

ORIGINAL ARTICLE

Comparative evaluation of Comfilcon A and Senofilcon A bandage contact lenses after transepithelial photorefractive keratectomy



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KEYWORDS

Photorefractive keratectomy;
Transepithelial PRK;
Silicone hydrogel contact lens;
Bandage contact lens

Abstract

Purpose: To evaluate and compare Comfilcon A and Senofilcon A silicone hydrogel contact lenses used as a therapeutic bandage following transepithelial excimer laser photorefractive keratectomy (PRK).

Methods: Patients undergoing transepithelial PRK for myopia were prospectively recruited. Included patients had a Comfilcon A silicone hydrogel lens inserted in one eye, with a Senofilcon A lens in the contralateral eye. Postoperative assessment of subjective pain, epithelial healing and visual recovery was at day 1, 3 and 7. Contact lens factors including centration, movement and deposits were assessed.

Results: 48 eyes of 24 patients were included in the study. Mean age was 31 years (SD 11) and mean refractive error -4.5 D (SD 1.8). Mean pain score at day 1 was significantly higher in the Comfilcon group at 4.6 (SD 2.7) vs. 1.5 (SD 2.5) in the Senofilcon group ($P < 0.005$). Mean time to healing was 3.17 days (SD 0.37) in the Comfilcon group, and 3.21 days (SD 0.4) in the Senofilcon group, with no difference in defect size. There was a pronounced central raphe in 1 eye in the Comfilcon group vs. 5 eyes in the Senofilcon group ($P = 0.19$). Significantly more eyes demonstrated no lens movement in the Senofilcon group (18 vs. 4, $P = 0.0001$).

Conclusion: The variation in material characteristics and lens geometry of different silicone hydrogel lenses affects their clinical characteristics in therapeutic roles. Other factors than oxygen permeability may affect pain and epithelial healing, with superior pain relief from the less permeable Senofilcon lens in this study.

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PALABRAS CLAVE

Queratectomía
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PRK transepitelial;
Lentes de contacto
de hidrogel de
silicona;
Lentes de contacto
como apósito

Evaluación comparativa de las lentes de contacto Comfilcon A y Senofilcon A como apósito, tras queratectomía fotorrefractiva transepitelial

Resumen

Objetivo: Evaluar y comparar las lentes de contacto de hidrogel de silicona Comfilcon A y Senofilcon A, utilizadas como apósito terapéutico tras la realización de una queratectomía fotorrefractiva (PRK) mediante láser excimer transepitelial.

Métodos: Se incluyó prospectivamente a pacientes sometidos a PRK transepitelial para corrección de miopía. Los pacientes incluidos en el estudio utilizaban una lente de hidrogel de silicona Comfilcon A en un ojo, y una lente Senofilcon A en el ojo contralateral. La evaluación postoperatoria del dolor subjetivo, reepitelización y recuperación visual se realizó los días 1, 3 y 7. Se evaluaron factores sobre lentes de contacto tales como centrado, movimiento y depósitos.

Resultados: El estudio se completó incluyendo 48 ojos de 24 pacientes. La edad media fue de 31 años (DE 11) y el error refractivo medio de -4,5 D (DE 1,8). El índice medio de dolor en el día 1 fue considerablemente superior en el grupo de Comfilcon, con un valor de 4,6 (DE 2,7), frente a 1,5 (DE 2,5) en el grupo de Senofilcon ($P < 0,005$). El tiempo medio de reepitelización fue de 3,17 días (DE 0,37) en el grupo de Comfilcon, y de 3,21 días (DE 0,4) en el grupo de Senofilcon, sin diferencia en cuanto a tamaño de defecto. Se produjo erosión central pronunciada en un ojo del grupo de Comfilcon, frente a cinco ojos en el grupo de Senofilcon ($P = 0,19$). Los ojos del grupo de Senofilcon reflejaron una considerable diferencia en cuanto a ausencia de movimiento de la lente (18 frente a 4, $P = 0,0001$).

Conclusión: La variación en cuanto a características del material y geometría de las distintas lentes de hidrogel de silicona afecta a sus características clínicas de función terapéutica. En este estudio, otros factores diferentes a la permeabilidad del oxígeno pueden afectar al dolor y a la curación epitelial, con un alivio superior del dolor en el grupo de las lentes Senofilcon, que son menos permeables.

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Introduction

Photorefractive keratectomy (PRK) is an established procedure for the excimer laser correction of refractive errors.¹ One of the major limitations of PRK is the postoperative pain and delay of visual recovery while epithelial healing takes place. Soft hydrogel contact lenses have been widely used as a bandage to reduce pain and optimise the epithelial healing process, both in PRK and subsequently LASEK.² In this situation, the bandage contact lens is typically inserted at the time of surgery and used on a continuous extended wear basis for 2–7 days. The most commonly used type of lens for this indication has been conventional hydrogel material. However these lenses are not intended for extended wear, and do not provide adequate corneal oxygenation, which may inhibit the healing process.^{3–6} Following the introduction of silicone hydrogel lenses, there is a superior option to maintain oxygenation during epithelial regeneration. Existing studies support this notion, with faster re-epithelialisation and reduced discomfort using silicone hydrogel lenses compared to conventional hydrogel lenses.^{3,4,7–10} Different silicone hydrogel lenses vary considerably with respect to a number of characteristics in addition to oxygen permeability alone (Table 1). These include water content, material stiffness (Young's modulus), thickness, edge profile, surface smoothness, surface treatments, hydrophobicity, size and base curve. The effect of these characteristics individually on pain and epithelial healing is not well elucidated; nevertheless there may be considerable variation in the performance of different silicone lenses. In the current study

we determined the relative merits of two silicone hydrogel lenses when used as a bandage following transepithelial photorefractive keratectomy.

Methods

This was a prospective double masked contralateral eye study carried out at a refractive surgical centre in Heraklion, Greece. After undergoing transepithelial PRK for myopia, a Comfilcon A (Biofinity, Coopervision, Fairport, NY) silicone hydrogel lens was inserted in one eye, with a Senofilcon A (Acuvue Oasys, Johnson & Johnson, Jacksonville, FL) lens in the contralateral eye. Patients were included subject to informed consent, and prior ethical approval was obtained from the institutional review board. The research conformed to the tenets of the Declaration of Helsinki. Patients were excluded in the presence of any ocular or systemic condition which might affect epithelial healing or pain perception. In particular patients with significant dry eye or any corneal epithelial abnormality were excluded.

Surgical technique

Preoperative topical anaesthesia was with 0.5% preservative free proxymetacaine. All treatments were carried out with single step transepithelial PRK using the Schwind Amaris 750 excimer laser platform. This excimer laser platform uses a 750 Hz flying spot laser of 0.54 mm super Gaussian profile, with 6-axis eye tracking. In the transepithelial treatment,

Table 1 Characteristics of Silicone hydrogel lenses used in study and comparison with other commonly available Silicone hydrogel lenses.

Material	Comfilcon A	Senofilcon A	Lotrafilcon A	Lotrafilcon B	Balafilcon A	Galyfilcon A
Proprietary name	Biofinity	Acuvue OASYS	Night & Day	O2 Optix/Air Optix	PureVision	Acuvue Advance
Manufacturer	CooperVision	Johnson & Johnson	CIBA Vision	CIBA Vision	Bausch & Lomb	Johnson & Johnson
Water content	48%	38%	24%	33%	36%	47%
Dk	128	103	140	110	91	60
Dk/t	160	147	175	138	101	86
Surface treatment	None	None (internal wetting agent)	Plasma Coating	Plasma Coating	Plasma Oxidation	None (internal wetting agent)
Modulus (Mpa)	0.8	0.72	1.5	1.0	1.1	0.43

Dk: Oxygen permeability ($\times 10^{-11}$); Dk/t: oxygen transmissibility ($\times 10^{-9}$).

excimer ablation of the epithelium and refractive ablation of the stroma are carried out in one continuous ablation profile. In contrast to mechanical or alcohol assisted epithelial removal during conventional PRK, laser epithelial removal in tPRK results in an epithelial defect of consistent size, with a regular smooth edge matching the stromal ablated area. This predictable regular defect avoids the inherent variability of epithelial defects present in conventional PRK, and thus provides a better experimental context for the evaluation of postoperative recovery than conventional PRK. A Comfilcon A silicone hydrogel lens (base curve 8.6 mm, diameter 14.0 mm, power -0.50 D) was inserted in one eye, with a Senofilcon A lens (base curve 8.4 mm, diameter 14.0 mm, power -0.50 D) in the contralateral eye. The laterality of insertion was determined from a custom randomisation table.

Clinical assessment

Preoperative history, refraction, vision and absence of corneal and ocular surface pathology were noted. Although patients were aware from the consent process that different contact lenses would be used in either eye, no information as to the nature of the specific lenses was provided. The operating surgeon was also not aware of the random lens choice selected by the assistant at the time of the procedure. The assessing doctor postoperatively was not aware of the laterality of lens used. Postoperative study assessment was at day 1, 3 and 7 with an additional visit at 5 days if epithelialisation was incomplete at day 3. Postoperative assessment at each visit was carried out by the same experienced practitioners, and included vision and slitlamp biomicroscopy. Subjective pain perception was assessed using a 0–10 numeric pain rating scale administered verbally, with 0 representing no discomfort and 10 representing severe pain. The epithelial defect was quantified by measuring the horizontal and vertical size at the slitlamp using the scale provided with a narrow slit beam. The area was calculated using the major and minor radii of the ellipse thus obtained. Contact lens centration was noted and degree of movement of the lens on blinking classified into none, minimal moderate or large. Contact lens deposits were noted, and graded according to the scheme of Gil-Cazorla et al.³

Results

The study population consisted of 50 eyes of 25 patients. One patient was excluded due to inadequate follow-up, leaving 48 eyes for analysis. The mean age was 31 years (SD 11: 22–41). Mean spherical equivalent error was -4.5 D (SD 1.8: -8 to -1.5). Baseline comparison of Comfilcon and Senofilcon groups demonstrated no significant difference in spherical equivalent (-4.5 D vs. -4.5 D), astigmatism (-0.7 D vs. -0.8 D) or preoperative vision (0.00 vs. 0.00 LogMAR), with $P < 0.001$ for all variables. All other factors were constant due to the contralateral eye design, and in all cases utilising Mitomycin C this was in both eyes. Raw clinical data for each patient are provided as [Appendix 1](#) to this paper.

Pain

Mean pain score at day 1 was 4.6 (SD 2.7: range 0–10) in the Comfilcon group, and 1.5 (SD 2.5: range 0–7) in the Senofilcon group ([Fig. 1](#)). The pain scores in the Comfilcon group were statistically significantly greater ($P < 0.005$ paired 2 tailed *T*-test). For each patient, the pain score was greater in the eye with Comfilcon lens in 18 (75%), equal in both eyes in 5 (21%) and greater in the eye with Senofilcon

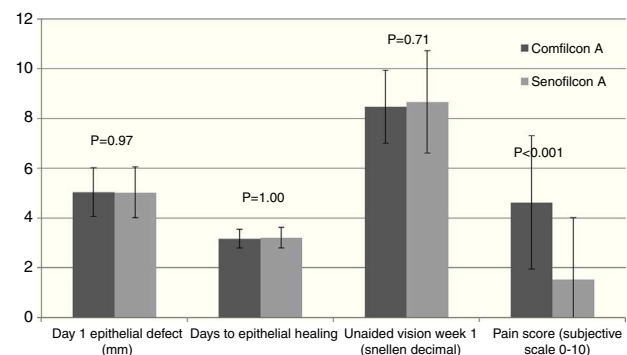


Figure 1 Mean postoperative characteristics of epithelial healing, vision and pain comparing a second and third generation silicone hydrogel bandage contact lens after transepithelial photorefractive keratectomy. Pain was significantly higher in the Comfilcon A group ($P < 0.05$).

difference was not statistically significant, and appeared to be of little clinical relevance. There was also no difference in early visual recovery following healing.

The main finding of this study was of superior pain control following PRK with the Senofilcon compared to the Comfilcon lens, with significantly lower pain scores. This is consistent with the findings of other previous studies. Razmjoo et al. compared a Senofilcon A to lotrafilcon A lens for PRK and found better pain and discomfort control with Senofilcon.¹¹ Qu et al. compared a Galyfilcon A lens to Balafilcon A for LASEK and found better pain control with Galyfilcon.⁷ This is notable because Galyfilcon is a precursor to the development of Senofilcon, and both share many chemical characteristics. Grentzelos et al. compared lotrafilcon A and lotrafilcon B for PRK and found no difference in pain.¹⁵ Plaka et al. compared Asmofilcon A to Lotrafilcon B following PRK, but did not assess subjective pain.¹⁶

The mechanism of pain relief by contact lenses is not well elucidated. Although the Comfilcon lens appears equivalent or superior in a range of material characteristics, there may be other factors involved in pain relief. It is notable that the Senofilcon lens in this study demonstrated significantly reduced movement compared to the Comfilcon lens. The potential benefits of tight fit for bandage use has been suggested by some authors.^{15,17} In this study, a base curve of 8.4 was used for the Senofilcon A lens, while the Comfilcon A lens was of base curve 8.6. It is thus possible that a tighter fit with reduced mobility of the Senofilcon lens accounts for

the greater relief of pain, by preventing blink effects on the exposed nerve endings or friable nascent epithelium.⁶

Conclusion

The expected performance of various silicone hydrogel lenses when used in a therapeutic role for pain or epithelial healing depends on a number of factors, and may not easily be predicted on the basis of material or geometry characteristics. Thus, clinical evaluation of lens types is valuable to determine the optimal lens choice in this setting.

Ethical approval

The study was performed with informed consent and approval by the Local Ethics Committee. Research adhered to the tenets of the Declaration of Helsinki.

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Conflict of interest

None of the authors have conflict of interest with the submission. No author has any financial interest in the subject matter.

Appendix 1. Raw clinical data for each patient

Patient	Eye with Comfilcon A lens				Contralateral eye with Senofilcon A lens			
	Day 1 epithelial defect (mean diameter mm)	Time to epithelial healing (days)	Unaided Vision (LogMAR)	Day 1 pain score	Day 1 epithelial defect (mean diameter mm)	Time to epithelial healing (days)	Unaided Vision (LogMAR)	Day 1 pain score
1	4.5	4	0.05	9	5	4	0.05	0
2	3.6	3	0.00	7	5.7	3	0.00	0
3	6.7	3	0.05	3	4.2	3	0.00	0
4		3	0.00	2		3	0.00	0
5	5.8	4	0.05	0	5.5	3	0.10	0
6	4.2	3	-0.08	5	4	3	-0.08	0
7		3	-0.08	0		3	-0.08	0
8	5.2	3	0.00	6	5.8	3	0.05	0
9	6	3	0.10	8	6.2	3	0.10	6
10	6	3	0.00	3	6.2	3	0.00	1
11	4	3	-0.08	7	5	3	-0.08	3
12	4.5	4	0.05	7	2	4	-0.08	7
13	4	3	0.00	10	5	3	0.00	5
14	4	3	-0.08	3	5.2	3	-0.08	0
15	5.5	4	0.10	2	6	4	0.10	0
16	4.5	3	0.00	2	4	3	0.00	0
17	5.5	3	-0.08	8	5.5	3	-0.08	0
18	6	3	-0.08	4	6	Persistent defect	0.05	7
19	3.2	3	0.00	6	4.7	3	0.00	0
20	5.5	3	-0.08	5	5.7	3	-0.08	0
21	4.2	3	0.10	5	3	3	0.00	5
22	6.5	3	0.00	3	5	4	0.00	0
23	5.5	3	0.00	3	5.5	3	0.05	3
24	6	3	0.05	3	5.5	3	0.05	0

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