

Contra-angle Handpieces Instruction For Use

1 Product introduction

1.1 Overview

The CA161, CA361-H, CA161-20, CA001-G, CA1471, CA221, CA161-RR, Contra- angle Handpieces and kinds of geared angle handpieces according to ISO 7785-2 and are used to hold and drive rotary instruments for the purposes of rotary processing, intended for the endodontic treatment. Features

a) Smooth transmission, low vibration, low noise and high cutting efficiency.

b) Long service life.

1.2 Technical specifications

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Model	CA161	СА361-Н	CA161-20	CA001-G	CA1471	CA221	CA161-RR	
Output	100~1200	100~1500	20~2500	100~1000	100~3300	100-1000	100-2500	
Speed Range	rpm	rpm	rpm	rpm	rpm	rpm	rpm	
Output	0.4~5.0	0.4~4.2	0.4~5.0	0.6~4.0	0.4~4.0	0.4~5.0	0.6~4.2	
Torque Range	Ncm	Ncm	Ncm	Ncm	Ncm	Ncm	Ncm	
Chuck type		Push-button						
Gear ratio	6:1	6:1	6:1	1:1	4.7:1	1.8:1	6:1	
Rotation mode	Continuous forward, continuous reverse and reciprocating rotation (Axia reciprocating rotation) (Axia reciprocating rotation) (Axia reciprocating rotation) (Axia reciprocating rotation)							
File shank attachment	Ф2.35mm ISO1797-1 Туре1							
Minimum fitting length of shank	11mm							
Maximum overall length of the rotary instrument	46mm							

1.3 Structure and compositions

The product consists of handle, driving medium and head.

1.4 Configuration

Please refer to packing list for more details of configuration.

1.5 Indications for use

The CA161, CA361-H, CA161-20, CA001-G, CA1471, CA221, CA161-RR Contra- angle Handpieces are medical devices and designed for holding and driving dental root canal instruments in continuous and reciprocating movement. These devices must only be used in hospital environments, clinics or dental offices by qualified dental personnel.

1.6 Operation conditions

- 1.6.1 Environment temperature:10°C~40°C
- 1.6.2 Relative humidity:30%~75%
- 1.6.3 Atmospheric pressure:70kPa~106kPa

1.7 Warnings

1.7.1 Please check the operation outside oral cavity before treatment. Pay full attention to check whether there is vibration, noise, or heating. If there is any unexpected problem, please stop operation.

1.7.2 Avoid strong impact and dropping;

1.7.3 Do not arbitrarily disassemble or modify the Contra- angle Handpieces; 1.7.4 Please prepare a backup Contra-angle Handpieces to cope with the case that Contra-angle Handpieces malfunction during operation;

1.7.5 Do not use non- recommended shanks/files. Otherwise, the running shanks/ files may bend or fly out;

1.7.6 Please keep the shanks/files handle clean. Otherwise, the stains on the surface of handle will affect the correct fixation of shanks/files, causing the shanks/files to fly out during operation;

1.7.7 Do not use the Contra-angle Handpieces without shanks/files.
 1.7.8 Do not press the button while operating. Otherwise, it may cause head

overheating or flying shanks/files;

1.7.9 For safety, please wear protective goggles or dust mask while cutting;

1.7.10 Do not exert too much force to treatment site;

1.7.11 Please carry out regular maintenance and inspection;

1.7.12 If the Contra-angle Handpieces has not been used for a period of time, before operation, please maintain it as per Clause 4 Maintenance. 1.7.13 Damage or serious wear will lead to increased transmission resistance,

abnormal sound and so on. As a result, the vibration of Contra-angle Handpieces handpiece will be strong; the drill will shake greatly; and there will be abnormal sound, etc. The operator should timely replace the Contra-angle Handpieces according to clinical conditions. If the Contra-angle Handpieces wears to a certain extent, please replace that with a new one

1.7.14 Împroper use will damage this product, and then hurt the patients, users and the third parties.

1.8 Contraindications and warnings

1.8.1 The hemophilia patient is forbidden to use this equipment.

1.8.2 The patients or doctors with heart pacemaker are forbidden to use this equipment.

1.8.3 Patients with allergic constitution and a history of drug allergy are forbidden to use this equipment;

1.8.4 Patients with oral and maxillofacial infections, various types of oral mucosal diseases that have not been cured, periapical disease, gingivitis, periodontal disease and oral cancer are forbidden to use this equipment.

1.8.5 Patients with heart disease and children should be cautious to use the

equipment.

1.8.6 People with mental disorders should be cautious to use this equipment; 1.8.7 Patients with severe systemic infection or systemic diseases such as diseases in heart, liver, kidney, hematopoietic system, digestive system and endocrine system should be cautious to use this equipment.

1.8.8 Pregnant women, lactating women, or women of childbearing age who have a plan of pregnancy should be cautious to use this equipment.

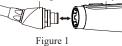
2 Operation instructions

2.1 Connecting and Disconnecting the Contra-angle Handpieces

2.1.1 Connecting

The Contra-angle Handpieces can be connected at 3 or 6 adjustable head positions. Adjust to a suitable angle during use. Align the positioning pins of the Contraangle Handpieces with the positioning slots of the motor handpiece and insert the head until it clicks (see Figure 1). When performing the APEX measurement, mount the insulating sleeve and pull it fully over the Contra-angle Handpieces. We recommend performing the treatment with a cofferdam and wear appropriate insulated gloves, while using an insulating sleeve. This will prevent inaccurate measurements caused by undesired leakage currents. During the measurement, the instrument must not come into contact with the patient's mucosa, metallic tooth restorations, or the mucosal electrode.

When removing the Contra-angle Handpieces, pull it straight out (see Figure 1).



2.1.2 Disconnecting

CAUTIONS

a) The Contra-angle Handpieces can be connected or disconnected only after the motor handpiece has completely stopped running. b) Connect only to the motor CA161 Contra-angle handpiece with a maximum

speed of less than 1,200 rpm.

c) Connect only to the motor CA001-G Contra-angle handpiece with a maximum speed of less than 1,000 rpm.

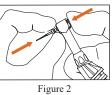
d) Check the Contra-angle Handpieces is securely assembled to the motor handpiece.

2.2 Inserting and Removing the shanks/files

2.2.1 Insertion See Figure 2.

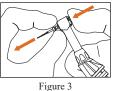
1) Lightly hold down the push-button and insert the shank/file.

2) Turn the shank/file back and forth until it engages with the latch mechanism. 3) Push the shank/file inwards to clickand release the push-button to lock the shanks/files.



2.2.2 Removal

See Figure 3. Press the push-button and pull out the shank/file.



a) When connecting and disconnecting the shank/file, turn the power off beforehand.

b) If it is necessary to use the insulating sleeve, first mount the insulating sleeve, then insert the shank/file. Remove the shank/file before removing insulating sleeve. c) After the shanks/files is locked, lightly pull out the shank/file to ensure that the shanks/files is locked.

d) Always clean the shank of the file to be installed. Allowing dirt to enter the chuck could cause deterioration of chucking force.

e) If the shank/file are not installed securely, it may fly out or be hard to be removed.

3 Cleaning, disinfection and sterilization

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Steps	Contra-angle handpiece
Warnings	1. The contra-angle handpieces are not suitable for ultrasonic cleaning. 2. The contra-angle handpiece should not be immersed in strong detergents or disinfectants (alkaline pH-9 or acid pH <5).
Resistance to sterilizing procedure	The contra-angle handpiece could be capable of being subjected to 600 cycles of sterilizing procedure.
Preparation at the Point of Use	The post-operative process must be carried out immediately, no later than 30 minutes after the completion of the operation. The steps are as follows: Remove the shanks/files and disconnect the contra-angle handpiece from Motor handpiece. Remove gross soiling of the instrument with cold water (<40°C) immediately after use. Don't use hot water (>40°C) as this can cause the fixation of residuals which may influence the result of the reprocessing process.
Transportati on	The contra-angle handpiece should be safety stored and transported to the point of reprocessing to avoid any damage and environment pollution.
Preparation for reprocessing	The products must be reprocessed in a disassembled state.
Pre-cleaning	Tools: tray, soft brush, clean and dry soft cloth Perform manual pre-cleaning until the handpice is visually clean. Rinse the chuck of bur with running water for at least 10 seconds. Clean the surface with a soft bristle brush. Note: The water temperature should not exceed 40°C during the washing stage, otherwise the protein will solidify and be difficult to remove.

Cleaning	Regarding cleaning/disinfection, rinsing and drying, it is to distinguish between manual and automated reprocessing methods, sepscially due to the better standardizing potential and industrial safety. Automatic cleaning The washer-disinfector should meet the requirements of the ISO 15883. Place the contra-angle handpiece in the washer-disinfector carefully. Ensure that contra-angle handpieces can not move freely in the washer-disinfector. The contra-angle handpiece are not permitted to contact with each other. Start the program: 4-min pre-washing with cold water(<40°C); 5-min washing with a mild alkaline cleaner at 55°C; 5- Emptying 5- S min meutralizing with warm water(>40°C); 5- Emptying 5- S min intermediate rinsing with warm water(>40°C); 5- Emptying 5- S min intermediate rinsing with warm water(>40°C); 6- Emptying 6- Drying the device at 80°C for 15min The automated cleaning processes have been validated by using 0.5% neodisher MediClean forte(Dr. Weigert)				
Disinfection	Automated Thermal Disinfection in washer/disinfector under consideration of national requirements in regards to A0 value (see EN 15883). A disinfection cycle of 5 min disinfection at 93°C has been validated for the device to achieve an A0 value of 3000.				
Drying	Drying of outside of instrument through drying cycle of washer-disinfector. If necessary, additional manual drying can be performed through lint free towel. Insufflate cavities of instruments by using sterile compressed air. If your washer-disinfector does not have an automatic drying function, please dry the device after cleaning and disinfection. The drying method as below: 1)Spread a clean white paper (white cloth) on the flat table, place the contra-angle handpiece on the white paper (white cloth), and then dry the contra-angle with filtered dry compressed air (maximum pressure 3 bar). When no liquid is sprayed on the white paper (white cloth), it indicates that the contra-angle handpiece is completely dry. 2)The device can also be dried directly in a medical drying cabinet (or oven). The recommended drying temperature is 80 °C and the time should be 15minutes. Note: 1)Dry the contra-angle handpiece repeatedly if necessary (refer to section "Drying"). 2)The device should be dried in a clean area. 5)The drying temperature should not exceed 137 °C.				
Maintenance	 Functional Test and Visual inspection Visually inspect the cleanfiness of the Handpiece. Perform Functional test according to instructions of use. If there is still visible stain on the device after cleaning, the entire cleaning process must be repeated. Before packaging and sterilization, make sure that the contra-angle handpiece has been maintained acc. to manufacturer's instruction. If the device is obviously damaged, smashed, detached, corroded or bent, it must be discarded and not allowed to continue to be used. If the accessories are found to be damaged, please replace it before use. And the new accessories for replacement must be cleaned, disinfected and dried. Use lubricant to lubricate the handpiece and dried it prior to sterilization. Aim the nozzle of lubricant bottle to the air hole at the end of the contra-angle handpice to inject oil for 1-2 seconds. 				
Packaging	The contra-angle handpiece should be quickly packaged in a medical sterilization bag (or special holder, sterile box). Precautions 1) Only use a legally marketed or a FDA cleared sterilization pouch; 2) The package should withstand high temperature of 137 °C and has sufficient stean permeability; 3) The packaging environment and related tools must be cleaned regularly to ensure cleanliness and prevent the introduction of contaminants; 4) Avoid contact with different metals when packaging.				
Sterilization	Sterilization of instruments by applying a fractionated pre-vacuum steam sterilization process (according to EN 285/EN 13060/EN ISO 17665) under consideration of the respective country requirements. Minimum requirements: at least 4 min at 132°C/134 °C (in EU: 5 min at 134 °C, in US: 4 min at 132 °C) Maximum sterilization temperature: 137°C Flash sterilization is not allowed on lumen instruments!				
Storage	Sterilized devices should be stored in a dry, clean and dust-free environment, refer to label and instructions for use.				
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5 Storage and transport conditions 5.1 Storage and transport conditions

- 5.1.1 Environment temperature: -20°C ~ +55°C 5.1.2 Relative humidity:10% ~ 93% 5.1.3 Atmospheric pressure:70kPa ~ 106kPa

- 5.2 This product should be placed or stored in a dry and clean environment which is away from acid, alkali and other harmful chemicals and gases.

6 Environment protection

This product contains no harmful ingredients and can be disposed and destroyed according to local regulations.

7 After-sales service

After the sale of the product, if there is any quality problem and the device cannot work normally during warranty period, Woodpecker is responsible for the maintenance.

Please refer to warranty card for the warranty period and other specific items. This product is aprecision equipment. If there is problem that needs to berepaired, returned to Woodpecker or handled by professionals is recommended.

8 Symbol description



Follow instructions for use

	Date of manufacture	
	Manufacturer	
†	Type B applied part	
IPX0	Ordinary equipment	
	Used indoor only	
X	Appliance compliance WEEE directive	
-20°C +55°C	Temperature limitation for storage	
70kPa	Atmospheric pressure for storage	
10%	Humidity limitation for storage	
Ť	Keep dry	
Ţ	Handle with care	
•	Recovery	
134°C 5555	Can be autoclaved	
SN	Serial number	
CE 0197	CE marked product	
ECREP	Authorised Representative in the EUROPEAN COMMUNITY	

9 Statement

The Figures are only for reference. The final interpretation rights belong to Guilin Woodpecker Medical Instrument Co., Ltd. (Please refer to product packaging label for date of manufacture.Service life of

product: ten years) **10.Contact information**



Guilin Woodpecker Medical Instrument Co., Ltd. Information Industrial Park, Guilin National High-Tech Zone, Guilin, Guangxi, 541004 P. R. China Sales Dept.: +86-773-5873196 Http://www.glwoodpecker.com E-mail: woodpecker@glwoodpecker.com ECREP MedNet EC-Rep GmbH Borkstrasse 10 · 48163 Muenster · Germany

ZMN-SM-025 V1.6-20220511