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MEDICAL ELECTRIC SAW DRILL

Operating Manual **E Series**

RJ-010007-Rev:02



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Please read this manual carefully before use and keep it properly

FOREWORD

Dear user

Thank you for using this product! This manual contains important contents that must be understood for safe and proper use of the product.

The instruction manual is a part of the product. Therefore, this manual should be placed in a proper position for easy reference at any time for the whole lifetime. The product must be operated by trained personnel with relevant knowledge and experience. All personnel must read this instruction carefully before using this product.

This manual shall be transmitted to subsequent owners or users of the product. Any duplication, photogrammetry, digital post-processing or duplication is prohibited without consent of Wuhu Ruijin Medical Instrument & Device Co., Itd.

This product is safe and reliable to use, except for the risk s arising from special factors like operated by non-professionals or used for other purposes, etc. Therefore, please follow strictly the following rules to avoid accidental use. The product must be operated by professionals.

Product maintenance must be carried out by professional technicians or skilled users of Wuhu Ruijin Medical Instrument & Device Co.,ltd and the authorized person.

If there are any operation problems not mentioned, please contact WuhuRuijin Medical Instrument & Device Co..ltd.

Instructions are subject to change without prior notice. Please put the manual in a proper position for reference.

Product introduction

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Product introduction

I.Product information

Product Name: Medical Electric Saw Drill
Product Model No.: RJ-MP RJ-PD RJ-PS

Product technical requirement No.: W.X.Z.Z No.:20162040284

II.Corporate information

Registrant/manufacturer name: Wuhu Ruijin Medical Instrument & Device Co., Ltd. Registrant's Residence/ Production Address: 33rd , East Wanchun Road, Wuhu Economic and Technological Development Zone, Wuhu , anhui Province, P.R.C. After-sales service unit: Wuhu Ruijin Medical Instrument & Device Co., Ltd TEL: 0553-2673688 FAX: 0553-2672513 Zin Code: 241000

Production License: W.X.Z.Z.XU NO.: 20160013

Medical Device Registration Certificate: W.X.Z.Z. No.: 20162040284

III. Main performance parameters

Sequence	Model No.:	Performance parameters
1	RJ-MP	Main performance parameters: Non-loading rotation speed or frequency of electricsaw and drill: i. Drill rotating speed 120rpm ii. Saw frequency: >6000 times/min
2	RJ-MP	 iii.Output power: ≥50W iv. Temperature rise: the temperature rise of the shell is no more than 50°C after 5min of non-loading operation;
3	RJ-PS	 v. Non-loading noise: drill saw non-loading noise ≤75dB(A); vi. Electric saw & drill saw blades shall be heat treated and its hardness shall be no less than 30 HRC.

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IV.The structural composition

This product is mainly composed of handpiece, drill components, saw components, accessories, battery and charger.

i.RJ-MP

(i).EM100

Note

All instructions in this manual are for the purpose of explanation only. If the appearance and function are inconsistent with the real object, the real object shall prevail.



ii.RJ—PS

Note

All instructions in this manual are for the purpose of explanation only. If the appearance and function are inconsistent with the real object, the real object shall prevail.



ii.RJ-PD

Note

All instructions in this manual are for the purpose of explanation only. If the appearance and function are inconsistent with the real object, the real object shall prevail.



V.The scope of application

Applied in bone drilling and cutting in orthopedic surgeries for medical organization.

VI. Contraindications, precautions, warnings and explanatory notes

i.Contraindications

NO

ii.Precautions

(i) Check whether the radial run-out of the bit is intact before use. The bit with large radial run-out cannot be used. Check the sharpness of the drill bit. The drill body should be fixed during drilling without loosening according to conventional surgical methods.

(ii). The drill bit must be firmly fixed well with drill chuck key to avoid accidents. Products and accessories should be placed in a dry place to prevent corrosion after use.

(iii). The battery shall not be disinfected by high temperature, high pressure or stay in the machine for a long time, take out battery after the operation. If power is cut off during the operation (the handpiece does not work), the backup battery should be replaced immediately.

(iv). If there is abnormal sound when using stop using it immediately. Please contact with the manufacturer or distributor immediately and return the machine to the manufacturer for repairing.

(v). Charge the battery immediately after use. If the battery is not used, charge it every three months to prevent battery loss.

iii.Warning and suggestive notes

The medical electric drill saw is classified according to the protection of electronic equipment from shock and vibration. The motor belongs to B-type equipment of internal power supply, and the charger belongs to class II B-type ordinary equipment; The equipment shall not be used with a mixture of flammable anesthetic gas or carbon oxide: Sterilization must be carried out before use.

Supplementary external mark: A Working schedule: Short time operation Each starup time shall not exceed 5min.

Marking pattern description: 🛧 B-type equipment

Rotation mark: F: Forward R: Reverse S: Stop ■ Operation

Battery cover marking: Unlock Lock

Recycling marks for electronic products at the end of their life:

VII.Installation and dis-assembly

This product is professional medical device, the installation and dis-assembly must be performed by medical personnel with appropriate technology or with appropriate professional training.

i.Installation and dis-assembly of drill bits

- (i).Loosen the chuck by turning counterclockwise with the drill chuck key;
- (ii). Insert the prepared bit into the three-iaw hole
- (iii). Turn the drill chuck key clockwise and tighten it vigorously.
- (vi). Conversely, the drill bit can be removed



ii.Installation and removal of oscillating saw blade

Press the button first, then put the saw blade into the interface of the saw blade. align the position, the saw blade hole falls into the circular column of the interface of the saw blade, then release the button to lock the saw blade.



iii.Installation and removal of drill saw adapters.

- (i). Align the assembly installation position with the handpiece, and pay attention to the phase position:
- (ii). Turn the knob on the handpiece;
- (iii). Release the knob after insertion, jam the drill or saw adapter and complete installation:
- (iv). Converse to detach components.



iv.Battery installation

- (i). The battery can not be directly connected with the handpiece to prevent pollution.
- (ii). Before installing the battery, install the battery aseptic transfer ring onto the battery box, insert thebattery into the battery compartment through the battery aseptic transfer ring, press it into place, remove transfer ring, and close the bottom cover.



VIII.Operating instructions

This product must be disinfected before it can be used and it must be tested before use after disinfection. The method is: join the corresponding battery for the handpiece, press the trigger gently, the motor should rotate, switch forward and reverse, the motor should work, or else, the handpiecehas a problem, then please stop using immediately, contact the manufacturer or distributor to send product back to the manufacturer for maintenance.

i.Battery charging

- (i). Plug the DC port of the charger terminal into the DC port of the charging base, and insert the battery into the charging base to charge.
- (ii). When the charger is connected to the power supply, the indicator is green. After inserting battery, the blue light indicates charging, and the green light shows fully charged.



ii. Opening and closing of bottom cover



iii.Mode adiustment

- (i). When it is necessary to turn forward, move the lever to the corresponding F mark:
- (ii).To reverse, use the same method to move the lever to the R mark;
- (iii). Move the lever to position S when installing or removing the drill bits or saw blade and while not in use.



Maintenance and cleaning

IX.Maintenance

This product is maintenance-free. It contains no parts that require maintenance by the user or manufacturer. However, the manufacturer recommends that the function and safety of the product be checked regularly by a professional or hospital technician. It must be checked once before each operation, and be recorded, analyzed and evaluated in time to ensure that the product is in good condition and guarantee the use quality.

X.Cleaning and sterilization

- A.Cleaning: the protection of cleaning after use is same as sterilization, and the handpiece can be wiped clean with gauze soaked in water or alrohol and then store
- B.Disinfection: Product can be used only when passing inspection after disinfection and sterilization according to the hospital standards.

Note: Do not soak any part of the medical electric saw drill in liquid.

Recommended handpiece disinfection method

Disinfection method	Operating conditions and environment	Scope of application	Limitation
Moist heat sterilization	Sterilization conditions are generally are: 121°C*15min,121°C*30min or 116°C*40min	High temperature and humid product sterilization	High temperature High humidity It is easy to breed bacteria again when the product is wet

Commitment

- i:This instrument has no parts that can be disassembled and repaired by non-professionals. If it fails, please contact the company's after-sales service department.
- ii: If you need technical information about this medical electric drill, please write to us.

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Service Guide

XI.Transport and storage conditions

Attention

Parameters in this table are for reference only. In case of any conflict between this table and the information on the nameplate of the machine, the nameplate prevails.

Transport and	Ambient temperature range	-10°C∼+40°C
Transport and storage	Relatively moderate range	≤90%
conditions	Atmospheric pressure range	500hPa∼1060hPa
	Ambient temperature range	5°C~40°C
	Relatively moderate range	≤70%
Equipment operating	Atmospheric pressure range	860hPa~1060hPa
conditions	Charger power	100-240V±10%;50/60Hz±1Hz
	Main power supply (DC)	7.4V±10%

Note: According to YY0904-2013 battery-powered bone tissue surgery equipment

Production date: Refer to product certificate Service life: two years short-time running 1000 times

XII.Accessories list

Prepare standard configuration and accessories according to product model number and product customer need, such as such as saw blade, drill bits, battery aseptic transfer ring, drill chuck keys,etc.

Parts List					
No.	Name	Model No.	Quantity	Note	
1	Handpiece	XXXX	1pcs	According to customer needs	
2	Battery	XXXX	2pcs	According to the handpiece model	
3	Charger	XXXX	1pcs	According to the handpiece model	
4	Charger base		1pcs	According to the handpiece model	
5	Manual		1pcs		
6	Certificate of conformity		1pcs		
7	Three certificates		1pcs		

XIII. Electromagnetic compatibility

Attention .

Parameters in this table are for reference only. In case of any conflict between this table and the information on the nameplate of the machine, the nameplate prevails.

Guidance and manufacturer's declaration – electromagnetic emission

The RJ-MP(RJ-PS;RJ-PD) is intended for use in the electromagnetic environment specified below. The customer or the user should assure that it is used in such an environment.

Emissions test	Compliance	Electromagnetic environment - guidance
RF emissions GB4842	Group 1	The RJ-MP(RJ-PS;RJ-PD) uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emissions GB4842	Class A	The RJ-MP(RJ-PS;RJ-PD) is suitable for use in all
Harmonic emissions GB17625.1	Not applicable	establishments other than domestic and those directly connected to the public low-voltage power supply network
Voltage fluctuations flicker emissions GB17625.1	Not applicable	that supplies buildings used for domestic purposes.

Guidance and manufacturer's declaration – electromagnetic immunity

The RJ-MP(RJ-PS;RJ-PD) is intended for use in the electromagnetic environment specified below. The customer or the usershould assure that it is used in such an environment.

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Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment -guidance
Electrostatic discharge GB/T 17626.2	± 6 kV contact ± 8 kV air	± 6 kV contact ± 8 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30 %.
Electrostatic transient / burst GB/T 17626.4	± 2 kV for power supply lines	± 2 kV for power supply lines	Mains power quality should be that of a typical commercial or hospital environment.
Surge GB/T 17626.5	± 1 kV wire to wire	± 1 kV wire to wire	Mains power quality should be that of a typical commercial or hospital environment.
Voltage dips, short interruptions and voltage variations on power supply input lines GB/T 17626.11	0 % UT; 0,5 cycle g) At 0°, 45°, 90°, 135°, 180° 225°, 270° and 315° 0 % UT; 1 cycle and 70 % UT; 25/30 cycles Single phase: at 0° 0 % UT; 250/300 cycle	0 % UT; 0,5 cycle g) At 0°, 45°, 90°, 135°, 180° 225°, 270° and 315° 0 % UT; 1cycle and 70 % UT; 25/30 cycles Single phase: at 0° 0 % UT; 250/300 cycle	Mains power quality should be that of a typical commercialor hospital environment. If the user of the RI-MP (RJ-PS,RI-PD) requires continued operation during power mains interruptions, it is recommended that the RI-MRI(RI-PS,RI-PD) be powered from an uninterruptible power supply or a battery.
Power frequency (50 Hz) magnetic field	3 A/m	3 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.

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A Attention

Parameters in this table are for reference only. In case of any conflict between this table and the information on the nameplate of the machine, the nameplate prevails,

Guidance and manufacturer's declaration – electromagnetic immunity

The RJ-MP(RJ-PS:RJ-PD) is intended for use in the electromagnetic environment specified below. The customer or the user should assure that it is used in such an environment.

Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment -guidance
Conducted RF GB/T 17626.6 Radiated RF GB/T 17626.3	3 V(Effective value) 150 kHz to 80 MHz 3 V(Effective value) 80 MHz to 2.5 GHz	3 V (Effective value) 3 V/m	Portable and mobile radio frequency communication equipment should not be used in any part of RJ-MP (RJ-PS, RJ-PD), including cables, than the recommended isolation distance. The distance should be calculated by the formula corresponding to the transmitter frequency. Recommended isolation distance d=1.2 d=1.2 80MHz-800MHz d=1.2 80MHz-800MHz d=1.2 80MHz-800MHz d=1.2 800MHz-800MHz d=1.2 800MHz-800MHz d=1.2 80MHz-800MHz d=1.2 800MHz-800MHz d=1.2 80MHz-800MHz d=1.2 80
			and in each frequency range b should be lower than the compliance level.Interference may occur near equipment marked with symbols as beside.

NOTE 1 At 80 MHz and 800 MHz, the higher frequency range applies.

NOTE 2 These guidelines may not apply in all situations. Electromagnetic is affected by absorption and reflection from structures, objects and people.

b Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio. AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measuredfield strength in the location in which the RJ-MP(RJ-PS;RJ-PD) is used exceeds the applicable RF compliance level above, the RJ-MP (RJ-PS:RJ-PD) should be observed to verify normal operation. If abnormal performance is observed. additional measures may be necessary, such as reorienting or relocating the RJ-MP(RJ-PS;RJ-PD).

c Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3V/m.

A Attention

Parameters in this table are for reference only. In case of any conflict between this table and the information on the nameplate of the machine, the nameplate prevails,

Recommended isolation distance betweenportable and mobile video communication equipment and RJ-MP (RJ-PS, RJ-PD)

RJ-MP (RJ-PS, RJ-PD) is expected to be used in an electromagnetic environment with controlled radio frequency radiation disturbance. According to the maximum rated output power of communication equipment, the purchaser or user can prevent electro magnetic waves by maintaining the minimum distance between portable and mobile radio frequency communication equipment (transmitter) and R.J-MP (R.J-PS, R.J-PD) as recommended below interference.

	Interference test	IEC60601 test level	Coincidence level	Electromagnetic	
	Maximum rated	Corresponding to the isolati	ion distance of the transmitte	r at different frequencies/m	
	output power of the transmitter	150kHz∼80MHz d=1.2	80MHz~800MHz d=1.2	800MHz~2.5GHz d=2.3	
	0.01	0. 12	0. 12	0.23	
	0.1	0.38	0.38	0.73	
	1	1.2	1.2	2.3	
	10	3.8	3.8	7.3 ((🕶))	
	100	12	12	23	

For the maximum rated output power of the transmitter not listed in the above table, the recommended isolation distance d, in meters (m), can be determined by the formula in the corresponding transmitter frequency column, where P is the emission provided by the transmitter manufacturer The maximum rated output power of the machine, in watts (w).

NOTE 1 At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.

NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and from structures, objects and people.