

F2 Labs 16740 Peters Road Middlefield, Ohio 44062 United States of America www.f2labs.com

| Manufacturing Address:                            | Bellus Medical, LLC DBA Crown Aesthetics<br>5005 Lyndon B. Johnson Fwy STE 370<br>Dallas, Texas 75244 USA |
|---|---|
| Applicant:  | Same as Above   |
| Product Name:                                     | SkinPen® Precision Elite Handpiece  |
| Product Description:                              | Microneedling device, 13.56 RFID  |
| Operating Voltage/Freq.<br>of EUT During Testing: | Battery-Operated (3.6VDC)   |
| Model(s):   | REF 200: SkinPen® Precision Elite Handpiece   |
| FCC ID:   | 2AGLK-REF-200   |
| IC:   | 21314-REF200  |

Standard(s):

- KDB447498 D04 Interim General RF Exposure Guidance v01
- FCC Rule Part 1.1307(b)(3)(i)(a)
- RSS-102: Issue 5, March 2015, Section 2.5 Radio Frequency (RF) Exposure Compliance of Radio Communication Apparatus (All Frequency Bands)

flindlithd

Report Constructed by:

Julius Chiller, Senior Wireless Project Engineer

The Ithe

Ken Littell, Vice President of Operations

F2 Labs 26501 Ridge Road Damascus, MD 20872 Ph 301.253.4500

**Report Reviewed by:** 

F2 Labs 16740 Peters Road Middlefield, OH 44062 Ph 440.632.5541 F2 Labs 8583 Zionsville Road Indianapolis, IN 46268 Ph 317.610.0611

This test report may be reproduced in full; partial reproduction only may be made with the written consent of F2 Labs. The results in this report apply only to the equipment tested.

# **TABLE OF CONTENTS**

- 1 ADMINISTRATIVE INFORMATION
- 2 <u>SUMMARY OF TEST RESULTS/MODIFICATIONS</u>
- 3 ENGINEERING STATEMENT
- 4 EUT INFORMATION AND DATA



# 1 ADMINISTRATIVE INFORMATION

### **1.1 Measurement Location:**

- F2 Labs in Middlefield, Ohio. Site description and attenuation data are on file with:
- FCC's Sampling and Measurement Branch at the FCC Laboratory in Columbia, MD.
- Certification and Engineering Bureau, Industry Canada, Site Number 4730B.

## 1.2 Document History

| Document Number  | Description Issue Date |            | Approved<br>By |
|------------------|------------------------|------------|----------------|
| F2P30652C-C5-06E | First Issue            | 2024-04-22 | K. Littell     |

# 2 SUMMARY OF RESULTS

|                  | Standard(s)  | Results  |
|------------------|--|----------|
| SAR<br>Exemption | <ul> <li>KDB447498 D04 Interim General RF Exposure<br/>Guidance v01</li> <li>FCC Rule Part 1.1307(b)(3)(i)(a)</li> <li>RSS-102, Section 2.5.1</li> </ul> | Complies |

| Modifications Made to the Equipment |
|-------------------------------------|
| None                                |



### **3 ENGINEERING STATEMENT**

This report has been prepared on behalf of **Bellus Medical, LLC DBA Crown Aesthetics**, to provide documentation for the SAR Exclusion herein. This equipment has been found to comply with the SAR Exclusion levels listed in KDB 447498 D01, FCC Rule Part 1.1307(b)(3)(i)(a) and RSS-102, Section 2.5.1

"(3) Determination of exemption.

(i) For single RF sources (i.e., any single fixed RF source, mobile device, or portable device, as defined in paragraph (b)(2) of this section): A single RF source is exempt if:

(A) The available maximum time-averaged power is no more than 1 mW, regardless of separation distance. This exemption may not be used in conjunction with other exemption criteria other than those in paragraph (b)(3)(ii)(A) of this section. Medical implant devices may only use this exemption and that in paragraph (b)(3)(ii)(A);"

The minimum distance from the antenna of the RFID Reader is 5mm. The maximum Field Strength of the transmitter was  $0.71dB\mu$ V/m at 3m which converts to -94.5 dBm, or  $0.0000004\mu$ W, which is well below the exemption limit of 1mW. For Canada, the Exemption limit specified in Table 11 of RSS-102, is 45mW for frequencies under 300 MHz.

| Frequency<br>(MHz) | ≤ 5 mm<br>(mW) | 10 mm<br>(mW) | 15 mm<br>(mW) | 20 mm<br>(mW) | 25 mm<br>(mW) | 30 mm<br>(mW) | 35 mm<br>(mW) | 40 mm<br>(mW) | 45 mm<br>(mW) |
|--------------------|----------------|---------------|---------------|---------------|---------------|---------------|---------------|---------------|---------------|
| ≤ 300              | 45             | 116           | 139           | 163           | 189           | 216           | 246           | 280           | 319           |
| 450                | 32             | 71            | 87            | 104           | 124           | 147           | 175           | 208           | 248           |
| 835                | 21             | 32            | 41            | 54            | 72            | 96            | 129           | 172           | 228           |
| 1900               | 6              | 10            | 18            | 33            | 57            | 92            | 138           | 194           | 257           |
| 2450               | 3              | 7             | 16            | 32            | 56            | 89            | 128           | 170           | 209           |
| 3500               | 2              | 6             | 15            | 29            | 50            | 72            | 94            | 114           | 134           |
| 5800               | 1              | 5             | 13            | 23            | 32            | 41            | 54            | 74            | 102           |



# EUT INFORMATION AND DATA

### 4.1 Equipment Under Test:

Product:SkinPen® Precision Elite HandpieceModel(s):REF 200: SkinPen® Precision Elite HandpieceFirmware Version:V0.61Software Version:N/ASerial No.:EP00005FCC ID:2AGLK-REF-200IC:21314-REF200

4.2 Trade Name: Bellus Medical, LLC DBA Crown Aesthetics

### 4.3 **Power Supply:**

Rechargeable Lithium Battery (3.6VDC) Device is battery-operated, rechargeable; cannot be used while charging.

- 4.4 Applicable Rules: KDB447498
- 4.5 Equipment Category: Radio Transmitter
- 4.6 Antenna: Internal Coil
- 4.7 Accessories: N/A