



CERTIFICATE OF REGISTRATION

This certifies that:

PARLEX (EUROPE), LTD.

Taylor Road

Newport, Isle Of Wight, UNITED KINGDOM PO30 5LG

is registered and has listed the following medical device with the U.S. Food and Drug Administration pursuant to Title 21, 807 et seq. of the United States Code of Federal Regulations, such registration and listing having been verified as currently effective on, and as of the date hereof, by FDA Registrar Corp.

Establishment Registration:	3003875378
Device Classification Name:	GLUCOSE OXIDASE, GLUCOSE
Product Code:	CGA
Regulation Number:	862.1345
Official Correspondent and U.S. Agent:	FDA Registrar Corp. 144 Research Drive, Hampton, Virginia, 23666, USA Telephone: +1-757-224-0177 • Fax: +1-757-224-0179

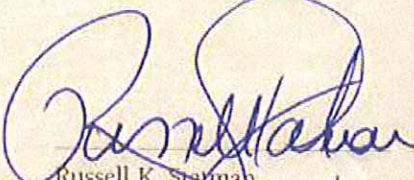
FDA Registrar Corp. will confirm that such registration remains effective upon request and presentation of this certificate until the end of the calendar year stated above, unless said registration is terminated after issuance of this certificate. FDA Registrar Corp. makes no other representations or warranties, nor does this certificate make any representations or warranties to any person or entity other than the named certificate holder, for whose sole benefit it is issued. This certificate does not denote endorsement or approval of the certificate-holder's device or establishment by the U.S. Food and Drug Administration. FDA Registrar Corp. assumes no liability to any person or entity in connection with the foregoing.

Pursuant to 21 CFR 807.39, "Registration of a device establishment or assignment of a registration number does not in any way denote approval of the establishment or its products. Any representation that creates an impression of official approval because of registration or possession of a registration number is misleading and constitutes misbranding."

FDA Registrar Corp. is a private registration agent not affiliated with the U.S. Food and Drug Administration.

FDA Registrar Corp. 

144 Research Drive, Hampton
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Russell K. Staman
Executive Director
FDA Registrar Corp.
Dated: *March 20, 2008*



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Establishment Registration: **3003875378**

Device Classification Name: **SYSTEM, TEST, BLOOD GLUCOSE, OVER THE COUNTER**

Product Code: **NBW**

Regulation Number: **862.1345**

Official Correspondent **FDA Registrar Corp.**

and U.S. Agent: **144 Research Drive, Hampton, Virginia, 23666, USA
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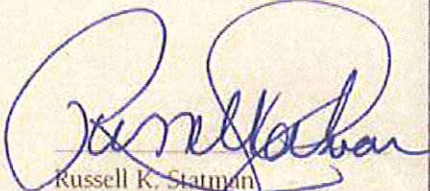
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