

## **Class II Permissive Change Request Letter**

November 05, 2020

Federal Communications Commission  
7435 Oakland Mills Road  
Columbia, MD 21046

**Attention: Application Engineer / Review Engineer**  
**Subject: Request for Class II Permissive Change**  
**Reference FCC ID: 2AVN910631**

This is to request a Class II permissive change for FCC ID: **2AVN910631**, originally granted on 3/4/2020.

This module has a grant note restricting operation to mobile applications. This is a Class II Permissive Change request to allow operation in a portable host device manufactured by **ClearTrac Technologies, LLC**.

Manufacturer: **ClearTrac Technologies, LLC**  
Product Name: CarePath® Urine Flow Device  
Model Numbers: DCD-001F, DCD-001M  
FCC ID: **2AVN910631**

No changes to the design, circuitry or construction have been made to the transmitter module. A SAR report documenting exposure compliance in a portable application is being submitted with this request. A limited modular approval for use only in the CarePath® Urine Flow Device is sought with this request.

The CarePath® Urine Flow Device is Verizon certified for LTE Cat-M1 Band 4 and Band 13. All other LTE Cat-M1 bands associated with Nordic Semiconductor ASA, FCC ID: 2ANPO00NRF9160 have been disabled through software and exclusive use of Verizon LTE-M1 SIM cards. Therefore, the EMC and SAR reports only show test data for Band 4 and Band 13.

Please contact me if there are questions or additional information required concerning this request.

The following files are electronically submitted as attachments:

- Cover Letter
- External and Internal Photos
- Label Information
- Test Reports
- Test Setup Photographs

Sincerely,



Brent Laing, MD  
CEO  
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