

DIRECTIONS FOR USE

EDGEV-TAPER™ and EDGEV-TAPER™ GLIDEPATH Heat Treated Fire-Wire™

Intended Use

Endodontic files and reamers are single use surgical instruments used for performing root canal treatment to mechanically shape and prepare the root canals during endodontic therapy or to remove the root canal obturating material when performing retreatment. The device is intended to be used sterile and single use only.

Intended Users

The device is designed to be used by a Dental or Endodontic specialist trained in endodontic techniques. No additional training is required for the safe use of the device by the treating physician.

Intended Patient Population

Adolescent to adult population. People with permanent teeth in need of endodontic pulpectomy.

COMPOSITION

The instrument is made of a nickel-titanium blade, handle, the stop, and the color-coded band.

Contraindications

- Mechanically driven endodontic instruments should not be used in cases with very severe and sudden curvatures.
- This product contains nickel and should not be used for individuals with known allergic sensitivity to this metal.

Warnings

- Endodontic files are for single use only, in order to avoid file separation.
- The product has not been designed or tested for reuse. The ability to effectively clean and re-sterilize this single use device and subsequent reuse may adversely affect the clinical performance, safety and/or sterility of the device.
- Endodontic files are sharp, and caution should be used if touching the blade directly.
- After use, this product may be a potential biohazard. Handle and dispose of in accordance with accepted medical practice and applicable local, state, and federal laws and regulations.
- If the packaging is damaged, please dispose of the damaged product and utilize an undamaged product instead as the former may be contaminated.

Precautions for Use

As with all products, use carefully until you become proficient with use. Always determine working length using radiographs and/or apex locator to properly use endodontic files. Important points to remember:

1. A rubber dam system should be used.
2. Use only in an electric motor and hand piece designed for endodontic (rotary/reciprocating) files.
3. Straight-line access is imperative for proper file use and endodontic treatment.
4. Do not force the files down canals, use minimal apical pressure.

5. Clean the flutes frequently and at least after removing the files from the canal.
6. Irrigate and lubricate the canal frequently throughout the procedure.
7. Take each file to length only one time and for no more than one second.
8. In apical areas and curved canals exercise caution.
9. Once file is used do not reuse. If file is reused and used on a different patient infection can be introduced. Performance of the file can also be reduced.
10. When instrumenting the canal, do not over enlarge the coronal portion of the canal.
11. Too large a file taken to length increases the risk of canal transportation and file separation.
12. Do not exceed the handpiece recommended maximum torque or speed. Exceeding settings may cause the device to fail.
13. Endodontic files undergo our proprietary Annealed Heat Treatment (AHT) forming our branded **Fire-Wire™** NiTi which increases cyclic fatigue resistance and torque strength. With this proprietary processing, the files may be slightly curved. This is not a manufacturing defect. While the file can be easily straightened with your fingers, it is not necessary as once they are inside the canal, endodontic files will follow and conform to the natural canal anatomy and curvatures.
14. Do not use after the expiration date on the label.

Adverse Reactions

- Device fracture/breakage
- Complications usually associated with endodontic procedures including:
 - Pain
 - Instrument fracture/breakage
 - Soft tissue damage/bleeding
- Infection – Do not use if package is damaged or open, due to risk of infection occurring.

Safe Unwinding

- As a safety feature the files are designed to unwind. They may be used until the files unwind backwards.

INSTRUCTIONS FOR USE

Sterilisation

No sterilisation steps are needed for product provided sterile.

Step-By-Step Instructions

1. Create straight line access to all canal orifices.
2. Irrigate and fill the chamber with **EDGELUBE™** 17% Liquid EDTA and establish the Glide Path using small hand files. Determine the Working Length and verify canal patency.
3. Always irrigate after each file with EdgeLube and if needed, expand the GlidePath using **EDGEV-TAPER™ GLIDEPATH** NiTi files 13/03 and/or 14/03.
4. With the canal filled with EdgeLube and following the **EDGEV-TAPER™ GLIDEPATH** NiTi files, use **EDGEV-TAPER™ HT** NiTi 17/04. Advance the file 2mm further down the canal each time using multiple inward-strokes with moderate pressure while brushing the sides of the canal on the out-stroke until the working length is reached.

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5. Next use the **EDGEV-TAPER™ HT** NiTi 20/06 as described in Step 4 until the working length is passively reached.
6. If the apical flutes of the 20/06 is loaded with dentin, the canal may be fully shaped and ready to be Apically Gauged.
7. Apically Gauge the apical foramen of the canal foramen with a #20 size hand file. If the file reaches the working length and has a snug fit, the canal is shaped and ready for disinfection.
8. If the #20 handfile is loose at the Working Length, then continue shaping with **EDGEV-TAPER™ HT** NiTi 25/06 and if needed, the larger **EDGEV-TAPER™ HT** NiTi files gauging after each instrument with a #30, #35, #40, #45 or #50 hand files respectively. Irrigate and recapitulate with a small-sized hand file after each **EDGEV-TAPER™ HT** instrument.

Motor Settings

- Speed: 300-500 RPM
- Torque: 4.0 - 5.2 Ncm/ 408-530 gcm

Disinfecting

- After each canal is fully shaped, rinse the canals for 1 minute with 17% Liquid EDTA to remove the canal Smear Layer.
- Rinse the canals for 5 minutes with 5% NaOCl to remove debris and bacteria.
- Rinse the canals for 1 minute with 17% Liquid EDTA to rinse out the 5% NaOCl.
- Rinse the canals for 5 minutes with 2% chlorohexidine or EDTA to kill bacteria.

Obturation of Canal Systems

- When using thermal carrier system use size verifiers to determine the proper sized carrier.
- When using a master gutta percha cone that matches the largest file taken to length, remember sometimes you may need to drop down in cone tip size if the corresponding gutta percha to your final rotary file does not go to length.



















Disposal

- Recommended file disposal: Place used files in Biohazard Sharps container.

Reporting of Incidents to manufacturer and competent authorities

- In case any patient/user faces a serious incident, the entirety of the incident will be reported to the following:
 - The manufacturer of the device: US ENDODONTICS
 - The competent authority of the country where the user/patient resides

Symbol Table

Symbol	Meaning (Standard, if Applicable)
	Manufacturer: Indicates the medical device manufacturer (ISO 15223-1)
	Authorized Representative: Indicates the AR in the EU
	Importer: Indicates the entity importing the medical device into the locale (ISO 15223-1)
	Conformité Européene. EU mandatory conformity marking.
	Translation: Indicates that the original information has been translated and replaced (ISO 15223-1)
	Medical Device: Indicates the item is a medical device (ISO 15223-1)
	Catalogue number: indicates the Medical Device SKU (ISO 15223-1)
	Batch Code: Manufacturer's batch code so batch or lot can be identified (ISO 15223-1)
	Indicates the date after which the medical device should not be used (ISO 15223-1)
	Do not reuse: Indicates a medical device that is intended for one single use only (ISO 15223-1)
	Do not resterilise: Indicates medical devices that is not to be resterilised (ISO 15223-1)
	Medical device sterilized using irradiation and packaged with a single outer sterile barrier system (ISO 15223-1)
	Medical device should not be used if package is damaged and consult instructions for use (ISO 15223-1)
	Consult IFU: consult the Instructions For Use, and eIFU website listed (ISO 15223-1)
	Caution is necessary when operating device. Align cautions (ISO 15223-1)
	Unique Device Identifier: Indicates a carrier that contains UDI information (ISO 15223-1)
	Date and Country of Manufacture: To identify the country of manufacture of products next to date of manufacture (ISO 15223-1)
	Prescription Use Only: Caution: Federal law restricts this device to sale by or on the order of a dentist (21CFR 801.109)