against a global control cohort of 453 genomes of 54 other sheep breeds (Naval-Sanchez et al. 2018) resulted in only four variants left (Online Resource 4). Among these four, only one variant on chromosome 15 (g.50640254C > T) was homozygous in both affected offspring and heterozygous in all sequenced parents fitting to simple recessive inheritance. As this variant is neither located in a NAD-linked genome region nor a shared homozygosity segment was identified before, we considered this as a less likely candidate, in addition to the fact that the affected LOC101121706 is an uncharacterized gene. Another missense variant in the HERC1 gene, known to be associated with macrocephaly, dysmorphic facies, and psychomotor retardation in humans (OMIM 605109), was found heterozygous in both cases and their sire, whereas the second missense variant in the HERC1 gene appeared also in the global control cohort. We finally excluded these two variants based on the absence of detected genetic linkage to the genome region on chromosome 7 (Online Resource 2). The two other remaining private variants were both located on chromosome 3 (g.23076713T > C and g.230750869G > A) and were predicted as loss-of-function variants of the PLA2G6 gene (Fig. 2). Variants in PLA2G6 are known to be associated with autosomal recessive early-onset forms of neurodegeneration, such as infantile neuroaxonal dystrophy 1 or Parkinson disease 14 (OMIM 603604). Both PLA2G6 variants were heterozygous in the two sequenced NAD-affected offspring, while the two dams were heterozygous for only the variant in PLA2G6 intron 2 (c.336-2A > G) and the sire was heterozygous for the variant in PLA2G6 exon 8 (c.1312C > T; Fig. 2a). Additionally, these two PLA2G6 variants map to one of the identified NAD-linked regions on chromosome 3.

PLA2G6 genotyping

To investigate how the two identified PLA2G6 variants segregate in the family of the NAD-affected sheep, we genotyped all 71 available Swaledale sheep. The sire of the five cases was heterozygous for the exon 8 variant, and all the
dams of the five NAD-affected lambs were heterozygous for the intron 2 variant (Fig. 2a). Interestingly, only the five affected animals carried both variants in a heterozygous state, which was consistent with a compound heterozygous inheritance (Fig. 2a). From the 13 clinically normal lambs, three were homozygous wild type for both PLA2G6 variants, seven were heterozygous for the exon 8 variant and three were heterozygous for the intron 2 variant. In total, out of 52 genotyped ewes, 25 were homozygous wild type for both variants, and 27 were heterozygous for the intron 2 variant (Fig. 2a). Although, we had no sample of the reported common maternal grandsire, we concluded that either he was a carrier or a germline mosaic for the PLA2G6 variant in intron 2.

**Functional confirmation of detected PLA2G6 variants**

To assess the putative impact of the PLA2G6 variant in intron 2 and the variant in exon 8, we experimentally analyzed both annotated PLA2G6 transcripts using mRNA extracted from the nervous tissues of three NAD-affected lambs with the compound heterozygous genotypes. The presence of both PLA2G6 variants in cDNA of three affected sheep was verified by Sanger sequencing after successful RT-PCR amplification. For the splice site variant at the end of intron 2 (c.210-2A > G) using primers located in the exon 2 and exon 4/5 boundary (Fig. 2c; Online Resource 5), a cDNA fragment of the expected size was obtained in all three studied tissues (spinal cord, sciatic nerve, and cerebellum). This single-nucleotide variant affects the canonical dinucleotide sequence for the U2-type GT-AG acceptor splice site at the end of intron 2 (Fig. 2b) and, therefore, was in silico predicted to disrupt splicing. Sanger sequencing of the RT-PCR amplicon confirmed the loss of the evolutionary strongly conserved splice site leading to a 10 bp deletion at the 5’-end of exon 3 by activation of a cryptic splice site (Fig. 2b). As a consequence of the mutant transcript, if translated, the PLA2G6 protein is predicted to be significantly truncated containing only 72 amino acids lacking more than 90% of the normal protein (p.Leu71TrfpsTer3; Fig. 2b). For the variant in PLA2G6 exon 8 (c.1186C > T), primers located at the exons 6/7 and exons 12/13 boundaries were used and resulted in amplification of two cDNA fragments of the expected sizes in all three studied tissues (Fig. 2c; Online Resource 5). Sanger sequencing revealed that the obtained RT-PCR products correspond to the two annotated PLA2G6 transcripts (XM_027968104.1 and XM_012175630.3; Fig. 2c, d). The identified exon 8 variant is predicted as a missense variant for the longer transcript (XM_012175630.3) leading to an exchange of leucine to phenylalanine in the longer protein isoform (XP_012031020.2: p.Leu396Phe; Fig. 2d). However, in the shorter transcript (XM_027968104.1), this variant is predicted as nonsense creating a premature stop codon (XP_027823905.1: p.Gln396*: Fig. 2d) that results in ~50% truncation of the resulting protein (Fig. 2e). Finally, both variants are assumed to severely affect the structure of PLA2G6 by truncating the functionally important domains (Fig. 2e).

**Discussion**

As a result of characterization of the clinicopathological phenotype and evaluating the relatedness of the observed cases of lambs suspicious for an inherited neurodegenerative disorder, we were able to unravel the most likely genetic cause of a novel form of recessive NAD in Swaledale sheep. The two identified loss-of-function variants in intron 2 and exon 8 of PLA2G6 are inherited in a compound heterozygous way. The intron 2 variant affects a highly conserved acceptor splice site leading to a premature stop codon. Similar splicing defect in the bovine MFN2 gene is known to cause a recessively inherited form of degenerative axonopathy in cattle (Drögemüller et al. 2011). PLA2G6 function is important in regulation of physiological processes, such as calcium homeostasis, inflammation, and cell death (Ramanadham et al. 2015). More than half of the amino acid sequence builds protein interaction domains and motifs: the N-terminal domain, the ankyrin repeat domain, and the catalytic domain (Fig. 2e) (Ramanadham et al. 2015). The long isoform includes an additional 54-residue proline-rich sequence at position 396, where it substitutes glutamine in the short isoform. The long isoform is membrane-bound, while the short isoform is found in the cytoplasm (Malley et al. 2018). The herein presented evidence suggests that the affected lambs, which carry a copy of each mutant shorter transcript version of ovine PLA2G6, are not able to express this isoform of the encoded protein. We hypothesize that both identified loss-of-function variants in PLA2G6 transcripts are likely to be degraded by nonsense-mediated decay. Malik et al. (2008) described Pla2g6-null mice showing age-dependent neurologic impairment by 13 months of age. The neuropathological analysis of Swaledale sheep revealed axonal spheroids in the brain similar to those observed in mice (Malik et al. 2008) and human INAD (Gregory et al. 2008). Taken together, the two described pathogenic variants (PLA2G6: c.210-2A > G and c.1186C > T) are considered disease-causing for this novel form of ovine NAD.

**Conclusion**

Based on the known function of PLA2G6 and its role in human and dog neurodegenerative disease, the rarity of the two identified variants in sheep and the perfect co-segregation of the variant alleles with the disease phenotype in the studied pedigree, we conclude that inherited NAD in Swaledale sheep is caused by compound heterozygosity...
for the two identified loss-of-function variants in PLA2G6. Thereby, this study provides another example for the occurrence of allelic heterogeneity causing Mendelian disorders in a livestock species. Our study is the first report of the precise underlying pathogenesis of ovine NAD and demonstrates the success of the WGS-based precision medicine approach using the parent–offspring trios in detecting pathogenic variants associated with rare neurodegenerative disease.

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Author contributions AL, performed all genetic analyses and drafted the manuscript. IMH carried out bioinformatics. JMP assisted in the genetic analyses. KW and BS collected samples and drafted parts of the manuscript. SS performed neuropathology and drafted parts of the manuscript. CD and SS designed the study, supervised the project and finalized the manuscript. All authors participated in writing the manuscript and have read and approved the final version.

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Data availability The whole-genome data of our group has been made freely available under study accession number PRJEB30931 in the European Nucleotide Archive (https://www.ebi.ac.uk/ena). All accession numbers of the WGS are available in the Online Resource 3. Sheep genomes project variant database of further 453 samples is deposited in ENA under accession number PRJEB14685. All references to the ovine PLA2G6 gene correspond to the accessions NC_040254.1 (NCBI accession), XM_027968104.1 and XM_012175630.3 (mRNA), and XP_027823905.1 and XP_012031020.2 (protein).

Compliance with ethical standards

Conflict of interest The authors have no relevant financial or non-financial interests to disclose.

Ethical approval This study did not require official or institutional ethical approval as ‘non-experimental clinical veterinary practices’ are specifically excluded from being considered regulated procedures under The Animals (Scientific Procedures) Act (ASPA), 1986, Section 2(8) (https://www.rcvs.org.uk/settings/standards/advice-and-guidance/code-of-professional-conduct-for-veterinary-surgeons/supporting-guidance-recognised-veterinary-practice/). The sheep were handled according to good ethical standards and all live animals were blood sampled by a veterinary surgeon (BS) for diagnostic purposes on the farm to determine the cause of the neurological disease. All other sampling was carried out postmortem on affected animals after euthanasia on humane grounds because of progressive neurological signs. The affected lambs were euthanized by intravenous injection of an overdose of sodium pentobarbital, performed by BS. The aim was to identify the cause of the congenital disorder and thereby improve the animal welfare situation on the farm by identifying the underlying genetic cause and preventing breeding of further cases. Written consent was obtained from the sheep owner.

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