Vitreous cytokine levels following the administration of a single 0.19 mg fluocinolone acetonide (ILUVIEN®) implant in patients with refractory diabetic macular edema (DME)—results from the ILUVIT study

Svenja K. Deuchler1 · Ralf Schubert3 · Pankaj Singh2 · Adonis Chedid2 · Ninel Kenikstul2 · Julia Scott1 · Thomas Kohnen2 · Hanns Ackermann4 · Frank Koch1,2

Received: 7 June 2021 / Revised: 29 October 2021 / Accepted: 11 January 2022 / Published online: 3 March 2022 © The Author(s) 2022

Abstract

Purpose To investigate the changes in vitreous inflammatory and angiogenic cytokine levels, primarily interleukin-(IL)-6, following intravitreal injection of the 0.19 mg fluocinolone acetonide (FAc, ILUVIEN®) implant in patients with diabetic macular edema.

Methods A single-center phase IV study involving 12 patients’ eyes with diabetic macular edema. Vitreous fluid samples were obtained prior to intravitreal injection of the fluocinolone acetonide implant and then again over a 6-month period. Vitreous samples were examined using a cytometric bead array to measure IL-6, IL-8, IP-10, MCP-1, VEGF, and CD54. PIGF and PEDF were measured using an enzyme-linked immunosorbent assay. Changes in the cytokine and chemokine expression patterns were analyzed. Clinical parameters such as BCVA and center point thickness (CPT) were also examined.

Results There were mean reductions in all parameters between baseline and month 6. Significant changes (p < 0.05 versus baseline) were observed in the expression of IL-6, IP-10, MCP-1, and CD54 following the administration of fluocinolone acetonide implant. VEGF and PIGF increased at month 1 before declining at month 6, though this trend was not significant. CPT decreased rapidly between screening and the first follow-up visit, and this decrease was sustained. BCVA remained relatively stable throughout.

Conclusion This study demonstrated changes in vitreous inflammatory and angiogenic cytokine levels following intravitreal injection of the FAc implant in patients with diabetic macular edema. Data show that the fluocinolone acetonide implant led to rapid and sustained reductions of some inflammatory cytokines with improvement of the overall clinical picture.

Keywords Cytokines · Diabetic macular edema · Diabetic retinopathy · Fluocinolone acetonide · Growth factors · Inflammatory markers
Key messages

- Diabetic macular edema is the leading cause of vision loss in diabetic patients. Options for treatment include intravitreal anti-vascular endothelial growth factor (VEGF) agents (ranibizumab, aflibercept and off-label injections of bevacizumab) and intravitreal corticosteroids (0.2 µg/day fluocinolone acetonide [FAc] implant, dexamethasone implant and off-label injections of triamcinolone acetonide).

- IL-6, IL-8, IP-10, MCP-1, VEGF, CD54, PIGF, and PEDF were measured in the vitreous of patients suitable for an FAc implant for treatment of chronic diabetic macular edema.

- Treatment with the FAc implant led to significant changes in inflammatory and angiogenic vitreous cytokine levels in IL-6, IP-10, MCP-1 and CD54.

- The use of the FAc implant for treatment of diabetic macular edema also improves clinical parameters, as evidenced by the reduction of macular edema on OCT.

Introduction

Diabetic macular edema (DME) is a complication of diabetic retinopathy (DR) and is the leading cause of vision loss in diabetics. It is estimated that between 14 and 25% of patients with diabetes develop DME within 10 years of being diagnosed and that the risk of development is influenced by both the type of diabetes and its duration [1].

Current treatment options for DME include intravitreal anti-vascular endothelial growth factor (VEGF) agents (ranibizumab, aflibercept and off-label injections of bevacizumab) and intravitreal corticosteroids (0.2 µg/day fluocinolone acetonide [FAc] implant, dexamethasone implant, and off-label injections of triamcinolone acetonide).

The FAc ILUVIEN® implant was designed to provide continuous treatment with fluocinolone acetonide for up to 3 years [2] and is delivered via an intravitreal injection for the treatment of DME that persists or recurs despite therapy. The outcomes from both clinical trials [3] and real-world practices [4, 5] have consistently shown that therapy with the FAc implant leads to rapid and sustained improvements of visual acuity and center point thickness (CPT).

Corticosteroids are multi-factorial in nature and target VEGF and several inflammatory cytokines relating to the development of DR and DME [6]. However, the actual effect of FAc on particular inflammatory cytokines still remains unclear due to limited research on humans in this area [7] and extrapolation of findings from other agents such as triamcinolone acetonide [8]. Previous investigations have shown that the pharmacokinetics of the FAc implant are unique compared to other commonly injected intravitreal corticosteroids as it delivers a lower daily dose of corticosteroid over a 3-year period [2], meaning that other preparations have a limited durability of release compared to the FAc implant [9].

Several cytokines have been documented to play a role in disease pathogenesis. VEGF is considered central to DME due to its pro-inflammatory, pro-angiogenic, and pro-vascular permeability effects [10]. The importance of VEGF in the development of DME is also reflected by the array of anti-VEGF therapies available and their use as first-line therapies. DME trials testing the effect of anti-VEGF agents have shown that around 40% of patients do not respond sufficiently to anti-VEGF agents, which suggests that other mediators play a role in the pathogenesis of DME [11]. In such cases, therapies that target inflammatory cytokines for example intercellular adhesion molecule-1 (CD54) and interleukin (IL)-6, may be a more effective therapeutic strategy [12]. The benefits of this approach seem to be reflected in clinical trials where it has been shown that corticosteroid agents, such as the FAc implant, are effective when used as a second-line therapy [4, 5].

A number of studies have investigated the effect of anti-VEGFs (ranibizumab [13, 14], aflibercept [15] and bevacizumab [8, 16]) on intraocular cytokine concentrations. There are no studies, however, that have investigated cytokine concentrations after the intravitreal injection of the FAc implant. The objective of this study was to determine its action on the expression of inflammatory and angiogenic cytokines, focusing on the change in factor IL-6 which plays a key role in the development of DME [6, 17–20] prior to and 6 months after the administration of the FAc implant in patients with DME that persisted or recurred despite treatment.
Methods

Ethics statement and patient recruitment

This pilot study was approved by the Bundesinstitut für Arzneimittel und Medizinprodukte (BfARM EudraCT-Number, 2016–004,488-38) and the ethics committee of the medical department of the Goethe University Frankfurt (E145/17). It was conducted at the Goethe University Hospital Frankfurt in Germany. The study was performed in accordance with the ethical standards as laid down in the 1964 Declaration of Helsinki and its later amendments. The study was designed as a phase IV trial involving 12 patients with DME that persisted or recurred despite treatment and therefore were eligible for treatment with the FAc implant. This was based on clinical judgment and decided prior to study initiation. Written informed consent was obtained from patients who wished to participate. Patient recruitment was from September 2018 to February 2019.

Data collection

Initial data collection and screening took place 0–4 weeks prior to start of treatment with FAc implant. Patients were excluded if they (1) had a prior vitrectomy (3-port pars plana vitrectomy); (2) had a prior partial vitrectomy with drug administration; (3) had been treated with an anti-angiogenic agent; (4) had been treated with laser coagulation during the 90 days prior to screening; (5) had been treated with a long-acting corticosteroid during the 90 days prior to screening; (6) had undergone cataract extraction during the 90 days prior to screening; (7) had secondary cataract treatment in the 90 days prior to screening; (8) presented with other eye diseases at the time of screening; (9) had uncontrolled intraocular pressure (≥30 mmHg); (10) had uncontrolled hypertension (>160/90 mmHg); (11) had poorly controlled diabetes (HbA1c >10%); and (12) if they were previously diagnosed with an autoimmune disease.

All study participants had an indication for and planned implantation of FAc. Additional inclusion criteria included age ≥ 18 years old and a diagnosis of either type 1 or 2 diabetes mellitus with DR and DME. A best corrected visual acuity (BCVA) between 19 and 78 letters using the Early Treatment Diabetic Retinopathy Study (ETDRS) scoring system was required, as well as a CPT ≥ 250.

Diabetic disease status (date of first diagnosis, diabetes type, current medication, last HbA1c value) were collected at T-1. A blood sample (5–10 ml blood) was also taken to determine blood sugar and HbA1c levels. Other parameters that were assessed included disease and treatment history (which included the diagnoses of DR and DME), prior treatments (laser, intravitreal injections, vitrectomies) and relevant concomitant diseases. Optical coherence tomography (OCT) (Topcon 3D-OCT-2000) and fluorescein angiography (FA) (Zeiss FF 450+) were also obtained at screening and were used to confirm presence of DME.

Surgical procedure

A minimally invasive partial pars plana vitrectomy (cPPV) was performed at baseline (FAc implantation) to extract vitreous from the eye for lab analysis. A minimum volume of 0.5 ml of pure vitreous was aspirated before the infusion was activated and the pressure was regulated towards normal interoperative levels (see video, supporting information 1, which shows a limited 2-port pars plana vitrectomy with vitreous probe). This procedure has been reported previously [21] and was associated with minimal complications (0.44%) which were comparable to those following the administration of intravitreal medications. Vitreous fluid samples were obtained prior to intravitreal injection of the FAc implant and repeated at 1 month and 6 months after insertion.

Primary and secondary parameters

Extracted vitreous samples were examined using a Cytometric Bead Array (CBA) and Flex sets [22–24] to determine IL-6, IL-8, IP-10 (pro-inflammatory cytokines), MCP-1 (pro-inflammatory chemokine), VEGF (growth factor), and CD54 concentrations (intercellular adhesion molecule). Placental growth factor (PIGF) and platelet-derived growth factor (PEDF) were measured using an enzyme-linked immunosorbent assay (ELISA; R&D Systems). The primary parameter in this study was the change in IL-6 and other cytokines were deemed secondary parameters.

Other (tertiary) parameters measured prior to and following intravitreal injection of the FAc implant included visual acuity (VA; measured using an ETDRS letter score), contrast VA (measured using a Central Vision Analyzer), CPT and average retinal thickness (ART) (both measured using OCT). Biomicroscopy was used to assess the lens and fundus state. FA was utilized to assess DR and color fundus photography was used to classify DR according to ETDRS. Finally, safety was determined from measurements of intraocular pressure (IOP) which was recorded during the day at both pre- and post-operative (i.e. post-PPV) time points. Post-operative measurements were taken 30 to 60 min following PPV.

Statistical analysis

The Friedmann test was used to check whether there were differences in the position parameters between the samples. In some cases, a Student’s t-test was used to compare mean values between individual time points. A p-value < 0.05
was taken as representing a statistically significant difference. Variables were recorded at screening, baseline: the time the FAc implant was administered (first vitreous probe), 1 week after the implant: this time point was used for the recording of some of the study variables, 1 month after the FAc implant, second vitreous probe: 3 months after FAc implant, and 6 months after the FAc implant: third vitreous probe. These time points are presented throughout. Values are reported as mean ± standard deviation throughout unless otherwise stated.

Results

Twelve patients underwent cPPV after screening. Thirteen patients were screened initially and assessed to be eligible to participate, but 1 declined to participate. Table 1 reports patient demographics and treatment history at screening for the 12 patients enrolled. The mean age of treated patients was 60.9 (range, 43–78) years. Eleven of the patients had type II diabetes and 3 of 12 study eyes were already pseudophakic at screening (Table 1). Prior therapies included laser with 4 patient eyes receiving panretinal laser coagulation, 2 receiving focal laser coagulation, and one receiving both focal and panretinal laser coagulation. Four patient eyes received panretinal cryocoagulation (Table 1). Intravitreal injections were given in 8 patient eyes (an average of 9.8 injections) with 6 eyes receiving aflibercept, 4 eyes receiving ranibizumab (7.0 injections) and 2 eyes receiving a dexamethasone implant (1.5 injections). Table 2 contains individual patient demographics and prior treatment details.

Volume of vitreous probes

In \( n = 23 \) cases, 1.2 ml volume could be extracted from the posterior compartment of the vitreous. In \( n = 13 \) cases, the minimum volume (0.5 ml) was reached with extraction volumes of 0.5 to 0.615 ml at baseline \( [n = 5] \), month 1 \( [n = 5] \), and month 6 \( [n = 3] \).

Changes in IL-6 (primary objective)

One month after administration of the FAc implant, IL-6 decreased from \( 84.30 \pm 126.81 \) pg/ml to \( 49.35 \pm 57.07 \) pg/ml at month 1 and \( 46.93 \pm 79.63 \) pg/ml at month 6 (see Table 3 and Fig. 1) \( (p = 0.0498) \).

Changes in IL-8, IP-10, MCP-1, VEGF, CD54, PIGF, and PEDF (secondary objectives)

Following administration of the FAc implant there were significant overall changes in mean MCP-1 \( (p = 0.0005) \), CD54 \( (p = 0.0388) \), PIGF \( (p = 0.0092) \), and IP-10 \( (p = 0.0131) \) between baseline and month 6. In contrast there were no significant \( (p > 0.05) \) changes in IL-8, VEGF, or PEDF (Table 3).

Visual acuity and contrast visual acuity outcomes (tertiary objectives)

At screening, mean visual acuity was \( 67.0 \pm 9.3 \) ETDRS letters and remained stable at month 6 \( (66.4 \pm 10.2 \) letters; \( P > 0.05 \), Student’s paired \( t \)-test). Contrast visual acuity also remained relatively stable with mean values of \( 29.5 \pm 3.6 \) letters and \( 27.1 \pm 3.3 \) letters \( (P > 0.05 \), Student’s paired \( t \)-test) at screening and at month 6. Table 4 shows individual patient results for visual acuity and contrast visual acuity.
OCT outcomes

The CPT measured during the study is displayed in Fig. 2 and shows a rapid decline by the first follow-up visit (1 week), versus the value recorded at screening (0–4 weeks prior to the administration of the FAc implant), and that this was sustained throughout ($p < 0.05$ for all points versus screening). Table 4 includes individual patient CPT and ART results.

Intraocular pressure outcomes

Intraocular pressure was measured during the study and mean values are reported in Fig. 3. The lowest mean intraocular pressure (9.1 mmHg) was measured postoperatively at month 1 and the highest mean value (17.8 mmHg) was obtained preoperatively at month 6. There was no significant intraocular pressure change when mean preoperative or non-operative (no probe taken) measurements were compared with mean screening measurements, with the exception of 1 week after implant ($p < 0.05$). For individual results, see Table 5.

Safety

Four events (16.7%) in 3 patients (25.0%) recorded during the study were related to the FAc implant. These were iris adhesions, increased intraocular pressure, a medical device problem (implant was stuck in the needle during application) and a surgery-related problem (extrusion of implant through trocar during vitreous probe insertion). However, none of the adverse events that occurred were serious and all could be resolved without sequelae.

Discussion

The majority of studies measuring cytokine levels in the eye have sampled fluid from the aqueous humor rather than the vitreous. The reasons for this relate to the fact that obtaining vitreous samples may lead to higher complications compared to samples obtained from the aqueous humor, which are relatively easier and safer to perform. Another reason is that cytokine levels in the aqueous humor have been correlated with levels in the vitreous fluid [25], at least in the case of VEGF and IL-6. However, the same correlation has not been proven for other cytokines and the study by Funatsu et al. [25] did not show a perfect correlation in cytokine levels taken from the 2 different sampling sites in the eye. With this in mind, the current study was designed to assess inflammatory and angiogenic cytokine expression in the vitreous fluid during a partial vitrectomy and in doing so had the
advantage of measuring the activity at the site of delivery and sites of action in the posterior segment of the eye.

The current study evaluated the influence of the FAc implant therapy on levels of IL-6 as well as different vitreous inflammatory and angiogenic cytokine levels (including IL-8, IP-10, MCP-1, VEGF, CD54, PIGF and PEDF) in patients with DME. IL-6, as well as MCP-1, VEGF, and CD54, have been reported to be significantly higher in patients with DME than in nondiabetic patients [20]. These findings demonstrate the importance of inflammatory cytokines, and not solely the VEGF cytokine, in the breakdown of the blood retinal barrier and the pathogenesis of DME.

It is documented that patients with diabetes, in contrast to non-diabetics, have elevated levels of aqueous and vitreous IL-6 [26]. In the present study, an average concentration of 84.30 ± 126.81 pg/ml was measured in the vitreous at baseline. At 1 and 6 months after the FAc implant, it decreased significantly to 49.3 ± 57.1 pg/ml and 46.9 ± 79.6 pg/ml, respectively. The result of this reduction of inflammatory parameters in the vitreous cavity is consistent with previous work showing inflammatory molecules such as IL-6 in the aqueous humor can be significantly reduced by intravitreal corticosteroid therapy [26]. Ecker et al. [27] failed to show a correlation between inflammatory parameters in the anterior chamber and the vitreous chamber and others [28] have reported that only a small minority of the proteins in both chambers can be correlated. These findings suggest that vitreous samples at the site of disease manifestation may be

| Cytokine | Baseline       | 1 month after implant | 6 months after implant | 6-months minus baseline | P-value* |
|---------|----------------|-----------------------|------------------------|-------------------------|----------|
| IL-6    | 84.3 ± 126.8   | 49.3 ± 57.1           | 46.9 ± 79.6            | -37.4 ± 83.8            | 0.0498   |
| IL-8    | 39.9 ± 35.3    | 37.1 ± 26.7           | 33.8 ± 25.0            | -6.1 ± 25.7             | 0.3679   |
| IP-10   | 134.8 ± 241.7  | 76.5 ± 125.2          | 64.1 ± 88.9            | -70.7 ± 155.3           | 0.0131   |
| CD54    | 839. ± 793.8   | 532.3 ± 354.7         | 513.8 ± 343.1          | -325.6 ± 610.0          | 0.0388   |
| MCP-1   | 866.5 ± 401.0  | 545.6 ± 235.6         | 541.9 ± 241.0          | -324.6 ± 305.5          | 0.0005   |
| VEGF    | 280.3 ± 351.5  | 286.4 ± 337.9         | 179.6 ± 164.0          | -100.7 ± 313.0          | 0.6136   |
| PIGF    | 95.8 ± 108.4   | 128.5 ± 157.7         | 75.8 ± 63.2            | -20.1 ± 67.7            | 0.0092   |
| PEDF    | 4696.2 ± 315.8 | 4743.0 ± 270.4        | 4711.5 ± 266.6         | 15.3 ± 439.6            | 0.3385   |

Notes: Data presented as mean ± SD. Concentrations reported in pg/ml. * Friedmann test

Fig. 1 Vitreous inflammatory and angiogenic cytokine levels during therapy with FAc. Notes: Individual data plots are presented with mean values shown as a solid line. P-values reported where P < 0.05 and were calculated using a Friedmann test.
Table 4  Individual patient OCT and visual acuity results

| Patient | Center Point Thickness (µm) | Average Retinal Thickness (µm) | Best Corrected Visual Acuity (letters) | Best Corrected Contrast Visual Acuity (letters) |
|---------|----------------------------|-------------------------------|----------------------------------------|-----------------------------------------------|
|         | screening | 1 week after implant | 1 month after implant | 3 months after implant | 6 months after implant | screening | 1 week after implant | 1 month after implant | 3 months after implant | 6 months after implant | screening | 1 week after implant | 1 month after implant | 3 months after implant | 6 months after implant |
| ILV1-01 | 368   | 285   | 253   | Not measured | 223 | 391 | 373 | 338.3 | Not measured | 292.7 | 77  | 77  | 83  | 76  | Not measured | 79 | 34  | 28  | 30  |
| ILV1-02 | 467   | 441   | 441   | 446 | 330 | 362.9 | 349 | 337.5 | 344.3 | 304.8 | 73  | 73  | 78  | 79  | 75  | 73  | 30  | 25  | 22  |
| ILV1-03 | 351   | 344   | 325   | 323 | 309 | 329 | 326.3 | 312.8 | 300.9 | 290.1 | 75  | 75  | 73  | 78  | 78  | 57  | 31  | 31  | 32  |
| ILV1-04 | 383   | 348   | 338   | 338 | 354 | 294.5 | 294.3 | 290.6 | 284.2 | 280.5 | 61  | 62  | 64  | 66  | 70  | 67  | 29  | 29  | 29  |
| ILV1-05 | 566   | 422   | 399   | 365 | 413 | 400.6 | 383.5 | 369.7 | 343.6 | 363.1 | 71  | 71  | 74  | 69  | 71  | 69  | 34  | 34  | 30  |
| ILV1-06 | 236   | 237   | 241   | 169 | 275 | 365.2 | 363.1 | 339.9 | 347.8 | 370.2 | 47  | 47  | 51  | 47  | 48  | 43  | 35  | 31  | 29  |
| ILV1-07 | 266   | 259   | 249   | 234 | 215 | 274.9 | 271.2 | 275.4 | 272.3 | 266.4 | 66  | 69  | 69  | 68  | 70  | 72  | 28  | 29  | 23  |
| ILV1-08 | 317   | 209   | 174   | 209 | 191 | 383.2 | 338.5 | 322.5 | 325.3 | 315.2 | 64  | 64  | 65  | 64  | 68  | 68  | 25  | 25  | 28  |
| ILV1-09 | 492   | 282   | 301   | 452 | 487 | 404.4 | 347.1 | 348.4 | 384.8 | 431.1 | 77  | 76  | 78  | 84  | 80  | 76  | 24  | 25  | 26  |
| ILV1-10 | 564   | 314   | 381   | 318 | 271 | 444.6 | 348 | 350.1 | 335.4 | 316.8 | 74  | 74  | 70  | 69  | 72  | 69  | 26  | 28  | 27  |
| ILV1-11 | 335   | 273   | 239   | 241 | 241 | 298.1 | 282 | 270.7 | 270.2 | 267.6 | 56  | 56  | 64  | 54  | 53  | 54  | 28  | 27  | 27  |
| ILV1-12 | 599   | 526   | 506   | 508 | 576 | 425.6 | 369.6 | 376.6 | 376.6 | 401.7 | 63  | 63  | 67  | 73  | 73  | 70  | 30  | 30  | 22  |
| Mean    | 413.5 | 328.3 | 320.6 | 327.5 | 323.8 | 364.5 | 337.13 | 327.7 | 325.9 | 325.0 | 67  | 67.3 | 69.7 | 68.9 | 63.2 | 66.4 | 29.5 | 28.5 | 27.1 |

Notes: Patient ILV-09 declined to participate in the study
necessary to reliably demonstrate a response to intravitreal therapy [27–29].

Song et al. [30] postulated that elevated levels of CD54 play a central role in the progression of DR and the severity of disease. Likewise, Funatsu et al. [20] demonstrated a correlation between DME severity and CD54 levels. In this study, there was a statistically significant decrease in CD54 after FAc therapy.

VEGF levels have also been reported to be elevated in aqueous and vitreous humor in patients with diabetes as compared to non-diabetic patients [26], although there are mixed findings concerning the relationship between

---

**Fig. 2** Mean center point thickness recorded at baseline and at time points between 1 week and 6 months after the administration of the FAc implant.

**Fig. 3** Mean intraocular pressure recorded prior to (screening) and after (1 week to 6 months) the administration of the FAc implant. Notes: Preoperatively refers to the measurement of pressure prior to and postoperatively refers to the measurement of pressure after 2 port pars plana vitrectomy for vitreous probes.
VEGF and DR or DME severity [20, 26, 30]. At the beginning of therapy with the FAc implant, this study showed mean VEGF concentrations of 280.31±351.49 pg/ml. After 1 month, the concentration increased slightly to 286.42±337.93 pg/ml and by month 6 had decreased (P>0.05) by 100.71±312.95 pg/ml to a level of 179.61±163.94 pg/ml. The overall tendency was for VEGF to decrease, but this was not statistically significant as VEGF levels increased in 4 eyes (range, 37.0 to 206.3 pg/ml) and remained unchanged in 1 eye.

The previous work of Song et al. [30] and Dong et al. [26] showed that the amount of MCP-1 in aqueous humor or vitreous humor also correlates with the severity of DR or DME. At the beginning of the present study the MCP-1 concentration was 866.46±400.97 pg/ml. At month 1, it had decreased to 545.63±235.58 pg/ml and at month 6 to 541.85±241.04 pg/ml. The decrease in MCP-1 values was statistically significant (p=0.0005) and indicates an effect of the FAc implant on the expression of MCP-1.

IL-8 levels are also associated with the severity of DR [30]. At baseline, the average IL-8 concentration was 39.90±35.33 pg/ml. Slight decreases but non-significant changes were observed at month 1, (to 37.09±26.74 pg/ml) and month 6 (to 33.80±24.97 pg/ml). Therefore, an effect of FAc implant therapy on IL-8 concentration could not be shown.

The mean PIGF concentration before treatment with the FAc implant was 95.86±108.35 pg/ml. At month 1 it increased to 128.49±157.67 pg/ml and at month 6 decreased to 75.79±63.17 pg/ml. Overall, a statistically significant decrease in PIGF concentration of 20.07±67.71 was observed by month 6.

The IP-10 concentration was 134.81±241.71 pg/ml on the day of FAc implantation. A statistically significant decrease in IP-10 concentration was observed during the course of the study. One month after transplantation, the IP-10 concentration was 76.50±125.21 pg/ml. A further decrease (to 64.13±88.91 pg/ml) was observed by month 6. Overall, there was a decrease of 70.68±155.34 pg/ml by month 6.

At baseline the average PEDF concentration was 4696.15±315.84 pg/ml. During the course of the study, the concentration increased slightly at month 1 (4742.98±270.44 pg/ml). At month 6, the mean concentration was 4711.48±266.64 pg/ml, which was similar to the baseline value and only changed by 15.33±439.60 pg/ml (p>0.05).

Regarding clinical results, the FAc implant is a sustained delivery device but still, the decrease in center point thickness occurred mostly in the first month after implantation of the FAc implant. Over the 6-month follow-up period, this decrease was sustained. This is adventitious to patients as they do not have to receive ongoing injections to obtain a similar result [5]. Only 1 patient had a mild steroid response resulting in increased (up to 25 mmHg) intraocular pressure following device implantation. This was resolved with topical antiglaucoma therapy. As seen in Table 5, due to the vitreous probe, there was an induced preliminary hypotony in 7 of 36 postoperative measurements but without clinical complications such as corneal changes or choroidal folds. This preliminary IOP fluctuation might have an impact on the cytokine concentrations due to the potential blood retinal barrier and blood aqueous barrier breakdown [31]. This should be considered as a limitation of the study.

Table 5 Individual patient intraocular pressure levels (mmHg)

| Patient | Screening (preoperatively) | Baseline (postoperatively) | 1 week after implant | 1 month after implant (preoperatively) | 1 month after implant (postoperatively) | 3 months after implant | 6 months after implant (preoperatively) | 6 months after implant (postoperatively) |
|---------|-----------------------------|-----------------------------|---------------------|--------------------------------------|---------------------------------------|-----------------------|----------------------------------------|------------------------------------------|
| ILV1-01 | 15                          | 18                          | 4                   | 10                                   | 18                                    | 12                    | 13                                     | 14                                       |
| ILV1-02 | 17                          | 18                          | 14                  | 19                                   | 12                                    | 6                     | 21                                     | 22                                       |
| ILV1-03 | 15                          | 14                          | 9                   | 14                                   | 14                                    | 4                     | 16                                     | 18                                       |
| ILV1-04 | 16                          | 9                           | 10                  | 8                                    | 9                                     | 11                    | 8                                      | 10                                       |
| ILV1-05 | 17                          | 15                          | 15                  | 10                                   | 14                                    | 12                    | 15                                     | 14                                       |
| ILV1-06 | 20                          | 23                          | 17                  | 16                                   | 16                                    | 10                    | 19                                     | 22                                       |
| ILV1-07 | 18                          | 16                          | 14                  | 14                                   | 13                                    | 9                     | 21                                     | 14                                       |
| ILV1-08 | 13                          | 13                          | 10                  | 12                                   | 13                                    | 9                     | 23                                     | 18                                       |
| ILV1-09 | 16                          | 20                          | 16                  | 18                                   | 13                                    | 10                    | 21                                     | 18                                       |
| ILV1-11 | 16                          | 15                          | 8                   | 22                                   | 20                                    | 7                     | 23                                     | 19                                       |
| ILV1-12 | 20                          | 13                          | 10                  | 17                                   | 22                                    | 14                    | 18                                     | 25                                       |
| ILV1-13 | 18                          | 15                          | 9                   | 12                                   | 13                                    | 5                     | 10                                     | 19                                       |
| Mean    | 17.1                        | 15.6                        | 11.3                | 14.3                                 | 14.8                                  | 9.1                   | 17.3                                   | 17.8                                     | 11.2
An additional limitation of the current study is that it included a cohort of 12 patients from a single clinical center. Hence, it needs to be confirmed in a larger cohort of patients across multiple centers. Furthermore, the FAc implant has been designed to provide sustained drug delivery for up to 3 years and some parameters (e.g., PIGF and PEDF) only started to decrease by month 6 or decreased slowly over the study period. From clinical data [2] we know that the FAc implant has an effect for up to 3 years but to our knowledge, cytokine levels have not been evaluated past 6 months post implantation up to this point. A single case reported by Singh et al. [32], showed that edema recurred approximately 3 years after implantation and it is speculated that this was driven by the underlying inflammatory cascade. Variables such as HbA1c which reflect the lifestyle of the patient might also play a role in cytokine changes. A study with a longer study period (i.e., 12 months or longer) with an additional vitreous probe would be valuable. Regarding statistical analysis, the study was designed to investigate whether the intra-vitreal FAc implant has an influence on IL-6 at baseline and month 1 and 6 (primary parameter). The other cytokines, evaluated at the same time points, were designated as secondary parameters. A further study could consider performing multiple comparisons over time if multiple cytokines are evaluated from the same vitreous samples.

This study reports changes in vitreous inflammatory and angiogenic cytokine levels following intravitreal injection of the FAc implant in patients with DME. Data show that the FAc implant altered the expression of vitreous inflammatory and angiogenic cytokine levels, which were maintained 6 months after administration of a single FAc implant. These effects were also accompanied by improvements in visual acuity and center point thickness.

Supplementary Information The online version contains supplementary material available at https://doi.org/10.1007/s00417-022-05564-2.

Acknowledgements This Investigator Initiated Trial was supported by Alimera Sciences [grant number 590/15]. The abstract of this paper was presented at Euretina 2020.

Funding This Investigator Initiated Trial was supported by Alimera Sciences [grant number 590/15].

Data availability Data is available on request from the corresponding author.

Declarations

Ethics approval This study was performed in line with the principles of the Declaration of Helsinki. Approval was granted by the Bundesinstitut für Arzneimittel und Medizinprodukte (BfARM EudraCT-Number 2016–004488-38) and by the ethics committee of the medical department of the Goethe University Frankfurt (E145/17).

Consent to participate. Informed consent was obtained from all individual participants included in the study.

Consent for publication. The authors affirm that the featured human research participant provided informed consent for publication of the supplementary information video.

Conflict of interest Frank Koch and Pankaj Singh are working as consultants for Alimera Sciences. All other authors have no financial interests to declare.

Open Access This article is licensed under a Creative Commons Attribution 4.0 International License, which permits use, sharing, adaptation, distribution and reproduction in any medium or format, as long as you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons licence, and indicate if changes were made. The images or other third party material in this article are included in the article’s Creative Commons licence, unless indicated otherwise in a credit line to the material. If material is not included in the article’s Creative Commons licence and your intended use is not permitted by statutory regulation or exceeds the permitted use, you will need to obtain permission directly from the copyright holder. To view a copy of this licence, visit http://creativecommons.org/licenses/by/4.0/.

References

1. Klein R, Klein BEK, Moss SE, Cruickshanks KJ (1995) The Wisconsin Epidemiologic Study of Diabetic Retinopathy XV: The Long-term Incidence of Macular Edema. Ophthalmology 102:7–16. https://doi.org/10.1016/s0161-6420(95)31052-4
2. Campochiaro PA, Nguyen QD, Hafiz G et al (2013) Aqueous levels of fluocinolone acetonide after administration of fluocinolone acetonide inserts or fluocinolone acetonide implants. Ophthalmology 120:583–587. https://doi.org/10.1016/j.ophtha.2012.09.014
3. Campochiaro PA, Brown DM, Pearson A et al (2012) Sustained delivery fluocinolone acetonide vitreous inserts provide benefit for at least 3 years in patients with diabetic macular edema. Ophthalmology 119:2125–2132. https://doi.org/10.1016/j.ophtha.2012.04.030
4. Panos GD, Arruti N, Patra S (2020) The long-term efficacy and safety of fluocinolone acetonide intravitreal implant 190 μg (ILUVIEN®) in diabetic macular oedema in a multi-ethnic inner-city population. Eur J Ophthalmol 31:620–629. https://doi.org/10.1177/1120672119898414
5. Augustin AJ, Bopp S, Fechner M et al (2020) Three-year results from the Retro-IDEAL study: Real-world data from diabetic macular edema (DME) patients treated with ILUVIEN (0.19 mg fluocinolone acetonide implant). Eur J Ophthalmol 30(2):382–391. https://doi.org/10.1177/1120672119834474
6. Rubsam A, Parikh S, Port P (2018) Role of Inflammation in Diabetic Retinopathy. Int J Mol Sci 19:942. https://doi.org/10.3390/ijms19040942
7. Kane FE, Green KE (2015) Ocular pharmacokinetics of fluocinolone acetonide following Iluvien implantation in the vitreous humor of rabbits. J Ocul Pharmacol Ther 31:11–16. https://doi.org/10.1080/joph.2014.0100
8. Sohn HJ, Han DH, Kim IT et al (2011) Changes in aqueous concentrations of various cytokines after intravitreal triamcinolone versus bevacizumab for diabetic macular edema. Am J Ophthalmol 152:686–694. https://doi.org/10.1016/j.ajo.2011.03.033
9. Eichenbaum D (2017) Fluocinolone acetonide (FAc): Pharmacokinetics and clinical relevance. Invest Ophthalmol Vis Sci 58:5791
10. Hillier RJ, Ojaimi E, Wong DT et al (2017) AQUEOUS HUMOR CYTOKINE LEVELS AS BIOMARKERS OF DISEASE SEVERITY IN DIABETIC MACULAR EDEMA. Retina 37:761–769. https://doi.org/10.1097/IAE.000000000001210
11. Gonzalez VH, Campbell J, Holekamp NM et al (2016) Early and Long-Term Responses to Anti-Vascular Endothelial Growth Factor Therapy in Diabetic Macular Edema: Analysis of Protocol I Data. Am J Ophthalmol 172:72–79
12. Whitcup SM, Cidlowski JA, Csaky KG, Ambati J (2018) Pharmacology of Corticosteroids for Diabetic Macular Edema. Invest Ophthalmol Vis Sci 59:1. https://doi.org/10.1016/j.ajo.2016.09.012
13. Hillier RJ, Ojaimi E, Wong DT et al (2018) Aqueous Humor Cytokine Levels and Anatomic Response to Intravitreal Ranibizumab in Diabetic Macular Edema. JAMA Ophthalmol 136:381. https://doi.org/10.1001/jamaophthalmology.2018.0179
14. Shiraya T, Kato S, Araki F, Ueta T, Miyaji T, Yamaguchi T (2017) Aqueous cytokine levels are associated with reduced macular thickness after intravitreal ranibizumab for diabetic macular edema. PLoS ONE 12:e0174340. https://doi.org/10.1371/journal.pone.0174340
15. Mastropasqua R, D’Aloisio R, Di Nicola M et al (2018) Relationship between aqueous humor cytokine level changes and retinal vascular changes after intravitreal aflibercept for diabetic macular edema. Sci Rep 8:16548. https://doi.org/10.1038/s41598-018-35036-9
16. Yu S-Y, Nam DH, Lee DY (2018) Changes in aqueous concentrations of various cytokines after intravitreal bevacizumab and subtenon triamcinolone injection for diabetic macular edema. Graefes Arch Clin Exp Ophthalmol 256:39–47. https://doi.org/10.1007/s00417-017-3819-2
17. Kim M, Yonguk Y, Lee S (2015) Comparison of aqueous concentrations of angiogenic and inflammatory cytokines based on optical coherence tomography patterns of diabetic macular edema. Indian J Ophthalmol 63:312–317. https://doi.org/10.4103/0301-4738.158069
18. El-Ghrably I, Steel DHW, Habib M, Vaideanu-Collins D, Manvikar S, Hillier R (2017) Diabetic macular edema outcomes in eyes treated with fluocinolone acetonide 0.2 µg/d intravitreal implant: real-world UK experience. Eur J Ophthalmol 27:357–362. https://doi.org/10.5301/ejo.5000929
19. Jonas JB, Jonas RA, Neumaier M, Flineisen P (2012) Cytokine concentration in aqueous humor of eyes with diabetic macular edema. Retina 32:2150–2157. https://doi.org/10.1097/IAE.0b013e3182576d07
20. Funatsu H, Noma H, Mimura T, Eguchi S, Hori S (2009) Association of vitreous inflammatory factors with diabetic macular edema. Ophthalmology 116:73–79. https://doi.org/10.1016/j.ophth.2008.09.037
21. Koch FH, Koss MJ (2011) Microincision vitrectomy procedure using Intrector technology. Arch Ophthalmol 129:1599–1604. https://doi.org/10.1001/archophthalmol.2011.341
22. Eickmeier O, Huebner M, Herrmann E et al (2010) Sputum biomarker profiles in cystic fibrosis (CF) and chronic obstructive pulmonary disease (COPD) and association between pulmonary function. Cytokine 50:152–157. https://doi.org/10.1016/j.cytot.2010.02.004
23. Chen R, Lowe L, Wilson JD et al (1999) Simultaneous Quantification of Six Human Cytokins in a Single Sample Using Microparticle-based Flow Cytometric Technology. Clin Chem 45:1693–1694
24. Carson RT, Vignali DA (1999) Simultaneous quantitation of 15 cytokines using a multiplexed flow cytometric assay. J Immunol Methods 227:41–52. https://doi.org/10.1016/s0022-1759(99)00069-1
25. Funatsu H, Yamashita H, Noma H et al (2005) Aqueous humor levels of cytokines are related to vitreous levels and progression of diabetic retinopathy in diabetic patients. Graefes Arch Clin Exp Ophthalmol 243:3–8. https://doi.org/10.1007/s00417-004-0950-7
26. Dong N, Xu B, Wang BCL (2013) Study of 27 aqueous humor cytokines in patients with type 2 diabetes with or without retinopathy. Mol Vis 19:1734–1746
27. Ecker SM, Hines JC, Pihliah SM, Glaser BM (2011) Aqueous cytokine and growth factor levels do not reliably reflect those levels found in the vitreous. Mol Vis 17:2856–2863
28. Noma H, Funatsu H, Mimura T, Harino S, Hori S (2010) Aqueous humor levels of vasoactive molecules correlate with vitreous levels and macular edema in central retinal vein occlusion. Eur J Ophthalmol 20:402–409. https://doi.org/10.1177/112067211002000222
29. Funatsu H, Yamashita H, Noma H, Mimura T, Yamashita Hori S (2002) Increased levels of vascular endothelial growth factor and interleukin-6 in the aqueous humor of diabetics with macular edema. Am J Ophthalmol 133:70–77. https://doi.org/10.1016/s0002-9394(01)01269-7
30. Song S, Yu X, Zhang P, Dai H (2020) Increased levels of cytokines in the aqueous humor correlate with the severity of diabetic retinopathy. J Diab Comp 34:107641. https://doi.org/10.1016/j.jdiacomp.2020.107641
31. Schmack I, Völcker HE, Grossniklaus H (2010) Phthisis bulbi. In: Levin A, Albert D (eds) Ocular Disease: Mechanisms and Management, 1st edn. Saunders Elsevier, Amsterdam, p 421
32. Singh P, Chedid A, Deuchler SK, Kohnen T, Müller M, Koch FH (2018) The efficacy and safety outcomes of the 0.19 mg fluocinolone acetonide implant after prior treatment with the 0.7 mg dexamethasone implant in patients with diabetic macular edema. Int Med Case Rep J 11:265–269. https://doi.org/10.2147/IMCRJ.S174461

Publisher’s Note Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.