

Australian Paediatric Surveillance Unit STUDY PROTOCOL Acute Flaccid Paralysis



The AFP Study Group announces a briefer questionnaire for the collection of clinical data.

Other study details remain unchanged.

BACKGROUND

Acute flaccid paralysis (AFP) is defined as the acute onset of flaccid paralysis in one or more limbs or acute onset of bulbar paralysis. In the past, acute poliomyelitis was the most common cause of AFP, now Guillian-Barre syndrome and transverse myelitis are the two leading causes of non-polio AFP (for other causes see table 1).

This study commenced in March 1995 at the request of the Commonwealth Department of Health in order to meet the certification requirements of the World Health Organisation for poliomyelitis eradication. In October 2000 the Western Pacific region, which includes Australia, was certified free of circulating wild poliovirus. However, AFP surveillance must continue to document maintenance of Australia's polio-free status and to detect imported cases and cases of vaccine associated paralytic polio.

STUDY OBJECTIVES

- 1. To determine whether AFP is caused by poliovirus infection and if so, whether it is 'wild' or vaccine strain poliovirus.
- 2. To document the reported rate of AFP over time and to assess the adequacy of AFP surveillance according to WHO criteria
- 3. To monitor the aetiology, clinical features and outcome at 60 days of non-polio AFP.

Table 1: Causes of AFP

Peripheral neuropathy Muscle disorders Disorders of neuromuscular transmission Guillain-Barre syndrome Polymyositis, dermatomyositis Myaesthenia gravis Trichinosis Acute axonal neuropathy Botulism Periodic paralyses Neuropathies of infectious Insecticide eq: organophosphate Corticosteroids and blocking diseases poisoning agents Acute toxic neuropathies (heavy Tick bite paralysis Mitochondrial diseases (infantile metals, snake toxin) Snake bite type) Focal mononeuropathy Post viral myositis Systemic disease Anterior horn cell disease Acute myelopathy Acute anterior poliomyelitis Cord compression eg: Tumor, Acute porphyrias trauma Demyelinating diseases Other neurotropic viruses Critical illness neuropathy eg: transverse myelitis, acute Acute myopathy in ICU Patients disseminated encephalomyelitis (ADEM) Ischaemic cord damage eg: anterior spinal artery syndrome Peri-operative complication

CASE DEFINITION

Please report any child less than 15 years of age with acute flaccid paralysis in one or more limbs or acute onset of bulbar paralysis.

All cases are reviewed by the National Polio Expert Panel and classified as: confirmed poliomyelitis; non-polio AFP, polio-compatible or non-AFP.

REPORTING INSTRUCTIONS

- Telephone/Email reporting: Please report cases immediately by telephone to VIDRL on (03) 9342 9607 or email enterovirus@mh.org.au
- APSU reporting: report cases on the monthly APSU report card. Please keep patient details in your blue APSU folder
- It is important that stool specimens be collected from all patients with AFP, even when an alternative definitive diagnosis has been confirmed.

COLLECTION OF STOOL SAMPLES

- Please collect 2 stool samples 24 hours apart within 2 weeks of onset of paralysis and send them to your local laboratory who will forward them to the National Enterovirus Reference Laboratory, VIDRL, Doherty Institute, 792 Elizabeth Street, Melbourne Victoria 3000
- On the request form the patient must be identified as having AFP.
- Please inform the laboratory that the specimens must be forwarded to VIDRL for exclusion of poliovirus.
- All costs for transport and analysis will be borne by VIDRL. Information regarding specimen transport can be obtained from the National Enterovirus Reference Laboratory at VIDRL on 03 9342 9607, or at
- http://www.vidrl.org.au/laboratories/poliovirus-reference/specimen-referral/
- VIDRL will send results to your local laboratory

INVESTIGATORS

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