

EMF TEST REPORT

Test Report No. : OT-213-RWD-011

Reception No. : 2102000509

Applicant : Tianjin Empecs Medical Device Co.,Ltd.

Address : No.35,37, Yingcheng Street, Hangu, Binhai New Area, 300480 Tianjin, China

Manufacturer : Tianjin Empecs Medical Device Co.,Ltd.

Address : No.35,37, Yingcheng Street, Hangu, Binhai New Area, 300480 Tianjin, China

Type of Equipment : Blood Beta-Ketone Monitoring System

FCC ID. : 2AFE8BKQBT

Model Name : KT26 BT

Multiple Model Name: N/A

Serial number : N/A

Total page of Report : 7 pages (including this page)

Date of Incoming : February 09, 2021

Date of issue : March 04, 2021

SUMMARY

The equipment complies with the regulation; FCC PART 15 SUBPART C Section 15.247

This test report only contains the result of a single test of the sample supplied for the examination.

It is not a generally valid assessment of the features of the respective products of the mass-production.

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Reviewed by

Approved by

/ Su-Min You / Assistant Manager

/ Ha-Ram Lee / Manager

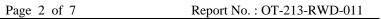
/ Ki-Hong, Nam/ General Manager

ONETECH Corp.

Tested by

ONETECH Corp.

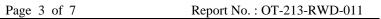
ONETECH Corp.





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Revision History

| Rev. No. | Issue Report No. | Issued Date | Revisions | Section Affected |
|----------|------------------|----------------|-----------------|------------------|
| 0 | OT-213-RWD-011 | March 04, 2021 | Initial Release | All |
| | | | | |
| | | | | |





1. VERIFICATION OF COMPLIANCE

Applicant : Tianjin Empecs Medical Device Co.,Ltd.

Address : No.35,37, Yingcheng Street, Hangu, Binhai New Area, 300480 Tianjin, China

Contact Person: Dongeon Cho/Senior Research Engineer

Telephone No.: 82(0)70-7124-0476

FCC ID : 2AFE8BKQBT

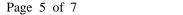
Model Name : KT26 BT
Brand Name : Medisign

Serial Number : N/A

Date : March 04, 2021

| EQUIPMENT CLASS | DTS – DIGITAL TRNSMISSION SYSTEM |
|---|--------------------------------------|
| E.U.T. DESCRIPTION | Blood Beta-Ketone Monitoring System |
| THIS REPORT CONCERNS | Original Grant |
| MEASUREMENT PROCEDURES | ANSI C63.10: 2013 |
| TYPE OF EQUIPMENT TESTED | Pre-Production |
| KIND OF EQUIPMENT | |
| AUTHORIZATION REQUESTED | Certification |
| EQUIPMENT WILL BE OPERATED | FOG DART 15 SURDART OF CALL 15 247 |
| UNDER FCC RULES PART(S) | FCC PART 15 SUBPART C Section 15.247 |
| Modifications on the Equipment to Achieve | Nama |
| Compliance | None |
| Final Test was Conducted On | 3 m, Semi Anechoic Chamber |

^{-.} The above equipment was tested by ONETECH Corp. for compliance with the requirement set forth in the FCC Rules and Regulations. This said equipment in the configuration described in this report, shows the maximum emission levels emanating from equipment are within the compliance requirements.





2. GENERAL INFORMATION

2.1 Product Description

The Tianjin Empecs Medical Device Co.,Ltd., Model KT26 BT (referred to as the EUT in this report) is a Blood Beta-Ketone Monitoring System. The product specification described herein was obtained from product data sheet or user's manual.

| Device Type | Blood Beta-Ketone Monitoring System | |
|--|-------------------------------------|--|
| Operating Frequency | 2 402 MHz ~ 2 480 MHz | |
| RF Output Power | -17.53 dBm | |
| Number of Channel | 40 Channels | |
| Modulation Type GFSK | | |
| Antenna Type | PCB Antenna | |
| Antenna Gain | -4.34 dBi | |
| Rated Supply Voltage | DC 3.0 V | |
| List of each Osc. or crystal Freq.(Freq. >= 1 MHz) | 32 MHz | |

2.2 Alternative type(s)/model(s); also covered by this test report.

-. None

3. EUT MODIFICATIONS

-. None





4. RF EXPOSURE EVALUATION

4.1 RF Exposure Calculation

According to the FCC rule 1.1310, the limit for General Population/Uncontrolled exposure is 1 mW/cm^2 for the device operating $1.500 \sim 100\,000 \text{ MHz}$.

4.2 EUT Description

| Kind of EUT | Blood Beta-Ketone Monitoring System | | |
|-----------------------------|-------------------------------------|--|--|
| | ■ Portable (< 20 cm separation) | | |
| Device Category | ☐ Mobile (> 20 cm separation) | | |
| | □ Others | | |
| | □ MPE | | |
| Exposure Evaluation Applied | □ SAR | | |
| | ■ SAR Test Exclusion Evaluation | | |

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4.3 Test Result of SAR Exclusion for Devices

According to the procedure, KDB 447498 D01, the standalone SAR test exclusion threshold is [(Max. Power of channel, including tune-up tolerance, mW)/(Min. test separation distance, mm)] X [$\sqrt{f(GHz)}$] < 3 = (0.02/5) X $\sqrt{2.402}$ =0.01

Conclusion: The SAR test exclusion threshold is less than 3, so the device meets the RF Exposure Requirement and excluded SAR Test.

| Mode Frequency (MHz) | Target Power W/tolerance | Max tune up power | Max tune up power | Separation distance | RF exposure | |
|----------------------|--------------------------|-------------------|-------------------|---------------------|-------------|------|
| | (MHZ) | (dBm) | (dBm) | (mW) | (mm) | |
| BLE | 2 402 | -17.53 ± 0.5 | -17.03 | 0.02 | 5 | 0.01 |