

July 13, 2018

Dongguan Prestige Sporting Goods Co., Ltd. % Cassie Lee
Manager
Guangzhou GLOMED Biological Technology Co., Ltd.
Suite 306, Kecheng Mansion, No.121 Science Road
Guangzhou Science Park
Guangzhou, Guangdong, 510663 CN

Re: K172440

Trade/Device Name: Solax Electric Scooter, Models: S302121, S302131, S302141, S302151

Regulation Number: 21 CFR 890.3800

Regulation Name: Motorized three-wheeled vehicle

Regulatory Class: Class II

Product Code: INI Dated: April 29, 2018 Received: May 3, 2018

Dear Cassie Lee:

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.

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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.

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 comprehensive regulatory information about medical devices and radiation-emitting products, including information about labeling regulations, please see Device Advice (https://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/) and CDRH Learn (http://www.fda.gov/Training/CDRHLearn). Additionally, you may contact the Division of Industry and Consumer Education (DICE) to ask a question about a specific regulatory topic. See the DICE website (http://www.fda.gov/DICE) for more information or contact DICE by email (DICE@fda.hhs.gov) or phone (1-800-638-2041 or 301-796-7100).

Sincerely,

Vivek J. Pinto -S

for Carlos L. Peña, PhD, MS
Director
Division of Neurological
and Physical Medicine Devices
Office of Device Evaluation
Center for Devices and Radiological Health

Enclosure

DEPARTMENT OF HEALTH AND HUMAN SERVICES Food and Drug Administration

Indications for Use

510(k) Number (if known)

Form Approved: OMB No. 0910-0120 Expiration Date: January 31, 2017 See PRA Statement below.

K172440			
Device Name Solax Electric Scooter, Models: S302121, S302131, S302141, S302151			
ndications for Use (Describe) t is a motor driven, indoor and outdoor transportation vehicle with the intended use to provide mobility to a disabled or elderly person limited to a seated position.			
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)		

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

CONTINUE ON A SEPARATE PAGE IF NEEDED.

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 PRAStaff@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."

510(k) Summary

This summary of 510(K) safety and effectiveness information is being submitted in accordance with the requirement of 21 CFR 807.92.

1. Submitter's Information

- ♦ 510(k) Owner's Name: Dongguan Prestige Sporting Goods Co., Ltd.
- Establishment Registration Number: 3008841035
- ♦ Address: 3rd industrial, Qiaotou Area, Houjie Town, Dongguan City, Guangdong province, China
- ♦ Tel: 13763128800
- ♦ Fax:86-769-85922505
- ♦ Contact Person: Zhang Zhao (General Manager)
- ♦ E-mail: leon@wisefame.com

2. Application Correspondent:

- ♦ Contact Person: Ms. Cassie Lee
- Guangzhou GLOMED Biological Technology Co., Ltd.
- Address: Suite 306, Kecheng Mansion, No.121 Science Road, Guangzhou Science Park, Guangzhou 510663, China
- ◆ Tel:+86-20-61099984
- ♦ Email: regulatory@glomed-info.com

3. Subject Device Information

Trade Name:	Solax Electric Scooter Model: S302121, S302131, S302141, S302151	
Common Name:	Scooter	
Classification name:	Vehicle, Motorized 3-Wheeled	
Review Panel:	Physical Medicine	
Product Code:	INI	
Regulation Class:	2	
Regulation Number:	890.3800	

4. Predicate Device Information

Sponsor	Dongguan Prestige Sporting Goods Co., Ltd.
Device Name	MOBIE scooter

510(k) Number	K122749
Product Code	INI
Regulation Number	890.3800
Regulation Class	2

5. DeviceDescription

The Solax Electric Scooter (Models: S302121, S302131, S302141, S302151) is an indoor/outdoor electric scooter that is intended to be used by individuals that are able to walk, but suffer from mobility limitations. It has a base with aluminum alloy frame, two front wheels, two rear wheels, two anti-tip wheels, a seat, an adjustable steering column, a control panel, an electric motor, an electromagnetic brake, a remote for folding/unfolding the device and a Lithium battery with an off-board charger. The movement of the scooter is controlled by the rider who operates the direction control lever, speed control switch and handle on the control panel. The device is installed with an electromagnetic brake that will engage automatically when the scooter is not in use and the brake cannot be used manually.

6. Intended Use / Indications for Use

It is a motor driven, indoor and outdoor transportation vehicle with the intended use to provide mobility to a disabled or elderly person limited to a seated position.

7. Tests Summary

The following performance and safety tests were conducted with Solax Electric Scooter:

- ISO 7176-1: 2014, Wheelchairs Part 1: Determination of static stability
- ISO 7176-2:2001, Wheelchairs Part 2: Determination of dynamic stability of Powered Wheelchairs
- ISO 7176-3: 2012, Wheelchairs Part 3: Determination of effectiveness of brakes
- ISO 7176-4, Third edition 2008-10-01, Wheelchairs Part 4: Energy consumption of electric wheelchairs and scooters for determination of theoretical distance range
- ISO 7176-5, Second edition 2008-06-01, Wheelchairs Part 5: Determination of overall dimensions, mass and manoeuvring space
- ISO 7176-6: 2001, Wheelchairs Part 6: Determination of maximum speed, acceleration and deceleration of Powered Wheelchairs
- ISO 7176-7, First Edition 1998-05-15, Wheelchairs Part 7: Measurement of seating and wheel dimensions
- ISO 7176-8:2014, Wheelchairs Part 8: Requirements and test methods for static, impact and fatigue strengths
- ISO 7176-9:2009, Wheelchairs Part 9: Climatic tests for Powered Wheelchairs
- ISO 7176-10:2008, Wheelchairs Part 10: Determination of obstacle-climbing ability of electrically powered wheelchairs

- ISO 7176-11, Second edition 2012-12-01, Wheelchairs Part 11: Test dummies
- ISO 7176-13, First edition 1989-08-01, Wheelchairs Part 13: Determination of coefficient of friction of test surfaces
- ISO 7176-14:2008, Wheelchairs Part 14: Power and control systems for electrically powered wheelchairs and scooters - Requirements and test methods
- ISO 7176-15:1996, Wheelchairs Part 15: Requirements for information disclosure, documentation and labeling
- ISO 7176-16, Second edition 2012-12-01, Wheelchairs Part 16: Resistance to ignition of postural support devices
- ISO 7176-21 Second edition 2009-04-01 Wheelchairs Part 21: Requirements and test methods for electromagnetic compatibility of electrically powered wheelchairs and scooters, and battery chargers
- IEC 62304: 2006 (First Edition), Medical device software, Software life- cycle processes.
- IEC 60601-1-6 Medical electrical equipment—Part1-6: General requirements for safety—Collateral Standard: Usability Edition 3.0, 2010
- IEC 62366 Medical devices Application of usability engineering to medical devices Edition 1.0,
 2007

The materials and manufacturing used for the Solax Electric Scooter are identical to those of the predicate device, which were demonstrated to conform with the following biocompatibility standards:

- ISO 10993-5:2009, Biological evaluation of medical devices Part 5: Tests for In Vitro cytotoxicity
- ISO 10993-10: 2009, Biological evaluation of medical devices Part 10: Tests for irritation and delayed-type hypersensitivity

8. Comparison to predicate device and conclusion

The technological characteristics, features, specifications, materials, mode of operation, and intended use of Solax Electric Scooter is substantially equivalent to the predicate devices quoted above.

The differences between the subject device and predicate devices do not raise new issues of safety or effectiveness.

Elements of Comparison	Predicate Device	Subject Device	Remark
Proprietary name	MOBIE scooter	Solax Electric Scooter	
Model	Mobie	S302121, S302131, S302141, S302151	
510K Number	K122749	K172440	
Common or Usual name	Electrical scooter	Electrical scooter	

Intended use	It is a motor driven, indoor and outdoor transportation vehicle with the intended use to provide mobility to a disabled or elderly person limited to a seated position.	It is a motor driven, indoor and outdoor transportation vehicle with the intended use to provide mobility to a disabled or elderly person limited to a seated position.	SE
Size (unfold)	980*450*985mm	930*450*865mm	SE Note
Tires	6 inches for front wheel (solid wheel) 7 inches for rear wheel (solid wheel)	6 inches for front wheel (solid wheel) 7 inches for rear wheel (solid wheel)	SE Note
Speed	6 km/h (3.7mph)	6 km/h (3.7mph)	SE
Safe Gradient / Maximum Gradient	0-13º	0-12°	SE Note
Range	15 km (9.32mile)	15 km (9.32mile)	SE
Turning circle	1.4 m	1.55 m	SE note
Base weight(not including battery)	25 kg	24 kg	SE Note
Battery weight	1.84 kg	1.84 kg	SE
Brake	Electromagnetic	Electromagnetic	SE
Drive system	PG 45A / Rear wheel drive	PG 45A / Rear wheel drive	SE
Maximum capacity	136 kg Approx.	125 kg Approx.	SE Note
Ground clearance	36 mm	36 mm	SE
Obstacle Climbing Ability	38 mm	38 mm	SE
Battery	Lithium battery 24V/10AH	Lithium battery 24V/10AH	SE
Motor	24V 120W	24V 120W	SE
Battery charger	DC24V/2A	DC24V/2A	SE
Biocompatibility	All user directly contacting materials are compliance with ISO10993-5 and ISO10993-10 requirements.	All user directly contacting materials are compliance with ISO10993-5 and ISO10993-10 requirements.	SE
EMC	ISO 7176 - 21	ISO 7176 - 21	SE
Performance	ISO 7176 series	ISO 7176 series	SE

Comparison in Detail(s):

Note:

The design and technological characteristics of the Solax Electric Scooter is basically similar to the predicate device chosen. There are minor differences between the devices including overall dimensions, Safe Gradient, Turning circle, Maximum capacity, Ground clearance, Battery and motor. There is no deleterious effect on safety and effectiveness due to the differences, and these minor differences do not influence the intended use function and use of the device. Moreover, the non-clinical tests and the predicate comparisons demonstrate that these differences in their technological characteristics do not

raise any questions as to the safety and effectiveness. Therefore, the Solax Electric Scooter is substantially equivalent to the MOBIE (K122749).

9. Date of the summary prepared: July 11, 2018