



December 24, 2013

IMMUNODIAGNOSTIC SYSTEMS LTD.
MICK FENTON
GLOBAL RA MANAGER
10 DIDCOT WAY
BOLDEN BUSINESS PARK
BOLDON, TYNE & WEAR NE35 9PD
UK

Re: K123763

Trade/Device Name: IDS iSYS Direct Renin Assay, IDS iSYS Direct Renin Control Set, and
IDS iSYS Direct Renin Calibration Verifiers

Regulation Number: 21 CFR 862.1085

Regulation Name: Angiotensin I and renin test system

Regulatory Class: II

Product Code: CIB

Dated: November 28, 2013

Received: December 2, 2013

Dear Mr. Fenton:

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. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. 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); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the

electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulations (21 CFR Parts 801 and 809), please contact 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/MedicalDevices/ResourcesforYou/Industry/default.htm>. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR 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/MedicalDevices/ResourcesforYou/Industry/default.htm>.

Sincerely yours,

Carol C. Benson -S for

Courtney H. Lias, Ph.D.
Director
Division of Chemistry and Toxicology Devices
Office of In Vitro Diagnostics
and Radiological Health
Center for Devices and Radiological Health

Enclosure

Indications for Use

510(k) Number (if known)

k123763

Device Name

IDS iSYS Direct Renin Assay, IDS-iSYS Direct Renin Control Set, and IDS-iSYS Direct Renin Calibration Verifiers

Indications for Use (Describe)

The IDS-iSYS Direct Renin assay is intended for the quantitative determination of Direct Renin in human EDTA plasma on the IDS iSYS Multi-Discipline Automated System. Renin measurements may aid in the diagnosis and treatment of certain types of hypertension.

The IDS-iSYS Direct Renin Control Set is used for quality control of the IDS-iSYS Direct Renin Assay on the IDS-iSYS Multi-Discipline Automated System.

The IDS-iSYS Direct Renin Calibration Verifier is a device intended for medical purposes for use in the quantitative verification of calibration of the IDS-iSYS Direct Renin Assay when performed on the IDS-iSYS Multi-Discipline Automated System.

Type of Use (Select one or both, as applicable)

Prescription Use (Part 21 CFR 801 Subpart D)

Over-The-Counter Use (21 CFR 801 Subpart C)

PLEASE DO NOT WRITE BELOW THIS LINE – CONTINUE ON A SEPARATE PAGE IF NEEDED.

FOR FDA USE ONLY

Concurrence of Center for Devices and Radiological Health (CDRH) (Signature)

Yung  Chan -S