October 21, 1999

Mr. Thomas W. Phillips Federal Communications Commission Equipment Authorization Division 7435 Oakland Mills Road Columbia, Maryland 21046

Subject:FCC ID DV8LX5120AApplicant:Fukuda Denshi Co LtdCorrespondence Number:10281Reference:Attached Test Plan approved by George Tannahill 2/13/97 for a previous application of
this device (FCC ID DV8LX5120)

Dear Mr. Phillips:

Thank you for your e-mail this morning regarding the subject application. The following response and attachments address the requested information:

1. "Please explain how the upper frequency range and the frequency tolerance were determined."

The frequency range was selected based upon the licensing guidelines found in 90.267(a5). This section allows the operation of a biomedical telemetry device with an output power of not more than 20mW in the frequency range of 450 MHz to 470 MHz.

The frequency tolerance was calculated based upon the guidelines found in 90.213 and 90.217(b). The calculations are detailed in the attached test plan in the last paragraph of Section 2.6 (bottom of page 3).

2. "Also, please determine the necessary bandwidth and include the emission designator."

The necessary bandwidth calculation is detailed in the attached excerpt from the test report. Please see Section 5.0.

The emission designator "F1D" was selected based upon the guidelines in 2.201: "F" designates an emission in which the main carrier is frequency modulated. "1" designates a single channel containing digital information without the use of modulating sub-carrier (the applicant confirmed that no sub-carriers are used). "D" designates data transmission, telemetry. As detailed in the user manual, the device is used to transmit one way, non-voice, biomedical telemetry. Please see the attached excerpt from the test report, Section 4.0.

Your attention to this matter is greatly appreciated.

Sincerely,

AU.K.P

Greg Kiemel Director of Engineering

Attachments

4.0 Modulation Characteristics Data

4.1 Test Setup

The reference device uses a digital, frequency shift keying modulation scheme with no sub-carriers. The patient data (analog signals) are digitally encoded. The digital signal is used to frequency modulate the carrier with a maximum modulation frequency of 3.5 kHz and a maximum data rate of 7000 bits per second.

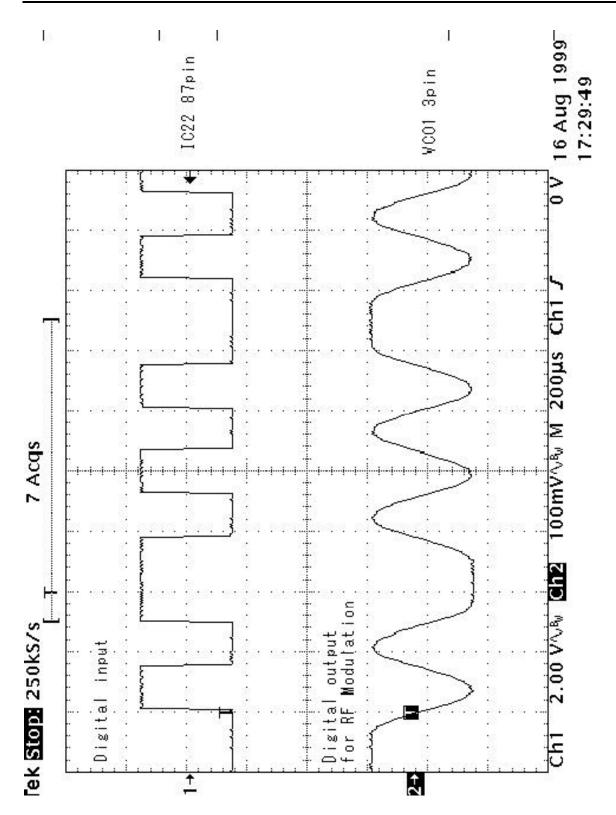
The emission designator "F1D" was selected based upon the guidelines in CFR 2.201: "F" designates an emission in which the main carrier is frequency modulated. "1" designates a single channel containing digital information without the use of a modulating sub-carrier (the applicant confirmed that no sub-carriers are used). "D" designates data transmission, telemetry. As detailed in the user manual, the device is used to transmit one way, non-voice, biomedical telemetry.

A waveform of the modulating signal as stored on the oscilloscope screen was recorded on a printer.

For telemetry operations, only A1D, A2D, F1D, and F2D emissions are authorized. The EUT utilizes F1D emissions.

4.2 Test Results

See data sheet on next page.



5.0 Necessary Bandwidth

The applicant specifies a maximum modulation frequency of 3.5 kHz. The necessary bandwidth, calculated using Carson's Rule: 2M + 2D = 2(3500) + 2(1750) = 10.5 kHz.

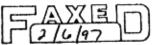
The applicant specifies a maximum data rate of 7kBpS (7000 bits per second).

Appendix I Test Plan and Faxed Approval from the FCC

FEDERAL COMMUNICATIONS COMMISSION

Equipment Authorization Division, Application Processing Branch 7435 Cakland Mills Road, Columbia, MD 21045

Telephone: (301) 725-1686, Facsimile: (301) 344-2050 197 Date: Time: From: George Tannahill Extension: 237 Kiene Grea To: EMC Organization: Telephone: (56 728 Facsimile: (< 31537 -0735 This cover sheet is page 1 Please direct inquiries to the sender at above extension. x dale Reference: Your The items indicated below must be submitted before processing can continue on the above referenced application, Failure to provide the requested information within 60 days may result in application dismissal pursuant to Section 2.917(c) and forfeiture of the filing fee pursuant to Section 1.1106. Block diagram(s) showing all clocks/oscillators or schematic diagrams \Box Additional detailed photographs of the described proce nOO 1 he \sim mes



Preliminary Test Plan - Subject to FCC Approval

Revision: 2/02/97

This device will be seeking type acceptance from the Commission under 90.267(a5) and 90.217(b).

1.0 EUT Specifications

Classification: Modulation Method: RF Output Power: Transmission Frequency Band: Channel Spacing: Occupied Bandwidth: One way, non voice, biomedical telemetry device Digital, Frequency Shift Keying (F1D) 1mW 460 to 470 MHz (approx. 474 channels) 12.5 kHz 8.5 kHz

2.0 Required Measurements

All measurements will be made with the EUT set to the lowest, a mid-range, and the highest frequency channels.

2.1 RF Power Output (2.985(a), 90.205, 90.217, 90.267(a5))

Power output will be measured at the RF output terminals with the transmitter biased for normal operation. The output power level is factory set (≈ 1 mW), and not adjustable by the user.

The output terminals will be directly coupled to a spectrum analyzer input through a 20dB (2W) attenuator. The characteristic impedance of the attenuator and the front end of the analyzer are 50 Ω . Power measurements will be made with the carrier unmodulated using both peak and average detectors on the spectrum analyzer. Resolution bandwidth will be set to 120kHz.

Maximum permissible power is 20mW or the minimum required for satisfactory operation. This measurement will verify 1mW of power output as specified by the manufacturer.

2.2 Modulation Characteristics (2.987(d))

The modulation characteristics will be measured at the RF output terminals with the EUT set for normal operation. A typical load (antenna or resistive load) will be attached to the output terminals. A high speed oscilloscope (\geq 500 MHz) will be connected to the output terminals through a high impedance, low capacitance probe (e.g. 10M Ω , 8pF probe). The carrier will be modulated (typical of normal operation).

A waveform of the modulating signal as stored on the oscilloscope screen will be recorded on a plotter.

For telemetry operations, only A1D, A2D, F1D, and F2D emissions are authorized. This measurement will verify F1D emissions.

2.3 Occupied Bandwidth (2.989(h), 2.202(c), 90.217(b))

The occupied bandwidth will be measured at the RF output terminals with the EUT set for normal operation. The output terminals will be directly coupled to a spectrum analyzer input through a 20dB (2W) attenuator.

The spectrum analyzer will be set with a peak detector, a resolution bandwidth of 10kHz, and a frequency span of 125kHz. No video filtering will be used. The carrier will modulated (typical of normal operation). The peak envelope of the out-of-band emissions, as stored on the spectrum analyzer screen, will be recorded on a plotter.

"...the sum of the bandwidth occupied by the emitted signal plus the bandwidth required for frequency stability shall be adjusted so that any emission appearing on a frequency 25 kHz or more removed from the assigned frequency is attenuated at least 30 dB below the unmodulated carrier."

2.4 Spurious emissions at antenna terminals (2.991, 2.997, 90.217(b))

No testing is required for spurious emissions at the antenna terminals <u>if the antenna is integral to the</u> construction of the EUT. If the antenna is detachable, then measurement data is required:

Spurious emissions will be measured at the RF output terminals with the EUT set for normal operation. The output terminals will be directly coupled to a spectrum analyzer input through a 20dB (2W) attenuator. The carrier will be modulated (typical of normal operation).

The spectrum will be scanned from 30MHz to 1GHz with a 10kHz resolution bandwidth and a peak detector. The spectrum will also be scanned from 1GHz to 5GHz (10th harmonic of carrier is 4.6 to 4.7 GHz) with a resolution bandwidth of 1MHz and a peak detector. No video filtering will be used. The peak envelope of the spurious emissions, as stored on the spectrum analyzer screen, will be recorded on a plotter. The magnitude and frequency of any spurious emission attenuated less than 40 dB (within 20 dB of the limit) below the power of the unmodulated carrier will be recorded.

For up to 5GHz, any spurious emission appearing on a frequency 25 kHz or more removed from the assigned frequency should be attenuated at least 30 dB below the unmodulated carrier.

2.5 Field strength of spurious radiation (2.993, 90.217(b))

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The field strength of spurious emissions that may be radiated from the EUT will be measured in the far field. The EUT will be taken to an Open Area Test Site on file with the FCC and set for normal operation. The carrier will be modulated by patient data from the host device (typical of normal operation). The EUT's antenna will be attached to the transmitter.

The measurement antenna to EUT test distance will be 3 meters. With the spectrum analyzer connected to the appropriate measurement antenna, the spectrum will be scanned from 30MHz to 1GHz with a 10kHz resolution bandwidth and a peak detector. The spectrum will also be scanned from 1GHz to 5GHz with a resolution bandwidth of 1MHz and a peak detector. No video filtering will be used.

The measurement antenna will be scanned in height from 1 to 4 meters and the orientation of the EUT varied to maximize the level of any spurious emissions. Each spurious emission will be maximized with the magnitude and frequency recorded for any signal attenuated less than 40 dB (within 20 dB of the limit) below the power of the unmodulated carrier. Measurement procedure ANSI C63.4 (1992) will be referenced.

For up to 5 GHz, the field strength of any radiated spurious emission appearing on a frequency 25 kHz or more removed from the assigned frequency should be attenuated at least 30 dB below the unmodulated carrier.

2.6 Frequency Stability (2.995(a1)(b)(d), 90.213, 90.217(b))

The frequency stability of the EUT's unmodulated carrier will be measured. The transmitter will be biased for normal operation. The output power level is factory set (≈ 1 mW), and not adjustable by the user. Although 2.995(a1) specifies that the ambient temperature will be varied from -30° to +50° centigrade, the EUT is specified to be used from +10° to +40° centigrade (hospital or clinic environment). The test will be destructive if performed from -30° centigrade. Therefore, frequency measurements will be made at the extremes of the operating range (+10° to +40° centigrade), and at intervals of not more than 10° centigrade through the range. A sufficient period of time will be allowed at each temperature interval to allow the components of the oscillator / clock circuitry to stabilize.

Frequency stability of the unmodulated carrier will also be measured for variations in supply voltage. For AC powered devices, the primary supply voltage will be varied from 85% to 115% of the nominal value. Frequency measurements will be made at the nominal supply voltage and at each extreme.

For battery powered devices, the primary supply voltage will be reduced to the battery operating end point specified by the manufacturer. Frequency measurements will be made at the nominal battery voltage and at the end point.

The carrier frequency must be within the assigned frequency by a margin no greater than the sum of: the bandwidth occupied by the emitted signal, plus the bandwidth required for frequency stability (2.5ppm \approx 1.15kHz), plus 25 kHz. This is equal to 12.5 kHz + 1.15 kHz + 25 kHz, which is equal to 38.65 kHz.