

Be sure to read this instruction manual before using this product.  
Keep this manual available for reference when needed.

## Instructions for Use

### Trade Name: Spring Scalp Hook Retractors Ti

#### Warning

Please read these instructions carefully prior to using this product. Product should be used according to these instructions and pay close attention to the safety of patients. Not doing so may give rise to serious problems or adverse events.

#### For the US market

Do not reuse the device when it is used on a patient with or suspected of having Creutzfeldt-Jakob Disease (CJD) or variant CJD (vCJD).

#### For the market outside the US

When the device is used on a patient with or suspected of having CJD or vCJD, make sure to adhere to the most recent and updated restrictions available in each country and/or region for its reuse. Be cautious of the possibility of secondary infection. Refer to [www.a-k-i.org](http://www.a-k-i.org) or AAMI Standard ST79 for more information related to cleaning and sterilization.

#### Contraindication / Prohibition

1. Use for intended purpose only  
Use devices for their intended purposes only. Incorrect use could cause this product to break.
2. Prohibition of use of chemicals  
Avoid exposing this product to chemicals. Doing so could damage this product due to corrosion.
3. Prohibition of secondary processing of this product  
Do not apply any secondary processing to this product. For example, do not apply impacts or vibration markings to the surface of this product. Doing so could break this product.
4. Handle with care  
Handle this product with care, as it can be deformed or damaged. Rough handling could significantly reduce the service life of devices and appliances.
5. Prohibition of use of polishing powder and wire wool  
When cleaning this product, do not attempt to polish its surfaces with rough polishing powder or wire wool. This could cause scratches on the surface of this product and result in rust or corrosion.
6. Prohibition of use of alkaline, acid, and household detergents  
Use only medical neutral detergents (pH 6 to 8) to clean this product. Do not use any alkaline, acid, or household detergent. Washing this product with an improper detergent could result in discoloration or corrosion.
7. Do not use hydrogen peroxide gas plasma sterilization  
This product is not suitable for sterilization with hydrogen peroxide low temperature gas plasma. It may discolor the surface of the product or affect the feature of the product.

#### Specifications



Code No.	Product Description
07-954-00Ti	Spring Scalp Hook Retractors, Ti

Material: stainless steel, titanium alloy

#### Intended Purpose

This product is a surgical instrument intended to withdraw an incised scalp to expose the skull during a surgical procedure.

#### Intended user

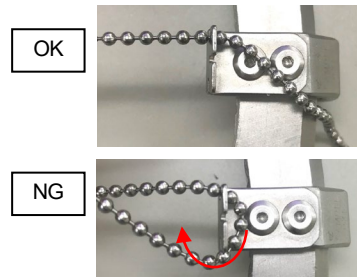
This product is to be used by health care professionals, including but not limited to surgeons, nurses and biomedical technicians.

#### Instructions for Use

Before using this product, inspect, wash, and sterilize in accordance with these instructions.

#### Warning/Caution

1. Important fundamental cautions
  - Prior to use the device must be sterilized under the standard sterilization conditions or the validated sterilization conditions for which validity has been proven by medical organizations in each country or region.
  - When using the scalp traction fish hook, do not wrap or fix the ball chain around the spring hook adjuster attachment while applying tension [The ball chain may break because of the bending load].



2. Defect/Adverse event

- Defect
- Corrosion and pitting caused by use of chemicals
  - Damage or breakage caused by the corrosion or pitting
- Adverse event
- Broken pieces of metal from the damaged instrument falling into the patient.

#### Storage/Life

1. Please store the device in normal ambient temperature areas. Do not store in areas of high humidity where the temperature may dramatically vary causing condensation. Do not store on or near chemicals as the chemicals may cause damage to the device.
2. Service life of this product: 1 year  
(Subject to the specified maintenance, inspection and proper storage)

#### Maintenance / Inspection

1. Check prior to each use  
Operational and functional checks  
Conduct daily and pre-operation checks of this product to make sure that it functions properly.
2. Check after each use
  - 2.1 Immediately wash with clean water  
If exposed to bleach or antiseptic solutions, immediately wash: Wash and rinse with clean water immediately and immerse in neutral enzyme detergent to remove any bleach or antiseptic solution, which may contain chlorine or iodine and can damage the instrument. Manually remove contaminated matter by hand or with an ultrasonic-cleaner.
  - 2.2 Further remove any remaining contamination with a plastic brush.

- 2.3 Select a proper detergent for each decontamination method and maintain appropriate density and handling.
- 2.4 Use a soft towel, a plastic brush, or a water gun for cleaning.
- 2.5 To avoid damage, do not use a metal brush, coarse polishing agents, or apply excessive force when handling the device.
- 2.6 Reverse osmosis water is recommended to wash this product.
- 2.7 Only use reverse osmosis water for the final rinse.  
It is recommended to use a washer-disinfector for this device. Thermal Disinfection can be used by following the manufacturer's defined parameters.:  
Thermal Disinfection Band: 90-93 °C/194.0-199.4°F, 5-10 minutes (A0 value: 3000-12000) (reference EN ISO 15883-1)
- 2.8 Fully dry this product immediately after washing it.  
Do not leave it wet for a longer time than necessary as residual water may damage the instrument.
- 2.9 Use distilled water or reverse osmosis water at least  
Use distilled water or reverse osmosis water to wash and sterilize this product at least. Residual chlorine and organic matters in tap water may cause stainings and/or rust and may damage the instrument.
- 2.10 Use a water-based anticorrosive lubricant  
Lubricating oil is completely removed by washing. After washing this product, apply a water-based anticorrosive lubricant prior to sterilization.

3. Sterilization

Device must be sterilized by users in accordance with validated sterilization procedures, such as an autoclave, that are regulated by medical organizations in each country or region.

Sterilization of the device may be accomplished by steam. The recommended sterilization parameters are as follows,

ISO/TS17665-2	
Temp.	Minimum exposure time
121°C / 250°F*	15 Min
126°C / 259°F*	10 Min
134°C / 273°F*	3 Min

\*According to the Guideline for Disinfection and Sterilization in Healthcare Facilities, 2008 (Update: May 2019)

**For the US market**

FDA has not approved or cleared medical devices, including sterilizers, for the intended use of reducing the infectivity of TSE agents (i.e., prions).

**For the market outside the US**

When the device is used on a patient with or suspected of having CJD or vCJD, make sure to adhere to the most recent and updated restrictions available in each country and/or region for its reuse.

**Maintenance and check by agents**

For safety use of this instrument, conduct periodic checks by the manufacturer or the agent recognized by the manufacturer. Maintenance and check by other agents could cause the adverse events and the decrease of the performance and the function. To schedule the periodic check, contact your local distributor or the manufacturer.

**Packing**

1 piece per pack

**Warranty**

MIZUHO Corporation will repair defective parts of this product without charge for one year from the date of delivery/installment except for cases of damage caused by a third party's repair, act of nature, improper use or damage on purpose. All other warranty terms and conditions are subject to regulations of MIZUHO Corporation.

**Disposal**

This device must be disposed of in accordance with local regulations.

Please contact your local distributor for proper disposal.

**Notice**

Any serious incident that has occurred in relation to the device should be reported to the manufacturer and the competent authority in which the user and/or patient is established.



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