**Ultrastructural features of canine neuroaxonal dystrophy in a Papillon dog**

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**ABSTRACT.** Neuroaxonal dystrophy (NAD) is a neurodegenerative disease characterized by severe axonal swelling (spheroids) throughout the nervous system. In dogs, NAD has been reported in several breeds and a missense mutation in \( PLA2G6 \) gene has recently been identified in the Papillon dog NAD. Here we performed ultrastructural analysis to clarify the detailed ultrastructural features of the Papillon dog NAD. Dystrophic axons consisted of accumulation of filamentous materials, tubulovesicular structures, and swollen edematous mitochondria with degenerated inner membranes were often observed in the central nervous system. At axonal terminals, degeneration of presynaptic membrane was also detected. As reported in \( Pla2g6 \) knockout mice, mitochondrial and presynaptic degeneration may be related with the pathogenesis of NAD in Papillon dogs.

**KEY WORDS:** electron microscopy, mitochondria, neuroaxonal dystrophy, papillon, \( PLA2G6 \)
2.5% glutaraldehyde in 0.1 M phosphate buffer (PB; pH 7.4), post-fixed with 1% osmic tetraoxide at 4°C overnight, and embedded in epoxy resin. One-μm semi-thin sections were cut and stained with toluidine blue (TB) for light microscopy. Ultrathin sections were stained with uranyl acetate and lead citrate and examined in a Hitachi H-7500 electron microscope (Hitachi, Tokyo, Japan).

Histopathologically, dystrophic swollen axons were found throughout the CNS, and the spheroids showed varied sizes and heterogeneous morphology; homogeneous and granular appearance with or without clefts and vacuoles (Figs. 1 and 2). Some spheroids had a densely stained or granular central core structure in TB stained semi-thin sections (Fig. 2). A number of axonal spheroids were predominantly localized in the dorsal horn of spinal cord, cerebellum and medulla oblongata including nuclei cuneatus, nuclei gracilis, nuclei olivaris, nuclei spinalis nervi trigemini and lemniscus medialis. These spheroids showed strong immunoreactivity for synaptophysin and NFs markers (data not shown) similar to previous reports [13]. From these histological features, this case was diagnosed as NAD. As previously reported by Tsuboi et al., a missense mutation in the patatin domain of PLA2G6 gene (c.1579G>A) was identified in this dog by whole exome sequencing analysis and TaqMan genotyping assays [20].

Transmission electron microscopic observation revealed that axonal spheroids consisted of accumulation of filamentous and granular materials, tubulovesicular structures (Figs. 3a, 3b and 3c: white asteriks) and densely packed mitochondria, as well as edematous vacuoles, vesicular structures and electron-dense bodies (Fig. 3a–d) in the CNS. It is noteworthy that numerous swollen edematous mitochondria with degenerated inner membranes were often observed in the spheroids and axons (Fig. 3d–g: arrowheads); some abnormal mitochondria were also detected in presynapses (Fig. 3e and 3f). In these abnormal mitochondria, cristae were diffusely degenerated, branching and tubular appearance. Abnormally degenerated mitochondria were also detected in neurons (Fig. 3h: Purkinje cell, arrowheads). Some abnormal mitochondria were surrounded by membrane structures, suggesting mitophagy (Fig. 3h: inset). There were only a few mitochondria or mitochondria-like structures containing dense granules (data not shown). In addition, the presynaptic membranes were degenerated and expanded irregularly in axonal terminals (Figs. 3e and 3f).

Canine NAD has previously been reported in several breeds [2–4, 6–9, 13–15, 18, 21], but both ultrastructural and genetic/molecular analyses have been performed for a few cases. For example, spheroids consisted of the accumulation of autophagosomes in Spanish water dogs with a tectonin beta-propeller repeat-containing protein 2 (TECPR2) mutation [9]. In laboratory dogs (crossbreed of Giant Schnauzer and Beagle) with a mitofusin 2 (MFN2) mutation, dystrophic axon showed accumulations of membrane-bound vesicles containing variably electron-dense materials and organelles such as fragmented degenerating mitochondria (mitophagy) [5, 6]. In our case, there were no accumulated autophagosome, but we found some abnormal mitochondria or dense bodies surrounded by membrane structures. Immunohistochemical approach will be helpful to investigate involvement of autophagy or mitophagy in the pathogenesis of Papillon dog NAD with a PLA2G6 mutation.

In this report, we performed detailed ultrastructural analysis on the CNS in a Papillon dog with NAD and revealed that ultrastructural features of axonal spheroid were closely similar to human INAD [3, 24] and INAD mouse models [1, 19, 22, 23]. In addition to the accumulation of tubulovesicular structures in spheroids (a pathological hallmark of INAD), it is noteworthy that densely packed mitochondria and numerous swollen edematous mitochondria with degenerated inner membranes were detected in the spheroids and axons. These result suggested that the Papillon dog NAD with a missense mutation in the PLA2G6 gene seems to have similar mitochondrial pathology with human and mouse INAD.

In mice, two types of INAD models have been established: Pla2g6-KO (null mutation) mouse and Pla2g6-inad mouse with a point mutation in the ankyrin domain of Pla2g6. Both of INAD mouse models have no phospholipase enzymatic activity, but the clinical course and ultrastructural findings are different. In Pla2g6-KO mouse, the clinical onset was critically late and neurological abnormalities developed slowly [1, 17, 19]. On the other hand, Pla2g6-inad mouse showed early onset of clinical symptoms and progressed rapidly, and then all of the homozygous mice died before 18 weeks of age [22, 23]. Ultrastructurally, typical tubulovesicular structures were detected in both mouse models [1, 19, 23]. In Pla2g6-INAD mouse, mitochondrial and presynaptic membrane abnormalities also developed from an early stage of disease (presymptomatic stage); mitochondrial inner membranes degenerated into granular materials, and mitochondria with tubulovesicular cristae were frequently observed in late stage of disease. This indicated that specific degeneration of inner mitochondrial and presynaptic membranes underlie INAD pathology [1, 19]. Most cases of Papillon dog NAD are considered to be an early onset [20], and the present case already had prominent mitochondrial abnormalities in the CNS, which correspond to those in later stage of Pla2g6-INAD mouse. In our Papillon dog (intermediate to late clinical stage) and Pla2g6-inad mouse, no or only a few abnormal mitochondria containing dense granules considered to be degenerated inner membrane were observed. This may reflect the differences in the disease stage or rate of progression. It would be important to study the relationship between the ultrastructural features of neuronal/axonal abnormalities and the disease stage or severity. Furthermore, phenotypic differences among these INAD cases could be caused by the pattern of the existence of mutated PLA2G6 protein and variations of mutated positions.

In human INAD, axonal spheroids contain tubulovesicular or tubulomembranous structures, aggregation of mitochondria, vesicles and membranous bodies [3, 24]. Detailed mitochondrial changes, such as degenerating changes and branching tubular cristae, were also mentioned [1, 3]. These mitochondrial abnormalities were similar to our case and INAD mouse models, but further accumulation of ultrastructural analysis data about human INAD cases is needed for detailed comparative analysis with animal models.

Taken together, our detailed ultrastructural study revealed that degeneration of mitochondrial inner membrane and presynaptic membrane may be related with the pathogenesis of NAD in Papillon dogs, and NAD-affected Papillon dogs could be valuable animal model for human INAD. Establishment of new rodent model that bears a missense mutation in the same domain as Papillon dog NAD might be also a favorable tool for further studies. The comparative analyses of different mutated types of INAD animal models may clarify the pathogenesis of PLAN including INAD.
Fig. 1. The spheroids (arrows) vary in sizes from 10 to 50 µm and show various morphologies. Some spheroids contain clefts or vacuoles. Medulla oblongata. HE. Bar=50 µm.

Fig. 2. The spheroids show homogenous appearance or contain granular material and dense central core with thin myelin (black arrows) or without myelin (white arrows). Spinal cord, gray (a) and white (b) matter. TB, semi-thin section. Bar=25 µm.

Fig. 3. Axonal spheroids are filled with tubulovesicular structures (a, b and c: white asterisks), granular or filamentous materials (d), edematous vacuoles (c, d and g: black asterisks), dense bodies (c and g: white arrows) and densely packed mitochondria (a). Swollen mitochondria with degenerated inner membranes, characterized by degenerated branching and tubular cristae, are also observed (d–g: arrowheads). In the axonal terminals, presynaptic membranes expand irregularly, and membranous degeneration is observed (e and f: black arrows). Abnormal mitochondria, sometimes surrounded by membrane structures (h: inset), are also detected in the Purkinje cells (h: arrowheads). (a–f) spinal cord, (g) cerebrum, (h) cerebellum. Transmission electron microscopy. Bar=1 µm.
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