K032124

JAN 16 2004

SSL Americas, Inc.

Office of Regulatory Affairs
3585 Engineering Drive, Suite 200
Norcross, GA 30092-9214
Tel: 770-582-2222
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Durex Play™, Premarket approval [510(k)] Summary

Section VIII.1 Submitter Information
SSL Americas
3585 Engineering Dr.
Suite 200
Norcross, GA 30092-9214
Phone: 770 – 582 – 2222
Fax: 770 – 582 – 2233
Contact person: Kathleen Harris, Regulatory Affairs Manager, SSL Americas
Date of Summary: July 4, 2003

Section VIII.2 General Device Information
Device Trade Name: Durex Play™
Device Common Name: Personal lubricant
Classification: Patient lubricant

Section VIII.3 Predicate Devices
K-Y Jelly Personal Lubricant (K955648)
AstroGlide (K935299)

Section VIII.4 Device Description
Durex Play™ is a clear colorless personal lubricant composed of purified water, Hydroxyethyl cellulose (Natrosol 250), propylene glycol, benzoic acid, sodium hydroxide.

Section VIII.5 Intended Use
Indications: Durex Play™ is intended as a moisturizer for vaginal dryness and personal lubrication of the vaginal entry to enhance condom use and to facilitate ease and comfort during intimate sexual activity.

Section VIII.6 Substantial Equivalence
Durex Play™ is substantially equivalent in intended use to K-Y Jelly personal lubricant and AstroGlide and similar in composition. All products are marketed as personal lubricants, safe to use with condoms, water soluble and sold Over-the-Counter.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

JAN 16 2004

Ms. Kathleen Harris
Regulatory Manager
SSL Americas, Inc.
Office of Regulatory Affairs
3585 Engineering Drive, Suite 200
NORCROSS GA 30092-9214

Re: K032124
Trade/Device Name: Durex Play Personal Lubricant
Regulation Number: 21 CFR 884.5300
Regulation Name: Condom
Regulatory Class: II
Product Code: 85 HIS
Dated: October 16, 2003
Received: October 21, 2003

Dear Ms. Harris:

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 Office of Compliance at one of the following numbers, based on the regulation number at the top of the letter:

8xx.1xxx	(301) 594-4591
876.2xxx, 3xxx, 4xxx, 5xxx	(301) 594-4616
884.2xxx, 3xxx, 4xxx, 5xxx, 6xxx	(301) 594-4616
892.2xxx, 3xxx, 4xxx, 5xxx	(301) 594-4654
Other	(301) 594-4692

Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. 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 may be obtained 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,

Nancy C. Brogdon

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

Enclosure

Indications for Use

510(k) Number (if known): K032124

Device Name: Durex Play Personal Lubricant

Indications For Use: Durex Play™ is intended as a moisturizer for vaginal dryness and personal lubrication of the vaginal entry to enhance condom use and to facilitate ease and comfort during intimate sexual activity.

Prescription Use _____ ~~AND~~/OR Over-The-Counter Use X
(Part 21 CFR 801 Subpart D) (21 CFR 807 Subpart C)

(~~PL~~EASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE IF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Nancy C. Brogdon

(Division Sign-Off)
Division of Reproductive, Abdominal,
and Radiological Devices
510(k) Number K032124