Case Report

Ramsay Hunt syndrome and zoster laryngitis with multiple cranial nerve involvement

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A R T I C L E   I N F O

Article history:
Received 15 February 2015
Received in revised form 24 February 2015
Accepted 24 February 2015

Keywords:
Ramsay Hunt syndrome
Laryngitis
Polyneuropathy

A B S T R A C T

Ramsay Hunt syndrome is characterized by varicella zoster virus infection affecting the geniculate ganglion of the facial nerve. It typically presents with vesicles in the external auditory canal associated with auricular pain and peripheral facial paralysis. Although vestibulocochlear nerve is frequently co-involved during the course of Ramsay Hunt syndrome, multiple lower cranial nerve involvement has rarely been described in the literature. In addition, laryngitis due to varicella zoster virus is a diagnostic challenge due to its unfamiliarity among clinicians. We report a case of Ramsay Hunt syndrome with laryngitis involving multiple lower cranial nerves.

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Introduction

Ramsay Hunt syndrome (RHS) is characterized by varicella zoster virus (VZV) infection in the geniculate ganglion of the facial nerve. It typically presents with ipsilateral facial palsy and vesicles in the external auditory canal [1]. Recognizing atypical clinical manifestations of RHS, though not frequently reported, is of clinical importance for clinicians. For instance, RHS without vesicles, condition known as zoster sine herpeticus [2], is difficult to distinguish itself from Bell’s palsy. Although the vestibulocochlear nerve is frequently co-involved during the course of RHS, multiple lower cranial nerve involvement has rarely been described in the literature [5–12]. Additionally, laryngitis due to VZV associated with RHS is not a well-recognized clinical entity. We herein report a case of Ramsay Hunt syndrome with zoster pharyngitis involving multiple lower cranial nerves.

Case report

A 66-year-old woman with a history of hypertension was hospitalized because of progressive dizziness, right-side facial weakness and dysphagia. The patient was in her usual state of health until 6 days before admission, when sore throat and odynophagia developed, associated with subjective fevers and anorexia. Two days before admission, the patient reported dizziness, right-side facial weakness associated with pain radiating to the right ear, and dysphagia. The patient denied headache, photophobia, neck stiffness, nausea or vomiting. On the morning of admission, the patient developed right-side tinnitus, unsteady gait and inability to close the right eye.

On physical examination, the patient appeared to be in mild distress. The temperature was 101.8 °F, blood pressure 170/93 mm Hg, pulse 102 beats per minute, respiration 20 breaths per minute and oxygen saturation 97% on room air. Her heart sounds were regular without murmurs and her lungs were clear to auscultation. On otologic examination, a few vesicles were observed in the external auditory canal of the right ear with a normal tympanic membrane. Neurological examination revealed peripheral facial nerve palsy (VII), with involvement of the left vestibulocochlear nerve (VIII), glossopharyngeal nerve (IX) and vagnus nerve (X) (Table 1). Spinal accessory nerve (XI) and hypoglossal nerve (XII) were intact. Flexible laryngoscopic examination revealed erythema and erosive lesions on the right side of the epiglottis and arytenoid. There was an asymmetry of the soft palate. A biopsy of the erosive lesions was not performed. A non-contrast computed tomography of the head and temporal bone revealed no abnormalities.

Ramsay Hunt syndrome with laryngitis due to varicella zoster virus was suspected and the patient was started on acyclovir and steroid therapy. After the completion of a 7-day course of the combination therapy, the patient’s neurological symptoms improved except for the facial nerve palsy. Healed erosions on the...
laryngeal mucosa were also noted on a follow-up flexible laryngoscopic examination. Three months after discharge, her residual facial nerve palsy had completely recovered.

Discussion

We describe a case of RVS with laryngitis due to VZV complicated by multiple lower cranial nerve involvement. RVS is characterized by the involvement of the geniculate ganglion of the VII nerve and typically presents with a facial nerve palsy and vesicles in the ipsilateral external auditory canal. Although the VII nerve is frequently co-involved, resulting in the development of dizziness and hearing loss, multiple lower cranial nerve involvement due to RVS has rarely been described [3]; available reports in the literature include zoster laryngitis related to RVS with cranial polyneuropathy [4–6], RVS involving the facial and superior laryngeal nerves in a patient with chronic renal disease [7], RVS involving the VII, VIII and X cranial nerves in an immunocompetent individual [8], severe dysphagia due to RVS involving the cranial nerves V, VII, VIII, X, and XII [9], and RVS complicated by seven cranial nerve involvement (V, VII, VIII, IX, X, XI, and XII) [10].

The mechanism of multiple lower cranial nerve involvement associated with RVS has not been entirely elucidated. Proposed mechanisms include VZV dissemination associated with diffuse lymphocytic infiltration along the course of the VII nerve beyond the geniculate ganglion [6], the X nerve anatomic communication with the VII and VIII nerves [11]. It is also proposed that inflammatory spread along the vascular system may result in polyneuropathy [12,13], since the nerves VIII, X, XI, and XII are supplied by the ascending pharyngeal artery [14]. Kiyokawa et al. demonstrated various communications among nerve branches around the acoustic meatus and the auricle before reaching the central nervous system; the X nerve distributing to the posterior wall of the external acoustic meatus had communications with the VII nerve and the IX nerve [15].

Acute laryngitis is a common clinical syndrome encountered by clinicians, typically associated with common respiratory pathogens, such as rhinovirus, coronavirus and Group A β-hemolytic streptococcus [16,17]. Herpesviruses, including VZV, are uncommon causes of acute laryngitis. Clinical findings in patients with laryngitis due to herpesviruses include edema, ulcerative lesions, and vesicles in the glottic or supraglottic region [18]. Although the diagnosis of acute laryngitis caused by common respiratory pathogens can often be made by history alone, laryngoscopic examination of the larynx may be necessary to differentiate zoster laryngitis from other rare etiologies of laryngitis, such as laryngeal tumor, in the presence of multiple cranial nerve involvement.

Table 1: Cranial nerve functions and clinical manifestations (VIII–X).

| Cranial nerve function | Clinical manifestations |
|------------------------|-------------------------|
| Vestibulocochlear (VIII) | Equilibrium, Vertigo, loss of balance |
| Cochlear (V) | Hearing, Tinnitus, hearing loss |
| Glosso-pharyngeal (IX) | Swallowing, Dyphagia, aspiration/cough/choking |
| Vagus (X) | Swallowing, Taste from the posterior one third of tongue |
| | Dysphagia, hoarseness, aspiration/cough/choking |

Conclusions

RVS is characterized by the involvement of the geniculate ganglion of the VII nerve by VZV. Although not common, RVS may involve multiple lower cranial nerves, presenting with a wide variety of neurological symptoms. Moreover, VZV is an uncommon cause of acute laryngitis which requires heightened awareness among clinicians to differentiate it from other etiologies in the presence multiple cranial nerve involvement.

Conflict of interest statement

None.

Sources of funding

None.

Ethical approval

Written informed consent was obtained from the patient for publication of this case report and accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal on request.

Acknowledgement

None.

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