



## MD<sup>®</sup> 1-CN0029

### LED Optimized Catheter Assembly Adhesive

#### APPLICATIONS

- Balloon/Lumen
- Hub/Lumen
- Manifold Bond Joint

#### FEATURES

- LED Optimized 385 nm
- Adhesion to a range of difficult to bond substrates

#### RECOMMENDED SUBSTRATES

- Nylon 12
- PEBA
- PC
- ABS
- TPU

#### BIOCOMPATIBILITY

- ISO 10993-4 Hemolysis
- ISO 10993-5 Cytotoxicity
- ISO 10993-6 Implantation
- ISO 10993-10 Intracutaneous
- ISO 10993-11 Systemic Toxicity

Dymax MD<sup>®</sup> 1-CN0029 is an LED Optimized for light curing at 385nm. Designed for bonding and assembly of next-generation catheter designs using Nylon 12, PEBA and other difficult to bond substrates. Dymax MD<sup>®</sup> Medical Device Adhesives contain no nonreactive solvents and cure upon exposure to light. Their ability to cure in seconds enables faster processing, greater output, and lower processing costs. When cured with Dymax light-curing spot lamps, focused-beam lamps, or flood lamps, they deliver optimum speed and performance for medical device assembly. Dymax lamps offer the optimum balance of UV and visible light for the fastest, deepest cures. This product is in full compliance with RoHS2 directives 2015/863/EU and 2011/65/EU.

#### UNCURED PROPERTIES \*

Property	Value	Test Method
Solvent Content	No Nonreactive Solvents	N/A
Chemical Class	Acrylated Urethane	N/A
Appearance	Light yellow Transparent	N/A
Soluble in	Organic Solvents	N/A
Viscosity, cP (20 rpm)	295 (nominal)	DSTM 520
Shelf Life @RT (22°C to 25°C) from Date of Manufacture	8 months	N/A

#### OTHER CURED PROPERTIES \*

Property	Value	Test Method
Boiling water absorption, % (2h)	1.03	ASTM D570
Water Absorption, % (25°C, 24 h)	0.87	ASTM D570
Linear Shrinkage, %	0.24	ASTM D2566
Glass Transition Tg, °C	59.12	ASTM D5418
CTE <sub>α1</sub> , μm/m/°C	20.78	DSTM 610 <sup>‡</sup>
CTE <sub>α2</sub> , μm/m/°C	317.48	DSTM 610 <sup>‡</sup>

#### CURED MECHANICAL PROPERTIES \*

Property	Value	Test Method
Durometer Hardness	D61	ASTM D2240
Tensile at Break, MPa [psi]	10.6 [1,538]	ASTM D638
Elongation at Break, %	363	ASTM D638
Modulus of Elasticity, MPa [psi]	255 [36,985]	ASTM D638

#### ADHESION

Substrate	Recommendation
ABS	✓
PC	✓
PVC	✓
Stainless Steel	✓
PMMA	✓
PS	✓
PA66	✓
TPU	✓
PBT	o
PEI	o

✓ Recommended      o Limited Applications  
st Requires Surface Treatment (e.g. plasma, corona treatment, etc.)

\* Not Specifications

N/A Not Applicable

‡ DSTM Refers to Dymax Standard Test Method

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## CURING GUIDELINES

Dymax Curing System (Intensity)	Fixture Time or Belt Speed <sup>A</sup>
BlueWave® 200 (9.5 W/cm <sup>2</sup> ) <sup>B</sup>	0.3 s
BlueWave® MX-150 385 nm PrimeCure® (15 W/cm <sup>2</sup> ) <sup>C</sup>	1 s

Fixture time is defined as the time to develop a shear strength of 0.1 N/mm<sup>2</sup> [10 psi] between glass slides. Actual cure time typically is 3 to 5 times fixture time.

A. Fixture times/belt speeds are typical for curing thin films through 100% UV and light-transmitting substrates. Light-obstructing substrates may require longer cure time

B. Intensity was measured over the UVA range (320-395 nm) using a Dymax ACCU-CAL™ 50 Radiometer.

C. Intensity was measured over the UVA/Visible range (350-450 nm) using a Dymax ACCU-CAL™ 50-LED Radiometer.

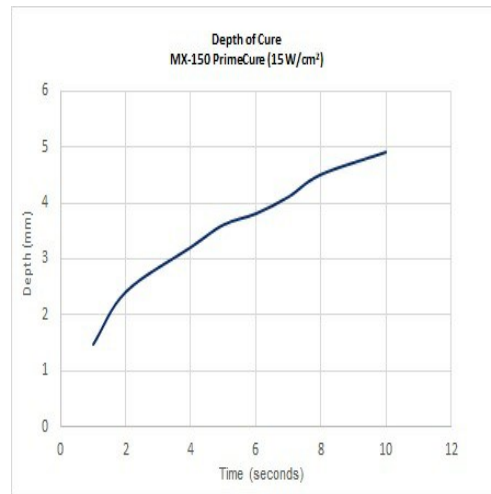
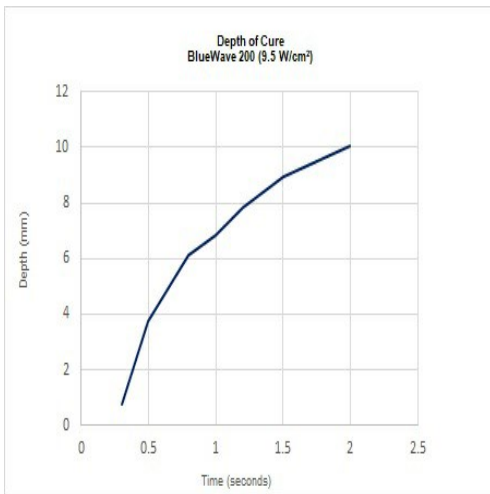
Full cure is best determined empirically by curing at different times and intensities, and measuring the corresponding change in cured properties such as tackiness, adhesion, hardness, etc. Full cure is defined as the point at which more light exposure no longer improves cured properties. Higher intensities or longer cures (up to 5x) generally will not degrade Dymax light-curable materials.

Dymax recommends that customers employ a safety factor by curing longer and/or at higher intensities than required for full cure. Although Dymax Application Engineering can provide technical support and assist with process development, each customer ultimately must determine and qualify the appropriate curing parameters required for their unique application.

## DEPTH OF CURE

The graphs below show the increase in depth of cure as a function of exposure time with two different lamps at different intensities.

A 9.5 mm [0.37 in] diameter specimen was cured in a polypropylene mold and cooled to room temperature. It was then released from the mold and the cure depth was measured.





### OPTIMIZING PERFORMANCE AND HANDLING

1. This product cures with exposure to UV and visible light. Exposure to ambient and artificial light should be kept to a minimum before curing. Dispensing components including needles and fluid lines should be 100% light blocking, not just UV blocking.
2. All bond surfaces should be clean and free from grease, mold release, or other contaminants prior to dispensing the adhesive.
3. Cure speed is dependent upon many variables, including lamp intensity, distance from the light source, required depth of cure, bond gap, and percent light transmission of the substrate.
4. Oxygen in the atmosphere may inhibit surface cure. Surfaces exposed to air may require high-intensity UV light to produce a dry surface cure. Flooding the bond area with an inert gas, such as nitrogen, can also reduce the effects of oxygen inhibition.
5. Parts should be allowed to cool after cure before testing and subjecting to any loads or electrical testing.
6. In rare cases, stress cracking may occur in assembled parts. Three options may be explored to eliminate this problem. One option is to heat anneal the parts to remove molded-in stresses. A second option is to open the gap between mating parts to reduce stress caused by an interference fit. The third option is to minimize the amount of time the liquid adhesive remains in contact with the substrate(s) prior to curing.
7. Light curing generally produces some heat. If necessary, cooling fans can be placed in the curing area to reduce the heating effect on components.
8. At the point of curing, an air exhaust system is recommended to dissipate any heat and vapors formed during the curing process.

### DISPENSING THE RESIN

This material may be dispensed with a variety of manual, semi-automated, and fully automated fluid delivery systems. Small area applications, including beads and small dots, can be achieved using hand-held Dymax dispensing systems like our SD-100 syringe dispenser and our Model 400 needle valve systems. The valve system can be used in a manual, semi-automated or fully automated system. Dymax has several other dispensing systems that may be suitable for use with our materials. Questions relating to and defining the best fluid delivery system and curing equipment for specific applications should be discussed with the Dymax Application Engineering Team.

### STORAGE AND SHELF LIFE

Store the material in a cool, dark place when not in use. Do not expose to light. This product may polymerize upon prolonged exposure to ambient and artificial light. Keep covered when not in use. This material has a 240-day shelf life from date of manufacture, unless otherwise specified, when stored between 10°C [50°F] and 32°C [90°F] in the original, unopened container.

### STERILIZATION

Compatible sterilization methods include gamma irradiation and ethylene oxide. Sterilization by autoclaving may be limited to certain applications. It remains the user's obligation to ascertain the effect of sterilization on the cured adhesive.

### CLEANUP

Uncured material may be removed from dispensing components and parts with organic solvents. Cured material will be impervious to many solvents and difficult to remove. Cleanup of cured material may require mechanical methods such as ultrasonic bath, water jet, vacuum tweezers, air knife and/or warming to aid in the removal.

### BIOCOMPATIBILITY

Polymerized Dymax MD® Medical Device Adhesives are biocompatibility tested in accordance with ISO 10993 and/or USP Class VI. The completed tests are listed on each product data sheet. Copies of the test reports are available upon request. In all cases, it is the user's responsibility to determine and validate the suitability of these adhesives in the intended medical device. These adhesives have not been tested for prolonged or permanent implantation, and are only intended for use in short-term (<29 days) or single-use disposable-device applications. Dymax does not authorize their use in long-term implant applications. Customers using these materials for such applications do so at their own risk and take full responsibility for ensuring product safety and biocompatibility.



## GENERAL INFORMATION

This product is intended for industrial use only. Keep out of the reach of children. Avoid breathing vapors. Avoid contact with skin, eyes, and clothing. Wear impervious gloves. Repeated or continuous skin contact with uncured material may cause irritation. Remove material from skin with soap and water. Never use organic solvents to remove material from skin and eyes. For more information on the safe handling of this material, please refer to the Safety Data Sheet before use.

The data provided in this document are based on historical testing that Dymax performed under laboratory conditions as they existed at that time, and are for informational purposes only. The data are neither specifications nor guarantees of future performance in a particular application. Dymax does not guarantee that this product's properties are suitable for the user's intended purpose.

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