Medtronic

EC DECLARATION OF CONFORMITY

DoC No:	2020-004-01
Manufacturer:	Medtronic MiniMed 18000 Devonshire Street Northridge CA, 91325 USA
EC Representative:	Medtronic BV Earl Bakkenstraat 10 6422 PJ Heerlen The Netherlands
Conformity Assessment Procedure:	MDD 93/42/EEC Medical Devices Directive Annex II, except Section 4
EC Certificate number and Expiry Date: (Annex II.3)	Certificate No. 8857 (Expiry: May 26th, 2024)
Notified Body:	GMED 1, rue Gaston Boissier 75015 PARIS FRANCE
Description of Device Concerned:	**MiniMed 740G Insulin Pump** Pump (MMT-1811, MMT-1812) Pump Kit (MMT-1861, MMT-1862)
Classification:	Class IIb (Rule 11 of Annex IX)
GMDN Code:	35983

Statement:

I, the undersigned, hereby declare that the Medical Device(s) specified above and provided with CE marking, conform with the Essential Requirements of the EC Directive 93/42/EEC Medical Device Directive. This declaration is supported by the EC Quality System Certificate(s) according to the provisions of relevant Annex(es) of above Directive. This declaration applies to all devices specified above distributed from the signature date forward.

Tamar Jaghasbanian
Sr. Regulatory Affairs Manager
Medtronic Minimed
18000 Devonshire Street
Northridge, CA 91325 USA

12 February 2020
Date

Valid from signature date