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Article type : Lessons from practice

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Title	Polyneuritis cranialis from varicella zoster virus reactivation
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This is the author manuscript accepted for publication and has undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the [Version of Record](#). Please cite this article as [doi: 10.1002/MJA2.50791](#)

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Primary Keywords [Office use only]	Infectious diseases; Nervous system diseases; Diagnostic techniques and procedures
Secondary keywords [Office use only]	Varicella; Virus diseases; Cranial nerve diseases; Neuropathology
Notes:	

Article details (press ctrl – 9 to enter details):

Article type	Medical education
Blurb	

Office use

<i>Ms. Number</i>	mja20.00157.R2
<i>Medical editor</i>	Christine Gee/Robyn Godding
<i>Medical editor email</i>	rgodding@mja.com.au
<i>Structural editor</i>	Laura Teruel
<i>Structural editor email</i>	lteruel@mja.com.au
<i>Section/Category</i>	Medical education
<i>Strapheading</i>	Medical education
<i>Substrap</i>	Lessons from practice

Wiley – file data:

Filename for copyediting	sch_mja20.00157_ms.docx
Accompanying graphics	sch_mja20.00157_gr.jpg, sch_mja20.00157_gr.tif
Stock images	None
Appendices	None

Office use – history:

Event	Date
Original submission received	13/02/2020

Event	Date
Accept	20/04/2020

Proof sent to author	
Proof returned by author	
Published (date format xx/xx/xx)	
Issue	
Vol	
DOI	
Journal	
Original article DOI (for response)	

Polyneuritis cranialis from varicella zoster virus reactivation

Clinical record

A 68-year-old man with chronic obstructive pulmonary disease, type 2 diabetes mellitus and stage II chronic kidney disease, presented to hospital with a 24-hour history of right-sided facial paralysis and 3 days of horizontal diplopia. He had received 4 days of oral ciprofloxacin and oral prednisolone (37.5 mg) for 2 weeks of progressive right-sided facial swelling with otalgia, initially thought to be otitis externa.

Examination revealed right facial cranial neuropathies. He had a sixth cranial nerve palsy and lower motor neuron seventh cranial nerve palsy. Audiometry confirmed eighth cranial nerve sensorineural involvement without vestibular dysfunction. Ninth and tenth cranial nerve palsies manifested as an absent gag reflex, asymmetric palatal movement and right vocal cord paralysis.

The patient was systemically well without fevers and had normal mentation. Examination of the remaining cranial nerves, cerebellar function, and upper and lower limbs was normal. Examination of the right external auditory meatus showed small, painful papules within the external auditory canal, with normal examination on the left. There was palpable, tender right submandibular swelling and lymphadenopathy without warmth or erythema.

Initially thought to have malignant otitis externa with base of skull osteomyelitis, he was commenced on intravenous piperacillin/tazobactam. Corticosteroids were discontinued. Brain magnetic resonance imaging (MRI) (with contrast) was normal. Both a computed tomography scan of the base of skull and a gallium bone scan revealed no malignant otitis externa or skull base osteomyelitis.

Varicella zoster virus (VZV) DNA was detected by polymerase chain reaction (PCR) from a swab of the lesions within the external auditory canal. Bacterial cultures revealed no significant growth. An antibody test for human immunodeficiency virus (HIV) — for which herpes zoster is an indicator condition — was negative. Further testing of immune status was not justified in the setting of known diabetes mellitus and steroid use. Antibiotic therapy was ceased and a 3-week course of intravenous acyclovir was commenced. The right external auditory canal vesicles dried and crusted (Box 1, A). The patient had no previous history of primary VZV infection or shingles vaccination.

The diagnosis was revised to polyneuritis cranialis of right cranial nerves VI to X due to VZV reactivation. A lumbar puncture was not performed, as VZV DNA was found on the external auditory canal swab. Oral prednisolone commencing at 60 mg daily was

slowly weaned over 3 weeks.

After 4 months, there has been negligible neurological recovery. The patient underwent a right tarsorrhaphy due to poor eyelid closure, and a thyroplasty for persisting right vocal cord paralysis (Box 1, B). Poor swallowing required the placement of a percutaneous endoscopic jejunostomy feeding tube.

Discussion

Polyneuritis cranialis is a clinical pattern of mononeuritis multiplex restricted to cranial nerves and of various aetiologies (Box 2). Differentiation of cranial nerve pathologies requires detailed history, examination and investigation. Time course can differentiate hyperacute (seconds to hours) vascular or traumatic causes from acute (days to weeks) inflammatory or infective aetiologies and subacute (months to years) malignant or infiltrative disease processes. Systemic features such as fever and constitutional symptoms may indicate inflammatory or infectious causes. A history of recent illness or environmental, toxin or medication exposure influences the probability of each differential diagnosis. Assessment of headache can differentiate diseases causing raised intracranial pressure or cranial nerves inflammation. Certain signs may suggest specific diagnoses, such as cranial nerve III, IV, V1, V2 and VI palsies in cavernous sinus pathology. Neuroimaging and cerebrospinal fluid analysis should be considered in all patients.

The lifetime risk of VZV reactivation is about 50%.² Cranial nerve syndromes from VZV commonly involve the trigeminal nerve, presenting as herpes zoster ophthalmicus, or the facial nerve, presenting as Ramsay Hunt syndrome.^{3,4} Polyneuritis cranialis is a rare and severe manifestation of VZV that may be considered as part of a spectrum of complicated VZV infections that includes Ramsay Hunt syndrome.³⁻⁵ Seventh and eighth cranial nerve involvement and the presence of a characteristic vesicular “shingles” rash may distinguish VZV reactivation from conditions such as oculopharyngeal Guillain–Barré syndrome, yet skin lesions are only present if the involved nerve has a cutaneous distribution.^{3,4}

MRI is important to investigate brainstem involvement and exclude other causes of peripheral cranial nerve palsies (Box 2). Enhancement of the cranial nerves is non-specific, and does not exclude VZV reactivation when absent, yet may correlate with pathological findings in the acute phase of VZV reactivation.⁴ Cerebrospinal fluid analysis may show an elevated, predominantly lymphocytic cell count, elevated protein, VZV antibodies and detectable VZV on PCR.³⁻⁵

There is a paucity of data to guide therapy for polyneuritis cranialis. Suggested treatment involves a multiweek course of intravenous acyclovir and systemic corticosteroids. Longer courses have not shown superiority to shorter ones when long term recovery is the outcome measure.⁵

Prognosis is variable and complete recovery is rare.^{4,5} VZV vaccination is likely to be an important preventive consideration in an ageing population.²

Lessons from practice

- Differentiation of cranial nerve lesions relies on accurate discernment of the anatomical location, time course, and associated features of the pathology, aided by neuroimaging and cerebrospinal fluid analysis.
- The time course of illness can help differentiate hyperacute (vascular or trauma) from acute (inflammatory or infective) and subacute disease processes (malignant or infiltrative).
- Headache with “red flags” may suggest raised intracranial pressure or cranial nerves inflammation, while the presence of systemic features raises suspicion of inflammatory or infectious causes. Any recent illness or environmental, toxin or chemotherapy exposure should be considered when differentiating possible causes.
- The presence of skin vesicles with positive polymerase chain reaction (PCR) can distinguish varicella zoster virus (VZV) reactivation from other differentials. There is limited evidence to guide management and prognostication of severe VZV reactivations, with typical treatment involving a combined multiweek course of intravenous acyclovir and corticosteroids. Full recovery is rare.

Competing interests: No relevant disclosures.

Provenance: Not commissioned; externally peer reviewed.

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doi: 10.5694/mja20.00157

References

- 1 Carroll CG, Campbell WW. Multiple cranial neuropathies. *Semin Neurol* 2009; 29: 53-65.
- 2 Zoster (Herpes Zoster): Australian Government Department of Health: Australian Immunisation Handbook; 2018. <https://immunisationhandbook.health.gov.au/vaccine-preventable-diseases/zoster-herpes-zoster> (viewed Jan 2020).
- 3 Tateno F, Sakakibara R, Kishi M, et al. Zoster lower cranial neuropathy: A case report and a review of literature.

Am J Case Rep 2011; 12: 137-139.

- 4 Nisa L, Landis BN, Giger R, Leuchter I. Pharyngolaryngeal involvement by varicella-zoster virus. *J Voice* 2013; 27: 636-641.
- 5 Kim YH, Chang MY, Jung HH, et al. Prognosis of Ramsay Hunt syndrome presenting as cranial polyneuropathy. *Laryngoscope* 2010; 120: 2270-2276.

[Insert Boxes]

[Box 1; sch_mja20.00157_gr]

1 Crusted varicella zoster virus rash in right external acoustic meatus at 2 weeks after admission (A). Persistent right facial nerve palsy and abducens nerve palsy at 3 weeks after admission (B)*

[Box 1 foot]

* Note that the patient's partially opened right tarsorrhaphy did not prevent abduction.

[Box 2]

2 Differential diagnosis of multiple cranial nerve palsies*;¹

Infectious	<ul style="list-style-type: none">▪ Viral: herpes zoster, cytomegalovirus, Epstein–Barr virus, HIV[†]▪ Bacterial: invasive sinusitis/otitis, Lyme disease, syphilis, tuberculosis, botulism▪ Fungal: cryptococcosis, mucormycosis, aspergillosis▪ Parasite: Chagas disease, neurocysticercosis
Inflammatory	<ul style="list-style-type: none">▪ Systemic vasculitis:[†] polyarteritis nodosa, temporal arteritis, ANCA-associated vasculitis▪ Systemic lupus erythematosus[†]▪ Sarcoidosis▪ Guillain–Barré (Miller–Fisher) syndrome▪ Idiopathic cranial polyneuropathy▪ Idiopathic hypertrophic cranial pachymeningitis▪ Tolosa–Hunt syndrome▪ Behçet disease▪ Thyroid eye disease
Neoplastic	<ul style="list-style-type: none">▪ Leptomeningeal disease — primary or metastatic▪ Metastatic — compressive or infiltrative▪ Chemotherapy-induced (eg, vincristine, cisplatin, immune-checkpoint inhibitor)
Vascular	<ul style="list-style-type: none">▪ Microvascular: diabetes[†], sickle cell disease▪ Macrovascular: intracranial aneurysm, carotid artery dissection, cavernous sinus pathology▪ Traumatic: carotid endarterectomy
Bone disease	<ul style="list-style-type: none">▪ Hereditary: osteopetrosis, Paget disease, hyperostosis cranialis interna▪ Traumatic: closed head injury, base of skull fractures

ANCA = antineutrophil cytoplasmic antibody; HIV = human immunodeficiency virus. * Central nervous system pathology involving the brainstem causing multiple cranial nerve palsies requires consideration, but is usually associated with concurrent sensorimotor deficits in the limbs and/or cerebellar dysfunction. [†] Via complication of vasculitis/mononeuritis multiplex.

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Title:

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Date:

2020-09-18

Citation:

Schnall, J. A., Khan, S. F., Zolio, L., Ray, J. C. & Jenney, A. W. J. (2020). Polyneuritis cranialis from varicella zoster virus reactivation. MEDICAL JOURNAL OF AUSTRALIA, 213 (8), pp.352-+. <https://doi.org/10.5694/mja2.50791>.

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