# NT2





endo★star



# Instruction for use

## Endostar NT2 NiTi Two Rotary System

In Important information regarding the system
 Endostar NT2 NTI Two Rotary System is an economical nickel-titanium file system for simple and fast mechanical
 processing of root canals, mainly with the traditional method. The system consists of 6 files. It is characterized
 by a constant. C2 uper and a non-cutting tip. The set is most commonly used as an extension of the Endostar E3
 Rotary System.

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- A handpiece, which can provide 150-300 rpm, should be used The operating speed of the handpi be constant throughout the shaping process. Do not apply excessive force. An up-and-down motion should be used when operating the files. Shaping time should be as short as possible. Always use a lubricating agent when shaping the canal. The files are very sharp and should be used very carefully, with little force and without excessiv down the canal. sive "pu The files are very sharp and should be used very caefully, with little force down the canal. Operate the instruments and handpiezes according to their operating instri speed settings). Use the type and number of instruments that is truly needed in a given of Control the amount of times that the instrument was used. Before using the instruments, be sure to see them working outside the ore and/or racks. Dispose of as medical waste. ins (especia lly to
- ded in a given clinical situation.
- > > ity to check

# Recommended movements Rotary movement - the instruction

- ily 360° in a clo n (CW - Clock W 3. Recommended torque settings
- File number Torque (Ncm) File number Torque (Ncm) 1 (02/15) 0.3 2 (02/20) 0.3 3 (02/25) 0.3-0.4 4 (02/30) 5 (02/35) 6 (02/40) 0.4-0.5 0.5-0.6 0.6-0.7

The torque settings indicated in the table above are only suggestions and may vary according to each user preferences and endodontic motor capabilities. Do not excreed the upper torque limit which is different for each instrument. If precise torque settings cannot be set, and only manufacturer-specific torque levels are available, be sure to select one that does not excred the recommended limit.

Recommended number of usage winnum of 5 times, provided that visual inspection performed by the pactitioner prior to use shows that the trument remains undamaged, is not bent, deformed, does not show signs of blade wear and can be secuely tached to the handpiece. If the file has been subjected to high torsion foce, especially in highly curved cana instrument should be used only once.

> Prolonging the life of the instrument more the > Dispose the file which appear to be defective

cal instruction for use

Rinse the canal each time after the file is used. Clean the files of any debris frequently.

- A. Prepare the cavity. Use a nubber dam.
  B. Locate all canal orffices. Fill the canal orffice with a lubricant.
  C. Specify the working length of the canal orffice with a lubricant.
  C. Specify the working length of the canal with your method of choile.
  Make sure all canals are patent up to a depth of 2-3 mm from the apex with the help of the K15 file.
  For very curved and narrow canals, use a different hand instrument, size 06, 08, or 10.
  E. Create access to the canal orffice using rotary files with a greater taper (06, 08) or with the use of Ge
  Gidden drils.
  F. Start working with the 02/15 or 02/20 file until you each full working length
- working with the 02/15 or 02/20 file until you mach full working length and then switch to a ments (02/25, 02/30, etc) until you mach the desired canal size. instru

Warnings is product is for profes nal dental use onlu

 Cleaning and disinfection
 Detailed instructions for cleaning, disinfection an and www.endostar.eu in the download tab. ilization can be fo

Sterlization is is a non-sterile product. Sterlize before use. The instru-ments can be sterilized in a steam sterilizer stockave) at 13/4°C. Recommended sterilization time: 3 minutes at 2.1 bar overpessure. Instruments can be infected with mild disinfectants and washed in ultasonic cleaners.

## 9. In

Storage struments

### Product claims

a roooccurrent with the distributor and manufacturer of any claims or adverse events which occurred as a result of perating this device. Each <u>serious</u> incident connected with this product should be reported to the manufacture d the competent authority of the Member State in which the user is established.

Cross-section	CE mark and	MD Medical device	134°C	Non-sterile	Used for root
LOT	identification number of the notified body	<b>NITI</b>	autoclave at 134°C	REF	canal preparation
Serial number Lot number	Consult instruction for use	Nickel-titanium	Date of manufacture	Catalogue numer	Packaging unit

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