

III. 510(k) SUMMARY

Submitter's Name and Address: Church & Dwight Co., Inc.
469 North Harrison Street
Princeton, NJ 08543

Contact Person: Stephen C. Kolakowsky
Director, Regulatory Affairs

Date Prepared: October 2007

Proprietary Name: TROJAN® THINTENSITY™ brand
with ULTRASMOOTH™ Lubricant, and
with WARM SENSATIONS™ Lubricant

Common Name: Male Latex Condom

Classification Name: Condom

Predicate Device: TROJAN® THINTENSITY™ brand Male Latex Condom
with Silicone Lubricant
Church & Dwight Co., Inc.; K071272

Description of the Device: The condoms are made of a natural rubber latex sheath, which completely covers the penis with a closely fitted membrane. The condom is smooth surfaced, straight-walled with a contoured-bulbous section near the closed-end, which terminates with a reservoir tip. The condom design is within the ASTM standard specifications D-3492, e.g., minimum length 160 mm, maximum width 54 mm, and minimum thickness of 30 µM. Each condom is lubricated with either of two varieties of lubricant, ULTRASMOOTH™ lubricant or WARM SENSATIONS™ lubricant.

Intended Use of the Device: This latex condom product has the same intended use as the predicate. It is used for contraception and for prophylactic purposes (to help prevent pregnancy and the transmission of sexually transmitted diseases).

Technological Characteristics: The 510(k)-subject condom products have the same technological characteristics as the predicate condom products identified above. The design is in conformance with ASTM Latex Condom Standard D3492 and the condoms are made of natural rubber latex.

NOV 09 2007



Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

NOV 09 2007

Mr. Stephen C. Kolakowsky
Director, Regulatory Affairs
Church & Dwight Co., Inc.
469 North Harrison Street
Law Department
Building 100
PRINCETON NJ 08543

Re: K073016

Trade Name: TROJAN® THINTENSITY™ brand male latex condom lubricated
with Ultrasmooth™ lubricant and with Warm Sensations™
lubricant

Regulation Number: 21 CFR §884.5300

Regulation Name: condom

Regulatory Class: II

Product Code: HIS

Dated: October 23, 2007

Received: October 25, 2007

Dear Mr. Kolakowsky:

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 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); good manufacturing practice requirements as set 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.

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 advice for your device on our labeling regulation (21 CFR Part 801), please contact the Center for Devices and Radiological Health's (CDRH's) Office of Compliance at one of the following numbers, based on the regulation number at the top of this letter.

21 CFR 876.xxxx	(Gastroenterology/Renal/Urology)	240-276-0115
21 CFR 884.xxxx	(Obstetrics/Gynecology)	240-276-0115
21 CFR 892.xxxx	(Radiology)	240-276-0120
Other		240-276-0100

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR Part 807.97). For questions regarding postmarket surveillance, please contact CDRH's Office of Surveillance and Biometric's (OSB's) Division of Postmarket Surveillance at 240-276-3474. For questions regarding the reporting of device adverse events (Medical Device Reporting (MDR)), please contact the Division of Surveillance Systems at 240-276-3464. 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 (240) 276-3150 or at its Internet address <http://www.fda.gov/cdrh/industry/support/index.html>.

Sincerely yours,



Nancy C. Brogdon
Director, Division of Reproductive,
Abdominal, and Radiological Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure

VIII. INDICATIONS FOR USE STATEMENT

510(k) Number: K073016

Device Name: TROJAN® THINTENSITY™ brand male latex condom lubricated with Ultrasmooth™ lubricant and with Warm Sensations™ lubricant

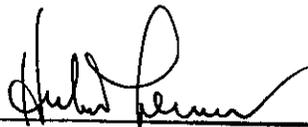
Indications For Use: The TROJAN® THINTENSITY™ brand male latex condom is used for contraception and for prophylactic purposes (to help prevent pregnancy and the transmission of sexually transmitted diseases).

Prescription Use _____
(Per 21 CFR §801.109)

Over-The-Counter Use

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

Concurrence of CDRH, Office of Device Evaluation (ODE)



(Division Sign-Off)

Division of Reproductive, Abdominal, and
Radiological Devices

510(k) Number K073016