



# MedWatch

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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**.

