



PDS – FDA MedWatch

Adverse Drug Reaction (ADR) Reporting Form



Patient Information

Name:

Age:

Gender:

Address (if available):

Contact number (if available):

Hospital/facility seen or admitted:

Details of Adverse Reaction

Date of onset:

Duration:

Do you consider the reaction to be serious?

- Yes, if yes indicate why: (tick/mark one)

- Patient died due to reaction

- Involved or prolonged in-patient hospitalization

- Life threatening Involved persistent or significant disability

- Congenital anomaly in the newborn

- Other outcome, please give details: _____

- No

Can this be due to Medication Error?(encircle one) No Yes

*Suspected drug product(s) Indicate brand name	Daily dose	Route	Reason(s) for using the product (indication)	Date started	Date stopped (if applicable)

Reporter's Details

Name:

Contact number:

Email: