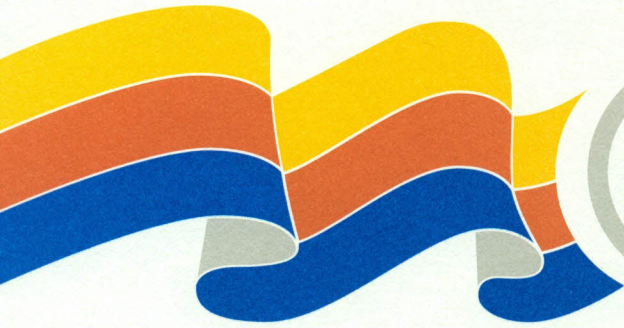


CERTIFICATE OF REGISTRATION



**Quality
System
Registrar**



Having been audited in accordance with requirements of

ISO 13485:2003

SRI Quality System Registrar, 300 Northpointe Circle, Seven Fields, Pennsylvania, 16046, USA, hereby grants to:

AceCo Precision Manufacturing

Registration of the management system at its location:

**4095 S. Gekeler Lane
Boise, Idaho, 83716, USA**

The conditions for maintaining this certificate of registration are set forth in the SRI registration agreements R20.3 and R20.4. Further clarifications regarding the scope of this certificate and the applicability of ISO 13485:2003 requirements may be obtained by consulting the organization.

Scope of ISO 13485:2003 registration: "Manufacture of non-active implantable devices, and general non-active, non-implantable medical devices, such as cuff repair plates and machining of complex geometrics and features of implants, and instruments."

Exclusions: Design and Development; Control of Production and Service Provision - Installation Activities; Control of Production and Service Provision - Servicing Activities; Control of Production and Service Provision - Particular Requirements for Sterile Medical Devices; Validation of Processes for Production and Service Provision - Particular Requirements for Sterile Medical Devices

Initial SRI registration date: March 31, 2009

Current registration period: March 30, 2015 through March 29, 2018

Signed for SRI:



Christopher H. Lake, President & COO

Release Date: March 30, 2015
Certificate Number: 015359
Registration Number: 3753-01

