KEMA Quality

CERTIFICATE

Number: 35235

John Zorich

has been qualified for following roles, indicated by the grey marked fields:

	Generally qualified for the role	Auditor Specifications					Product Categories					
		ISO 13485 (obligatory)	ISO 9001:2008 (additional)	EAC code 12: Chemicals, chemical products and fibres	EAC code 14: Rubber and plastic products	EAC code 19: Electrical and optical equipment	Non-Active Medical Devices	Non-Active Medical Devices - Animal Tissue	Active Medical Devices	Active Implantable Devices	In Vitro Diagnostic Medical Devices - Active Equipment	In Vitro Diagnostic Medical Devices - Reagents and Kits
Auditor												
Lead Auditor												
CMDCAS ISO 13485 Auditor												
Taiwan GMP Auditor												
Japan PAL Auditor												
Review Panel Member												
Low Risk Dossier Review												
High Risk Dossier Review												
510K Reviewer												
510K Supervisor												
Senior Peer Reviewer												
Notified Body Auditor												
Notified Body Lead Auditor												
Project Manager												
Certification Manager												
Product Expert												

Restrictions and/or extensions of the qualification:	also qualified for EAC codes 4 and 13
Applicable NBOG scopes Low Risk:	to be defined
Applicable NBOG scopes High Risk:	to be defined

Arnhem, 18 May 2010

KEMA Quality B.V.

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Mrs N.M. Pol,
Qualification Manager

KEMA Quality B.V.

A.J. Knipmeijer

Mr A.J. Knipmeijer,
Certification Manager