CASE REPORTS

Ocular reaction to timolol maleate

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Summary

A patient prescribed timolol maleate for the control of hypertension developed dryness of the eyes. The symptoms improved on withdrawal of the drug.

Vigilance for such adverse reactions must be observed in the use of any β-adrenoreceptor blocking agent.

Introduction

Adverse reactions affecting the eyes have been described with several β-adrenoreceptor blocking agents, principally practolol. Both propranolol and oxprenolol have been incriminated in causing ocular symptoms but there are no reported instances of ocular damage due to timolol. The patient reported here developed dryness of the eyes, which improved subjectively on withdrawal of timolol.

Case report

A 40-year-old Caucasian fireman was discovered to have raised blood pressure on medical examination to assess fitness for use of breathing apparatus. He had no symptoms, nor had he suffered serious ill health in the past. Examination and investigation did not show any underlying cause for his hypertension. He was treated with chlorthalidone 100 mg daily and timolol in doses of from 15 to 75 mg daily, the BP falling satisfactorily from 180/130 to 120/95 mmHg.

He first complained of dryness of the eyes and nose 14 months after starting treatment with timolol and later said that he had noticed his eyes had been dry and tender since starting the treatment. Examination showed poor tear production with punctate corneal staining after fluorescein. Some areas of the bulbar and palpebral conjunctivae were noted as staining with Rose Bengal.

His symptoms improved immediately on withdrawal of timolol. When seen again 3 months later the bulbar conjunctival staining with Rose Bengal had cleared all but for the slightest staining in the left eye. Schirmer’s test still showed poor tear production in both eyes. The raised BP was subsequently controlled with chlorthalidone 50 mg and labetalol 600 mg daily.

Comment

The adverse effects of practolol on the eyes were first described 4 years after the introduction of the drug, as were the cutaneous manifestations of hypersensitivity (Wright, 1974; Felix, Ive and Dahl, 1974). Holt and Waddington (1975) reported an oculo-cutaneous reaction to oxprenolol, and propranolol has been reported as causing an ocular reaction which resolved on changing the β-blocker (Cubey and Taylor, 1975). Timolol has not been reported as producing such side effects, although there are single reports to the Committee on Safety of Medicine (COSM) of morbilliform and psoriasiform rashes, as well as exfoliative dermatitis.

The patient reported here developed irritating dryness of the eyes and nose shortly after starting treatment with timolol. Conjunctival epithelial and corneal changes developed and there was a marked reduction in tear volume. Subjectively, he was improved on withdrawal of the drug. Objectively, tear production was not improved but the epithelial changes had nearly resolved and the corneal abnormalities were completely healed. Rahi et al. (1976) described a fibrosing polymerositis affecting the eyes, with acanthosis, and thickening of, and loss of goblet cells from, the epithelium, and chronic inflammatory changes during treatment with β-blockers. The cornea showed epithelialysis and stromal ulceration, inadequate tear production leading to epidermalization of the conjunctival epithelium – similar changes to those found in the present patient.

These changes have been found in association with an autoantibody with affinity for intercellular zones of squamous epithelium. A significant increase in antinuclear antibody titre has been described in patients taking practolol, as well as thyroid cytoplasmic antibody. These serological changes did not, however, correlate with the development of the oculo-cutaneous syndrome (Jachuck et al., 1977). In the present patient thyroglobulin antibody tests were negative as were fluorescent antibody tests, although RA latex test was positive.

Although the ocular side effects described here are not as well developed as those seen with practolol, the resemblance and the onset and partial clearing in
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relation to treatment with timolol suggest that the drug or a metabolite was responsible. In this patient there is no evidence that the adverse effects were of immunological cause. Such ocular damage may possibly occur with any of the β-adrenoreceptor blocking drugs. The absence of reports may be due to the failure of the patient to report and the doctor to enquire about symptoms, and the failure of doctors to notify the COSM.

References


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