Intravitreal ganciclovir in CMV retinitis: case report

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SUMMARY A case is reported of a 35-year-old male with established acquired immune deficiency syndrome who developed cytomegalovirus retinitis. When intravenous treatment with ganciclovir had to be withdrawn because of toxicity, one eye was treated successfully by intravitreal injections of the drug. After 15 weeks of intravitreal treatment this eye developed endophthalmitis.

A frequent ophthalmological complication of the acquired immune deficiency syndrome (AIDS) is cytomegalovirus (CMV) retinitis.1–4 Until recently this blinding condition was untreatable. However, the acyclovir analogue ganciclovir given intravenously has been shown to be effective in CMV retinitis,5 with an 88% response rate.6 There are disadvantages to its use. Firstly, it is a virostatic agent and maintenance treatment is required.6 Second, it is toxic, and neutropenia occurs in up to 38% of patients on maintenance treatment.8 Thirdly, 'breakthrough' disease is encountered on maintenance therapy in up to 50% of patients.6 We report on a patient who had intravitreal injections of ganciclovir when systemic treatment had to be abandoned owing to neutropenia.

Case report

A 35-year-old male homosexual with established AIDS presented in March 1987 with painless loss of vision in the left eye of three days' duration. Ocular examination showed a corrected visual acuity of 6/5 right and 6/9 left, with extensive loss of temporal visual field in the left eye. Biomicroscopy revealed flare in the aqueous of the left eye, and on funduscopy there were the typical changes of CMV retinitis involving the nasal retina of the left eye up to and surrounding the optic nerve head. In addition there was a haemorrhagic patch of retinitis just inside the inferior temporal arcade of the right retina 2–3 disc diameters in area (Fig. 1).

The patient was admitted to hospital and started on intravenous ganciclovir 5 mg/kg daily. On admission his white cell count was 1.8 x 10⁹/l, and rectal biopsy showed mild proctitis with CMV inclusion particles. After an initial decline in the vision of the left eye the disease came under control, and maintenance treatment of 5 mg/kg on three days per week was started. During this period his white cell count fell, the dosage of maintenance treatment was halved, but after 12 weeks of treatment his white cell count had fallen to 0.8 x 10⁹/l and intravenous ganciclovir had to be discontinued. His visual acuity at this time was 6/12 right, 6/24, left.

We decided to treat the right eye by intravitreal injections of ganciclovir. Injections of 200 μg ganciclovir in 0.1 ml of normal saline were prepared in a sterile cabinet and passed through a Micropore filter giving a near maximum soluble solution of pH 10.14. After topical anaesthesia with amethocaine 1% drops the injections were made through the inferior pars plana via a 27 gauge needle on a tuberculin syringe. The circulation at the optic nerve head was checked after each injection; we did not find it necessary to perform an anterior paracentesis. Topical chloramphenicol 0.5% drops were given four times a day, on the days preceding and following as well as on the day of each injection. The treatment was given twice weekly for three weeks and weekly thereafter. The injections were generally well tolerated. The retinitis in the right eye showed a good response to this treatment while the retinitis in the left eye showed rapid progression. By the thirteenth week of intravitreal therapy the visual acuity of the untreated left eye was hand movements, that of the right eye remained at 6/12, but funduscopy showed new haemorrhages and exudates extending on both sides of the fovea (Fig. 2).

After 15 weeks of intravitreal treatment (total dose 30 mg) the patient returned to the clinic with a painful, red right eye. He was admitted to hospital for treatment of a right endophthalmitis. A specimen from diagnostic vitreous biopsy grew a mixture of two
coagulase-negative staphylococci, both resistant to chloramphenicol and one also to methicillin. His endophthalmitis responded to treatment, the vision recovering from perception of light to counting fingers. Concurrently he had developed a recurrent pneumocystis pneumonia and died 18 days later. Permission for post-mortem examination was not obtained.

Discussion

In 1985 Pulido et al. found no ophthalmoscopic or histological changes on the retina after single-dose injections of ganciclovir into rabbit vitreous. Henry et al. have reported the use of this agent via intra-vitreal injection in one case of CMV retinitis. Using an ID$_{50}$ (dose sufficient to reduce viral replication to
50% of 2-87 μg Henry et al. calculated that the intravitreal concentration of the drug remained above this level for approximately 62 hours after a single intravitreal injection of 200 μg. They were able to control the retinitis in their patient with weekly injections. Our case confirms that intravitreal administration of ganciclovir can be used as an alternative route in cases where the use of intravenous therapy is limited by the occurrence of leucopenia. The late recurrence of disease raises the question of whether maintenance treatment of weekly 200 μg is sufficient to control the retinitis adequately. A larger dosage may be desirable, but 200 μg in 0.1 ml is close to the maximum solubility for the drug. Injecting larger volumes intravitreally would certainly require anterior paracentesis to prevent increased intraocular pressure. It remains to be proved that repeated injections of this alkaline agent are not retinotoxic.

This case was complicated by the endophthalmitis, which was always a possibility in this situation. Staphylococci are the commonest cause of endophthalmitis and are remarkable for their ability to develop resistance to antibiotics. This makes the choice of prophylactic agents difficult.

In conclusion, ganciclovir has been found to be an effective agent when used intravitreally in CMV retinitis. A maintenance dose of 200 μg weekly does not exclude the possibility of breakthrough of the disease. Endophthalmitis is a constant threat when repeated intravitreal injections are given.

References


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