Dear Sir,

Vocal cord abductor paralysis (VCAP), frequent in the later stage of multiple systemic atrophy (MSA) [1–5], is rare in Parkinson’s disease (PD) [5, 8, 9] and exceptional in dementia with Lewy bodies (DLB). Here we report the first autopsy-verified case of DLB with VCAP.

The patient felt anxiety at the age of 71, and consulted the psychiatrist of our hospital; mild dementia was diagnosed and minor tranquilizer was prescribed. At the age of 75, he experienced difficulty in playing tennis. Gradually, his walking slowed down. At the age of 76, he had visual hallucination. A neurologist of our hospital noted akinesia, mask-like face and small steppage gait and apparent dementia based on mini-mental state examination (MMSE 21/30). He was diagnosed as having probable DLB based on the clinical criteria [10]. L-DOPA was administered, but his symptoms progressed gradually. He had a percutaneous endoscopic gastrostomy because of aspiration pneumonia at the age of 81. He became bedridden at the age of 83. At the age of 87, he exhibited inspiratory stridor, and endoscopic examination of the larynx disclosed bilateral VCAP. The ambiguous nucleus was at most slight (fig. 1c, d). Lewy pathology were also seen in the limbic system (hippocampus, amygdala, cingulate gyrus) moderately (fig. 1e), in the neocortex (superior temporal, insula, parietal, occipital gyri) mildly, and in the spinal cord (cervical to sacral segment) moderately. These were consistent with DLB of the neocortical type in pathological criteria [10]. Alzheimer pathology of Braak NFT stage II and amyloid stage B were also seen. There were no findings associated with old age hippocampal sclerosis. Intrinsic laryngeal muscles showed mild neurogenic change including atrophic fibers, angulated fibers, and small group atrophy (fig. 1f).

Anatomical tracing based on autopsy samples of MSA patients from ambiguous nucleus, myelinated fibers in the motor divisions of the laryngeal branch of the recurrent laryngeal nerve [6], neuromuscular junctions of the posterior cricoarytenoid muscle [7] and intrinsic laryngeal muscles indicate neurogenic paralysis of intrinsic laryngeal muscles as the underlying mechanism for VCAP in MSA. In contrast with frequent VCAP in MSA, VCAP is very rare in clinically diagnosed PD patients [5, 8, 9].
and has not yet been reported in clinically diagnosed DLB. Among them, autopsy-verified PD with VCAP was limited to a single report [5]. Because no remarkable changes were detected in the intrinsic laryngeal muscles of these PD patients with VCAP, sustained tonic state of intrinsic laryngeal muscles was considered to be responsible for VCAP in PD. In this DLB patient as well, there was only mild neurogenic change in intrinsic laryngeal muscles and neuronal loss was not evident in ambiguous nucleus, which alone cannot explain VCAP. We speculated that the same mechanism induced VCAP in PD and DLB.

This is the first autopsy report of DLB with VCAP. VCAP is a life-threatening complication, which may cause sudden death in patients. Therefore, even if VCAP is exceptionally rare in DLB, it is the condition of a patient that we should be careful about.

Disclosure Statement

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References