Abbott

26 August 2025

Federal Communications Commission
7435 Oakland Mills Road
Columbia, MD 21046

Re: Applicant: Thoratec Corporation dba Abbott Medical
FCC ID: **2BQFL-FLEXCTRL**
Part 2.933(b) Change in Identification of Equipment

Thoratec Corporation dba Abbott Medical requests a change in identification of a currently certified device under Part 2.933(b).

The original equipment identification is FCC ID: **QOQGM210P**
The original Grantee: Silicon Laboratories Finland Oy. was granted approval on 9/26/19.

The original test data filed by Silicon Laboratories Finland Oy continues to be representative of the newly identified device. There are no changes to the hardware or software.

This change in identification is requested for the newly identified device, FCC ID: **2BQFL-FLEXCTRL**, which is physically and electrically identical as defined by Part 2.907, to the originally approved module.

Photographs as required under Part 2.933(5) and FCC ID label are submitted with this request.

Please let us know if you might require anything further to assist with your review of this Change in Identification of Equipment.

Sincerely,

RANGAAX3 Digitally signed by RANGAAX3
Date: 2025.09.10 16:31:44 -07'00'

Archana Ranganathan
Senior Manager, Regulatory Affairs
Abbott Medical