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ADVERSE DRUG REACTIONS REPORT FORUM

. IF YOU SUSPECT THAT ADVERSE REACTION MAY, BE RELATED TO A CERTAIN DRUG OR COMBINTION OF DRUGS YOU SHOULD COMPLETE THIS FORUM AND SEND TO THE ADDRESS SHOWN AT THE END OF THE REPORT.

1.Patient details

Name/(initials)\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_Sex Male Female Weight Kg Age\_\_\_\_\_\_

|  |  |  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| 2.SUSPECTED  DRUG S | generic | Trade name | Concentration | Used  for | Dose | Rout of  administration | Data  started | | Data stopped | Batch number | Company name | How to get |
|  |  |  |  |  |  |  |  | |  |  |  |
|  |  |  |  |  |  |  |  | |  |  |  |
|  |  |  |  |  |  |  |  | |  |  |  |
| 3.OTHER DRUG S  CONCOMITANT |  |  |  |  |  |  |  |  | |  |  |  |
|  |  |  |  |  |  |  |  | |  |  |  |
|  |  |  |  |  |  |  |  | |  |  |  |

4. Suspected Reaction

Please describe the reaction(s): \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Data reaction (s) started: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_Date Reaction(S): \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Dose the reaction stooped after stopping the drug? Yes No Don’t know

Dose the reaction stooped after retaking the drug? Yes NO Don’t know

Was the reaction serious?

If serious specify; Patient Died Life threatening hospitalization

Prolonged hospitalization congenital anomaly Permanents Disability

Required intervention to prevent damage other specify\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

5.Reporter Details

The one who fill this form Patient Physician pharmacist nurse other specify

Name \_\_\_\_\_\_\_\_\_\_\_Specialty (if physician) \_\_\_\_\_\_\_\_\_ Address\_\_\_\_\_\_\_\_\_Mobile\_\_\_\_\_\_\_\_

FEX E-mail date signature

6-Any More Comments: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

. The information in this report is confidential and totally protected including both the patient and reporter identity.

. You can send voluntarily the adverse drug reaction (ADRs) reports to the pharmacovigilance department.

. Reporting for ADRs is vital for safely usage of drugs. enough information will help the department to evaluated the safety of the drugs marketed in our country

Pharmacovigilance department: ministry of heath

Behind Tripoli medical center

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