

“United” Implant Card Instructions for HCP

Introduction

- Based on the EU 2017/745 regulatory requirements, the implant device should provide the Implant Card to the patient.
- Healthcare Professionals filling out the “United” Implant card following the Instructions.
- Implant Card should be obtained from your “United” distributor.



1 Filling the Date of Implantation

2 Filling the Name or ID of the Patient

3 Filling the Name and Address of Healthcare Institute

4 Paste one “Implant card label” from each product packaging

(EN)Knee implant/ (FR)prothèse de genou/ (ES) implante de rodilla/
(IT)impianto di ginocchio/ (PT) implante de joelho

MD USTAR II UDI-DI:04711605584487
REF 2715-7314
LOT XXXXXXXX
UDI 

Manufacturer **MADE IN TAIWAN**
United Orthopedic Corporation
 No. 57, Park Ave. 2, Science Park
 Hsinchu, Taiwan

UNITED ORTHOPEDIC®

<https://www.unitedorthopedic.com> 

UNITED ORTHOPEDIC®

- EN** Implant Card
- FI** Implanttikortti
- FR** Carte d'implant
- EL** Κάρτα εμφυτεύματος
- IT** Scheda implantare
- PT** Cartão de Implante
- ES** Tarjeta de implante

Symbol glossary



Patient Name or patient ID



Date of implantation



Name and Address of the implanting healthcare institution/provider



Catalogue number



Name and Address of the manufacturer



Information website for patients



Lot Number / Batch Code



Unique Device Identifier



Device Name