

# Patient Implant Card

# Geistlich

Pharma

 Manufacturer  
 Geistlich Pharma AG  
 Bahnhofstrasse 40  
 6110 Wolhusen  
 Switzerland  
 Tel.: +41 41 492 55 55  
 info@geistlich.com  
 www.geistlich-pharma.com

ENGLISH

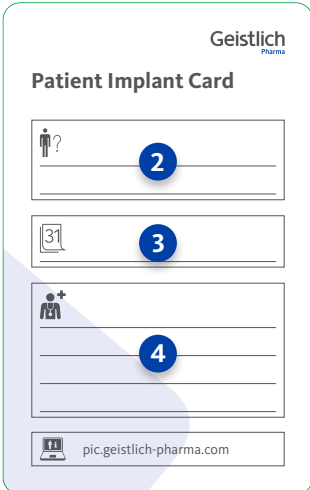
## Background

The Patient Implant Card is required by the new European Medical Device Regulation. In accordance with this regulation, Geistlich Pharma AG as legal manufacturer provides the Patient Implant Card together with its implantable devices.

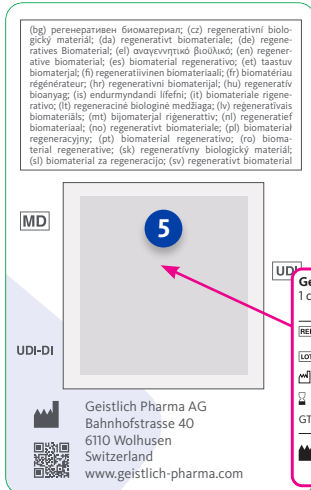
The Patient Implant Card must be filled with the required information by the healthcare institution/provider and handed to the patient. This document describes the content of the Patient Implant Card and instructs the healthcare provider on how to fill it out.

## The Geistlich Patient Implant Card

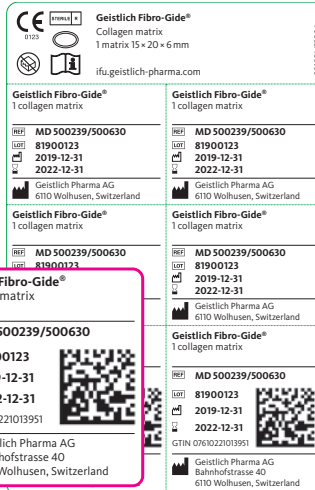
*Front*



*Back*















*Patient labels on the sterile packaging*



## Instructions

- 1** Use the language spoken by the patient to fill out the Patient Implant Card.
- 2** Fill in the name and surname of the patient (or patient ID).
- 3** Fill in the date of implantation.
- 4** Fill in the name and address of the healthcare institution/ provider which performed the implantation.
- 5** Remove **one** of the patient labels that includes the data matrix code from the sterile packaging and stick it on the designated field.
- 6** Once the Patient Implant Card has been filled out completely, it must be handed over to the patient. The purpose of the Patient Implant Card and the information contained therein must be explained to patients in a language that they understand.

## Explanations of the symbols use

-  Patient name and surname (or patient ID)
-  Date of implantation
-  Name and address of the implanting healthcare institution
-  Patient information website
-  Medical device name
-  Unique device identification
-  Unique device identification – device identifier (=GTIN code)
-  Catalog number
-  Batch code
-  Date of manufacture
-  Use-by date
-  Name and address of the manufacturer

# Patient Implant Card


# Geistlich Pharma

 Manufacturer  
Geistlich Pharma AG  
Bahnhofstrasse 40  
6110 Wolhusen  
Switzerland  
Tel.: +41 41 492 55 55  
info@geistlich.com  
www.geistlich-pharma.com

ENGLISH


## Examples for how to fill out the patient implant card completely


To help you fill out the card, see the following examples with explanations. These examples should help you check the content of the patient implant card for completeness and accuracy prior to giving it to the patient.





**Geistlich**  
Pharma

**Patient Implant Card**


 **John Doe**  
or patient ID 1234

 **07/29/2020**

 **Any dental practice**  
Any road 1  
1000 Any town  
Switzerland


 pic.geistlich-pharma.com


Front





**Geistlich**  
Pharma

**Patient Implant Card**

 **John**

 **07/29/2020**

 **Any practice**  
Any town

 pic.geistlich-pharma.com

## A fully completed patient implant card contains:

- > the patient's name in full or patient ID
- > date of implantation
- > **full** name and address of the healthcare facility performing the implantation

## An incompletely filled out patient implant card, for example, contains:

- > illegible and partial details
- > abbreviations which are not customary
- > **incomplete** name and address of the healthcare facility performing the implantation



Back



## A fully completed patient implant card contains:

- > the right translation of the **device type** in the respective national language
- > a patient label containing the data matrix code in the designated field

## An incompletely filled out patient implant card, for example, contains:

- > areas covered by the patient label and so no longer legible