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ABSTRACT

Percutaneous coronary intervention is a coronary revascularization procedure that may rarely result in thromboembolic events. Although infrequent, ophthalmological complications of percutaneous interventions include a wide range of clinical presentations, with differing severity and outcomes. In this case report, an 83-year-old woman, with multiple cardiovascular risk factors, presents with horizontal diplopia after a percutaneous transluminal coronary angioplasty. After ophthalmological evaluation and a head computed tomography scan, the diagnosis of isolated ischemic internuclear ophthalmoplegia was established. After six months of follow-up, the patient showed complete recovery of her symptoms and ocular movements. We discuss the post-percutaneous intervention ophthalmic complications that, although uncommon, must be recognized by health care providers.

Keywords: Diplopia; Myocardial Revascularization; Ocular Motility Disorders; Percutaneous Coronary Intervention

RESUMO

A intervenção coronária percutânea é um procedimento de revascularização coronária que pode, raramente, resultar em eventos tromboembólicos. Apesar de incomuns, as complicações oftalmológicas da intervenção percutânea incluem um largo espectro de manifestações clínicas com diferentes prognósticos. Reportamos um caso clínico de uma mulher de 83 anos, com múltiplos fatores de risco cardiovascular, que apresenta diplopia horizontal após uma angioplastia coronária percutânea. O diagnóstico de oftalmoplegia internuclear isquémica isolada foi estabelecido após avaliação oftalmológica e realização de tomografia computadorizada cranioencefálica. Seis meses após a apresentação, a doente apresentava recuperação completa dos sintomas e dos movimentos oculares. Os autores discutem as complicações oftalmológicas da intervenção percutânea que, apesar de incomuns, devem ser reconhecidas pelos profissionais de saúde.

Palavras-chave: Diplopia; Intervenção coronária percutânea; Revascularização Miocárdica; Transtornos da Motilidade Ocular

INTRODUCTION

Cardiovascular disease is the leading cause of mortality worldwide.¹ Percutaneous coronary intervention (PCI), or coronary angioplasty, is performed to diagnose and treat significant coronary artery disease (CAD).² Although less invasive than coronary artery bypass grafting, PCI is associated with some complications, mainly thromboembolic events, leading to organ infarction.^{3,4} Ophthalmological complications of PCI include a wide spectrum of clinical manifestations, ranging from transient visual disturbances to permanent and devastating situations. These complications tend to occur in the immediate postoperative period and urgent treatment may prevent permanent vision loss.^{3,4}

We report a case of post-PCI acute diplopia resulting from internuclear ophthalmoplegia (INO) and discuss the risks and management of post-PCI ophthalmological complications.

CASE REPORT

An 83-year-old woman with coronary heart disease, type 2 diabetes mellitus, hypertension and dyslipidaemia, underwent elective PCI with everolimus-eluting stent placement in the anterior descending artery. Within 24 hours after the procedure, the patient developed isolated horizontal diplopia on the right gaze. Upon ophthalmologic evaluation, the patient presented a best-corrected visual acuity of 5/10 in both eyes and intraocular lens posterior capsule opaci-

fication. Neuro-ophthalmic evaluation revealed adduction limitation of the left eye and abduction nystagmus of the right eye (Fig. 1). Convergence and other eye movements were intact. There was no anisocoria, pupillary light reflex deficits or ptosis. No other neurological signs were found.

Emergent computed tomography (CT) scan of the brain showed normal findings for the patient's age. The clinical findings were consistent with left INO, probably caused by an ischemic embolic event following PCI. The patient was discharged with the recommended double antiplatelet therapy after drug-eluting stent implantation and kept under observation with regular consultations.

During the first six months of follow up, the patient's symptoms improved progressively with a complete resolution of the diplopia. Ophthalmological re-evaluation revealed normal ocular movements without nystagmus (Fig. 2).

DISCUSSION

The conjugate horizontal movements of the eyes are mediated by the paramedian pontine reticular formation (PPRF), which receives information from the contralateral frontal lobe and projects to the ipsilateral abducens nucleus (nVI). Motor neurons of the nVI, are responsible for the stimulus of the lateral rectus muscle. Interneurons of the nVI cross the midline, travel through the medial longitudinal

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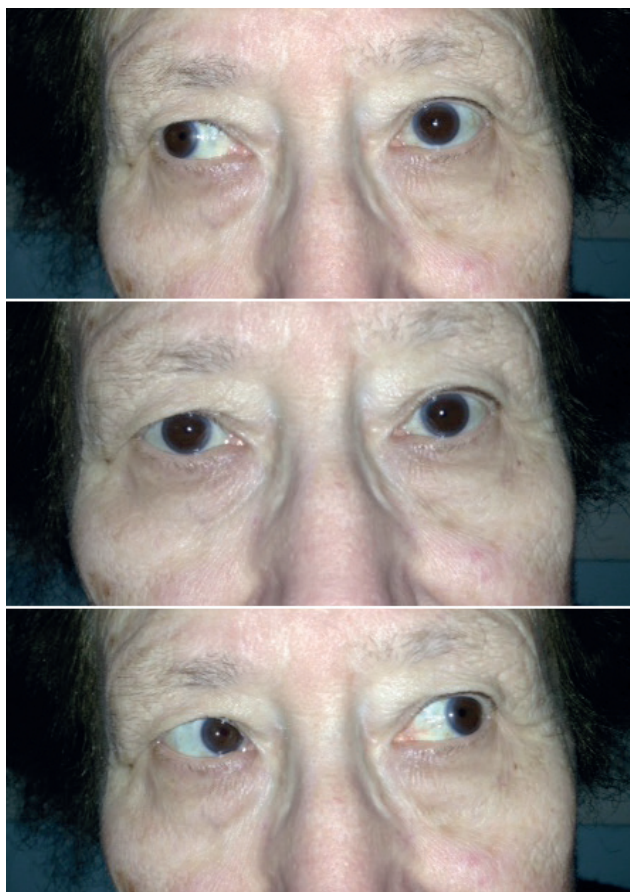


Figure 1 – Photographs of ocular movements at presentation

fasciculus (MLF) in the dorsomedial brainstem, and project in the contralateral medial rectus subnucleus of the oculomotor nuclear complex (nIII).⁵ Therefore, the activation of the nVI generates ipsilateral horizontal gaze (see Fig. 3).

A lesion in the MFL affects the conjugate movements and causes INO. This is one of the most localized brainstem syndromes, and manifests with ipsilateral adduction deficit and abduction nystagmus of the contralateral eye when gazing to the side opposite the lesion. The most common symptom is horizontal diplopia on conjugate lateral gaze and other clinical findings may be present according to the extent of the lesion and its location.^{5,6} The MFL is irrigated almost exclusively by end-arteries from small perforating arteries originating at the top of the basilar artery. Because of this vascular supply, it is particularly vulnerable to ischemia and microemboli.⁷

Isolated INO related to PCI accounted for 4.5% of a series of 110 patients diagnosed with INO.⁸ In this setting, INO may result from ischemia in the context of microembolization or thrombosis related with hypotension, pre-existing vascular disease or dehydration. INO related with PCI showed good prognosis with resolution of diplopia in a few months.⁸

Several other ophthalmic complications arising from PCI have been reported in the literature. These events include retinal complications and neuro-ophthalmic complications.

Retinal complications range from asymptomatic findings

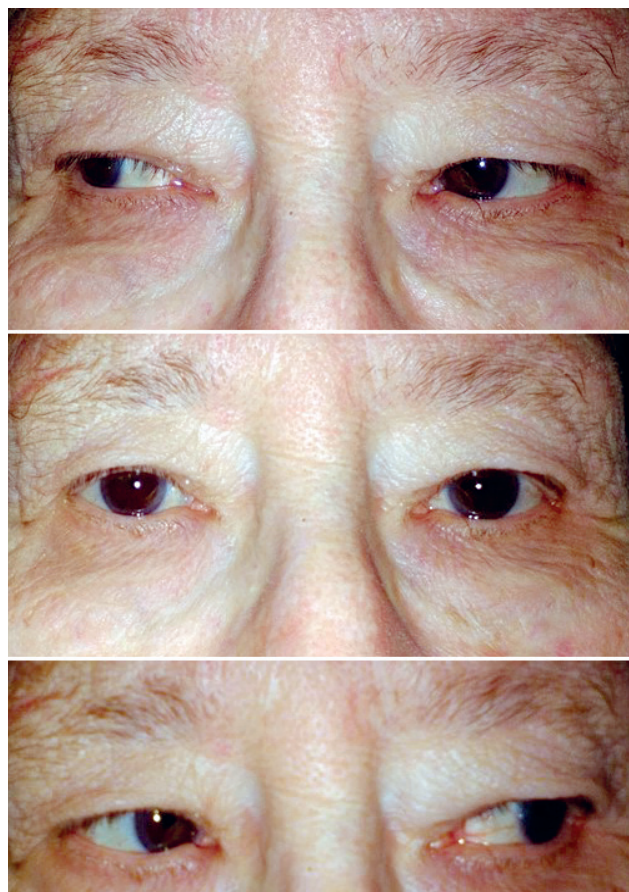


Figure 2 – Photographs of ocular movements six months after presentation

(cotton wool spots and superficial haemorrhages) to severe retinal thromboembolic events with vision loss.^{2,4,5} New retinal microemboli have been reported with an incidence ranging from 0% to 15% after heart catheterization procedures.² Kojuri *et al* reported an incidence of 6.33% in new post-PCI retinal emboli in asymptomatic patients. With the ophthalmic artery being the first major branch of the internal carotid artery, emboli originating from atheromatous plaques, thrombus of the catheter tip, and, rarely, foreign materials from the catheter or guide wire, may travel from the heart towards the eyes.⁴ Severe thromboembolic events in the eyes include branch retinal artery occlusion (BRAO) and central retinal artery occlusion (CRAO). Retinal artery occlusion is an ophthalmological emergency since immediate reperfusion of retinal vessels, within six hours, is believed to reverse or reduce tissue damage and prevent permanent lesions.³

Neuro-ophthalmological complications of PCI may arise from thromboembolic events in several important nuclei of ocular motility,² including isolated partial third nerve palsy;⁹ partial lateral rectus palsy;¹⁰ and bilateral ophthalmoplegia.¹¹ Supranuclear ophthalmoplegia¹² and skew deviation¹³ have also been reported following PCI.

Another important neuro-ophthalmological complication is cortical blindness, with an estimated incidence of 0.05% - 1%, which manifests as an acute loss of vision

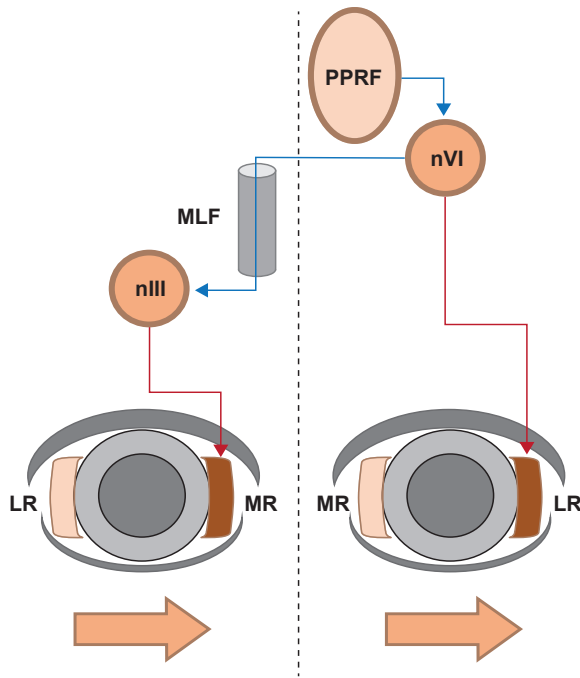


Figure 3 – MLF pathway

PPRF: paramedian pontine reticular formation; nVI: abducens nucleus; nIII: oculomotor nucleus; MLF: medial longitudinal fasciculus; LR: lateral rectus muscle; MR: medial rectus muscle.

immediately after the procedure, usually with complete recovery in the following days. Although its pathophysiology remains uncertain, it is thought to result from occipital cortex damaged by contrast agents due to a focal permeability of the blood-brain barrier.^{14,15}

Diplopia or other ocular symptoms in patients that underwent a recent coronary revascularization procedure warrant urgent ophthalmological consultation. Ophthalmological and systemic evaluation, as well as a CT scan of the brain, are indicated to rule out differential diagnoses, namely other cerebral vascular events. CT scan is also essential for the diagnosis of contrast-induced neurotoxicity.^{2,3}

Special attention must be given to high risk cases, namely: older patients, the presence of systemic comorbidities

(for example, hypertension, diabetes mellitus) and specific heart conditions (associated arrhythmia, hypokinetic wall, intramural thrombus and extension of left main stem artery).^{3,9} Additional risk factors, including operator expertise³ and prolonged and technically difficult procedures,⁹ have been identified in some studies. Specific risk factors, including impaired renal function and hyperosmolar iodinated contrast agents have been associated with cortical blindness.¹⁵

CONCLUSION

In conclusion, post-PCI ophthalmological complications are infrequent and vary from a wide range of presentations and severity. Patients and their health care providers must be alert to any visual disturbances after cardiac catheterization. In most cases, the prognosis is generally favourable, with complete recovery from the initial signs or symptoms.

PROTECTION OF HUMANS AND ANIMALS

The authors declare that the procedures were followed according to the regulations established by the Clinical Research and Ethics Committee and to the 2013 Helsinki Declaration of the World Medical Association.

DATA CONFIDENTIALITY

The authors declare having followed the protocols in use at their working center regarding patients' data publication.

PATIENT CONSENT

Obtained.

CONFLICTS OF INTEREST

All authors report no conflict of interest.

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Ex-vivo Surgical Repair of a Renal Artery Aneurysm with Kidney Autotransplantation

Cirurgia Ex-vivo para Tratamento de Aneurisma da Artéria Renal com Autotransplante Renal



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ABSTRACT

Renal artery aneurysms are rare. They are most commonly degenerative, congenital or due to medial fibroplasia. Proximal aneurysms can be repaired by endovascular and *in-situ* surgical techniques. However, aneurysms of the distal renal artery and its branches require *ex-vivo* surgical repair, also known as auto-transplantation: the kidney is removed, dissected and reconstructed in cold ischemia, and put back in place. A 69-year-old woman, with hypertension, presented with bilateral renal artery aneurysms with a diameter of 3.4 cm on the right kidney and 1 cm on the left kidney. The right renal artery aneurysm, which was due to medial fibroplasia, was successfully repaired using the *ex-vivo* surgical technique. Patency was confirmed by postoperative computed tomography angiography.

Keywords: Aneurysm/surgery; Kidney Transplantation; Renal Artery/surgery; Transplantation, Autologous

RESUMO

Os aneurismas das artérias renais são raros. As etiologias mais comuns são a degenerativa, a fibroplasia da média e os defeitos congénitos. O tratamento endovascular e a cirurgia *in-situ* são adequados para os aneurismas proximais. O envolvimento da artéria renal distal e seus ramos requer tratamento cirúrgico *ex-vivo* que consiste num autotransplante: o rim é removido, dissecado e reconstruído em isquemia fria, e finalmente reimplantado. Uma mulher de 69 anos, apresentava hipertensão arterial e aneurismas bilaterais das artérias renais com 3.4 cm de diâmetro à direita e 1 cm à esquerda. O aneurisma da artéria renal direita, de etiologia fibrodissplásica, foi tratado por cirurgia *ex-vivo* com sucesso. A angiotomografia computadorizada pós-operatória revelou a permeabilidade da reconstrução renal.

Palavras-chave: Aneurisma/cirurgia; Artéria Renal/cirurgia; Transplante Autólogo; Transplante de Rim

INTRODUCTION

Renal artery aneurysms are rare. They are most commonly degenerative, congenital or due to medial fibroplasia.¹ Proximal aneurysms can be repaired by endovascular and *in-situ* surgical techniques. However, aneurysms of the distal renal artery and its branches require *ex-vivo* surgical repair, also known as auto-transplantation: the kidney is removed, dissected and reconstructed in cold ischemia, and put back in place.²

CLINICAL CASE

A 69-year-old Caucasian female presented with controlled hypertension (on four anti-hypertensive drugs), dyslipidemia, acute myocardial infarction (with a normal coronary angiography one year before), obesity, obstructive sleep apnea syndrome treated with continuous positive airway pressure (CPAP). She had a 40 pack-year smoking history until the age of 44. In the workup for bilateral low back pain, a computed tomography angiography (CTA) revealed bilateral renal artery aneurysms (Fig. 1). Glomerular

filtration measured by renal scintigraphy with ^{99m}Tc-DTPA was 54 mL/min (right), 46 mL/min (left) and there was right excretory delay without significant mechanical obstruction. Serum creatinine was 0.90 mg/dL and hemoglobin was 10.3 g/dL. The ultrasound scan of the carotid and vertebral arteries showed no changes.

Ex-vivo surgical repair of the right renal aneurysm was carried out because the aneurysm involved the branches of the renal artery. This technique has been described in detail.²⁻⁵ A right retroperitoneal flank lazy-S shaped incision, extending from the tip of the 11th rib to an area just medial to the McBurney point was used. The kidney was removed without dividing the ureter, encompassing a small ellipse of vena cava patch with the renal vein (which was closed longitudinally) to allow for a patch anastomosis, placed on the patient's abdomen, immersed in ice slush and perfused with Celsior® solution at 4°C. The hilum was dissected and three branches distal to the aneurysm were identified. The two cephalic branches were syndactylized and the combined

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