Froin's syndrome associated with spinal tuberculosis

Kripa Maharjan,¹ Sudeep Adhikari,² Buddha Basnyat³

¹Patan Academy of Health Sciences, Kathmandu, Nepal

²Department of Internal Medicine, Patan Hospital, Kathmandu, Nepal

³Oxford University Clinical Research Unit, Patan Hospital, Kathmandu, Nepal

Correspondence to

Professor Buddha Basnyat, buddhabasnyat@gmail.com

Accepted 30 November 2018

DESCRIPTION

A 25-year-old male patient from Sarlahi district in Southern Nepal presented to the emergency department of Patan Hospital in Kathmandu, Nepal with an inability to move both lower limbs for 8 days. Prior to this, he had a low-grade fever, dry cough, decreased appetite, weight loss and tingling sensation on the lower limbs for 3 months. He was diagnosed with sputum-positive pulmonary tuberculosis (TB) 1 month ago and category 1 antitubercular therapy (ATT) was started with daily three tablets of HRZE, each containing 75 mg isoniazid (H), 150 mg rifampicin (R), 400 mg pyrazinamide (Z) and 275 mg ethambutol (E) but as fever subsided, he stopped taking the ATT after about 10 days. Ten days later after stopping his medicines, he developed an inability to move his lower limbs and had urinary retention in addition.

Nervous system examination revealed a spastic type of paraplegia with bilateral upgoing plantar reflexes and exaggerated lower limb deep tendon reflexes. Other examinations were normal.

Lumbar puncture revealed xanthochromic cerebrospinal fluid (CSF) which coagulated within

Figure 2 MRI of a thoracic spine showing compression of T7 and T8 vertebrae along with discitis at T7–T8 level, prevertebral abscess extending from T6 to T9 level, and contrast enhancing soft-tissue density (epidural abscess) in the anterior aspect of cord from T6 to T8, compressing the spinal cord.

Figure 1 Xanthochromic cerebrospinal fluid.

an hour while awaiting for laboratory analysis (figure 1). Blood counts could not be done, but the protein level was more than 1500 mg/dL. This triad (CSF xanthochromia, hypercoagulability and very high protein count) was first described by Froin, a French physician.¹ This classic combination has been termed Froin's syndrome. It occurs with blockage of CSF flow by a spinal cord mass such as a tumour or with meningeal irritation from inflammation.² Distal to the block stagnation of CSF occurs leading to higher protein transudation, and resulting xanthochromia (due to bilirubin binding to albumin) and hypercoagulability (due to high fibrinogen level).³

The presence of paraplegia in the setting of pulmonary TB prompted us to consider Pott's paraplegia in our patient, and also the possibility of some obstructive cord lesion was considered due to the presence of Froin's syndrome. MRI was suggestive of Pott's spine with the presence of spondylitis at thoracic T3–T9 vertebrae, compression of T7 and T8 vertebrae along with discitis at T7–T8 level,

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To cite: Maharjan K, Adhikari S, Basnyat B. *BMJ Case Rep* 2018;**11**:e228367. doi:10.1136/bcr-2018-228367

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prevertebral abscess extending from T6 to T9 level, and contrast enhancing soft-tissue density in the anterior aspect of cord from T6 to T8, compressing the spinal cord, suggestive of tuberculous epidural abscess (figure 2). This explained the presence of Froin's syndrome in our patient with Pott's spine.

He was restarted on ATT. The abscess was drained. On follow-up after 2 weeks of discharge, he had marked improvement in his paraplegia, though still unable to function independently.

The importance of reporting about Froin's syndrome is that its presence in our patient prompted us to think of some obstructive

cord lesion and hence helped to clinch the proper diagnosis with an MRI. In our part of the world, we can use this syndrome as a clue towards the diagnosis when diagnostic modalities like MRI are not available or are unaffordable.

Acknowledgements To orthopedics department, Patan Hospital for their help in patient management. To radiology department, Patan Hospital for providing MRI image.

Contributors SA conceived the write-up, KM wrote the first draft, SA and BB reviewed the manuscript, KM, SA and BB took care of patient and decided the final draft.

Funding The authors have not declared a specific grant for this research from any funding agency in the public, commercial or not-for-profit sectors.

Competing interests Not required.

Patient consent Obtained.

Provenance and peer review Not commissioned; externally peer reviewed.

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Learning points

- Pott's spine usually takes the form of spondylodiscitis leading to paraplegia, and sometimes can develop epidural abscess compressing the spinal cord and the central cerebrospinal fluid (CSF) canal.
- Yellow and hypercoagulable CSF with high protein is a classic triad called Froin's syndrome.
- The presence of Froin's syndrome should prompt us to think of some obstruction to CSF flow such as epidural abscess as in this case.

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