

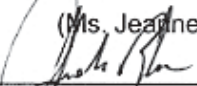
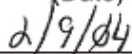


Declaration of Conformity

Manufacturer: WR Medical Electronics Co.	Address: 123 North Second Street Stillwater, MN 55082-5047 USA
Product Group: Computer Aided Sensory Evaluation - Neurological	
Product Family: CASE IV™	
Device Name: CASE IV™	
Product Part Number(s): 5000, 5569	
Device Classification Per MDD: Class I - Measuring, per Rule 1	
Year of Manufacture: 2003	
European Representative: Medical Device Safety Service GmbH, Burckhardtstr 1, D-30163, Hannover, Germany	
Annex V Notified Body: SEMKO (Sweden) (0413)	
Technical File No: RA-2, Revision A, 03 November 2003	
Declaration: WR Medical Electronics Co. hereby declares that the medical device specified above, to which this declaration relates, is in conformance with the essential requirements of Council Directive 93/42/EEC Medical Device Directive under Annex V (EC Declaration of Conformity; Production Quality Assurance), and with Swedish National Legislation under LVFS 2001:6.	
Declaration Based On: Annex V of the Directive 93/42/EEC on Medical Devices	
Certificate No.: 41314493	Issued by: Intertek Semko AB
Declaration of Conformance Issued By: Ms. Jeanne Anderson, Director of Regulatory and Clinical Affairs, Mr. Jack Blais, Executive Vice President, WR Medical Electronics Co. 123 North Second Street, Stillwater, MN 55082-5047 USA	
Prepared By: Quality Steering Team and Director of Regulatory and Clinical Affairs	
 _____ (Ms. Jeanne Anderson)	 _____ (Date)
 _____ (Mr. Jack Blais)	 _____ (Date)