



2010

CERTIFICATE OF REGISTRATION

This certifies that:

**PPI-TIME ZERO INC.
262 Buffalo Avenue
Paterson, NJ 07503**

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 Registrar Corp.

Establishment Registration:	2244698
Device Classification Name:	GONIOMETER, AC-POWERED
Product Code:	KQX
Regulation Number:	888.1500

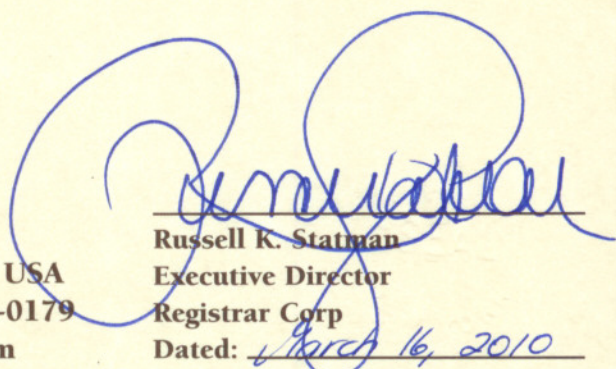
Registrar Corp. will confirm that such registration remains effective upon request and presentation of this certificate until the end of the calendar year stated above, unless said registration is terminated after issuance of this certificate. Registrar Corp. makes no other representations or warranties, 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. 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."

The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. Food and Drug Administration recognize a certificate of registration. Registrar Corp. is not affiliated with the U.S. Food and Drug Administration.

Registrar Corp 

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Russell K. Statman
Executive Director
Registrar Corp

Dated: March 16, 2010