

Shenzhen SEM.Test Technology Co., Ltd.

1/F, Building A, Hongwei Industrial Park, Liuxian 2nd Road, Bao'an District, Shenzhen, P.R.C. (518101)

CERTIFICATE OF CONFORMITY

Certificate No.: SEM18118330

The following product has been tested by Shenzhen SEM. Test Technology Co., Ltd. with the listing standards and found in conformity with the **EU Directive 2014/30/EU & 2014/53/EU Directive.** It is possible to use CE marking to demonstrate the conformity with this **EMC & RED Directive.**

Report No. : STR18118126E
Applicant : GlobTek, Inc.

Address : 186 Veterans Dr. Northvale, NJ 07647 USA

Manufacturer : 1.GlobTek, Inc.

2.GlobTek (Suzhou) Co., Ltd

Address : 1.186 Veterans Dr. Northvale, NJ 07647 USA

2.Building 4, No. 76, Jin Ling East Rd., Suzhou Industrial

Park, Suzhou, Jiang Su 215021, China

Description of Product: ITE / Medical Power supply

Model No. : See Model List

Trade Name : GlobTek, Inc.

Test Standards : EN 55032:2015/AC:2016-07

EN 61000-3-2:2014 EN 61000-3-3:2013 EN 55035:2017

EN 60601-1-2:2015

The referred test report(s) show that the product complies with the essential requirements in the above listed standards. The applicant is authorized to use this certificate in connection with the EU declaration of conformity according to Annex 1 of the Directive.



Jandy So Supervisor

Date of Issue: Nov. 20 2018

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This certificate of conformity is based on a single evaluation of the submitted sample(s) of the above mentioned product. It does not imply an assessment of the whole production and other relevant Directives have to be observed.

Tel.: +86-755-33663308 Fax.: +86-755-33663309 E-mail: <u>sem@semtest.com.cn</u> Website: <u>www.semtest.com.cn</u>



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ANNEX 1 TO ADDING MODEL

Certificate No. : SEM18118330 Model List

GT*46161-**-USB
The 1st "*" can be "M" or "-"or "H" for market identification and not related to safety.
The 2nd "*" denote the rated output wattage designation, which can be "01" to "16", with interval
The 3rd "*" denote the standard rated output voltage designation, which can be "5.0" to "5.5" or
"05.5" with interval of 0,1.

This certificate of conformity is based on a single evaluation of the submitted sample(s) of the above mentioned product. It does not imply an assessment of the whole production and other relevant Directives have to be observed.

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