

Airofit Pro TM (Medical Device)

000

COVER LETTER Airofit AF003 - FCC Part 15 Subpart B EKTOS Clarification

### Product

ID: -Name: -Model: -Version: AF003 Manufacturer: Airofit A/S

### Airofit AF003 COVER LETTER Test Report FCC Part 15 Subpart B

#### Author's Signature:

The signature indicates that this document has been prepared in accordance with expectations from the Quality Manual, applicable SOPs, and that Good Documentation Practices have been followed.

Author:

Meaning associated with the Signature, Date and Signature

Kenn Milton Consultant Airofit A/S

CONFIDENTIALITY:	PAGE:	LOCATION:		
Business Use Only	Only 1 of 5 MyBlueLabel			
Any print-out of this document is considered an "Uncontrolled Copy".				



Airofit Pro TM (Medical Device)

COVER LETTER Airofit AF003 - FCC Part 15 Subpart B EKTOS Clarification

## **Document History**

Version	Author	Date (DD-MMM-YYYY)	Comments
1.0	Kenn Milton	25-MAY-2022	This is the first approved version of this document

CONFIDENTIALITY: PAGE: LOCATION:			
Business Use Only 2 of 5 MyBlueLabel			
Any print-out of this document is considered an "Uncontrolled Copy".			



### Airofit Pro TM (Medical Device)

COVER LETTER Airofit AF003 - FCC Part 15 Subpart B EKTOS Clarification

## **Table of Contents**

1	Purpose	4
2	Clarification	5

CONFIDENTIALITY:	IFIDENTIALITY: PAGE:		
Business Use Only	3 of 5	MyBlueLabel	
Any print-out of this document is considered an "Uncontrolled Copy".			

	Α		0	F	1	Т
--	---	--	---	---	---	---

#### Airofit Pro TM (Medical Device)

COVER LETTER Airofit AF003 - FCC Part 15 Subpart B EKTOS Clarification

## **1** Purpose

The document is additional comments to the FCC Part 15 subpart B application.

 Test Report - P21-0064-2 rev 1.
 Date: 23-May-2022

 Test Report - P21-0064-2 rev 1 Appendix 1.
 Date: 23-May-2022

## 2 Radio State in Charging Mode

During normal charging conditions the BLE transmitter is disabled via software and therefore no AC mains conducted emissions is required.

Please see futher clarification below.

CONFIDENTIALITY:	PAGE:	LOCATION:	
Business Use Only	4 of 5	MyBlueLabel	
Any print-out of this document is considered an "Uncontrolled Copy".			



3.0

TITLE:

#### Airofit Pro TM (Medical Device)

000

COVER LETTER Airofit AF003 - FCC Part 15 Subpart B EKTOS Clarification

# 3 Clarification

From: David Busk <<u>dbu@ektos.net</u>>

Sent: Monday, May 16, 2022 7:56 AM

To: 'Jennifer Sanchez' <jennifers@acbcert.com>

**Cc:** 'Kenn Milton' <<u>kenn.milton@mybluelabel.com</u>>; <u>c@formika.dk</u>; 'Rudy' <<u>betronic@mail.dk</u>> **Subject:** RE: Airofit, yderlig spørgsmål fra ACB vedr. "ATCB028560, FCC ID: 2ATQX-AF003, ISED ID: 25191 -AF003"

Dear Jennifer,

Airofit has asked me to answer. I will explanation of our rational of testing below. 😊

The radio is turned off during battery charging mode. When the charger is connected, the SW makes the radio turn off.

This is why 15.207 was not tested in the 15.247 test report.

At the same time that we performed FCC 15B testing, EU immunity tests were also performed. The battery discharged so quickly from the high usage that a special software was implanted where the charger and radio were active at the same time.

This same setup, that was used for immunity tests, was also used for the FCC 15B testing. Thus, the BLE transmitter was operational during 15.109 and 15.107.

We agree that the BLE operation should have been completely shut off during FCC 15B measurements.

But the EUT version we had did transmit.

We deemed it to be OK for the measurements performed.

Best regards,

David Busk Lab. Manager, M.Sc. EE Phone: <u>+45 28 83 17 01</u> Email: <u>dbu@ektos.net</u>

### **EKTOS Testing & Reliability Services A/S**

Hammerholmen 45A, DK-2650 Hvidovre Internet: <u>www.ektos.net</u>

ONFIDENTIALITY: PAGE:		LOCATION:	
Business Use Only	Only <b>5 of 5</b> MyBlueLabel		
Any print-out of this document is considered an "Uncontrolled Copy".			