

# Customer Feedback Form

The inclusion of as many details as possible greatly aids the complaint investigation process as well as provides useful information for continuous improvement. **Missing information will delay processing. All fields are required.**

## A. Customer Information

Date of Report: \_\_\_\_\_

City, State, Zip, Country: \_\_\_\_\_

Account # \_\_\_\_\_

Phone #: \_\_\_\_\_

Account Name/Dr: \_\_\_\_\_

E-mail: \_\_\_\_\_

Address: \_\_\_\_\_

Submitted By: \_\_\_\_\_

**B. Product Information:** One form should be used per patient or event. If more than one device is associated with a single reported event, multiple item numbers may be included below.

### Component Type:

Implant  Abutment  Screw  Instrument/Tool  Other \_\_\_\_\_

Order # (if available): \_\_\_\_\_

Item Number	Lot/Serial #	Qty.

## C. Product Replacement

**Are we replacing the same product?**  Yes  No - please provide preferred length and diameter \_\_\_\_\_

(if not specified, same size implant will be sent as a replacement). Store Credit will not be issued in lieu of replacement.

Patient/Case ID: \_\_\_\_\_

## D. Event Information

**Implantation Date** (mm/dd/yy): \_\_\_\_\_ **Loading Date** (mm/dd/yy): \_\_\_\_\_ **Removal Date** (mm/dd/yy): \_\_\_\_\_

### Description of the Event (check all that apply):

No Primary Stability  Failure to integrate (Before or coincident with loading)  Loss of Integration (After Loading)  
 Packaging or Labeling  Fit  Fracture  Customer Service  Other (please describe): \_\_\_\_\_

### Discovered During:

Receiving  Unpacking  Clinical Procedure  Laboratory Procedure  
 Other: \_\_\_\_\_

### Procedure Completed in the Same Visit:

Yes  No

**Provide a detailed description of the reported problem (including procedure being performed, related products and settings used):** \_\_\_\_\_

### Outcomes Attributed to Event:

Death  N/A  
 Life Threatening  Required Intervention to Prevent Permanent Impairment  
 Hospitalization  Disability or Permanent Damage  
 Other Serious (Important Medical Events)

**Describe Any Signs or Symptoms present in patient as a result of the event (check all that apply):**

- No Patient Impact  Allergic Reaction  Aspiration  Hemorrhage  Nerve Damage  Pain  Bone Loss  
 Dehiscence  Delayed Healing  Edema  Hyperesthesia  Hyperplasia  Infection  Inflammation  
 Other: \_\_\_\_\_

**E. Patient Information**

**Gender:**  Male  Female

**Age at Time of Event:** \_\_\_\_\_

**Tooth Number:** \_\_\_\_\_

**Bone Density Type:**  I  II  III  IV  Unknown

**Oral Hygiene:**  Excellent  Good  Fair  Poor

**Other Relevant Patient History (check all that apply):**

- Smoker/Tobacco use  Bruxism /Clenching  Osteoporosis  Diabetes  Biphosphonate Therapy  Steroid Therapy  
 Chemotherapy  Radiation Therapy  Periodontitis  Other: \_\_\_\_\_

**Instructions for US/Canada Customers and International Distributors:**

1. Please complete each field of this form.
2. Submit form to [customer.claims@implantdirect.com](mailto:customer.claims@implantdirect.com)
3. Within 3 business days of submitting this form, Customer Claims department will issue an RMA and a UPS label through UPS.com for the return of the product.
4. A replacement order will be processed after the UPS label has been received/scanned by UPS  
(some complaint cases requiring additional evaluation may delay replacement orders.)

**Instructions for Europe Customers:**

1. Please call 00800 4030 403 or email [inside.sales@implantdirect.eu](mailto:inside.sales@implantdirect.eu) to receive an RMA and Feedback Form .
2. RMA #: \_\_\_\_\_
3. Complete each field of this form.
4. Submit form to [inside.sales@implantdirect.eu](mailto:inside.sales@implantdirect.eu)
5. Customer Service will arrange a pickup on your preferred date.
6. A replacement order will be processed after receipt and inspection of the product.

**When returning product the following guidelines must be followed:**

- Used product MUST be sterilized in pouches which show sterility with color change or other indications prior to shipping.
- Please write claim number on the outside of the box. Must include a copy of the Return Material Authorization.

**United States**

Implant Direct Manufacturing  
Attn: Complaints Department  
3050 E. Hillcrest Drive  
Thousand Oaks, CA 91362  
Phone: 1.888.649.6425  
Email: [customer.claims@implantdirect.com](mailto:customer.claims@implantdirect.com)

**Europe**

Implant Direct AG  
Basicweg 20  
BR Amersfoort, 3821 Netherlands  
Phone: 00800 4030 403  
Email: [inside.sales@implantdirect.eu](mailto:inside.sales@implantdirect.eu)

**Canada**

Implant Direct Sybron Intl.  
55 Laurier Drive  
Morrisburg, Ontario  
Phone: 1.888.649.6425  
Email: [customer.claims@implantdirect.com](mailto:customer.claims@implantdirect.com)

**Australia**

Implant Direct Oceania  
Unit 10 112-118 Talavera Road  
North Ryde NS 2113, North, 2113 Australia  
Phone: 02.8870.3099  
Email: [cbrown@implantdirect.com.au](mailto:cbrown@implantdirect.com.au)