

Stroke as a first manifestation of Takayasu disease I.A. Ionescu¹, O. Rujan¹, A-M. Enachi¹, V. Bucica¹, A. Pavel¹, G. Bododea³, C. Baetu¹, G. Mihailescu^{1,2}, I. Buraga^{1,2} 1. Department of Neurology, "Colentina" Hospital, Bucharest, Romania 2. Department of Neurology, "Carol Davila" University of Medicine and Pharmacy, Bucharest, Romania 3. Department of Neurology, Ramnicu Valcea County Hospital, Romania

Objectives:

•chronic inflammatory disease of unknown etiology, which affects the aorta and its branches •incidence of stroke in patients with TA is 10-20%

•our case consists in a patient afflicted by TA who developed a stroke in territory irrigated by the medial cerebral artery (MCA), without previous symptoms



- TA a nonspecific chronic arteritis, diagnosed mostly in young women (2nd or 3dr decade of life).
- thickening of the arterial wall due to inflammation leads to fibrosis, stenosis, occlusion, thrombosis and forming of aneurysms
- clinical from minimal changes like a diminished pulse to sever neurological complications
- Neurological manifestations appear in 50% of the cases (headache, vertigo, amaurosis, seizures)

Case description:

- a 40 year old female with left hemiplegia (involving the face also) and left hemianesthesia.
- CT scan : an acute infarction in the right MCA territory.
- Clinicaly: a difference in the arterial tension values measured on both arms and an absence in the peripheral pulse in both legs (pedis artery)
- transthoracic echography : the absence of subclavian and carotidian flux on the right side
- a chest and cervical CT scan confirmed the occlusion in the right subclavian and carotid artery and revealed the thickening of the thoracic aorta and a low pulse in the left carotid artery
- the blood tests : normal parameters (except a medium elevated VSH)
- a temporal artery biopsy normal
- The patient received the specific treatment for acute stroke and corticotherapy
- an angiography was performed and two stents were inserted in the left carotid artery.

CONCLUSIONS:

What sets this case aside is the fact that our patient developed a cerebral infarction in the whole MCA territory due to vascular changes secondary to Takayasu disease which was otherwise asymptomatic till then.

References:

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