BRIDGE ELECTRODES
FOR TUBULAR ELASTIC EEG HEAD CAPS
RECORDING AREA: Ø 10 mm
REUSABLE

Suitable for tubular EEG head caps.
The metallic part of the electrode is made of massive silver and coated in Ag/AgCl.
The final part of these bridge electrodes is a pin (which requires an alligator clip).
The electrodes come with a cotton pad on the recording area.

Packaging: box containing 5 pieces.

CODE	SIZE	ELECTRODE END	
CUPT10C	small	pin	
CUPT20C	medium	pin	
CUPT30C	large	pin	

ALLIGATOR CLIP CONNECTION CABLE FOR EEG BRIDGE ELECTRODES

REUSABLE

This cable is suitable for quick connection to the bridge electrodes with a pin.
A plastic part located between cable and alligator clip makes the cable tangle-free.

Packaging: box containing 1 piece.

CODE	*CABLE LENGTH*	*ELECTRODE END*
CUCDP1-09	0.9 m	alligator clip
CUCDP1-12	1.2 m	alligator clip

COLOUR: Yellow	*More available colours*

Connection cable: extra-flexible antistatic TINSEL

CONNECTION
electrode end: alligator clip
EEG box end: Ø 1.5 mm touchproof DIN42802

SEI EMG srl	**DECLARATION OF CONFORMITY**
DEVICE: BRIDGE *FOR EEG RECORDING*	**DC_009en**

Declaration of conformity of the device referred as "BRIDGE for EEG recording" manufactured by the company SEI EMG srl (already STRUMENTAZIONE ELETTRONICA INDUSTRIALE di Didonè Ing. Giorgio & C. snc), to the **essential requirements of Annex I and Annex VII of Directive 93/42/EEC as amended by Directive 2007/47/EC.**

The writing firm **SEI EMG s.r.l.** established in Cittadella, Via S. Chiara 10,
VAT Number: IT-01597140282
manufacturer of the device family called **CUPTxxx**
declare under our sole responsibility that this medical device satisfies all applicable provisions in the Medical Device Directive 93/42/EEC amended by Directive 2007/47/EC.

For this purpose, the undersigned represent and warrant under his own responsibility:

- The device in question meets the essential requirements of Annex I to Directive 93/42/EEC updated by Directive 2007/47/EC;
- the device is considered in Class I;
- the device is sold in packs NOT STERILE;
- the device is not an instrument of measure;
- the device is not intended for clinical investigations;
- the manufacturer must undertake to establish a procedure to review experience gained, as required by Annex VII, paragraph 4;
- The manufacturer shall undertake to preserve and make available the technical documentation of the Notified Body, as specified in Annex VII of Directive 93/42/EEC for a period of five years after the last date of manufacture of the product..

He therefore declares that the medical device is in accordance with the requirements of Directive 93/42/EEC and which will be marketed with the CE marking, as required by Article 17 of Directive 93/42/EEC.

First issue: 16/06/1998

Replace the certificate issued on : 08/01/2009

Current issue: 22/01/2010

Authorized signature

SEI EMG s.r.l.
Ing. Giorgio Didonè

This is a translation of the Italian text, which prevails in case of doubts

SEI EMG srl	**DECLARATION OF CONFORMITY**
DEVICE: CONNECTING CABLES *FOR EEG RECORDING*	**DC_011en**

Declaration of conformity of the device referred as "CONNECTING CABLES for EEG recording" manufactured by the company SEI EMG srl (already STRUMENTAZIONE ELETTRONICA INDUSTRIALE di Didonè Ing. Giorgio & C. snc)**, to the essential requirements of Annex I and Annex VII of Directive 93/42/EEC as amended by Directive 2007/47/EC.**

The writing firm **SEI EMG s.r.l.** established in Cittadella, Via S. Chiara 10,
VAT Number: IT-01597140282
manufacturer of the device family called **CUPLxx, CUMLxx CUCDxx CUEMxx**
declare under our sole responsibility that this medical device satisfies all applicable provisions in the Medical Device Directive 93/42/EEC amended by Directive 2007/47/EC.

For this purpose, the undersigned represent and warrant under his own responsibility:

- The device in question meets the essential requirements of Annex I to Directive 93/42/EEC updated by Directive 2007/47/EC;
- the device is considered in Class I;
- the device is sold in packs NOT STERILE;
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He therefore declares that the medical device is in accordance with the requirements of Directive 93/42/EEC and which will be marketed with the CE marking, as required by Article 17 of Directive 93/42/EEC.

First issue: 16/06/1998

Replace the certificate issued on : 08/01/2009

Current issue: 25/01/2010

Authorized signature

SEI EMG s.r.l.
Ing. Giorgio Didonè

This is a translation of the Italian text, which prevails in case of doubts