



CONTACT LENSES

*Materials, Chemicals,
Methods and Applications*



Johannes Karl Fink

 Scrivener
Publishing

WILEY

Contact Lenses

Scrivener Publishing

100 Cummings Center, Suite 541J
Beverly, MA 01915-6106

Publishers at Scrivener

Martin Scrivener (martin@scrivenerpublishing.com)
Phillip Carmical (pcarmical@scrivenerpublishing.com)

Contact Lenses

**Materials, Chemicals,
Methods and Applications**

Johannes Karl Fink
Montanuniversität Leoben, Austria



WILEY

This edition first published 2022 by John Wiley & Sons, Inc., 111 River Street, Hoboken, NJ 07030, USA and Scrivener Publishing LLC, 100 Cummings Center, Suite 541J, Beverly, MA 01915, USA

© 2022 Scrivener Publishing LLC

For more information about Scrivener publications please visit www.scrivenerpublishing.com.

All rights reserved. No part of this publication may be reproduced, stored in a retrieval system, or transmitted, in any form or by any means, electronic, mechanical, photocopying, recording, or otherwise, except as permitted by law. Advice on how to obtain permission to reuse material from this title is available at <http://www.wiley.com/go/permissions>.

Wiley Global Headquarters

111 River Street, Hoboken, NJ 07030, USA

For details of our global editorial offices, customer services, and more information about Wiley products visit us at www.wiley.com.

Limit of Liability/Disclaimer of Warranty

While the publisher and authors have used their best efforts in preparing this work, they make no representations or warranties with respect to the accuracy or completeness of the contents of this work and specifically disclaim all warranties, including without limitation any implied warranties of merchantability or fitness for a particular purpose. No warranty may be created or extended by sales representatives, written sales materials, or promotional statements for this work. The fact that an organization, website, or product is referred to in this work as a citation and/or potential source of further information does not mean that the publisher and authors endorse the information or services the organization, website, or product may provide or recommendations it may make. This work is sold with the understanding that the publisher is not engaged in rendering professional services. The advice and strategies contained herein may not be suitable for your situation. You should consult with a specialist where appropriate. Neither the publisher nor authors shall be liable for any loss of profit or any other commercial damages, including but not limited to special, incidental, consequential, or other damages. Further, readers should be aware that websites listed in this work may have changed or disappeared between when this work was written and when it is read.

Library of Congress Cataloging-in-Publication Data

ISBN 978-1-119-85735-8

Cover image: Pixabay.com

Cover design by Russell Richardson

Set in size of 11pt and Minion Pro by Manila Typesetting Company, Makati, Philippines

Printed in the USA

10 9 8 7 6 5 4 3 2 1

Contents

Preface	xi
1 Types of Lenses	1
1.1 History of Contact Lenses	1
1.2 Materials	3
1.3 Monomers	3
1.3.1 Monomers for Block Copolymers	3
1.3.2 Silicone Acrylamides	7
1.4 Soft Lenses	13
1.4.1 Hydrogels	13
1.4.2 PVA Hydrogel	48
1.4.3 Clear Contact Lenses	48
1.5 Water Absorbable Formulations	49
1.6 Bandage Contact Lenses	53
1.6.1 Antimicrobial Bandage Contact Lens	53
1.7 Functional Contact Lenses	56
1.7.1 Remote Health Monitoring	56
1.7.2 Graphene Oxide Nanocolloids	60
1.7.3 Diabetic Diagnosis	61
1.7.4 Target Analyte Sensing	66
1.7.5 Adaptive Tuning	71
1.7.6 Wireless Communication	72
1.7.7 Glucose Biosensors	76
1.7.8 Cancer Detection	78
1.8 Scleral Contact Lenses	78
1.8.1 Fabrication of Scleral Lenses	79
1.8.2 Scleral Lens Fitting	82
1.8.3 Ocular Drug Delivery Systems	83
1.9 Multifocal Contact Lenses	83
1.9.1 Bifocal Contact Lenses	83
1.9.2 Silicone Hydrogels	85

1.9.3	Non-Silicone Hydrogels	89
1.9.4	Tilted-Wear Type Contact Lenses	93
1.9.5	Neutral Density Filters	94
1.10	Augmented Reality Contact Lens Systems	95
1.10.1	Electronic Contact Lenses	96
1.10.2	Smart Contact Lenses	96
1.10.3	Wearable Smart Contact Lenses	97
1.10.4	Collimated Light-Emitting Diodes	98
1.11	Siloxane Macromers	99
1.11.1	Silicone Urethane Polymers	102
1.12	Oxygen-Permeable Lenses	107
1.12.1	Extended Wear Lenses	107
1.12.2	Structures for Thick Payloads	115
1.13	Natural Protein Polymer Contact Lenses	118
1.14	Ultrathin Coating	119
1.15	Anti-Biofouling Contact Lenses	121
1.15.1	Phosphorylcholine	121
1.15.2	2-Hydroxyethyl methacrylate	125
1.15.3	Chitosan	127
1.16	Drug Delivery via Hydrogel Contact Lenses	129
1.16.1	Hydrogels with Phosphate Groups	129
1.16.2	Ophthalmic Drug Delivery	131
1.17	Simulation Methods	133
1.17.1	Ocular Topography Parameters	133
1.17.2	Rigid Gas-Permeable Lenses	134
1.17.3	Computerized Videokeratography	134
	References	135
2	Fabrication Methods	149
2.1	Computer-Aided Contact Lens Design and Fabrication	149
2.1.1	Spline-Based Mathematical Surfaces	149
2.1.2	Corneal Refractive Therapy Program	152
2.2	Contact Lenses with Selective Spectral Blocking	154
2.3	Colored Contact Lenses	156
2.3.1	Hard Colored Contact Lenses	157
2.4	Decentered Contact Lenses	161
2.5	Stabilized Contact Lenses	162
2.6	Additive Manufacturing	163
2.7	Mold Process	164
2.7.1	Injection Molding	164
2.7.2	Cast Molding	166

2.7.3	Two-Part Mold Assembly	168
2.8	Reactive Ion Etching	170
2.9	Electrospinning	172
2.9.1	Creating Electrospun Contact Lens Structures	172
2.9.2	Electrospinning Controlled Polymer Fibril Matrices	174
2.9.3	Electrospinning of a Prepolymer Solution	175
2.10	Rigid Plastic Lenses	183
2.10.1	Rigid Gas-Permeable Contact Lenses	183
2.11	Soft Plastic Lenses	184
2.11.1	Layer-by-Layer Deposition	184
2.11.2	Electron-Beam Irradiation Polymerization	191
2.11.3	Shaping and Cutting	192
2.12	Coating Methods	195
2.12.1	Zwitterionic Coating	195
2.12.2	Antibacterial Nanocoating	196
2.13	Disinfection of Contact Lenses	196
2.13.1	Hydrogen Peroxide and Fibrous Catalyst	197
2.13.2	Hydrogen Peroxide and Metal Catalyst	197
2.13.3	Removing Hydrogen Peroxide	199
2.14	Integrated Microtubes	201
2.15	Injection Molding	201
2.15.1	Aspheric Contact Lenses	201
2.16	Handling Tools	202
2.16.1	Insertion Tool	202
2.16.2	Insertion Tool	205
	References	205
3	Properties	211
3.1	Ophthalmic Compatibility Requirements	211
3.2	Standards	212
3.2.1	Tensile Properties of Plastics	212
3.2.2	Tear-Propagation Resistance	215
3.2.3	Oxygen Gas Transmission Rate	215
3.2.4	Biomaterials	215
3.2.5	Eye Protectors	216
3.3	Eye Model with Blink Mechanism	217
3.4	Assessment of Cytotoxic Effects	219
3.4.1	Draize Eye Irritation Test	219
3.4.2	Acute Eye Irritation Testing	220
3.4.3	Benzalkonium Chlorides	220
3.4.4	Residual Monomer Content	221

3.5	Special Functions	223
3.5.1	Intraocular Pressure	224
3.5.2	Coating Thickness	229
3.6	Cleaning of Contact Lenses	229
3.7	Biofouling	234
3.8	Wettability	234
3.8.1	Blister Pack Solutions	236
3.8.2	Captive Bubble Method	237
3.8.3	Tethered Hyaluronic Acid-Based Coatings	239
3.9	Material Properties and Antimicrobial Efficacy	240
3.10	Microscopic Examination	242
3.10.1	X-Ray Photoelectron Spectroscopy	243
3.10.2	Atomic Force Spectroscopy	244
3.10.3	Electrochemical Impedance Spectroscopy	246
3.10.4	Scanning Electron Microscopy	247
3.11	Schirmer Tear Test	248
3.12	Ocular Surface Disease Index Test	248
3.13	Corneal Fluorescein Staining Test	249
3.14	Ion Permeability	250
3.14.1	Ionoflux Technique	250
3.14.2	Ionoton Measurement Technique	252
3.15	Hydrodell Water Permeability Technique	253
3.16	Oxygen Permeability and Transmissibility	253
3.16.1	Contact Lens Solutions	254
3.17	Optical Biometer	255
3.17.1	Ophthalmologic Apparatus	255
3.17.2	Ophthalmologic Information Processing	259
3.17.3	Swept-Source Optical Coherence Tomography	259
	References	260
4	Drug Delivery	271
4.1	Basic Issues	272
4.2	Methodologies for the Design of Therapeutic Contact Lenses	273
4.2.1	Soaking Method	273
4.2.2	pH-Sensitive Lenses	273
4.2.3	Magnetic Micropump	275
4.2.4	Molecular Imprinting	275
4.2.5	Colloidal Nanoparticles	276
4.2.6	Polymeric Nanoparticles	276
4.2.7	Cyclodextrins	277

4.2.8	Liposomes	277
4.2.9	Microemulsion and Micelles	278
4.2.10	Vitamin E	278
4.2.11	Supercritical Fluid Technology	278
4.2.12	Hydrophobic Drug Loading	279
4.2.13	Cationic Drugs	279
4.3	Hydrogels	281
4.3.1	Salt-Induced Modulation	283
4.3.2	Polymeric Hydrogels	284
4.3.3	Colloid-Laden Hydrogels	285
4.3.4	Ligand-Containing Hydrogels	285
4.3.5	Amphiphilic Polymers	286
4.3.6	Silicone Hydrogel Contact Lenses	289
4.3.7	Zwitterionic Hydrogels	290
4.3.8	Surface-Modified Hydrogels	291
4.3.9	Cyclodextrin-Hyaluronan Hydrogels	293
4.3.10	Bioinspired Hydrogels	293
4.3.11	Tobramycin Release	294
4.4	Contact Lens Gels	297
4.5	Molecularly Imprinted Contact Lenses	298
4.5.1	Molecular Imprinting Technology	298
4.5.2	Molecularly Imprinted Contact Lenses	299
4.5.3	Hydrogels	301
4.5.4	Supercritical Fluid-Assisted Preparation	302
4.6	Special Drugs	303
4.6.1	Timolol	303
4.6.2	Dexamethasone	308
4.6.3	Ketotifen Fumarate	312
4.6.4	Ciprofloxacin	315
4.6.5	Ofloxacin	318
4.6.6	Polymyxin B and Vancomycin	321
4.6.7	Epinastine	323
4.6.8	Lactoferrin	323
4.6.9	Bimatoprost	324
4.6.10	Dipicolylamine	325
4.6.11	Gatifloxacin	326
4.6.12	Hydroxypropyl Methylcellulose	328
4.6.13	Dorzolamide	328
4.6.14	Ethoxzolamide	329
4.6.15	Hyaluronic Acid	331
4.6.16	Lifitegrast	335

x CONTENTS

4.6.17	Diclofenac Sodium	336
4.6.18	Moxifloxacin	339
4.6.19	Norfloxacin	340
4.6.20	Sparfloxacin	341
4.6.21	Latanoprost	342
4.6.22	Loteprednol	343
4.6.23	Release of Multiple Therapeutics	344
	References	347
5	Medical Problems	363
5.1	Eye Diseases	363
5.2	Corneal Edema	363
5.2.1	PMMA Lenses	365
5.2.2	Thickness Changes	365
5.2.3	Corneal Swelling	366
5.2.4	Acanthamoeba Keratitis	367
5.3	Presbyopia and Myopia Control	368
5.4	Toxic Soft Lenses	369
5.4.1	Allergic and Toxic Reactions	370
5.5	Disinfection Agents	374
5.5.1	Polymeric Biguanide and Vinylimidazole	375
5.5.2	Saccharides	376
5.5.3	Amphipathic Peptides	381
5.5.4	Antibacterial Properties	384
5.6	Silicone Hydrogels	385
5.7	Limbal Stem Cell Deficiency	385
5.8	Computer Vision Syndrome	387
5.8.1	Tests and Analysis	388
5.8.2	Pathophysiology	388
5.8.3	Problems for Radiologists	389
5.9	Dry Eye Problems	390
5.9.1	Ions in Tears	390
5.9.2	Treatment Methods	392
5.9.3	Comparative Study of the Reasons for Dry Eyes	394
5.10	Orthokeratology	395
5.10.1	Myopia	398
	References	401
Index		413
	Acronyms	413
	Chemicals	416
	General Index	429

Preface

This book focuses on the chemistry and properties of contact lenses and their fabrication methods.

The text starts with a chapter in which a detailed history of contact lenses spanning over almost 500 years is presented

Next, common materials that are used for the fabrication of contact lenses are listed and explained, including both the monomers and polymers that are used in their production. Special issues regarding soft lenses, clear contact lenses, and functional contact lenses are also discussed.

Functional contact lenses can be used for remote health monitoring and ocular drug delivery systems. Besides the materials used here, these issues are detailed in further separate chapters. Also, special fabrication methods are discussed, e.g., the fabrication of multifocal contact lenses and the fabrication of ultrathin coatings.

There is also an important discussion on additives that can be used, e.g., for oxygen-permeable materials or anti-biofouling materials.

The chapter ends with a discussion of simulation methods for contact lenses, such as ocular topography parameters, gas-permeable lenses, and computerized videokeratography.

In the second chapter, several common fabrication methods for contact lenses are discussed. Here, computer-aided contact lens design, methods for the fabrication of colored contact lenses, and the fabrication of decentered contact lenses are detailed.

Also, special processes are reviewed, including mold processes, reactive ion etching, electrospinning and others.

Another chapter discusses the properties of contact lenses and methods of measurement. Here, a lot of standard methods are discussed. Besides standard methods, other issues are discussed such as the assessment of cytotoxic effects, the Schirmer tear test and others.

A chapter is devoted to drug delivery of contact lenses, a comparatively new issue.

Finally, a chapter details the possible medical problems related to contact lenses and how to avoid them. These are eye diseases, allergic and toxic reactions. Also, disinfection agents that can be used and methods for the medical treatment of such problems are detailed.

The text focuses on the literature of the past decade. Beyond education, this book will serve the needs of industry engineers and specialists who have only a passing knowledge of the plastics and composites industries but need to know more.

How to Use This Book

Utmost care has been taken to present reliable data. Because of the vast variety of material presented here, however, the text cannot be complete in all aspects, and it is recommended that the reader study the original literature for more complete information.

The reader should be aware that mostly US patents have been cited where available, but not the corresponding equivalent patents in other countries. For this reason, the author cannot assume responsibility for the completeness, validity or consequences of the use of the material presented herein. Every attempt has been made to identify trademarks; however, there were some that the author was unable to locate.

Index

There are three indices: an index of acronyms, an index of chemicals, and a general index.

In the index of chemicals, compounds that occur extensively, e.g., “acetone,” are not included at every occurrence, but rather when they appear in an important context.

Acknowledgements

I am indebted to our university librarians, Dr. Christian Hasenhüttl, Margit Keshmiri, Friedrich Scheer, Christian Slamenik, Renate Tschabuschnig, and Elisabeth Groß for their support in literature acquisition. I also want to express my gratitude to all the scientists who have carefully published their results concerning the topics dealt with herein. This book could not have been otherwise compiled.

In addition, I am very grateful to the ophthalmologists Dr. Anna Schlanitz-Bolldorf and Dr. Ferdinand Schlanitz, who inspired me to write

this text. Last, but not least, I want to thank the publisher, Martin Scrivener, for his abiding interest and help in the preparation of the text. In addition, my thanks go to Jean Markovic, who made the final copyedit with utmost care.

Johannes Fink
Leoben, December 2, 2021

1

Types of Lenses

1.1 History of Contact Lenses

A history of contact lenses spanning over almost 500 years has been detailed (1, 2). It is based on historical works, scientific papers and journal articles and looks at both the modern disposable lens as well as the hard and soft lenses that came before. Some important events are collected in Table 1.1.

Table 1.1 History of contact lenses (2).

Year	Inventor	Issue
1508	Leonardo da Vinci	Corneal neutralization
1637	René Descartes	Fluid-filled tube
1685	Philippe de La Hire	Neutralization of cornea
1801	Thomas Young	Three color theory of perception
1827	George Biddell	Theory of astigmatism
1845	Sir John F. W. Herschel	Convex lenses
1846	Carl Zeiss	Optical instruments
1851	Johann Nepomuk Czermak	Water-filled goggle
1887	Adolf Eugen Fick	First successful contact lens
1961	Otto Wichterle	Soft contact lenses
1979	Kyoichi Tanaka	Silicone hydrogel materials

In 1508, Leonardo da Vinci first had the idea of placing a corrective lens directly onto the surface of the eye (3–5). In 1637, René Descartes proposed another idea in which a glass tube filled with liquid is placed in direct contact with the cornea.

In 1887, Adolf Eugen Fick, a German physiologist, created the first successful contact lens (6). Glass-blown scleral lenses remained the only form of contact lens until 1938, when poly(methyl methacrylate) (PMMA) was developed, and Mullen and Obring used the plastic to manufacture scleral lenses. Obring developed the Plexiglass series in New York in 1940 (4).

In 1961, the Czech chemist Otto Wichterle invented soft contact lenses (7, 8). In 1970, rigid gas-permeable contact lenses were developed, and widely accepted for the advantages of small diameter (about 9 *mm*) and gas permeability. Silicone hydrogel materials were developed in 1979 (9). In 1999, an important development was the launch of the first silicone hydrogels onto the market. These new materials showed an extremely high oxygen permeability with comfort performance (4).

The factors that influenced the development of special materials have been reported (10). Accounts of early attempts to improve vision by use of a lens contacting the eye are limited to a few isolated observations (11). Practical success was not realized until techniques for fabrication of lenses from glass were sufficiently developed (12). PMMA replaced glass in the late 1930s. This material is more durable, more readily fabricated and was claimed by some authors to show a better ocular compatibility (13). During the same broad period of time, there was also a change in emphasis from scleral to corneal contact lenses, which placed different demands on material design and development.

More is demanded from ophthalmic treatments using contact lenses, which are currently used by over 125 million people around the world (14). Improving the material of contact lenses is currently a rapidly evolving discipline (10).

A search has been performed of the titles of papers in the Scopus database to identify contact lens-related articles published this century (15). The ten most highly cited papers were determined from the total list of 4,164 papers found. Rank-order lists by count were assembled for the *top 25* in each of four categories: authors, institutions, countries and journals. A 20-year subject-specific contact lens h-index was derived for each author, institution, country and journal to serve as a measure of impact in the field. The top 10 constituents (of the top 25) of each category were ranked and tabulated (15).

1.2 Materials

Contact lens materials (10) are typically based on polymer- or silicone-hydrogel, with additional manufacturing technologies employed to produce the final lens. These processes are simply not enough to meet the increasing demands for contact lenses and the ever-increasing number of contact lens users (14).

An advanced perspective on contact lens materials has been presented, with an emphasis on materials science employed in developing new contact lenses (14, 16). The future trends for contact lens materials are to graft, incapsulate, or modify the classic contact lens material structure to provide new or improved functionality. Also, some of the fundamental material properties are discussed, and the outlook for related emerging biomaterials is presented.

Contact lens materials and lens types, treatment for contact lens and tear film complications, and myopia correction and contact lenses for abnormal ocular conditions have been detailed (17). Current topics in this field are miniscleral lenses, keratoconus, corneal crosslinking, and pediatric, cosmetic and prosthetic contact lenses. Furthermore, simulation programs for scleral lens fitting, sagittal values, soft toric mislocation, front vertex power, orthokeratology and rigid lens design are discussed.

1.3 Monomers

The monomers that can be used for contact lenses, which are described in the following sections and in both tables and references, are collected in Table 1.2.

These issues will be detailed in the following sections of this chapter.

1.3.1 *Monomers for Block Copolymers*

A block copolymer that contains both hydrophobic and hydrophilic blocks with amino acid groups has been described (18).

The principal monomers for such block copolymers are a combination of two monomers capable of forming a hydrogel; such monomers are collected in Table 1.3.

Table 1.2 Monomers for contact lenses.

Monomers and monomer types	Usage	References
2-Hydroxyethyl methacrylate <i>N</i> -Vinyl-2-pyrrolidone Methyl methacrylate Isobornyl methacrylate <i>tert</i> -Butylcyclohexyl methacrylate	Soft lenses	(19)
Hydrophobic monomers	Strengthening agents	Table 1.8
Hydrophilic monomers		Table 1.8
Hydrophilic monomers	Hydrogels	Table 1.10
Azlactones	Surface treatment	Table 1.12
Acrylamide <i>N</i> -Hydroxyethyl acrylamide <i>N</i> -Isopropyl acrylamide 2-Acrylamido-2-methylpropane-sulfonic acid 2-Hydroxyethyl methacrylate 2-Hydroxyethyl acrylate	Macromers	(20)
Acryl monomers	Water absorbable	Table 1.16
Poly(siloxane)	Water absorbable	Table 1.17
4-(Phenyldiazenyl) phenyl methacrylate	Blue-light blocking	(21)
Acrylates	UV-blocking	Table 1.22
Silicone hydrogel	Multifocal lenses	Table 1.23
Acrylates	Non-silicone hydrogel	Table 1.25
Crosslinking agents	Non-silicone hydrogel	Table Table 1.26
Oxyperm	Oxygen permeable	Table 1.32
Ionoperm	Oxygen permeable	Table 1.32

Table 1.3 Monomers (18).

Monomer	Monomer
2-Ethylphenoxy acrylate	2-Ethylphenoxy methacrylate
2-Ethylthiophenyl acrylate	2-Ethylthiophenyl methacrylate
2-Ethylaminophenyl acrylate	2-Ethylaminophenyl methacrylate
Phenyl acrylate	Phenyl methacrylate
Benzyl acrylate	Benzyl methacrylate
2-Phenylethyl acrylate	2-Phenylethyl methacrylate
3-Phenylpropyl acrylate	3-Phenylpropyl methacrylate
3-Propylphenoxy acrylate	3-Propylphenoxy methacrylate
4-Butylphenoxy acrylate	4-Butylphenoxy methacrylate
4-Phenylbutyl acrylate	4-Phenylbutyl methacrylate

Side-chain-linked amino acids are collected in Table 1.4. Some of these compounds are shown in Figure 1.1.

Table 1.4 Side-chain-linked amino acids (18).

Monomer	Monomer
Acryloyl- <i>L</i> -lysine	Acryloyl- <i>L</i> -serine
Acryloyl- <i>L</i> -threonine	Acryloyl- <i>L</i> -tyrosine
Acryloyl- <i>L</i> -amino-phenylalanine	Acryloyl- <i>L</i> -cysteine
Acryloyl- <i>L</i> -oxy-proline	N ϵ -acryloyl-N α -Oelityl- <i>L</i> -Lysine

The synthesis of a variety of such monomers has been detailed (18). For example, the synthesis of (*S*)-6-acrylamido-2-amino-hexanoic acid monomers is performed via a copper complex (18):

Preparation 1-1: L-lysine (14.62 g; 100 mmol) was dissolved in 150 ml deionized water and heated to about 80°C. Copper carbonate (16.6 g; 75 mmol) was added in portions over a period of 30 min. The reaction was stirred for an additional 30 min. The hot, deep-blue suspension was filtered through silica gel. The filter was washed with a small amount of water. On the following day, the lysine copper complex containing the combined filtrate was cooled in an ice bath, and 100 ml tetrahydrofuran was added. A solution of acryloyl chloride in methyl-*tert*-butylether (8.9 ml, 110 mmol) was added dropwise during a period of 1 h. The pH was initially maintained between 8 and 10 by parallel, dropwise addition of 10% sodium hydroxide solution. After half of the acryloyl chloride solution had been added, the product began to precipitate. When most of the acryloyl chloride had been added, addition of sodium hydroxide was

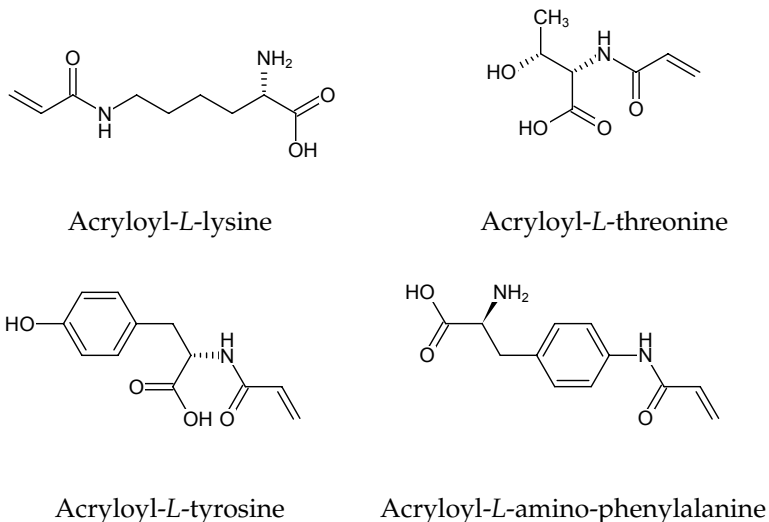


Figure 1.1 Monomers for side-chain-linked amino acids.

slowed down to allow the pH to drop to about 6 and the temperature of the reaction mixture was allowed to reach room temperature. The blue suspension was stirred for an additional 2 h and was then filtered. The solid material retained on the filter was washed with water and acetone and then dried. A yield of 6.5 g of acryloyl-L-lysine copper complex was obtained. Acryloyl-L-lysine copper complex (29.5 g) was suspended in 300 ml deionized water and cooled in an ice bath. H₂S gas was bubbled into the suspension until copper sulfide precipitation was complete; then 3 g of active charcoal was added to the suspension. The suspension was heated briefly to 100°C. After cooling to room temperature, 500 ml acetone was added to the suspension which was then filtered on silica gel. The clear filtrate was put in a rotary evaporator. After evaporation of the solvent, the solid product was recrystallized from 200 ml of 50% aqueous acetone. A yield of 17.76 g (69.76%) of white powder was obtained. The structure of the compound was verified by nuclear magnetic resonance spectroscopy and LC-MS spectroscopy.

The preparation of a block copolymer containing a hydrophilic cellophil polymer and a lipid-like copolymer was done by raft polymerization as follows (18):

Preparation 1-2: Step A

In a 50 ml round-bottom flask, a solution of 2-(dodecylthiocarbonothioylthio)-2-methylpropionic acid (6.13 mg, 0.017 mmol), *N,N*-Dimethyl acrylamide (0.624 ml, 6.05 mmol), and *iso*-decyl acrylate (0.163 ml, 0.673 mmol) in 10 ml *N,N*-dimethylformamide was degassed using ultrasonic treatment. Subsequently, 2-benzyl-2-(dimethylamino)-4'-morpholinobutyrophenone (6.40 mg, 0.017 mmol) was added, and polymerization was induced by UV light. After 4 h of polymerization under stirring, the reaction mixture was purified by extensive dialysis against deionized water using a membrane with a 3.5 kDa MWCO. The mixture was subsequently lyophilized. The average molecular weight (12 kDa) and PDI (1.19) of the block copolymer was verified by GPC measurement.

Step B

The lyophilized macro-CTA prepared in step A (300 mg, 6.82 μ mol) was mixed with acryloyl-*L*-lysine (100 mg, 0.499 mmol) in 10 ml deionized water. The mixture was degassed using ultrasonic treatment. 2,2'-Azobis(2-methylpropionamidine) dihydrochloride (4.62 mg, 0.017 mmol) was added to the mixture. The polymerization was induced by heating the mixture in a reaction vessel to 50°C. After 4 h of polymerization at 50°C, the resulting block copolymer was purified by extensive dialysis against deionized water using a membrane with a 3.5 kDa MWCO. The cellophil block copolymer was subsequently lyophilized. The average molecular weight (18 kDa) and PDI (1.25) of the block copolymer were verified by GPC measurement. Larger block copolymers (32 kDa, PDI 1.28; 58 kDa, PDI 1.24) were obtained by decreasing the ratio of CTA to monomers in step A from 1/100 to 1/200 (32 kDa) and 1/400 (58 kDa), respectively, whereas the molar ratios of *i*-decylacrylate (7.5 mol of 5), DMA (63.9 mol of 5) and AK (28.6 mol of 5) were kept constant.

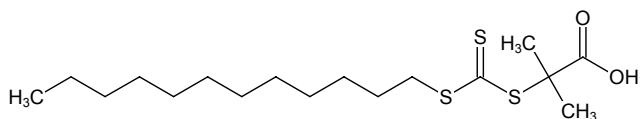
Some of the compounds mentioned in Preparation 1–2 are shown in Figure 1.2.

1.3.2 Silicone Acrylamides

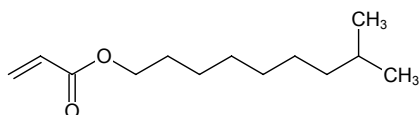
Examples of hydrophilic methacrylamide monomers are collected in Table 1.5. Some of these compounds are shown in Figure 1.3. Also, several other similar monomers have been detailed (22).

These alkyl and aryl groups can be straight or branched. Of these monomers, the *N*-(2-hydroxyethyl)methacrylamide monomer is preferable from a perspective of increasing the transparency of the so obtained polymer.

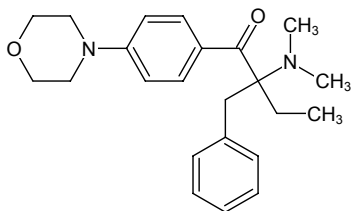
The monomer mixture for synthesizing the polymer additionally may contain between about 1% and about 30% of a hydrophilic



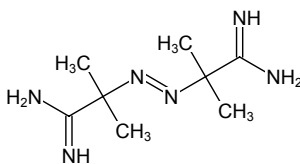
2-(Dodecylthiocarbonothioylthio)-2-methylpropionic acid



iso-Decyl acrylate



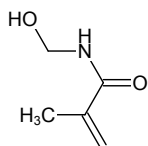
2-Benzyl-2-(dimethylamino)-4'-morpholinobutyphenone



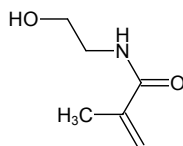
HCl

2,2'-Azobis(2-methylpropion- amide) dihydrochloride

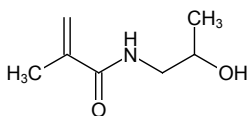
Figure 1.2 Compounds for a block copolymer.



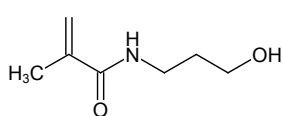
N-Hydroxymethyl methacrylamide



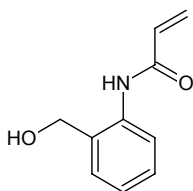
N-(2-Hydroxyethyl) methacrylamide



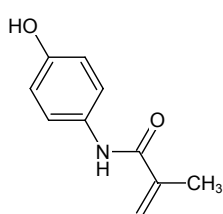
N-(2-Hydroxypropyl) methacrylamide



N-(3-Hydroxypropyl) methacrylamide



N-(2-Hydroxymethylphenyl) methacrylamide



N-(4-Hydroxymethylphenyl) methacrylamide

Figure 1.3 Hydrophilic methacrylamide-based monomers.

Table 1.5 Hydrophilic methacrylamide-based monomers (23).

Compound
<i>N</i> -Hydroxymethyl methacrylamide
<i>N</i> -(2-Hydroxyethyl) methacrylamide
<i>N</i> -(2-Hydroxypropyl) methacrylamide
<i>N</i> -(3-Hydroxypropyl) methacrylamide
<i>N</i> -(2-Hydroxybutyl) methacrylamide
<i>N</i> -(3-Hydroxybutyl) methacrylamide
<i>N</i> -(4-Hydroxybutyl) methacrylamide
<i>N</i> -(2-Hydroxymethylphenyl) methacrylamide
<i>N</i> -(3-Hydroxymethylphenyl) methacrylamide
<i>N</i> -(4-Hydroxymethylphenyl) methacrylamide

polymer with a molecular weight of about 1000 *Dalton* or higher in the monomer and polymer component of the monomer mixture in order to enhance the wettability, resistance to adhesion of proteins, resistance to adhesion of lipids and combinations thereof.

Examples of hydrophilic polymers that can be used in the polymer are shown in Table 1.6. Some of the monomers of these compounds are shown in Figure 1.4.

Hydrophilic polymers selected from poly(vinyl pyrrolidone), poly(*N,N*-dimethyl acrylamide), poly(acrylic acid), and poly(vinyl alcohol) may be particularly effective for enhancing the wettability of silicone hydrogels (23). Poly(vinyl pyrrolidone) and poly(*N,N*-dimethyl acrylamide) provide a balance between the wettability and the compatibility of the polymerization mix in certain formulations.

The polymer can also include a monomer with two or more reactive groups as a copolymerization component. In this case, the polymer becomes solvent resistant.

Preferable monomers with two or more vinyl groups include bifunctional and polyfunctional acrylates. Examples are shown in Table 1.7. Some bisacrylamide monomers are shown in Figure 1.5. Polyfunctional methacrylate compounds are shown in Figure 1.6.

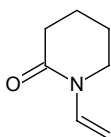
A polymerization initiator may be added to enhance the polymerization traction. Suitable initiators include thermal polymerization initiators, such as a peroxide compound or an azo compound, or

Table 1.6 Hydrophilic polymers (23).

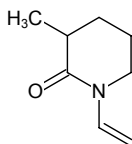
Polymer compound
Poly(<i>N</i> -vinyl pyrrolidone)
Poly(<i>N</i> -vinyl-2-piperidone)
Poly(<i>N</i> -vinyl-2-caprolactam)
Poly(<i>N</i> -vinyl-3-methyl-2-caprolactam)
Poly(<i>N</i> -vinyl-3-methyl-2-piperidone)
Poly(<i>N</i> -vinyl-4-methyl-2-piperidone)
Poly(<i>N</i> -vinyl-4-methyl-2-caprolactam)
Poly(<i>N</i> -vinyl-3-ethyl-2-pyrrolidone)
Poly(<i>N</i> -vinyl-4,5-dimethyl-2-pyrrolidone)
Poly(2-vinylimidazole)
Poly(<i>N</i> -vinyl formamide)
Poly(<i>N</i> -vinyl acetamide)
Poly(<i>N</i> -methyl- <i>N</i> -vinyl acetamide)
Poly(<i>N,N</i> -dimethyl acrylamide)
Poly(<i>N,N</i> -diethyl acrylamide)
Poly(<i>N</i> -isopropyl acrylamide)
Poly(vinyl alcohol)
Poly(acrylate)
Poly(ethylene oxide)
Poly(2-ethyl oxazoline)
Heparine Polysaccharide
Poly(acryloyl morpholine)

Table 1.7 Multifunctional monomers (23).

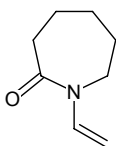
Compound	Compound
Ethylene glycol acrylate	Ethylene glycol dimethacrylate
Diethylene glycol diacrylate	Diethylene glycol dimethacrylate
Triethylene glycol diacrylate	Triethylene glycol dimethacrylate
Neopentyl glycol diacrylate	Neopentyl glycol dimethacrylate
Tetraethylene glycol diacrylate	Tetraethylene glycol dimethacrylate
Glyceryl triacrylate	Glyceryl trimethacrylate
Pentaerythritol tetraacrylate	Pentaerythritol tetramethacrylate
Trimethylol propane triacrylate	Trimethylol propane trimethacrylate
<i>N,N'</i> -Methylene bisacrylamide	<i>N,N'</i> -Ethylene bisacrylamide
<i>N,N'</i> -Propylene bisacrylamide	



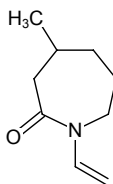
N-Vinyl-2-piperidone



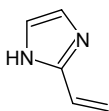
N-Vinyl-3-methyl-2-piperidone



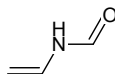
N-Vinyl-2-caprolactam



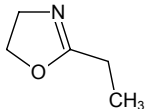
N-Vinyl-4-methyl-2-caprolactam



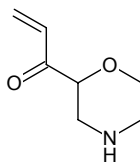
2-Vinylimidazole



N-Vinyl formamide

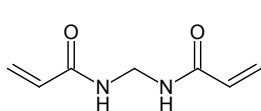
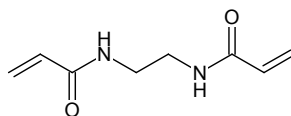
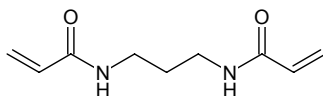


2-Ethyl oxazoline



Acryloyl morpholine

Figure 1.4 Monomers for hydrophilic polymers.

*N,N'*-Methylene bisacrylamide*N,N'*-Ethylene bisacrylamide*N,N'*-Propylene bisacrylamide**Figure 1.5** Bisacrylamide monomers.

photopolymerization initiators. Also, photoinitiators can be added in order to enhance the polymerization.

Several examples of the polymerization procedure have been detailed (23).

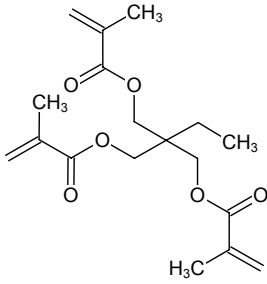
1.4 Soft Lenses

1.4.1 Hydrogels

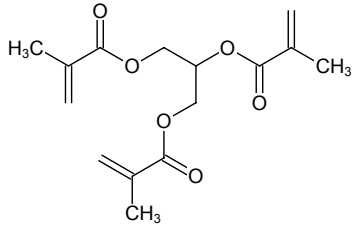
Soft contact lens materials are made by polymerizing and crosslinking hydrophilic monomers such as 2-hydroxyethyl methacrylate, *N*-vinyl-2-pyrrolidone, and combinations thereof (19).

The polymers produced by polymerizing these hydrophilic monomers exhibit significant hydrophilic character themselves, and are capable of absorbing a significant amount of water in their polymeric matrices. Due to their ability to absorb water, these polymers are often referred to as *hydrogels*.

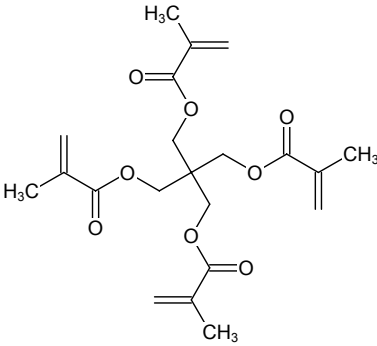
These hydrogels are optically clear and, due to their high levels of water of hydration, are particularly useful materials for making soft contact lenses. However, the high levels of water of hydration of hydrogels contributes to their relative lack of physical strength, which results in hydrogel contact lenses being relatively easy to tear (19).



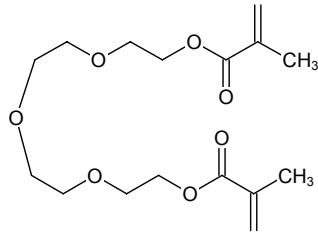
Trimethylol propane
trimethacrylate



Glycerol
trimethacrylate



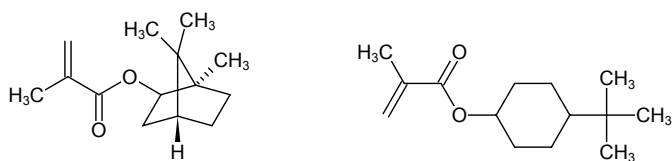
Pentaerythritol
tetramethacrylate



Tetraethylene glycol
dimethacrylate

Figure 1.6 Polyfunctional methacrylate compounds.

Various hydrophobic monomers have been copolymerized with these hydrophilic monomers in order to obtain polymers with an improved physical strength. Such hydrophobic monomers include styrene, and various acrylates and methacrylates such as methyl methacrylate, isobornyl methacrylate, and *tert*-butylcyclohexyl methacrylate. The last two compounds are shown in Figure 1.7.



Isobornyl methacrylate *tert*-Butylcyclohexyl methacrylate

Figure 1.7 Hydrophobic monomers.

While these hydrophobic monomers do increase the physical strength of hydrogel polymers, they also produce polymers with lower levels of water of hydration than the unmodified hydrogels.

So, an attempt was made to provide polymeric materials with an increased physical strength and high levels of water of hydration (19).

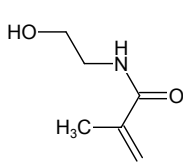
It has been found that certain hydrophobic monomers act as strengthening agents when copolymerized with hydrophilic monomers and crosslinkers. Examples of these monomers are shown in Table 1.8 and in Figures 1.8, 1.9, and 1.10.

Most preferred contact lenses have an oxygen transport rate of at least about $2 \times 10^{-6} \text{ cm}^3 \text{ sec}^{-1} \text{ cm}^{-2} \text{ atm}^{-1}$, which makes them hydrolytically stable, biologically inert, transparent, and resilient. Furthermore, they should preferably have a softness of about 60 or below on the Shore hardness A scale when hydrated. The more preferred materials have a Shore hardness between 25 to 35 on the A scale.

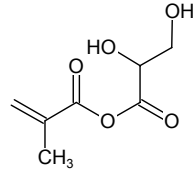
The tensile modulus of elasticity of these hydrated polymers is at least about 50 g mm^{-2} , preferably from about 75 g mm^{-2} to about 100 g mm^{-2} , and the tear strength is at least about 2.0 g mm^{-1} thickness, preferably from about 2.0 g mm^{-1} to about 250 g mm^{-1} thickness.

Table 1.8 Monomers (19).

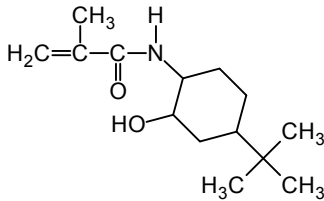
Hydrophilic monomers
2-Hydroxyethyl methacrylate
<i>N</i> -(2-Hydroxy ethyl)-methacrylamide
<i>N</i> -Vinyl-2-pyrrolidone
Glyceryl methacrylate
<i>N</i> -Methacryloyl glycine
2-Hydroxyl-3-methacryl(propyl)-4-methoxy phenyl ether
2-Hydroxy cyclohexyl methacrylate
Hydrophobic strengthening agent monomers
4- <i>tert</i> -Butyl-2-hydroxycyclohexyl methacrylate
4- <i>tert</i> -Butyl-2-hydroxycyclopentyl methacrylate
Methacryloylamino-4- <i>tert</i> -butyl-2-hydroxycyclohexane
6-Isopentyl-3-hydroxycyclohexyl methacrylate
Methacryloylamino-2-isohexyl-5-hydroxy cyclopentane
Crosslinking monomers
Allyl methacrylate
Ethylene glycol dimethacrylate
Divinyl ethylene urea
1,3-Bis(4-methacryloxybutyl) tetramethyl disiloxane



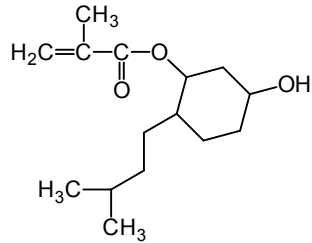
N-(2-Hydroxy ethyl)-methacrylamide



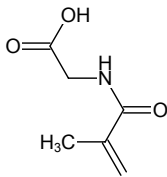
Glyceryl methacrylate



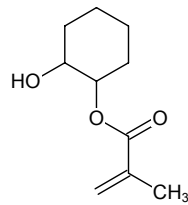
Methacryloylamino-4-*tert*-butyl-2-hydroxycyclohexane



6-Isopentyl-3-hydroxy cyclohexyl methacrylate

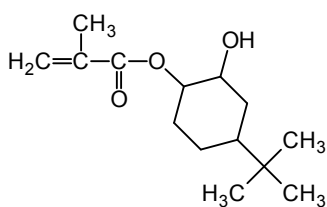


N-Methacryloyl glycine

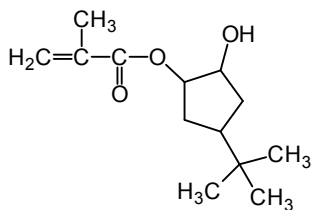


2-Hydroxy cyclohexyl methacrylate

Figure 1.8 Hydrophilic monomers.

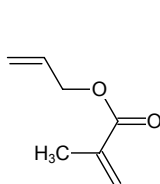


4-*tert*-Butyl-2-hydroxy-
cyclohexyl methacrylate

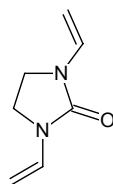


4-*tert*-Butyl-2-hydroxy-
cyclopentyl methacrylate

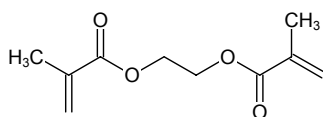
Figure 1.9 Hydrophobic strengthening monomers.



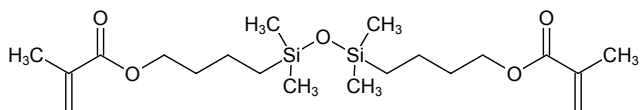
Allyl methacrylate



Divinyl ethylene urea



Ethylene glycol dimethacrylate



1,3-Bis(4-methacryloxybutyl) tetramethyl disiloxane

Figure 1.10 Crosslinking monomers.

High tensile modulus of elasticity is desirable for strength and durability. High tear strength is desirable in order to prevent damage to the contact lens due to tearing during patient use, i.e., the removing and placing of the lens on the eye, and to prevent damage to the lens during cleansing and disinfecting.

A soft contact lens formulation is illustrated in the following example (19):

Preparation 1-3: A mixture was made containing 77.0 g of glyceryl methacrylate, 22.5 g of 4-*tert*-butyl-2-hydroxycyclohexyl methacrylate, and 0.5 g of ethylene glycol dimethacrylate. To this mixture was added 0.5 g of benzoin methyl ether, an ultraviolet-induced polymerization initiator. The solution was cast between glass plates separated by a Teflon perfluoro polymer gasket, 0.3 mm thick and cured. After curing, the film was released from the glass plates and hydrated and extracted in hot distilled water for 4 h.

Then, the film was placed in a borate buffered saline solution for testing. The resultant material was optically clear and had a water content of 53% and an oxygen permeability of $18 \times 10^{-11} \text{ cm}^3 \text{ cm sec cm}^2 \text{ mm Hg}^{-1}$.

The mechanical properties were measured according to the following test methods and gave the results shown in Table 1.9.

Table 1.9 Mechanical properties (19).

Standard	Ref.	Name	Value
ASTM-D 1708	(24)	Young's modulus of elasticity	60 $g \text{ mm}^{-2}$
ASTM-D 1708	(24)	Tensile strength	84 $g \text{ mm}^{-2}$
ASTM-D 1708	(24)	% Elongation	164 %
ASTM-D 1938	(25)	Tear Initiation	3.8 $g \text{ mm}^{-1}$

1.4.1.1 Polymerizable Contact Lens Formulations

Hydrogel contact lenses, polymerizable compositions useful for making such lenses, packaging systems for use with such lenses and methods of producing such lenses have been discovered (26). These contact lenses have a relatively low surface friction and are able to release hydrophilic polymers present in the contact lenses for prolonged periods of time.

The contact lenses have a lens body. The lens body is the reaction product of a polymerizable composition containing one or

more monomers, and a crosslinker that crosslinks the monomers during a polymerization reaction to form a first polymer component. The polymerizable composition also contains a hydrophilic polymer component, which is substantially unreactive during the polymerization.

Thus, the resulting lens body includes a first polymer component formed from the one or more monomers present in the polymerizable composition, and the second polymer component, the hydrophilic polymer component that is physically entangled with the first polymer component in the lens body.

The hydrophilic polymer component is unreactive or substantially unreactive during the polymerization process. Thus, the resulting hydrogel lens body can be understood to be a network of a first polymeric component, formed from the monomers present in the polymerizable composition, and a second polymeric component, the hydrophilic polymer component, in which the hydrophilic polymer component is substantially physically entrapped by the first polymer component. Although there may be some small amounts of reactivity of the hydrophilic polymer component, the reactivity is not sufficient to prevent leaching or release of the hydrophilic polymer from the lens body. The present contact lenses can be understood to consist of an interpenetrating polymer network where the formation of the first polymer component occurs in the presence of the hydrophilic polymer component. However, as discussed herein, in the present contact lenses, it is possible for the hydrophilic polymer component to be released from the lens body even though it is entrapped by the first polymer component.

Examples of these monomers are shown in Table 1.10 and in Figure 1.11.

Polymerization initiators can be used in the polymerizable composition. Thermal initiators that may be used are azo or peroxide compounds, such as those having a half-life at the polymerization temperature of at least 20 *min*. Examples of initiators are shown in Table 1.11 and in Figure 1.12.

A tinting agent can be any agent that imparts a visibility to the otherwise clear hydrogel lens body. The tinting agent may be a water-soluble dye, or particles of pigment, or combinations thereof. Some examples of tinting agents include copper phthalocyanine