infoscan
telemedical systems

6/03/2017

Federal Communications Commission
7435 Oakland Mills Road
Columbia, MD 21046

To whom it may concern:

This letter is to inform you that we, Infoscan S.A., are requesting Change in FCC ID for the new certification(s), FCC ID: 2ALKP-MED-RECORDER. The device to be certified uses identical in design and construction originally approved product.

In order to ensure continued compliance of the product, we have employed the following:

The originally approved product is still manufactured by the same company using the same design specifications.

The same PCB design, layouts and parts list of the originally approved product will be used to manufacture our product.

Cellular module installed in our Device is being tested after production phase of each unit according to the ISO 13485 standards which our company has applied. Furthermore, according to internal procedures, each unit is tested after each 12-month period during technical check-up.

Sincerely,

Bartosz Turczynowicz
Vice-President
Infoscan S.A.

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