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Reporting Adverse Events to MedWatch

Adverse events, product quality problems, product use errors, or reports of therapeutic inequivalence or failure should be reported to **MedWatch**, The FDA Safety Information and Adverse Event Reporting Program.



Complete and submit the report online:
[MedWatch Online Voluntary Reporting Form](#).

Download [form](#) at or call **1-800-332-1088**
to request a reporting form,
or submit by fax to **1-800-FDA-0178**.

