FDA

MedWatch

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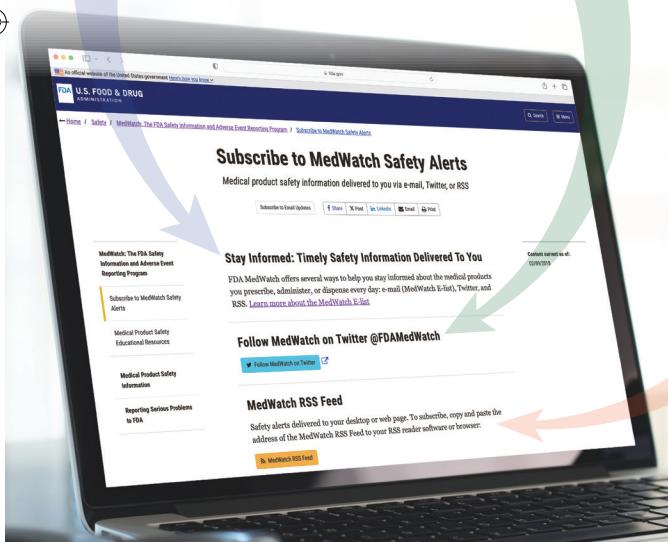
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Safety alerts delivered directly to your device or website.

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**.