

K973807

DEC 23 1997

*510(k) Premarket Notification
DuoDERM® Hydroactive® Paste*

ITEM 8: 510(k) SUMMARY OF SAFETY AND EFFECTIVENESS

The purpose of this 510 (k) Premarket Notification is to request clearance for claims specific to the moist wound healing environment created by DuoDERM Hydroactive Paste.

The concept of DuoDERM Hydroactive Paste is not significantly different from other commercially available absorbent dressings intended for use in the management of exudating wounds. DuoDERM Hydroactive Paste is indicated for use on exudating dermal ulcers, including full thickness wounds such as leg ulcers, pressure ulcers and diabetic ulcers. It is used in association with other DuoDERM® dressings. DuoDERM Hydroactive Paste is substantially equivalent to ConvaTec's SignaDRESS Hydrocolloid Dressing. The two products are equivalent in intended use and dressing characteristics.

DuoDERM Hydroactive Paste is contraindicated for use in patients with active vasculitis, such as, periarteritis nodosa, systemic lupus erythematosus, and cryoglobulinemia and on ulcers resulting from infection such as tuberculosis, syphilis or deep fungal infections.

Data/information supporting the safety of DuoDERM Hydroactive Paste was presented in Premarket Notification K881302 . All testing was performed in accordance with Good Laboratory Practice Regulations.



Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

DEC 23 1997

Ms. Adrienna McNally
Senior Manager, Regulatory Affairs
ConvaTec
PO Box 5254
Princeton, New Jersey 08543-5254

Re: K973807
DuoDERM® Hydroactive™ Paste
Regulatory Class: Unclassified
Product Code: MGP
Dated: September 29, 1997
Received: October 6, 1997

Dear Ms. McNally:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to 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 (the Act). You may, therefore, market your device subject to the general controls provisions of the Federal Food, Drug, and Cosmetic Act (Act) and the following limitations:

1. This device may not be labeled for use on third degree burns.
2. This device may not be labeled as having any accelerating effect on the rate of wound healing or epithelization.
3. This device may not be labeled as a long-term, permanent, or no-change dressing, or as an artificial (synthetic) skin.
4. This device may not be labeled as a treatment or a cure for any type of wound.

The labeling claims listed above would be considered a major modification in the intended use of the device and would require a premarket notification submission (21 CFR 807.81). The general controls provisions of the Act include requirements for annual

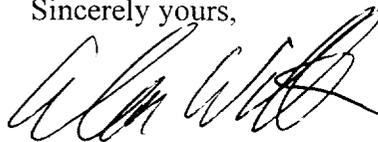
registration, listing of devices, good manufacturing practices, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval) it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations (CFR), Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Good Manufacturing Practices (GMP) for Medical Devices: General GMP regulation (21 CFR Part 820) and that, through periodic GMP inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531 through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 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 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4595. 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" (21 CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or 301-443-6597 or at its internet address <http://www.fda.gov/cdrh/dsmamain.html>.

Sincerely yours,



Celia M. Witten, Ph.D., M.D.

Director

Division of General and

Restorative Devices

Office of Device Evaluation

Center for Devices and

Radiological Health

Enclosure

K973807

510(k) Premarket Notification
DuoDERM® Hydroactive® Paste

1J: INDICATIONS FOR USE STATEMENT

510(k) Number (if known): Not Known

Device Name: DuoDERM Hydroactive Paste

Indications for Use:

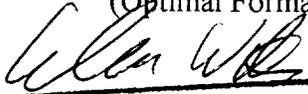
DuoDERM Hydroactive Paste is designed to be used with a secondary dressing such as DuoDERM Dressings for the local management of exudating dermal ulcers, including full thickness wounds, such as leg ulcers, pressure ulcers and diabetic ulcers.

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

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use X
(Per 21 CFR 801.109)

OR Over the Counter Use _____
(Optimal Format 1-2-96)



(Division Sign-Off)
Division of General Restorative Devices
510(k) Number K973807

*ConvaTec - A Division of E.R. Squibb & Sons, Inc.
DuoDERM® Hydroactive® Paste - A trademark of ConvaTec
SignaDRESS™ Hydrocolloid Dressing - A trademark of ConvaTec*