

DUTCH ACCREDITATION COUNCIL RvA



PO Box 2768 NL-3500 GT Utrecht

The Dutch Accreditation Council RvA, operating as accreditor for certification bodies,
hereby declares that

**Orion Registrar Inc.
Arvada (Colorado)
USA**

has demonstrated to operate management system certification in a competent,
consistent and impartial manner.

This statement is based on an assessment against the requirements specified in
ISO/IEC 17021:2006 and in the RvA regulations.

The accreditation covers the certification schemes and working areas specified in the
authorized annex bearing the accreditation number.

The accreditation is valid provided that the certification body demonstrable continues
to meet the requirements.

This accreditation statement with accreditation number:

C 147

is granted on 25 June 2008 and is valid until

30 August 2010

The accreditation has been granted for the first time on

30 August 1995

for: ISO/IEC Guide 62 (C147)

The Chief Executive

Ir. J.C. van der Poel

ACCREDITATION CERTIFICATE

of **Orion Registrar Inc.**
Arvada (CO)
USA

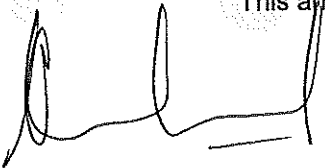
Valid from: **25-06-2008 to 30-08-2010**

Replaces annex dated: **19-12-2006**

Premises: **N.A.**

Standard / Normative document	Certification scheme (including sectors as applicable)
ISO 9001	Quality management system for the scopes (reference to EA/IAF-codes and NACE Rev. 1.1) : 7 pulp, paper and paper products 9 printing companies 12 chemicals, chemical products and fibres limited to "blending and purification activities through filtering" 13 pharmaceuticals, limited to "Ayurvedic drugs and Ayurvedic therapeutic products" 14 rubber and plastic products 17 basic metals and fabricated metal products 18 machinery and equipment 19 electrical and optical equipment 22 other transport equipment 29 wholesale and retail trade; repair of motor vehicles, motorcycles and personal and household goods 32 financial intermediation; real estate; renting 33 information technology 35 other professional services 38 health and social work, limited to Health, NACE 85.1 "Human Health Activities"
ISO 13485	Medical Devices - Quality Management Systems - Requirements for regulatory purposes

This annex has been approved by:



Ir. J.C. van der Poel
Chief Executive