


INTERNATIONAL IMPLANT CARD

General Information


The HumanTech Spine International Implant Card must be completed by the healthcare professional with the required information and given to the patient. This document describes the content of the HumanTech Spine International Implant Card and instructs the healthcare provider on how to complete it.


The HumanTech Spine International Implant Card


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International Implant Card

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 www.humantech-spine.de



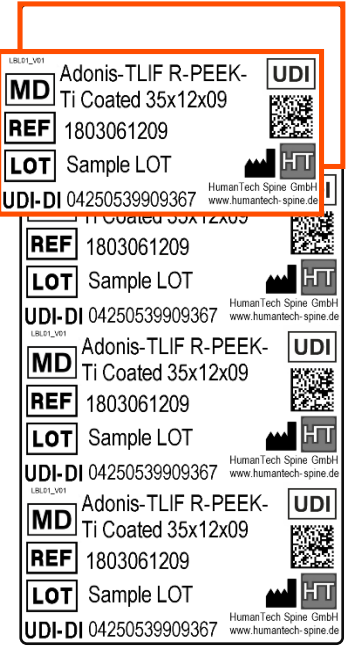
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MD	4
REF	5
LOT	6
UDI	7

MD	
REF	
LOT	
UDI	

 HumanTech Spine GmbH
Gewerbestraße 5
71144 Steinenbronn
Germany

Patient labels on or inside the product packaging



Instruction for completion: Fill in...

- 1 Name and surname of the patient (or patient ID)
- 2 Date of implantation
- 3 Name and address of the healthcare institution/ provider that performed the implantation
- 4 Name of the implanted medical device
- 5 Catalogue number of the manufacturer
- 6 Lot number of the manufacturer
- 7 Unique Device Identifier












For sterile products: Use our sticker: You can find the patient label either inside or on the back of the product packaging, as the design may vary depending on the date of manufacture.

Rev. date: 2025-06-23

Rev. No.: 01

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Explanation of the symbols

	Patient Name or patient ID
	Date of Implantation
	Name and Address of the implanting healthcare institution/ provider
	Information website for patients
	Name and Address of the manufacturer
	MR conditional (device may safely enter the MRI scanner room only under the very specific conditions provided in the labeling)
	Medical Device Name
	Catalogue Number
	Lot Number
	Unique Device Identifier
	Unique Device Identifier - Device Identifier

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