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FOR IMMEDIATE RELEASE

Phillips Plastics Corporation® Receives EU GMP Registration

Menomonie, Wisconsin – Phillips Plastics Corporation recently received the European Union's Good Manufacturing Practices (EU GMP) Certification for a high volume drug product currently manufactured at its Medical Molding and Assembly facility. To achieve the certification, a superior level of performance was required in order to comply with the strict rules of Europe's Medicine and Healthcare Products Regulatory Agency (MHRA). A very unique distinction for contract manufacturing, the EU GMP certification will allow Phillips' combination product and drug delivery device customers to tap directly into the European pharmaceutical markets. As explained by Dave Thoreson, Plant Manager at Medical Molding and Assembly, "This EU GMP Certification, combined with our 21 CFR 820 and 21 CFR 210 and 211 registrations allow us to provide world class combination product capabilities to our customers. This new certification opens up a new realm of business possibilities for Phillips and its ability to comply with European regulations in the pharmaceutical market."

About Phillips Plastics

Phillips Plastics Corporation is a forty-three year-old custom injection molder of plastic and metal with annual sales of over \$260 million. The Company employs 1,600 people in 15 locations throughout the United States, occupying over 620,000 square feet, including design centers in Wisconsin and California, and provides complete services from concept design, rapid prototyping, and tooling through production, assembly, packaging and distribution to virtually every market. www.phillipsplastics.com

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