



A REFERENCE GUIDE

HIGH-PERFORMANCE MATERIALS FOR HEALTHCARE APPLICATIONS

Version 2.0



PATIENTS DEPEND ON DOCTORS

Doctors and other healthcare professionals depend on medical devices.

Those who design medical devices depend on Covestro, one of the leading producers of high-performance plastics around the world.

COVESTRO — A MARKET LEADER WITH MORE THAN 50 YEARS' EXPERIENCE IN HEALTHCARE

From devices to treat disease like glucose monitors and drug delivery devices, to devices used to treat acute medical emergencies like intravenous access components and surgical instruments — Covestro materials are essential.

Covestro is a global market leader in supplying resins to the healthcare market, and our continued success is built on five principles:

1. Over 50 years' experience serving the healthcare market, backed by a deep understanding of the industry's needs.
2. One-stop shop for our customer's polycarbonate (and blends) needs.
3. Demonstrated supplier reliability with resin production plants around the world, able to produce globally consistent healthcare grades.
4. Committed to the future of healthcare customers, offering value-added services and developing new products.
5. Dedicated to sustainability, utilizing renewable feedstocks and circular design principles to drive a more sustainable healthcare industry.

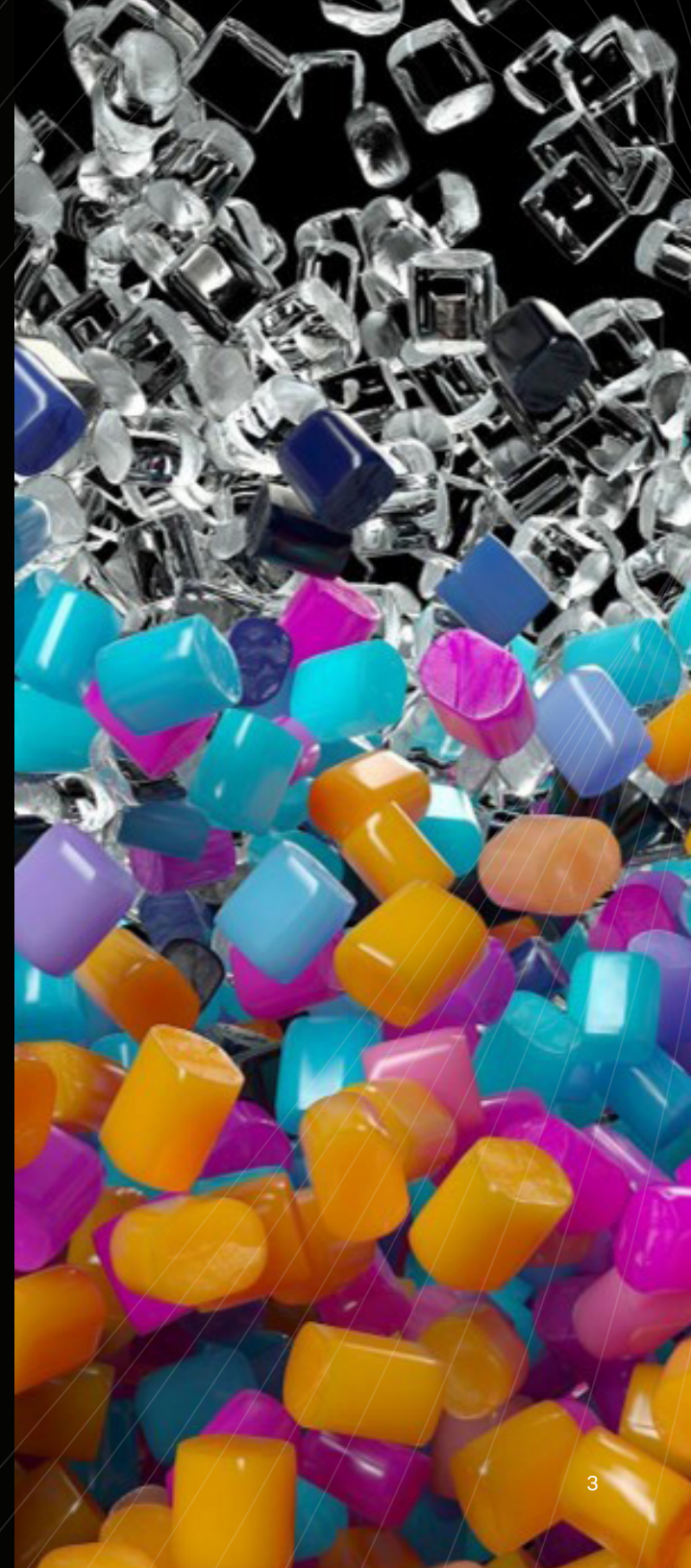




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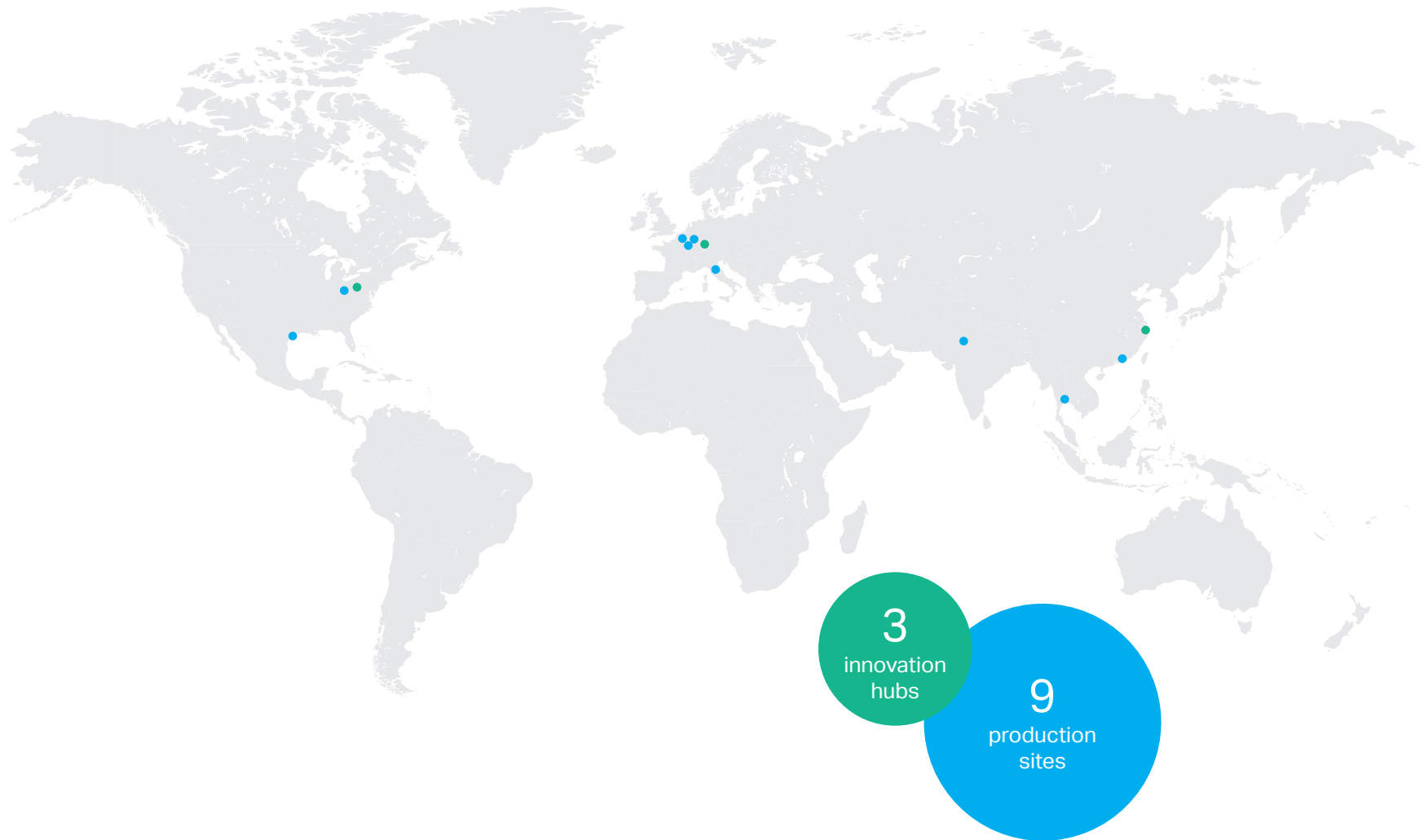
This brochure reviews our polycarbonate-based and thermoplastic polyurethane materials designed for use in healthcare applications. It includes typical applications, their performance requirements and the Covestro materials that may meet those requirements.

Our intent is to provide a general overview of these materials and their performance characteristics specific to the healthcare industry.

COVESTRO LOCATIONS AROUND THE GLOBE

Covestro operates a robust network of production sites and innovation centers worldwide.

We take pride in the global consistence, quality and reliable supply of our healthcare polycarbonate materials.





COVESTRO MEDICAL GRADES AND REGULATORY COMPLIANCE

Our market-leading portfolio meets the following criteria:

- Biocompatibility: ISO 10993 and USP Class VI for contact of 30 days or less.
- Compliance to material requirements of Regulation (EU) 2017/745 on medical devices and (EU) No 722/2012 regarding medical devices manufactured utilizing tissues of animal origin.
- Adhere to disclosures in FDA Device Files (MAF) and Drug Master Files (DMF). Letters of authorization are available.
- Manufactured at ISO 9001 certified sites that follow Good Manufacturing Practices (GMP) standards.

This brochure presents Covestro's polycarbonate and TPU medical-grade portfolios. Covestro also offers a large portfolio of standard resins, which may be used for applications which do not need the stringent requirements of change notification and biocompatibility of a medical-grade product.

Please contact your local Covestro representative for more details. To connect with an expert today, email plastics@covestro.com.

COVESTRO HEALTHCARE PORTFOLIO

Makrolon[®], Bayblend[®], Makroblend[®], and Apec[®] Medical Grades

With our family of polycarbonate resins and resin blends, you can find the material that best suits the requirements of your medical device or healthcare application. These products are biocompatible per certain ISO 10993 test requirements. Below is an overview of these products and their properties. The products are available in granule form and are typically processed by injection molding, extrusion or blow molding.

Makrolon[®] Polycarbonate

This lightweight resin offers a unique combination of impact resistance and rigidity, and is available in transparent, translucent and opaque colors. With a range of grades to meet the needs of medical applications, Makrolon[®] medical grades can be sterilized by all common methods.

Apec[®] High-Heat Polycarbonate

This copolymer is characterized by its high transparency, toughness and strength, with a higher heat resistance than Makrolon[®] resin. It is suitable for high-heat steam autoclaving up to 143°C.

Bayblend[®] PC+ABS Blend

This opaque resin is a blend of polycarbonate (PC) and acrylonitrile butadiene styrene polymer (ABS). It offers an excellent combination of mechanical and thermal properties, exhibiting high toughness, rigidity, dimensional stability and easy processing.

Makroblend[®] PC + Polyester Blend

This opaque is a blend of polycarbonate and polyester (PBT and PET) that offers an excellent combination of toughness and chemical resistance.

Flame-retardant Resins

Flame retardant medical Makrolon[®], Bayblend[®] and Makroblend[®] grades are available with UL94 V-0 ratings down to 1.5mm.

Covestro Also Offers Texin[®] Rx Thermoplastic Polyurethane Medical Grades

Biocompatible per certain ISO 10993 test requirements, these resins are tough, high-tensile materials with good tear strength and excellent resistance to abrasion, fuels, oils and greases. Elastic and resilient, they also offer great cold temperature flexibility, flex fatigue properties, excellent adhesion to various substrates and a Shore hardness range from 70A to 85D. The products are available in granule form and are typically processed by injection molding and extrusion.

ADVANCING HEALTHCARE SUSTAINABILITY

Sustainability is a core element of our vision to work toward becoming more circular, and an integral part of our strategy. We live and breathe innovation and continue to offer new products and innovative technologies that benefit society while reducing the impact on certain aspects of our ecological systems.

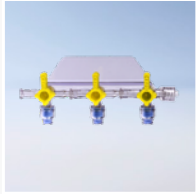
We support our healthcare customers with diverse solutions to contribute to a more circular economy:

- **Renewable feedstock:** Up to 89% renewable content via mass balance approach, with several ISCC Plus certified production sites
- **Recycling solutions:** Mechanical recycling for select pure waste streams and chemical recycling to convert certain waste types into base chemicals for high-quality plastics
- **Circular design:** Application development support for transitioning to mono-material solutions, optimizing material deployment, and improving recycling at end-of-life
- **Joint solutions:** Help close material loops while increasing transparency and maintaining device performance

To address climate change and resource constraints, we offer select sustainable engineering plastics containing up to 89% renewable content via mass balancing, alongside innovative solutions toward a more circular, lower-carbon healthcare future compared to conventional alternatives.



APPLICATION CASE STUDIES



IV Access

Customer: Elcam

Product: Marvelous™ 3-way stopcock

Material: **Makrolon® Rx1805 resin**

Benefits: Transparent; biocompatible; dimensionally stable; radiation resistant

[More information on Covestro Solution Center](#)



Drug Delivery

Customer: QS Medical Technology

Product: Quinnocare QS-P insulin pen

Material: **Makrolon® Rx1805 resin (ampoule)**

Bayblend® FR3050 resin (housing)

Benefits: Transparent; radiation resistant (ampoule); chemical resistant, paint elimination (housing)

[More information on Covestro Solution Center](#)



Surgical

Customer: Metric Medical Devices, Inc.

Product: Super Staple™ Classic bone fixation device

Material: **Makrolon® Rx2530 resin**

Benefits: Tough; dimensionally stable

[More information on Covestro Solution Center](#)



Healthcare Electronics

Customer: Howard Technology Solutions

Product: HI-Care E 2-tier cart

Material: **Makroblend® M525 resin**

Benefits: Chemical resistant; tough; processability; biocompatible

[More information on Covestro Solution Center](#)



Renal Therapy

Customer: Chengdu OCI Medical Device Co., Ltd.

Product: Dialyzer housing

Material: **Makrolon® Rx2440 resin**

Benefits: Designed for low-oxygen radiation sterilization; biocompatible; good processability

[More information on Covestro Solution Center](#)



Respiratory

Customer: Breathe Technologies

Product: Breathe Pillow Interface™

Material: **Makrolon® 2458 resin**

Benefits: Medium viscosity; easy release; EtO and steam sterilizable; biocompatible

[More information on Covestro Media](#)

APPLICATION CASE STUDIES



Wellness

Customer: Apollo Neuroscience

Product: Apollo

Material: Makroblend® M525 resin

Benefits: Impact resistant; chemical resistant; biocompatible

[More information on Covestro Solution Center](#)



Cardiovascular

Customer: Wego New Life Medical Devices

Product: Oxygenator

Material: Makrolon® 2458 resin

Benefits: Transparent; durable; biocompatible; heat resistant; excellent processability

[More information on Covestro Solution Center](#)



Clinical Communication

Customer: Beam Mobile

Product: Mobile device cases

Material: Makrolon® polycarbonate

Benefits: Impact resistant; chemical resistant; biocompatible; design freedom

[More information on Covestro Solution Center](#)



Optical

Customer: Occuity

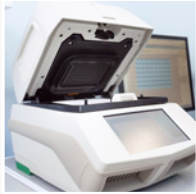
Product: AX1 Axiometer™ for myopia management

Material: Makrolon® 2458 resin

Benefits: Durable; custom color and finish; biocompatible; flame retardant

[More information on Covestro Solution Center](#)

MATERIAL SHOWCASE



Makroblend® M5005 FR Resin

A FR polycarbonate/polyester blend specifically designed for the healthcare market.

Benefits: Chemically resistant; advanced phosphorous based FR UL94 V-0 @ 2.4mm; excellent processability; biocompatible

[More information on Covestro Solution Center](#)



Makrolon® GF Resin Series

A family of six healthcare grades with glass content ranging from 10-30%. Available as high-flow and high-performance options.

Benefits: Superior rigidity; dimensionally stable; biocompatible; globally available

[More information on Covestro Solution Center](#)

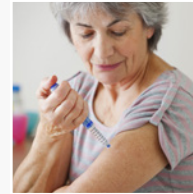


Makrolon® M6011 FR Resin

A FR polycarbonate resin specifically designed for the healthcare market.

Benefits: Chemically resistant; advanced phosphorous based FR UL94 V-0 @ 1.5mm; biocompatible

[More information on Covestro Solution Center](#)



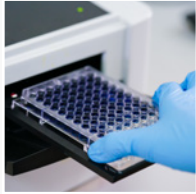
Makrolon® LF Resin Series

A family of opaque healthcare grades with lower surface friction properties. Options for reinforcement with glass fibers are also available.

Benefits: Lower deployment forces; dimensionally consistent; biocompatible; globally available

[More information on Covestro Solution Center](#)

MATERIAL SHOWCASE



Makrolon® Rx2235 Resin

A transparent polycarbonate resin for the healthcare market with outstanding high-flow properties.

Benefits: Excellent processability; transparent; radiation stabilized; biocompatible

[More information on Covestro Press](#)



Makrolon® 3638

A polycarbonate resin with a unique combination of properties maintained over a wide temperature range.

Benefits: Durable; chemically resistant; biocompatible; sterilizable

[More information on Covestro Press](#)



Makrolon® Rx3440 Resin

A transparent polycarbonate resin for the healthcare market with best-in-class chemical and oncology drug resistance.

Benefits: Chemically resistant; radiation stabilized; transparent; biocompatible

[More information on Covestro Solution Center](#)



Apec® 2045

A transparent polycarbonate offering the highest heat resistance in our medical polycarbonate portfolio.

Benefits: High heat resistance; sterilizable; biocompatible; easy release

[More information on Covestro Press](#)

DESIGN STUDIES



Drug Delivery

A drug delivery demonstrator showcases how polycarbonates can improve the functionality and sustainability of medical devices.

Benefits: Recyclable; low friction; cosmetic flexibility; sterilizable via radiation

[More information on Covestro Solution Center](#)



Drug Delivery

A connected drug delivery concept device highlighting the benefits of using Covestro materials.

Benefits: Tough and durable; biocompatible; sterilizable; dimensional stability; design flexibility

[More information on Covestro Solution Center](#)



Wellness

Smart pill dispenser concept demonstrates the potential of medical polycarbonate solutions with smart sensor technology.

Benefits: Design flexibility; chemically resistant; biocompatible

[More information on Covestro Solution Center](#)



Diagnostics

Demonstrating the convergence of healthcare and consumer electronics, this wearable Continuous Glucose Monitor concept combines functionality and fashion

Benefits: Tough and durable; colorable; translucent; design flexibility; processability; biocompatible

[More information on Covestro Solution Center](#)

PRODUCT OVERVIEW OF MEDICAL GRADES



PRODUCT OVERVIEW OF MEDICAL GRADES

Standard	Radiation Stabilized	Glass Filled	Low Friction	Flame Retardant Blends	Non-Flame Retardant Blends	High Heat
Makrolon® 2258	Makrolon® Rx2235	Makrolon® M410 GF	Makrolon® M204 LF	Makrolon® M6011 FR^{1,2}	Bayblend® M750	Apec® 1745
Makrolon® 2458	Makrolon® Rx2430	Makrolon® M420 GF	Makrolon® M402 LF	Bayblend® M301 FR²	Bayblend® M850 XF	Apec® 2045
Makrolon® Rx1452	Makrolon® Rx2440	Makrolon® M430 GF	Makrolon® M404 LF	Bayblend® M303 FR²	Makroblend® M525²	
Makrolon® Rx1851	Makrolon® Rx2435	Makrolon® M810 GF		Makroblend® M4000 FR²		
Makrolon® 2558	Makrolon® Rx2530	Makrolon® M820 GF		Makroblend® M5005 FR^{1,2}		
Makrolon® 2658	Makrolon® Rx2635	Makrolon® M830 GF				
Makrolon® 2858	Makrolon® Rx1805					
Makrolon® 3158	Makrolon® Rx3440					
Makrolon® 3258						
Makrolon® 3638						

¹ Advanced phosphorous based flame retardant formulation

² Limited biocompatibility, meets the biocompatibility requirements of ISO 10993 testing for contact with uncompromised skin.

PRODUCT OVERVIEW OF MEDICAL GRADES¹

Makrolon® 2258 resin (MVR 34 cm³ / 10 min) is a very low viscosity polycarbonate with good release characteristics, suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® 2458 resin (MVR 19 cm³ / 10 min) is a medium/low viscosity polycarbonate with good release characteristics, suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® Rx1452 resin (MVR 16 cm³ / 10 min) is a medium/low viscosity polycarbonate with enhanced release and surface lubricity characteristics, suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® Rx1851 resin (MVR 23 cm³ / 10 min) is a low viscosity polycarbonate with enhanced release and surface lubricity characteristics, suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® 2558 resin (MVR 14 cm³ / 10 min) is a medium viscosity polycarbonate with good release characteristics, suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® 2658 resin (MVR 12 cm³ / 10 min) is a medium viscosity polycarbonate with good release characteristics, suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® 2858 resin (MVR 9 cm³ / 10 min) is a medium viscosity polycarbonate with good release characteristics, suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® 3158 resin (MVR 6 cm³ / 10 min) is a high viscosity, high-performance polycarbonate with good release characteristics, suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® 3258 resin (MVR 5 cm³ / 10 min) is a high viscosity, high-performance polycarbonate with good release characteristics, suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® 3638 resin (MVR 2.0 cm³/10 min) is a high viscosity polycarbonate with low temperature ductility and good release characteristics, suitable for EtO, radiation and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements. Suitable for injection and blow molding.

Makrolon® Rx2235 resin (MVR 37 cm³ / 10 min) is a very low viscosity polycarbonate with easy-release characteristics. It is particularly suitable for sterilization by radiation but may also be sterilized with EtO and steam at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® Rx2430 resin (MVR 19 cm³ / 10 min) is a medium/low viscosity polycarbonate. It is particularly suitable for sterilization by radiation but may also be sterilized with EtO and steam at 121°C. Biocompatible according to certain ISO 10993 test requirements.

¹ See datasheets for MVR test conditions

PRODUCT OVERVIEW OF MEDICAL GRADES¹

Makrolon® Rx2440 resin (MVR 19 cm³ / 10 min) medium/low viscosity polycarbonate. It is stabilized for radiation sterilization of devices in oxygen-free packaging, but may also be sterilized with EtO and steam at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® Rx2435 resin (MVR 23 cm³ / 10 min) is a low viscosity polycarbonate with easy-release characteristics. It is particularly suitable for sterilization by radiation but may also be sterilized with EtO and steam at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® Rx2530 resin (MVR 15 cm³ / 10 min) is a medium viscosity polycarbonate. It is particularly suitable for sterilization by radiation but may also be sterilized with EtO and steam at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® Rx2635 resin (MVR 12 cm³ / 10 min) is a medium viscosity polycarbonate with easy-release characteristics. It is particularly suitable for sterilization by radiation but may also be sterilized with EtO and steam at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® Rx1805 resin (MVR 6 cm³ / 10 min) is a high viscosity, lipid-resistant polycarbonate. It is particularly suitable for sterilization by radiation but may also be sterilized with EtO and steam at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® Rx3440 resin (MVR 4.5 cm³ / 10 min) is a high viscosity, lipid-resistant polycarbonate with improved oncology drug resistance. It is particularly suitable for sterilization by radiation but may also be sterilized with EtO and steam at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® M410 GF resin (MVR 12 cm³ / 10 min) is a medium/low viscosity 10% glass filled polycarbonate. It is suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® M420 GF resin (MVR 9 cm³ / 10 min) is a medium/low viscosity 20% glass filled polycarbonate. It is suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® M430 GF resin (MVR 7 cm³ / 10 min) is a medium/low viscosity 30% glass filled polycarbonate. It is suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® M810 GF resin (MVR 4 cm³ / 10 min) is a high viscosity 10% glass filled polycarbonate. It is suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® M820 GF resin (MVR 3.5 cm³ / 10 min) is a high viscosity 20% glass filled polycarbonate. It is suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® M830 GF resin (MVR 3 cm³ / 10 min) is a high viscosity 30% glass filled polycarbonate. It is suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® M204 LF resin (MVR 34 cm³ / 10 min) is a very low viscosity polycarbonate with excellent low friction properties. It is suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

¹ See datasheets for MVR test conditions

PRODUCT OVERVIEW OF MEDICAL GRADES¹

Makrolon® M402 LF resin (MVR 19 cm³ / 10 min) is a medium/low viscosity polycarbonate with good low friction properties. It is suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® M404 LF resin (MVR 19 cm³ / 10 min) is a medium/low viscosity polycarbonate with excellent low friction properties. It is suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Makrolon® M6011 FR resin (MVR 25 cm³ / 10 min) is an opaque, uv stabilized, advanced phosphorous based flame retardant (V0 @ 1.5mm) polycarbonate with excellent chemical resistance and easy release characteristics suitable for EtO sterilization and steam sterilization at 121°C. Biocompatible according to certain ISO 10993 test requirements.

Bayblend® M301 FR resin (MVR 25 cm³ / 10 min) is an opaque, low viscosity, advanced phosphorous based flame retardant (UL94 V-0 @ 1.5mm), PC+ABS blend. This resin meets the biocompatibility requirements of certain ISO 10993 testing for contact with uncompromised skin.

Bayblend® M303 FR resin (MVR 11 cm³ / 10 min) is an opaque, medium viscosity, advanced phosphorous based flame retardant (UL94 V-0 @ 1.5mm), PC+ABS blend. Suitable for extrusion. This resin meets the biocompatibility requirements of certain ISO 10993 testing for contact with uncompromised skin.

Makroblend® M4000 FR resin (MVR 25 cm³ / 10 min) is an opaque, medium/low viscosity, PC+PBT blend. UL94 listed V-0 @ 2.0mm. This resin meets the biocompatibility requirements of certain ISO 10993 testing for contact with uncompromised skin.

Makroblend® M5005 FR resin (MVR 35 cm³ / 10 min) is an opaque, medium/low viscosity, PC+PBT blend. Advanced phosphorous based flame retardant UL94 listed V0 @ 2.4mm. UV-stabilized and meets the biocompatibility requirements of certain ISO 10993 testing for contact with uncompromised skin.

Bayblend® M750 resin (MVR 11 cm³ / 10 min) is an opaque, medium viscosity, PC+ABS blend. This resin is biocompatible according to certain ISO 10993 test requirements.

Bayblend® M850 XF resin (MVR 25 cm³ / 10 min) is an opaque, low viscosity, PC+ABS blend. This resin is biocompatible according to certain ISO 10993 test requirements.

Makroblend® M525 resin (MVR 21 cm³ / 10 min) is an opaque, low viscosity, PC+PBT blend with exceptional low temperature impact. This resin meets the biocompatibility requirements of certain ISO 10993 testing for contact with uncompromised skin.

Apec® 1745 resin (MVR 17 cm³ / 10 min) is a high-heat copolycarbonate suitable for repeated steam sterilization up to 143°C. This resin has easy-release characteristics, high softening temperature (VST/B 120) = 170 °C, and good hydrolysis resistance. Biocompatible according to certain ISO 10993 test requirements.

Apec® 2045 resin (MVR 8 cm³/10 min) is a high-heat copolycarbonate suitable for superheated steam sterilization up to 143°C. This resin has easy-release characteristics and high softening temperature (VST/B 120) = 203 °C. Biocompatible according to certain ISO 10993 test requirements.

¹ See datasheets for MVR test conditions

PRODUCT OVERVIEW OF STANDARD GRADES

Standard grades for medical applications (non-biocompatible)

The following flame-retardant Bayblend® grades are used in applications such as device housings where flame-retardancy is needed, yet ISO 10993 biocompatibility is not required.

Please contact your local Covestro representative to discuss these and other alternative materials.

Bayblend® FR3005 HF resin (PC+ABS) Blend; flame retardant; very easy-flowing; Vicat/B 120 temperature = 96 °C; UL recognition 94 V-0 at 1.5 mm

Bayblend® FR3010 resin (PC+ABS) Blend; flame retardant; Vicat/B 120 temperature = 110 °C; increased heat resistance; UL94 V-0 at 1.5 mm; glow wire temperature (GWFI): 960 °C at 2.0 mm; improved chemical resistance and stress cracking behavior

Bayblend® FR3010 HF resin (PC+ABS) Blend; flame retardant; easy flowing; Vicat/B 120 temperature = 108 °C; UL94 V-0 at 1.5 mm; glow wire temperature (GWFI): 960 °C at 2.0 mm; optimized processability



STERILIZATION METHODS

Medical devices typically require sterilization before use. There are three sterilization methods most prevalent in the medical industry:

1. Ethylene oxide (EtO gas)
2. Steam autoclave
3. Irradiation with high-energy radiation (gamma or e-beam)

The table below shows which Makrolon®, Bayblend®, Apec® or Makroblend® grades are suitable for each sterilization method. However, the resistance to the different sterilization methods and the number of sterilization cycles a medical device can withstand will vary depending upon the type/grade of Makrolon®, Bayblend®,

Makroblend® and Apec®, part design, processing parameters and other factors. The manufacturer of the medical device must determine the suitability of the sterilization method in each case.

Sterilization Method	Makrolon® 2258 Rx1851 2458 Rx1452 2558 2658 2858 3158 3258 3638	Makrolon® Rx2235 Rx2435 Rx2430 Rx2440 Rx2530 Rx2635 Rx1805 Rx3440	Makrolon® M410 GF M420 GF M430 GF M810 GF M820 GF M830 GF	Makrolon® M204 LF M402 LF M404 LF	Apec® 1745 2045	Bayblend® M750 M850 XF	Makroblend® M525
Ethylene oxide	Yes	Yes	Yes	Yes	Yes	Yes	Yes
Steam 121°C	Yes	Yes	Yes	Yes ²	Yes	No	No
Steam 134°C	No	No	No	No	Yes	No	No
Steam 143°C	No	No	No	No	Yes	No	No
Gamma radiation	Yes ¹	Yes	Yes ¹	Yes ¹	Yes ¹	Yes ¹	Yes ¹
E-beam	Yes ¹	Yes	Yes ¹	Yes ¹	Yes ¹	Yes ¹	Yes ¹

¹See information under "Sterilization by high energy radiation (gamma/e-beam) on page 21"

²Single sterilization cycle only

STERILIZATION METHODS

Sterilization by ethylene oxide (EtO gas)

Ethylene oxide can be used for sterilization either undiluted or in an inert gas mixture containing between 10 to 20% EtO.

During sterilization the temperature should not exceed 65°C. Tests have shown that frequent sterilization can lead to slight brittleness and crack formation. Tests carried out on specimens treated with pure ethylene oxide at 55°C showed that, after 50 cycles of 6 hours each, the impact strength is unchanged compared with the starting level despite slight crack formation.

Sterilization by steam (saturated steam)

The sterilization temperature should not exceed 125°C for Makrolon® resin, otherwise the molded parts can become deformed. Parts made of Apec® resin may be sterilized in steam up to 143°C. Care must also be taken to ensure that the Makrolon® or Apec® resin part is not damaged by any substances added to the boiler feed water, such as alkaline corrosion inhibitors, and that the article is positioned correctly so that no condensation can accumulate inside it.

As a rule, it is possible to sterilize molded parts made of Makrolon® and Apec® resins many times before gradual chemical decomposition reduces the mechanical strength to a level where it is no longer adequate for certain applications. Sterilization tests on on Apec® test specimens have shown that even



after 100 cycles of 30 minutes each at 120 to 125°C, the parts retain comparatively good impact strength. Medical articles made of Makrolon® or Apec® resins, which are intended for single use, are not suitable for multiple use.

Sterilization by high-energy radiation (gamma/e-beam)

Makrolon® and Apec® resins have a high resistance to the effects of high-energy radiation. Assuming that 28 kGy of energy is required to sterilize Makrolon® or Apec® resins, the resin can be sterilized 10 to 20 times before any appreciable reduction in mechanical strength occurs. Standard Makrolon® and Apec, however, become more yellow with each sterilization cycle. Makrolon® Rx2235, Rx2435, Rx2430, Rx2440, Rx2530, Rx2635, Rx1805 and Rx3440 resins are high-energy radiation stable grades. The color of these grades

shifts to a neutral tint after a typical sterilization dose of high-energy radiation. The degree of color shift depends on the sterilization dose. Makrolon® Rx2440 resin has been designed for devices sterilized by radiation in oxygen-free packaging. Parts made from Bayblend® or Makroblend® resins can become more yellow and exhibit decreased impact resistance after sterilization with high-energy radiation.

Sterilization techniques approved for Texin® Rx TPU

Texin® Rx TPU can be sterilized using ethylene oxide gas, gamma irradiation and E-beam. Gamma sterilization may cause some color shifting. Hot air (dry heat) may also be considered. Do not use steam autoclaving or boiling water methods for sterilization.

CHEMICAL RESISTANCE

Parts molded from Makrolon®, Bayblend®, Apec® and Makroblend® resins are resistant to mineral acids, including a large number of organic acids (e.g., carbonic acid, lactic acid, oleic acid and citric acid), to oxidizing and reducing agents, neutral and acidic saline solutions, a range of greases and oils, saturated aliphatic and cycloaliphatic hydrocarbons and also alcohols, with the exception of methanol. Makrolon®, Bayblend®, Apec® and Makroblend® resins are severely degraded by alkaline/caustic solutions, ammonia gas and its solution and amines. Makrolon®, Bayblend®, Apec® and Makroblend® resins will dissolve in a number of industrial solvents such as dichloromethane or tetrahydrofuran (THF).

Other organic compounds, such as benzene or acetone, cause them to swell. A number of chemical substances may cause stress cracking. Chemical resistance depends both on the internal molded-in stresses and on the external stresses to which the part is exposed (see “[Makrolon® - Stress Crack Test](#)”). In the case of Makrolon® resins, the higher-viscosity grades generally offer better resistance to chemicals. Makroblend® grades can provide even higher levels of chemical resistance.

The following tables list test results after exposure to various chemicals that medical devices may encounter.

As with any compatibility test, the results depend on variables such as concentration, time, temperature, part design and residual stresses, and should serve only as a guideline. It is imperative that production parts be evaluated under actual application conditions prior to commercial use.

In addition to the resins listed on the next page, industrial resins may be used in some applications where neither biocompatibility nor formulation control is required. Some of these grades are listed in the tables at the end of this guide. For more information about the properties of these and other grades, including chemical compatibility, please contact your local Covestro representative for more details. To connect with an expert today, email plastics@covestro.com.



CHEMICAL RESISTANCE

Resins for Medical Devices

Medical devices often come into contact with various substances from medical tubing, drugs, IV-fluids and antiseptics. The table below summarizes the resistance of medical-grade products to examples of these media.

The method used to determine degree of chemical resistance reported in the tables below was

as follows: ISO 527 tensile specimens were continuously exposed to chemicals for 24 hours or repeatedly wiped (10× wet-to-dry) at fixed flexural strains of 1.0% and 0.6%.

Pass criteria: (1) no visible cracking upon close inspection, (2) tensile yield strength fully retained: >98%, and (3) yield behavior preserved with >10% nominal strain at break.

R "resistant": passed evaluation at 1.0% strain

L "limited resistance": passed evaluation at 0.6% strain, failed at 1.0% strain

N "not resistant": failed evaluation at 0.6% strain

Media	Makrolon®			Apec®
	Rx3440	2858	Rx2530	1745
20% lipid solution	R	L	L	L
1% saline solution	R	R	R	R
Deionized water	R	R	R	R
Betadine	R	R	R	R
3% hydrogen peroxide	R	R	R	L



Resistant



Limited resistance



No resistance

RESINS FOR ELECTROMECHANICAL HOUSINGS

Housings encounter a greater variety of substances than medical devices. In addition to media described in the table above, the surfaces of housings are often cleaned with aggressive disinfectants. The following table summarizes the chemical resistance of our medical resins for electro-mechanical devices.¹

Disinfectants	Bayblend®				Makroblend®			Makrolon®			
	M301 FR	M303 FR	M750	M850 XF	M4000 FR	M525	M5005 FR	2458	2858	Rx3440	M6011 FR
CaviCide™	R	R	R	R	R	R	R	R	R	R	R
Clorox Healthcare® Bleach Wipes	R	R	R	R	R	R	R	R	R	R	R
Lysol® Disinfecting Wipes (Lemon & Lime)	R	R	R	R	R	R	R	R	R	R	R
Opti-Cide® ³ Surface Wipes	R	R	R	R	R	R	R	R	R	R	R
Oxivir® Tb	N	N	L	N	R	R	R	L	L	R	R
Sporicidin®	R	R	R	R	R	R	R	R	R	R	R
Super Sani-Cloth® Germicidal Wipe	R	R	R	R	R	R	R	R	R	R	R
Virex® II 256	R	R	R	R	R	R	R	R	R	R	R

R Resistant
L Limited resistance
N No resistance

Disinfection with UVC light

An emerging disinfection technology for housings in hospitals is to use UVC light (wavelength 200 – 280 nm). UV light in this range is absorbed by the pathogen's DNA/RNA, disrupting their cell replication processes and inhibiting their spread. UV light can also affect the appearance and mechanical performance of plastics.

Depending on amount of UVC exposure, the color and mechanical properties of the material may be affected. Preliminary testing has demonstrated that when exposed to a typical cumulative dose of 120 J/m², mechanical properties of our resins are retained. Please consult with your local Covestro representative for more information.

¹ See also the white paper *Compatibility with disinfectants used against SARS-CoV-2*

RESINS FOR WEARABLES

The emergence of wearable medical devices presents unique challenges for material selection. Parts often need to be durable enough to withstand being worn 24 hours a day while being exposed to chemicals such as sunscreen and lotions. The table below summarizes the effect of these substances on our products.

Consumer Products	Bayblend®				Makroblend®			Makrolon®			
	M301 FR	M303 FR	M750	M850 XF	M4000 FR	M525	M5005 FR	2458	2858	Rx3440	M6011 FR
Aveeno® Daily Moisturizing Lotion	L	R	R	L	R	R	R	R	R	R	R
Banana Boat® Sunscreen, Ultra Sport™, SPF 50+	N	L	R	L	L	L	N	R	R	R	R
Goo Gone® Original Goo & Adhesive Remover	N	R	R	N	R	R	R	L	R	R	R
Hand Soap (Softsoap®)	R	R	R	R	R	R	R	R	R	R	R
Isopropanol, 70% (v/v)	R	R	R	R	R	R	R	R	R	R	R
Purell® Advanced Hand Sanitizer	R	R	R	R	R	R	R	R	R	R	R
Sebum (synthetic, ASTM D4265)	L	R	R	L	R	R	R	R	R	R	R
Skin Tac™ Adhesive Barrier Wipes	L	R	R	R	R	R	R	R	R	R	R

R Resistant
 L Limited resistance
 N No resistance

COVESTRO MEDICAL GRADES & REGULATORY INFORMATION

Guidance on Use of Covestro Products in a Medical Application

All Covestro polycarbonate thermoplastics, sheets, and films (herein after “products”) that are designated as “Medical Grade” meet certain biocompatibility test requirements of USP Plastics Class VI and/ ISO 10993 (table on right).

These tests are conducted under Good Laboratory Practices as defined by the FDA in 21 CFR Part 58. Prior to testing, skin contact Medical Grades were treated with alcohol swabs; all other medical products were sterilized by ethylene oxide and gamma radiation. Only Medical Grade products may be considered candidates for applications requiring biocompatibility. No “Medical Grade” product will be available for sale until successfully assessed for biocompatibility.

“Medical Application” means all applications of medical devices wherein the medical device is manufactured with a Covestro product(s) and is intended under normal use to be brought into direct contact with the patient’s body (e.g., skin, body fluids or tissues, including indirect contact to blood).

Covestro products designated as “Medical Grade” shall not be considered candidates for bioabsorbable or long-term (greater than 30 days) implant applications unless Covestro explicitly agrees, in writing, to sell such products for such applications. The biocompatibility testing referenced above cannot assure the biocompatibility of final or intermediate products made from Covestro products or the suitability of such products for their use in a Medical Application, i.e., the test data cannot be used to conclude that any medical devices manufactured from the Covestro products meet the necessary requirements of ISO Standard 10993. It is the sole responsibility of the manufacturer of final end-use product to conduct all necessary tests (including biocompatibility tests) and inspections and to evaluate the final product under actual end-use requirements.

The designation as “Medical Grade” does not mean that Covestro has determined the product is suitable for use in any particular Medical Application. Covestro makes no representations regarding the suitability of a Covestro product for a particular Medical Application or final end-use product.

A determination that the Covestro product is suitable for use in a particular Medical Application or final end-use product can only be made by the purchaser of the Covestro product who utilizes it in a Medical Application and conducts all necessary testing and evaluation to support such a determination.

Only virgin Medical Grade Covestro products have been assessed for biocompatibility. Any use of regrind, for example, runners from mold flow channels or trim pieces, must be evaluated by the medical device manufacturer for suitability.

“Biological Evaluation of Medical Devices” Selected Tests May Include

- | | |
|-----------------------------|------------------------------------|
| 1. Cytotoxicity | 7. Implantation |
| 2. Sensitization | 8. Hemolysis (Direct and Indirect) |
| 3. Intracutaneous Injection | 9. In-vitro Hemocompatibility |
| 4. Acute Systemic Toxicity | 10. USP Physicochemical Test |
| 5. Pyrogenicity | 11. Heavy Metals Analysis |
| 6. Genotoxicity | (Acid Digest and Extraction) |

COVESTRO MEDICAL GRADES & REGULATORY INFORMATION

Appropriate Use of Covestro Products In a Medical Application

Covestro has not performed clinical medical studies concerning the use of Covestro products. Moreover, Covestro has neither sought nor received approval from the United States Food and Drug Administration (FDA) or other competent authorities from other regions for the use of Covestro products in a Medical Application.



Covestro makes no representations or warranty regarding (and accepts no responsibility for determining) either: (a) the suitability of a Covestro product for a particular Medical Application or final end-use product or (b) the adequacy of any warning relating to a Covestro product or particular Medical Application or final end-use product.

The suitability of a Covestro product in a given end-use environment is dependent upon various conditions, including, without limitation, chemical compatibility, method of manufacture, temperature, part design, sterilization method, residual stresses, and external loads. It is the sole responsibility of the manufacturer of the final end-use product to determine the suitability (including biocompatibility) of all raw materials and components, including any Covestro products, in order to ensure that the final product:

- meets relevant biocompatibility requirements and is otherwise safe for its end-use
- performs or functions as intended
- is suitable for its intended use
- complies with all applicable FDA and other regulatory requirements.

It also is the sole responsibility of the manufacturer of the final end-use product to conduct all necessary tests and inspections and to evaluate the final product under actual end-use requirements and adequately advise and warn purchasers, users and/or learned intermediaries (such as physicians) of pertinent risks and fulfill any post-market surveillance obligations.

Any decision regarding the appropriateness of a particular medical product in a particular clinical or Medical Application should be based on the judgment of the manufacturer, seller, the competent authority, and the treating physician. Covestro cannot weigh the benefits against the risks of a medical device and cannot offer a medical or legal judgment on the safety or efficacy of the use of a Covestro product in a specific Medical Application.

Sterilization: The sterilization method and the number of sterilization cycles a medical device made from Texin® resins resin can withstand will vary depending upon type/grade of product, part design, processing parameters, sterilization temperature and chemical environment. Therefore, the manufacturer must evaluate each device to determine the sterilization method and the number of permissible sterilization cycles appropriate for actual end-use requirements and must adequately advise and warn purchasers and users thereof.

TYPICAL PRODUCT VALUES



TYPICAL PRODUCT VALUES

				Biocompatible Makrolon® Resins for Medical Applications Requiring EtO and Steam Sterilization									
				2258	Rx1851	2458 ¹	Rx1452	2558	2658 ²	2858 ³	3158 ⁴	3258 ⁵	3638
MVR	ISO 1133	300°C/1.2 kg	cm³/10 min	34	23	19	16	14	12	9	6	5	2.0
Molding Shrinkage, Parallel	ISO 294-4	60x60x2 mm; 500 bar	%	0.65	0.70	0.65	0.60	0.65	0.70	0.70	0.70	0.70	0.75
Molding Shrinkage, Normal	ISO 294-4	60x60x2 mm; 500 bar	%	0.65	0.70	0.70	0.65	0.70	0.75	0.75	0.75	0.75	0.8
Tensile Modulus	ISO 527-1, -2	1 mm/min	MPa	2400	2400	2400	2400	2400	2400	2400	2400	2400	2300
Yield Stress	ISO 527-1, -2	50 mm/min	MPa	65	65	65	68	66	66	66	66	66	64
Yield Strain	ISO 527-1, -2	50 mm/min	%	6.0	6.0	6.1	6.0	6.1	6.1	6.1	6.2	6.2	6.6
Notched Izod Impact (3mm)	b.o. ISO 180-A	23°C	kJ/m²	65	65	75	65	80	80	85	70	75	65P
Notched Izod Impact (3mm)	b.o. ISO 180-A	-30°C	kJ/m²	12	12	14	12	14	14	14	15	20	55P(C)
Temperature of Deflection under Load	ISO 75-1, -2	1.80 MPa	°C	124	121	125	120	124	124	125	126	127	132
Temperature of Deflection under Load	ISO 75-1, -2	0.45 MPa	°C	137	134	139	132	136	137	137	138	139	145
Coefficient of Linear Thermal expansion, Parallel	ISO 11359-1, -2	23 to 55°C	10 ⁻⁴ /K	0.65	0.65	0.65	0.65	0.65	0.65	0.65	0.65	0.65	0.7
Coefficient of Linear Thermal Expansion, Transverse	ISO 11359-1, -2	23 to 55°C	10 ⁻⁴ /K	0.65	0.65	0.65	0.65	0.65	0.65	0.65	0.65	0.65	0.65
Density	ISO 1183-1	—	kg/m³	1200	1200	1200	1200	1200	1200	1200	1200	1200	1200

¹Makrolon® 2408 resin also available — equivalent grade without release agent

²Makrolon® 2608 resin also available — equivalent grade without release agent

³Makrolon® 2808 resin also available — equivalent grade without release agent

⁴Makrolon® 3108 resin also available — equivalent grade without release agent

⁵Makrolon® 3208 resin also available — equivalent grade without release agent

TYPICAL PRODUCT VALUES

				Biocompatible Makrolon® Resins for Medical Applications Requiring Gamma and E-beam Sterilization							
				Rx2235	Rx2435	Rx2430	Rx2440	Rx2530	Rx2635	Rx1805	Rx3440
MVR	ISO 1133	300°C/1.2 kg	cm³/10 min	34	23	19	19	15	12	6	4.5
Molding Shrinkage, Parallel	ISO 294-4	60x60x2 mm; 500 bar	%	0.5-0.7 ¹	0.65	0.60	0.60	0.60	0.6-0.8 ¹	0.70	0.70
Molding Shrinkage, Normal	ISO 294-4	60x60x2 mm; 500 bar	%	0.5-0.7 ¹	0.65	0.65	0.65	0.65	0.6-0.8 ¹	0.70	0.70
Tensile Modulus	ISO 527-1, -2	1 mm/min	MPa	2400	2400	2400	2400	2400	2400	2400	2300
Yield Stress	ISO 527-1, -2	50 mm/min	MPa	65	67	67	67	67	67	67	65
Yield Strain	ISO 527-1, -2	50 mm/min	%	5.9	6.1	6.1	6.1	6.1	6.1	6.3	6.4
Notched Izod Impact (3mm)	b.o. ISO 180-A	23°C	kJ/m²	10	75	70	75	70	65	80	80
Notched Izod Impact (3mm)	b.o. ISO 180-A	-30°C	kJ/m²	8	12	14	12	14	12	15	14
Temperature of Deflection under Load	ISO 75-1, -2	1.80 MPa	°C	117	120	122	120	122	121	126	126
Temperature of Deflection under Load	ISO 75-1, -2	0.45 MPa	°C	131	132	134	132	134	135	138	139
Coefficient of Linear Thermal expansion, Parallel	ISO 11359-1, -2	23 to 55°C	10 ⁻⁴ /K	0.65	0.65	0.65	0.65	0.65	0.65	0.65	0.65
Coefficient of Linear Thermal Expansion, Transverse	ISO 11359-1, -2	23 to 55°C	10 ⁻⁴ /K	0.65	0.65	0.65	0.65	0.65	0.65	0.65	0.65
Density	ISO 1183-1	—	kg/m³	1200	1200	1200	1200	1200	1200	1200	1200

¹ Value range based on general practical experience

TYPICAL PRODUCT VALUES

				Biocompatible Low Friction Makrolon® Resins for Medical Applications		
				M204 LF	M402 LF	M404 LF
MVR	ISO 1133	300°C/1.2 kg	cm³/10 min	34	19	19
Molding Shrinkage, Parallel	ISO 294-4	60x60x2 mm; 500 bar	%	0.7	0.7	0.7
Molding Shrinkage, Normal	ISO 294-4	60x60x2 mm; 500 bar	%	0.7	0.7	0.7
Tensile Modulus	ISO 527-1, -2	1 mm/min	MPa	2200	2150	2100
Yield Stress	ISO 527-1, -2	50 mm/min	MPa	60	60	60
Yield Strain	ISO 527-1, -2	50 mm/min	%	5.8	6.1	6.0
Notched Izod Impact (3 mm)	b.o. ISO 180-A	23°C	kJ/m²	50	60	60
Notched Izod Impact (3 mm)	b.o. ISO 180-A	-30°C	kJ/m²	10	12	12
Temperature of Deflection under Load	ISO 75-1, -2	1.80 MPa	°C	122	122	122
Temperature of Deflection under Load	ISO 75-1, -2	0.45 MPa	°C	137	138	137
Coefficient of Linear Thermal Expansion, Parallel	ISO 11359-1, -2	23 to 55°C	10 ⁻⁴ /K	0.65	0.65	0.65
Coefficient of Linear Thermal Expansion, Transverse	ISO 11359-1, -2	23 to 55°C	10 ⁻⁴ /K	0.65	0.65	0.65
Density	ISO 1183-1	—	kg/m³	1200	1200	1200

TYPICAL PRODUCT VALUES

				Biocompatible Glass Filled Makrolon® Resins for Medical Applications					
				M410 GF	M420 GF	M430 GF	M810 GF	M820 GF	M830 GF
MVR	ISO 1133	300°C/1.2 kg	cm³/10 min	12	9	7	4	3.5	3
Molding Shrinkage, Parallel	ISO 294-4	60x60x2 mm; 500 bar	%	0.5	0.3	0.2	0.55	0.3	0.2
Molding Shrinkage, Normal	ISO 294-4	60x60x2 mm; 500 bar	%	0.5	0.5	0.5	0.55	0.55	0.55
Tensile Modulus	ISO 527-1, -2	1 mm/min	MPa	3800	5800	8000	3600	5600	7800
Yield Stress	ISO 527-1, -2	5 mm/min	MPa	80	105	128 ¹	77	100 ¹	121 ¹
Yield Strain	ISO 527-1, -2	5 mm/min	%	3.8	3.0	2.8 ²	4.1	4.0 ²	3.3 ²
Notched Izod Impact	b.o. ISO 180-A	23°C	kJ/m²	7	12	14	12	16	18
Notched Izod Impact	b.o. ISO 180-A	-30°C	kJ/m²	6	11	13	10	14	16
Temperature of Deflection under Load	ISO 75-1, -2	1.80 MPa	°C	135	138	140	136	140	142
Temperature of Deflection under Load	ISO 75-1, -2	0.45 MPa	°C	142	143	144	145	146	147
Coefficient of Linear Thermal Expansion, Parallel	ISO 11359-1, -2	23 to 55°C	10 ⁻⁴ /K	0.4	0.3	0.3	0.4	0.3	0.2
Coefficient of Linear Thermal Expansion, Transverse	ISO 11359-1, -2	23 to 55°C	10 ⁻⁴ /K	0.4	0.4	0.4	0.4	0.4	0.4
Density	ISO 1183-1	—	kg/m³	1270	1340	1420	1270	1340	1420

¹Stress at break

²Yield at break

For more information on the grades listed above, check the [Product Finder](#) on the Covestro Solution Center

TYPICAL PRODUCT VALUES

				Biocompatible High Heat Apec® for Medical Applications	
				1745	2045
MVR	ISO 1133	330°C/2.16 kg	cm³/10 min	17	8
Molding Shrinkage, Parallel	ISO 294-4	60x60x2 mm; 500 bar	%	0.8	1.05
Molding Shrinkage, Normal	ISO 294-4	60x60x2 mm; 500 bar	%	0.8	1.05
Tensile Modulus	ISO 527-1, -2	1 mm/min	MPa	2400	2400
Yield Stress	ISO 527-1, -2	50 mm/min	MPa	70	75
Yield Strain	ISO 527-1, -2	50 mm/min	%	6.8	6.8
Charpy impact strength	ISO 179/1eU	23°C	kJ/m²	Non-Break	Non-Break
Charpy impact strength	ISO 179/1eU	-30°C	kJ/m²	Non-Break	Non-Break
Temperature of Deflection under Load	ISO 75-1, -2	1.80 MPa	°C	148	173
Temperature of Deflection under Load	ISO 75-1, -2	0.45 MPa	°C	160	192
Coefficient of Linear Thermal Expansion, Parallel	ISO 11359-1, -2	23 to 55°C	10⁻⁴/K	0.65	0.65
Coefficient of Linear Thermal Expansion, Transverse	ISO 11359-1, -2	23 to 55°C	10⁻⁴/K	0.65	0.65
Density	ISO 1183-1	—	kg/m³	1170	1130

TYPICAL PRODUCT VALUES

				Biocompatible Resins and Blends for Medical Applications							
				Makrolon®	Makroblend®			Bayblend®			
				M6011 FR ¹	M4000 FR ¹	M5005 FR ¹	M525 ¹	M301 FR ¹	M303 FR ¹	M750 ²	M850XF ²
MVR	ISO 1133	300°C/5 kg	cm ³ /10 min	25	—	—	—	—	—	—	—
MVR	ISO 1133	240°C/5 kg	cm ³ /10 min	—	—	—	—	25	—	—	—
MVR	ISO 1133	260°C/5 kg	cm ³ /10 min	—	18	35	21	—	11	11	25
Molding Shrinkage, Parallel	b.o. ISO 2577	Value range based on practical experience	%	0.6-0.8	0.7 - 0.9	0.7 - 0.9	0.7 - 0.9	0.5 - 0.7	0.5 - 0.7	0.7 - 0.9	0.55-0.75
Molding Shrinkage, Normal	b.o. ISO 2577	Value range based on practical experience	%	0.6-0.8	0.7 - 0.9	0.7 - 0.9	0.7 - 0.9	0.5 - 0.7	0.5 - 0.7	0.7 - 0.9	0.55-0.75
Tensile Modulus	ISO 527-1, -2	1 mm/min	MPa	2300	2300	2300	2000	2600	2650	2000	2500
Yield Stress	ISO 527-1, -2	50 mm/min	MPa	61	56	55	55	60	69	47	62
Yield Strain	ISO 527-1, -2	50 mm/min	%	5.9	4.3	4	4.5	4	5	4.8	4.9
Notched Izod Impact	ISO 180-A	23°C	kJ/m ²	70	40	45	60	35	40	45	48
Notched Izod Impact	ISO 180-A	-30°C	kJ/m ²	20	12	13	20	8	10	35	15
Temperature of Deflection under Load	ISO 75-1, -2	1.80 MPa	°C	113	85	77	75	85	98	104	109
Temperature of Deflection under Load	ISO 75-1, -2	0.45 MPa	°C	128	115	89	100	95	115	127	127
Coefficient of Linear Thermal Expansion, Parallel	ISO 11359-1, -2	23 to 55°C	10 ⁻⁴ /K	0.7	0.8	0.9	0.9	0.8	0.7	0.85	0.7
Coefficient of Linear Thermal Expansion, Transverse	ISO 11359-1, -2	23 to 55°C	10 ⁻⁴ /K	0.7	0.8	0.9	0.9	0.8	0.7	0.85	0.7
Density	ISO 1183-1	—	kg/m ³	1190	1340	1250	1220	1190	1190	1120	1140
Burning behavior UL 94 [UL recognition]	UL94	V-0	mm	1.5	2.0	2.4	—	1.5	1.5	—	—
Burning behavior UL 94 [UL recognition]	UL94	5VA	mm	—	3.0	—	—	3.0	3.0	—	—

¹Skin contact biocompatibility for uncompromized skin

²Biocompatible for non-implant applications

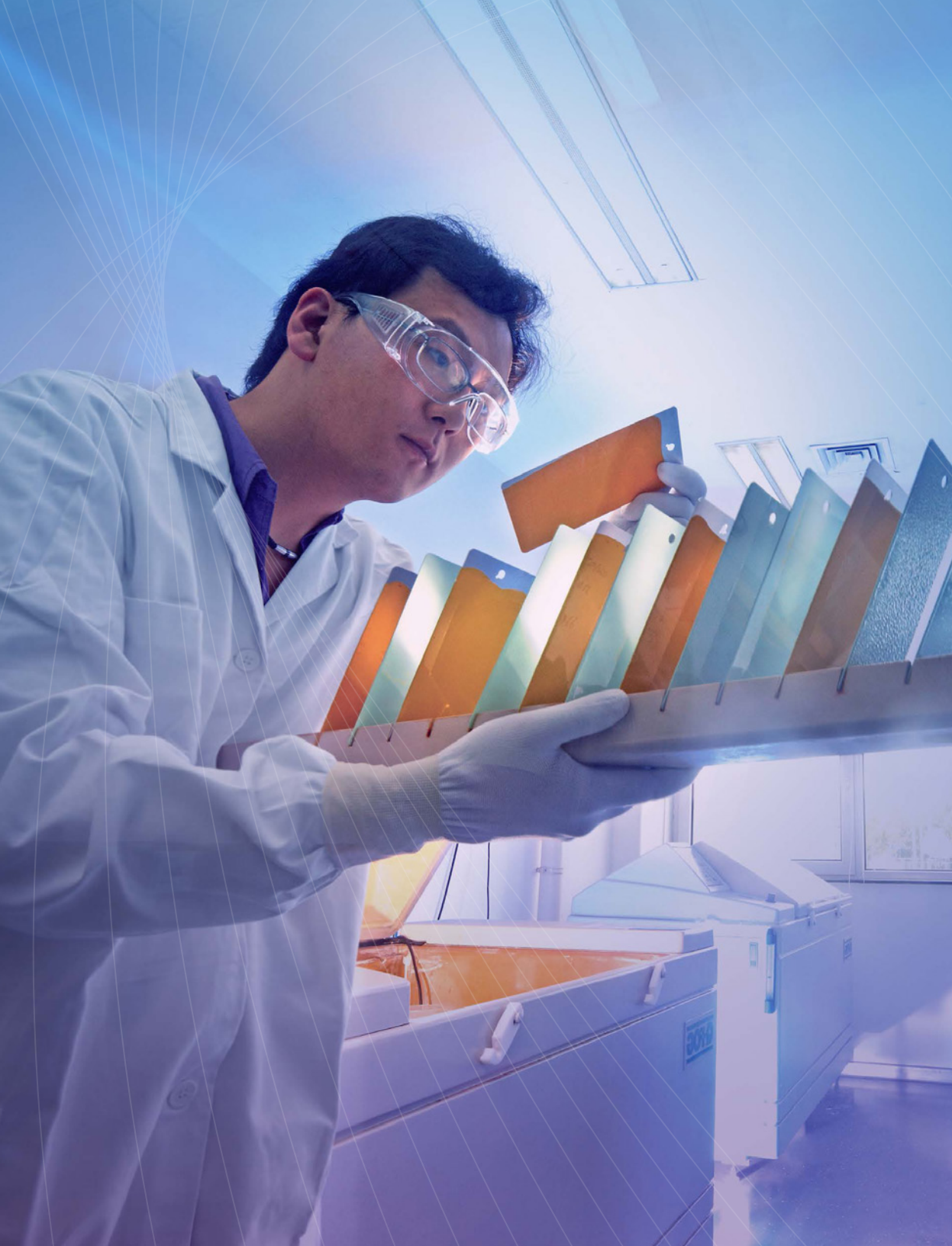
For more information on the grades listed above, check the [Product Finder](#) on the Covestro Solution Center

TYPICAL PRODUCT VALUES

				General Purpose Resins for Medical Applications (Non-biocompatible)		
				Bayblend®		
				FR3010	FR3010 HF	FR3005 HF
MVR	ISO 1133	240°C/5 kg	cm³/10 min	15	25	40
Molding Shrinkage, Parallel	b.o. ISO 2577	Value range based on practical experience	%	0.5-0.7	0.5-0.7	0.5-0.7
Molding Shrinkage, Normal	b.o. ISO 2577	Value range based on practical experience	%	0.5-0.7	0.5-0.7	0.5-0.7
Tensile Modulus	ISO 527-1, -2	1 mm/min	MPa	2700	2600	2700
Yield Stress	ISO 527-1, -2	50 mm/min	MPa	60	60	60
Yield Strain	ISO 527-1, -2	50 mm/min	%	4	4	3.5
Notched Izod Impact	b.o. ISO 180-A	23°C	kJ/m²	35	35	13
Notched Izod Impact	b.o. ISO 180-A	-30°C	kJ/m²	10	10	8.0
Temperature of Deflection Under Load	ISO 75-1, -2	1.80 MPa	°C	90	90	81
Temperature of Deflection Under Load	ISO 75-1, -2	0.45 MPa	°C	100	100	90
Coefficient of Linear Thermal Expansion, Parallel	ISO 11359-1, -2	23 to 55°C	10 ⁻⁴ /K	0.76	0.76	0.76
Coefficient of Linear Thermal Expansion, Normal	ISO 11359-1, -2	23 to 55°C	10 ⁻⁴ /K	0.8	0.8	0.8
Density	ISO 1183-1	—	kg/m³	1180	1180	1180
Burning behavior UL 94 [UL recognition]	UL94	V0	mm	1.5	1.5	1.5
Burning behavior UL 94 [UL recognition]	UL94	5VA	mm	3.0	3.0	3.0

TYPICAL PRODUCT VALUES

			Biocompatible High-Performance Thermoplastic Polyurethanes for Healthcare Applications								
			Texin®								
			RxT70A	RxT80A	RxT85A	RxT90A	RxT50D	RxT65D	RxT76D	RxS285	RxS292
Shore Hardness	ISO 868	—	70A	84A	85A	90A	50D	65D	76D	85A	92A
Tensile Stress at 100% Elongation	ISO 37	MPa	3.4	5.3	5.5	7.6	14.0	25.0	29.0	5.3	9.7
Tensile Strength	ISO 37	MPa	26	27	37	41	48	52	46	38	40
Tear Strength	ISO 34	kN/m	60	88	88	136	131	210	257	88	129
Vicat A	ISO 306	°C	75	80	80	106	128	138	133	91	88
Flex Modulus	ISO 178	MPa	14	27	27	41	114	421	1467	28	73
Taber Abrasion	ISO 4649	mg loss	7	30	30	25	75	75	62	35	27
Clarity	—	—	Good	Excellent	Excellent	Excellent	Excellent	Excellent	Good	Excellent	Excellent



APPLICATION DEVELOPMENT

Our Capabilities:
Engineering and Application
Development Support

- Material selection
- CAE design and analysis
- Mold and injection system design, analysis
- Component and system testing
- Part and mold design reviews
- On-site technical support
- Advanced processing feasibility and support
- Concept and feasibility studies

ADDITIONAL RESOURCES

Access these and other downloads on the [Covestro Solutions Center](#)

POLYCARBONATES IN HEALTHCARE APPLICATIONS

- Adhesion of Thermoplastic elastomers (TPEs) on polycarbonates for medical wearables
- Balanced filling in thermoplastic medical molding
- Compatibility with disinfectants used against SARS-CoV-2
- Enhancing impact resistance and toughness in molded medical parts
- Eye protection for welders
- Influence of UVC LED disinfection on polycarbonate materials
- Photo-elastic stress analysis of polycarbonate medical parts
- Thermoplastic polyurethanes for medical applications
- Understanding flow hesitation in molded medical parts
- Radiopacity of Makrolon®

Injection Molding & Mold Design

- Calculating the mold-filling process for thin-walled injection moldings
- Determining the dryness of Makrolon® by the TVI test
- Gate design for high-quality surface finish
- Injection molding - production equipment and machinery
- Injection molding of high-quality molded parts-drying
- Optimized mold temperature control
- Overmolding with polycarbonate and polycarbonate blends
- Part and mold design brochure
- Process variables - injection molding
- Purging compounds for use when molding thermoplastics
- Shrinkage and deformation of glass fiber reinforced thermoplastics
- The fundamentals of shrinkage in thermoplastics
- Understanding and optimizing weld lines in thermoplastic molding

Secondary Options

- Joining techniques design guide
- Laser marking thermoplastics
- Laser transmission welding
- Marking products made of technical thermoplastics
- Self-tapping screws for thermoplastics
- The insertion of connecting elements using ultrasound

Part Performance

- Addressing molded-in stresses and part durability
- Environmental stress cracking - Bend strip test
- Makrolon® - stress crack test
- Makrolon® - chemical resistance

Part Design

- Design with Makrolon® thermally conductive polymers
- Designing with light
- Snap-fit joints for plastic - A design guide

Material Selection

- Makrolon® for LED lighting
- Makrolon® for optical data storage
- Materials and packaging solutions for energy storage systems used in electromobility
- Optical properties of Makrolon® and Apec®
- Polycarbonate and polycarbonate blends for the electrical and electronics industries
- Shaping LED diffuser performance



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