Intracerebral Haemorrhage And Angiographic Beading Following Ingestion Of Catecholaminergics

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SUMMARY We report two cases of serious intracerebral haemorrhage occurring in young women following their first use of oral medications containing catecholaminergic agents (phenylpropanolamine in combination with ephedrine or pseudoephedrine). Both women were previously well, and there was no evidence for systemic vasculitis, coagulopathy, aneurysm or arteriovenous malformation. Angiography in both cases, performed approximately forty hours following drug ingestion, revealed the beading pattern typical of that seen in previously reported cases of presumed amphetamine-induced "vasculitis." We believe that this arteriographic abnormality is non-specific and should not be construed as necessarily indicative of arteritis.

THE INGESTION OF AMPHETAMINES by either oral or parenteral route has been associated with intracranial haemorrhage and with the angiographic appearance of "beading." Following the demonstration by Citron et al. of a syndrome in drug abusers that is clinically and pathologically reminiscent of polyarteritis nodosa, it has been usual to attribute such occurrences to drug-induced cerebral vasculitis. In many instances, the pathological support for this contention has been weak. We report two cases seen recently in which haemorrhage following ingestion of amphetamine congeners could not reasonably be attributed to an inflammatory angiopathy.

Case Histories

Patient 1
This 20 year old woman, previously in good health, ingested two capsules of a purported amphetamine look-alike twelve hours prior to her admission to hospital. There was no history of previous drug intake. Two hours later, she experienced severe headache and nausea, and after another hour, she developed right hemiparesis and aphasia, followed by progressive drowsiness. On admission, her blood pressure was 130/70 and the general physical examination was normal. She was drowsy and mute, and her eyes opened in response to voice. She had a right homonymous hemianopsia, deviation of her eyes to the left and a dense right spastic hemiparesis.

Normal laboratory investigations included complete blood count (CBC), sedimentation rate, prothrombin time (PT), partial thromboplastin time (PTT), anti-nuclear antibody (ANA), rheumatoid factor, hepatitis B surface antigen, C3, C4, CH50, Clq binding, urinary vanillylmandelic acid (VMA) metanephrine and free catecholamines. Blood and urinary toxic screens were negative. A computerized tomographic (CT) scan (fig. 1) revealed a left external capsular haemorrhage with shift of the midline structures. Cerebral angiography (fig. 2), performed 36 hours following drug ingestion, showed alternating narrowing and dilatation of several branches of the middle cerebral artery. Analysis of the capsules demonstrated the presence of ephedrine, phenylpropanolamine and caffeine. She was managed conservatively and made a slow but incomplete recovery. Three years later, she had minimal word-finding difficulty, normal visual fields and was fully ambulatory with a moderately severe spastic hemiparesis.

Patient 2
This 23 year old woman was previously in good health. Fifteen hours prior to admission, she ingested a single "black beauty" as a "wake-up" pill. She had not used these or similar agents in the past. One hour later, she experienced sudden headache, nausea and vomiting, and this was followed within an hour by aphasia and right hemiparesis. On admission, her BP was 130/90 and the general physical examination was unremarkable. She was obtunded and had a non-fluent aphasia, a right homonymous hemianopsia and a dense spastic right hemiparesis.

Normal laboratory values included CBC, sedimentation ratio, PT, PTT, Bleeding Time, immunoglobulins, ANA, rheumatoid factor and VDRL. A CT scan (fig. 3), showed a left ganglionic haemorrhage with mass effect. Angiography (fig. 4), performed approximately 40 hours after drug ingestion, revealed an avascular mass and patchy areas of spasm in branches of the middle and anterior cerebral arteries. Analysis of the ingested capsules revealed caffeine, phenylpropanolamine, pseudoephedrine and a barbiturate.

This woman was also managed conservatively and made a modest improvement. One year later, her language was normal, but she still had a moderate right arm paresis.

Discussion
The chronic abuse of drugs has been clearly associated with the development of a systemic necrotizing vasculitis that may be difficult to distinguish from polyarteritis nodosa, and amphetamines have been identified as common agents among the multitude of preparations ingested. Intracranial haemorrhage and the cerebral angiographic appearance of alternating narrowing and dilatation of vessels "(beading)") have been reported in association with the chronic use of amphet-
amines and have been attributed to cerebral vasculitis.1-3

More recently, intracranial haemorrhage has been seen following the use of phenylpropanolamine1-6 and as a complication of pseudoephedrine overdose.7 Reports have also implicated the chronic abuse of nasal decongestant sprays8 and abuse of ephedrine.9 These drugs are used as decongestants, stimulants and appetite suppressants, and are readily available without prescription. Angiography has again demonstrated beading;7-9 in one case, this was seen following single-time use.7

The appearance of beading is non-specific, and has been noted in meningitis and encephalitis,10 and following aneurysmal subarachnoid haemorrhage.11 In adult rhesus monkeys, transient angiographic changes were seen within ten minutes of intravenous administration of methamphetamine.12 The abnormalities reverted to normal by 24 hours, but were again noted at one week. Further analysis suggests, however, that such animal studies are not truly representative of the human disorder, in that all of the monkeys experienced significant hypertension, and there was a high incidence of cardiac dysrhythmia, focal cerebral ischaemia and diffuse cerebral edema, as well as gross subarachnoid haemorrhage. Angiographic evidence of
spasm has also been seen acutely following the intracisternal injection of norepinephrine.\(^1\)

Although extracerebral angiitis has been well demonstrated in drug abusers,\(^4,14\) few attempts have been made to correlate cerebral arteriographic changes with histopathology. In 16 amphetamine-related haemorrhages reviewed by Delaney and Estes,\(^3\) 10 patients underwent angiography and beading was demonstrated in only 4. Of these, pathological study was lacking in 3, and necrotizing cerebral angiitis was demonstrated in the remaining patient.\(^5\) All 4 patients had subarachnoid haemorrhage.

In the case of ephedrine-related haemorrhage recently reported by Wooten and colleagues,\(^9\) beading was not seen on the initial angiogram, but was found at followup study. The second angiogram was, however, complicated by symptoms of ischaemia in the distribution of the injected artery (left carotid), and the beading may therefore have been related to contrast-induced spasm.

**Editor's Note:**
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**References**

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