



CERTIFICATE OF APPROVAL

This is to certify that the Quality Management System of:

Immucor, Inc.
3130 Gateway Drive
Norcross, Georgia 30071, USA

has been approved by Lloyd's Register Quality Assurance
to the following Quality Management System Standards:

ISO 13485:2003

The Quality Management System is applicable to:

Design, Manufacture and Technical Support of In Vitro Diagnostic Reagents and Design, Manufacture and Servicing of Instruments/Analyzers used for Diagnosis or Management of Blood Grouping, Compatibility Testing, Transmissible Agents, Prenatal and Donor Screening. The In Vitro Diagnostic Reagents include Monoclonal, Polyclonal and Lectin Blood Grouping Reagents for Direct and Indirect Hemagglutination, Anti-Human Globulin Reagents and Control Red Cells, Reagent Red Blood Cells for Serum Grouping and Detection/Identification of Red Cell Antibodies, Bovine Albumin and Potentiating/Enhancement Reagents, Enzyme Solutions, Phosphate Buffer and Red Cell/Platelet Preservative Solutions, Reagents for Red Cell and Platelet Antibody Removal, Fetal Bleed Screening, Capture Assays and Components for the Detection/Identification of Red Cell, Platelet, Syphilis and CMV Antibodies and Control Reagents for ABO, Rh and Red Cell Reagents. The In Vitro Diagnostic Instrumentation includes Immunohematology Instruments/Analyzers Intended for use with Immucor In Vitro Diagnostic Reagents.

Approval
Certificate No: UQA 0113238/B

Original Approval: February 13, 2003
Effective Date: March 1, 2009
Certificate Expiry: February 28, 2012

Cheryl Distefano

Cheryl Distefano, Client Services Certificates Coordinator
Issued by Lloyd's Register Quality Assurance, Inc.



This document is subject to the provision on the reverse
1401 Enclave Parkway, Suite 200, Houston, Texas 77077, USA

This approval is carried out in accordance with the LRQA assessment and certification procedures and monitored by LRQA.

Macro Revision 13