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America

CERTIFICATE

No. U8 18 02 01888 001

Holder of Certificate: **Amico Clinical Solutions, Corp**
55 East Wilmot St
Richmond Hill ON L4B 1A3
CANADA

Production Facility(ies): 95914

Certification Mark:

C US

Product: **General Medical Devices**
Medical Pendant and Console

Model(s):
LP-L-xxxx-N-E
LP-L-xxxx-R-E
LP-L-PLP-xxxx-R-E
LP-H-NNNN
LP-H-NNNN-R-E
LP-V-xxxx-N-E-90
LP-V-TM-xxxx-N-E-90
Where xxxx represents the length of the arm, each x can be 1,2,8 or N

Parameters:

Rated voltage:	120V
Rated frequency:	50/60Hz
Rated input current:	20A
Protection Class:	I
Duty Cycle:	1 min. ON, 9 min. OFF

Tested according to:
CAN/CSA-C22.2 No. 60601-1:2014
excluding Biocompatibility (clause 11.7)
and EMC (clause 17).
ANSI/AAMI ES60601-1:2005/A1:2012-08
excluding Biocompatibility (clause 11.7)
and EMC (clause 17)

The product was voluntarily tested according to the relevant safety requirements noted above. It can be marked with the certification mark above. The mark must not be altered in any way. This product certification system operated by TÜV SÜD America Inc. most closely resembles system 3 as defined in ISO/IEC 17067. Certification is based on the TÜV SÜD "Testing and Certification Regulations". TÜV SÜD America Inc. is an OSHA recognized NRTL and a Standards Council of Canada accredited certification body.

Test report no.: 7169002865-000

Date, 2018-03-01

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