K030213

510(k) Summary

APR 11 2003

Introduction

According to the requirements of 21 CFR 807.92, the following information provides sufficient detail to understand the basis for a determination of substantial equivalence.

1) Submitter name, address, contact

Roche Diagnostics Corporation
9115 Hague Rd.
Indianapolis, IN 46250
(317) 521-7637

Contact Person: Kerwin Kaufman

Date Prepared: January 17, 2003

2) Device name

Proprietary name: ONLINE DAT II Cannabinoids II

Common name: Cannabinoids test system

Classification name: Enzyme immunoassay, cannabinoids

3) Predicate device

We claim substantial equivalence to the currently marketed Abuscreen OnLine Cannabinoids assay (K983701).

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4) Device Description

The ONLINE DAT II Cannabinoids II (THC II) assay is an in vitro diagnostic test for the qualitative and semi-quantitative detection of cannabinoids in human urine on automated clinical chemistry analyzers at cutoff concentrations of 20 ng/ml, 50 ng/ml and 100 ng/ml. Semi-quantitative test results may be obtained that permit laboratories to assess assay performance as part of a quality control program.

Principal of procedure

The ONLINE DAT II Cannabinoids II assay is based on the kinetic interaction of microparticles in a solution (KIMS) as measured by changes in light transmission. In the absence of sample drug, soluble drug-polymer conjugates bind to antibody-bound microparticles, causing the formation of particle aggregates.

When a urine sample containing the drug in question is present, this drug competes with the conjugate-bound drug derivative for microparticle-bound antibody. Antibody bound to sample drug is no longer available to promote particle aggregation, and subsequent particle lattice formation is inhibited.

As the aggregation reaction proceeds in the absence of sample drug, the absorbance increases. Conversely, the presence of sample drug diminishes the increasing absorbance in proportion to the concentration of drug in the sample. Sample drug content is determined relative to the value obtained for a known cutoff concentration of drug.

Negative Sample

drug-polymer conjugate + antibody-bound microparticle = particle aggregates
(↑ absorbance)

Positive Sample

sample drug + antibody-bound microparticle = particle aggregation inhibited
drug-polymer conjugate + antibody bound microparticle = particle aggregates

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510(k) Summary, Continued

5.) Intended Use

The ONLINE DAT II Cannabinoids II (THC II) assay is an in vitro diagnostic test for the qualitative and semi-quantitative detection of cannabinoids in human urine on automated clinical chemistry analyzers at cutoff concentrations of 20 ng/ml, 50 ng/ml and 100 ng/ml. Semi-quantitative test results may be obtained that permit laboratories to assess assay performance as part of a quality control program.

6.) Comparison to the Predicate Device

The Roche ONLINE DAT II Cannabinoids II assay is substantially equivalent to other products in commercial distribution intended for similar use. Most notably, it is substantially equivalent to the currently marketed Roche Abuscreen OnLine Cannabinoids assay (K983701).

The Roche ONLINE DAT II Cannabinoids II assay utilizes a modified KIMS technology relative to the currently marketed Abuscreen OnLine Cannabinoids assay. Differences between this application and the cleared assay include:

- use of a new cannabinoids monoclonal antibody (mouse) attached to microparticles in solution,
- a soluble drug-polymer conjugate, and
- use of new calibrators and unassayed controls.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Food and Drug Administration
2098 Gaither Road
Rockville MD 20850

APR 11 2003

Mr. Kerwin Kaufman
Regulatory Affairs Consultant
Roche Diagnostics Corporation
9115 Hague Road
P.O. Box 50457
Indianapolis, IN 46250-0457

Re: k030213
Trade/Device Name: Roche Diagnostics ONLINE DAT II Cannabinoids II
Regulation Number: 21 CFR 862.3870
Regulation Name: Cannabinoid test system
Regulatory Class: Class II
Product Code: LDJ
Dated: January 17, 2003
Received: January 21, 2003

Dear Mr. Kaufman:

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 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,

Steven Gutman

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

Indications for Use Statement

510(k) Number (if known): K030213

Device Name: Roche Diagnostics ONLINE DAT II Cannabinoids II

Indications for Use: Cannabinoids II (THC II) is an in vitro diagnostic test for the qualitative and semiquantitative detection of cannabinoids in human urine on automated clinical chemistry analyzers at cutoff concentrations of 20 ng/ml, 50 ng/ml and 100 ng/ml. Semi-quantitative test results may be obtained that permit laboratories to assess assay performance as part of a quality control program.

(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)

Dan Conde

(Division Sign-Off)
Division of Clinical Laboratory Devices
510(k) Number K030213