

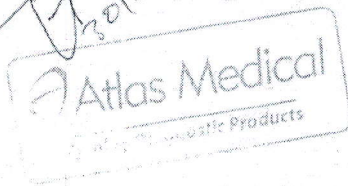
Date: 30/06/2018

STATEMENT

We, **Atlas Medical** having a registered office at William James House, Cowley Road, Cambridge, CB4 0WX, UK assign SRL Sanmedico having a registered office at A. Corobceanu street 7A, apt. 9, Chişinău MD-2012, Moldova , as authorized representative in correspondence with the conditions of directive 98/79/EEC.

We declare that the company mentioned above is authorized to register, notify, renew or modify the registration of medical devices on the territory of the Republic of Moldova.

On behalf of the Manufacturer
General Manager
Haya Amawi



Head Office William James House, Cowley Rd, Cambridge, CB4 0WX, United Kingdom.
Tel: +44 (0) 1223 858910, Fax: +44 (0) 1223 858524

Middle East Site : King Abdullah the Second Industrial Estate, Street 19, Sahab Free Zone Area, P.O. Box: 204, Amman 11521 Jordan





Declaration Ref No: DC11-0051

CE Declaration of Conformity

According to Annex III of the IVD Directive 98/79/EC

We,

Atlas Medical

Head office: William James House, Cowley Road, Cambridge, CB4 0WX, UK

Tel: +44 1223 858 910

Fax: +44 1223 858 524

Email: info@atlas-site.co.uk

Middle East Site: Sahab Free Zone Area, P. O. Box 212555, Amman, Jordan.

Tel.: +962 6 4026468

Fax: +962 6 4022588

Email: info@atlas-medical.com

Declare our responsibility that the following product:

See Attached list

- Comply with all essential requirements (Annex I) of the IVD Directive 98/79/EC. This compliance has been properly documented and covers the items listed in Annex I of the IVD Directive.
- This product is produced under Atlas quality system (ISO13485:2003) issued by Lloyd's Register Quality Assurance.
- Comply with the essential requirements of following standards (EN 18113-1, -2, -4:2011, EN ISO 15223:2012, EN ISO 13532: 2002, EN ISO 14971:2012, EN ISO 13640:2002, ISO 2859/1:1999, EN ISO 13612:2002, EN ISO 13641:2002.

And

Intended for In-Vitro Professional use only.

Manufacturer
Atlas Medical
 William James House, Cowley Rd.,
 Cambridge, CB4 0WX, UK

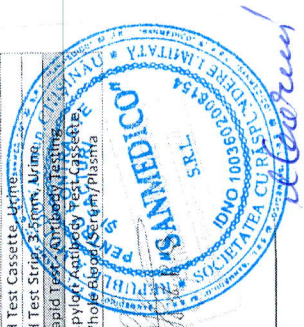
Atlas Medical	Issue date December, 2011	Date of review 21st of March, 2018	Management approval 	MRXD010F.10 08.02.2011
---------------	------------------------------	---------------------------------------	-------------------------	---------------------------

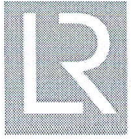


CE Declaration of Conformity

According to Annex III of the IVD Directive 98/79/EC

Catalogue No	Description	Catalogue No	Description
8.00.00	CRP latex Kits	8.02.48	Calcium Chloride
8.00.01	CRP latex Kits with buffer	8.02.69	Fibrinogen Reagent
8.00.02	ASO latex Kits	Hemoglobin Reagents	
8.00.03	ASO latex Kits with buffer	8.02.46	Drabkins Reagent, 40x
8.00.04	RF latex Kits	8.02.50	Hemoglobin Standard, 15g/dL
8.00.05	RF latex Kits with buffer	Sickle Cell Kits	
8.00.07	HCG latex Kits	8.02.67	Sickle Cell Kit
8.00.08	IM (Horse Stroma) latex Kits	8.02.68	Sickle Cell positive & negative control set
8.00.11	SLE latex kits	Urine Reagent Strips	
8.00.12	Staphylococcus Latex Kits	8.03.00	URS 1 Parameter: Glucose
8.00.13	Streptococcus Latex kits	8.03.01	URS 1 Parameter: Protein
8.00.15	E.Coli Latex Kits	8.03.02	URS 1 Parameter: Ketone
8.00.16	Rota Virus Latex Kits	8.03.03	URS 2 Parameters: Glucose, Ketone
8.00.17	D-Dimer Latex kits	8.03.04	URS 2 Parameters: Glucose, Protein
8.00.21	Waaler rose Latex Kits	8.03.05	URS 2 Parameters: Urobilinogen, Bilirubin (Liver Function Test)
Febrile Antigen Kits		8.03.06	URS 3 Parameters: Protein, pH, Glucose
8.01.00	Brucella Rose Bengal	8.03.07	URS 3 Parameters: Glucose, Protein, ketone
8.01.01	Salmonella OA Reagent	8.03.15	URS 9 Parameters: Nitrite, Protein, pH, Blood, Specific Gravity, ketone, Bilirubin, Glucose
8.01.02	Salmonella OB Reagent	8.03.16	URS 10 Parameters: Leucocytes, Nitrite Urobilinogen, Protein, pH, Blood, Specific Gravity, Ketone, Bilirubin, Glucose
8.01.03	Salmonella OC Reagent	8.03.17	URS 10 Parameters: Nitrite, Urobilinogen, Protein, pH, Blood, Specific Gravity, Ketone, Bilirubin, Glucose
8.01.04	Salmonella OD Reagent	8.03.18	URS 11 Parameters: Leucocytes, Nitrite, Urobilinogen, Protein, pH, Blood, Specific Gravity, Ketone, Bilirubin, Glucose, Ascorbic Acid
8.01.05	Salmonella HA Reagent	Fertility Rapid Tests	
8.01.06	Salmonella HB Reagent	8.04.00	hCG Test Cassette, Urine
8.01.07	Salmonella HC Reagent	8.04.01	hCG Test Cassette, Urine/Serum
8.01.08	Salmonella HD Reagent	8.04.04	hCG Test Strip, 5.0mm, Urine
8.01.10	Brucella Abortus Reagent	8.04.05	hCG Test Strip, 3.5mm, Urine
8.01.11	Brucella Melitensis Reagent	8.04.06	hCG Test Strip, 2.5mm, Urine/Serum
8.01.12	Proteus OX2 Reagent	8.04.10	hCG Test Strip, 5.0mm, Urine/Serum
8.01.13	Proteus OX19 Reagent	8.04.12	hCG Test Strip, 2.5mm, Urine/Serum
8.01.14	Proteus OXK Reagent	8.04.88	hCG Test Strip, 3.5 mm, Urine/Serum
8.01.15	Brucella Antigen Kits	8.04.90	hCG Test Strip, 2.5 mm, Urine/Serum
8.01.16	Salmonella Antigen Sets	8.04.14	LH Test Cassette - Urine
8.01.17	Febrile Antigen Set (10 Antigens)	8.04.15	LH Test Strip - 5.0mm, Urine/Serum
8.01.17	Febrile Antigen Set (10 Antigens) With controls	8.04.20	Infectious Disease Rapid Tests - Urinary Testing
8.01.18	Salmonella Antigen Set, Widal Kit (6 Antigens: OA, OB, OD, HA, HB, HD)	Rapidity Quality Test Cassette	
8.01.18	Salmonella Antigen Set, Widal Kit (6 Antigens: OA, OB, OD, HA, HB, HD) with controls	Whole Blood #7/Plasma	
8.01.19	Febrile Antigen Positive Control		
8.01.20	Febrile Antigen Negative Control		
Co-agulation Reagents			
8.02.40	PT Calcium Rabbit Brain		
8.02.41	Thromboplastin, liquid		
8.02.41	APTT (PTT) Micronised Silica Platelet Substitute, Liquid		
8.02.60	Normal Coagulation Control		
8.02.61	Abnormal Coagulation Control		
8.02.44	PT Kit		
8.02.45	APTT (PTT) Kit		





Lloyd's Register

Certificate of Approval

This is to certify that the Management System of:

Atlas Medical

King Abdullah II Industrial Estate, Street No. 19, Sahab Free Zone Area, Amman, 11512, Jordan

has been approved by LRQA to the following standards:

ISO 13485:2003

Basem Obaid - Area Operations Manager

Issued By: Lloyd's Register EMEA

for and on behalf of: Lloyd's Register Quality Assurance Limited

Current Issue Date: 23 March 2018
Expiry Date: 31 March 2019
Certificate Issue Number: 10067833

Original Approvals:
ISO 13485 28 February 2009

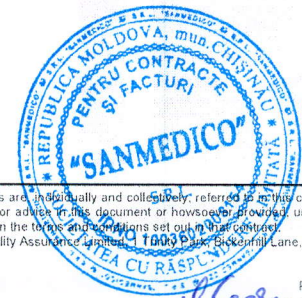
Approval Certificate Number: ISO 13485 – 0046833

The scope of this approval is applicable to:
ISO 13485:2003

Design Manufacturing and Supply of Medical
Diagnostic Reagents and Kits



001





LumiQuick Diagnostics, Inc.
2946 Scott Blvd., Santa Clara, CA 95054, USA

Tel: 1-408-855-0061
Fax: 1-408-855-0063
E-mail: info@lumiquick.com
Website: www.lumiquick.com

Date: February 13, 2018

LETTER OF AUTHORIZATION

To whom it may concern:

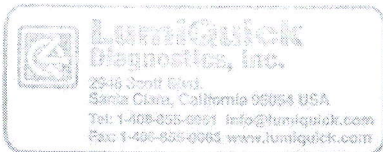
We, LumiQuick Diagnostics Inc. having a registered office at 2946 Scott Blvd, Santa Clara, CA 95054, USA, assign Sanmedico SRL having a registered office at str. A. Corobceanu 7A, apt. 9, Chişinău MD-2012, Moldova , as authorized representative in correspondence with the conditions of directive 98/79/EEC.

We declare that the company mentioned above is authorized to register, notify, renew or modify the registration of medical devices on the territory of the Republic of Moldova.

This authorization letter is valid until February 28, 2020.

Best regards,

Charles Yu
President



bsi.



By Royal Charter

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016

This is to certify that:

LumiQuick Diagnostics, Inc.
2946 Scott Blvd
Santa Clara
California
95054
USA

Holds Certificate No:

FM 574919

and operates a Quality Management System which complies with the requirements of ISO 13485:2016 for the following scope:

The design, development, manufacture and distribution of in vitro diagnostics test kits and reagents used in the diagnosis and management of disease status, including Infectious Diseases tests, Drugs of Abuse tests, Cardiac Monitor tests, Cancer Marker tests, Fertility Hormone tests, ELISA tests & Urine Chemistry tests.

For and on behalf of BSI:

Stewart Brain, Head of Compliance & Risk - Medical Devices

Original Registration Date: 2011-10-20

Latest Revision Date: 2018-11-21

Effective Date: 2017-10-20

Expiry Date: 2020-10-19

Page: 1 of 1



...making excellence a habit.™

This certificate remains the property of BSI and shall be returned immediately upon request.
An electronic certificate can be authenticated [online](#). Printed copies can be validated at www.bsigroup.com/ClientDirectory
To be read in conjunction with the scope above or the attached appendix.

Americas Headquarters: BSI Group America Inc., 12950 Worldgate Drive, Suite 800, Herndon, VA 20170-6007 USA
A Member of the BSI Group of Companies.





LumiQuick Diagnostics, Inc.
2946 Scott Blvd., Santa Clara, CA 95054, USA

Tel: 408-855-0061
Fax: 408-855-0063
E-mail: info@LumiQuick.com
Web: www.LumiQuick.com

Declaration of Conformity

PRODUCT IDENTIFICATION		
Product name	Model/number	
Cardiac Marker Test Devices		
QuickProfile Troponin I Serum Test Card	75001	
QuickProfile Troponin I Whole Blood Test Card	75002	
QuickProfile Cardiac Panel Serum Test Card	75003	
QuickProfile Cardiac Panel Whole Blood Test Card	75004	
QuickProfile Myoglobin Serum Test card	75005	
QuickProfile Myoglobin Whole Blood Test Card	75006	
QuickProfile CK-MB Serum Test Card	75007	
QuickProfile CK-MB Whole Blood Test Card	75008	
QuickProfile Troponin I Strip	75009	
QuickProfile CK-MB Strip	75010	
QuickProfile Myoglobin Strip	75011	
MANUFACTURER		
Name of company	Address	Representative
LumiQuick Diagnostics, Inc.	2946 Scott Blvd. Santa Clara, CA 95054 USA	Jeff Wang
AUTHORIZED REPRESENTATIVE		
Name of company	Address	Telephone/email
Emergo Europe	Prinsessegracht 20 2514 AP The Hague, Netherlands	+31.70.345.8570 - phone +31.70.346.7299 - fax europe@emergogroup.com
CONFORMITY ASSESSMENT		
Device classification	Route to compliance	Standards applied
Class: Self-Certify	Annex III of IVDD 98/79/EC Council Directive	ISO 13485:2003

LumiQuick Diagnostics, Inc. declares that the above mentioned products meet the provision of the Council Directive 98/79/EC for In Vitro Diagnostic Medical Devices and Directive 98/79/EC as transposed in the national laws of the Member States.

COMPANY REPRESENTATIVE: Jeff Wang

TITLE: Quality Systems Manager

SIGNATURE: 

DATE: 28/04/2017

EC_Declaration_Letter_Emergo_E2R0_NewAddress





Your **Trusted** Partner

San Diego July 11th, 2018

We, ACON Laboratories Inc. having a registered office at 10125 Mesa Rim Road. San Diego, CA 92121, USA assign SRL Sanmedico having a registered office at A. Corobceanu street 7A, apt. 9, Chişinău MD-2012, Moldova , as authorized representative in correspondence with the conditions of directive 93/42/EEC, 98/79/EEC and 90/385/EEC.

We declare that the company mentioned above is authorized to register, notify, renew or modify the registration of medical devices on the territory of the Republic of Moldova.

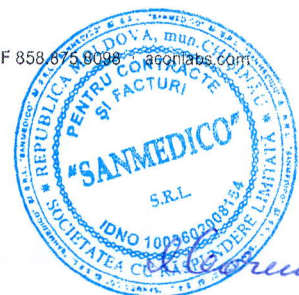
ACON reserves the right to cancel this authorization at any time with a one month notice. If this is the case, ACON will honor any obligation to supply to our representative SanMedico SRL all the products distribution acquired or in the process of being acquired in Public Price bids and Public Tenders process.

Sincerely,


Jassy Alvarenga
Account Manager, International Sales



ACON Laboratories





Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical
Devices (MDD), Annex II excluding (4)
(Devices in Class Ila, I Ib or III)
No. G1 080997 0018 Rev. 00

Manufacturer: ACON Laboratories, Inc.

10125 Mesa Rim Road,
San Diego CA 92121
USA

EC-Representative: Medical Device Safety Service GmbH
Schriftgraben 41, 30175 Hannover, GERMANY

Product Category(ies): Lancets, Safety Lancets

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: SH18743EXT01

Valid from: 2018-09-07
Valid until: 2023-09-06

Date, 2018-09-05
S. Preis
Stefan Preis



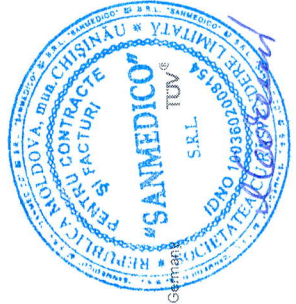
EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical
Devices (MDD), Annex II excluding (4)
(Devices in Class Ila, I Ib or III)
No. G1 080997 0018 Rev. 00

Facility(ies):

ACON Laboratories, Inc.
10125 Mesa Rim Road, San Diego CA 92121, USA

AZURE Institute, Inc.
10125 Mesa Rim Road, San Diego CA 92121, USA



ZERTIFIKAT ◆ CERTIFICATE ◆ CERTIFICADO ◆ CERTIFICAT ◆ СЕРТИФИКАТ ◆ 認証証書 ◆ CERTIFICATE ◆ CERTIFICATE ◆ CERTIFICATE



Product Service

CERTIFICATE

No. Q1N 16 05 42074 027

Holder of Certificate: Acon Biotech (Hangzhou) Co., Ltd.

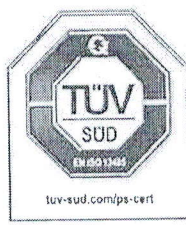
No.210 Zhenzhong Road
West Lake District
310030 Hangzhou
PEOPLE'S REPUBLIC OF CHINA

Facility(ies):

Acon Biotech (Hangzhou) Co., Ltd.
No.210 Zhenzhong Road, West Lake District,
310030 Hangzhou, PEOPLE'S REPUBLIC OF CHINA



Certification Mark:



Scope of Certificate: Design and Development,
Production and Distribution of
In Vitro Diagnostic Test Kits
and Related Instruments,
Lancet and Lancing Device

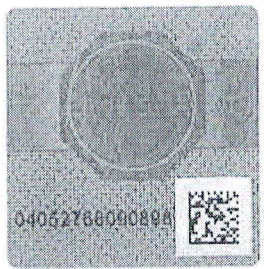
Applied Standard(s):

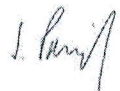
EN ISO 13485:2012 + AC:2012
Medical devices - Quality management systems -
Requirements for regulatory purposes
(ISO 13485:2003 + Cor. 1:2009)
DIN EN ISO 13485:2012

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). See also notes overleaf.

Report No.: SH1610619

Valid from: 2016-07-15
Valid until: 2019-07-14

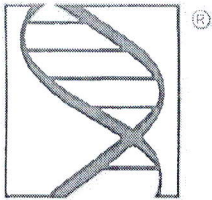


Date, 2016-07-08

Stefan Preis

Page 1 of 1

DAKKS





SYNTESYS



SYNTESYS S.A.S. DI RINALDO R. & C.

VIA G. GALILEI, 10/3
35037 Z.I. SELVE DI TEOLO (PD)
TEL. +39 049 9903866 R.A. FAX +39 049 9903867
COD. FISCALE P.IVA N.REG.IMP. PADOVA 03573950288
E-MAIL INFO@SYNTESYS.IT · WEB WWW.SYNTESYS.IT

AUTHORIZATION LETTER

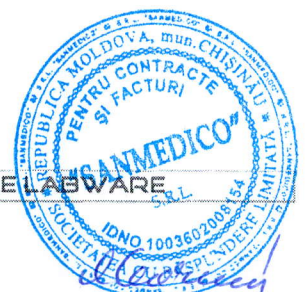
We, **Syntesys S.A.S.** having a registered office at Via G. Galilei 10/3, 35037 Selve di Teolo - PD - Italy, assign **Sanmedico SRL** having a registered office at A. Corobceanu str., apt. 9, Chişinău MD-2012, Moldova, as authorized representative in correspondence with the conditions of directive 98/79/CE and 93/42/CE.

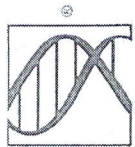
We declare that the company mentioned above is authorized to register, notify, renew or modify the registration of medical devices on the territory of the Republic of Moldova.

Teolo, 02.01.2018

 **SYNTESYS S.A.S.**
DI RINALDO R. & C.
Via G. GALILEI, 10/3
Z.I. SELVE 35037 TEOLO (PD) - CF 03573950288
TEL. 049 9903866 FAX 049 9903867


Rinaldo Ruggero
CEO and Legal Representative
SYNTESYS S.A.S.





SYNTESYS



SYNTESYS S.A.S. DI RINALDO R. & C.
VIA G. GALILEI, 10/3
35037 ZI. SELVE DI TEOLO (PD)
TEL. +39 049 9903666 R.A. FAX. +39 049 9903667
COD. FISCALE P.IVA N. REG. IMP. PADOVA 03573950288
E-MAIL: INFO@SYNTESYS.IT - WEB: WWW.SYNTESYS.IT

DICHIARAZIONE DI CONFORMITA'
Conformity declaration



Il sottoscritto, Rinaldo Ruggero, legale rappresentante della ditta:
The undersigned, Rinaldo Ruggero, legal representative of the company:

produttore/manifatturere

SYNTESYS S.a.s. di Rinaldo Ruggero & C.
indirizzo/address

Via G. Galilei, 10/3 35037 Zona Industriale SELVE DI TEOLO (PADOVA) ITALY

o rappresentante il mandatario autorizzato entro la Unione Europea o representing the
authorized mandatary within the European Community

Mandatario autorizzato/authorized mandatary

indirizzo/address

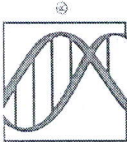
Dichiara sotto la propria responsabilità che il prodotto/declares under his own
responsibility that the product:

Denominazione degli
articoli
prodotti/Description of
Manufacturer

Contenitori per urina, contenitori per feci,
contenitori universali, pipette Pasteur, piastre di
Petri, Anse Sterili per batteriologia, Aste a "L",
Puntali Eppendorf gialli e blue, cuvette per
spettrofotometro, tazzine per campionamento siero-
bacchette per distacco ed estrazione del coagulo,
pinzette in polistirolo monouso, provette monouso in
plastica, tappi alettati per provetta diam. 12 mm e
16mm, provette con granuli ed acceleratore, provette
sottovuoto per prelievo, Sistema SEDIPLAST,
Microprovette, portavetrini, Vetrini precolorati,
Portaprovette, supporti per microprovette, bottiglie
per raccolta urina.

Urine container, faeces container, universal
container, Pasteur pipette, Petri dishes, Sterile
loops, Sterile loops open "L", Eppendorf tips yellow
and blue, cuvettes for spectrophotometer, samples
cups, Rod to detach clot, disposable forceps,
disposable plastic tubes, winged stoppers for tubes
diam. 12mm & 16mm, Test tube with granules and clot
activator, vacuum test tube, SEDIPLAST system,
micro test tubes, Slides holder, TESTSIMPLETS slide,
rack for test tubes, rack for micro test tubes,
Bottles for urine collection.

PRODOTTI E STRUMENTAZIONE PER ANALISI DI LABORATORIO - DISPOSABILE LABWARE



SYNTESYS

Materiale/Material

Polipropilene, Polistirolo, Polietilene e
Polimetilmetacrilato
Polypropylene, Polystyrene, Polyethylene and
Polymethylmetacrylate

È conforme alle disposizioni della direttiva 98/79/CE concernente i dispositivi medici
diagnostici in vitro e recepito in Italia con D.L. del 06/04/2000 n° 332 allegato I
(requisiti essenziali) ed è fabbricato in accordo ai requisiti di cui all'Allegato III della
sopra citata direttiva / It meets the CE Directive 98/79 CE about in vitro diagnostic device
specifications established by the Italian law n. 332, dated 6th September 2000. The device is
made according to the specifications of the III attached of the above-mentioned directive.

Dichiara inoltre che la documentazione tecnica di supporto alla presente dichiarazione di
conformità è conservata presso gli uffici dell'azienda e sarà posta alla disposizione di chi
la richiede/declares that all technical documents attached to this conformity statement are
filed in our company and can be consulted by any authorized body on demand.

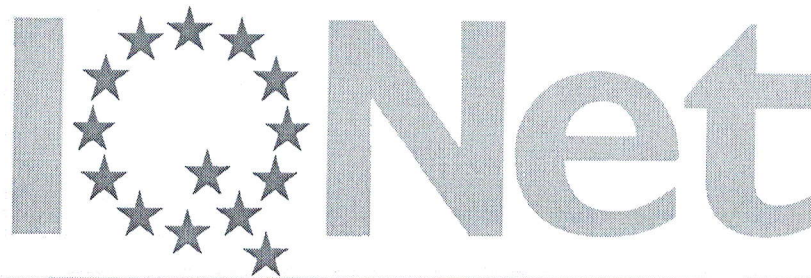
Data 07/01/2016
Issued on January 7th 2016

SYNTESYS S.a.s.
Il legale rappresentante
Rinaldo Ruggero



PRODOTTI E STRUMENTAZIONE PER ANALISI DI LABORATORIO - DISPOSABILE LABWARE

®



THE INTERNATIONAL CERTIFICATION NETWORK

CERTIFICATE

CISQ/ICIM SPA has issued an IQNet recognized certificate that the organization:

SYNTESYS S.a.s. di Rinaldo Ruggero e C.

Via G. Galilei, 10/3 - Zona Industriale - I-35037 Selve di Teolo (PD)

has implemented and maintains a

Quality Management System

for the following scope:

Trading of products for laboratory analysis. Design, manufacturing and sale of products for laboratory analysis and sanitary products. Sale agency of instruments, reagents and consumable products for laboratory diagnostic.

which fulfils the requirements of the following standard:

ISO 9001:2015

Issued on: 2018-06-04

First issued on: 2013-06-05

Expires on: 2019-06-04

This attestation is directly linked to the IQNet Partner's original certificate and shall not be used as a stand-alone document.

Registration Number: IT-83562



Alex Stoichitoiu
President of IQNET



Ing. Claudio Provetti
President of CISQ

IQNet Partners*:

AENOR Spain AFNOR Certification France APCER Portugal CCC Cyprus CISQ Italy
CQC China CQM China CQS Czech Republic Cro Cert Croatia DQS Holding GmbH Germany FCAV Brazil
FONDONORMA Venezuela ICONTEC Colombia Inspecta Sertifiointi Oy Finland INTECO Costa Rica
IRAM Argentina JQA Japan KFQ Korea MIRTEC Greece MSZT Hungary Nemko AS Norway NSAI Ireland
NYCE-SIGE México PCBC Poland Quality Austria Austria RR Russia SII Israel SIQ Slovenia
SIRIM QAS International Malaysia SQS Switzerland SRAC Romania TEST St Petersburg Russia TSE Turkey YUOS Serbia
IQNet is represented in the USA by: AFNOR Certification, CISQ, DQS Holding GmbH and NSAI Inc.



* The list of IQNet partners is valid at the time of issue of this certificate. Updated information is available under www.iqnet-certification.com

®



THE INTERNATIONAL CERTIFICATION NETWORK

CERTIFICATE

CISQ/ICIM SPA has issued an IQNet recognized certificate that the organization:

SYNTESYS S.a.s. di Rinaldo Ruggero e C.

Via G. Galilei, 10/3 - Zona Industriale - I-35037 Selve di Teolo (PD)

has implemented and maintains a

Quality Management System

for the following scope:

Trading of products for laboratory analysis. Design, manufacturing and sale of products for laboratory analysis and sanitary products. Sale agency of instruments, reagents and consumable products for laboratory diagnostic.

which fulfils the requirements of the following standard:

UNI CEI EN ISO 13485:2016

Issued on: 2018-06-04

First issued on: 2014-06-21

Expires on: 2019-06-04

This attestation is directly linked to the IQNet Partner's original certificate and shall not be used as a stand-alone document.

Registration Number: IT-93779



Alex Stoichitoiu
President of IQNET



Ing. Claudio Provetti
President of CISQ

IQNet Partners*:

AENOR Spain AFNOR Certification France APCER Portugal CCC Cyprus CISQ Italy
CQC China CQM China CQS Czech Republic Cro Cert Croatia DQS Holding GmbH Germany FCAV France
FONDONORMA Venezuela ICONTEC Colombia Inspecta Sertifiointi Oy Finland INTECO Costa Rica
IRAM Argentina JQA Japan KFQ Korea MIRTEC Greece MSZT Hungary Nemko AS Norway NSAI Ireland
NYCE-SIGE Mexico PCBC Poland Quality Austria Austria RR Russia SII Israel SIQ Slovenia
SIRIM QAS International Malaysia SQS Switzerland SRAC Romania TEST St Petersburg Russia TSE Turkey UNIOS Serbia
IQNet is represented in the USA by: AFNOR Certification, CISQ, DQS Holding GmbH and NSAI Inc.



* The list of IQNet partners is valid at the time of issue of this certificate. Updated information is available under www.iqnet-certification.com