

# Certificate of Registration

# Intertek

This is to certify that the quality management system of

## Medical Systems

**Main Site: 3715 Laird Road, Unit #13, Mississauga, Ontario L5L 0A3 Canada**

has been assessed and registered by Intertek, a **CMDCAS recognized registrar**, as conforming to the requirements of

## ISO 13485:2003

The quality management system is applicable to

Manufacturing and distribution of products listed below which are applicable to health care institutions and to OEM products: sterile plastic transfer sets and extension sets in various configurations utilizing standard bore, minibore and microbore tubing sizes with and without male/female luer lock connectors, luer slip connectors, back check valves, needleless valves, injection ports, vented or nonvented infusion chambers with or without solution filter, stopcocks and ganged manifolds, drainage bags and related accessories. Chemotherapy reconstruction devices; and a variety of kits with or without the above listed transfer sets together with accessories such as syringes, hypodermic needles, blades, scalpels, plastic drapes, table covers, plastic equipment covers, wound dressing, gowns, gloves, dressing, steristrip, conductivity gel, needles with blades, surgical skin marker, bowls, sponge with forceps, labels, needles, hand towels, for the areas of general hospital requirements, cardiac cath lab, angiographic department, radiology, urology, vein clinics, operating room and general surgery.

Certificate Number: 9394-9  
Initial Certification Date: 18 August 2006  
Certificate Effective Date: 18 August 2015  
Certificate Expiry Date: 17 August 2018



*Calin Moldovean, President*

*Intertek Testing Services NA, Ltd. – 1829, 32nd avenue, Lachine, QC, H8T 3J1, Canada*



In the issuance of this certificate, Intertek assumes no liability to any party other than to the Client, and then only in accordance with the agreed upon Certification Agreement. This certificate's validity is subject to the organization maintaining their system in accordance with Intertek's requirements for systems certification. Validity may be confirmed via email at [certificate.validation@intertek.com](mailto:certificate.validation@intertek.com) or by scanning the code to the right with a smartphone.

The certificate remains the property of Intertek, to whom it must be returned upon request.

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