



Food and Drug Administration
10903 New Hampshire Avenue
Building 66
Silver Spring, MD 20993

EPS BIO Technology Corp.
c/o Y. C. Lei
General Manager
No. 8 R & D Road III
Hsinchu Science Park
Hsinchu City, China (Taiwan) 30077

JUN 21 2011

Re: k102345
Trade Name: EasyMax TI Self Monitoring Blood Glucose System,
EasyMax TI Pro Self Monitoring Blood Glucose System
Regulation Number: 21 CFR 862.1345
Regulation Name: Glucose Test System.
Regulatory Class: Class II
Product Codes: CGA, NBW
Dated: May 20, 2011
Received: May 23, 2011

Dear Y. C. Lei:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

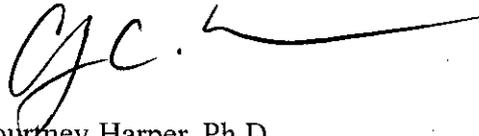
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in Title 21, Code of Federal Regulations (CFR), Parts 800 to 895. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Parts 801 and 809); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); and good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820).

If you desire specific advice for your device on our labeling regulation (21 CFR Parts 801 and 809), please contact the Office of *In Vitro* Diagnostic Device Evaluation and Safety at (301) 796-5450. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to <http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm> for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Small Manufacturers, International and Consumer Assistance at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address <http://www.fda.gov/cdrh/industry/support/index.html>.

Sincerely yours,

A handwritten signature in black ink, appearing to read 'CHC', with a long horizontal line extending to the right.

Courtney Harper, Ph.D.
Director
Division of Chemistry and Toxicology
Office of *In Vitro* Diagnostic Device
Evaluation and Safety
Center for Devices and Radiological Health

Enclosure

Indications for Use

510(k) Number (if known): K102345

Device Name: EasyMax T1 Pro Self Monitoring Blood Glucose System

Indications for Use:

The **EasyMax T1 Pro Self Monitoring Blood Glucose System** is intended for the quantitative measurement of glucose in fresh capillary whole blood samples drawn from the fingertips. Testing is done outside the body (In Vitro diagnostic use). It is indicated for multiple-patient use in a professional healthcare setting, as an aid to monitor the effectiveness of diabetes control. The system is only used with single-use lancing devices. The system is not to be used on neonates, nor for the diagnosis of, or screening for diabetes mellitus. Alternative site testing can be only used during steady-state blood glucose conditions.

The system consists of the **EasyMax T1 Pro Meter** and the **EasyMax T1 Pro Blood Glucose Test Strips**. The EasyMax T1 Pro Meter is only used with the EasyMax T1 Pro Blood Glucose Test Strips to quantitatively measure glucose in fresh capillary whole blood samples drawn from fingertips.

EasyMax T1 Glucose Control Solutions

For use with the EasyMax T1 Pro Self Monitoring Blood Glucose System as a quality control check to verify the accuracy of blood glucose test results.

Prescription Use V AND/OR Over-The-Counter Use V
(Part 21 CFR 801 Subpart D) (21 CFR 807 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of In Vitro Diagnostic Devices (OIVD)



Division Sign-Off

Office of In Vitro Diagnostic
Device Evaluation and Safety

510(k) K102345

1 of 2

Indications for Use

510(k) Number (if known): K102345

Device Name: EasyMax T1 Self Monitoring Blood Glucose system

Indications for Use:

The **EasyMax T1 Self Monitoring Blood Glucose System** is intended for the quantitative measurement of glucose in fresh capillary whole blood samples drawn from the fingertips or forearm. Testing is done outside the body (In Vitro diagnostic use). It is indicated for use at home (over the counter [OTC]) by a single patient with diabetes and should not be shared, as an aid to monitor the effectiveness of diabetes control. The system is not to be used on neonates, nor for the diagnosis of, or screening for diabetes mellitus. Alternative site testing can be only used during steady-state blood glucose conditions.

The system consists of the **EasyMax T1 Meter** and the **EasyMax T1 Blood Glucose Test Strips**. The EasyMax T1 Meter is only used with the EasyMax T1 test strips to quantitatively measure glucose in fresh capillary whole blood samples drawn from fingertips or forearm.

EasyMax T1 Glucose Control Solutions

For use with the EasyMax T1 Self Monitoring Blood Glucose System as a quality control check to verify the accuracy of blood glucose test results.

Prescription Use _____
(Part 21 CFR 801 Subpart D)

AND/OR

Over-The-Counter Use V
(21 CFR 807 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of In Vitro Diagnostic Devices (OIVD)



Division Sign-Off

Office of In Vitro Diagnostic
Device Evaluation and Safety

510(k) K102345

2 of 2