Report an Adverse Event

THIS SITE IS NOT INTENDED OR DESIGNED FOR REPORTING OF ADVERSE EVENTS

Roche is required to document reports of adverse events, both for comprehensive safety monitoring of its products and also to fulfil reporting requirements to Regulatory Authorities. It allows continued monitoring of the benefit/risk balance of the medicinal product. Adverse events should be reported.

Reporting of suspected adverse events or reactions (for healthcare professionals)

Reporting suspected adverse events or reactions after authorisation of the medicinal product is important. It allows continued monitoring of the benefit/risk balance of the medicinal product. Healthcare professionals are asked to report any suspected adverse reactions (see details below).

Where possible, healthcare professionals should report adverse events or reactions by brand name and batch number.

In the event of a suspected adverse event,
please report it to:

The Drug Surveillance Centre
Roche Products (Ireland) Limited
3004 Lake Drive, Citywest,
Naas Road, Dublin 24
Telephone: (01) 4690700
Email: ireland.drug_surveillance_centre@roche.com

Alternatively, suspected adverse reactions should be reported to:

HPRA Pharmacovigilance
Website: www.hpra.ie