December 15, 2015

Oculus Innovative Sciences Incorporated  
Mr. Brian Martin  
Director of Regulatory Affairs and Quality Control  
1129 North Medowell Boulevard  
Petaluma, California 94954

Re: K152945  
Trade/Device Name: Alevicyn SG Antipruritic Gel  
Regulatory Class: Unclassified  
Product Code: FRO  
Dated: October 5, 2015  
Received: October 6, 2015

Dear Mr. Martin:

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. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

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 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 Part 801), please contact the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm. 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 Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm.

Sincerely yours,

Joshua C. Nipper -S

For Binita S. Ashar, M.D., M.B.A., F.A.C.S.
Director
Division of Surgical Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure
Indications for Use

Rx Indication: Under the supervision of a health care professional, Alevicyn SG Gel is indicated to manage and relieve the burning, itching, erythema, scaling, and pain experienced with various types of dermatoses, including atopic dermatitis, radiation dermatitis, and seborrhea and seborrheic dermatitis. Alevicyn SG Gel may be also used to relieve the pain of first and second degree burns. Alevicyn SG Gel helps to relieve dry waxy skin by maintaining a moist wound & skin environment, which is beneficial to the healing process.

Type of Use (Select one or both, as applicable)

☒ Prescription Use (Part 21 CFR 801 Subpart D) ☐ Over-The-Counter Use (21 CFR 801 Subpart C)

CONTINUE ON A SEPARATE PAGE IF NEEDED.

This section applies only to requirements of the Paperwork Reduction Act of 1995.

*DO NOT SEND YOUR COMPLETED FORM TO THE PRA STAFF EMAIL ADDRESS BELOW.*

The burden time for this collection of information is estimated to average 79 hours per response, including the time to review instructions, search existing data sources, gather and maintain the data needed and complete and review the collection of information. Send comments regarding this burden estimate or any other aspect of this information collection, including suggestions for reducing this burden, to:

Department of Health and Human Services
Food and Drug Administration
Office of Chief Information Officer
Paperwork Reduction Act (PRA) Staff
PRASStaff@fda.hhs.gov

“An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB number.”
1 510(k) SUMMARY
The following is a summary of 510(k) safety and effectiveness information in accordance with 21 CFR 807.92.

I. SUBMITTER
Oculus Innovative Sciences, Inc.
1129 North McDowell Blvd.
Petaluma, CA 94954
Phone: (707) 283-0550
Fax: (707) 283-0551
Contact Person: Brian W. Martin, Director of Regulatory Affairs and Quality Control
Date Prepared: October 5, 2015

II. DEVICE
Name of Device: Alevicyn SG Antipruritic Gel
Common or Usual Name: Antipruritic Gel
Classification Name: Dressing, Wound, Drug
Regulatory Class: Unclassified, Pre-amendment status
Product Code: FRO

III. PREDICATE DEVICE
Alevicyn SG Gel (K143590) manufactured by Oculus Innovative Sciences, Inc.,
TL Triseb Cream (K121134), manufactured by Trigen Laboratories, Inc.,
Dermiseb Cream (K123724) manufactured by IGI Laboratories, Inc., and
Loutrex Topical Cream (K120730) manufactured by Acella Pharmaceuticals, LLC

IV. DEVICE DESCRIPTION
The Alevicyn SG Gel is an opaque gel, slightly chlorinated odor, low viscosity spray gel. The product has a pH range of 5.5-7.0 and a viscosity target of 500-5000 cP and will be supplied in polyethylene terephthalate (PET) round-bottles with polypropylene (PP) screw-top closure and a finger pump sprayer.

V. INDICATIONS FOR USE
Rx Indication: Under the supervision of a health care professional, Alevicyn SG Gel is indicated to manage and relieve the burning, itching, erythema, scaling, and pain experienced with various types of dermatoses, including atopic dermatitis, radiation dermatitis, and seborrhea and seborrheic dermatitis. Alevicyn SG Gel may be also used to relieve the pain of first and second degree burns. Alevicyn SG Gel helps to relieve dry
waxy skin by maintaining a moist wound & skin environment, which is beneficial to the healing process.
VI. COMPARISON OF TECHNOLOGICAL CHARACTERISTICS WITH THE PREDICATE DEVICE

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<td>Indications for Use</td>
<td>Rx Indication: Alevicyn SG Gel is indicated to manage and relieve the burning, itching, erythema, scaling, and pain experienced with various types of dermatoses, including atopic dermatitis, radiation dermatitis, and seborrhea and seborrheic dermatitis. Alevicyn SG Gel may be also used to relieve the pain of first and second degree burns. Alevicyn SG Gel helps to relieve dry waxy skin by maintaining a moist wound &amp; skin environment, which is beneficial to the healing process.</td>
<td>Rx Indication: Alevicyn SG Antipruritic Gel is indicated to manage and relieve the burning, itching and pain experienced with various types of dermatoses, including radiation dermatitis and atopic dermatitis. Alevicyn SG Antipruritic Gel may be also used to relieve the pain of first and second degree burns. Alevicyn SG Antipruritic Gel helps to relieve dry waxy skin by maintaining a moist wound &amp; skin environment, which is beneficial to the healing process.</td>
<td>Rx Indication: Under the supervision of a healthcare professional, TL Triseb Cream is indicated to manage and relieve the signs and symptoms of seborrhea and seborrheic dermatitis such as itching, erythema, scaling and pain. TL Triseb Cream helps relieve dry waxy skin by maintaining a moist wound &amp; skin environment, which is beneficial to the healing process.</td>
<td>Rx Indication: Under the supervision of a healthcare professional, Dermiseb Cream is indicated to manage and relieve the signs and symptoms of seborrhea and seborrheic dermatitis such as itching, erythema, scaling and pain. Dermiseb Cream also aids to relieve dry, waxy skin by maintaining a moist wound and skin environment. A moist wound and skin environment is beneficial to the healing process.</td>
<td>Rx Indication: Loutrex Topical Cream is indicated to manage and relieve the signs and symptoms of seborrhea and seborrheic dermatitis such as itching, erythema, scaling and pain. Loutrex Topical Cream helps relieve dry, waxy skin by maintaining a moist wound and skin environment, which is beneficial to the healing process.</td>
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<td>Mechanism of Action</td>
<td>Skin barrier emollient and moisturizer</td>
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<td>Delivery System</td>
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<td>Shelf Life</td>
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<td>Sponsor/Manufacturer</td>
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VII. PERFORMANCE DATA

The following performance data were provided in support of the substantial equivalence determination.

Biocompatibility Testing

Bench Testing
The following tests were reviewed to support the performance of Alevicyn SG Gel: package integrity, visual inspection, pH, Free Available Chlorine (FAC). The Alevicyn SG Gel meets specification and performance characteristics and is substantially equivalent to the predicate device.

VIII. CONCLUSION

Alevicyn SG Gel is substantially equivalent in intended use, technological characteristics, safety and effectiveness to Alevicyn SG Gel (K143590), TL Triseb Cream (K121134), Dermiseb Cream (K123724), and Loutrex Topical Cream (K120730). Therefore, the Oculus Alevicyn SG Gel is substantially equivalent to the predicate devices.