Visual acuity, endothelial cell density and polymegathism after iris-fixated lens implantation

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Purpose: The purpose of this study was to evaluate the visual acuity as well as endothelial cell density (ECD) and polymegathism after iris-fixated lens (Artiflex\textsuperscript{\textregistered} AC 401) implantation for correction of moderate to high myopia.

Patients and methods: In this retrospective cross-sectional study, 55 eyes from 29 patients undergoing iris-fixated lens implantation for correction of myopia (~5.00 to −15.00 D) from 2007 to 2014 were evaluated. Uncorrected visual acuity, best spectacle-corrected visual acuity, refraction, ECD and polymegathism (coefficient of variation [CV] in the sizes of endothelial cells) were measured preoperatively and 6 months postoperatively.

Results: In the sixth month of follow-up, the uncorrected vision acuity was 20/25 or better in 81.5\% of the eyes. The best-corrected visual acuity was 20/30 or better in 96.3\% of the eyes, and more than 92\% of the eyes had a refraction score of ±1 D from the target refraction. The mean corneal ECD of patients before surgery was 2,803±339 cells/mm\textsuperscript{2}, which changed to 2,744±369 cells/mm\textsuperscript{2} six months after surgery (p=0.142). CV in the sizes of endothelial cells before the surgery was 25.7\%±7.1\% and six months after surgery it was 25.9\%±5.4\% (p=0.857).

Conclusion: Artiflex iris-fixated lens implantation is a suitable and predictable method for correction of moderate to high myopia. There was no statistically significant change in ECD and polymegathism (CV in the sizes of endothelial cells) after 6 months of follow-up.

Keywords: cornea, endothelial cell, density, Artiflex, refractive surgery

Plain language summary
We studied the visual acuity as well as endothelial cell density (ECD) and polymegathism after Artiflex\textsuperscript{\textregistered} iris-fixated lens implantation among patients with moderate to high myopia. Six months postoperatively, there was no statistically significant change in corneal ECD (p=0.142) and also the coefficient of variation for endothelial cells’ size (p=0.857). Our results indicate that Artiflex iris-fixated lens implantation is a suitable and predictable method for correction of moderate to high myopia, causing no statistically significant change in ECD and polymegathism after 6 months of follow-up.

Introduction
A variety of refractive surgeries are used to correct refractive errors among those who do not desire to use glasses or contact lenses. Since corneal methods have some limitations in correcting high myopia, phakic intraocular lens (IOL) implantation has been used to correct moderate and high myopia since 1986.\textsuperscript{1} However, one of the perceived complications of phakic IOL is damage to corneal tissues, especially endothelial cells.\textsuperscript{2,3} Since endothelial cells in the human eye do not have the ability to reproduce,
any damage to these cells after infection, impact, surgical procedures and aging will affect their function.\textsuperscript{4,5} In various studies, it has been shown that endothelial cells naturally decrease with age.\textsuperscript{6,7} In addition, eye surgeries, especially intraocular surgeries, such as cataract surgery,\textsuperscript{9,10} posterior chamber phakic IOL implantation,\textsuperscript{10,11} or iris-fixated IOL (iris claw lens),\textsuperscript{12,13} can reduce the density of these cells. Preoperative endothelial cell density (ECD) examination provides a good predictor of corneal condition after phakic IOL surgery.\textsuperscript{14}

Many studies have indicated that iris-fixated IOL surgery is a predictable and effective method for correction of refractive errors.\textsuperscript{15,16} A study by Tehrani and Dick reported that ECD, 6 months after iris-fixated IOL implantation, had no significant difference with the preoperative readings.\textsuperscript{17} Pop and Payette\textsuperscript{18} reported that the reduction in endothelial cells was less than 1\% at 6, 12 and 24 months after the surgery, which was not statistically significant. In addition, Tahzib et al\textsuperscript{19} reported that ECD did not show a significant reduction compared to preoperative readings in 1 month and 6 months as well as 10 years after the Artisan phakic IOL implantation. However, contrary to these studies, in 1997, Menezo et al\textsuperscript{20} reported that the reduction in ECD was 5.8\% in the first 6 months after the surgery and 7.9\% after 12 months, which was statistically significant. In addition, in a study by Doors et al\textsuperscript{21} in 2008 with a mean follow-up of 34.1±24.7 months ECD was significantly reduced after iris-fixated IOL implantation, and in another study by Koss and Kohnen in 2009, a 10.5\% reduction in endothelial cells was observed 22 months after secondary iris-claw IOL surgery in aphakic individuals.\textsuperscript{22}

Therefore, regarding the different reports of changes in the count of endothelial cells after phakic IOL surgery, we decided to perform the present study to evaluate the changes in endothelial cells after Artiflex\textsuperscript{9} iris-fixated lens implantation.

**Patients and methods**

In this retrospective study, 55 eyes of 29 patients referred to Vanak Ophthalmology Clinic, Tehran, Iran, from 2007 to 2014 were evaluated. This study was approved by the ethics committee of Shahid Beheshti University of Medical Sciences and adhered to the rules established by the Declaration of Helsinki. All participants gave a written and informed consent before their data were used in this study. The subjects had a myopic score ranging from $-5.00$ to $-15.00$ D, and their myopic score was stable for 18 months prior to surgery (change of $<0.5$ D). In selecting the patients for Artiflex lens implantation, we followed the minimum requirements for ECD based on the recommendations by the manufacturer and patients’ age. All surgeries were performed by the same surgeon, and the same lens (Artiflex AC 401; Ophtee BV, Groningen, Netherlands, overall diameter: 8.5 mm/body diameter: 6.0 mm, convex–concave/hydrophobic polysiloxane foldable design) was used in all eyes. Patients were asked to stop using soft contact lens 1 week and hard contact lens 3 weeks before the operation. In all patients, the anterior segment was normal and had no pathological defects. Anterior chamber angle in the Orbscan (Bausch & Lomb, Rochester, NY, USA) assessment was $\geq 3.2$ mm, and patients did not have any systemic or ocular pathology other than their refractive error.

**Patient evaluation before and after the surgery**

In all subjects, basic examinations, including best spectacle-corrected visual acuity (BSCVA) and uncorrected visual acuity (UCVA), were performed using an auto chart projector (Nidek CP 670 [Nidek Co., Ltd.; Gamagori, Japan]). Manifest and cycloplegic refraction, the pupil size in scotopic light condition (pupil size in scotopic condition should be less than lens optic zone), pachymetry (DGH Technology, Inc., Exton, PA, USA), topography, keratometry, precision examinations using slit lamp, tonometer applanation, indirect ophthalmoscopy and examination of endothelial cells were also performed.

**Endothelial cells’ evaluation**

Corneal endothelial cells were studied in terms of cell density and cell size diversity (polymegathism) using specular microscopy (non-contact specular microscope: Topcon SP-3000P; Topcon Corp., Tokyo, Japan). All evaluations were performed by the same examiner, and three measurements in each eye were taken. Finally, the mean of these three measurements was used in statistical analysis. Evaluation of endothelial cells was performed before and 6 months after the surgical procedure. The van der Heijde’s formula was used to calculate the IOL power.\textsuperscript{23}

**Surgical method**

An iridectomy was performed 2 days before surgery by a neodymium: YAG laser in all patients. All patients underwent general anesthesia. The dosage of the anesthetic agent was proportional and varied depending on the patient’s condition, including weight and drug sensitivity. The duration of anesthesia was between 20 and 60 minutes. To place the Artiflex foldable lens, a 3.2 mm clear corneal incision at the 12 o’clock position was made and two stab incisions were also placed at 10 and 2 o’clock positions in the direction of the
enclavation sites. After injection of an ophthalmic cohesive viscosurgical substance (sodium hyaluronate 1%, Healon), the phakic intraocular lens was inserted into the anterior chamber using a specially designed implantation device for Artiflex lens (Operaid Artiflex Implantation Spatula; Ophtec BV). After positioning, the iris tissue was grasped and enclaved into the haptics at 3 and 9 o’clock with the aid of an Ophtec Artiflex enclavation needle (Ophtec BV). The viscoelastic agent was removed by manual irrigation.

Statistical methods
To present data, we used mean, SD, median and range and interquartile range. To assess the changes, we used paired t-test. All statistical analysis was performed using SPSS version 22.0 software (IBM Corporation, Armonk, NY, USA). p-values <0.05 were considered as statistically significant. To have a power of 90% to detect 5% change of in the cell density when the SD of the changes was assumed to be 300 cells, a minimum sample size of 49 eyes was calculated.

Results
In total, 55 eyes from 29 patients with a refractive error of −5 to −15 D were included in the study. The average age of patients was 24.9 years with a range of 20–39 years. In all, 20 patients (75.9%) were female and seven patients (24.1%) were male. Uncorrected visual acuity (UCVA), best-corrected visual acuity (BCVA), spherical equivalent, corneal ECD and coefficient of variation (CV) in the sizes of endothelial cells before the surgery are shown in Table 1.

The ECD before lens implantation was 2,809±339 cells/mm², which changed to 2,744±369 cell/mm² six months after surgery (p=0.142; Table 2). The CV in the sizes of endothelial cells before surgery was 25.7%±7.1%, which changed to 25.9%±5.4% six months after surgery, showing no statistically significant change (p=0.857; Table 2).

Discussion
In the present study, the visual acuity and changes in endothelial cells before and after Artiflex iris-fixated lens implantation were evaluated. The implantation of Artiflex lens is considered as an appropriate alternative to glasses and contact lens, especially in patients younger than 40 years with a high refractive error.

According to the results of examinations, 6 months after lens implantation, visual acuity in both eyes was significantly improved, so that the mean BCVA before surgery was 0.11±0.23 LogMAR, which improved to 0.06±0.19 LogMAR postoperatively (p<0.001), and the mean UCVA before the surgery was 1.68±0.16 LogMAR, which improved to 0.10±0.22 LogMAR (p<0.001). After the surgery, UCVA was 20/40 or better in 93% of the eyes, and BCVA was 20/30 or better in 94% of the eyes. In a study by Akcay et al21 in 2012, from 31 patients undergoing phakic anterior chamber lens implantation, 80% of the eyes showed improved BCVA 18 months after the surgery. In a study by Dick et al22 in 2009 on 191 patients receiving Artiflex phakic IOL, 37.7% of patients after 6 months and 41.8% of patients after 1 year of Artiflex implantation showed a BSCVA improvement of one line or better. In addition, in their study, 97.4% of the eyes after 6 months and 97.4% of the eyes after 1 year had a UCVA of >20/40.23 In another study by Yuan et al,24 among 43 patients undergoing Artisan iris-fixated IOL implantation, 79.7% of the eyes had a UCVA of 20/25 or better after 6 months and 71.4% of the eyes had a BCVA improvement of one line or better 5 years after surgery.

In our study, 94% of the eyes had a refraction of ±1 D from target refraction within 6 months after the surgery. Similarly, Yuan et al26 reported that 93.2% of the eyes had a refraction of ±1 D from target refraction after 5 years of follow-up. In a study by Tahzib et al,19 74.2% of the eyes had refraction of ±1 D from target refraction after 1 year of Artisan phakic IOL implantation. Also in a study by Coullet et al,16 83.9% of the eyes had refraction of ±1 D from target refraction after 1 year of iris-supported phakic lens implantation.

According to our results, the ECD did not show a statistically significant change six months after refractive surgery.
which is more likely to increase the corneal manipulation compared to baseline count. Similar to our findings Dick et al\(^{23}\) reported no statistically significant changes in the density of endothelial cells 6 months, 1 year and 2 years after Artiflex lens implantation and contributed this to lack of direct contact between the endothelial cells and lens surface. In addition, Tahzib et al\(^{19}\) did not observe a statistically significant reduction in the endothelial cell count 1 month, 6 months and 10 years after Artificial phakic IOL implantation for the correction of moderate to high myopia. In another study by Pop and Payette,\(^ {18}\) the reduction in the endothelial cell count in Artificial lens implantation was <1% in 6, 12 and 24 months after the surgery, which was not statistically significant.

Our findings are however in contrast with the findings of Akcay et al,\(^ {24}\) who evaluated ECD 1 month and 18 months after phakic anterior chamber lens implantation to correct very high myopia and reported an 8.61% reduction 18 months postoperatively, which was statistically significant. Also in a study by Menezo et al\(^ {20}\) in 1997, which evaluated ECD in 6-month intervals for 5 years, the cell density reduction after iris claw phakic IOL implantation for correction of high myopia was −5.8% in the first 6 months and 7.9% after 12 months, which was statistically significant. In this study, Worst–Fechner iris claw lens with biconcave or concave–convex design was used and some factors such as surgical technique, lens position, lens design, lens material and viscoelastic materials used during surgery had a relation with endothelial cell damage.\(^ {20}\) In a study by Budo et al\(^ {13}\) in year 2000, ECD after Artisan lens implantation was reduced by 4.8% in 6 months and 2.4% in 1 year after the surgery. Benedetti et al\(^ {12}\) also evaluated ECD after Artisan lens implantation and reported a cell loss of 2.3% after 4 months and 3.5% after 12 months, which was statistically significant. It should be noted that in the last three abovementioned studies reporting a significant endothelial cell loss, the surgical incision was −5.5 to 6.5 mm, which is more likely to increase the corneal manipulation and surgical trauma compared with Artiflex lens implantation, with an incision of −3.2 mm. In addition, the mean age of subjects in all three studies was higher than the average age of our patients, which might explain part of the difference in reported endothelial cell loss. It should also be noted that although the mean percentage of decrease in ECD at 6 months in the present study was only 2.1% and not statistically significant, with a longer follow-up, this may become statistically significant compared to the normal age-related decline.

In the present study, the CV in the sizes of endothelial cells (polymegathism) before the surgery was 25.7%±7.1%, which changed to 25.9%±5.4% 6 months postoperatively, showing no statistically significant change (p=0.857; Table 2). Similarly in 2007, Benedetti et al\(^ {12}\) did not find a significant change in the CV in the sizes of endothelial cells 5 years after Artisan lens implantation. In addition, Menezo et al\(^ {20}\) found no significant change in the CV after iris claw phakic IOL implantation for correction of high myopia.

Some limitations of the present study were the retrospective nature of the study, a relatively low number of participants and a short period of follow-up.

**Conclusion**

Artiflex iris-fixated lens implantation is a suitable and predictable method for correction of moderate to high myopia. There was no statistically significant change in cell density and CV in the sizes of endothelial cells after 6 months of follow-up.

**Disclosure**

The authors report no conflicts of interest in this work.

**References**


