Functional Recovery of the Injured Optic Chiasma after Omental Transplantation

Hasarlı Optik Chiasmanın Omental Transplantasyon Sonrası Fonksiyonel Düzelmesi

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Abstract: The authors describe a woman with visual loss and epilepsy that were caused by removal of a meningioma on the right sphenoidal ridge in an operation 5 years prior to presentation. The patient started to improve neurologically after omental transplantation to the injured optic chiasma and right temporal lobe. The epileptic seizures virtually disappeared after this surgery, and vision gradually improved over 3 months. These results indicate that placing omentum directly on injured nervous tissue can lead to neurological improvement.

Key Words: Epilepsy, omentum, omental transplantation, optic chiasma, visual loss

INTRODUCTION

Many different lesions can cause chiasmal compression (16,24), including pituitary adenoma, suprasellar meningioma, basal leptomeningitis, craniopharyngioma, aneurysm, intrasellar cysticercosis, chordoma, and metastases. In addition, the chiasma and optic nerves can be damaged by ischemia (5,23) and trauma (12). However, aside from rehabilitation, to date there is no specific treatment aimed at improving the function of the chiasma and optic nerves after chronic injury. We report the unusual case of a young woman with visual loss and epilepsy caused by ischemia of the optic chiasma. The injury occurred during removal of a meningioma in the right sphenoidal ridge. Five years after this surgery, the patient underwent omental transplantation to the affected region, and our results suggest that the omentum improved the function of the residual nervous tissue in the visual pathway.
CASE REPORT

A 30-year-old right-handed woman from Lima, Peru was admitted for surgery. In February 1990, she presented with headache and complex partial seizures evolving to tonic-clonic generalized seizures. Despite these problems, the patient's motor and visual functions were normal. On May 22, 1990 the woman underwent surgery via a pterional approach to remove a meningioma in the medial portion of the right sphenoidal ridge. She was extubated the morning after the operation, and manifested binocular blindness at that time. Two months postsurgery, the patient began to experience two or three partial seizures per month and one tonic-clonic generalized seizure every 3 months. On admission, the patient was taking daily doses of 600 mg carbamazepine and 200 mg diphenylhydantoin.

Examination:

On physical examination, the patient's appearance was appropriate for her age and she was lucid. The ocular fundi revealed optic atrophy of the papillae, loss of retinal nerve fibers, and macular pigment epitheliopathy. She exhibited blindness and absence of a pupillary light response in her right eye. In the left eye, her visual acuity and visual field (Fig. 1A) were reduced to distinguishing shades at a distance of 30 cm, she was unable to identify colors, and she lacked the ability for central fixing. The remainder of the neurological examination was normal. Magnetic resonance imaging (MRI) scans demonstrated an infarct and a cystic formation in the right temporal pole (Fig. 2A). The visual evoked response (VER) revealed absence of waves in the right eye, and P100 waves with poor amplitude and latency of 152.3 ms in the left eye. An electroretinogram (ERG) showed absence of waves in the right eye, and “a” waves with low amplitude and latency of 37.6 ms, as well as absence of “b” waves.

Surgery:

Omental transplantation was performed on February 15, 1995 without complication. We opted to expose the chiasmatic region through the same right pterional approach that had been used previously. The surgical findings were as follows: 1) multiple adhesions between the dura mater and right temporal lobe; 2) infarct and cyst in the temporal pole; 3) hypotrophy of the chiasma and both optic nerves (left optic nerve reduced to 80% and right optic nerve to 50%); 4) marked pallor of the chiasma; 5) chiasma in normal position; 6) right ophthalmic artery of normal caliber; and 7) marked decrease in blood vessels on the dorsal surface of the chiasma.

The neurosurgical procedures for the omental transplantation were carried out according to the technique we described in earlier reports (19,20). Briefly, an end to-end anastomosis by invagination between occipital vessels and the gastroepiploic vessels were performed. Afterwards, a small segment of omentum was passed between the right optic nerve and the ophthalmic artery, and placed above the prechiasmal space and chiasma. Another omental...
segment was placed into the cyst cavity and spread over the lateral surface of the damaged temporal lobe. Finally, the surgical wound was closed in standard fashion.

Postoperative Course:

Two days after surgery, the patient had photophobia, headache, and left facial paresis. She was able to walk with assistance, and day 2 postoperative computed tomography (CT) scans confirmed the presence of omentum above ("covering" or "over") the chiasma, temporal fossa, and right orbital lobe (Fig. 2B), as well as revascularization of the underlying brain tissue.

Three months later, the patient's condition was good and she had had no epileptic seizures since the surgery. She was taking daily doses of 400 mg carbamazepine and 2 mg clonazepam. Her ophthalmologic examination revealed photophobia, finger-counting to 50 cm in the left eye, and some reactivity to light in the right pupil. At 8 months postsurgery, there was visual improvement in both eyes, and her direct and consensual pupillary light responses were intact. Using her left eye, the patient was able to identify some white, black, and gray objects at a distance of 60 cm. Postoperative MRI scans showed hypotrophy of both optic nerves and confirmed the presence of omentum above the chiasma. During this period, the patient was taking only a nightly dose of 3 mg clonazepam. The last automated perimetry (January 23, 1996) showed fixation and increase of some threshold in the left eye (Fig. 1B). Furthermore, VER testing showed P100 waves with poor amplitude and latency of 192.7 ms in the right eye, and distorted, low amplitude P100 waves with a latency of 125.6 ms in the left.

At present, 26 months after omental transplantation, the patient's findings are as follows: pupillary light response intact and can distinguish shades at a distance of 20 cm with the right eye; able to write numbers and recognize the shape and size of objects at a distance of 100 cm with the left eye. As well, she occasionally recognizes the color blue. The patient's condition is good, she is having no epileptic seizures, and is taking 3 mg clonazepam nightly.

**DISCUSSION**

We decided to transplant omental tissue to the chiasma and right temporal lobe of our patient for several reasons. First, based on the patient's clinical and neuroophthalmological findings. Second, the optic nerve is a fasciculus of white matter that is embryologically, morphologically, and physiologically similar to the central nervous system (7,18). Third, other authors have achieved neurological improvement using the omental transplantation for patients with traumatized spinal cord (1,15,17), capsular hemiparesis (19, 20), and Dejerine-Roussy syndrome (21) due to lacunar stroke. We chose the omentum because it is the best tissue for developing vascular connections with the cerebral and cerebellar cortex, and with the spinal cord and cauda equina (1,15,20). Additionally, omental tissue releases neurotransmitters and neurotrophic factors.
that are transported to the underlying nervous tissue through the penetrating omental neovessels (4).

We believe that the patient's visual improvement was primarily due, initially to the revascularization of residual vital axons in the ischemic regions and ischemic penumbra in the chiasma and optic nerves, and in the right temporal loop of visual radiation. Based on the fact that experimental (3,6,10) and clinical (15,17,20) findings have proven there is a direct relationship between revascularization and recovery of demyelinated and/or degenerated axons, we also attribute her improvement to functional recovery of other axons in different stages of demyelination and degeneration. In addition, the visual improvement suggests functional recovery of some optic fibers from the retina up to the level of the lateral geniculate nuclei, as well as the presence of geniculocalcarine axons in the right Flechsig-Meyer loop (7,18). Evidence of this is the direct light response in the patient's right pupil, as well as the improvement in visual acuity and visual field in her left eye.

These preliminary results demonstrate that there are axons in the injured optic pathway that can recover if circulation is regained. They also confirm previous experience (1,17,25) that large numbers of axons can be remyelinated or regenerated under selected conditions (3,6,10). Therefore, contrary to the opinions of other authors (2,8,11), we believe that the revascularization of the injured chiasma should be attempted even years after the damage has occurred. We also believe that omental transplantation may be beneficial in patients with nonarteritic anterior ischemic optic neuropathy (NAION). Although the etiology of NAION is unknown, this technique may help revascularize the anterior optic nerve in these cases since ischemic optic neuropathy is involved (23).

Moreover, our patient's case confirms that the placement of omental tissue on the epileptic focus aborts seizures. It does so in the manner observed previously (since May 1988) in eight patients with epilepsy and occlusive type cerebrovascular disease (20), and in a 10-year-old girl with epilepsia partialis continua (13). We believe that this excellent improvement in seizure status is due to revascularization of the epileptogenic foci in the temporal lobe, which are characterized by loss of neurons, astrocytic gliosis, loss of GABAergic axon terminals, and severe hypoxia/ischemia (9,14,22).

Similar histopathologic changes have been noted in lacunar stroke of cerebrovascular disease (20). Therefore, increasing the flow of blood, oxygen, and neurotrophic factors to the damaged hippocampal formation, and improving the function of neurons in chronic ischemia reduce the excessive excitability of the dendrites and favor neuronal regeneration. In conclusion, our patient's recovery from binocular blindness and seizures shows that placing omental tissue directly on epileptogenic foci and injured optic chiasma tissue produces significant neurological improvement.

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