



APR - 5 2004

Food and Drug Administration  
2098 Gaither Road  
Rockville MD 20850

Mr. Mark J. Kopnitsky  
Vice President of Research Development  
Zeus Scientific, Inc.  
200 Evans Way  
Branchburg, NJ 08876

Re: k040111  
Trade/Device Name: AtheNA Multi-Lite™ MPO/ PR3 IgG Test System  
Regulation Number: 21 CFR 866.5660  
Regulation Name: Multiple autoantibodies immunological test system  
Regulatory Class: Class II  
Product Code: MOB  
Dated: January 13, 2004  
Received: February 13, 2004

Dear Mr.Kopnitsky:

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.

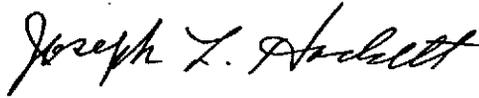
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in Title 21, Code of Federal Regulations (CFR), Parts 800 to 895. In addition, FDA may publish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

Page 2

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,



Joseph L. Hackett, Ph.D.  
Acting Director  
Division of Immunology and Hematology Devices  
Office of In Vitro Diagnostic Device Evaluation and Safety  
Center for Devices and Radiological Health

Enclosure

Indications for Use Form

(Appendix E)

510(k) Number (if known): K04011

Device Name: AtheNA Multi-Lyte™ MPO/PR3 Test System

Indications for Use:

The Zeus Scientific, Inc. AtheNA Multi-Lyte™ MPO/PR3 Test System is for the qualitative and semi-quantitative detection of IgG autoantibodies to human myeloperoxidase (MPO) and/or human proteinase 3 (PR3) in human serum. The results of this serological test together with other clinical findings may aid in the diagnosis of systemic vasculitides (SV). This test is for in vitro diagnostic use.

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

\_\_\_\_\_

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use

OR

Over-The-Counter Use

(Per 21 CFR 801,109)

(Optional Format 1-

2-96)

*Maria Chan*

\_\_\_\_\_  
Division Sign-Off

Office of In Vitro Diagnostic Device  
Evaluation and Safety

510(k) K04011