

# Fast Forward to the Future

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1

## Lyndon Jones PhD, DSc, FRSC, FCAHS, FCOptom, FAAO Financial Disclosures

*Over the past three years, CORE has received research funding and/or honoraria from the following 22 companies:*

- |                        |                            |                         |
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| • Allied Innovations   | • i-Med Pharma             | • Oté Pharma            |
| • Aurinia Pharma       | • Johnson & Johnson Vision | • PS Therapy            |
| • Azura Ophthalmics    | • Lubris                   | • Santen                |
| • Bausch Health Canada | • Menicon                  | • SightGlass            |
| • BHVI                 | • Nature's Way             | • SightSage             |
|                        |                            | • Visioneering Tech Inc |

2

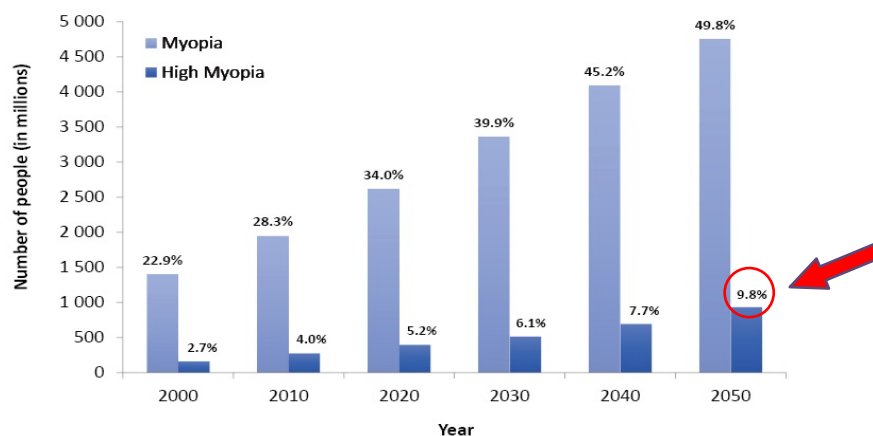
## CLEAR: Contact lens technologies of the future

- Diagnosis & screening for systemic disease
- Diagnosis & screening for ocular disease
- Treatment of ocular conditions
- Drug delivery to the ocular surface
- Antimicrobial lenses
- Theranostics
- Optical enhancements
- Contact lens packaging
- Storage cases



**Available Now!**

## Increase in high myopia



6

## Myopia - what are the risks

Emmetropia = 1x	Cataract (PSCC)	Retinal Detachment	Myopic Maculopathy
-1.00 to -3.00	2.1	3.1	2.2
-3.00 to -6.00	3.1	9.0	9.7
-6.00 or greater	5.5	21.5	40.6



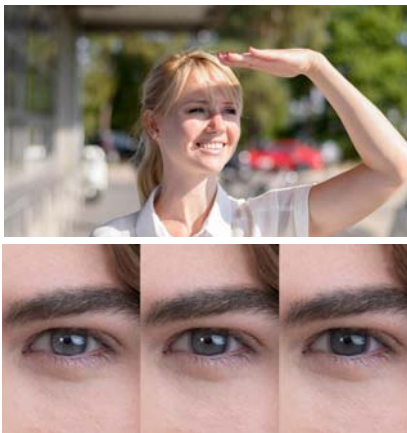
7

# Myopia Management Products Available



8

## Contact lens innovation



- ACUVUE® OASYS with Transitions™
- Launched in US & Canada, April 2019
- First-of-its-kind photochromic contact lens
- Filters light both outdoors and indoors



<https://www.jnj.com/latest-news/acuvue-oasys-with-transitions-contact-lenses-make-2018-times-best-inventions-list>  
<https://www.jnjvisionpro.com/acuvue-transitions>

9

# Diagnosis & Screening for Systemic Disease

10

10

## Wearable Sensor Market

- Market for wearable technology is expected to reach \$US 31.27 Billion by 2025, at annual growth rate of 17.8%
  - smartwatches
  - wristbands
- Major players
  - Adidas; Apple; Fitbit; Garmin; Google; Nike; Samsung; Sony



A MUCH More Diversified Market Than Investors Realize



CREDIT SUISSE

11

# Wearable Sensor Market



Menu Weekly edition Q Search



Review

## Wearable Health Devices—Vital Sign Monitoring, Systems and Technologies

Duarte Dias <sup>1,\*</sup> and João Paulo Silva Cunha <sup>1,2</sup>

<sup>1</sup> Biomedical Research and INnovation (BRIN), Centre for Biomedical Engineering Research (C-BER), INESC Technology and Science, Porto 4200-465, Portugal; jpcunha@fe.up.pt

<sup>2</sup> Faculty of Engineering, University of Porto, Porto 4200-465, Portugal  
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Received: 31 May 2018; Accepted: 21 July 2018; Published: 25 July 2018



**Abstract:** Wearable Health Devices (WHDs) are increasingly helping people to better monitor their health status both at an activity/fitness level for self-health tracking and at a medical level providing more data to clinicians with a potential for earlier diagnostic and guidance of treatment. The technology revolution in the miniaturization of electronic devices is enabling to design more reliable and adaptable wearables, contributing for a world-wide change in the health monitoring approach. In this paper we review important aspects in the WHDs area, listing the state-of-the-art of wearable vital signs sensing technologies plus their system architectures and specifications. A focus on vital signs acquired by WHDs is made: first a discussion about the most important vital signs for health assessment using WHDs is presented and then for each vital sign a description is made concerning its origin and effect on health, monitoring needs, acquisition methods and WHDs and recent scientific developments on the area (electrocardiogram, heart rate, blood pressure, respiration rate, blood oxygen saturation, blood glucose, skin perspiration, capnography, body

## Wearable technology promises to revolutionise health care

Do not let bureaucracy delay matters



May 10th 2018

Share

**I**T is a stealthy killer. When the heart's chambers beat out of sync, blood pools and clots may form. Atrial fibrillation causes a quarter of more than 100,000 strokes in Britain each year. Most of those would never happen if the heart arrhythmia were treated, but first it has to be found. Tests are costly and inaccurate, but Apple Watches, and soon Fitbits, can detect it, are far cheaper and can save those whose lives are in danger.

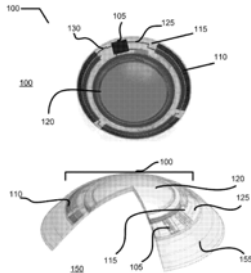
Dias & Cunha. Wearable Health Devices-Vital Sign Monitoring, Systems and Technologies. Sensors (Basel) 2018; 18:8: 2414.

12

## Patents: Samsung; Sony; J&J; Medella Health

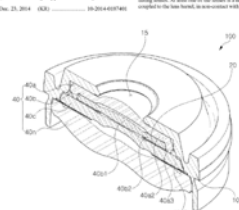
**United States**  
Patent Application Publication  
(44) Pub. No.: US 2014/0248014 A1  
(45) Pub. Date: Sep. 18, 2014

(54) **OPTICAL AND ELECTRIC DEVICE FOR MONITORING PHYSIOLOGICAL DATA**  
(71) Applicant: **AMERSON, A. ROBERTSON** (US)  
(72) Inventor: **Amerson, A. Robert**, Jacksonville, FL (US)  
(73) Assignee: **Amerson, A. Robert**, Jacksonville, FL (US)  
(57) **ABSTRACT**  
This invention describes a portable device with multiple sensors that can monitor physiological data in real-time. The device is designed to be worn on the wrist and can be used to monitor heart rate, blood pressure, and other physiological data. The device is designed to be used by a user and can be used to monitor physiological data in real-time.



**United States**  
Patent Application Publication  
(44) Pub. No.: US 2016/078074 A1  
(45) Pub. Date: Jun. 23, 2016

(54) **LENS ASSEMBLY AND OPTICAL DEVICE INCLUDING THE SAME**  
(71) Applicant: **Samsung Display Co., Ltd.**, Suwon (KR)  
(72) Inventor: **Kim, Min-Gook**, Suwon (KR)  
(73) Assignee: **SAMSUNG ELECTRONICS CO., LTD.**, Suwon (KR)  
(57) **ABSTRACT**  
A lens assembly including a lens and a lens holder. The lens holder includes a lens holder body and a lens holder cap. The lens holder body includes a lens holder body and a lens holder cap. The lens holder cap is configured to cover the lens holder body.



>250 patents since 2012

**United States**  
Patent Application Publication  
(44) Pub. No.: US 2017/0042480 A1  
(45) Pub. Date: Feb. 16, 2017

(54) **FUNCTIONAL CONTACT LENS AND RELATED SYSTEMS AND METHODS**  
(71) Applicant: **Medella Health Inc., Kitchener (CA)**  
(72) Inventors: **Harry Gandhi, Kitchener (CA); Hanyu Gan, Kitchener (CA); Haerig Bulg, Kitchener (CA); Ray Chen, Kitchener (CA)**  
(73) Assignee: **Medella Health Inc., Kitchener, ON (CA)**  
(57) **ABSTRACT**  
A functional contact lens and a lens holder. The functional contact lens includes a lens and a lens holder. The lens holder includes a lens holder body and a lens holder cap. The lens holder body includes a lens holder body and a lens holder cap. The lens holder cap is configured to cover the lens holder body.

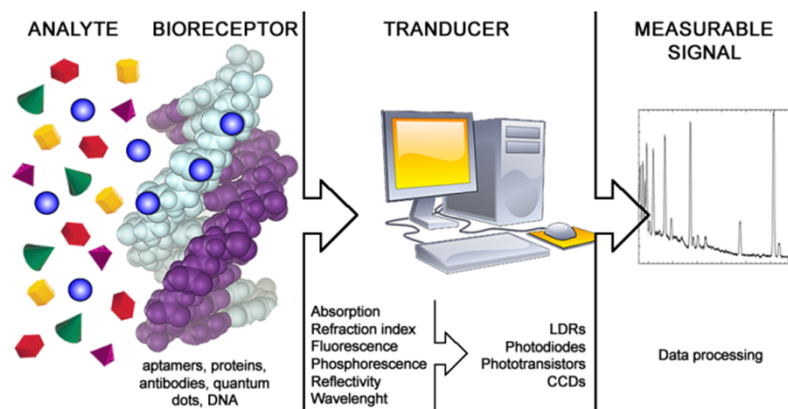
13

## “Smart” CL System

### Technology Overview



## Biosensing Contacts





## Systemic Disease Biomarkers

**Table 1**  
Systemic disease biomarkers found within the tear film.

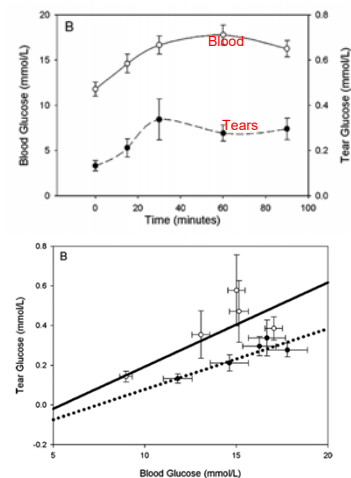
Disease	Potential tear biomarkers
Alzheimer's disease	Increased levels of dermcidin, lacritin, lipocalin-1 and lysozyme-C [2]
Cancer	Increased levels of lacryglobin [3,4], changes in combination of specific proteins [5]
Cystic fibrosis	IL-8 and IFN- $\gamma$ [6], MIP-1 $\alpha$ [7] and MIP-1 $\beta$ [8]
Diabetes	Increased levels of glucose [9], advanced glycation end products [10], cytokine changes [11]
Multiple sclerosis	Oligoclonal bands of IgG [12,13] and $\alpha$ -1-antichymotrypsin [14]
Parkinson's disease	TNF- $\alpha$ [15] and oligomeric alpha-synuclein [16]
Thyroid disease	IL-1 $\beta$ , IL-6, IL-17, TNF- $\alpha$ [17] and IL-7 [18]

IL – Interleukin; IFN – Interferon; MIP – Macrophage inflammatory protein; TNF – tumor necrosis factor; IgG – Immunoglobulin G.

16

## Blood vs Tear Glucose

- Poor correlation between blood and tears for glucose levels in **non-diabetics**
- Excellent correlation for **diabetics** ( $R^2 > 0.8$ )
- Unable to develop a viable CL-based detection product at that time due to technical limitations



17



# Glucose Biosensors for CL

**Table 2**  
Examples of glucose biosensors developed for contact lenses.

Mode of detection	Glucose sensor	Reader
Fluorescence [60]	Boronic acid, Concavalin A	External detector
Colourimetric [47]	Boronic acid	Colour chart
Fluorescence [61]	Boronic acid	Photodetector
Fluorescence, colourimetric [62]	Boronic acid, Concavalin A	External detector
Fluorescence, colourimetric [63]	Boronic acid, Concavalin A	Photodetector
Fluorescence [64]	Boronic acid, Concavalin A	External detector
Light emitted [65]	Boronic acid	Photodetector
Electrochemical [45]	Boronic acid	Electrode
Fluorescence, luminescence [66]	Boronic acid	External reader
Light emitted [31]	Boronic acid	Smart phone
Optical [33]	Boronic acid	External reader
Absorbance [50]	Concavalin A	Spectrophotometer
Fluorescence [49]	Concavalin A	Handheld photofluorometer
Fluorescence [32]	Concavalin A	Handheld photofluorometer
Electrochemical [46]	Glucose oxidase	Electrode
Electrochemical [29]	Glucose oxidase	Smart phone
Electrochemical [30]	Glucose oxidase	Handheld reader or smart phone
Electrochemical [67]	Glucose oxidase	External receiver
Electrochemical [38]	Glucose oxidase	On lens display
Electrochemical [68]	Metal oxides	External receiver

Google reveals smart contact lens prototype that tracks glucose for diabetics



Verily (Google Life Sciences): 2013 - 2018

After five years, Verily shelves project to create glucose-sensing contact lens



Jones et al.: CLEAR - Contact lens technologies of the future. Cont Lens Anterior Eye 2021; 44:2: 398-430.

18

# TF Substance P: Marker for Diabetic Peripheral Neuropathy?



Original Research

Tear film substance P: A potential biomarker for diabetic peripheral neuropathy

Shyam Sunder Tummanapalli<sup>1,2,\*</sup>, Mark D.P. Willcox<sup>3</sup>, Tushar Issar<sup>3</sup>, Ajmy Yan<sup>3</sup>, Jana Pisarcikova<sup>3</sup>, Natalie Kwai<sup>3</sup>, Ann M. Poynten<sup>3</sup>, Arun V. Krishnan<sup>3</sup>, Maria Markouli<sup>3</sup>

<sup>1</sup> School of Optometry & Vision Science, University of New South Wales, Australia

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<sup>3</sup> Department of Ophthalmology, Prince of Wales Hospital, Australia

## ARTICLE INFO

**Keywords:**  
Diabetic peripheral neuropathy  
Contact lens wear  
Tear composition  
Substance P  
Calcitonin gene-related peptide  
Type 1 diabetes  
Type 2 diabetes

## ABSTRACT

**Objective:** To explore the changes that occur in the concentrations of substance P (SP) and calcitonin gene-related peptide (CGRP) in tears as a result of corneal denervation and its association with diabetic peripheral neuropathy (DPN).

**Methods:** Forty-three individuals with type 1 diabetes, type 2 diabetes (T1D/T2D) and 34 age-matched healthy controls underwent a detailed assessment of neuropathy using the Total Neuropathy Score (TNS). The concentration of SP and CGRP in tears was measured by enzyme-linked immunosorbent assay. The corneal sub-basal nerve plexus was imaged using confocal microscopy. Corneal nerve fibre length, fibre density, branch density, total branch density, nerve fascial diameter and intra-corneal length were quantified.

**Results:** In T1D, the median [IQR] concentration of SP in tears was significantly reduced to those with DPN (130 [91-160] pg/mL) compared to both control subjects (TSD [165-153] pg/mL,  $P < 0.01$ ) and to those without DPN (91 [41-138] pg/mL,  $P = 0.01$ ). The concentration of CGRP was not changed. In T2D, there was no difference in concentrations between participants with diabetes and controls, regardless of neuropathy status. In T1D and T2D, corneal nerve parameters were significantly different between those with DPN or without DPN and controls. A significant correlation was noted between the concentration of tear film SP and TNS in T1D ( $r = -0.49$ ,  $P = 0.001$ ) and corneal nerve fibre density ( $r = 0.40$ ,  $P = 0.003$ ). The concentration of tear film CGRP was correlated significantly with the reduction of corneal nerve fibre density ( $r = 0.41$ ,  $P = 0.01$ ) in T1D.

## Conclusion

Tear film Substance P may provide a potential non-invasive biomarker for assessing neuropathy in Type 1 Diabetes



Tummanapalli et al. Tear film substance P: A potential biomarker for diabetic peripheral neuropathy. Ocul Surf 2019; 17:4: 690-698

19

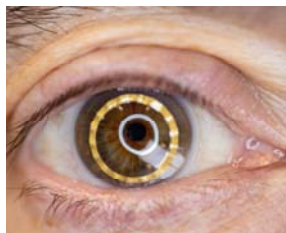
# Diagnosis & Screening for Ocular Disease

20

20

## CL Monitoring Devices: Glaucoma

- Sensimed Triggerfish®
- "Smart" contact lens with tiny embedded strain gauge to monitor curvature of the eye over a period of 24 hours
- Looks at relative changes in IOP
  - not absolute values
- Silicone-based (Dk~350)
- Thickness ~ 600µm
  - Dk/t ~ 60
  - hypoxia with overnight wear
- Single-use only



21

## CL Monitoring Devices: Other Opportunities

- Dry eye diagnosis and monitoring
  - osmolarity <sup>1</sup>
  - inflammatory cytokines <sup>2</sup>
    - TNF- $\alpha$ , IL-6, IL-17a and IL-8
- Blink monitoring <sup>3</sup>
- Ocular surface temperature <sup>4</sup>
- Ocular surface vasculature responses <sup>5</sup>

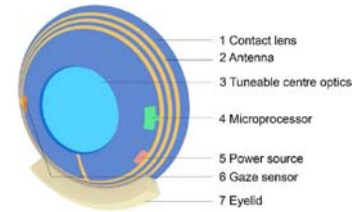


Fig. 3. Schematic design of an electronic presbyopic contact lens [425]. The sensor monitors (6) the gaze and sends the information to a microprocessor (4), which controls the tuneable centre optics (3). The optics can be tuned using a responsive polymer [435] or liquid crystals [424,425,436]. The entire system is supported by a power source (5) and an antenna (2).

## Treatment of Ocular Conditions

## Topical Drops

- >95% of current market for disease management
  - poor insertion technique in >50%
    - over-spill
  - poor compliance in 50%
  - rapid tear flow drainage
  - drug diluted by blinking
- Substantial systemic absorption
- <5% of drug gets to target

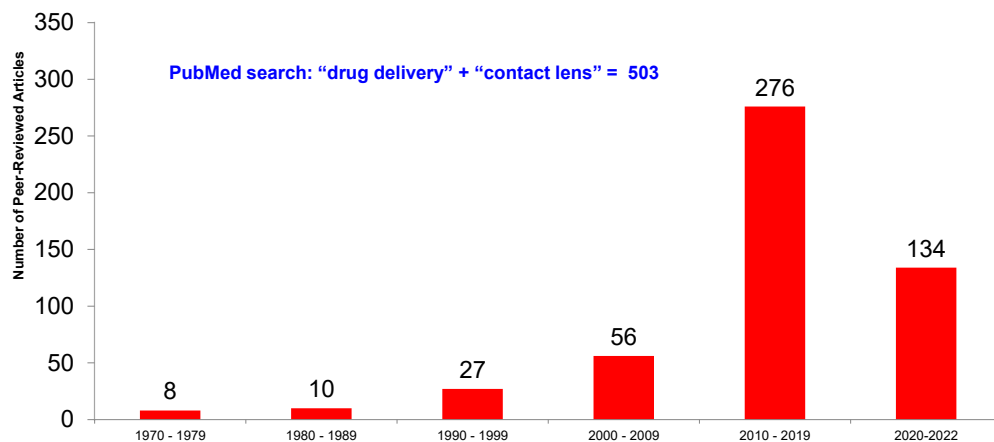


## What about using CL as a delivery vehicle?

- Soaked CL **should release drugs much slower**
- First suggested in original Wichterle hydrogel patents in 1960's <sup>1</sup>
  - first published manuscript in 1971 <sup>2</sup>
- 93% of surveyed clinicians would be interested in using a drug delivering CL <sup>3</sup>
- Clinical success depends on
  - drug loading
  - drug release

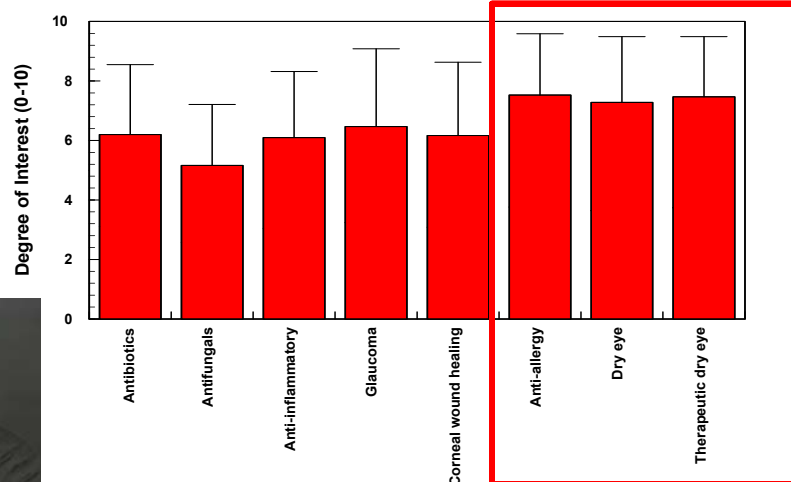


## Plenty of academic interest...



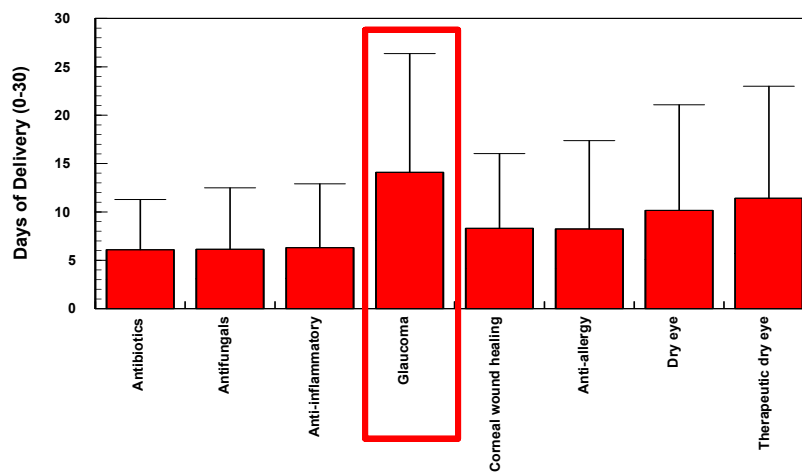
26

## Practitioner Interest?



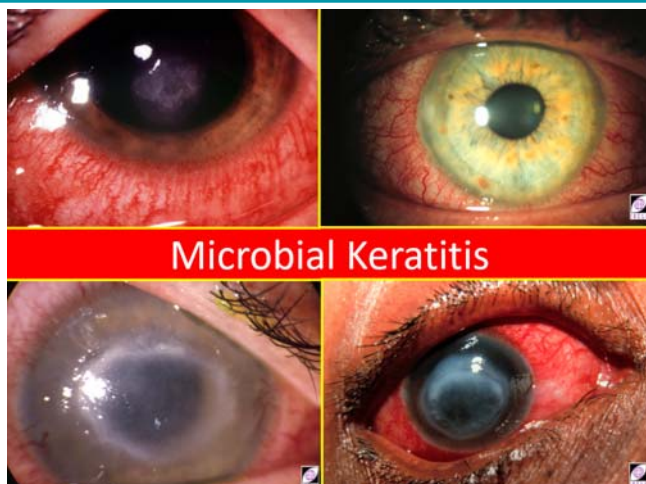
27

## Days of Continuous Delivery?



28

## Relevant diseases: short-term therapy



29

## Severe corneal abrasions



30

## Relevant diseases: long-term therapy

- Ocular allergy
- Glaucoma



31



# Why not just use existing commercial lenses?

32

32

## Interactions between topical drugs & CL materials: Many publications...

*Eye & Contact Lens* 20(2): 13-18, 2003

© 2003 Contact Lens Association of Ophthalmologists, Inc.

Available online at [www.sciencedirect.com](http://www.sciencedirect.com)

**ELSEVIER**

International Journal of Pharmaceutics 297 (2005) 169–175

**international journal of pharmaceutics**

**Ciprofloxacin Interaction with Silicon-Based and Conventional Hydrogel Contact Lenses**

C.C.S. Kallgari<sup>a</sup>, N.S. Wong<sup>a,b</sup>, L.W. Jones<sup>a</sup>, F.C. Ojima<sup>a</sup>, and C. Moresoli<sup>a,c</sup>, D. de Tsch

<sup>a</sup> Department of Chemical Engineering, University of Waterloo, 200 University Avenue, R. Waterloo, Ont. Canada N2L 3G1

<sup>b</sup> Department of Biomedical Engineering, University of Waterloo, 200 University Avenue, R. Waterloo, Ont. Canada N2L 3G1

<sup>c</sup> Contact Lens Research, Inc.

Received 11 September 2002; in final form 11 September 2002; accepted 11 September 2002

**ARTICLE**

**Uptake and Release of Dexamethasone Phosphate From Silicone Hydrogel and Group I, II, and IV Hydrogel Contact Lenses**

Adrienne Boone, Ph.D., Alex Hui, and Lyndon Jones, Ph.D., F.C.Optom.

**ORIGINAL ARTICLE**

**Delivery of Ketotifen Fumarate by Commercial Contact Lens Materials**

Anthony Schulz<sup>a</sup>, Alex Hui<sup>a</sup>, and Lyndon Jones<sup>a</sup>

**ABSTRACT**

Objectives: To investigate the uptake and delivery of the anti-allergy drug ketotifen fumarate (KF) by commercially available contact lenses.

**Methods:** A total of 18 different commercially available contact lenses were investigated, including five biconvex soft hydrogel lenses, five biconvex silicone hydrogel lenses, five biconvex rigid gas permeable lenses, and three biconvex rigid gas permeable lenses. The lenses were soaked in a 0.01% KF solution for 24 h, and the concentration of KF in solution over time was determined by ultraviolet spectrophotometry at 290 nm. After the 24 h soaking period, lenses were placed in fresh media containing freshly buffered saline, and the release of KF was monitored over time.

**Results:** The uptake and release of KF by the lenses was significantly different. The silicone hydrogel lenses showed the highest uptake and release of KF, followed by the rigid gas permeable lenses. The hydrogel lenses showed the lowest uptake and release of KF.

**Conclusions:** The uptake and release of KF by the lenses was significantly different. The silicone hydrogel lenses showed the highest uptake and release of KF, followed by the rigid gas permeable lenses. The hydrogel lenses showed the lowest uptake and release of KF.

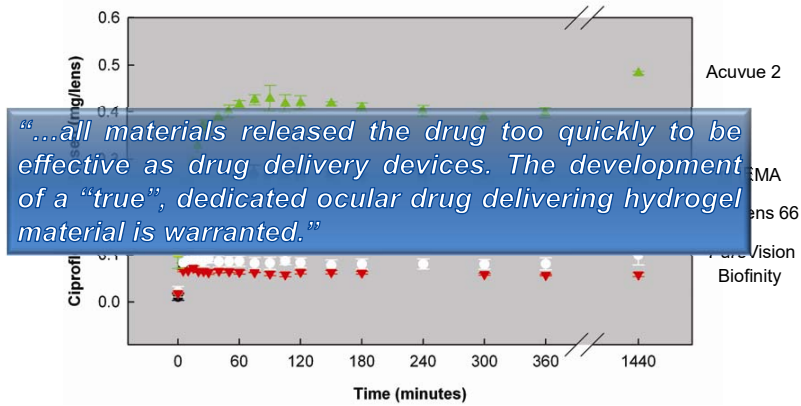
**KEYWORDS:** ciprofloxacin; silicone; hydrogel; contact lenses; drug delivery; ketotifen fumarate; dexamethasone phosphate; natamycin

**CORE**  
Centre for Ocular Research & Education

CORE – published >30 papers on this topic

33

## Release Rates



34

**What slow release technologies have been investigated?**

35

35

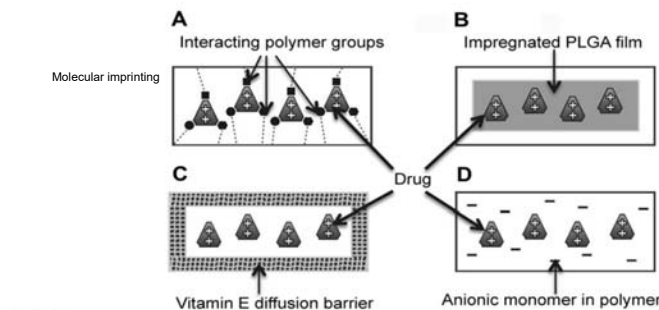
# CL as Drug Delivery Devices

- 144 review papers!
- 1982-2022
- 7 already this year



36

# CL Slow-Release Drug Delivering Concepts



**FIGURE 1.** Schematic representation of different strategies used to modify drug release from contact lens materials. Researchers have investigated (A) molecular imprinting to create noncovalent interactions between a drug and polymer groups, (B) incorporation of drug-impregnated PLGA films to serve as a drug reservoir within the center of the lens, (C) the addition of diffusion barriers to slow down the diffusion of drugs from the contact lens matrix, and (D) cationic/anionic monomer incorporation to generate an overall net charge within the material to interact with charged drug molecules, all in the hopes of modifying drug release kinetics and the amount of drug released.

37

## Nanoparticle Technologies

**Table 3**  
Examples of nanoparticle technologies for contact lens drug delivery.

Drug	Nanoparticle	Synthesis method	Loading method	Average size (nm)	Release Duration
Ciprofloxacin [249]	Pullulan-PCL micelles	Dropwise addition of water to DMSO	Dispersion in pre-polymer solution and soaking	142 ± 12	3–4 days
Cyclosporine [250]	Brij surfactants micelles	Dissolution in water	Dispersion in pre-polymer solution	<40	>15 days
Cyclosporine [243]	C-HA micelles	Dissolution in water and DMSO	Dispersion in pre-polymer solution	300	12 days
Ketotifen [242]	silica shell	Microemulsion	Dispersed in pre-polymer solution	104.2–126.54	10 days
Lidocaine [221]	DMPC liposomes	Microemulsion	Dispersed in pre-polymer solution	20	8 days
Loteprednol etabonate [251]	PCL/HEMA/PEG-DA	Surfactant-free mini-emulsion polymerisation	Dispersed in pre-polymer solution	52.3–83.4	12 days
Natamycin [252]	Dex- $\beta$ -PLA micelles	Nanoprecipitation (DMSO to water)	Soaking	26.1–26.6	12–24 h
Prednisolone [253]	PLGA	Emulsion-solvent evaporation	Dispersed in pre-polymer solution	294.5 ± 1.8	24 h
Timolol [254]	PVP-PNIPAAm	Electrohydro-dynamic atomisation	Dissolved in polymeric solution	52 % of nano-structures < 200	24 h
Timolol [241]	EC	Double emulsion	Dispersed in pre-polymer solution	261–340	160 h

C-HA, cholesterol-hyaluronic acid; DA, diacrylate; EC, ethyl cellulose; Dex, Dextran; DMPC, dimyristoylphosphatidylcholine; DMSO, dimethylsulfoxide; HEMA, poly (2-hydroxyethyl methacrylate); PEG, polyethylene glycol; PCL, polycaprolactone; PLA, polylactic acid; PLGA, poly (lactic-co-glycolic acid); PNIPAAm, poly (N-isopropylacrylamide); PVP, poly(vinylpyrrolidone).

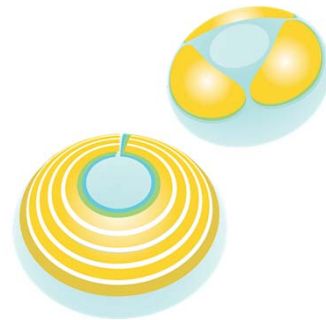


Jones et al.: CLEAR - Contact lens technologies of the future. Cont Lens Anterior Eye 2021; 44;2: 398-430.

38

## Drug “Printing” Concept

- MediPrint™ Ophthalmics
- A proprietary process allows for printing of drug and barrier layers on the contact lens surface to control drug diffusion release kinetics
- Early phase study on LL-BMT1
  - drug-eluting CL for glaucoma treatment using bimatoprost



39

## In vitro vs In vivo?

Journal of Controlled Release 343 (2022) 672–702

Contents lists available at ScienceDirect

Journal of Controlled Release

journal homepage: [www.elsevier.com/locate/jconrel](http://www.elsevier.com/locate/jconrel)

Review article

Testing drug release from medicated contact lenses: The missing link to predict *in vivo* performance

Ana F. Pereira-da-Mota<sup>a</sup>, Chau-Minh Phan<sup>b,c</sup>, Angel Concheiro<sup>a</sup>, Lyndon Jones<sup>b,c</sup>, Carmen Alvarez Lorenzo<sup>b,c</sup>

<sup>a</sup> Departamento de Farmacología, Farmacia y Tecnología Farmacéutica, I+D+i Farmacia Group (G1-1643), Facultad de Farmacia, Instituto de Materiales (IMATLUS) and Health Research Institute of Santiago de Compostela (ISCI), Universidade de Santiago de Compostela, 15702 Santiago de Compostela, Spain

<sup>b</sup> Centre for Ocular Research & Education (CORE), School of Optometry and Vision Science, University of Waterloo, Waterloo, ON, Canada

<sup>c</sup> Centre for Eye and Vision Research (CEVR), 179c, Hong Kong Science Park, Hong Kong

**ARTICLE INFO**

**Keywords:**  
Drug-eluting contact lens  
in vitro release tests  
in vivo release  
Release rate specification  
Therapeutic response  
in vitro-in vivo correlations

**ABSTRACT**

Contact lenses (CLs) offer a wide variety of advantages as ocular drug-releasing platforms, but the feasibility of medicated CL development is constrained by numerous scientific, technological, and regulatory challenges. One main difficulty is the setting of release rate specifications for each drug, since at present there are no standardized *in vitro* release models that can appropriately predict the performance of drug-eluting CLs once placed onto the eye. CL-adapted release tests may provide knowledge on how the drug release pattern should perform *in vivo* to trigger and maintain the therapeutic effects for both anterior and posterior ocular issues. Moreover, *in vitro* release tests are valuable tools for quality assessment during production and to investigate the effect of a change in composition or process variables. This review aims to shed light on bioequivalent ways of evaluating *in vitro* drug

**CORE**  
Centre for Ocular Research & Education

Pereira-da-Mota et al.: Testing drug release from medicated contact lenses: The missing link to predict *in vivo* performance. J Control Release 2022; 343 672-702.

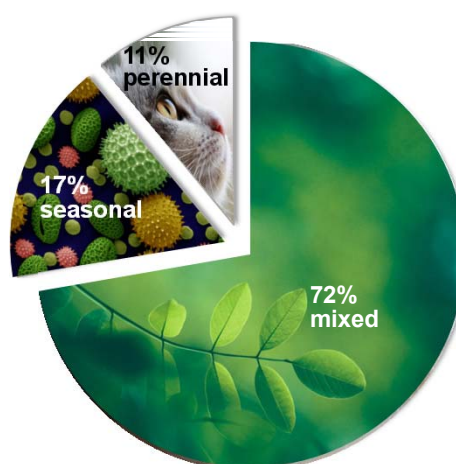
40

## Allergic Rhino-Conjunctivitis (Hay Fever)

### Classification by Sensitization\*1

- **Seasonal 17% (airborne pollens)**
  - Symptoms more irritative, more conjunctivitis (64.9%)
  - Peak suffering March - September
- **Perennial 11% (mites, pets)**
  - Symptoms more obstructive
  - Peak suffering October - February
- **Mixed 72% (indoor & pollen)**
  - Both irritative & obstructive symptoms
  - Suffering year round

12% overall but rising year over year



\*19,475 consecutive subjects

1. Ciprandi G, Cirillo L, Vizzaccaro A et al. Season and perennial allergic rhinitis: is this classification adherent to real life? Allergy 2005; 60:882-887.

41

## Contact Lens Patients with Allergies



ocular allergy patients are contact lens wearers<sup>1</sup>



of allergy days result in symptoms that might require drops<sup>1</sup>



contact lens wearers switch to glasses during allergy season<sup>1</sup>

**Disruptive and Frustrating!**

1. JJV Data on File 2019. JJV US & Canada Ocular Allergy Key Insights Review, March 2019

42

## Licensed in Japan & Canada May 2021; USA Mar 2022

- J&J Acuvue Theravision with Ketotifen
  - topical antihistamine
- “Itchy eye relief”
  - allergic conjunctivitis
- Daily disposable platform
  - etafilcon A
  - hydrogel; 58% water content
  - preservative free
  - releases product over 5 hrs
  - provides up to 12 hr relief



43

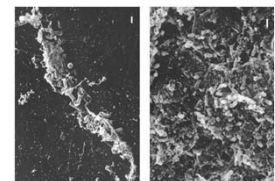
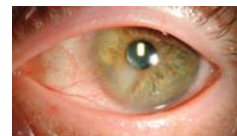
# Antimicrobial Lenses

44

44

## Antimicrobial lenses

- Microbial adhesion to CL is a risk factor for developing MK, CLARE & CLPU <sup>1</sup>
  - reduction in bacterial adhesion to CL using antimicrobial coatings/treatments could thus be a viable means of reducing these potentially sight threatening complications
- Technologies investigated include
  - silver <sup>2</sup>
  - antimicrobial peptides melimine, Mel4 and Esculentin-1a <sup>3</sup>
  - fimbrolides (furanones) <sup>4</sup>



45



# Storage Cases

46

46

## CL Storage Cases

- Implicated in MK
  - can be contaminated in as little as 7 days <sup>1</sup>
  - 80% contamination possible in 14 days <sup>2</sup>
  - mature biofilms by 30 days <sup>1</sup>
- Overcome by
  - in-built reminders to replace <sup>3</sup>
  - controlled obsolescence <sup>4</sup>
  - silver or selenium containing cases <sup>5,6</sup>
  - in-built colorimetric biofilm sensors <sup>7,8</sup>



Fig. 4. Microbiosensor in a contact lens case with the bottom blue colour indicating microbial contamination [525].

47

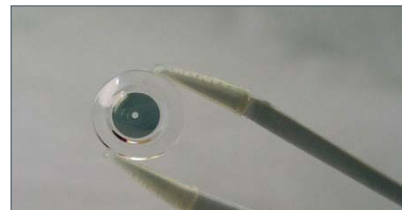
# Optical Enhancements

48

48

## Visual Augmentation

- Six Million Dollar Man (1974-78)
  - “bionic” eye
    - 20x zoom lens
    - camera
    - IR filter (night vision & heat detection)
- Contact lens is a key component of any such modern-day concept
  - permits focus on content close to the eye
  - or carries the information display



49

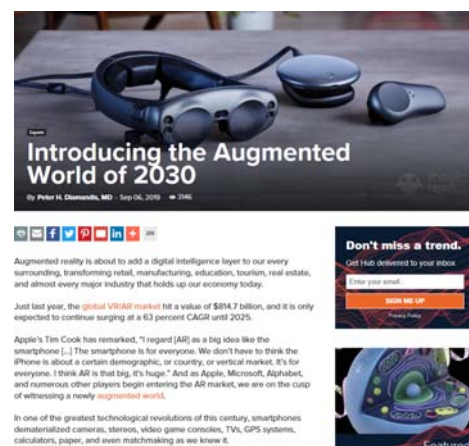
## Mission Impossible: Ghost Protocol (2011)



50

## Augmented & Virtual Reality

- VR - completely immerses users in digital realities
- AR - allows users to remain engaged with their physical surroundings, serving as a visual enhancement rather than replacement
- In 2018 the global VR/AR market hit a value of \$814.7 billion
  - expected to continue surging at a **63% annual growth** until 2025



51

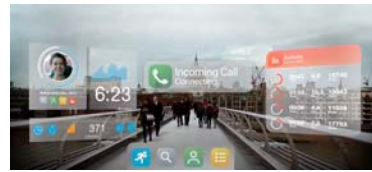
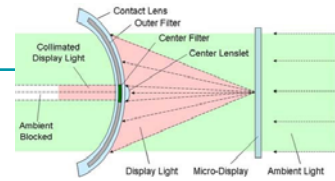
## AR & VR “Players”

- Well-known players
  - Microsoft, Apple, Alphabet, Intel have made tremendous strides
  - Samsung
    - recently granted US patent to develop smart lenses capable of streaming text, capturing videos, and even beaming images directly into a wearer’s eyes
    - contacts are designed to include a motion sensor (for eye movement tracking), hidden camera, and display unit
- Start-ups remain competitive (>1800)
  - Magic Leap - raised a total of \$2.6 billion since its founding in 2010
    - head-mounted virtual retinal display, Magic Leap projects a digital light field into users’ eyes to superimpose 3D computer-generated imagery over set environments
  - Mojo Vision - raised ~\$150 million in its efforts to develop and produce an AR contact lens

## Approaches: Electronic and Optical

## Innovega

- No electronics in the lens
- Uses a two-state linear polarizer to separate a display path and a non display normal vision path
- Filters coupled with a very high-power 1mm center add (+65 to +85 D) allows the ability to see a display in the spectacle plane and see the real world with normal contact lens refractive correction
- Display eyewear may then be directly viewed with no additional optics and may have transparent displays for AR or occluded displays for VR

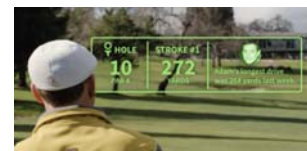
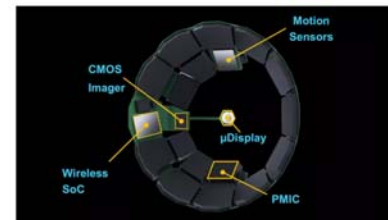


<http://innovega-inc.com>

54

## Mojo Vision

- Scleral lens containing electronics
- The Mojo Lens centerpiece is a hexagonal display less than half a millimeter wide
  - a "femtoprojector" expands the imagery optically and beams it to a central patch of the retina
- Mojo is developing its lenses with Japanese contact lens maker Menicon



<https://www.mojo.vision/>

55

## Contact Lens “Zooming” Under Ocular Control

News

### Robotic lens zooms-in with blink of an eye

Movements of soft robotic lens are controlled by the eyes

September 6, 2019  
by Liozel Labios, UC  
San Diego

#### Topics

Electronics

#### Tags

contact lens

A research team led by the University of California San Diego has developed a soft robotic lens whose movements are controlled by the eyes — blink twice and the lens zooms in and out; look left, right, up or down and the lens will follow.

The lens is the first example of an interface between humans and soft machines. “The human-machine interface, as we know it, features classical machines: computers, wheelchairs, and rigid robotics, for example. The innovation here is the interface with soft robotics. This can really open up new opportunities in the field,” said Shengqiang Cai, a professor of mechanical and aerospace engineering at UC San Diego who led the research.

Potential applications of the system include visual prostheses, adjustable glasses, VR, and soft robots that can see.

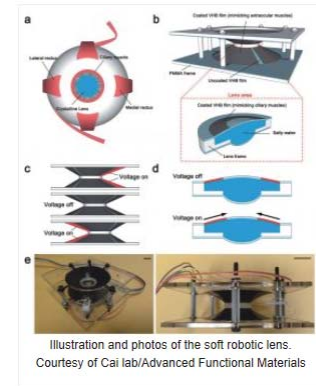


Illustration and photos of the soft robotic lens.  
Courtesy of Cai lab/Advanced Functional Materials



Li et al. A Biomimetic Soft Lens Controlled by Electrooculographic Signal. *Advanced Functional Materials* 2019; 29: 1903762.

56

## Augmented Reality for Education



57







# Summary

61

61

## Summary

- Incredible diversity of new technologies under development that will shape the future for CL
- Rapid growth in novel biomaterials and the development of powered CL through advancements in nanotechnology will enable the commercialisation of lenses that can both detect and treat ocular and systemic disease
- Novel optical designs will provide enhanced vision for patients with low vision and a wide variety of other optical considerations
  - head-up displays etc
- Contact lenses have been around for over 100 years - and their future remains bright



62

# THANK YOU