



CERTIFICATE OF APPROVAL

This is to certify that the Quality Management System of:

Harrison Clinical Research Group GmbH
Albrechtstr. 43, 80636 Munich
Germany

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

ISO 9001:2008

The Quality Management System is applicable to:

Head Office activities. Design and evaluation of clinical trials for the pharmaceutical, biotechnology, medical device and food industries, including conduct of phase I/II/III/IV trials. Secondary packaging and labelling operations of IMP and devices for use in clinical trials. Quality management consultancy and auditing services for clinical research organisation. Provision of subcontract clinical trial specialists to the pharmaceutical, biotechnology and medical device industries. Provision of clinical research and associated training courses.

This certificate is valid only in association with the certificate schedule bearing the same number on which the locations applicable to this approval are listed.

Approval
Certificate No: KLN 0206475

Original Approval: 26 March 1998

Current Certificate: 23 February 2010

Certificate Expiry: 31 March 2013

Issued by: Lloyd's Register Quality Assurance GmbH



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This document is subject to the provision on the reverse.

Innere Kanalstr. 15, 50823 Köln, Germany, Handelsregister Nr. B 34587

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

The use of the UKAS Accreditation Mark indicates Accreditation in respect of those activities covered by the Accreditation Certificate Number 001

Macro Revision 13



CERTIFICATE SCHEDULE

Harrison Clinical Research Group GmbH Albrechtstr. 43, 80636 Munich Germany

Head Office

Harrison Clinical Research Group GmbH
Albrechtstr. 43
D-80636 Munich

Activities

Head Office activities.

Locations

Harrison Clinical Research Deutschland GmbH
Albrechtstr. 14
D-80636 Munich

Activities

Design and evaluation of clinical trials for the pharmaceutical, biotechnology, medical device and food industries, including conduct of phase I/II/III/IV trials. Secondary packaging and labelling operations of IMP and devices for use of in clinical trials. Quality management consultancy and auditing services for clinical research organisations.

Harrison Clinical Research Benelux NV.
Keizer Karellaan 576
B-1082 Brussels

Management and monitoring of phase I/II/III/IV trials for the pharmaceutical, biotechnology, medical device and food industries. Provision of subcontract clinical trial specialists to the pharmaceutical, biotechnology and medical device industries.

Harrison Clinical Research France SAS
3 rue Bellanger
F-92300 Levallois-Perret

Management and monitoring of phase I/II/III/IV trials for the pharmaceutical, biotechnology, medical device and food industries.





CERTIFICATE SCHEDULE

Harrison Clinical Research Group GmbH Albrechtstr. 43, 80636 Munich Germany

Locations

Harrison Clinical Research Iberica SL
Calle Princep Jordi, 21-23, Esc. B
Entresuelo 1º B
E-08014 Barcelona

Harrison Clinical Research Ltd.
Gemini House, Bartholomew's Walk
Angel Drove
UK-Ely Cambridgeshire CB74EA

European Centre
for Clinical Research Training, ECCRT
Keizer Karellaan 576
B-1082 Brussels

Activities

Management and monitoring of phase I/II/III/IV trials for the pharmaceutical, biotechnology, medical device and food industries. Provision of subcontract clinical trial specialists to the pharmaceutical, biotechnology and medical device industries.

Management and monitoring of phase I/II/III/IV trials for the pharmaceutical, biotechnology, medical device and food industries.

Provision of clinical research and associated training courses.

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