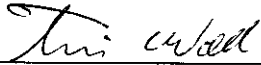
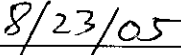
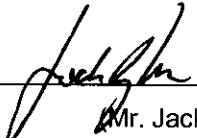
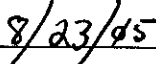
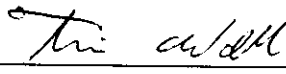
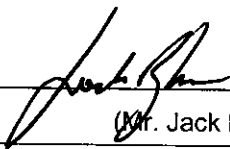


## Declaration of Conformity

<b>Manufacturer:</b> WR Medical Electronics Co.	<b>Address:</b> 123 North Second Street Stillwater, MN 55082-5047 USA
<b>Product Group:</b> Nerve Stimulators	
<b>Product Family:</b> Facial Nerve Stimulator	
<b>Device Name:</b> Accessories for the Hilger™; Facial Nerve Stimulator	
<b>Product Part Number(s):</b> 3302, 3119	
<b>Device Classification Per MDD:</b> Class I, per Rule 1	
<b>Year of Manufacture:</b> 2005	
<b>European Representative:</b> Medical Device Safety Service GmbH, Burckhardtstr 1, D-30163, Hannover, Germany	
<b>Technical File No:</b> RA-5, Revision A, November 9, 2004	
<b>Declaration:</b> 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.	
<b>Declaration Based On:</b> Annex VII (EC Declaration of Conformity) of the Directive 93/42/EEC on Medical Devices	
<b>Declaration of Conformance Issued By:</b> Mr. Jack Blais, President, and Mr. Tim Wall, Engineer, WR Medical Electronics Co. 123 North Second Street, Stillwater, MN 55082-5047 USA	
<b>Prepared By:</b> Quality Steering Team and Top Management	
 _____ (Mr. Tim Wall)	 _____ (Date)
 _____ (Mr. Jack Blais)	 _____ (Date)

# Declaration of Conformity

<b>Manufacturer:</b> WR Medical Electronics Co.	<b>Address:</b> 123 North Second Street Stillwater, MN 55082-5047 USA
<b>Product Group:</b> Nerve Stimulators	
<b>Product Family:</b> Facial Nerve Stimulator	
<b>Device Name:</b> Hilger™; Facial Nerve Stimulator	
<b>Product Part Number(s):</b> 3912	
<b>Device Classification Per MDD:</b> Class IIa, Non-Invasive, Transient; Rule 10	
<b>Year of Manufacture:</b> 2005	
<b>European Representative:</b> Medical Device Safety Service GmbH, Burckhardtstr 1, D-30163, Hannover, Germany	
<b>Annex II Notified Body:</b> SEMKO (Sweden) (0413)	
<b>Technical File No:</b> RA-5, Revision A, November 9, 2004	
<b>Declaration:</b> 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.	
<b>Declaration Based On:</b> Annex II (EC Declaration of Conformity; Full Quality Assurance System) of the Directive 93/42/EEC on Medical Devices	
<b>Certificate No.:</b> 41314493	<b>Issued by:</b> Intertek Semko AB
<b>Declaration of Conformance Issued By:</b> Mr. Jack Blais, President, and Mr. Tim Wall, Engineer, WR Medical Electronics Co. 123 North Second Street, Stillwater, MN 55082-5047 USA	
<b>Prepared By:</b> Quality Steering Team and Top Management	
 _____ (Mr. Tim Wall)	<u>8-23-05</u> _____ (Date)
 _____ (Mr. Jack Blais)	<u>8-23-05</u> _____ (Date)