Infarcts in the territory of the lateral branch of the posterior inferior cerebellar artery

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Abstract
The territory of the lateral branch of the posterior inferior cerebellar artery (1PICA) supplies the anterolateral region of the caudal part of the cerebellar hemisphere. Because infarcts in the territory of the 1PICA have rarely been studied specifically, 10 patients with this type of infarct are reported. An 1PICA infarct was isolated in only three patients, whereas it was associated with brainstem infarct in four, with occipital infarct in one, and with multiple infarcts in two patients. The most common symptom at onset was acute unsteadiness and gait ataxia without rotatory vertigo (six patients). Unilateral cerebellar dysfunction was found in all patients, with limb ataxia (nine patients), dysdiadochokinesia (five patients), and ipsilateral body sway (four patients), but dysarthria and primary position nystagmus were notably absent. In the patients with a coexisting infarct in the brainstem, cranial nerve and sensorimotor dysfunction was prominent and often masked the signs of cerebellar dysfunction. Unlike other infarcts in the PICA territory, 1PICA territory infarcts were mainly associated with vertebral artery atherosclerosis (six patients), whereas cardiac embolism was less common (three patients). Unilateral limb ataxia without dysarthria or vestibular signs suggests isolated 1PICA territory infarction and should allow its differentiation from other cerebellar infarcts.

(Cerebellar infarcts in the territory of the lateral branch of the posterior inferior cerebellar artery (1PICA) have rarely been reported. A few pathological findings and one clinical report exist. We had the opportunity to study 10 patients with an infarct in the 1PICA territory documented on MRI. This allowed us to delineate suggestive clinical features and to evaluate the potential causes of infarcts in the 1PICA territory.

Patients and methods
Ten patients (two women aged 77 and 81 years; eight men, mean age 64 (SD 10) years) were selected from the Lausanne Stroke Registry between 1988 and 1992. The characteristics of this registry, which records information on first ever stroke patients consecutively admitted to a population based primary care centre, have been reported. All patients had had the systematic investigations of the registry (CT, Doppler sonography, ECG, 3-lead ECG monitoring for at least 24 hours, standard blood tests). Conventional cerebral angiography was performed in one patient and echocardiography in four patients. MRI was performed in all patients with a Siemens Magnetom 63 SP 1.5 tesla between three and seven days after stroke. The MRI studies included T1, T2, and proton weighted images. MR angiography of the vertebral and basilar arteries was performed in eight patients and consisted of a three dimensional time of flight sequence sensitive to arterial flow with saturation of the veins (TR 36 ms, TE 7 ms, FA 15°, 64 slices of 0.8 to 1 mm). The angiographic data were reconstructed with a multiple intensity projection algorithm and displayed in three dimensions.

The lateral branch territory of the PICA was determined on MRI after recent anatomical and pathological studies. The PICA arises from the end portion of the vertebral artery and supplies the caudal part of the cerebellar hemisphere and vermis, as well as the dorsolateral region of the medulla oblongata. It divides into two main branches: (a) the medial branch (mPICA), which supplies the dorsomedial part of the cerebellar PICA territory and the dorsal medullary territory; (b) the lateral branch (1PICA), which supplies the anterolateral part of the cerebellar PICA territory (fig 1).

Because of possible variations of the PICA calibre and the possible supply of its lateral territory by the anterior inferior cerebellar artery (AICA) in 13% to 40% of the population, we considered a diagnosis of infarct in
the 1PICA territory only when the infarct spared the middle cerebellar peduncle and the flocculus, which are classically supplied by the AICA.

**Results** (figure 2)
Among 10 patients, only three had an isolated infarct in the 1PICA territory. Seven patients had associated infarcts: four in the brainstem, one occipital, and two multiple (supratentorial and infratentorial). The 1PICA territory infarcts were equally distributed on the right and left side. They were never bilateral. Figure 3 gives an example of an 1PICA territory infarct on MRI.

**CLINICAL FEATURES**
The most common symptom at onset was axial unsteadiness without rotatory vertigo, being present in six patients including all three with isolated 1PICA territory infarcts. In these patients, stroke onset was typically subacute with symptoms progressing over several hours. Acute rotatory vertigo was present at onset in three patients, all of whom had associated infarct(s) in the brainstem. One patient with a concomitant pontine infarct presented with diplopia and hemiparesis. Dysarthria, nausea and vomiting, and headaches were present only in patients with associated infarcts. All patients had pronounced signs of cerebellar dysfunction with ipsilateral limb ataxia in nine patients (defined as movements with irregular accelerations and decelerations producing oscillations on getting close to the target on the finger to nose and heel to knee tests). Limb ataxia involved arms and legs to the same extent. Dysdiadochokinesia was present in five patients and ipsilateral body sway in four. There was no case with primary position nystagmus. Multidirectional gaze evoked nystagmus was present in five patients, always in association with other infarcts. In all four patients with a coexisting infarct in the brainstem, cranial nerve and sensorimotor dysfunction was prominent and often masked the signs of cerebellar dysfunction. The table summarises these data.

**CAUSES OF STROKE**
Arterial hypertension (six patients) and cigarette smoking (four patients) were the most frequent risk factors. Two patients had fasting hypercholesterolaemia. Only one patient had chronic non-valvar atrial fibrillation. No patients had diabetes mellitus.

Local atherosclerosis with narrowing of the
ipsilateral distal (V3-V4) vertebral artery was the most frequent vascular pathology associated with IPICA territory infarct (six patients). Local occlusion of the PICA was not visualised with certainty on MR angiographic sequences, probably because of current technical limitations (fig 4). Conventional angiography performed in one patient showed a proximal occlusion of the PICA associated with advanced atherosclerotic narrowings of the vertebrobasilar arteries. Three patients had a potential cardiac source of embolism (atrial fibrillation in one patient and left ventricular akinetic segment secondary to ischaemic heart disease in two patients). In another patient with an IPICA territory infarct and distal vertebral artery thrombosis, IgM anticoagulant antibodies were highly positive (40 MPL units).

**EVOLUTION**

In the three patients with an infarct limited to the IPICA territory, the evolution was favourable, with complete regression of the cerebellar dysfunction within one month after stroke. Four patients with associated brainstem infarct(s) only had partial improvement of cerebellar ataxia and brainstem dysfunction. In one patient with a coexisting occipital infarct, the cerebellar ataxia resolved within four weeks, but the patient still had hemianopsia and difficulties in visuospatial orientation. One month after stroke, two patients with multiple supratentorial and infratentorial infarcts still had residual hemiparesis and gait difficulties, but no signs of cerebellar ataxia.

**Discussion**

Our findings suggest that IPICA territory infarction is probably not as rare as previously assumed. In 1936 Goodhart and Davison reported a pathological finding (case 4). In a necropsy series of 15 cases of infarct limited to the PICA territory, Amarenco et al. found five cases with selective involvement of the lateral branch territory within the cerebellum. Symptoms in two patients were masked by associated supratentorial infarcts and in the other three patients IPICA infarct was an incidental necropsy finding, so that no clinical correlation could be made. In 13 additional cases of PICA territory infarcts associated with multiple cerebellar infarcts, selective
involvement of the lateral branch was not reported. Recently Amarenco et al described a patient with a transient infarct in the IPICA territory that evolved to infarction in the full territory of the PICA after five days. The clinical presentation included acute vertigo, vomiting, unsteadiness, and dysmetria. In our series of 10 patients, only three had an isolated infarct in the IPICA territory. All other patients had associated infarcts that partially or completely masked the manifestations of infrolateral cerebellar involvement, mainly in the four patients with a coexisting brainstem infarct, who had prominent signs of cranial nerve and long tract dysfunction. This probably explains why IPICA territory infarcts have not been recognised clinically yet. They are regularly overlooked on CT and the advent of MRI may allow this type of cerebellar infarct to be recognised more often. We suggest that IPICA territory infarcts may indeed be as common as those involving the territory of the medial branch of the PICA or the whole PICA territory.

All three patients with isolated IPICA territory infarct were only moderately ill at onset and developed a pure "hemispheric" cerebellar dysfunction with unsteadiness, and ipsilateral dysmetria and dysdiadochokinesia involving arms and legs to the same extent. Also, two patients had a moderate ipsilateral body sway when standing or walking. None of these three patients had eye movement disturbances or dysarthria. They had a good neurological outcome, with complete regression of cerebellar dysfunction, and persistent disability in the other patients seemed mainly related to coexisting brainstem infarct(s).

In its pure form, the clinical syndrome of IPICA territory infarct may well be distinguished from neurological dysfunction due to other cerebellar infarcts. Compared with infarcts involving the whole PICA territory or the medial branch of PICA territory, IPICA territory infarcts do not provoke a vestibular type of dysfunction with acute rotatory vertigo, primary position nystagmus, or truncal ataxia. As the lateral branch does not participate in the supply to the dorsal medullar territory, its occlusion never results in medullary infarction. Also, contrary to AICA territory infarcts, isolated IPICA territory infarcts are not associated with signs of latero-pontine involvement. Finally, IPICA territory infarcts do not produce dysarthria, which is one of the leading manifestations of SCA territory infarcts.

Our findings emphasise that, unlike other cerebellar infarcts, IPICA territory infarcts are characterised by isolated signs of hemispheric cerebellar dysfunction. This is in accordance with the purely hemispheric territory of the lower cerebellum which is supplied by the IPICA. Large artery atherosclerosis may be a more common aetiology than in other cerebellar infarcts.