
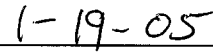
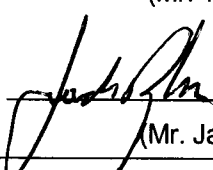
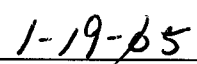


Declaration of Conformity

Manufacturer: WR Medical Electronics Co.	Address: 123 North Second Street Stillwater, MN 55082-5047 USA
Product Group: Nerve Stimulators	
Product Family: Silverstein™; Facial Nerve Monitor/Stimulator	
Device Name: Accessories for the Silverstein™; Facial Nerve Monitor/Stimulator	
Product Part Number(s): 3129, 3210, 3186EU	
Device Classification Per MDD: 3129=Class IIa, Rule 5; 3210=Class IIa, Rule 6, 3186EU=Class IIa, Rule 6	
Year of Manufacture: 2004	
European Representative: Medical Device Safety Service GmbH, Burckhardtstr 1, D-30163, Hannover, Germany	
Annex II Notified Body: SEMKO (Sweden) (0413)	
Technical File No: RA-4, Revision A, November 11, 2004	
Declaration: WR Medical Electronics Co. hereby declares that the medical device specified above, to which this declaration relates, is in conformance with the requirements of the Council Directive 93/42/EEC Medical Device Directive which apply to them and with Swedish National Legislation under LVFS 2003:11.	
Declaration Based On: Annex II (EC Declaration of Conformity; Full Quality Assurance System) of the Directive 93/42/EEC on Medical Devices	
Certificate No.: 41314493	Issued by: Intertek Semko AB
Declaration of Conformance Issued By: Mr. Jack Blais, President, and Mr. Tim Wall, Engineer, WR Medical Electronics Co. 123 North Second Street, Stillwater, MN 55082-5047 USA	
Prepared By: Quality Steering Team and Top Management	
 _____ (Mr. Tim Wall)	 _____ (Date)
 _____ (Mr. Jack Blais)	 _____ (Date)