Normal pressure hydrocephalus presenting as Parkinson’s syndrome

A. Miodrag, T. K. Das and R. J. Shepherd

Department of Geriatric Medicine, Leicester General Hospital, Gwendolen Road, Leicester LE5 4PW, UK.

Summary: Although extrapyramidal features in normal pressure hydrocephalus (NPH) are not uncommon, presentations with Parkinson’s syndrome as the predominant feature are rare and may give rise to diagnostic difficulties. Failure of patients with parkinsonism to respond to therapy, should alert one to the possibility of NPH.

Introduction

There is universal acceptance of Hakim & Adams' original description of the clinical features of normal pressure hydrocephalus (NPH); namely, the triad of gait disturbance, dementia and urinary incontinence. This is associated with dilatation of the ventricular system of the brain without clinical evidence of raised intracranial pressure. There are wide variations in the clinical manifestations of NPH, particularly in regard to the disturbance of gait. This is most often a spastic ataxia, although less commonly, parkinsonian features can occur. Rarely patients may present atypically with signs indistinguishable from Parkinson’s syndrome.

We report a case of normal pressure hydrocephalus in whom the initial manifestation was that of Parkinson’s syndrome, which resolved completely following the insertion of a ventriculo-peritoneal shunt.

Case report

A previously fit and active 71 year old woman was admitted with an 18 month history of tremor of the hands and progressive difficulty in walking. Parkinson’s syndrome had been diagnosed by a neurologist who commenced treatment with amantadine, but despite this her walking continued to deteriorate with numerous backward falls. She was virtually bed bound for 2 weeks before admission and was incontinent of urine for 6 months.

On examination she looked healthy, was a little forgetful but well orientated. Tendon reflexes were generally brisk, particularly in the legs, with bilateral extensor planter responses. Tone was increased in all limbs with cogwheeling rigidity of both wrists. There was slight weakness of the legs but sensation was normal. She was bradykinetic and could take only a few retropulsive, wide based shuffling steps between two nurses. The rest of the examination, including fundoscopy, was normal.

Haematological and biochemical screen and chest X-ray were normal. X-ray of spine revealed spondylitic changes in the cervical and dorso-lumbar spine. In view of her paraparesis a myelogram was done, which showed no evidence of cord compression. Cerebrospinal fluid (CSF) pressure and protein content were normal.

There was little improvement in her walking despite intensive physiotherapy and the addition of levodopa/carbidopa (Sinemet) and selegiline (Eldepryl) to her medication. Over the next 2 months she was intermittently confused, forgetful, aggressive and abusive. She remained parkinsonian, ataxic and incontinent of urine. A computed tomographic (CT) head scan (Figure 1) showed enlargement of the ventricles with normal cerebral sulci, consistent with normal pressure hydrocephalus. She was referred to a neurosurgical unit where a ventriculo-peritoneal shunt was inserted. CSF pressure at operation was normal. Four weeks post-operatively her urinary incontinence resolved and by 6 weeks her plantar responses were flexor. There were no signs of parkinsonism and her confusion cleared, with no recurrence of symptoms on stopping all her medication. By 8 weeks she was independent, walking well with a Zimmer frame and could climb stairs. She was discharged home.

Discussion

The clinical features of NPH are due to the selective involvement of periventricular regions of the brain consequent upon progressive ventricular enlargement.

© The Fellowship of Postgraduate Medicine, 1987
lateral ventricles; accounts for paraparesis and incontinence. Involvement of the terminal dopaminergic neurons bordering the lateral ventricles may produce extrapyramidal symptoms. Edvinsson et al. demonstrated a significant progressive decline in fore and midbrain dopamine and homovanillic acid content in rabbits one month after producing experimental normal pressure hydrocephalus. Sypert et al. reported three cases of NPH in which an extrapyramidal syndrome was the predominant manifestation, with resolution following shunt insertion. Mazza et al. reported a similar case, with mild improvement post shunt. Sypert et al. suggested that parkinsonian symptoms in NPH are the result of mechanical distortion of the basal ganglia and consequent vascular insufficiency in the nigrostriatal system. In the same way, local distortion of the basal ganglia and mid brain by tumour can produce parkinsonian symptoms that disappear after successful decompression.

The original report of NPH being an eminently treatable cause of dementia led to an explosion of shunt insertions in demented patients. This was often inappropriate due to the inclusion of patients with cerebral atrophy (hydrocephalus ex vacuo) who do not respond to shunting. Clinical features alone are insufficient to make a diagnosis of NPH, but can raise the possibility. Other causes of dementia, such as Alzheimer’s disease, can be associated with gait disturbance and incontinence, but in NPH gait is usually severely impaired before significant mental changes are apparent. Diagnostic tests include; CT scanning, isotopic cisternography, clinical response to test removal of CSF and intracranial pressure monitoring.

Overall, about 50% of patients improve after a shunt procedure with best results in subjects in whom gait disturbance is the predominant feature and vice versa where dementia is predominant.

References

9. Edvinsson, L., Nielsen, K.C., Owman, Ch., Rosengren, E. & West, K.A. Concomitant fall in brain dopamine and


Normal pressure hydrocephalus presenting as Parkinson's syndrome.
A. Miodrag, T. K. Das and R. J. Shepherd

doi: 10.1136/pgmj.63.736.113

Updated information and services can be found at:
http://pmj.bmj.com/content/63/736/113

These include:

Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/