VERIFICATION CERTIFICATE

NOT TRANSFERABLE

This Verification Certificate is hereby issued to the named GRANTEE and is VALID ONLY for the equipment identified hereon for use under the rules and regulations listed below:

GRANTEE:	**Communications & Power Industries Inc.**
Address:	45 River Drive Georgetown, Ontario Canada, L7G 2J4
Contact Person:	Ms. Katrina de Asis Phone #: 905-702-2222 x414 Fax #: (905) 877-5327 Email Address: Katrina.deAsis@cpii.com
Equipment Type:	Medical Electrical/Electronic Equipment
Product Name:	**Indico IQ® high frequency X-ray generators**
Models:	**VZW2558FJ8xx-yy** **VZW2558FD8xx-yy** **VZW2558FC8xx-yy**
The above product was tested by UltraTech Engineering Labs Inc. and found to comply with:	• **CISPR 11:2009+A1:2010 / EN 55011:2009+A1:2010, Class A, Group 1** • **EN 60601-1-2: 2015 /IEC 60601-1-2: 4th Edition**- Medical electrical equipment. General requirements for basic safety and essential performance. Collateral Standard. Electromagnetic disturbances. Requirements and tests

Note(s): *See attached report, UltraTech's File No.:18CPI262_EN60601-1-R1, dated* January 16, 2019 *for details and conditions of Verification Compliance.*

Approved by: Tri M. Luu BASc.
V.P. – Engineering

UltraTech

3000 Bristol Circle, Oakville, Ontario, Canada, L6H 6G4
Tel.: (905) 829-1570 Fax.: (905) 829-8050
Website: www.ultratech-labs.com, Email: vic@ultratech-labs.com, Email: tri@ultratech-labs.com

ANSI ACCREDITED #0685 ISO/IEC 17065 Product Certification Body

APEC TEL CA0001 | 1309 | CA 0001/2049 | AT-1945 | SL2-IN-E-1119R | Korea KCC-RRA CA2049

(Recommended Sample DoC for Manufacturer based on the attached Test report)

EU DECLARATION OF CONFORMITY

APPPLICATION OF COUNCIL DIRECTIVE(S): **93/42/EEC - The Medical Device Directive**

APPLICANT:	Communications & Power Industries Inc.
Equipment Type:	Medical Electrical/Electronic Equipment
Product Name:	**Indico IQ® high frequency X-ray generators**
Models:	**VZW2558FJ8xx-yy** **VZW2558FD8xx-yy** **VZW2558FC8xx-yy**

I, the undersigned, hereby, declare that the above device has been tested and found to comply with the following standard(s):

STANDARD(S) TO WHICH CONFORMITY IS DECLARED:	• **CISPR 11:2009+A1:2010 / EN 55011:2009+A1:2010, Class A, Group 1** • **EN 60601-1-2: 2015 /IEC 60601-1-2: 4th Edition** - Medical electrical equipment. General requirements for basic safety and essential performance. Collateral Standard. Electromagnetic disturbances. Requirements and tests
TEST LABORATORIES:	UltraTech Group of Labs Inc. 3000 Bristol Circle Oakville, Ontario Canada, L6H 6G4

Applicant:		**Legal Representative in Europe:**	
Signature:	_____	*Signature:*	_____
Full Name:	_____	*Full Name:*	_____
Title:	_____	*Title:*	_____
Full Address:	_____	*Full Address:*	_____
Phone No.:	_____	*Phone No.:*	_____
Email Address:	_____	*Email Address:*	_____