**Site Language**

MedCheck: Check Twice. Choose Once.

Intro Text

* Search for a medication and find related recalls and adverse events.
* MedCheck is an open-source web application exploring openFDA data to provide medication information and its relation to product recalls and adverse event reports.

Disclaimer

* It is not the intention of Octo Consulting to provide specific medical advice, but rather to provide users with information to better understand medication recalls and adverse events provided by the openFDA api. Please contact your physician for all medication related questions.

Drug Results

* The results of your search for [Medication Name] includes(s) [resultNumber] related items.

Seriousness

* This was a non-serous event.
* This event caused patient hospitalization.
* This event was life threatening to the patient.
* This event ended in the death of a patient.
* This event caused the disability of a patient.
* This event caused a congenital defect to the patient.
* This event has an unknown level of seriousness.

Search Alerts

* No results were found for your search.
* Your search could not be made at this time.

Recall Description

* Recalls are actions taken by a firm to remove a product from the market. Recalls may be conducted on a firm's own initiative, by FDA request, or by FDA order under statutory authority.

Adverse Reaction Description

* An adverse event is any undesirable experience associated while using a medication.

**Recall Data**

Recall Number

* Tracking number associated with recall.

Recall Initiation Date

* Date that the firm first began notifying the public or their consignees of the recall.

Reason for Recall

* Information describing how the product is defective and violates the FD&C Act or related statutes.

Distribution Pattern

* Information describing how the product is defective and violates the FD&C Act or related statutes.

Recalling Firm

* The firm that initiates a recall or, in the case of an FDA requested recall or FDA mandated recall, the firm that has primary responsibility for the manufacture and (or) marketing of the product to be recalled.

Quantity

* Amount of items recalled.

Product Description

* Information about the recalled medication

**Adverse Event Data**

Safety Report Id

* 8-digit Safety Report ID number, also known as the case report number or case ID.

Received Date

* Date that the FDA first received the report.

Seriousness

* The level of seriousness of the event and any reported outcome.

Reactions

* MedDRA term(s) for the reaction(s). Note that these terms are encoded in British English. For instance, “diarrhea” is recorded as “diarrohea.”.

Patient Age

* The age of the patient when the event first occurred.

Patient Sex

* The sex of the patient when the event first occurred.

Substance Name(s) = Other Substance(s) Taken

* Name of substances taken by the patient when the event first occurred.

**Chart Legends**

Top 5 Reactions

* Reaction 1
* Reaction 2
* Reaction 3
* Reaction 4
* Reaction 5

Outcomes

* Recovered/resolved.
* Recovering/resolving.
* Not recovered/not resolved.
* Recovered/resolved with sequelae.
* Fatal.
* Unknown.