



Food and Drug Administration
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CareRay Digital Medical System Co., Ltd.
% Mr. Xu Wei
Manager
B3-501, Biobay, 218 Xinghu Street
SuZhou Industrial Park
SuZhou, Jiangsu 215123
P. R. CHINA

December 1, 2016S

Re: K163019
Trade/Device Name: CareView 750Cw/750C X-ray Flat Panel Detectors
Regulation Number: 21 CFR 892.1680
Regulation Name: Stationary x-ray system
Regulatory Class: II
Product Code: MQB
Dated: July 1, 2016
Received: November 3, 2016

Dear Mr. Wei:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Division of Industry and Consumer Education at its toll-free number (800) 638 2041 or (301) 796-7100 or at its Internet address

<http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>. Also, please note the regulation entitled, “Misbranding by reference to premarket notification” (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to

<http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm> for the CDRH’s Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address

<http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>.

Sincerely yours,

A handwritten signature in blue ink that reads "Michael D. O'Hara". The signature is written over a large, faint, grey watermark of the FDA logo.

For

Robert Ochs, Ph.D.
Director
Division of Radiological Health
Office of In Vitro Diagnostics
and Radiological Health
Center for Devices and Radiological Health

Enclosure

Indications for Use

510(k) Number (if known)

K163019

Device Name

CareView 750Cw/750C X-ray Flat Panel Detectors

Indications for Use (Describe)

The CareView 750Cw/750C detector is indicated for digital imaging solution designed for providing general radiographic diagnosis of human anatomy. It is intended to replace radiographic film/screen systems in all general-purpose diagnostic procedures. This product is not intended for mammography applications.

Type of Use (Select one or both, as applicable)

☒ Prescription Use (Part 21 CFR 801 Subpart D)

☐ Over-The-Counter Use (21 CFR 801 Subpart C)

CONTINUE ON A SEPARATE PAGE IF NEEDED.

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510(k) Summary

510(k) Summary

[As required by 21 CFR 807.92]

1. Date Prepared [21 CFR807.92 (a) (1)]

Aug.01, 2016

2. Submitter's Information [21 CFR807.92 (a) (1)]

Company Name:	CareRay Digital Medical System Co., Ltd.
Company Address:	A2-201/B3-501, Biobay, 218 Xinghu Street, SuZhou Industrial Park, SuZhou 215123, P. R. China
Contact Person:	Mr. Xu
Phone Number:	(86) 512-86860288
Fax Number:	(86) 512-86860388
E-mail:	wei.xu@careray.com

3. Trade Name, Common Name, Classification [21 CFR807.92(a)(2)]

Trade Name:	CareView750Cw/750C X-ray Flat Panel Detectors (hereinafter: CareView750Cw/750C or CareView750Cw/750C detector , apply to the all submission documents)
Model Name:	CareView 750Cw/750C
Classification Name:	Stationary X-ray system
Regulation Number:	21 CFR 892.1680
Regulatory Class:	Class II
Product Code:	MQB

4. Identification of Predicate Device(s) [21 CFR 807.92(a)(3)]

The identified predicates within this submission are as follows:

Manufacturer:	CareRay Digital Medical System Co., Ltd.
Trade Name:	X-ray Flat Panel Detectors

Model Name:	CareView 1500Cw
Classification Name:	Stationary X-ray system
Regulation Number:	21 CFR 892.1680
Regulatory Class:	Class II
Product Code:	MQB
FDA 510(k) #:	K150929

5. Description of the Device [21 CFR 807.92(a)(4)]

CareView 750Cw/750C is a kind of wireless(750C does not have wireless, it has only wired function.) portable digital X-ray flat panel detectors which have 244 mm × 307 mm imaging area. The device communicates by not only the wireless communication but also wired communication feature (Giga-bit Ethernet communication mode by connecting the power box) optionally.

The device intercepts X-ray photons and then the scintillator emits visible spectrum photons that illuminate an array of photo detectors (a-Si) that create electrical signals. After the electrical signals are generated, it is converted to a digital value and an image will be displayed on the monitor.

The detector should be integrated with an operating PC and an X-ray generator to utilize as digitalizing X-ray images and transfer for radiography diagnostic.

The detector can't provide feedback to the generator to terminate the x-ray exposure.

System Software Version is API V4.5.0 (NDT V3.5.5) that based on V4.2.0. But V4.5.0 is more powerful than old version. Please see the comparison of section 7 Technological Characteristics.

6. Intended Use [21 CFR 807.92(a)(5)]

The CareView 750Cw/750C detector is indicated for digital imaging solution designed for providing general radiographic diagnosis of human anatomy. It is intended to replace radiographic film/screen systems in all general-purpose diagnostic procedures. This product is not intended for mammography applications.

7. Technological Characteristics [21 CFR 807.92(a)(6)]

Item	Proposed Device: CareView750Cw/750C X-ray Flat Panel Detectors	Predicate Device: CareView 1500Cw X-ray Flat Panel Detectors
510(K) Number	K163019	K150929

Model	CareView 750Cw/750C	CareView 1500Cw
Classification Name	Stationary X-ray system	Stationary X-ray system
Product Code	MQB	MQB
Regulation Number	892.1680	892.1680
Panel	Radiology	Radiology
Class	II	II
X-ray Absorber	CsI Scintillator	CsI Scintillator
InstallationType	Wireless (except 750C) ,Wired, Portable	Wireless, Wired, Portable
ReadoutMechanism	Thin Film Transistor	Thin Film Transistor
Image Matrix Size	2048 × 2560 pixels	2304 × 2816 pixels
Pixel Pitch	120μm	154μm
Effective Imaging Area	244 mm × 307 mm	355 mm × 434 mm
Grayscale	16 bit, 65536 grayscale	16 bit, 65536 grayscale
Spatial Resolution	Min. 4.2 line pair/mm	Min. 3.3 line pair/mm
Rated Power Supply ● Wired ● Wireless	DC +24 V, Max. 1.3A Powered by the switching power supply using interface cable Powered by the battery pack(except 750C)	DC +24 V, Max.1.5 A Powered by the power box using interface cable Powered by the battery pack
Power Consumption	Max. 31.2 W	Max. 36 W
Communications ● Wired ● Wireless	Gigabit Ethernet IEEE 802.11a/b/g/n (2.4 / 5 GHz)	Gigabit Ethernet IEEE 802.11a/b/g/n (2.4 / 5 GHz)
Imaging Plate	Carbon Fiber Plate	Carbon Fiber Plate
Cooling	Air cooling	Air cooling
Dimensions	282mm × 333mm × 15mm	384 mm × 460 mm × 15 mm
Operation	Temperature: +5 ~ +35℃ Humidity: 30 ~ 75% (Non-Condensing) Atmospheric pressure: 70 ~ 106 kPa Altitude: Max.3000 meters	Temperature: +5 ~ +35℃ Humidity: 30 ~ 75% (Non-Condensing) Atmospheric pressure: 70 ~ 106 kPa Altitude: Max.3000 meters
Storage and transportation	Temperature: -20 ~ +55℃ Humidity: 10 ~ 90% (Non-Condensing) Atmospheric pressure: 70 ~ 106 kPa Altitude: Max.3000 meters	Temperature: -20 ~ +55℃ Humidity: 10 ~ 90% (Non-Condensing) Atmospheric pressure: 70 ~ 106 kPa Altitude: Max.3000 meters
Intended Use	The CareView 750Cw/750C detector is indicated for digital imaging solution designed for providing general radiographic	The CareView 1500Cw detector is indicated for digital imaging solution designed for providing general radiographic diagnosis of

	diagnosis of human anatomy. It is intended to replace radiographic film/screen systems in all general-purpose diagnostic procedures. This product is not intended for mammography applications.	human anatomy. It is intended to replace radiographic film/screen systems in all general-purpose diagnostic procedures. This product is not intended for mammography applications.
MTF	~70%.....(@ 1lp/mm) ~40%.....(@ 2lp/mm) ~22%.....(@ 3lp/mm)	~70%.....(@ 1lp/mm) ~40%.....(@ 2lp/mm) ~22%.....(@ 3lp/mm)
DQE	(@RQA5, 30μGy) ~65%.....(@ 0lp/mm) ~20%.....(@ 3lp/mm)	(@RQA5, 30μGy) ~65%.....(@ 0lp/mm) ~20%.....(@ 3lp/mm)
Dynamic range	~82 dB	~82 dB
System Software Version	API V4.5.0 (NDT V3.5.5) Based on V4.2.0, V4.5.0 includes the following new enhancements: ➤ CarerayTouch handswitch ➤ Resume broken downloads ➤ One-shot calibration ➤ Small-window AED	API V4.2.0 (NDT V3.4.0)

8. System requirements to operate with other radiographic system components

1) Recommended Generator Specification:

Energy range: 40~150kVp

mA range: 10~1000mA(depending on the generator power)

ms range: 10~6300ms to produce 0.1~1000mAs (depending on the generator power)

Note: To our best knowledge, the detector is compatible with the X-ray generators with the specifications described above. If you have questions regarding the compatibility issue for other generators, please contact your distributor or CareRay.

2) Application Program Interface (API) for system integration manufacturer

Peripheral hardware: CareView detector connected via wired communication.

CPU: Intel (R) Core (TM) 2 Duo, 2.93GHz or above

RAM: 2 GB or higher

Hard disk: 160 GB or higher

Monitor: 1280 x 1024 or higher

OS: Windows XP or Windows 7

Development environment: MS Visual Studio 2005

3) X-ray exposure mode

The synchronous connection mode is the signal transfer mode between the X-ray generator which sends the X-ray and the detector which receives the X-ray.

CareView 750Cw/750C supported typical sync mode contains external sync, soft sync, manual sync and auto sync containing FFAED mode.

The detector can't provide feedback to the generator to terminate the x-ray exposure.

9. Substantial Equivalence [21 CFR 807.92(b) (1) and 807.92(b)(2)]

➤ Electrical safety and EMC testing

Electrical, mechanical, environmental safety and performance testing according to IEC/ES 60601-1 was performed, and EMC testing was also conducted in accordance with IEC/EN 60601-1-2. All test results are satisfactory.

➤ Performance Testing

The performance testing items included Offset Stability, Dark Noise, Spatial Resolution, Low-contrast Resolution, Dynamic Range&Sensitivity, Lag, Ghost, MTF and DQE etc. These testing conducted in accordance with IEC61223-3-1, IEC 62220-1 and FDA's SSXI guidance document (Guidance for the Submission of 510(k)s for Solid State X-ray Imaging Devices). The RF module that provided by supplier is in compliance with ANSI C63.4-2003 and 47 CFR FCC Part 15 Subpart C and get FCC ID.

➤ Nonclinical and clinical considerations

The proposed device (CareView 750Cw/750C) and predicate device (CareView 1500Cw) share most of primary product specifications including intended use, technology, material, and imaging principle, etc. Difference lies in the mechanical structure.

The differences of mechanical structure don't affect the technological parameters and clinical images.

10. Conclusion [21 CFR 807.92(b)(3)]

In accordance with the Federal Food, Drug and Cosmetic Act, 21 CFR Part 807 and based on the information provided in this premarket notification, the CareView 750Cw/750C X-ray flat panel detector is substantially equivalent to predicate device CareView 1500Cw (K150929). Both proposed and predicate devices are same in the intended use, the design principle, the applicable standards and specification. Some characteristics, for example, the mechanical structure are different. However the test reports in this submission documents

provide demonstration that these differences do not raise any new questions of safety and effectiveness. Therefore, CareRay Digital Medical System Co., Ltd. concludes the CareView750Cw/750C X-ray flat panel detector is substantially equivalent with the predicate device CareView1500Cw (K150929).