Section 3 510(k) Summary

This 510(k) Summary of 510(k) safety and effectiveness information is being submitted in accordance with requirements of SMDA 1990 and Title 21, CFR Section 807.92.

The assigned 510(k) Number: ____________

1. Date of Submission: 09/10/2013

2. Sponsor Identification

Wenzhou Wuzhou Import & Export Co., Ltd.
Room 1703 Fortune Center Chezhan Avenue
Wenzhou, Zhejiang Province, 325000, China

Establishment Registration Number: 9681901

Contact Person: Bingyi Xiang
Position: General Manager
Tel: +86-577-88868068
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3. Submission Correspondent

Ms. Diana Hong & Mr. Lee Fu
Mid-Link Consulting Co., Ltd
P.O. Box 120-119
Shanghai, 200120, China
Tel: +86-21-22815850
Fax: 240-238-7587
Email: info@mid-link.net
4. Proposed Device Identification

Proposed Device Name: Insulin Pen Needle
Proposed Device Common Name: Insulin Pen Needle

Regulatory Information:
Classification Name: Needle, Hypodermic, Single Lumen;
Classification: 2;
Product Code: FMI;
Regulation Number: 21 CFR 880.5570;
Review Panel: General Hospital;

Intended Use Statement:
The Insulin Pen Needle is intended for use with pen injector devices for the subcutaneous injection of insulin.

5. Predicate Device Identification

510(k) Number: K120690
Product Name: insulin pen needle
Manufacturer: Tiger Medical Products Ltd.

6. Device Description

The proposed device, Insulin Pen Needle, is a single-use device, which is designed for use with a pen injector for the subcutaneous injection of insulin. It consists of needle tube, hub, tube sheath, hub sheath and sealed paper. The hub can be connected screwed onto the insulin pen.

The Insulin Pen Needle is offered in various gauge sized and length.

They are provided sterilized with Sterility Assurance Level (SAL) of 10^6.

7. Non-Clinical Test Conclusion

Non clinical tests were conducted to verify that the proposed device met all design specifications as was Substantially Equivalent (SE) to the predicate device. The test results demonstrated that the proposed device complies with the following standards:

ISO 9626 First edition 1991-09-01, AMENDMENT 1 2001-06-01 Stainless steel needle tubing for the
manufacture of medical devices.
ISO 10993-7:2008(R) 2012, Biological evaluation of medical devices - Part 7: Ethylene oxide sterilization residuals

8. Substantially Equivalent (SE) Conclusion

The following table compares the DEVICE to the predicate device with respect to intended use, technological characteristics and principles of operation, etc.

<table>
<thead>
<tr>
<th>Item</th>
<th>Proposed Device</th>
<th>Predicate Device</th>
</tr>
</thead>
<tbody>
<tr>
<td>Product Code</td>
<td>FMI</td>
<td>Same</td>
</tr>
<tr>
<td>Regulation Number</td>
<td>21 CFR 880.5570</td>
<td>Same</td>
</tr>
<tr>
<td>Class</td>
<td>Class II</td>
<td>Same</td>
</tr>
<tr>
<td>Intended Use</td>
<td>The Insulin Pen Needle is intended for use with pen injector devices for the subcutaneous injection of insulin.</td>
<td>Same</td>
</tr>
<tr>
<td>Configuration</td>
<td>Needle Tube, Hub, Tube Sheath, Hub Sheath and Sealed Paper</td>
<td>Similar</td>
</tr>
<tr>
<td>Operation mode</td>
<td>Manual</td>
<td>Same</td>
</tr>
<tr>
<td>Needle Gauge</td>
<td>29G/30G/31G/32G</td>
<td>Same</td>
</tr>
<tr>
<td>Performance specification</td>
<td>Comply with ISO 7864, ISO 9626, and ISO 11608-2</td>
<td>Same</td>
</tr>
<tr>
<td>Material</td>
<td>Tube: 304 Stainless Steel</td>
<td>Same</td>
</tr>
<tr>
<td></td>
<td>Hub: Polypropylene</td>
<td>Same</td>
</tr>
<tr>
<td>Sterilization</td>
<td>SAL:10^6</td>
<td>Same</td>
</tr>
<tr>
<td></td>
<td>EO Sterilized</td>
<td>Same</td>
</tr>
<tr>
<td>Shelf Life</td>
<td>5 years</td>
<td>Same</td>
</tr>
<tr>
<td>Single Use</td>
<td>Yes</td>
<td>Same</td>
</tr>
<tr>
<td>Labeling and Labels</td>
<td>Meet FDA requirements</td>
<td>Same</td>
</tr>
</tbody>
</table>

The proposed device, Insulin Pen Needle, is determined to be Substantially Equivalent (SE) to the predicate device, insulin pen needle (K120690), in respect of safety and effectiveness.
February 25, 2014

Wenzhou Wuzhou Import & Export Company, Limited  
C/O Ms. Diana Hong  
General Manager  
Mid-Link Consulting Company, Limited  
P.O. Box 120-119  
Shanghai, 200120  
CHINA

Re: K133058  
Trade/Device Name: Insulin Pen Needle  
Regulation Number: 21 CFR 880.5570  
Regulation Name: Hypodermic Single Lumen Needle  
Regulatory Class: II  
Product Code: FMI  
Dated: September 18, 2013  
Received: September 27, 2013

Dear Ms. Hong:

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 Small Manufacturers, International and Consumer Assistance 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 Small Manufacturers, International and Consumer Assistance 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,

Kwame O. Ulmer, M.D.
Acting Director
Division of Anesthesiology, General Hospital, Respiratory, Infection Control and Dental Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure
Additional Information for K133058  Exhibit#2 Updated Indications for Use Form

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

Indications for Use

510(k) Number (If known)
K133058

Device Name
Insulin Pen Needle

Indications for Use (Describe)
The Insulin Pen Needle is intended for use with pen injector devices for the subcutaneous injection of insulin.

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)

PLEASE DO NOT WRITE BELOW THIS LINE – CONTINUE ON A SEPARATE PAGE IF NEEDED.

Digitally signed by
Richard C. Chapman
Date: 2014.02.25
12:54:03 -05'00'