

**Screwdriver** REF 6003-100-100  
**Universal Joint Screwdriver** REF 6003-100-110

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## Instructions For Use

### Intended Use

Screwdriver and Universal Joint Screwdriver are intended for adjustment and tightening/loosening operations of Stryker Navigation surgical instruments such as, but not limited to, the Pivotal Resection Guide (REF 6003-100-000), OrthoLock (REF 6007-003-000), the NavLock (REF 6000-999-002/-003/-004/-005), the Nav MIS Jig (REF 6003-200-010/-020/-030), and any other Stryker Navigation instruments which provide a compatible interface (refer to the respective instruction for use).

### Contraindications

None known.

### User Group

Healthcare professionals (surgeon/resident, nurse/professional caregiver) educated in computer-assisted surgery and thoroughly familiar with the instructions for use and with the operation of this product.

To request an additional in-service instruction, contact Stryker.

### WARNING

- Clean and sterilize the instrument before first and every use.
- Prior to each use, inspect the instrument for loose components and damage. Do not use if these conditions exist.
- Use only Stryker-approved products unless otherwise specified.
- For information related to compatible software applications, refer to the user manual of the software application

**NOTE:** The user and/or patient should report any serious product-related incident to both the manufacturer and the national competent authority where the user and/or patient is established.



### Cleaning, Disinfection, Sterilization and Inspection Instructions

Cleaning Group: IV

Refer to the Guide for Cleaning, Disinfection and Steam-Based Sterilization (TD6000005750) for cleaning safety and caution notes, cleaning equipment and detailed cleaning and inspection instructions.

### Disposal

Products that have been in contact with material of human origin may be infectious. Dispose of with the necessary precautionary measures in accordance with local regulations. Ensure infected products are decontaminated prior to recycling.

	<b>Medical device:</b> Indicates a medical device according to European Union harmonization legislation.
	Indicates that a device is in conformity with the applicable requirements set out in applicable European Union harmonization legislation providing for its affixing.

DE

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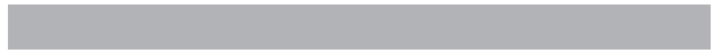
KO



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ZHS



JA



RO



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VI

