



DEPARTMENT OF HEALTH & HUMAN SERVICES

Food and Drug Administration
2098 Gaither Road
Rockville MD 20850

MAY 12 2003

Eric S. Hoy, Ph.D., AI(ASCP)
Chief Scientific Officer
Immuno Concepts Incorporated
9779 "D" Business Park Drive
Sacramento, California 95827

Re: k030775
Trade/Device Name: Auto I.D.® Jo-1 Positive Control
Auto I.D.® Scl-70 Positive Control
Auto I.D.® PCNA Positive Control
Regulation Number: 21 CFR § 866.5100
Regulation Name: ANA Immunological Test System
Regulatory Class: II
Product Code: LKJ
Dated: March 10, 2003
Received: March 11, 2003

Dear Dr. Hoy :

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.

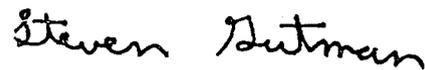
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in Title 21, Code of Federal Regulations (CFR), Parts 800 to 895. In addition, FDA may publish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

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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

510(k) number (if known): K030775

Device Name: AUTO I.D.® Positive Control Serum

Trade or Proprietary Name:

- AUTO-I.D.® Jo-1 Positive Control Serum (Catalog number 6004)
- AUTO-I.D.® Scl-70 Positive Control Serum (Catalog number 6005)
- AUTO-I.D.® PCNA Positive Control Serum (Catalog number 6006)

Indications for use:

These are qualitative calibrators/controls to confirm the presence of antibodies to autoantigens (Jo-1, Scl-70, or PCNA) in human serum. Antibodies to Jo-1 are found in patients with polymyositis and dermatomyositis, antibodies to Scl-70 are found in patients with scleroderma, and antibodies to PCNA are found in patients with systemic lupus erythematosus. These optional controls are intended for use with the Immuno Concepts AUTO-ID® Autoantibody Test System.

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

Concurrence of CDRH, Office of Device Evaluation (ODE)

J. Reeves for J. Bautista
 (Division Sign-Off)
 Division of Clinical Laboratory Devices
 510(k) Number K030775

Prescription Use
 (Per 21 CFR 801.109)

OR Over-The-Counter Use