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Project and Product Certification Representative Authorization Letter

Reason for Amendment (current / obsolete)	Revision History		Approved Date
	From	To	
Initial Release (obsolete)	1.0	1.0	Nov-14-2006
Revised wording (obsolete)	1.0	2.0	Sept-25-2007
Updated company template (obsolete)	2.0	3.0	Jan-31-2012
Updated letter information (obsolete)	3.0	4.0	May-23-2014
Added FCC ID field (current)	4.0	5.0	Sept 16 2014

Medtronic

3/27/2018

To: SIEMIC, Inc.
775 Montague Expressway,
Milpitas, CA 95035
USA

Dear Sir/Madam,

Re: Project and Product Certification Representative Authorization Letter

We, Medtronic, Inc. hereby authorize SIEMIC, Inc. to act as Certification Body for certifying for the following project(s):

InterStim[®] Model 3023 Neurostimulator
FCC ID: LF53023

We affirm that between SIEMIC, Inc. and Medtronic, Inc., any difference in understanding, including test plan, measurement methods, applicable standards and relevant procedures and processes have been resolved prior to commencement of testing activities.

Sincerely,



Guillaume Girard
EMC and RF Regulatory Compliance
Technical Fellow
Medtronic, Inc.
Tel 763-526-0652
g.guillaume@medtronic.com