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 Reg. Imprese Forlì-Cesena n. 3422 REA n. 72846 / n. FO 006868

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## DECLARATION OF CONFORMITY

of the device named "GELS & CREAMS", "ECG, EEG & TENSE ELECTRODES", "MOUTHPIECE  
 "PLATES FOR ELECTROSURGICAL", "SPECULUM" and "OTHER ACCESSORIES" produced  
 company Ceracarta on the basis of the essential requirements, see enclosure I of the di  
 93/42/CEE, as prescribed in enclosure VII of the above directive.

The writing company Ceracarta located in Via secondo Casadei, 14 Forlì, manufacturer of the product  
 named, " GELS & CREAMS", "ECG, EEG & TENSE ELECTRODES", "MOUTHPIECES",  
 "PLATES FOR ELECTROSURGICAL", "SPECULUM" and "OTHER ACCESSORIES", *declares*  
*under its own responsibility that such a device satisfies all the requirements of directive 93/42/CEE*  
*about medical devices and in particular that:*

- the Device in object satisfies the essential requirements as per in Enclosure I of Directive 93/42/CEE;
- the Device in object must be considered as belonging to Class I;
- the Device 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;
- The manufacturer has prepared and keeps the technical files updated in accordance with enclosure VII, section 3 of the directive itself.
- Such documentation is available at the headquarters of Ceracarta, for any reference by the entitled bodies.

Dr. F.A. Fusconi  
 Marketing & Sales Manager  
 Ceracarta Spa

**CERACARTA s.p.a.**