

K120747

510(k) Summary of Safety and Effectiveness

OCT 22 2012

510(k) Submitter: Streck
7002 South 109th Street
Omaha, NE 68128

Official Correspondent: Deborah Kipp, Quality Assurance Coordinator
(402)537- 5215

Date Prepared: Revised-September 21, 2012

Name of Device:
Trade Name: XN CAL™ PF
Common Name: Assayed Hematology Calibrator
Classification Name: Calibrator for Cell Indices (KRX-864.8150)

Predicate Device: X-CAL™ (K083200) Manufactured by Streck

Description:

XN CAL™ PF is an in-vitro diagnostic product that contains the following: stabilized red blood cell component(s), and stabilized platelet component(s) in a preservative medium. The product is packaged in polypropylene plastic vials with screw caps containing 3 ml. The vials will be packaged in (5) well or (1) well vacuum formed clamshell container with the Instructions for Use / assay sheet. The product must be stored at 2 - 8° C.

Intended Use:

XN CAL PF is used for calibration and calibration verification of Sysmex XN series (XN-10, XN-20) analyzers. The assayed parameter is:

PLT-F (10³/μL)

Comparison to Predicate Device:

| | X-CAL™-K083200 (Predicate Product) | XN CAL™ PF |
|-------------------------------|---|--|
| Intended Use Statement | X-CAL™ is used to calibrate and verify calibration of Sysmex hematology analyzers. Refer to product assay sheet. | XN CAL™ PF is used for calibration and calibration verification of Sysmex XN series (XN-10, XN-20) analyzers. The assayed parameter is: PLT-F (10 ³ /μL) |
| Open Vial Stability | 24 hours | 4 hours |
| Closed Vial Stability | 35 days | Same |
| Reagents | X-CAL™ is composed of a mixture of stabilized human and animal blood cells. The cells are suspended in a solution containing biological salts and anti-microbial preservatives. | XN CAL™ PF contains the following: stabilized red blood cell component(s), and stabilized platelet component(s) in a preservative medium. |
| Storage Conditions | 2 - 8°C | Same |

Discussion of Tests and Test Results:

The following studies were conducted to establish performance of XN CAL™ PF. The tests conducted were Open-Vial Stability, Closed-Vial Stability, and Precision Performance. All testing showed that XN CAL™ PF is consistently reproducible, substantially equivalent to the predicate product and stable for the shelf life claimed.

Conclusions Drawn From Tests:

Study results show XN CAL™ PF to be consistently reproducible, substantially equivalent to the predicate products, and stable for the entire product dating. XN CAL™ PF is a safe and effective product, which fulfills its intended use when used as instructed in the Instructions for Use.



10903 New Hampshire Avenue
Silver Spring, MD 20993

Streck
c/o Ms. Deborah S. Kipp
Quality Assurance Coordinator
7002 South 109th Street
Omaha, NE 68128

OCT 22 2012

Re: k120747
Trade/Device Name: XN-CAL™ PF
Regulation Number: 21 CFR 864.8150
Regulation Name: Calibrator for cell indices
Regulatory Class: Class II
Product Code: KRX
Dated: September 28, 2012
Received: October 01, 2012

Dear Ms. Kipp:

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 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 Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set

Page 2 – Ms. Deborah Kipp

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 regulation (21 CFR Parts 801 and 809), please contact the Office of *In Vitro* Diagnostics and Radiological Health at (301) 796-5450. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR 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/cdrh/industry/support/index.html>.

Sincerely yours,



Maria M. Chan, Ph.D.
Director
Division of Immunology and Hematology Devices
Office of *In Vitro* Diagnostics and Radiological Health
Center for Devices and Radiological Health

Enclosure

Indication for Use Form

510(k) Number (if known): K120747

Device Name: XN CAL™ PF

Indications For Use:

XN CAL PF is used for calibration and calibration verification of Sysmex XN series (XN-10, XN-20) analyzers. The assayed parameter is:

PLT-F ($10^3/\mu\text{L}$)

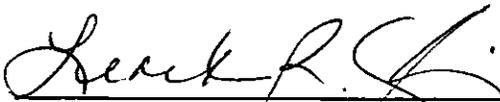
Prescription Use X
(Per 21 CFR 801 Subpart D)

AND/OR

Over-The-Counter Use
(21 CFR 807 Subpart C)

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

Concurrence of CDRH, Office of In Vitro Diagnostic Devices (OIVD)



Division Sign-Off
Office of In Vitro Diagnostic Device
Evaluation and Safety

510(k) K120747

Page 1 of 1