Case Report

Acute Visual Loss as a First Sign of Hyperhomocysteinaemia

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A 38-year-old man presented with an acute right homonymous visual field defect due to occipital lobe infarct caused by hyperhomocysteinaemia. Visual field testing, magnetic resonance imaging, laboratory studies, and genetic analysis were carried out. On magnetic resonance imaging, a left occipital lesion with bright signal on the diffusion-weighted and fluid-attenuated inversion recovery images suggested a diagnosis of an acute infarct. Blood tests revealed raised homocysteine of 52.08 μmol/L (reference range, <15 μmol/L) and genetic analysis showed the patient to be homozygote to 5, 10-methylenetetrahydrofolate reductase deficiency. Hyperhomocysteinaemia is a rare cause of acute visual loss due to cerebral ischaemia and should always be suspected and investigated with the appropriate tests to diagnose the condition and limit further vision deterioration.

Key words: Hyperhomocysteinemia, Magnetic resonance imaging, Visual fields


Introduction
Homocystinuria is a disorder of methionine metabolism, leading to abnormal accumulation of homocysteine and its metabolites such as homocysteine and homocysteine-cysteine complex in blood and urine. Normally, these metabolites are not found in appreciable quantities in blood or urine. Hyperhomocysteinaemia, a separate but related entity, is defined as elevation of homocysteine level in blood. The reported incidence of homocystinuria varies between 1 in 50,000 and 1 in 200,000. The prevalence of hyperhomocysteinaemia in the general population is not known. Studies investigating the prevalence of homozygosity for the thermodabile variant mutation in 5, 10-methylenetetrahydrofolate reductase (MTHFR) have found a prevalence of approximately 15% in European, Middle Eastern, and Japanese populations compared with a prevalence of ≤1.4% in African Americans. The prevalence of heterozygosity is 30% to 40%.

This report is of a 38-year-old man who, with no other risk factors, presented with an acute right homonymous visual field defect as the first sign of hyperhomocysteinaemia.

Case Report
A 38-year-old man presented to the eye casualty because of sudden-onset visual field loss of his right eye, accompanied by a mild non-localized headache. He had no prior medical or significant family history. He was not taking any medications and did not smoke. His visual acuity was 20/20 in the right eye and 20/20 in the left eye. Colour vision, anterior and posterior segment examination, intraocular pressure, and pupillary responses were normal. Neurological examination was normal except for the visual field, which revealed a right homonymous hemianopia that was densest in the superior quadrant (Figure 1). The differential
diagnosis included stroke (ischaemic or haemorrhagic), vascular malformation, space-occupying lesion, demyelinating disease, and migraine.\(^1\)\(^2\)

Urgent magnetic resonance imaging (MRI) showed an occipital infarct on diffusion-weighted and fluid-attenuated inversion recovery images (Figure 2). The patient was admitted to the Department of Ophthalmology, Agios Dimitrios General Hospital, Thessaloniki, Greece, and treated with antiplatelet drugs according to the international standards.\(^3\)\(^4\) Carotid artery triplex ultrasound and echocardiogram (transoesophageal and transthoracic) did not show any abnormalities. Laboratory studies, including evaluation of coagulation (prothrombin time, activated partial thromboplastin time, thrombin time, and bleeding time) and immunological status (antinuclear antibodies and antineutrophil cytoplasmic antibodies), revealed raised levels of homocysteine in the blood of 52.08 μmol/L (reference range, <15 μmol/L). Genetic analysis showed the patient to be homozygote to MTHFR deficiency, a known cause for raised homocysteine blood levels. The patient was treated with clopidogrel bisulphate 75 mg once daily, B6 and B12 vitamin supplements, and folic acid, and discharged.

At the 2-month follow-up examination, the patient felt his vision had improved. There was improvement in the visual field in the inferior and superior quadrants (Figure 1). Blood homocysteine levels had decreased and all other laboratory tests were normal.

Four months later, a new MRI revealed left occipital lobe hyperintensity of a smaller size with signs of laminar cortical necrosis (Figure 2). Visual field examination was stable with no further deterioration.

**Discussion**

This report is of a unique patient with an acute right homonymous visual field defect being the first sign of hyperhomocysteinaemia. Only one similar patient has been reported in the literature which, contrary to this patient, had a risk factor of atrial septal defect.\(^5\)

Homocysteine levels increase in a variety of situations including poor diet (low consumption of vitamins B6, B12, and folic acid), low levels of thyroid hormone, renal disease, psoriasis, and use of certain medicines such as fibrates. Increased homocysteine level is associated with a higher risk of strokes. Several mechanisms have been suggested as the possible cause of accelerated vascular disease, including endothelial cell damage, smooth muscle cell proliferation, lipid peroxidation, upregulation of prothrombotic factors (XII and V), and downregulation of antithrombotic factors or endothelial-derived nitric oxide.\(^6\)\(^\text{a}\)

Hyperhomocysteinaemia is one of the rare causes of acute visual loss due to cerebral ischaemia and should always be suspected and investigated with the appropriate tests because several studies have shown that early diagnosis and institution of treatment, as well as dietary restriction, is likely to slow the progression of disease and reverse some of the features.

This report illustrates the need for further laboratory studies in young patients who present with symptoms of acute stroke in the absence of typical risk factors such as hypertension, smoking, and diabetes, and highlights the need for neuroimaging studies for every patient with visual field loss.

**References**

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