

Injection Molding of MAGNUM™ 2642 MED ABS Resin

Drying Recommendations

Moisture Content: It is recommended to dry this resin to less than 0.02% moisture.

Dryer: A desiccant-type dryer is recommended for this material. Hopper dryer may also be needed.

Dew Point: -20°F (-30°C) or lower is desired.

Drying Time: A minimum of 2 hours at 180-185°F (82-85°C) is recommended.

Equipment Specifications

Shot Size: As required to fill part.

Clamp Tonnage: 2.0-3.0 tons/in² (0.28-0.42 tonnes/cm²) of projected surface area.

Barrel Capacity: Minimum of three times shot size, but no greater than six times shot size (i.e.: if shot is 10 oz., barrel capacity should be 30-40 oz., but no greater than 60 oz.). For short barrel machines, if shot size exceeds 50% of barrel capacity, increase "feed" and "transition" barrel temperatures by 15°F (8°C).

Process Conditions

Melt Temperature: Can be varied as required to fill part. See Table 1.

Mold Temperature: Can be changed to affect gloss or demolding characteristics. See Table 1.

Manifold/Drop Temperature: Range of 15-25°F (8-14°C), aim of 20°F (11°C) below melt temperature. See Table 1 for recommended melt temperatures and adjust manifold/drop temperature accordingly.

Purging/Shutdown: For short interruptions in production, empty the barrel and reduce barrel temperatures. For extended shutdowns, purge the machine with a purging compound or general purpose polystyrene before shutting down machine.

Table 1: Machine Temperature Recommendations

Temperatures	Range, °F (°C)	Aim, °F (°C)
Melt Temperature	450-480 (232-249)	465 (241)
Mold Temperature	80-120 (27-49)	100 (38)
Barrel – Feed	360-390 (182-199)	375 (191)
Barrel – Transition	410-440 (210-227)	425 (218)
Barrel – Metering	435-465 (224-241)	450 (232)
Nozzle	445-475 (229-246)	465 (241)

These are typical properties only and are not to be construed as specifications. Users should confirm results by their own tests.



The principles of Responsible Care® and Sustainable Development influence the production of printed literature for Styron LLC, Styron Hold Co B.V., and subsidiaries (“Styron”). As a contribution toward the protection of our environment, Styron’s printed literature is produced in small quantities on paper containing recovered/post-consumer fiber and using 100 percent soy-based ink whenever possible.

PRODUCT STEWARDSHIP

Styron and its subsidiaries have a fundamental concern for all who make, distribute, and use its products and for the environment in which we live. This concern is the basis for our Product Stewardship philosophy by which we assess the safety, health and environmental information on our products to protect employee and public health and our environment. The success of our product stewardship program rests with each and every individual involved with Styron products – from the initial concept and research, to manufacture, use, sales, disposal and recycle of each product.

CUSTOMER NOTICE

Customers are responsible for reviewing their manufacturing processes and their applications of Styron products from the standpoint of human health and environmental quality to ensure that Styron products are not used in ways for which they are not suitable. Styron personnel are available to answer questions and to provide reasonable technical support. Styron product literature, including safety data sheets, should be consulted prior to the use of Styron products. Current safety data sheets are available from Styron.

NOTICE REGARDING MEDICAL APPLICATION RESTRICTIONS

Styron requests that customers considering the use of Styron products in medical applications notify Styron so that appropriate assessments may be conducted.

Styron does not endorse or claim suitability of its products for specific medical applications and does not knowingly sell or sample its products or services for certain medical applications.

It remains the responsibility of the medical device or pharmaceutical manufacturer to determine that the Styron product is safe, lawful and technically suitable for the intended use.

GENERAL NOTICE

Any photographs of end-use applications in this document represent potential end-use applications but do not necessarily represent current commercial applications, nor do they represent an endorsement by Styron of the actual products. Further, these photographs are for illustration purposes only and do not reflect either an endorsement or sponsorship of any other manufacturer for a specific potential end-use product or application, or for Styron, or for specific products manufactured by Styron.

If products are described as “experimental” or “developmental”: (1) product specifications may not be fully determined; (2) analysis of hazards and caution in handling and use are required; (3) there is greater potential for Styron to change specifications and/or discontinue production; and (4) although Styron may from time to time provide samples of such products, Styron is not obligated to supply or otherwise commercialize such products for any use or application whatsoever.

For more information on products, innovations, expertise, and other services available from Styron, visit www.styron.com, or contact us as indicated below.

North America

U.S. & Canada	+1-888-STYRON1 +1-989-633-1718
Mexico	+1-800-441-4369

Latin America

Argentina	+54-11-4319-0100
Brazil	+55-11-2142-2355
Colombia	+57-1-219-6000
Mexico	+52-55-5201-4700

Europe/Middle East

	+800-444-11-444 +32-3-450-2967
Germany	+800-181-1361

Asia Pacific

	+800-7776-7776 +603-7965-5319
--	----------------------------------

www.styron.com

DISCLAIMER

THE INFORMATION PROVIDED IN THIS DOCUMENT IS BASED UPON THE INFORMATION AVAILABLE AT THE TIME THIS DOCUMENT WAS PREPARED AND IS SUBJECT TO CHANGE. WHILE STYRON WILL COMMIT TO MEET PRODUCT SPECIFICATIONS, IT IS THE RESPONSIBILITY OF THE MEDICAL DEVICE OR PHARMACEUTICAL MANUFACTURER TO DETERMINE THE SUITABILITY OF THE PARTS AND RAW MATERIALS, INCLUDING STYRON PRODUCTS, USED IN FINAL PRODUCTS TO ENSURE SAFE, SUITABLE, LAWFUL AND TECHNICAL COMPLIANCE FOR THE INTENDED END USE. STYRON MAKES NO WARRANTIES, EXPRESS OR IMPLIED, CONCERNING THE SUITABILITY OF ANY STYRON PRODUCT FOR USE IN MEDICAL APPLICATIONS. STYRON DISCLAIMS ANY AND ALL LIABILITY FOR LOSSES OR DAMAGES THAT MAY RESULT FROM THE USE OF STYRON PRODUCTS IN MEDICAL APPLICATIONS. STYRON MAKES NO WARRANTIES, EXPRESS OR IMPLIED, THAT THE USE OF ANY STYRON PRODUCT, INCLUDING, WITHOUT LIMITATION, USE IN MEDICAL APPLICATIONS, WILL BE FREE FROM ANY INFRINGEMENT CLAIMS.

For additional information not covered by the content of this document, please refer to the Customer Information Group contact information on our web site at www.styron.com/contact/. To ensure you have the latest version of this document available, please contact our Customer Information Group at www.styron.com/contact/.

Published 1/29/12. Printed in North America.
©2011 Styron LLC, Styron Hold Co B.V., and subsidiaries

™Trademark

®Responsible Care is a service mark of the American Chemistry Council.