Acute occlusion of a giant aneurysm of the internal carotid artery: Recanalisation of the middle cerebral artery through the contralateral carotid artery

Oclusión aguda de aneurisma gigante de arteria carótida interna: Recanalización de la arteria cerebral media a través de la arteria carótida contralateral

Dear Editor,

Early recanalisation of an occluded vessel is essential for good outcomes in patients with acute stroke. When access to the occlusion is impossible, the contralateral carotid artery may be approached via the anterior communicating artery (ACoA). We present the case of a patient with an occlusion in the distal segment of the internal carotid artery (ICA) (carotid-T occlusion) caused by an embolus in a giant aneurysm in the cavernous segment of the ICA which migrated cranially. Intracranial stenting was performed through the contralateral ICA and the ACoA.

Introduction

Occlusion of the cervical ICA resulting from thrombosis of a giant aneurysm in the cavernous segment of the ICA limits the viability of endovascular treatment. Several authors have used microcatheters to reach the emboli through ipsilateral vessels and administer intra-arterial fibrinolytic therapy. The literature also reports some cases of anterior-to-posterior circulation approach through the contralateral ICA using Penumbra devices.

We present the first case of stent placement through the ACoA in a patient with acute stroke due to carotid-T occlusion caused by an embolus from a giant aneurysm in the cavernous segment of the ICA.

Clinical case

Our patient was a 53-year-old left-handed man with no relevant history who visited our hospital due to sudden loss of consciousness. After the patient recovered consciousness, he displayed severe left-sided hemiplegia, left facial palsy, and dysarthria (NIHSS score of 14). Code stroke was activated upon arrival at the emergency department; time from symptom onset to arrival at the emergency department was 90 minutes. A brain CT scan detected an expansive process in the temporal region involving the ICA and early signs of infarction in the territory of the right middle cerebral artery (MCA) (ASPECTS score of 6). CT angiography revealed complete occlusion of the origin of the ICA resulting from thrombosis of a giant aneurysm in the ICA; occlusion extended towards segments A1 and M1 of the anterior cerebral artery (ACA) and MCA, respectively (carotid-T occlusion) (Fig. 1A and B).

Angiography was performed with the Seldinger technique; we studied right intracranial circulation through the left ICA. Contrast injected into the left carotid artery was observed to flow into the ACoA towards the right AZ segment (Fig. 1C and D); we therefore decided to use a microcatheter to navigate through the ACoA in order to place a stent between right A1 and M1 to open the right distal ICA occlusion. To this end, we placed a 7F sheath introducer measuring 80 cm (Super Arrow-Flex®) in the left common carotid artery (CCA); the left cervical ICA was catheterised using a Navien microcatheter of 0.072" inner diameter (Covidien®). After administering 10 mg abciximab intravenously, we placed 2 stents (Codman® Enterprise vascular reconstruction device) using a PROWLER SELECT Plus microcatheter (Codman®) and a Synchro-14 guidewire (Stryker®). An angiography (parenchymal phase) performed at the end of the procedure showed contrast passage through the branches of the right MCA (Fig. 1E-G). Recanalisation was achieved at 420 minutes after symptom onset. Following our hospital’s protocol, our patient began dual antiplatelet
Figure 1  (A) Hyperdense lesion in the region of the cavernous segment of the right ICA corresponding to a thrombosed giant aneurysm. (B) CT-angiography of the circle of Willis revealing no blood flow in the right MCA territory. (C) Digital subtraction angiography with contrast injection into the right CCA revealed no blood flow in the right ICA. (D) Angiography of the left CCA (intracranial projection) showing no abnormalities in the ACoA, adequate filling of the distal portion of right A1, right ACA elevation, and T occlusion in the terminal segment of the right ICA. (E) Selective catheterisation of the right ACA and MCA via the ACoA. (F) Use of Enterprise® stents between the right MCA and the right carotid artery. (G) Angiography (parenchymal phase) showing recanalisation type 2b according to the mTICI scale, associated with delayed venous drainage compared to the left hemisphere.
therapy (clopidogrel 75 mg plus acetylsalicylic acid 100 mg orally every 24 hours).

Within the following 24-48 hours, our patient showed a decreased level of consciousness; a brain CT scan revealed extensive oedema in the infarcted area and uncal and subfalcine herniation requiring decompressive craniectomy. Our patient was discharged after 30 days of hospital stay; he progressed favourably and experienced no complications. He scored 1 on the mRS. Mild distal paresis of the left arm and mild postural tremor persisted.

Four months later, our patient visited our department due to fronto-orbital oppressive headache, tearing, and conjunctival chemosis, suggesting cavernous sinus involvement. An MRI scan and MR angiography confirmed recanalisation of the right carotid artery and the giant aneurysm (Fig. 2). The case was discussed in a meeting of all members of our multidisciplinary team; a decision was made to perform endovascular occlusion of the aneurysm (Fig. 3).

Our patient progressed favourably and symptoms associated with cavernous sinus compression by the aneurysm resolved. At 6 months, another craniectomy was performed and dual antiplatelet therapy was continued. At one year, our patient scored 1 on the mRS; he displayed disempathy and mild hemiparesis of the left arm, which was treated with rehabilitation.

Discussion

Stroke due to acute ICA occlusion accounts for 15% to 25% of all cases of acute stroke. Prognosis in these patients is poor; the recanalisation rate after intravenous thrombolysis barely reaches 10%. Although recanalisation has traditionally been a controversial treatment in cases of acute ICA occlusion, several studies and randomised trials present it as a technically viable and clinically effective treatment alternative for these patients.

Moret et al. were the first authors to use an alternative pathway to access an occluded vessel; these authors used the posterior communicating artery. Based on this approach, Hui et al. treated a patient with acute stroke secondary to MCA occlusion using intraarterial thrombolysis and thrombectomy. These authors intended to navigate through the posterior communicating artery with a Penumbra device.
LETTERS TO THE EDITOR

In cases of non-recanalisable ipsilateral ICA, endovascular treatment of MCA occlusion may be performed via the contralateral carotid artery when the anatomical configuration of the ACoAs and ACAs is favourable.

Conflicts of interest

The authors have no conflicts of interest to declare.

References

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Recurrent diaphoresis, acute confusional state, and pleural mesothelioma

Diaforesis recurrente, estado confusional agudo y mesotelioma pleural

Dear Editor:

Daytime and night-time excessive, generalised recurrent sweating is an infrequent and under-reported symptom in elderly patients. When present, however, it may point to a number of conditions, including tumours. Acute confusional state (ACE) or delirium is frequent among elderly patients and may be the reason for institutionalisation or an intermittent condition in institutionalised patients. Cancer is a possible diagnosis for this manifestation. Both diaphoresis and delirium are regarded as syndromes due to the wide range of causes potentially triggering them.

We present the case of a 79-year-old man who was brought in the morning to the emergency department of our hospital due to behavioural changes, inattention, incoherent thinking, and visual hallucinations. The night before he had displayed normal behaviour and cognition. He had no history of vascular risk factors and had previously been diagnosed with right bundle branch block. Our patient had been taking 50 µg levothyroxine during fasting for the last 20 years and was not receiving any psychoactive drugs. Over the previous year he had experienced episodes of excessive sweating in the evening and night 2-3 times per week, which forced him to change clothes. These episodes were not associated with hyperthermia; complete blood counts performed on 2 occasions in the previous months revealed normal results. He displayed diaphoresis the night before admission. Physical examination revealed no abnormalities except for mild gait instability, increased base of support, and side stepping.

Our patient displayed cognitive and behavioural alterations and was inattentive; we were unable to perform more specific tests due to lack of patient cooperation. He met the diagnostic criteria for ACE. Results from the heart examination were normal except for right bundle branch block; he displayed normal heart rate and blood pressure level, and required no additional medications. A brain CT scan revealed moderate cortical and central atrophy, a typical finding for his age, and no lesions in or around the brain parenchyma. A complete blood count showed normal results except for low glucose levels (26 mg per cent). We suspected a typing error since our patient was not diabetic and was not taking any antidiabetic drugs. A blood sugar test revealed a glucose level of 28 mg per cent; we therefore started treatment with glucose 10%. Chest radiography revealed a large radiopaque mass on the left side (Fig. 1A). ACE resolved 2 hours after glucose infusion. An insulin test disclosed 7.5 mIU/mL (normal range: <10 mIU/mL). A lung biopsy revealed mesothelioma with typical cells and reactive fibrosis (Fig. 1B). We ruled out asbestos exposure based on reports from our patient’s relatives. Our patient was discharged one week later. We decided not to excise the mesothelioma in view of its proximity to the mediastinum and our patient’s advanced age. He received medications similar to those prior to admission, started a carbohydrate-rich diet, and was instructed to eat every 3 hours, even at night. Our patient was followed-up every 3 months for 5 years; he experienced no further episodes of diaphoresis or ACE, his cognitive function was normal, and he was independent for daily living activities. Follow-up chest radiographies were performed every 4 months and revealed no changes compared to the image shown in Fig. 1A. Our patient was diagnosed with benign pleural fibrous mesothelioma.

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