

TO WHOM IT MAY CONCERN

Whereas **ANTONIO MATACHANA, S.A** who are established and reputable manufacturers of Infection Control equipments, with headquarters in Calle Almogàvers 174-176, 08018 Barcelona (Spain),

do hereby certify that the company **Echipamed Plus SRL**, with business address at Valea Trandafirilor, 24B, of.80 - MD-2001 Chisinau, Republic of Moldova, operates as AUTHORIZED DISTRIBUTOR in the territory of Republic of Moldova.

They are fully authorised to perform after sales activities (quote, sell spare parts) as well carry out maintenance activities at IMSP SCM Sf. Treime" hospital.

This sole agency agreement will be valid until the 31st of December 2019.

And in proof of this, we hereunder sign the present certificate in the city of Barcelona, on the 21st of February 2019

ANTONIO MATACHANA, S.A



Antonio Matachana
Servicio Técnico Internacional
Service & Support
Baurell, 21-29 Nave 4
Polígono Industrial Camí Ral
08860-Castelldefels (Barcelona)
Ramón Torres
Technical Manager Europe & USA

DECLARACIÓN DE CONFORMIDAD CE

EC DECLARATION OF CONFORMITY / DECLARATION DE CONFORMITÉ CE

La empresa: **ANTONIO MATACHANA, S.A.**
The company: **Almogàvers, 174**
La société: **E-08018 – Barcelona**

Declara bajo su única responsabilidad que el equipo,
Declares that, in its own responsibility, the equipment
Déclare sous sa propre responsabilité, que l'appareil,

Marca: **Matachana** **Tipo:** **Esterilizador de vapor.**
Brand / Marque: **Matachana** *Type / Tipe:* **Steam sterilizer / Stérilisateur à la vapeur d'eau.**

Serie: **S1000 Modelos:** **S1004 V-1, S1006 V-1, S1008 V-1, S1010 V-1, S1012 V-1,**
Series / Série Models / Modèles: **S1004 E-1, S1006 E-1, S1008 E-1, S1010 E-1, S1012 E-1,**
S1004 V-2, S1006 V-2, S1008 V-2, S1010 V-2, S1012 V-2,
S1004 E-2, S1006 E-2, S1008 E-2, S1010 E-2, S1012 E-2

siendo productos en el ámbito de la Directiva Europea 93/42/CEE por la que se regulan los productos sanitarios y clasificado como clase IIb según la regla 15 del Anexo IX,

Being devices within the scope of the regulation of the European Directive 93/42/EEC concerning Medical Devices (MDD), and classified as Class IIb according to rule 15 of Annex IX therein,
étant dispositifs dedans le champ d'application de la Directive Européenne 93/42/CEE relative aux dispositifs médicaux, et classifié comme Classe IIb selon la règle 15 de l'Annexe IX,

se hallan en conformidad con:

conforms to / sont conformes à:

- **Directiva 93/42/CEE relativa a los Productos Sanitarios, Anexo I.**
- *Directive 93/42/EEC concerning Medical Devices, Annex I.*
- *Directive 93/42/CEE relative aux dispositifs médicaux, Annexe I.*

Evaluación de la conformidad según:

Conformity assessment according to: / Évaluation de la conformité selon:

- **Directiva 93/42/CEE relativa a los Productos Sanitarios, Anexo II excepto II.4**
- *Directive 93/42/EEC concerning Medical Devices, Annex II without II.4*
- *Directive 93/42/CEE relative aux dispositifs médicaux, Annexe II à l'exception du II.4*

Con número de registro HD 60029459 0001 del Organismo Notificado Europeo:

Under the Registration number
Avec registre numéro

from the European Notified Body number:
de l'Organisme Notifié Européen :

TÜV Rheinland LGA Products GmbH
An Grauen Stein
D-51105 Köln



Además de la Directiva de Productos Sanitarios 93/42/CEE los productos mencionados o sus componentes cumplen con los requisitos esenciales de las siguientes Directivas que son relevantes:

Additionally to the Medical Device Directive 93/42/EEC the a. m. devices or the included components fulfil the essential requirements of the following relevant Directives:

En addition à la Directive relative aux Dispositifs Médicaux 93/42/CEE les dispositifs mentionnés si dessus les exigences essentielles des suivantes Directives relevantes:

EMC 2004/108/EC, LVD 2006/95/EC, MD 2006/42/EC, PED 97/23/EC

Estando disponibles las correspondientes Declaraciones o Certificados de Conformidad emitidos por los fabricantes.

Respective Declarations or Certificates of Conformity issued by the manufacturers are available.

Étant disponibles les respectives Déclarations ou Certifications de Conformité livrés par les fabricants

Nombre:

Jose Carlos López

Sagué

Name / Nom:

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Cargo:

Director Industrial

Title / Position:

Director de Calidad

Lugar y fecha:

Barcelona, 14 de mayo de 2010

Place and date / lieu et date: