

# **AERO**mini<sup>®</sup>

IMPLANTED DEVICE PATIENT INFORMATION CARD



 **MERTMEDICAL**  
**ENDOTEK<sup>®</sup>**

# **AERO***mini*<sup>®</sup>

## Fully Covered Tracheobronchial Stent

Patient Name: \_\_\_\_\_ Stent Lot #: \_\_\_\_\_

Date of Implant: \_\_\_\_\_ Stent Size: \_\_\_\_\_

Implant Location: \_\_\_\_\_

Implanting Physician: \_\_\_\_\_

Hospital: \_\_\_\_\_ Contact Number: \_\_\_\_\_



## MR Conditional

### MRI Safety Information

Non-clinical testing has demonstrated that the AEROMini is MR Conditional. A patient with this device can be safely scanned in an MR system meeting the following conditions:

- Static magnetic field of 1.5T and 3.0T only
- Maximum spatial gradient magnetic field of 3,000 Gauss/cm (30 T/m) or less
- Maximum MR system reported, whole body averaged specific absorption rate (SAR) of 2 W/kg (Normal Operating Mode)

Under the scan condition defined above, the AEROMini is expected to produce a maximum temperature rise of 2.0°C after 15 minutes of continuous scanning.

In non-clinical testing, the image artifact caused by the device extends approximately 5 mm from the AEROMini when imaged with a gradient echo pulse sequence and a 3Tesla MRI system. The artifact does obscure the device lumen.



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EXPANDING THE POSSIBILITIES™

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