Visual Function and Structure Outcomes in a Patient with Diabetic Macular Oedema Insufficiently Responsive to Available Therapies – Treatment with Fluocinolone Acetonide Implant

Érica ABC Guerreiro Paulo and Claus Eckardt

1. Ophthalmology Resident, Department of Ophthalmology, Klinikum Frankfurt Höchst, Frankfurt am Main, Germany; 2. Chief, Department of Ophthalmology, Klinikum Frankfurt Höchst, Frankfurt am Main, Germany

Abstract

Importance: Early experience with intraocular, non-biodegradable fluocinolone acetonide (FAc) (0.2 µg/day) implant (ILUVIEN®), comparing visual function before and after treatment in an insufficiently responsive diabetic macular oedema (DMO) patient. Observations: This 62-year-old male patient with type 2 diabetes mellitus was first treated for DMO in 2004, when best corrected visual acuity (BCVA) was 0.8 for his right eye. He did not visit an ophthalmologist again until 2011 when BCVA had declined to 0.2. Three separate, monthly intravitreal (IV) ranibizumab injections and additional laser photocoagulation resulted in no improvement. Subsequently, 0.7 mg IV dexamethasone was administered, giving short-term DMO improvement. However, six further IV ranibizumab injections produced no effect and a combined injection of IV ranibizumab with 0.7 mg IV dexamethasone provided only short-term improvements. Following phacoemulsification, a 0.2 µg/day FAc implant was administered. Optical coherence tomography (OCT) indicated complete DMO regression after 3 weeks, sustained 6 months post-implant. BCVA improved to 0.25 and the patient reported greater vision-related quality of life. Intraocular pressure increased gradually but resolved with daily timolol/dorzolamide and tafluoprost eye drops. Conclusions and relevance: In a DMO patient showing insufficient response to IV ranibizumab and dexamethasone injections, FAc implant provided an effective therapeutic option with manageable side effects.

Keywords

Diabetic macular oedema, fluocinolone acetonide

Diabetic retinopathy (DR) is the most common cause of moderate-to-severe visual impairment among working class adults.1 Diabetic macular oedema (DMO), a common manifestation of DR, causes central vision loss,2 and is a consequence of vascular inner blood–retinal barrier breakdown.3 Global prevalence of DR among diabetic individuals is approximately 35 %, with DMO present in 6.8 %.4 DMO is difficult to manage due to its chronicity. Historically, the standard of care was laser therapy; however, a large number of patients continued to lose vision,5 with significant vision recovery in only 15 % of treated patients.6 In particular, patients presenting with diffuse DMO, where causal vascular abnormalities are not restricted to discrete foci, are likely to be refractory to laser therapy.7 Given the proportion of patients with DMO refractory to laser therapy, new therapies providing sustained benefit are needed.8

Alternative DMO treatments are evolving rapidly, with a number demonstrating promise in preventing visual acuity (VA) deterioration and improving vision.9 One such treatment is fluocinolone acetonide (FAc) 0.2 µg/day (ILUVIEN®), a non-biodegradable, intravitreal (IV) corticosteroid implant,10 which releases daily sustained, low-dose FAc for 3 years.11 The Fluocinolone Acetonide in Diabetic Macular Edema (FAME) study compared 0.2 µg/day FAc with sham injections in patients with DMO, a baseline best corrected VA (BCVA) in Early Treatment Diabetic Retinopathy Study (ETDRS) letter score between 19–68 (20/50–20/400), central foveal thickness (CFT) ≥250 µm, and ≥1 prior focal laser treatment.12 Patients receiving non-protocol treatments were included. Patients were eligible for rescue laser after 6 weeks or retreatment with assigned therapy after 1 year.

Three-year treatment with 0.2 µg/day FAc improved BCVA in 34.0 % of chronic (≥3 years median duration) DMO patients compared with 13.4 % sham-treated patients, with mean changes in BCVA score of 7.6 and 1.8 letters, respectively.10 CFT rapidly and substantially decreased from baseline in both FAc- and sham-treated chronic DMO patients.12 Although at 36 months FAc implant patients needed cataract surgery more frequently than those patients who received sham injection, their post-surgery visual benefit was similar to subjects pseudophakic at baseline. Surgery to manage elevated intraocular pressure (IOP) was also more frequent for FAc-treated patients compared with sham (4.8 % versus 0.5 %).10
Diabetic Macular Oedema Case Study

Figure 1: Best Corrected Visual Acuity (A) and Central Foveal Thickness (B) of the Right Eye with Chronic Diabetic Macular Oedema Measured Over Time

A. Fundus photography showing right eye (RE) diabetic maculopathy in a patient with diabetic macular oedema. B. Optical coherence tomography (OCT) scan of the RE before treatment with 0.2 µg/day fluocinolone acetonide (FAc) implant; best corrected visual acuity (BCVA) 0.2; central foveal thickness (CFT) 506 µm. C. 3 weeks after 0.2 µg/day FAc: BCVA 0.25; CFT 298 µm. D. 7 months after 0.2 µg/day FAc: BCVA 0.25; CFT 256 µm. Little long-term improvement was seen in foveal thickness from the time initial RE treatment began to just before 0.2 µg/day FAc implantation. Following 320 µm. D. 7 months after 0.2 µg/day FAc: BCVA 0.25; CFT 298 µm. E. 12 months after 0.2 µg/day FAc implant: BCVA 0.25; CFT 223 µm.

The Fluocinolone Acetonide for Macular Edema (FAME) study demonstrated a single 0.2 µg/day FAc implant can provide significant, long-term BCVA and CFT improvements in chronic DMO patients. In this case study, a single 0.2 µg/day FAc implant improved BCVA and CFT when the patient was insufficiently responsive to all previous treatments, including ranibizumab, laser photocoagulation and IV ranibizumab injections in the RE, administered between September and November 2011, did not improve BCVA and focal laser photocoagulation was provided: IOP was 16 mmHg. A 0.7 mg dexamethasone implant in December 2011 produced short-term CFT decrease (442 µm to 347 µm) without improving BCVA (see Figures 1A and B). Six further IV ranibizumab injections over 8 months had no effect on visual outcomes, with CFT worsening (347 to 488 µm; Figure 1B) and IOP increasing (22 mmHg). Following IV ranibizumab injection combined with 0.7 mg dexamethasone implant in October 2012, CFT decreased to 223 µm and BCVA improved from 0.1 to 0.2, lasting 4 months. Two further ranibizumab injections gave no visual improvement. After phacoemulsification in April 2013, and one further ranibizumab injection in May, BCVA (0.25 to 0.2) and CFT (350 to 506 µm) worsened. IOP was 19 mmHg.

In July 2013 the patient had a 0.2 µg/day FAc implant inserted. Within 3 weeks, OCT indicated complete DMO regression that was not achieved with any of the previous IV therapies, resulting in CFT decreasing by 58 % after 10 weeks (548 µm to 232 µm; Figure 1B and Figure 2). These improvements were still apparent 6 months following 0.2 µg/day FAc implantation, with a concurrent increase in BCVA to 0.3 at its highest point (a clinically meaningful change from the previous BCVA of 0.2). In January 2014, OCT showed DMO with a progression of CFT (289 µm). Bevacizumab was injected twice, 3 weeks apart in February 2014, leading to regression of DMO (CFT 282 µm; BCVA 0.25). In July 2014, FAc was seen to be further reduced to 214 µm, and another bevacizumab injection was administered. The patient reported greater vision-related quality of life and the ability to perform daily routines without problems, with no complaints regarding 0.2 µg/day FAc implant therapy.

Management of Intraocular Pressure

Following 0.2 µg/day FAc implantation, gradual elevation in IOP was observed over the first 2 months (from 16 mmHg to 28 mmHg). This was resolved with daily dorzolamide/timolol eye drops (21 mmHg at 6 months). In February 2014, tafluprost eye drops were prescribed in addition to further reduce the IOP. The daily application of eye drops was not a concern for the patient and did not interfere with his quality of life.

Comments

Randomized clinical trials indicate intraocular vascular endothelial growth factor (VEGF) inhibition provides significant BCVA and CFT improvements in patients with DMO. While this is a widely used therapeutic strategy in patients refractory to laser therapy, other studies have shown that anti-VEGF therapy alone is not effective in all patients with DMO, as shown in this case.

The Fluocinolone Acetonide for Macular Edema (FAME) study demonstrated a single IV injection FAc implant can provide significant, long-term BCVA and CFT improvements in chronic DMO patients. In this case study, a single 0.2 µg/day FAc implant improved BCVA and CFT when the patient was insufficiently responsive to all previous treatments, including ranibizumab, laser photocoagulation and CFT measured using spectral optical coherence tomography (OCT). IV ranibizumab injection in 2011 gave no benefit for LE BCVA and given the lack of response to therapy and the long history of DMO, no further treatment was administered to the LE at this time.

Report of a Case

A retired, 62-year-old male patient with type 2 diabetes mellitus since 1992 (insulin dependent since 2002) was first treated for DMO in 2004, for which he received left eye (LE) laser photocoagulation and triamcinolone injection when his BCVA was 0.2 for his LE and 0.8 for his right eye (RE). Lost to follow-up until 2011, BCVA had deteriorated to 0.2 in both eyes. From 2011, the patient’s response to therapy was evaluated using BCVA...
and dexamethasone implant. However, dexamethasone implant provides release of corticosteroid over a shorter timeframe compared with FAC implant,60 possibly accounting for the relatively short duration of CFT decrease and lack of VA response observed when administered.

FAC implant gave marked improvements of BCVA and with complete regression of MO after 3 weeks, consistent with the FAME study,7 which were still apparent after 6 months. Additional anti-VEGF therapy in February 2014 appeared to increase these benefits, reflecting the multifactorial nature of DMO.16 IOP was elevated with 0.2 µg/day FAC, but was effectively managed by twice-daily dorzolamide/timolol and once-daily tafluprost eye drops. Given the lack of response to other currently available therapies, and the improvement in vision, the benefit–risk ratio following the 0.2 µg/day FAC implant is positive for this patient and highlights the importance of early intervention when a patient is responding insufficiently, or not responding, to anti-VEGF therapy. The current outcomes also suggest that to achieve functional benefits from DMO resolution (e.g. BCVA of 0.5 to permit holding a standard driving licence), earlier treatment should be considered.

This case illustrates how 0.2 µg/day FAC implant, with its long-term, low-dose corticosteroid, can provide effective long-term therapy (up to 36 months with one injection) with manageable side effects and an improvement in quality of life and general daily living. It is rewarding to note that our patient reported greater vision-related quality of life and the ability to perform daily routines without problems following 0.2 µg/day FAC implant. In patients with chronic DMO, the relative functional benefits and manageable side-effect profile of 0.2 mg/day FAC implant support its use in patients that are insufficiently responsive to available therapies.

To summarise, 0.2 µg/day FAC implant could have been considered earlier in the treatment of DMO and once the patient had shown limited or no response to anti-VEGF therapy. The current case shows CFT improved following treatment with dexamethasone. If ILUVIEN had been licensed at the time, it may have been considered at this stage to help stabilise CFT and BCVA.