W66 N220 Commerce Ct.
Cedarburg, WI 53012 USA

Date: 1 July 2018

To:

To Whom It May Concern:

GE Medical Systems Information Technologies is requesting a Class II Permissive Change to convert from a Product Filing to a Limited Modular Approval.

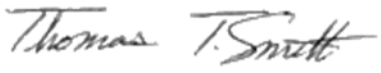
The original certification is under FCC ID: OU52102462001 and IC: 4048B-2102462001.

Product has not been modified or altered from the Original filing and certification. The intent of the permissive change is to allow for more implementation into future GE products without the need of obtaining a new FCC ID each time.

Product is not intended for handheld or body-worn applications. Radio module will be mounted inside a Medical Monitoring device with the sole purpose of providing secure access to the equipment via the RFID authentication protocol. The radio module is powered from the end-products power system and does not operate from a battery on its own.

GE does not intend to market the RFID module to other OEM's or end-customers. RFID module will be controlled in GE products. GE is aware for future products reassessment will be required for RF Exposure. In most cases GE would then do a Permissive Change to add necessary documentation.

Sincerely,



Thomas T. Smith
Director of EMC Compliance
Phone: 262.421.4986
Email: tom.smith@lairdtech.com