



CERTIFICATO CE

Certificato n. 1976/MDD

Dichiarazione di approvazione del sistema qualità

(Garanzia di qualità della produzione)

Visto l'esito delle verifiche condotte in conformità all'Allegato V, punto 3 e tenendo conto dell'Allegato VII, punto 5 della direttiva 93/42/CEE e s.m.i., si dichiara che la ditta:

CERACARTA SPA

47122 FORLI' (FC) - VIA SECONDO CASADEI 14 Z.I. VILLA SELVA (ITA) - Italy

mantiene nello stabilimento di:

47122 FORLI' (FC) - VIA GRAMADORA 12/14 (ITA) - Italy

47122 FORLI' (FC) - VIA SECONDO CASADEI 14 Z.I. VILLA SELVA (ITA) - Italy

un sistema qualità che assicura la conformità dei seguenti prodotti:

Carte per registrazione ad uso medico

Modd. come da documento "Allegato al Certificato CE no. 1976/MDD - Elenco dei Dispositivi" rev.0 del 07/10/2020, valido solo se provvisto di timbro IMQ; tale allegato costituisce parte integrante e sostanziale del presente certificato.

ai requisiti metrologici ad essi applicabili della direttiva suddetta (in tutte le fasi della fabbricazione) ed è sottoposta alla sorveglianza prevista dal punto 4 dell'Allegato V.

Riferimento pratiche IMQ:

DM17-0017248-01; DM20-0056104-01.

Questa Dichiarazione di approvazione è rilasciata dall'IMQ S.p.A. quale organismo notificato per la direttiva 93/42/CEE e s.m.i. Il numero identificativo dell'IMQ S.p.A. quale organismo notificato è: 0051.

Emesso il: 2017-11-18
Data aggiornamento: 2020-10-07
Sostituisce: 2017-11-18
Data scadenza: 2022-11-17

IMQ



EC CERTIFICATE

Certificate No 1976/MDD

Production Quality Assurance System Approval Certificate

On the basis of our assessment carried out according to Annex V, section 3 and considering the Annex VII, section 5 of the Directive 93/42/EEC and its revised version, we hereby certify that:

CERACARTA SPA

47122 FORLI' (FC) - VIA SECONDO CASADEI 14 Z.I. VILLA SELVA (ITA) - Italy

manages in the factory of:

47122 FORLI' (FC) - VIA GRAMADORA 12/14 (ITA) - Italy

47122 FORLI' (FC) - VIA SECONDO CASADEI 14 Z.I. VILLA SELVA (ITA) - Italy

a quality assurance system ensuring the conformity of the following products:

Medical recording chart paper

Type ref. as to document "Annex of EC Certificate no. 1976/MDD - Device List" rev. 0 dated 2020/10/07 valid only if provided with IMQ stamp; this annex is integral and substantial part of this certificate.

with the relevant metrological requirements of the aforementioned directive (as far as all the manufacturing stage is concerned) and it is subject to surveillance as specified in section 4 of Annex V.

Reference to IMQ files Nos:

DM17-0017248-01; DM20-0056104-01.

This Approval Certificate is issued by IMQ S.p.A. as Notified Body for the Directive 93/42/EEC and its revised version. Notified Body notified to European Commission under number: 0051.

Date: 2017-11-18
Updated: 2020-10-07
Substitution Date: 2017-11-18
Expiry Date: 2022-11-17

IMQ

Allegato al Certificato CE n. 1976/MDD- Elenco dei Dispositivi

EC Certificate Annex 1976/MDD - Device List

rev. 0 del/of 2020/10/07

Categoria di dispositivo: Device category:	Carte per registrazione ad uso medico Medical recording chart paper
Modello/i: Model(s):	22.01 Pacchi stampati medicali (per ECG,EEG,CTG,e laboratorio analisi); 21.01 Rotoli stampati medicali (per ECG,EEG,CTG,e laboratorio analisi); 32.01 Schede e dischi stampati medicali.
Marca/Marche: Trade mark(s):	Ceracarta
Altre informazioni rilevanti: Other relevant data:	Dispositivi medici in classe I con funzione di misura (IM) Medical Devices in class I with measuring function (IM)

Timbro IMQ
IMQ Stamp



2020-10-07

Carte diagrammate per tutte le apparecchiature di elettrodiagnostica.
Materiale di consumo ed accessori elettromedicali.
Carte per apparecchi registratori industriali.
Rotoli e pacchi speciali per sistemi esattoriali, di controllo, lotterie.
Etichette radiofrequenza e soluzioni integrate.

Chart Papers for all electrodiagnostic equipments.
Disposable and electromedical accessories.
Chart Papers industrial recording instruments.
Special rolls and fanfolds for tickets checking system, lottery.
Rfid labels and chain solutions.

Sede (Head office and works) :
Via Secondo Casadei, 14 - 47122 FORLÌ - ITALY
Tel : 0039 0543 780055 • Fax : 0039 0543 781404
http : // www.ceracarta.it • e-mail : info@ceracarta.it.
Capitale Sociale : € 1.000.000 int. vers.
Registro Imprese FORLÌ-CESENA
P.I. / C.F. / VAT.N. IT 00136740404
R.E.A. FORLÌ N. 72646 - N. MECC. FO 006863

Messrs

Medicines and Medical Devices Agency

Forlì, 3rd August 2022

We, CERACARTA SPA V having a registered office at Via Secondo Casadei 14, 47122 Forlì (FC)- Italy, assign "GBG-MLD" SRL, having a registered office at Str. Albisoara 64/2, Chisinau MD -2005, Moldova, as **Authorized representative** in correspondence with the conditions of LAW No. 102 from 09.06.2017 regarding medical devices.

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.

CERACARTA SPA
Bandini Alessandro

 CERACARTA S.p.A.

Dal 1960 al Vostro Servizio - Since 1960 at Your Service

DECLARATION OF CONFORMITY

Forlì, 3rd August 2022

Manufacturer: **Ceracarta S.p.A.**

Address: **Via Secondo Casadei, 14 47122 Forlì - ITALY**

DECLARES

that

THE RECORDING "THERMAL" AND/OR "INK" PAPERS IN THE MEDICAL FIELD
for "ECG", "EEG", "CTG" and papers for laboratory tests (including the model M 1911 A),
identified and classified in the

Technical file, comply with the directive about medical devices (DIRECTIVE 93/42/CEE as amended by 2007/47/EC).

In addition to this, we precise that:

- according to the Directive 93/42/EEC the listed products are medical devices belonging to class I with function of measure.
- they are subject to the regulations of the Attachment I of the above mentioned directive;
- the evaluation of the device was in accordance with Annex V of that Directive, by the certified body IMQ S.p.A. – CE0051;
- Ceracarta S.p.A. has implemented a quality system in accordance with the regulations UNI EN ISO 9001: 2015 and UNI CEI EN ISO 13485:2016.

CERACARTA SPA
Bandini Alessandro


Alessandro Bandini

DECLARATION OF CONFORMITY

Forlì, 3rd August 2022

The writing company Ceracarta S.p.a. located in Via Secondo Casadei n° 14, 47122 Forlì, manufacturer of the products named , "ECG GEL / ECO SUPERGEL ." (**BASIC UDI-DI 8059170GC004NN**), identified and classified in the technical file, declares under its own responsibility that such devices satisfies all the requirements of MDR 2017/745 , about medical devices and in particular that:

- in accordance with enclosure VIII-section III of the MDR the Dispositives in object must be considered as belonging to Class I;
- the Dispositives in object satisfy the essential requirements as in enclosure I of MDR 2017/745;
- the manufacturer has prepared and keeps the technical files updated in accordance with enclosure II of the MDR;
- such documentation is available at the headquarters of Ceracarta , for any reference by the entitled bodies;
- the Dispositive in object must be exclusively used together with electro-medical instruments for recording, diagnosis and therapy, which base their functioning upon the measuring of energy flows of electric, magnetic and ultrasound type;
- Ceracarta S.p.A. has implemented a quality system in accordance with the regulations UNI EN ISO 9001: 2015 and UNI CEI EN ISO 13485:2016.

The president

CERACARTA SPA
Bandini Alessandro


Alessandro Bandini

DECLARATION OF CONFORMITY

Forlì, 3rd August 2022

The writing company Ceracarta S.p.a. located in Via Secondo Casadei n° 14, 47122 Forlì, manufacturer of the products named , **TOP TRACE ECG ELECTRODES (BASIC UDI-DI 8059170EL001PR)**, identified and classified in the technical file, declares under its own responsibility that such devices satisfies all the requirements of MDR 2017/745 , about medical devices and in particular that:

- in accordance with enclosure VIII-section III of the MDR the Dispositives in object must be considered as belonging to Class I;
- the Dispositives in object satisfy the essential requirements as in enclosure I of MDR 2017/745;
- the manufacturer has prepared and keeps the technical files updated in accordance with enclosure II of the MDR;
- such documentation is available at the headquarters of Ceracarta , for any reference by the entitled bodies.

In addition:

- the device is tested according to the voluntary Association for the Advancement of Medical Instrumentation (AAMI) standard requirements for electrical performance for disposable ECG electrodes (ANSI/AAMI EC 12:2000) and test results meet or exceed these performance standards.

- the device is tested and it is found to be acceptable for use, according to:

UNI EN ISO 10993-1: "Biological evaluation of medical devices" ;

UNI EN ISO 10993-5 : "Biological evaluation of medical devices :tests for in vitro cytotoxicity";

UNI EN ISO 10993-10 : "Biological evaluation of medical devices :tests for irritation and sensitization".

- Ceracarta does not use any latex/PVC materials or ingredients in the manufacturing of our electrodes;
- Ceracarta S.p.A. has implemented a quality system in accordance with the regulations UNI EN ISO 9001: 2015 and UNI CEI EN ISO 13485:2016.

The president

CERACARTA SPA
Bandini Alessandro


CERACARTA S.p.A.

DECLARATION OF CONFORMITY

Forlì, 3rd August 2022

The writing company Ceracarta S.p.a. located in Via Secondo Casadei n° 14, 47122 Forlì, manufacturer of the products named , **“COMPATIBLE PAPER FOR SONY UPP 110 S / HG / HD ROLLS”**, (BASIC UDI-DI 8059170CA001LN), identified and classified in the technical file, declares under its own responsibility that such devices satisfies all the requirements of MDR 2017/745 , about medical devices and in particular that:

- in accordance with enclosure VIII-section III of the MDR the Dispositives in object must be considered as belonging to Class I;
- the Dispositives in object satisfy the essential requirements as in enclosure I of MDR 2017/745;
- the manufacturer has prepared and keeps the technical files updated in accordance with enclosure II of the MDR;
- such documentation is available at the headquarters of Ceracarta , for any reference by the entitled bodies.
- Ceracarta S.p.A. has implemented a quality system in accordance with the regulations UNI EN ISO 9001: 2015 and UNI CEI EN ISO 13485:2016.

The president

CERACARTA SPA
Bandini Alessandro

CERACARTA S.p.A.
Alessandro Bandini

SONY

Sony Corporation
1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan

EU DECLARATION OF CONFORMITY

(EN)

1. Model No.: UPC-21L
2. Name and address of the manufacturer's authorised representative: Sony Belgium, bijkantoor van Sony Europe B.V., Da Vincilaan 7-D1, 1930 Zaventem, Belgium
3. This declaration of conformity is issued under the sole responsibility of the manufacturer: Sony Corporation, 1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan
4. Object of the declaration: Color Printing Pack
5. The object of the declaration described above is in conformity with:
2017/745 (Medical Device Regulation)
6. Where applicable, references to the relevant harmonised standards used or references to the technical specifications in relation to which conformity is declared:
Medical: EN 60601-1:2006 + A1:2013
7. Where applicable, the notified body (name and number), description of intervention and certificate:
8. Additional information:
 - Classification: Class I, MDR Annex VIII
 - Basic UDI-DI: 4901780PrintMediaWR
 - Manufacturer SRN: JP-MF-000014553
 - Authorised Representative SRN: BE-AR-000007876
 - Intended Purpose: Print media for Sony's medical printer
 - Accessory(ies): -
 - Note(s): -

Signed for and on behalf of: Sony Corporation

Tokyo, 2021-11-04

Reference Number: 2021EU02261

Sony Corporation
1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan



Ryogo Katayama
Quality Officer
Quality Officer Group
Quality & Environmental Dept.

SONY

Sony Corporation
1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan

EU DECLARATION OF CONFORMITY

(EN)

1. Model No.: UPP-110S
2. Name and address of the manufacturer's authorised representative: Sony Belgium, bijkantoor van Sony Europe B.V., Da Vincilaan 7-D1, 1930 Zaventem, Belgium
3. This declaration of conformity is issued under the sole responsibility of the manufacturer: Sony Corporation, 1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan
4. Object of the declaration: Thermal Print Media
5. The object of the declaration described above is in conformity with: 2017/745 (Medical Device Regulation)
6. Where applicable, references to the relevant harmonised standards used or references to the technical specifications in relation to which conformity is declared:
Medical: EN 60601-1:2006 + A1:2013
7. Where applicable, the notified body (name and number), description of intervention and certificate:
8. Additional information:
 - Classification: Class I, MDR Annex VIII
 - Basic UDI-DI: 4901780PrintMediaWR
 - Manufacturer SRN: JP-MF-000014553
 - Authorised Representative SRN: BE-AR-000007876
 - Intended Purpose: Print media for Sony's medical printer
 - Accessory(ies): -
 - Note(s): -

Signed for and on behalf of: Sony Corporation

Tokyo, 2021-11-04

Reference Number: 2021EU02266



Ryogo Katayama
Quality Officer
Quality Officer Group
Quality & Environmental Dept.

Sony Corporation
1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan

SONY

Sony Corporation
1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan

EU DECLARATION OF CONFORMITY

(EN)

1. Model No.: UPP-110HG
2. Name and address of the manufacturer's authorised representative: Sony Belgium, bijkantoor van Sony Europe B.V., Da Vincilaan 7-D1, 1930 Zaventem, Belgium
3. This declaration of conformity is issued under the sole responsibility of the manufacturer: Sony Corporation, 1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan
4. Object of the declaration: Thermal Print Media
5. The object of the declaration described above is in conformity with: 2017/745 (Medical Device Regulation)
6. Where applicable, references to the relevant harmonised standards used or references to the technical specifications in relation to which conformity is declared:
Medical: EN 60601-1:2006 + A1:2013
7. Where applicable, the notified body (name and number), description of intervention and certificate:
8. Additional information:
 - Classification: Class I, MDR Annex VIII
 - Basic UDI-DI: 4901780PrintMediaWR
 - Manufacturer SRN: JP-MF-000014553
 - Authorised Representative SRN: BE-AR-000007876
 - Intended Purpose: Print media for Sony's medical printer
 - Accessory(ies): -
 - Note(s): -

Signed for and on behalf of: Sony Corporation

Tokyo, 2021-11-04

Reference Number: 2021EU02267

Sony Corporation
1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan



Ryogo Katayama
Quality Officer
Quality Officer Group
Quality & Environmental Dept.

SONY

Sony Corporation
1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan

EU DECLARATION OF CONFORMITY

(EN)

1. Model No.: UPP-110HD
2. Name and address of the manufacturer's authorised representative: Sony Belgium, bijkantoor van Sony Europe B.V., Da Vincilaan 7-D1, 1930 Zaventem, Belgium
3. This declaration of conformity is issued under the sole responsibility of the manufacturer: Sony Corporation, 1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan
4. Object of the declaration: Thermal Print Media
5. The object of the declaration described above is in conformity with: 2017/745 (Medical Device Regulation)
6. Where applicable, references to the relevant harmonised standards used or references to the technical specifications in relation to which conformity is declared:
Medical: EN 60601-1:2006 + A1:2013
7. Where applicable, the notified body (name and number), description of intervention and certificate:
8. Additional information:
 - Classification: Class I, MDR Annex VIII
 - Basic UDI-DI: 4901780PrintMediaWR
 - Manufacturer SRN: JP-MF-000014553
 - Authorised Representative SRN: BE-AR-000007876
 - Intended Purpose: Print media for Sony's medical printer
 - Accessory(ies): -
 - Note(s): -

Signed for and on behalf of: Sony Corporation

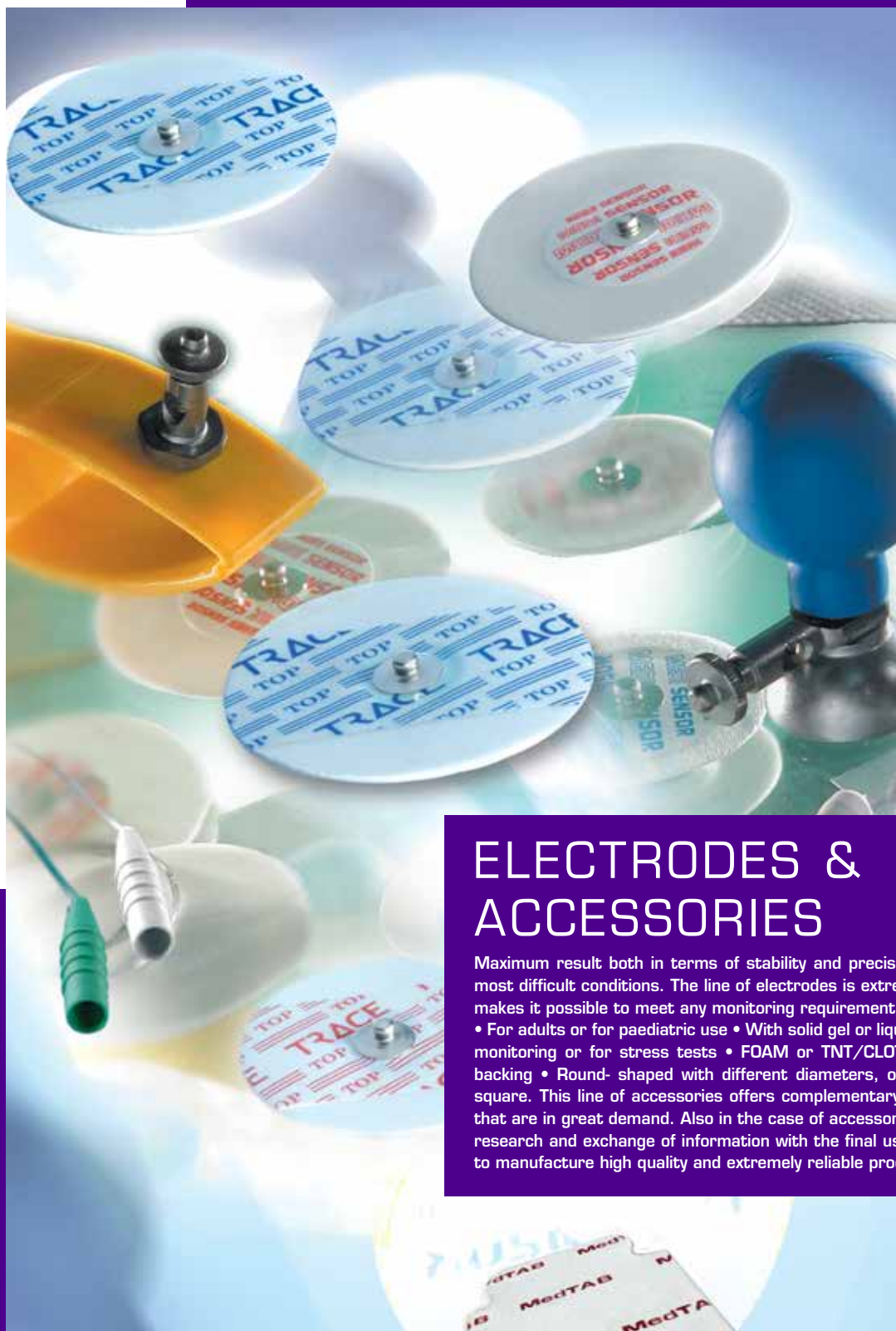
Tokyo, 2021-11-10

Reference Number: 2021EU02280

Sony Corporation
1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan



Ryogo Katayama
Quality Officer
Quality Officer Group
Quality & Environmental Dept.



ELECTRODES & ACCESSORIES

Maximum result both in terms of stability and precision, also under the most difficult conditions. The line of electrodes is extremely complete and makes it possible to meet any monitoring requirement.

- For adults or for paediatric use
- With solid gel or liquid gel
- For normal monitoring or for stress tests
- FOAM or TNT/CLOTH or MICROPORE backing
- Round- shaped with different diameters, oval, rectangular or square. This line of accessories offers complementary or similar articles that are in great demand. Also in the case of accessories, the continuous research and exchange of information with the final user have allowed us to manufacture high quality and extremely reliable products.

TOP TRACE LINE

MONITORING & RESTING ECG'S



NM 50 RFI

Ø 50 mm.	Dimensions	Ø 55 mm.
Foam	Backing	Foam
Solid	Gel	Liquid
Ag/AgCl	Sensor	Ag/AgCl



NM 55 RLI

40 pcs / pouch - 800 pcs / box - 9600 pcs / carton

Packaging

25 pcs / pouch - 500 pcs / box - 6000 pcs / carton

The most typical sizes. The most demanded supports. A proved quality always useful.



NM 42 DFI

45x42 mm.	Dimensions	42x36 mm.
Foam	Backing	Foam
Solid	Gel	Solid
Ag/AgCl	Sensor	Ag/AgCl



NM 3642 OFI

50 pcs / pouch - 1200 pcs / box - 14400 pcs / carton

Packaging

50 pcs / pouch - 1200 pcs / box - 14400 pcs / carton

Two multifunctional models: suitable for adult and pediatric ECG's.



NP 30 RFI

Ø 30 mm.	Dimensions	28x44 mm.
Foam	Backing	Foam
Solid	Gel	Solid
Ag/AgCl	Sensor	Ag/AgCl



NM 2844 RFS

30 pcs / pouch - 900 pcs / box - 10800 pcs / carton
The classical and largely used electrode for pediatric ECG's.
Excellent quality ensured also on "critical" patients.

Packaging

50 pcs / pouch - 1250 pcs / box - 15000 pcs / master carton



NM 3351 OFI

51x33 mm.	Dimensions	28x44 mm.
Foam	Backing	Foam
Solid	Gel	Solid
Ag/AgCl	Sensor	Ag/AgCl - Banana plug



NM 2844 FBI

30 pcs / pouch - 900 pcs / box - 10800 pcs / carton

Packaging

30pcs / pouch - 600pcs / box - 7200pcs / carton

TOP TRACE LINE

STRESS TEST



ST 50 RFI

Ø 50 mm.	Dimensions	Ø 50 mm.
Foam	Backing	Foam
Solid	Gel	Liquid
Ag/AgCl	Sensor	Ag/AgCl

40pcs/pouch-800pcs/box- 9600pcs/carton

Packaging

ST 50 RLI



30 pcs / pouch - 600 pcs / box - 7200 pcs / carton

The most suitable electrodes for ECG's under difficult conditions.
Easy to apply and use • Strong adhesiveness • Do not irritate the skin • Highly conductive gel

MRI NON - MAGNETIC



RT 50 RFI

Ø 50 mm.	Dimensions	23x34 mm.
Foam	Backing	Silicone coated pet film
Solid	Gel	Solid
Ag/AgCl	Sensor	Ag/AgCl

30pcs/pouch-600pcs/box- 7200pcs/carton

Packaging

MEDTAB



100 pcs / pouch - 500 pcs / box - 5000 pcs / carton

This electrode entirely replaces suction chests and clamps to patient's advantage both in terms of hygiene and health. To be attached to alligator clip connectors.

MONITORING / PRE-WIRED ELECTRODES



RE 30 LWF

30 mm. female 4 mm	Dimensions	
Foam	Backing	Foam
Solid	Gel	Solid
Ag/AgCl	Sensor	Ag/AgCl

3 pcs / pouch - 300 pcs / box

Available also with 22 mm diameter Model RE 22 LWF (with 4 mm female socket)

Packaging

RE 22 LWS

22 mm.
safety 1,55 mm

3 pcs / pouch - 300 pcs / box

Available also with 30 mm diameter Model RE 30 LWS (with 1,55 mm safety plug)



RT 30 LWF

30 mm. female 4 mm	Dimensions	
Foam	Backing	Foam
Solid	Gel	Solid
Carbon	Sensor	Carbon

3 pcs / pouch - 300 pcs / box

Available also with 22 mm diameter Model RT 22 LWF (with 4 mm female socket)

Packaging

RT 22 LWS

22 mm.
safety 1,55 mm

3 pcs / pouch - 300 pcs / box

Available also with 30 mm diameter Model RT 30 LWS (with 1,55 mm safety plug)



MEDI SENSOR LINE

MONITORING & RESTING ECG'S



MODEL F 50-S

Ø 50 mm.	Dimensions	Ø 26 mm.
Foam	Backing	Foam
Solid	Gel	Solid
Ag/AgCl	Sensor	Ag/AgCl

30 pcs / pouch – 1200 pcs / carton

Available also with liquid gel Model F 50-L

F 26-S



50 pcs / pouch – 2000 pcs / carton

HOLTER / LONG MONITORING



MODEL W 601-S

Ø 55 mm.	Dimensions	Ø 55 mm..
TNT	Backing	MICROPORE
Solid	Gel	Solid
Ag/AgCl	Sensor	Ag/AgCl

30 pcs / pouch – 1500 pcs / carton

Available also with Ø 50 mm Model W 601
and also with liquid gel Model W 601-L

MODEL T 601-S



30 pcs / pouch – 1500 pcs / carton

STRESS TEST



MODEL FS 50-L

Ø 50 mm.	Dimensions	Ø 55 mm.
Foam	Backing	Foam
Liquid	Gel	Liquid
Ag/AgCl	Sensor	Ag/AgCl

30 pcs / pouch – 1200 pcs / carton

MODEL F 55-L



30 pcs / pouch – 1200 pcs / carton



MODEL FS 501-S

Ø 35x45 mm	Dimensions	Ø 50 mm
Foam	Backing	Foam
Solid	Gel	Solid
Ag/AgCl	Sensor	Carbon

50 pcs / pouch – 2000 pcs / carton

MULTIFUNCTION

MODEL F 50-S BLUE

Ø 50 mm

Foam

Solid

Carbon

30 pcs / pouch – 1200 pcs / carton

Ideal for all ECG monitoring, radiolucent application included!
AVAILABLE ALSO FOR PEDIATRIC, Ø 26 mm, MODEL F 26-S BLUE



SUCTION CHEST ELECTRODES



Ag/AgCl sensor to guarantee maximum reliability in signal transmission.

- Available in three sizes: Ø15 mm for pediatric use / Ø 24 mm and Ø 30 mm for adult use.

RUBBER LIMB & THORACIC STRAPS



For attaching electrodes • for limb use - one hole (15 mm x 40 cm / 25 mm x 45 cm), - two holes (32mm x 45cm) - for thoracic use - one hole (32mm x 135cm), six holes (100mm x 135cm).

CLAMP ELECTRODES



Classical-designed clamps with an improved quality. Ag/AgCl to ensure excellent signal transmission • Silver component to prevent distortions and to increase efficiency • Universal connection.

ELECTRO- SURGICAL PLATES



ELECTROSURGICAL PLATES WITH A SUPERIOR QUALITY

• Made of FOAM suitable to obtain perfect results with all instruments • manufactured and tested in compliance with EN 60601-1/EN 60601-2/AAMI-HF 18 • available with one zone and two zones and REM alarm for adults (adhesive surface 215 sq cm), for children (surface 129 sq cm) and for new born (surface 63 sq cm). **All the models are available even equipped with cables.**

DEFIBRILLATOR PADS



A wide range of pads suitable for all the defibrillators available. Raw material quality and attention to details resulted into an absolutely reliable range of products. Universal without cable or equipped with cable, having a specific connector to comply with all instruments.



ELECTRODES FOR
TENS AND EMG

• Superior quality • available with single or double cable • lead wire with 2 mm socket or press-stud connection • manufactured in compliance with EN ISO 9001 and AAMI standards.

PULSE OXIMETER
SENSORS

A broad range of sensors to monitor and detect the arterial oxygen saturation. Excellent quality and proven accuracy are the characteristics that better describe a whole series of disposable and reusable sensors. Suitable for all the main fittings spread on the market.

ELECTRODES &
ACCESSORIES
FOR EEG & EMG

A) BRIDGE ELECTRODES Made of silver/silver chloride to optimize signal transmission. Slightly curved support to improve head adaption.

B) SILICONE HEADCAPS With 2 or 4 adjustable bands to fit for various requirements. Anatomically-designed earmuffs. Available in three sizes [Small-Medium-Large];

C) SPECIAL INK FOR EEG Suitable & excellent for all devices [Siemens excluded], allowing a perfect trace without blurs. Black colour- 1 litre bottle

D) DISC ELECTRODES Available in silver/silver chloride or gold - 10 mm diameter- With 120-150-200-250 cm lead wire & DIN 42802 safety plug - 10 electrodes per set with coloured lead wires (10 colours available);

E) DISPOSABLE CONCENTRIC NEEDLES FOR ELECTROMYOGRAPHY Ergonomic handgrip and extremely sharp point - Equipped with wings for automatic unfastening -Shielded, light & flexible lead wire with 5 pole DIN 45322 connector and entirely shielded handgrip. Different diameters (from 0.30 to 0.60 mm) and different lengths (from 25 to 75 mm) - 25 pcs per packing.

MOUTHPIECES



Made of material particularly resistant to saliva or manipulation, perfectly hygienic, without residues and non-allergic • can be used on most spirometers of the most important brands present on the market • available in the external diameters 22-26-28-30 and 34 mm, but in case of large orders the customer can request a specific diameter • boxes of 500 pcs • an extremely handy dispenser is available on customer's demand.



Via Secondo Casadei N.14
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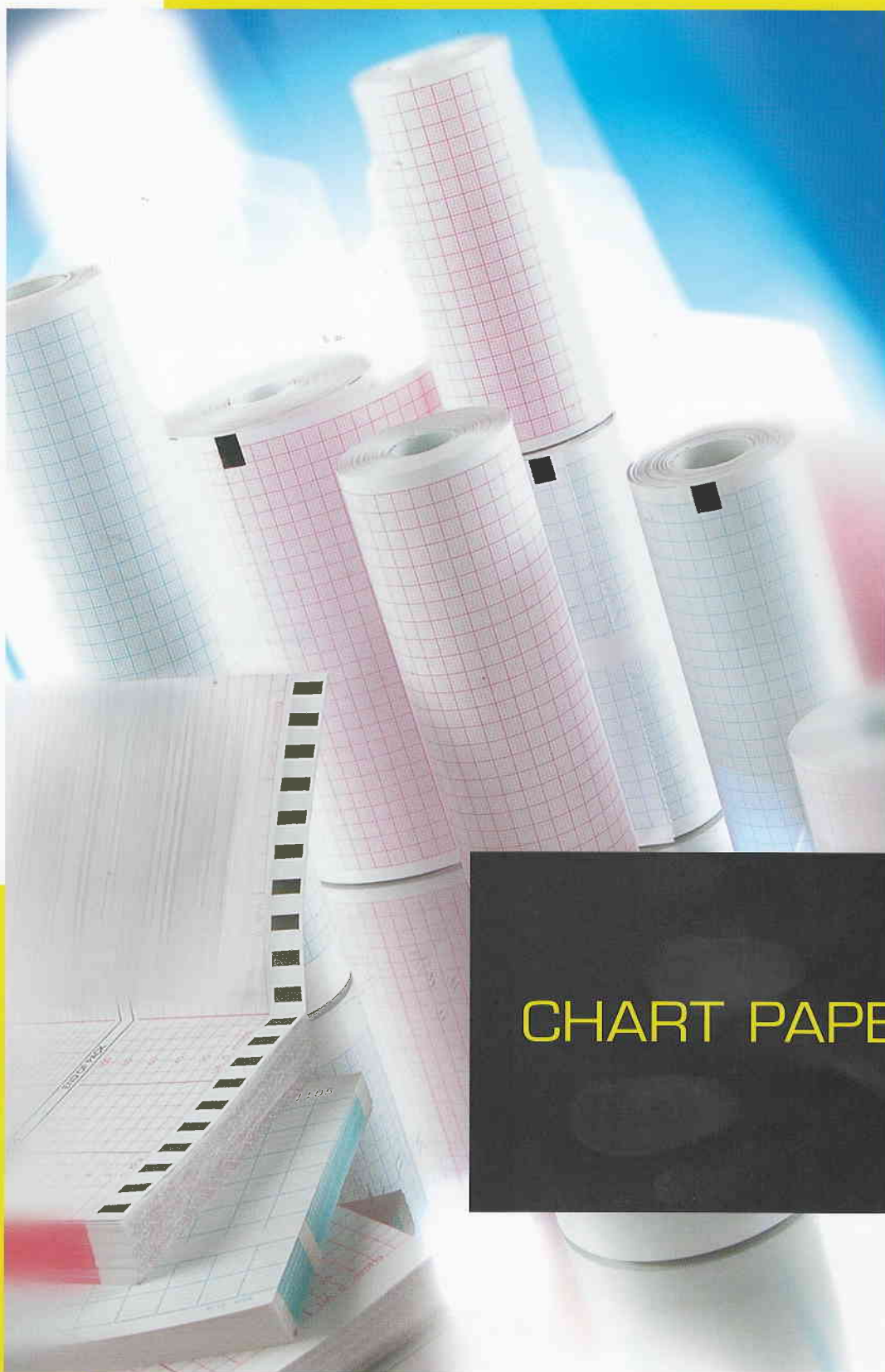
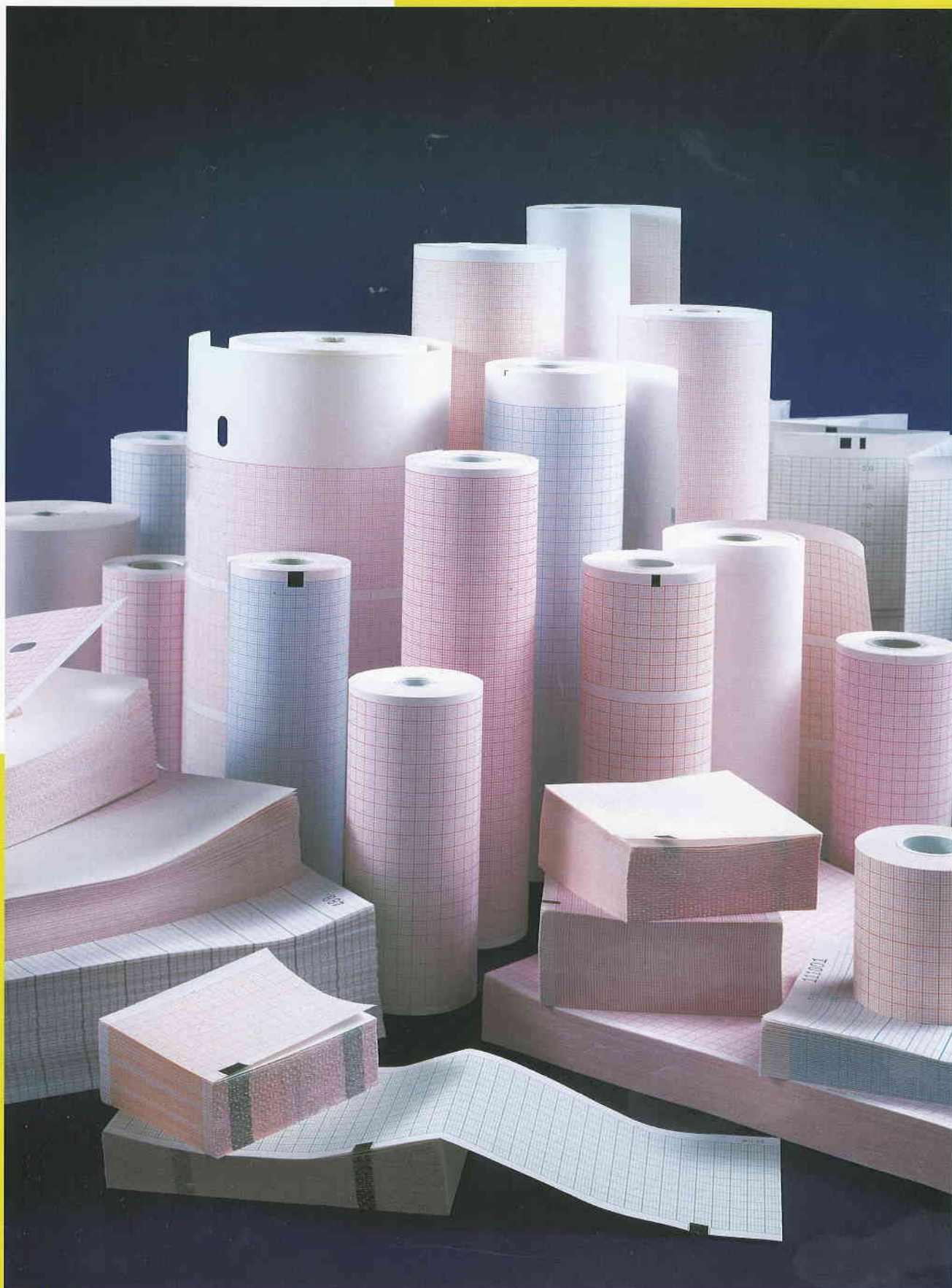


CHART PAPERS

 CEROCARTA



Widest choice of articles for a variety of sectors including: - ELECTROCARDIOGRAMS - ELECTROENCEPHALOGRAMS - FOETAL MONITORING - NEUROLOGY - GYNAECOLOGY - PNEUMOLOGY - INTENSIVE CARE UNITS - POLIGRAPHS - RESUSCITATION - ULTRASOUND SCANS - LABORATORY ANALYSES

If requesting an offer, please state: model and instrument code, where possible the paper code and, in any case, the sizes. For eeg z-folds colour and line distance.

PAPERS FOR ECG



FROM THE SIMPLEST SINGLE-CHANNEL INSTRUMENTS TO THE MOST COMPLEX APPLIANCES FOR STRESS TESTS OR FOR PARTICULAR MONITORING OF CORONARY UNITS, THE BEST SOLUTION FOR ALL QUALITY AND OPERATION REQUIREMENTS CAN BE FOUND IN THE ARTICLES PRODUCED BY CERACARTA. RESEARCH AND STUDY OF THE BEST SUPPORT FOR EACH SPECIFIC APPLIANCE DERIVES FROM AN IN-DEPTH KNOWLEDGE OF THE INSTRUMENTS ON WHICH VARIOUS TYPES OF PAPER ARE USED.

Furthermore constant checks, detailed control and regular technological updating ensure continuously high efficiency in recording. For particular requirements we can supply paper which guarantees conservation of the recording trace for more than 20 years. Ceracarta always guarantees the absolute reliability of its paper with respect to all the instruments available commercially, even the less common.



PAPERS FOR EEG



IN VIEW OF THE PARTICULAR NATURE OF EEG INSTRUMENTS AND THE DIVERSITY OF THEIR FUNCTIONS, PAPER MANUFACTURERS MUST BE EXTREMELY CAREFUL WHEN EVALUATING THE VARIOUS CHARACTERISTICS OF THE APPLIANCE AND BE VERY FLEXIBLE IN VARYING BASIC RAW MATERIALS.

A great priority must be given to the perfect running of the paper and to its excellent receptivity as for the writing pen. Through its paper Ceracarta constantly guarantees best results also in this sector, even during long recordings, thanks to its perfect adaptability, specifically and attentively researched, for all types of instruments.



PAPERS FOR CTG



THE EXTREMELY PRECISE PRINTING OF DIAGRAMS AND BASE SCALES AND THE GAUGED SENSITIVITY OF THERMAL PAPER ARE ESSENTIAL FOR THE BEST FUNCTIONING OF THE FOETAL MONITORING EQUIPMENT AND, THEREFORE, AN EXCELLENT DIAGNOSTIC READING.

Ceracarta has paid special attention to these details, manufacturing products which, also due to the perfection of the lateral holes and the numbering of each page, and owing to the accuracy of the regulating sensors, are among the best available commercially, where too often paper of insufficient efficiency is on offer. Ceracarta represents a guarantee, not found everywhere, also in the sector of foetal monitoring.



PAPERS FOR ANALYSIS LABORATORY AND LABELS



THE APPARENT SIMPLICITY OF PAPERS FOR INSTRUMENTS OF ANALYSIS HAS OFTEN INDUCED CONSUMERS TO BE EASILY SATISFIED TO THE DETRIMENT OF THE QUALITY OF THE REPORT AND, ESPECIALLY OF THE CONSERVATION OVER TIME OF

DOCUMENTATION RELATING TO THE ANALYSIS.

Also in this field, specialisation is crucial and the quality of the raw materials used is of a primary importance. Ceracarta is at your disposal with rolls and packs for all types of instruments used in laboratories, guaranteeing optimal functioning and a perfect result.

CERACARTA PRODUCES THERMAL PAPER LABELS, PAPERS FOR THERMAL TRANSFER PRINTERS AND POLYESTER FILMS FOR ANALYSIS LABORATORIES AND TRANSDUCTION CENTRES. MANUFACTURED WITH SPECIAL MATERIALS AND ADHESIVES, FOR ALL REQUIREMENTS.



Whatever the use (labels for test tubes, medical reports, radiological envelopes or "blood bags"), an excellent outcome is ensured. Ceracarta has implemented a division for the production of labels and tickets with a microchip (RFID technology) to be applied also to the medical field.





THE COMPANY CERACARTA

- We are leaders in Europe for the manufacturing of paper for electrical diagnostic instruments and industrial recording with more than 40 years' experience in the medical sector.
- Advanced technologies, equipment and methodical quality controls for the production of specific chart and non-chart paper for all diagnostic equipment available on the international market.
- Superior quality materials and absolute manufacturing precision for perfect diagnosis.
- Study and research centre to meet any specific need or customisation requirement and client consulting service.
- Documents and samples sent on request.
- ISO 9002 - EC certification - 2000 vision / ISO46002 / 93-42-CE



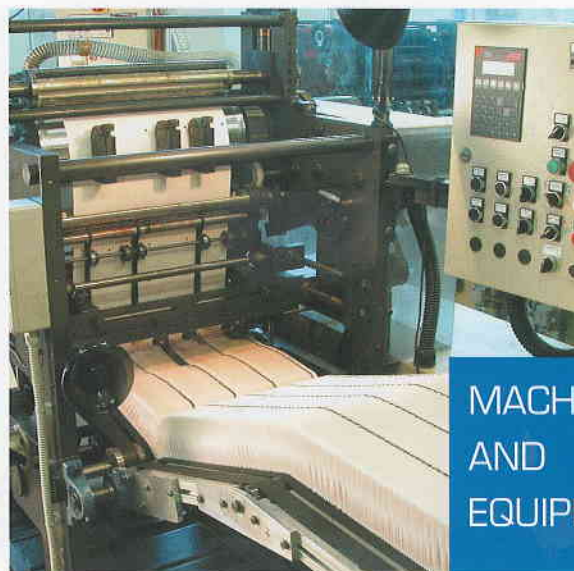
THE WAREHOUSE

WAREHOUSE

Perfect service offered to customers in terms of guaranteeing delivery times, the great care paid when choosing the most suitable form of packaging for the type of shipment, as well as the perfect correspondence between the product supplied and what is requested. • Optimal management of the stock related to production programmes: all this is guaranteed by a computerised system which now supports the traditional bar code, the revolutionary and brand-new radio frequency transponder system (TAG). • Professional skill and competence of our staff ensure an ideal service for customers.



Products EC compliant



MACHINES AND EQUIPMENT

MACHINES AND EQUIPMENT

After the new premises were set up, the machinery has been almost completely replaced or modified in order to improve the quality of the product even further, while at the same time speeding up production times which means lower costs for the customer. The whole production system has the advantage of a double check: from the specialised technical staff and the automated computer control.



Via Secondo Casadei N. 14
zona industriale Villa Selva - 47100 Forlì (Italy) tel. +39.0543.780055 - fax +39.0543.781404
www.ceracarta.it - e mail: info@ceracarta.it

COMPANY
WITH QUALITY SYSTEM
CERTIFIED BY DNV
ISO 9000



www.imq.it



IQNet, the association of the world's first class certification bodies, is the largest provider of management System Certification in the world. IQNet is composed of more than 30 bodies and counts over 150 subsidiaries all over the globe.

**CERTIFICATO N.
CERTIFICATE N. 9190.CRC3**

SI CERTIFICA CHE IL SISTEMA DI GESTIONE PER LA QUALITA' DI
WE HEREBY CERTIFY THAT THE QUALITY MANAGEMENT SYSTEM OPERATED BY

CERACARTA SPA
VIA SECONDO CASADEI 14 Z.I. VILLA SELVA - 47122 FORLI' (FC)

UNITA' OPERATIVE / OPERATIVE UNITS

Vedere gli Allegati per le Unità Operative (n. 2 pagine)
View the Annexes for the Operative Units (n. 2 pages)

E' CONFORME ALLA NORMA / IS IN COMPLIANCE WITH THE STANDARD

ISO 9001:2015

PER LE SEGUENTI ATTIVITA' / FOR THE FOLLOWING ACTIVITIES

Produzione e stampa di carte speciali e diagrammate per registrazione ad uso industriale, ferroviario, medicale e biglietteria anche conto terzi. Produzione e stampa di etichette e biglietti anche a lettura/scrittura in radio frequenza (RFID). Sviluppo e produzione di creme, gel sterile e non sterile per applicazioni elettrodiagnostiche ad ultrasuoni, anche conto terzi. Commercializzazione ed immissione in commercio di accessori per applicazioni elettrodiagnostiche, ad ultrasuoni e per strumenti elettromedicali. Produzione di elettrodi per ECG. Gestione della produzione ed immissione in commercio di elettrodi per ECG. Immissione in commercio di piastre per elettrobisturi e defibrillatori. Commercializzazione di video stampanti, carte fotografiche per video stampanti, stampanti e relativi materiali di consumo ed accessori

Manufacture and print of special recording chart papers for industrial, railway, medical use and ticketing also on behalf of third parties. Manufacture and print of labels and tickets also radio frequency reading/writing (RFID).

Development and manufacture of creams, gels sterile and not sterile for electromedical and ultrasound procedures also on behalf of third parties. Trade and placing on the market of accessories for electromedical and ultrasound diagnostic devices and for electromedical equipment. Manufacture of electrodes for ECG.

Production management and placing on the market of electrodes for ECG. Placing on the market of electrosurgical plates and defibrillation pads. Trade of videoprinters, photographic papers for videoprinters, printers and related consumable and accessories

Ulteriori informazioni riguardanti l'applicabilità dei requisiti ISO 9001:2015 possono essere ottenute consultando l'organizzazione
Further clarifications regarding the applicability of ISO 9001:2015 requirements may be obtained by consulting the organization

IL PRESENTE CERTIFICATO E' SOGGETTO AL RISPETTO DEL
REGOLAMENTO PER LA CERTIFICAZIONE DEI SISTEMI DI GESTIONE
THE USE AND THE VALIDITY OF THE CERTIFICATE SHALL SATISFY THE
REQUIREMENTS OF THE RULES FOR CERTIFICATION OF MANAGEMENT SYSTEMS

DATE:	PRIMA CERTIFICAZIONE FIRST CERTIFICATION 2002-11-26	EMISSIONE CORRENTE CURRENT ISSUE 2020-09-29	SCADENZA EXPIRY 2023-10-07
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IMQ S.p.A. - VIA QUINTILIANO, 43 - 20138 MILANO ITALY
Management Systems Division - Flavio Ornago



SGQ N° 005 A

Membro degli Accordi di Mutuo
Riconoscimento EA, IAF e ILAC
Signatory of EA, IAF and ILAC
Mutual Recognition Agreements

IAF: 07, 09, 19, 29, 12

La validità del certificato è subordinata a sorveglianza annuale e riesame completo
del Sistema di Gestione con periodicità triennale
The validity of the certificate is submitted to annual audit and a reassessment
of the entire management System within three years.



Organismo di Certificazione Federato CISQ
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CISQ è la Federazione Italiana di Organismi di
Certificazione dei sistemi di gestione aziendale.
CISQ is the Italian Federation of management
system Certification Bodies.



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ALLEGATO N. 9190.CRC3-1
ANNEX N.



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CERACARTA SPA

VIA SECONDO CASADEI 14 Z.I. VILLA SELVA - 47122 FORLÌ (FC)

Attività:
Activities:

Produzione e stampa di carte speciali e diagrammate per registrazione ad uso industriale, ferroviario, medicale e biglietteria anche conto terzi. Produzione e stampa di etichette e biglietti anche a lettura/scrittura in radio frequenza (RFID). Sviluppo e produzione di creme, gel sterile e non sterile per applicazioni elettrodiagnostiche e ad ultrasuoni, anche conto terzi. Commercializzazione ed immissione in commercio di accessori per applicazioni elettrodiagnostiche, ad ultrasuoni e per strumenti elettromedicali. Produzione di elettrodi per ECG. Gestione della produzione ed immissione in commercio di elettrodi per ECG. Immissione in commercio di piastre per elettrobisturi e defibrillatori. Commercializzazione di video stampanti, carte fotografiche per video stampanti, stampanti e relativi materiali di consumo ed accessori.

Manufacture and print of special recording chart papers for industrial, railway, medical use and ticketing also on behalf of third parties. Manufacture and print of labels and tickets also radio frequency reading/writing (RFID). Development and manufacture of creams, gels sterile and not sterile for electromedical and ultrasound procedures also on behalf of third parties. Trade and placing on the market of accessories for electromedical and ultrasound diagnostic devices and for electromedical equipment. Manufacture of electrodes for ECG. Production management and placing on the market of electrodes for ECG. Placing on the market of electrosurgical plates and defibrillation pads. Trade of videoprinters, photographic papers for videoprinters, printers and related consumable and accessories

IL PRESENTE ALLEGATO HA LO SCOPO DI ESPlicitARE LE ATTIVITA' SVOLTE PRESSO IL SINGOLO SITO/UNITA' OPERATIVA NELL'AMBITO DELLA CERTIFICAZIONE DEL SISTEMA DI GESTIONE RILASCIATA A CERACARTA SPA

THE AIM OF PRESENT ANNEX IS TO EXPLAIN THE ACTIVITIES PERFORMED IN EACH SITE/OPERATIVE UNIT OF THE MANAGEMENT SYSTEM CERTIFICATION ISSUED TO CERACARTA SPA

PER LA VALIDITA' RIFERIRSI AL CERTIFICATO N. 9190.CRC3
FOR THE VALIDITY PLEASE REFER TO CERTIFICATE N. 9190.CRC3

DATE:	PRIMA CERTIFICAZIONE FIRST CERTIFICATION	EMISSIONE CORRENTE CURRENT ISSUE	SCADENZA EXPIRY
	2002-11-26	2020-09-29	2023-10-07


IMQ S.p.A. - VIA QUINTILIANO, 43 - 20138 MILANO ITALY
Management Systems Division - Flavio Ornago



www.cisq.com



SGQ N° 005 A

Membro degli Accordi di Mutuo Riconoscimento EA, IAF e ILAC
Signatory of EA, IAF and ILAC
Mutual Recognition Agreements

Il presente documento integra il certificato n. 9190.CRC3
This document is a part of certificate n. 9190.CRC3

IAF: 07, 09, 19, 29, 12

La validità del certificato è subordinata a sorveglianza annuale e riesame completo del Sistema di Gestione con periodicità triennale
The validity of the certificate is submitted to annual audit and a reassessment of the entire management system within three years

Organismo di Certificazione Federato CISQ
www.imq.it

CISQ è la Federazione Italiana di Organismi di Certificazione dei sistemi di gestione aziendale.
CISQ is the Italian Federation of management system Certification Bodies.



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ALLEGATO N. 9190.CRC3-2
ANNEX N.



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CERACARTA SPA

VIA GRAMADORA 12/14 - 47122 FORLÌ (FC)

Attività:
Activities:

Produzione di creme, gel sterile e non sterile per applicazioni elettrodiagnostiche e ad ultrasuoni, anche conto terzi
Manufacture of creams, gels sterile and not sterile for electromedical and ultrasound procedures also on behalf of third parties

IL PRESENTE ALLEGATO HA LO SCOPO DI ESPLICITARE LE ATTIVITA' SVOLTE PRESSO IL SINGOLO SITO/UNITA' OPERATIVA NELL'AMBITO DELLA CERTIFICAZIONE DEL SISTEMA DI GESTIONE RILASCIATA A CERACARTA SPA

THE AIM OF PRESENT ANNEX IS TO EXPLAIN THE ACTIVITIES PERFORMED IN EACH SITE/OPERATIVE UNIT OF THE MANAGEMENT SYSTEM CERTIFICATION ISSUED TO CERACARTA SPA

PER LA VALIDITA' RIFERIRSI AL CERTIFICATO N. 9190.CRC3
FOR THE VALIDITY PLEASE REFER TO CERTIFICATE N. 9190.CRC3

DATE:	PRIMA CERTIFICAZIONE FIRST CERTIFICATION	EMISSIONE CORRENTE CURRENT ISSUE	SCADENZA EXPIRY
	2002-11-26	2020-09-29	2023-10-07

IMQ S.p.A. - VIA QUINTILIANO, 43 - 20138 MILANO ITALY
Management Systems Division - Flavio Ornago



SGQ N° 005 A

Membro degli Accordi di Mutuo Riconoscimento EA, IAF e ILAC
Signatory of EA, IAF and ILAC Mutual Recognition Agreements

Il presente documento integra il certificato n. 9190.CRC3
This document is a part of certificate n. 9190.CRC3

IAF: 12

La validità del certificato è subordinata a sorveglianza annuale e riesame completo del Sistema di Gestione con periodicità triennale
The validity of the certificate is submitted to annual audit and a reassessment of the entire management System within three years



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CISQ is the Italian Federation of management system Certification Bodies.



THE INTERNATIONAL CERTIFICATION NETWORK

CERTIFICATE

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

CERACARTA SPA

VIA SECONDO CASADEI 14 Z.I. VILLA SELVA - 47122 FORLÌ (FC)

VIA GRAMADORA 12/14 - 47122 FORLÌ (FC)

has implemented and maintains a

Quality Management System

for the following scope:

Manufacture and print of special recording chart papers for industrial, railway, medical use and ticketing also on behalf of third parties. Manufacture and print of labels and tickets also radio frequency reading/writing (RFID). Development and manufacture of creams, gels sterile and not sterile for electromedical and ultrasound procedures also on behalf of third parties. Trade and placing on the market of accessories for electromedical and ultrasound diagnostic devices and for electromedical equipment. Manufacture of electrodes for ECG. Production management and placing on the market of electrodes for ECG. Placing on the market of electrosurgical plates and defibrillation pads. Trade of videoprinters, photographic papers for videoprinters, printers and related consumable and accessories

Further clarifications regarding the applicability of ISO 9001:2015 requirements may be obtained by consulting the organization

which fulfills the requirements of the following standard:

ISO 9001:2015

Issued on: 2020 - 09 - 29

Expires on: 2023 - 10 - 07

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 - 112265



Alex Stoichitoiu
President of IQNET



Ing. Mario Romersi
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 EAGLE Certification Group USA
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 YUQS Serbia

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



www.imq.it

**CERTIFICATO N.
CERTIFICATE N. 9124.CRC4**

SI CERTIFICA CHE IL SISTEMA DI GESTIONE PER LA QUALITA' DI
WE HEREBY CERTIFY THAT THE QUALITY MANAGEMENT SYSTEM OPERATED BY

CERACARTA SPA

VIA SECONDO CASADEI 14 Z.I. VILLA SELVA - 47122 FORLI' (FC)

UNITA' OPERATIVE / OPERATIVE UNITS

Vedere gli Allegati per le Unità Operative (n. 2 pagine)
View the Annexes for the Operative Units (n. 2 pages)

E' CONFORME ALLA NORMA / IS IN COMPLIANCE WITH THE STANDARD
ISO 13485:2016

PER LE SEGUENTI ATTIVITA' / FOR THE FOLLOWING ACTIVITIES

Produzione e stampa di carte speciali e diagrammate per registrazione ad uso medicale anche conto terzi. Produzione e stampa di etichette ad uso medicale. Sviluppo e produzione di creme, gel sterile e non sterile per applicazioni elettrodiagnostiche e ad ultrasuoni, anche conto terzi. Commercializzazione ed immissione in commercio di accessori per applicazioni elettrodiagnostiche, ad ultrasuoni e per strumenti elettromedicali. Produzione di elettrodi per ECG. Gestione della produzione ed immissione in commercio di elettrodi per ECG. Immissione in commercio di piastre per elettrobisturi e defibrillatori. Commercializzazione di video stampanti, carte fotografiche per video stampanti, stampanti e relativi materiali di consumo ed accessori per uso medicale

Manufacture and print of special recording chart papers for medical use also on behalf of third parties. Manufacture and print of labels for medical use. Development and manufacture of creams, gels sterile and not sterile for electromedical and ultrasound procedures also on behalf of third parties. Trade and placing on the market of accessories for electromedical and ultrasound diagnostic devices and for electromedical equipment. Manufacture of electrodes for ECG. Production management and placing on the market of electrodes for ECG. Placing on the market of electrosurgical plates and defibrillation pads. Trade of videoprinters, photographic papers for videoprinters, printers and related consumable and accessories for medical use

Ulteriori informazioni riguardanti l'applicabilità dei requisiti ISO 13485:2016 possono essere ottenute consultando l'organizzazione
Further clarifications regarding the applicability of ISO 13485:2016 requirements may be obtained by consulting the organization

IL PRESENTE CERTIFICATO E' SOGGETTO AL RISPETTO DEL
REGOLAMENTO PER LA CERTIFICAZIONE DEI SISTEMI DI GESTIONE
THE USE AND THE VALIDITY OF THE CERTIFICATE SHALL SATISFY THE
REQUIREMENTS OF THE RULES FOR CERTIFICATION OF MANAGEMENT SYSTEMS

DATE:	PRIMA CERTIFICAZIONE FIRST CERTIFICATION	EMISSIONE CORRENTE CURRENT ISSUE	SCADENZA EXPIRY
	1999-07-20	2020-09-29	2023-10-07

IMQ S.p.A. - VIA QUINTILIANO, 43 - 20138 MILANO ITALY
Management Systems Division - Flavio Ornago



SGQ N° 005 A

Membro degli Accordi di Mutuo
Riconoscimento EA, IAF e ILAC
Signatory of EA, IAF and ILAC
Mutual Recognition Agreements

La validità del certificato è subordinata a sorveglianza annuale e riesame completo
del Sistema di Gestione con periodicità triennale
The validity of the certificate is submitted to annual audit and a reassessment
of the entire management system within three years



Organismo di Certificazione Federato CISQ
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Certificazione dei sistemi di gestione aziendale.
CISQ is the Italian Federation of management
system Certification Bodies.

CISQ is a member of



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certification bodies, is the largest provider of management
System Certification in the world.
IQNet is composed of more than 30 bodies and counts
over 150 subsidiaries all over the globe.*



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ALLEGATO N. 9124.CRC4-1
ANNEX N.



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CERACARTA SPA

VIA SECONDO CASADEI 14 Z.I. VILLA SELVA - 47122 FORLÌ (FC)

Attività:
Activities:

Produzione e stampa di carte speciali e diagrammate per registrazione ad uso medicale anche conto terzi. Produzione e stampa di etichette ad uso medicale. Sviluppo e produzione di creme, gel sterile e non sterile per applicazioni elettrodiagnostiche e ad ultrasuoni, anche conto terzi.

Commercializzazione ed immissione in commercio di accessori per applicazioni elettrodiagnostiche, ad ultrasuoni e per strumenti elettromedicali. Produzione di elettrodi per ECG.

Gestione della produzione ed immissione in commercio di elettrodi per ECG. Immissione in commercio di piastre per elettrobisturi e defibrillatori. Commercializzazione di video stampanti, carte fotografiche per video stampanti, stampanti e relativi materiali di consumo ed accessori per uso medicale

Manufacture and print of special recording chart papers for medical use also on behalf of third parties. Manufacture and print of labels for medical use. Development and manufacture of creams, gels sterile and not sterile for electromedical and ultrasound procedures also on behalf of third parties. Trade and placing on the market of accessories for electromedical and ultrasound diagnostic devices and for electromedical equipment. Manufacture of electrodes for ECG. Production management and placing on the market of electrodes for ECG. Placing on the market of electrosurgical plates and defibrillation pads. Trade of videoprinters, photographic papers for videoprinters, printers and related consumable and accessories for medical use

IL PRESENTE ALLEGATO HA LO SCOPO DI ESPlicitARE LE ATTIVITA' SVOLTE PRESSO IL SINGOLO SITO/UNITA' OPERATIVA NELL'AMBITO DELLA CERTIFICAZIONE DEL SISTEMA DI GESTIONE RILASCIATA A CERACARTA SPA

THE AIM OF PRESENT ANNEX IS TO EXPLAIN THE ACTIVITIES PERFORMED IN EACH SITE/OPERATIVE UNIT OF THE MANAGEMENT SYSTEM CERTIFICATION ISSUED TO CERACARTA SPA

PER LA VALIDITA' RIFERIRSI AL CERTIFICATO N. 9124.CRC4
FOR THE VALIDITY PLEASE REFER TO CERTIFICATE N. 9124.CRC4

DATE:	PRIMA CERTIFICAZIONE	EMISSIONE CORRENTE	SCADENZA
	FIRST CERTIFICATION	CURRENT ISSUE	EXPIRY
	1999-07-20	2020-09-29	2023-10-07

IMQ S.p.A. - VIA QUINTILIANO, 43 - 20138 MILANO ITALY
Management Systems Division - Flavio Ornaghi



www.cisq.com



SGQ N° 005 A

Membro degli Accordi di Mutuo
Riconoscimento EA, IAF e ILAC
Signatory of EA, IAF and ILAC
Mutual Recognition Agreements

Il presente documento integra il certificato n. 9124.CRC4
This document is a part of certificate n. 9124.CRC4

La validità del certificato è subordinata a sorveglianza annuale e riesame completo
del Sistema di Gestione con periodicità triennale
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of the entire management System within three years

Organismo di Certificazione Federato CISQ
www.imq.it

CISQ è la Federazione Italiana di Organismi di
Certificazione dei sistemi di gestione aziendale.
CISQ is the Italian Federation of management
system Certification Bodies.



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ALLEGATO N. 9124.CRC4-2
ANNEX N.



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CERACARTA SPA

VIA GRAMADORA 12/14 - 47122 FORLI' (FC)

Attività:
Activities:

Produzione di creme, gel sterile e non sterile per applicazioni elettrodiagnostiche e ad ultrasuoni, anche conto terzi
Manufacture of creams, gels sterile and not sterile for electromedical and ultrasound procedures also on behalf of third parties

IL PRESENTE ALLEGATO HA LO SCOPO DI ESPLICITARE LE ATTIVITA' SVOLTE PRESSO IL SINGOLO SITO/UNITA' OPERATIVA NELL'AMBITO DELLA CERTIFICAZIONE DEL SISTEMA DI GESTIONE RILASCIATA A CERACARTA SPA

THE AIM OF PRESENT ANNEX IS TO EXPLAIN THE ACTIVITIES PERFORMED IN EACH SITE/OPERATIVE UNIT OF THE MANAGEMENT SYSTEM CERTIFICATION ISSUED TO CERACARTA SPA

PER LA VALIDITA' RIFERIRSI AL CERTIFICATO N. 9124.CRC4
FOR THE VALIDITY PLEASE REFER TO CERTIFICATE N. 9124.CRC4

DATE:	PRIMA CERTIFICAZIONE <i>FIRST CERTIFICATION</i> 1999-07-20	EMISSIONE CORRENTE <i>CURRENT ISSUE</i> 2020-09-29	SCADENZA <i>EXPIRY</i> 2023-10-07
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IMQ S.p.A. - VIA QUINTILIANO, 43 - 20138 MILANO ITALY
Management Systems Division - Flavio Ornago



SGQ N° 005 A

Membro degli Accordi di Mutuo Riconoscimento EA, IAF e ILAC
Signatory of EA, IAF and ILAC Mutual Recognition Agreements

Il presente documento integra il certificato n. 9124.CRC4
This document is a part of certificate n. 9124.CRC4

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Organismo di Certificazione Federato CISQ
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CISQ is the Italian Federation of management system Certification Bodies.