GE Healthcare

GE Vingmed Ultrasound AS
Strandpromenaden 45
3183 Horten
Norway

T 3302 1100
F 3302 1320

Vscan Air CL – Label Information

Product Name : Vscan Air CL
FCC ID : YOM-VSCANAIR
IC: 9136A-VSCANAIR

A. Information to be displayed

1. The information that will be displayed on the e-label and information that will remain in the user manual is shown below:

Information	E-label	User Manual
FCC ID number	Yes	Yes
15.19 statement ("*This device complies with part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) This device may not cause harmful interference, and (2) this device must accept any interference received, including interference that may cause undesired operation.*")	No	Yes
Class A / B Digital Device user manual statements	No	Yes
Caution to the user that changes or modifications not expressly approved by the party responsible for compliance could void the user's authority to operate the equipment	No	Yes

Images of the e-label screen are provided below:

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B. Access to the required information on the e-label

1. Users can access the information in no more than three steps in a device's menu. The actual steps are:

To access the e-label information, in the device,

Choose: Menu > About > Regulatory

No special access codes or permissions are required to go through the above steps beyond entering a user-defined password(optional) to protect against unauthorized access to the application.

2. The information is stored on the device, no special accessories or supplemental plug-ins *(e.g.,* a SIM/USIM card) are required to access the information.

Yes.

3. Users are provided specific instructions on how to access the information.

The information to the user is provided in the product user manual.

C. Labeling for Importation and Purchasing

1. Products utilizing e-labels are required to have a physical label on the product at the time of importation, marketing and sales. For devices imported in bulk and not packaged individually, a removable adhesive label or, for devices in protective bags, a label on the bags is acceptable for this purpose. Any removable label shall be of a type intended to survive normal shipping and handling and must only be removed by the customer after purchase. For devices imported already in individual packages ready for sale, the information may alternatively be provided on the package. It shall contain:

    1. The FCC ID and/or the DoC logo (if applicable); and
    2. Any other information required by specific rule to be provided on the surface of the product unless such information is permitted to be included in the User's manual or other packaging inserts.

The FCC ID is present on the product packaging.

Model number is present on the product label, packaging label and shipping label.

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D. Other considerations

1. The above information must be programmed by the responsible party and the information must be secured in such a manner that third-parties cannot modify it.

The e-label information is pre-programmed by the grantee. The user cannot modify the e-label information.

2. All the applicable regulatory information required on the packaging or user instructions must be provided according to the rules even if it is displayed electronically. For example, hearing aid compatibility (HAC) ratings for the phones as specified in 47 C.F.R. § 20.19.

Yes, it is provided in User Manual.