



DEPARTMENT OF HEALTH & HUMAN SERVICES

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Food and Drug Administration  
2098 Gaither Road  
Rockville MD 20850

APR 11 2003

Joseph Lawlor, Ph.D.  
President  
Reference Diagnostics, Inc.  
19 Crosby Drive, Suite 30  
Bedford, MA 01730

Re: k030772  
Trade/Device Name: RDI Extended Range CRP Calibrator Set  
Regulation Number: 21 CFR 862.1150  
Regulation Name: Calibrator  
Regulatory Class: Class II  
Product Code: J1T  
Dated: March 10, 2003  
Received: March 11, 2003

Dear Dr. Lawlor:

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.

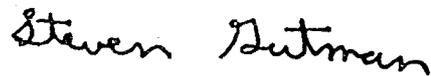
Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's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

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This letter will allow you to begin marketing your device as described in your Section 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific information about the application of labeling requirements to your device, or questions on the promotion and advertising of your device, please contact the Office of In Vitro Diagnostic Device Evaluation and Safety at (301) 594-3084. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). 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) 443-6597 or at its Internet address <http://www.fda.gov/cdrh/dsma/dsmamain.html>.

Sincerely yours,

A handwritten signature in black ink that reads "Steven Gutman". The signature is written in a cursive style.

Steven I. Gutman, M.D., M.B.A.  
Director  
Office of *In Vitro* Diagnostic Device  
Evaluation and Safety  
Center for Devices and  
Radiological Health

Enclosure



**REFERENCE DIAGNOSTICS, INC.**

Abbreviated 510(k): RDI Extended Range CRP Calibrator Set

19 Crosby Drive, Suite 30  
Bedford, Massachusetts 01730  
Tel 781 276-9000 Fax 781 276-9002

**Statement of Intended Use**

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510(k) Number (if known): K030772

Device Name: RDI Extended Range CRP Calibrator Set

**Indications For Use:**

For in vitro diagnostic use only.

The RDI Extended Range CRP Calibrator Set is intended for medical purposes for use with the RDI Extended Range hsCRP Kit, to establish points of reference that are used in the quantitative determination of C-reactive protein (CRP) in human serum or plasma, by immunoturbidimetric measurement.

**Summary and Explanation of the Test:**

Measurement of CRP helps in detecting and evaluating infection, tissue injury, inflammatory disorders, and associated diseases. The accuracy of the CRP measurement is insured by using calibrators that are traceable to the international reference preparation (CRM470).

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

Concurrence of CDRH, Office of Device Evaluation (ODE)

J. P. Reeves for J. Bantista  
(Division Sign-Off)  
Division of Clinical Laboratory Devices  
510(k) Number K030772

Prescription Use  
(Per 21 CFR 801.109)

OR

Over-The Counter Use