

High / Low Speed Handpiece for Dental

EMB-901 Built In

Fiber Optic Brushless Electric Micromotor Set

Operation Manual



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1.Component Instructions

Fiber Optic Brushless Electric Micromotor System for Dental Handpieces. Set Model: EMB-901 Built in. Consists of electric motor, control box, driver module and motor cord. Production date: Please check the motor control box label.

Brushless electric motor	Control Box	Driver Module	Motor Cord
Permanent magnet brushless motor supplies varying torque output. Model : EMB-901-MOTOR	Motor speed and direction control and digital RPM display. Model : EMB-901-Control Unit Built in	Execute commands to drive electric motor running. Model : EMB-901-Power Module Built in	Connect electric motor and control box, providing water, spray, and electricity. Model : EMB-901-Cable Built in
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2.Symbols

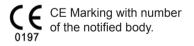


Ground Connection Protection earth.



Caution! Refer to accompanying documents.









Type B device.



Alternating Current



Recyclable Materials



Recyclable electrical and electronic materials.

3.Description of Products

This product is a comprehensive control equipment of dental electric motor, utilizing precise electric motor as the primary drive source with LED of high brightness as light source. This product has the following features: Free of regular maintenance, microvibration and low noise, ease of operation, and easy to use. Simply jointed water and spray pipes with motor, and connected air pipe with driver module, you can use the original air-powered control system with different speed-increasing and decreasing handpiece to complete most of the clinical therapy.

Brushless Electric Motor

Maximum speed : 40000±10%(RPM) Minimum speed : 2000±10%(RPM) Maximum torque : 3 Ncm Size : 50 mm Weight : 87g

Control Box

Class of insulation : I class BF Class of Protection : IP X class 1 (Protection against vertically falling drops of water) Size : 123mm X 63mm X 42mm

Operation Condition:

Temperature : 10°C - 40°C Relative humidity : 30% - 80%

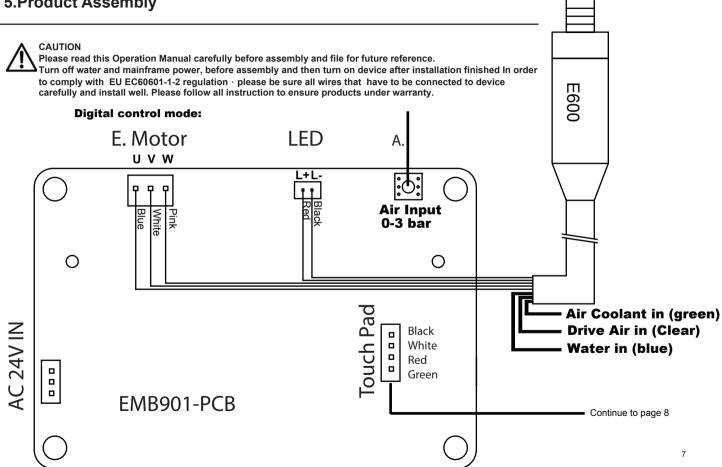
Transportation and storage condition: Temperature : -18°C - 70°C

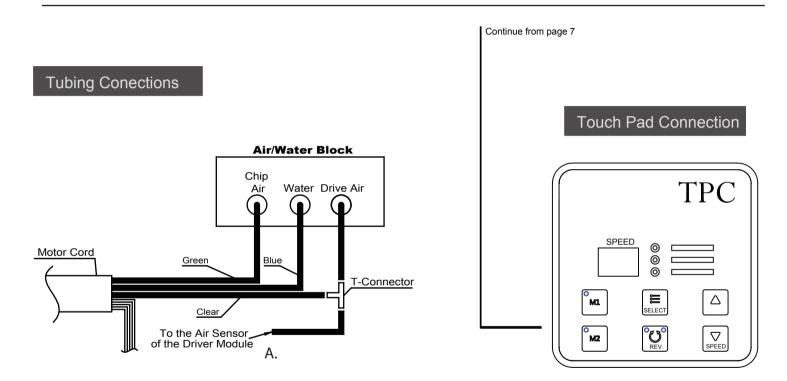
Relative humidity : 10% - 100% Atmospheric pressure : 500hPa - 1060hPa

Driver Module

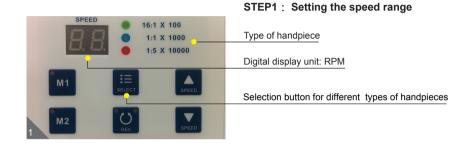
Input Voltage : AC 24V ±5% · 6.5A · 50/60 Hz 'Power : 130 W(Comply with IEC60601-1-2) 'Drive Air Pressure : 0 to 3 kg/cm² (0 to 3 bar)

5.Product Assembly





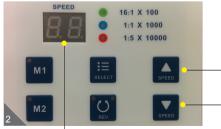
6.Operation Instructions



Handpiece Gear Ratio Indicator of SPEED Speed range (rpm) 16:1 01~25 100~2500 1:1 02~40 2000~40000 1:5 01~20 10000~200000

Push the SELECT switch to adjust the type of handpiece

* The SELECT button is functional only by the motor in static condition



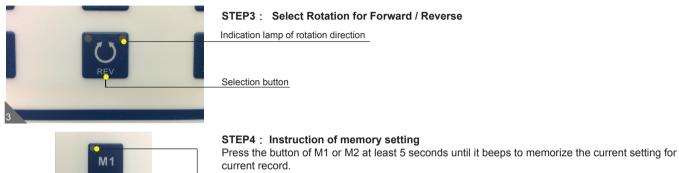
STEP2 : Speed Control Switches Set the speed of the motor.

The button for acceleration

The button for deceleration

Digital display unit: RPM

6.Operation Instructions



* The reaction triggers after push the button of select only when motor is still.

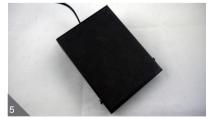
Light for showing the selected model

Button of Memory



STEP5 : Instruction of foot air pedal of delivery system

After all settings completed, step the foot air pedal to delivery the air into control box.



1.Brushless electric motor applied to dental handpieces in accordance to ISO3964-1982(E).

2.Be careful with handpiece hose avoiding any twist or break.

3.Water and air input shall comply with required quality standards.

4.After long-term use, please clean the control box by soft cloth to keep it free from pollution and surface erosion by chemical solvents.

5.Be cautioned the device is forbidden to run in the conditions of existing flammable anesthetics, oxygen or nitrous oxide mixture gases.

6.Motor should be kept dry, free of oil or any liquid internally or it will damage the motor and the control unit.

7. The device running may interfere with the cardiac pacemaker. If you have any questions with the patient having heart pacemaker, please consult cardiologists.

8. Please do not put off or touch power plug for fear of accidents or device breaking down.

Error Code

If the motor stops due to an breakdown such as a malfunction, overload, break or wrong use, it would automatically check the state of the control unit and detects the cause and displays an error code on the speed indicator. If an error code is displayed, please turn on the power again and check whether the same error code is displayed. If the same error code is displayed, please take action by referring to the instructions provided in the column of "Check/Remedy" in the following table.

Error Code	Trouble	Causes	Check/Remedy
EO	Overcurrent Error	Motor overload.	Restart. If ineffective, please contact your dealer.
E1	I Overneat Error	High temperature sate system is functioning, due to long-time usage at a high load.	Cool it down for a while, and try again.
E2	Inrush current Error	Detected inrush current.	Restart. If ineffective, please contact your dealer.
E3	Bulb voltage Error	Overvoltage is given from the internal circuit.	Restart. If ineffective, please contact your dealer.
E4	Motor start Error	 Motor didn't reach the preset speed in a prescribed period of time. Motor cord wire break down or internal circuit malfunction. 	Restart. If ineffective, please contact your dealer.
E5	Overvoltage Input Error	Overvoltage is given to the control box.	Restart. If ineffective, please contact your dealer.

Troubleshooting

If the motor stops due to an breakdown such as a malfunction, overload, break or wrong use, it would automatically check the state of the control unit and detects the cause and displays an error code on the speed indicator. If an error code is displayed, please turn on the power again and check whether the same error code is displayed. If the same error code is displayed, please take action by referring to the instructions provided in the column of "Check/Remedy" in the following table.

Error Code	Causes	Solution
	Power Switch is OFF.	Turn On Power Switch.
Pilot Lamp does not light	Power Plug is not connected correctly.	Check the connection.
	Internal Fuse is blown, due to some reason.	Contact your dealer.
	Tubing, Motor cord, driver module is not connected correctly.	Check the connection
Motor does not run	Air pressure is not given, or not properly from the delivery unit.	Check the air pressure of the delivery system.
	Check the ERROR CODE in the Speed indicator.	Refer to ERROR CODE.

Only allowable to use maintaining tools and parts provided by TPC Advanced Technology., LTD., If not it will void your warranty.

Overhaul

Do not disassemble the product, for all Adjustment and maintenance, it is recommended that you directly contact your dealers or TPC Advanced Technology. TPC requires users to overhaul the used equipment at least once a year.

Application

This product must to be used by the qualified personnel for dental treatment only.

Information

All technical specification, diagram and size in this manual are only for reference, and shall not be used as the basis for any claim. The manufacturer for compensation technical specification is subject to revise without notice. For other information please contact TPC Advanced Technology, mailing address can be found on the back of this manual.

Clean

Use a clean, moist cloth to disinfect.Do not press the light source.Do not use product which containing acetone, chlorine and bleach as a disinfectant.Do not immerse in the liquor.Do not use in the ultrasonic cleaner.

Sterilization

Autoclave sterilization is recommended. Autoclave sterilization is required for the first time use and after each patient as below notes.



CAUTION : Do not autoclave (or any other high temperature sterilization) Control Box, Driver module and Motor Cord.

General Principles

The product should only be performed by the qualified personnel in accordance with the existing relevant industrial safety, health and accident prevention, as well as the provisions of this job description. According to the above requirements, the operator:

"Use only perfect performance fault-free products: When the occurrence of abnormal operation, excessive vibration, overheating or other failure of signs, shall immediately stop working and contact TPC or its authorized repaired center.

Must ensure that the device only for its design purpose, and shall ensure that its own patients and other people's safety, the process of use should avoid pollution. The device can only be used for the purpose of dental treatment; prohibit this product from other purposes, otherwise it will be dangerous. The medical devices conform with the equirements of all existing EU law.



Do not use in the place which containing explosive gas (anesthetic gas).

Do not immerse in the disinfectants liquor.

This device is intended for recycling. Electrical and electric equipment may contain dangerous substances with harmful impact on health and the environment. The user is required to recover such equipment to distributor or recyclable such equipment directly to authorized personnel (EU Directive 2002/96/CE).

Warranty

TPC Advanced Technology Inc., warrants its products to the original purchaser against defects in material and workmanship under normal practices of installation use and servicing, The warranty time is 12 months from the invoice date of purchase. If claims proved, TPC Advanced Technology Inc., or its authorized agent has the obligations to provide free repair or replace of the product. Any other reasons for the claim, especially due to factors such as destruction or interests, are not covered under warranty. TPC Advanced Technology Inc., Is not responsible for damage or problem caused by the following factors

Excessive operation.

Incorrect operation.

Without following the manual instruction to install, operate, and maintain.

Influenced by the unusual chemical, electrical or electrolyte substance.

'Gas, water or electricity tube are mistakenly connected.

The "fiber optical" and any parts made from synthetic material are not under warranty, if the damage or the subsequent impact are caused due to incorrect operation or adjustment without permission from TPC Advanced Techlology Inc., the warranty term will be immediately invalidated. In order to meet the warranty and compensation term, any returned products must be indicated with the date of purchase, product type, invoices or shipping records.

Prevention measures for medical device and ElectroMagnetic Compatibility (EMC)

It must be in accordance with the user manual and the existing file of electromagnetic compatibility information for installation and using.

This product complies with IEC60601-1-2 of the EU on the provisions of the letter of the electromagnetic compatibility. It is forbidden to use radio transmitting equipment, mobile phones nearby the device, so as not to affect the performance of the instrument, specific precautions needs to be taken when using strong wave of radiation sources, such as high-frequency surgical equipment and similar equipment if example, is not available in the instrument winding high-frequency near the pipeline. If there is any question, please contact a qualified technician or TPC Advanced Technology Inc.

This product can not be closed to other equipment or stacked with other equipment. If it must be closed to or stacked using, please to set up and operate in a normal manner to the way they should be used.

Concerning replacement parts for internal components, accessories, transformers and hose fails to specify the manner or the use of TPC Advanced Technology Inc., the adapter and tubing will increase the amount of radiation waves or reduce the tolerance of this commodity.

Guidelines and manufacturer's declaration - electromagnetic emission waves

this product is intended for using in the electromagnetic environment.

Customers who use the product should ensure that operating environment meet standard condition.

Emissions test	Accordance	Electromagnetic environment standards	
RF Emissions CISPR11		This product which uses radio frequency energy for internal purposes only operation. Radio frequency (RF) radiation waves is quite low, and should not interfere with nearby electronic equipment.	
RF Emissions CISPR11	Class B		
Harmonic Emissions IEC61000-3-2		This product is applicable to all facilities, including household facilities, as well as directly connecte the public low-voltage power supply network, for general building household power supply facilities	
Voltage fluctuation / flicker radiation waveIEC61000-3-2	Section 5		

Guidelines and manufacturer's declaration - electromagnetic tolerance

This product can be used for the following specific electromagnetic environment. Customers or users of this product should ensure that operating in an appropriate environment.

Immunity test	IEC60601Test level	Compliance Level	Electromagnetic Environment - Guidance
Electrostatic discharge(ESD) IEC61000-4-2	±6KV · ground ±4KV · contact ±2KV · contact ±8KV · air ±4KV · air ±2KV · air		Floors must be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%.
Electrical fast / instant Burst IEC61000-4-4	± 2 KV \cdot Powered supply line ± 1 KV \cdot Applicable Input / out	put line	Main power quality should be at the same level with the location of the general commercial or hospital environment.
Surge wave IEC61000-4-5	±0.5KV · Tubing on tubing ±1KV · Tubing on tubing ±0.5KV · Tubing on ground ±1KV · Tubing on ground ±2KV · Tubing on ground		Main power quality should be at the same level with the location of the general commercial or hospital environment.
Voltage dips, short interruptions and voltage variations on the power supply input tubing IEC61000-4-11	< 5% UT(> 95% dips · at UT)Around 0.5 circles 40% UT(60% dips · at UT)Around 5 circles 70% UT(30% dips · at UT)Around 25ciercles < 5% UT(> 95% dips · at UT)Around 5 seconds		Power voltage should meet the level of the general public or hospital environment. The users of this product during the main power supply is interrupted continue operation of the proposed use of the supply of continuous electrical device- oriented products.
Power currency (50/60Hz) Magnetic filed IEC61000-4-8	3 A/m		Power frequency magnetic fields should be at the same level with the location of the general commercial or hospital environment.

12.Manufacturing company information and after sales service

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