

This authorizes the application of the Certification Mark(s) shown below to the models described in the Product(s) Covered section when made in accordance with the conditions set forth in the Certification Agreement and Listing Report. This authorization also applies to multiple listee model(s) identified on the correlation page of the Listing Report.

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|--|-------------------------------------|-----------------------|---|
| Applicant: | GlobTek, Inc. 186 Veterans Drive | Manufacturer: | GlobTek (Suzhou) Co., Ltd. Building 4 No. 76 JinLing East Road |
| Address: | NORTHVALE NJ 07647 USA | Address: | Suzhou Industrial Park SUZHOU Jiangsu 215021 |
| Country: | USA | Country: | China |
| Contact: | Hans Moritz | Contact: | Demon Zhou |
| Phone: | (201)784-1000 Ext.253 | Phone: | 86 512 6279 0301 Ext.189 |
| FAX: | (201)784-0111 | FAX: | 86 512 6279 0355 |
| Email: | Moritzh@globtek.com | Email: | demon.zhou@globtek.cn |
| Party Authorized To Apply Mark: | Same as Manufacturer | | |
| Report Issuing Office: | INTERTEK TESTING SERVICES SH LTD | | |
| Control Number: | <u>4007497</u> | Authorized by: |  for Dean Davidson, Certification Manager |



This document supersedes all previous Authorizations to Mark for the noted Report Number.

This Authorization to Mark is for the exclusive use of Intertek's Client and is provided pursuant to the Certification agreement between Intertek and its Client. Intertek's responsibility and liability are limited to the terms and conditions of the agreement. Intertek assumes no liability to any party, other than to the Client in accordance with the agreement, for any loss, expense or damage occasioned by the use of this Authorization to Mark. Only the Client is authorized to permit copying or distribution of this Authorization to Mark and then only in its entirety. Use of Intertek's Certification mark is restricted to the conditions laid out in the agreement and in this Authorization to Mark. Any further use of the Intertek name for the sale or advertisement of the tested material, product or service must first be approved in writing by Intertek. Initial Factory Assessments and Follow up Services are for the purpose of assuring appropriate usage of the Certification mark in accordance with the agreement, they are not for the purposes of production quality control and do not relieve the Client of their obligations in this respect.

Intertek Testing Services NA Inc.
545 East Algonquin Road, Arlington Heights, IL 60005
Telephone 800-345-3851 or 847-439-5667 Fax 312-283-1672

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| Standard(s): | Medical Electrical Equipment - Part 1: General Requirements For Basic Safety And Essential Performance [AAMI ES60601-1:2005 +A1] |
| | Medical Electrical Equipment - Part 1: General Requirements For Basic Safety And Essential Performance [CSA C22.2#60601-1:2014 Ed.3] |
| | Medical Electrical Equipment - Part 1-11: General Requirements For Basic Safety And Essential Performance - Collateral Standard: Requirements For Medical Electrical Equipment And Medical Electrical Systems Used In The Home Healthcare Environment [IEC 60601-1-11:2015 Ed.2] |
| Product: | Medical Power Supply |
| Brand Name: | GlobTek |

GT followed by M, - or H; followed by 91099-; followed by 01 to 60; followed by 09, 15, 24 or 48; may be followed by -; may be followed by 0.01 to 23.9; followed by -T2, -T2A, -T3, -T3A, -F, -FW, -P2 or -P3; may be followed by six characters.

GT followed by M, - or H; followed by 96600-; followed by 01 to 65; followed by 05 to 54; followed by -T2, -T2A, -T3, -T3A, -T2L, -T2AL, T3L, -T3AL, -R2, -R3A, -P2 or -P3; may be followed by six characters.

Models:

GT followed by M, - or H; followed by 96600-; followed by 01 to 65; followed by 5.0 to 54.0; followed by -T2, -T2A, -T3, -T3A, -T2L, -T2AL, T3L, -T3AL, -R2, -R3A, -P2 or -P3; may be followed by six characters.

GT followed by M, - or H; followed by 96600-; followed by 01 to 70; followed by 56; followed by -T2, -T2A, -T3, -T3A; followed by -AP, -PP, -SP; may be followed by six characters.