

Listing#: E115461  
Report #: 130411  
Original Certification Date: January 31, 2024  
Revised Certification Date:

This Certification is issued to:  
GlobTek, Inc.  
186 Veterans Dr. Northvale, NJ 07647 USA

Stating that the product(s):  
Medical Power Supply  
Series GTM46360-\*\*\*\*, GTM96183-\*PD\*-USB1C\*, GTM96181-\*PD\*\*\*

Product Rating(s):  
GTM46360-\*\*\*\*:

- Input: 100-240V~, 50-60Hz, Max. 0.75A,
- Output: 3.0-5.0Vdc, Max. 6.0A, Max. 30W

GTM96183-\*PD\*-USB1C\*, GTM96181-\*PD\*\*\*:

- Input: 100-240V~, 50-60Hz, 1.2A,
- Output: 5.0- 20.0Vdc, Max. 3.0A, Max. 36W

GTM96183-\*PD\*-PPS-USB1C\*, GTM96181-\*PD\*-PPS\*\*:

- Input: 100-240V~, 50-60Hz, 1.2A,
- Output: 5.0- 21.0Vdc, Max. 3.0A, Max. 36W

Achieved Certification to the following standard(s):

ANSI/ AAMI ES60601-1:2005, ES60601-1:2005/AMD1 1:2012, ES60601-1:2005/AMD 2:2021

CAN/CSA-C22.2 No. 60601-1:14 + A2:22 (R2022)

Medical electrical equipment— Part 1: General requirements for basic safety and essential performance

IEC 60601-1-6 Edition 3.2 2020-07

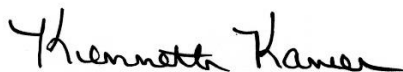
CAN/CSA-C22.2 NO. 60601-1-6:11 + A1:15 + A2:21 (R2021)

Medical electrical equipment - Part 1-6: General requirements for basic safety and essential performance - Collateral standard: Usability

ANSI/ AAMI HA60601-1-11:2015 [Including AMD1: 2021]

CSA C22.2 NO. 60601-1-11:15 (R2020) + A1:21

Medical Electrical Equipment -- Part 1-11: General requirements for basic safety and essential performance -- Collateral Standard: Requirements for medical electrical equipment and medical electrical equipment and medical electrical systems used in the home healthcare environment



Ken Kamer  
Eurofins Electrical and Electronic Testing North America, Inc.



All changes proposed in the previously identified product that affects the above information must be submitted to Eurofins for evaluation prior to implementation to assure continued NRTL Certification status. The covered product(s) shall be subject to follow-up inspections to ensure that the Certified product(s) are identical to the product sample evaluated by Eurofins E&E NA and that all responsibilities are being fulfilled as specified in the Applicants' Responsibility section of the Certification Report. The Applicant named above has been authorized Eurofins E&E NA to represent the product(s) listed in this record as "MET Certified" and to mark this/these product(s) according to the terms and conditions of the Eurofins E&E NA Applicant Contract, Listing Reports, and the applicable agreements. Only the product(s) bearing the MET Mark and under a follow-up service are considered to be included in this Certification program. This certification has been granted under a System 3 program as defined in ISO/IEC 17067.



*Eurofins E&E North America, Inc. is accredited by OSHA and the Standards Council of Canada.*

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