

MAY 03 2002

K020760

510(k) Summary of Safety and Effectiveness
(As Required by 21 C.F.R. §807.92)

Submitted by: Craig Bruns
Vice President
Danville Materials, Inc.
2021 Omega Dr.
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Date of summary February 14, 2002

Device name Starflow HV

Common name Tooth Shade Resin Material

Classification names Regulation Number Product Code
21 CFR 872.3690 EBF

Device Description Starflow HV is a light-cured, hybrid dental restorative which incorporates BIS/GMA resin along with Barium aluminosilicate glass filler (64% by wt) to yield a flowable dental restorative.

Predicate Device The device is substantially equivalent to other legally marketed devices in the United States including Revolution by Sybron Dental Specialties (K013647), StarFlow by Danville Materials Inc (K990108), and Tetric Flow (K 993783) by Ivoclar Vivadent. The primary differences are very minor changes to the material composition.

Intended Use Starflow HV is intended for use for the restoration of Class III, Class IV, and Class V cavities, small Class I cavities not in occlusion, repair of enamel defects, repair of porcelain restorations, minor occusal build-ups in non-stress bearing areas, pit and fissure sealant, cement for seramic/composite veneers, incisal abrasions, and core build-ups.



Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

MAY 03 2002

Mr. Craig R. Bruns
Vice President & General Manager
Danville Materials, Incorporated
2021 Omega Drive
San Ramon, California 94583

Re: K020760

Trade/Device Name: Starflow HV
Regulation Number: 872.3690
Regulation Name: Tooth Shade Resin Material
Regulatory Class: II
Product Codes: EBF
Dated: February 20, 2002
Received: March 07, 2002

Dear Mr. Bruns:

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.

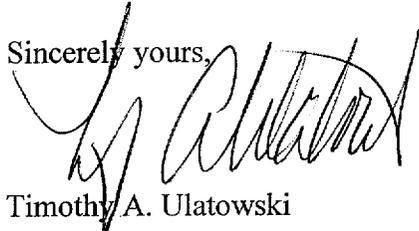
If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish 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 and additionally 21 CFR Part 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4613. 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). 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,


Timothy A. Ulatowski
Director
Division of Dental, Infection Control,
and General Hospital Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

510(k) Number: K020760

Device Name: Starflow HV

Indications for use:

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

Concurrence of CDRH, Office of Device Evaluation (ODE)

Susan P. [Signature]

(Division Sign-Off)
Division of Dental, Infection Control,
and General Hospital Devices
510(k) Number K020760

Prescription Use
(Per 21 CFR 801.109)

OR

Over-The-Counter Use

(Optional Format 1-2-96)