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EXTENDED REPORT

Incidence of keratitis of varying severity among contact lens wearers

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Aim: To determine the incidence of non-severe keratitis (NSK) and severe keratitis (SK) among wearers of current generation contact lenses.

Methods: A 12 month, prospective, hospital based epidemiological study was conducted by examining all contact lens wearers presenting with a corneal infiltrate/ulcer to a hospital centre in Manchester. A clinical severity matrix was used to differentiate between NSK and SK, based on the severity of signs and symptoms. The size of the hospital catchment population and the wearing modalities (daily wear (DW) or extended wear (EW)) and lens types being used were estimated from relevant demographic and market data.

Results: During the survey period, 80 and 38 patients presented with NSK and SK, respectively. The annual incidences (cases per 10 000 wearers) for each wearing modality and lens type were: DW rigid—NSK 5.7, SK 2.9; DW hydrogel daily disposable—NSK 9.1, SK 4.9; DW hydrogel (excluding daily disposable)—NSK 14.1, SK 6.4; DW silicone hydrogel—NSK 55.9, SK 0.0; EW rigid—NSK 0.0, SK 0.0; EW hydrogel—NSK 48.2, SK 96.4; EW silicone hydrogel—NSK 98.8, SK 19.8. The difference in SK between EW hydrogel and EW silicone hydrogel was significant ($p=0.04$).

Conclusions: A clinical severity matrix has considerable utility in assessing contact lens related keratitis. There is a significantly higher incidence of SK in wearers who sleep in contact lenses compared with those who only use lenses during the waking hours. Those who choose to sleep in lenses should be advised to wear silicone hydrogel lenses, which carry a five times decreased risk of SK for extended wear compared with hydrogel lenses.

The landscape of contact lens practice has changed dramatically over the past decade, with the introduction of new contact lens genres such as daily disposable lenses^{1,2} and silicone hydrogel materials.³ It might be expected that these new products will be associated with lower levels of ocular morbidity, as well as providing greater levels of comfort and convenience for contact lens wearers. For example, daily disposable lenses could reduce the risk of ocular complications^{1,2} because they obviate the need for lens maintenance and avoid problems related to long term surface spoilation. Silicone hydrogel lenses should minimise clinical problems relating to corneal hypoxia because of their greater capacity to transmit atmospheric oxygen to the ocular surface.⁴ Despite these significant advances, there have been recent case reports of severe microbial keratitis with daily disposable lenses⁵ and silicone hydrogel lenses.^{6,7}

The aim of this study was to undertake a prospective epidemiological investigation of the safety of all current forms of contact lens wear. We employed a previously described novel methodological paradigm capable of differentiating the severity of keratitis associated with contact lens wear.

METHODS

Study setting

A prospective survey was conducted between 25 January 2003 and 24 January 2004 of all patients, who were wearing contact lenses (whether or not this was the primary reason for presentation), attending the acute service of the Royal Eye Hospital, Manchester, United Kingdom (the study centre).

This research followed the tenets of the Declaration of Helsinki. Informed consent was obtained from all patients surveyed in this study. Ethical approval to conduct this study was obtained from the local research ethics committee (central) of Manchester Health Authority.

Survey procedure

During the survey period, all presenting patients were asked if they habitually wore contact lenses. Patients giving an affirmative response were asked to complete a green coloured patient survey questionnaire (fig 1) to provide information such as personal details, whether or not they sleep in lenses (wearing modality), lens type (defined in terms of the material from which the lens is fabricated and/or the lens replacement frequency), lens care system used, and their reason for attending the study centre.

When a patient was examined, the presence of any form of corneal infiltrate or ulcer was noted. If there were no evidence of such corneal involvement, no further information was captured for the purposes of this survey. If there were a corneal infiltrate or ulcer, the attending clinician completed a red coloured clinical survey form (on the reverse side of the patient survey questionnaire; fig 1) which gathered clinical information relevant to this study. The colour coding of the survey forms facilitated ease of implementation in a busy clinical environment.

A key feature of the clinical data captured was the use of a clinical severity matrix as described by Aasuri *et al*⁸ (with minor textual adaptations) (fig 1). This involved the clinician encircling a severity descriptor (on a 0 to 3 scale) with respect to 10 pertinent signs and symptoms, providing a mechanism for classifying corneal infiltrative events as non-severe or severe. The cumulative score for each event—the “clinical severity score”—was between 2 (the minimum score relating to the criterion for this study of only including patients exhibiting an infiltrative response) and 22, whereby the higher the score, the more clinically severe the event. Aasuri

Abbreviations: DW, daily wear; EW, extended wear; NSK, non-severe keratitis; PMMA, poly(methyl methacrylate); SK, severe keratitis

Contact lens problem survey: patient side

Name: _____ Sex: Male Female Age: _____ Occupation: _____ Date: _____

Do you smoke? Yes No List any health problems: _____ List current medications: _____

Do you have any other eye problems?
 No Dry eyes Blepharitis Other: _____

What sort of contact lenses do you currently wear?
 Rigid (hard) Soft daily disposable (lenses thrown out after each daily use)
 Soft disposable Soft other (please specify): _____

Do you deliberately sleep in your contact lenses?
 Yes No

Which brand of lens do you use? (tick)
 PureVision (Bausch & Lomb) Focus Night & Day (CIBA Vision) All Day All Night (Specsavers Easyvision) Acuvue (Johnson & Johnson)
 Other (please specify): _____

How often do you sleep in your lenses? 1 2 3 4 5 6 7 nights per week (circle one) How often do you wear your lenses? 1 2 3 4 5 6 7 days per week (circle one)

When did you last sleep in your lenses? _____ When did you last wear your lenses? _____

Which solutions do you use with your lenses (list brand names if you can remember them) _____

When did you notice something was wrong? _____ What did you notice was wrong? _____

What did you do to try and solve the problem? _____

When did you last go swimming? _____ Did you wear swimming goggles? Yes No

It might be helpful to us to contact you at a later stage to clarify some aspects of your contact lens wear. Also, we might need to contact your contact lens practitioner. If you are happy for us to do this, please provide details below, and sign to indicate that you agree to this.

Your daytime phone number: _____ Contact lens practitioner: _____ Signature: _____

Contact lens problem survey: clinician side

Is there a corneal abscess or any form of corneal infiltrate?
 Yes No

If yes, please complete the remainder of this form. If no, do not complete this form, but please file it with the other completed forms.

Patient name: _____ DOB: _____ Hospital number: _____ Date: _____ Nurse/practitioner/Examining doctor: _____

Circle one descriptor on each row, for the parameters listed in the table below.

Parameter	0	1	2	3
Symptoms	None	Mild	Moderate	Severe
Lid swelling	Absent		Present	
Conjunctival redness	Absent	Localised	Generalised	
Infiltrate shape*		Round		Irregular
Infiltrate size (largest diameter)**		≤1.0mm	1.0 - 2.0mm	≥2.0mm
Fluorescein staining	Absent	Present		
Surrounding cornea	Clear	Slight haze	Severe haze	
Endothelial debris	Absent	Present		
Hypopyon	Absent		Present	
Effect of lens discontinuation**	Resolving	No change	Slight worsening	Significant worsening

*Record details of largest infiltrate if more than one ** Circle 'No change' if unknown e.g. if patient presents wearing lenses

Please draw the condition

R

L

Was the cornea scraped? Yes No Was the eye photographed? Yes No Was the patient admitted? Yes No

What is your tentative diagnosis? _____ What is the prescribed treatment? _____

Figure 1 Survey forms completed by patient (green) and clinician (red). The clinical severity matrix referred to in the text can be seen in the middle of the clinician survey form.

*et al*⁸ have validated this scoring system indicating that any score greater than 8 should raise a high index of suspicion of clinically severe microbial keratitis.

This study required the attending clinician to indicate if a corneal scrape for culture was performed, a decision based on their own clinical judgment and unrelated to the conduct of this study. The results of any such scrapes were recorded. Only information relating to wearing modality, lens type, clinical severity score, and outcome of any corneal scrapes undertaken was analysed. Additional information gathered during this survey relating to potential risk factors and outcomes will be the subject of further analysis and is not reported here.

Terminology

In this paper, we use the term “severe keratitis” to describe any clinical presentation which attracted a clinical severity score of more than 8; in conventional clinical parlance, this is presumed to be a case of severe microbial keratitis (essentially the designation given by Aasuri *et al*⁸) which requires intensive antimicrobial treatment.

The term “non-severe keratitis” is used to describe any clinical presentation which attracted a clinical severity score of 8 or less; conditions with this range of scores do not typically require medical intervention. We have chosen to use this term rather than the term “sterile keratitis” adopted by Aasuri *et al*⁸ in view of uncertainty relating to the aetiology and pathogenesis of this category of events.^{9 10}

Patient follow up

To provide assurance of the quality of data collected, an attempt was made to telephone all patients and/or their contact lens practitioner (with permission from the patient). Any unclear, ambiguous, or incomplete data entries were clarified and any required amendments were recorded.

Hospital catchment population

Calculation of the incidence of infiltrative events necessitated an estimation of the catchment population of the study centre. We applied the technique of Bailey (reported by Senn and Sampson¹¹) to demographic details of the 24 630 patients with all forms of ocular problems presenting to the study centre during the survey period. Essentially, this involved determining in which of the 14 National Health Service primary care trusts (urban geographic regions) within Greater Manchester the patients were resident. The small number of patients resident outside Greater Manchester was also accounted for in this analysis. Assuming that the rates of ocular problems were similar for all areas of residence, a weighted population size was calculated for each area and the sum total of these was determined to be 1 071 503—the “hospital catchment population.”

Determination of lens wearing modality

We have conducted an annual survey on contact lens prescribing trends in the United Kingdom for the past 9 years.¹² This data set comprises information relating to

8833 contact lens fits by 977 practitioners. From these data, we could determine the proportion of each of the four lens types described below that are prescribed for daily wear versus extended wear.

Categorisation of lens type

We categorised contact lens types as follows:

- Rigid—lenses made from silicone-containing rigid gas permeable materials. (Virtually no wearers have been fitted with poly(methyl methacrylate) (PMMA) lenses over the past 9 years in the United Kingdom.¹²)
- Hydrogel daily disposable—single use hydrogel lenses (excluding silicone hydrogel lenses) which are discarded after each daily use
- Hydrogel—all forms of hydrogel lenses (excluding daily disposable and silicone hydrogel lenses). Over the past 9 years, 4% of wearers of this lens type have been prescribed lenses to be replaced fortnightly, 70% monthly, and 18% less frequently; 8% of wearers were advised to replace lenses as and when required (that is, on an unplanned replacement basis)¹²
- Silicone hydrogel—new generation silicone-containing hydrogel lenses; during the time of this survey, only two such materials were available in the United Kingdom (balafilcon A and lotrafilcon A). Both of these lens types are prescribed for monthly replacement.¹²

Determination of number of lens wearers

Information on the number of contact lens wearers in the United Kingdom in each of the above lens type categories was accessed from lens sales data collected by the UK Association of Contact Lens Manufacturers for 2003.¹³ Using the data outlined above on wearing modality and lens type, and assuming a total UK population of 59 050 800,¹⁴ we were able to estimate the number of wearers within the hospital catchment population using each lens type on a daily wear versus extended wear basis. The above analyses assume that lens use in the hospital catchment population is the same as that for the United Kingdom as a whole.

Data analysis

For each wearing modality/lens type combination investigated in this 12 month study, the annual incidence of non-severe and severe keratitis was determined by dividing the number of cases of each of these conditions by the estimated number of wearers for each wearing modality/lens type combination within the hospital catchment population. In common with previous practice in this field, we have expressed incidence data as the number of cases per 10 000 wearers per year. The 95% confidence intervals (CI) for all incidence estimates were calculated according to the methodology of Wilson.¹⁵ The data are also expressed in terms of “relative risk,” which is the ratio of annual incidence of a given condition to an arbitrary “referent” condition,¹⁶ which we have chosen to be daily wear hydrogel lenses.

The significance of differences between various combinations of annual incidence was determined using the χ^2 test. For comparisons involving small numbers of presenting patients (that is, less than five for any cell within the analysis), the Fisher’s exact test variant of the χ^2 test was employed.

RESULTS

Wearing modality and lens type

The number of wearers of each combination of wearing modality/lens type in the United Kingdom and the hospital catchment population is presented in table 1.

Survey outcome

Over the 12 month survey period, 415 patients who were wearing contact lenses presented to the study centre. None of these patients was wearing PMMA lenses. All of these patients except one agreed to complete a survey form. Five forms were unintelligible and therefore were not analysed in this study. The remaining usable forms provided details of 118 patients who were observed to have a corneal infiltrate or ulcer. For these patients, a clinical survey form was completed by the attending clinician while examining the patient. A clinical severity score for each case was subsequently determined from the sum of the gradings on the clinical severity matrix.

Figure 2 shows the distribution of clinical severity scores for the 118 patients included in this analysis, stratified by wearing modality and lens type. According to the scoring system, 80 patients presented with non-severe keratitis and 38 with severe keratitis. The number of patients who presented with non-severe and severe keratitis with respect to each of the two wearing modalities and four lens types is presented in table 2.

Table 2 documents one case of non-severe keratitis and two cases of severe keratitis in patients wearing hydrogel daily disposable lenses on an “extended wear” basis. Although such lenses are self evidently prescribed for daily wear, in these three cases the patients admitted to being non-compliant by way of sleeping in lenses immediately

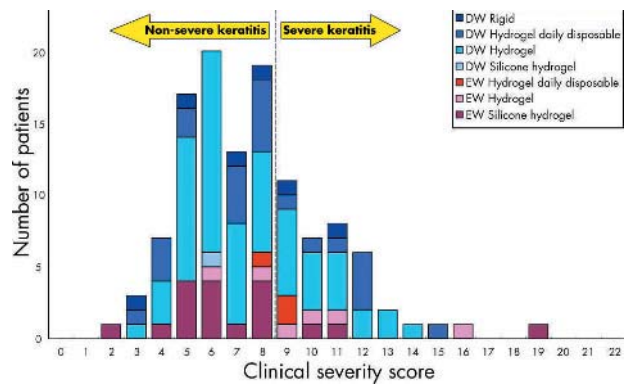


Figure 2 Distribution of clinical severity scores for non-severe and severe keratitis stratified by wearing modality and lens type.

Table 1 Contact lens demographics

Lens type	Wearers in UK	Wearers in HCP*	Daily wear	Extended wear	Daily wearers in HCP	Extended wearers in HCP
Rigid	386 186	7014	99.75%	0.25%	6996	18
Hydrogel daily disposable	903 765	16 413	100.00%	0.00%	16 413	0
Hydrogel	1 667 905	30 291	98.63%	1.37%	29 876	415
Silicone hydrogel	93 410	1696	10.55%	89.45%	179	1517

*Hospital catchment population.

Table 2 Number of patients with non-severe and severe keratitis*

Wearing modality	Lens type	Non-severe keratitis	Severe keratitis	Total
Daily wear	Rigid	4	2	6
	Hydrogel daily disposable	15	8	23
	Hydrogel	42	19	61
	Silicone hydrogel	1	0	1
Extended wear	Rigid	0	0	0
	Hydrogel daily disposable	1	2	3
	Hydrogel	2	4	6
	Silicone hydrogel	15	3	18
Total	80	38	118	

*Based on a threshold diagnostic score of non-severe keratitis $\leq 8 <$ severe keratitis.

before presentation at the hospital; data relating to these cases are therefore entered as extended wear incidents.

Patient follow up

Our quality assurance procedure resulted in 70 of these 118 patients and/or their contact lens practitioners being contacted directly. Of this subgroup, data relating to 67 patients were determined to be accurate and adjustments were needed with respect to the other three patients. For the 48 patients/practitioners who could not be contacted—or declined permission to be contacted—it was judged that the information supplied was sufficiently clear to be included in the analysis.

Incidence and relative risk of non-severe and severe keratitis

Incidence and relative risk data are presented in table 3. Comparisons of the statistical significance of differences in incidence of non-severe and severe keratitis between various pairs of combinations of wearing modality/lens type are displayed in table 4. Although caution must be exercised in interpreting p values with multiple comparison testing (because of the increased likelihood of type I statistical error), we feel that the comparisons in table 4 are instructive in that they offer useful guidance in the management of contact lens wearers.

Corneal cultures

Twenty five corneal scrapes were performed on the keratitis patients surveyed in this study; the culture results are presented in table 5. A clinical decision was taken to execute a corneal scrape in two of 80 cases which were classified by the scoring system to be non-severe keratitis, and bacterial growth was detected in both cases. A corneal scrape was performed in 23 of the 38 cases classified as severe keratitis, yielding a positive culture result for bacteria in nine cases and fungus in one case.

DISCUSSION

In the epidemiological studies of Poggio *et al*¹⁷ and Cheng *et al*,¹⁸ participating clinicians were required to include cases of microbial keratitis based on the following clinical criterion: "...a corneal stromal infiltrate with an overlying epithelial abnormality (ulceration) clinically diagnosed as microbial keratitis ... [and] ... antimicrobial treatment ...". In essence, this criterion relates to clinically severe microbial keratitis, which is broadly equivalent to the category of "severe keratitis" in the present study; as such, a comparison of results between these studies can be made.

The finding in this study of a higher overall incidence of severe keratitis in wearers who sleep in contact lenses compared with those who only use lenses during waking hours is consistent with previous reports.¹⁷⁻¹⁹ However, the annual incidence of severe keratitis per 10 000 wearers with hydrogel lenses worn on a daily wear basis reported here (6.4 (95% CI 4.1 to 9.9)) is about 1.7 \times greater than the figures of Poggio *et al*¹⁷ (4.1 (95% CI 2.9 to 5.2)) and Cheng *et al*¹⁸ (3.5 (95% CI 2.7 to 4.5)). Furthermore, we report a much higher annual incidence (about 4.7 \times) of severe keratitis for hydrogel lenses worn on an extended wear basis (96.4 (95% CI 37.5 to 245.2)) relative to the figures reported by Poggio *et al*¹⁷ (20.9 (95% CI 15.1 to 26.7)) and Cheng *et al*¹⁸ (20.0 (95% CI 10.3 to 35.0)).

There are various possible reasons for the higher incidence figures relating to severe keratitis reported in this study. The highly managed, single centre methodology adopted here is likely to have resulted in a higher rate of capture of severe keratitis cases compared with the multicentre approach of Poggio *et al*¹⁷ and Cheng *et al*,¹⁸ whereby reliance was placed upon full compliance in terms of case capture and reporting by 612 and 379 doctors, respectively, spread over large geographic areas. A higher capture rate will result in higher and necessarily more accurate incidence figures. Another reason for the above discrepancy relates to the choice of criterion adopted in relation to the classification of

Table 3 Annual incidence and relative risk of non-severe and severe keratitis in contact lens wear

Wearing modality	Lens type	Annual incidence*		Relative risk‡	
		Non-severe keratitis	Severe keratitis	Non-severe keratitis	Severe keratitis
Daily wear	Rigid	5.7 (2.2 to 14.7)†	2.9 (0.8 to 10.4)	0.4 (0.2 to 1.1)	0.5 (0.1 to 1.9)
	Hydrogel daily disposable	9.1 (5.5 to 15.1)	4.9 (2.5 to 9.6)	0.7 (0.4 to 1.2)	0.8 (0.3 to 1.8)
	Hydrogel	14.1 (10.4 to 19.0)	6.4 (4.1 to 9.9)	1.0	1.0
	Silicone hydrogel	55.9 (9.9 to 309.7)	0.0 (0.0 to 210.1)	4.0 (0.6 to 28.7)	§
Extended wear	Rigid	0.0 (0.0 to 1758.8)	0.0 (0.0 to 1758.8)	§	§
	Hydrogel	48.2 (13.2 to 174.0)	96.4 (37.5 to 245.2)	3.4 (0.8 to 14.1)	15.2 (5.2 to 44.4)
	Silicone hydrogel	98.8 (60.0 to 162.5)	19.8 (6.7 to 58.0)	7.0 (3.9 to 12.7)	3.1 (0.9 to 10.5)

*Number of cases per 10,000 wearers per year.

†95% confidence limits.

‡Calculated independently for non-severe and severe keratitis, taking "daily wear hydrogel" as the referent in each case.

§Indeterminable because of an annual incidence of zero.

Table 4 Statistical test results of comparisons of incidences of non-severe and severe keratitis

Modality of wear Lens type	Type of keratitis	Daily wear Hydrogel daily disposable		Daily wear Hydrogel		Daily wear Silicone hydrogel		Extended wear Hydrogel		Extended wear Silicone hydrogel	
		χ^2	p Value	χ^2	p Value	χ^2	p Value	χ^2	p Value	χ^2	p Value
Daily wear	Non-severe	0.4	0.46	3.2	0.09	6.3	0.12	8.7	0.04	48.6	0.0001*
Rigid	Severe	0.5	0.73	1.2	0.40	†	†	42.4	0.0001	6.1	0.04
Daily wear	Non-severe			2.1	0.15*	4.0	0.16	6.1	0.06	67.0	0.0001*
Hydrogel daily disposable	Severe			0.4	0.53*	†	†	47.6	0.0002	5.0	0.06
Daily wear	Non-severe					2.2	0.23	3.3	0.12	57.3	0.0001*
Hydrogel	Severe					†	†	43.7	0.0003	3.7	0.09
Daily wear	Non-severe							0.0	1.00	0.3	1.00
Silicone hydrogel	Severe							†	†	†	†
Extended wear Hydrogel	Non-severe									1.0	0.55
	Severe									5.3	0.04

*p value determined using χ^2 test (all other p values determined using the Fisher's exact variant of the χ^2 test).

†Indeterminable because of the absence of severe keratitis with daily wear silicone hydrogel lenses.

corneal infiltrative events as representing non-severe versus severe keratitis. The sensitivity of the assessment criterion in making this differential diagnosis can be adduced from inspection of figure 2, and is demonstrated with respect to the scoring system used in this study in table 6. As can be seen from this table, when the threshold increases, the incidence of non-severe keratitis increases and the incidence of severe keratitis decreases. In addition, the employment of a clinical severity matrix such as that used in the present work has the advantage of greatly reducing between observer variability in differentially diagnosing non-severe from severe keratitis.

The relative risk of developing severe keratitis for extended wear versus daily wear of hydrogel lenses was higher in this study (15.2 \times) compared with the data of Schein *et al*²⁰ (5.2 \times) and Cheng *et al*¹⁸ (5.7 \times). It is interesting to observe that, in the present study, relative risk is not markedly affected by the choice of threshold clinical severity score; for example, adopting threshold scores of 6, 8, and 10 gives rise to relative risk values for severe keratitis for extended wear relative to daily wear of hydrogel lenses of 10.9 \times , 15.2 \times , and 16.0 \times , respectively (see table 6).

Stapleton *et al*²¹ reported that the risk of sterile keratitis (which equates to "non-severe keratitis" in the present study) for extended wear hydrogel lenses, relative to daily wear hydrogel lenses, was 1.2 \times , whereas we have found a relative risk of 3.4 \times (although our value was not statistically significantly different from unity). The reasons for the discrepancies in relative risk reported in this study compared with the data of Schein *et al*,²⁰ Cheng *et al*,¹⁸ and Stapleton *et al*²¹ are unclear, but could be because of differences in assumptions relating to the proportion of wearers in the population using lenses on a daily versus extended wear basis.

A key finding of this study is the five times decreased risk of severe keratitis with silicone hydrogel lenses compared to that with hydrogel lenses when worn on an extended wear basis. The essential distinction between these lens types is the superior oxygen transmissibility of silicone hydrogel lenses.⁴ Our observation lends support to the notion that the pathogenesis of severe keratitis is significantly related to corneal oxygen availability. Solomon *et al*²² confirmed such an association in a rabbit eye model by demonstrating higher rates of severe microbial keratitis in animals subjected to

Table 5 Outcomes of cultures of corneal scrapes

Patient code	Age	Sex	Wearing modality	Lens type	Clinical severity score	Culture outcome
82	26	F	Daily wear	Hydrogel	7	<i>Staphylococcus</i> spp*
48	30	F	Extended wear	Silicone hydrogel	8	<i>Bacillus</i> spp
36	32	M	Extended wear	Hydrogel daily disposable	9	No growth 5 days
58	40	M	Daily wear	Hydrogel	9	No growth 5 days
197	23	M	Daily wear	Hydrogel	9	No growth 5 days
205	34	M	Daily wear	Hydrogel	9	No growth 5 days
209	35	F	Extended wear	Hydrogel	9	No growth 5 days
362	83	F	Daily wear	Rigid	9	<i>Stenotrophomonas maltophilia</i>
364	23	F	Daily wear	Hydrogel daily disposable	9	No growth 5 days
44	61	M	Extended wear	Silicone hydrogel	10	<i>Propionibacterium acnes</i>
185	20	F	Daily wear	Hydrogel	10	Mixed coliforms
206	51	F	Daily wear	Hydrogel daily disposable	10	<i>Candida albicans</i>
369	30	F	Daily wear	Hydrogel	10	<i>Acinetobacter</i> spp
92	24	F	Daily wear	Hydrogel	11	No growth 5 days
95	20	F	Extended wear	Hydrogel	11	<i>Staphylococcus</i> spp*
195	31	M	Daily wear	Rigid	11	No growth 5 days
291	60	F	Extended wear	Silicone hydrogel	11	No growth 5 days
41	27	F	Daily wear	Hydrogel daily disposable	12	No growth 5 days
213	48	M	Daily wear	Hydrogel	12	Mixed coliforms
295	24	F	Daily wear	Hydrogel daily disposable	12	No growth 5 days
365	19	M	Daily wear	Hydrogel	12	No growth 5 days
372	21	M	Daily wear	Hydrogel	13	No growth 5 days
375	47	M	Daily wear	Hydrogel	14	<i>Pseudomonas</i> spp
93	29	M	Extended wear	Hydrogel	16	<i>Pseudomonas</i> spp
273	25	M	Extended wear	Silicone hydrogel	19	<i>Pseudomonas</i> spp

*Coagulase negative.

Table 6 Incidence and relative risk of non-severe and severe keratitis with various threshold clinical severity scores

Wearing modality	Lens type	Threshold score* = 4						Threshold score = 6						Threshold score = 8						Threshold score = 10						Threshold score = 12					
		Non-severe keratitis			Severe keratitis			Non-severe keratitis			Severe keratitis			Non-severe keratitis			Severe keratitis			Non-severe keratitis			Severe keratitis			Non-severe keratitis			Severe keratitis		
		AI†	RR‡	AI	RR	AI	RR	AI	RR	AI	RR	AI	RR	AI	RR	AI	RR	AI	RR	AI	RR	AI	RR	AI	RR	AI	RR	AI	RR		
Daily wear	Rigid	1.4	1.1	7.1	0.4	2.9	0.3	0.5	5.7	0.4	2.9	0.4	0.4	7.1	0.4	1.4	0.5	8.6	0.4	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	0.0	
	Hydrogel daily disposable	2.4	1.8	11.6	0.6	3.7	0.4	10.4	0.9	9.1	0.7	4.9	0.8	10.4	0.6	3.7	1.2	13.4	0.7	0.6	0.6	0.6	0.6	0.6	0.6	0.6	0.6	0.6	0.6	0.6	
	Hydrogel	1.3	1.0	19.1	1.0	9.4	1.0	11.0	1.0	14.1	1.0	6.4	1.0	17.4	1.0	3.0	1.0	19.4	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0		
Extended wear	Silicone hydrogel	0.0	§	55.9	2.9	55.9	6.0	0.0	0.0	55.9	4.0	0.0	§	55.9	3.2	0.0	§	55.9	2.9	0.0	§	0.0	§	0.0	§	0.0	§	0.0	§	0.0	
	Rigid	0.0	§	0.0	§	0.0	§	0.0	§	0.0	§	0.0	§	0.0	§	0.0	§	0.0	§	0.0	§	0.0	§	0.0	§	0.0	§	0.0	§	0.0	
	Hydrogel	0.0	§	144.6	7.6	24.1	2.6	120.5	10.9	48.2	3.4	96.4	15.2	96.4	5.5	48.2	16.0	120.5	6.2	24.1	24.0	24.0	24.0	24.0	24.0	24.0	24.0	24.0	24.0		
Extended wear	Silicone hydrogel	13.2	9.8	105.4	5.5	65.9	7.0	52.7	4.8	98.8	7.0	19.8	3.1	105.4	6.1	13.2	4.4	112.0	5.8	6.6	6.6	6.6	6.6	6.6	6.6	6.6	6.6	6.6	6.6		

*A score equal to or less than the threshold score is classified as non-severe keratitis.
 †Annual incidence, expressed as the number of cases per 10 000 wearers per year.
 ‡Relative risk, calculated independently for non-severe and severe keratitis, taking "daily wear hydrogel" as the referent in each case.
 §Indeterminable because of an annual incidence of zero.

lower levels of corneal oxygenation. Ren *et al*²³ demonstrated that *Pseudomonas* bacteria are less likely to attach to corneal epithelial cells in the rabbit cornea (bacterial attachment to the cornea being a necessary precursor for infection) when contact lenses of higher oxygen transmissibility are worn.

The above discussion has concentrated on our findings with respect to silicone hydrogel lenses. Daily disposable lenses are another recent innovation in the contact lens field. Notwithstanding the advantages of this lens type in terms of wearer convenience and avoiding adverse effects related to long term deposit formation,^{1 2} we have found no statistically significant difference in the incidences of severe keratitis among wearers of daily wear daily disposable lenses versus daily wear hydrogel lenses that are replaced less frequently.

Of the 23 corneal scrapes performed on patients deemed to have severe keratitis (according to the scoring system used), 10 were culture positive. This is equivalent to a culture positive rate of 43%, which is consistent with that observed in other contact lens studies.^{18 20} *Pseudomonas* species, which are generally considered to be the most virulent bacterial pathogen in contact lens related keratitis,²⁴ were associated with the three highest clinical severity scores of the patients on whom a corneal scrape was performed.

We conclude that (a) a clinical severity matrix has considerable utility in assessing contact lens related keratitis, (b) there is a significantly higher incidence of severe keratitis in wearers who sleep in contact lenses compared with those who only use lenses during the waking hours and (c) those who choose to sleep in lenses should be advised to wear silicone hydrogel lenses as they carry a five times lower risk of severe keratitis for extended wear compared with hydrogel lenses.

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REFERENCES

- 1 Sankaridurg PR, Sweeney DF, Holden BA, *et al*. Comparison of adverse events with daily disposable hydrogels and spectacle wear: results from a 12-month prospective clinical trial. *Ophthalmology* 2003;**110**:2327–34.
- 2 Solomon OD, Freeman MI, Boshnick EL, *et al*. A 3-year prospective study of the clinical performance of daily disposable contact lenses compared with frequent replacement and conventional daily wear contact lenses. *Contact Lens Assoc Ophthalmol J* 1996;**22**:250–7.
- 3 Sweeney DF. *Silicone hydrogels. Continuous wear contact lenses*. 2nd ed. Oxford: Butterworth-Heinemann, 2004.
- 4 Alvord L, Court J, Davis T, *et al*. Oxygen permeability of a new type of high Dk soft contact lens material. *Optom Vis Sci* 1998;**75**:30–6.
- 5 Hingorani M, Christie C, Buckley RJ. Ulcerative keratitis in a person wearing daily disposable contact lenses. *Br J Ophthalmol* 1995;**79**:1138.
- 6 Lim L, Loughnan MS, Sullivan LJ. Microbial keratitis associated with extended wear of silicone hydrogel contact lenses. *Br J Ophthalmol* 2002;**86**:355–7.
- 7 Whiting MA, Raynor MK, Morgan PB, *et al*. Continuous wear silicone hydrogel contact lenses and microbial keratitis. *Eye* 2004;**18**:935–7.
- 8 Aasuri MK, Venkata N, Kumar VM. Differential diagnosis of microbial keratitis and contact lens-induced peripheral ulcer. *Eye Contact Lens* 2003;**29**(1 Suppl):S60–2.
- 9 Sweeney DF, Jalbert I, Covey M, *et al*. Clinical characterization of corneal infiltrative events observed with soft contact lens wear. *Cornea* 2003;**22**:435–42.
- 10 Baum J, Donshik PC. Corneal infiltrates associated with soft contact lens wear (Letter). *Cornea* 2003;**22**:435–42.
- 11 Senn SJ, Samson WB. Estimating hospital catchment populations. *Statistician* 1982;**31**:81–96.

- 12 **Morgan PB**, Efron N. Trends in UK contact lens prescribing 2003. *Optician* 2003;**225**(5904):34–5.
- 13 **Morgan PB**. Taking the pulse of the UK contact lens market. *Optician* 2004;**227**(5954):26–7.
- 14 www.statistics.gov.uk/ci/nugget.asp?id=6, accessed 4 May 2004.
- 15 **Wilson EB**. Probable inference, the law of succession, and statistical inference. *J Am Stat Assoc* 1927;**22**:209–12.
- 16 **Bland MJ**, Altman DG. The odds ratio. *BMJ* 2000;**320**:1468.
- 17 **Poggio EC**, Glynn RJ, Schein OD, et al. The incidence of ulcerative keratitis among users of daily-wear and extended-wear soft contact lenses. *N Engl J Med* 1989;**321**:779–83.
- 18 **Cheng KH**, Leung SL, Hoekman HW, et al. Incidence of contact-lens-associated microbial keratitis and its related morbidity. *Lancet* 1999;**354**:181–5.
- 19 **Schein OD**, Buehler PO, Stamler JF, et al. The impact of overnight wear on the risk of contact lens-associated ulcerative keratitis. *Arch Ophthalmol* 1994;**112**:186–90.
- 20 **Schein OD**, Glynn RJ, Poggio EC, et al. The relative risk of ulcerative keratitis among users of daily-wear and extended-wear soft contact lenses. A case-control study. Microbial Keratitis Study Group. *N Engl J Med* 1989;**321**:773–8.
- 21 **Stapleton F**, Dart J, Minassian D. Nonulcerative complications of contact lens wear. Relative risks for different lens types. *Arch Ophthalmol* 1992;**110**:1601–6.
- 22 **Solomon OD**, Loff H, Perla B, et al. Testing hypotheses for risk factors for contact lens-associated infectious keratitis in an animal model. *Contact Lens Assoc Ophthalmol J* 1994;**20**:109–13.
- 23 **Ren DH**, Petroll WM, Jester JV, et al. The relationship between contact lens oxygen permeability and binding of *Pseudomonas aeruginosa* to human corneal epithelial cells after overnight and extended wear. *Contact Lens Assoc Ophthalmol J* 1999;**25**:80–100.
- 24 **Fleiszig SM**, Evans DJ. The pathogenesis of bacterial keratitis: studies with *Pseudomonas aeruginosa*. *Clin Exp Optom* 2002;**85**:271–8.

ECHO

Familial keratoconus maps to chromosome 2p24



Please visit the *British Journal of Ophthalmology* website (www.bjophthalmol.com) for a link to the full text of this article.

Researchers have taken an important step to understanding keratoconus (KC), a common reason for corneal transplantation in the developed world. They did so with the first scan of the whole genome in different races, identifying a new locus likely to be responsible for a significant proportion of cases of isolated familial KC.

Genotyping followed by linkage analysis in a small sample of families suggested linkage on chromosome 2p for marker D2S305 but not to chromosomes 16q, 18p, or 21q, previously suggested for familial KC. Typing the full complement of 28 families for 36 other markers across an 80 cM region spanning D2S319 and D2S286 suggested linkage at D2S305. Further analysis suggested significant linkage to 2p24. Marker alleles segregated at 2p24 in affected members in 17 of the 28 families, and the locus for KC was placed in a 1.69 Mb region, within which were two intervals common to all 17. None of the five known genes in the two intervals yet has a known role in human disease, but research is continuing into the expression of all candidate genes

The families were of Caucasian, Arabic, or Caribbean African descent. Among the 253 family members at least two per family had the disorder. The cause of KC is still unknown. The familial form accounts for 6–10% of cases. Mostly it seems to show autosomal dominant inheritance, though autosomal recessive and, rarely, X linked inheritance has been claimed. Various other loci have been mooted—in restricted, homogeneous populations—maybe pointing to genetic heterogeneity.

▲ Hutchings H, et al. *Journal of Medical Genetics* 2005;**42**:88–94.