

PRODUCT COMPLAINT FORM



Please complete this form and send it to your regional Telix Customer Service by email or use the button [Submit Form](#) below.

CUSTOMER CATEGORY

DATE OF REPORTED INCIDENT

Does the complainant wish to be contacted regarding this complaint? Yes No

Customer's contact details

Site name	Contact name and role
Address	Contact phone
Country	Contact email

If the reporter is not the end user, add the end-user's details below. *(for clinical trials do not add patient details, see below)*

Site name	Contact name and role
Address	Contact phone
Country	Contact email

To be completed by the end-user

Product description
Batch number
Quantity
Expiry date

Please give a detailed description of the issue
[refer to attachment for more detail](#)

Location of the occurred issue

Role of operator who experienced the issue
(If other, please specify)

Product injected
*(If Yes, please report within 24hrs to
pharmacovigilance@telixpharma.com)*

Yes No

Complaint linked to Clinical Trial
(If Yes, please include trial number and patient number)

Yes No Clinical Trial IIT EAP/AS Compassionate Use
Trial Number: Patient Number:

For clinical trial products: Harm to patient
(If Yes, please describe)

Yes No

Supportive evidence of the incident available for
investigation by Telix (photos, reports, etc)
(If Yes, please send as attachment)

Yes No

Note. If email client does not launch once the button is pressed,
please submit form manually by email to the region the issue
occurred.

For Americas: customerservice.americas@telixpharma.com
For Asia Pacific (APAC): customerservice.apac@telixpharma.com
For Europe (EMEA): customerservice.emea@telixpharma.com

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AMERICAS
11700 Exit 5 Pkwy,
Suite 200 Fishers, IN
46037, USA
+1 463-235-2159

ASIA PACIFIC (HQ)
55 Flemington Road, North
Melbourne, VIC 3051
AUSTRALIA
+61 1800-0-83549

EUROPE
Rue de Hermée 255
4040 Herstal
BELGIUM
+32 2 616 40 00