

CORDIS PRECISE® PRO TRANSCAROTID STENT SYSTEM IMPLANT CARD

Cordis **PRECISE**® stents are MR Conditional.

Patients with **PRECISE** stents should consult with their healthcare provider and inform MRI site personnel prior to an MR exam that they have an MR Conditional medical device.

A patient with the **PRECISE** stent may be safely scanned under the following conditions. Failure to follow these conditions may result in injury to the patient.

- Static magnetic field of 1.5 T and 3.0 T
- Maximum spatial field gradient of 40 T/m (4000 Gauss/cm) or less
- Circularly polarized RF excitation
- Whole body and head transmit-receive coils, and any receive only coil may be used
- Maximum whole body SAR of 2.0 W/kg
- 15 minutes of continuous RF followed by a wait time of 10 minutes if this limit is reached
- The presence of this implant produced an image artifact of approximately 16 mm when imaged with a gradient echo pulse sequence and a 3.0 T MRI system.
- Non-clinical testing has demonstrated that the **PRECISE** Stent is MR Conditional in single and overlapped configuration up to a maximum of 60 mm as defined in ASTM F2503-13.

Persons with allergic reaction to nickel titanium (Nitinol) may suffer an allergic response to this implant. Maximum Stent Composition: Nickel 57.0% weight/weight (w/w), Titanium 45.5% w/w



MR Conditional

Cordis™

Patient's Name

Date of Implant

Site of Implant

Implanting Physician

Hospital

Address

Telephone No.

Primary Physician

100641754.1

INSTRUCTION

Please carry this card at all times and show it to any medical personnel who may be treating you.

Expected lifetime of the device: 10 years.

ATTACH LABEL



Medical
Device

Cordis US Corp.
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Miami Lakes, Florida 33014, USA
1-800-327-7714 (U.S.A.)
www.cordis.com

REF

Catalogue
number

LOT

Lot
number