ADVERSE DRUG REACTION REPORTING FORM

| A. | PATIENT DETAILS | | | |
|--------|--|--|--|--|
| | Patient name | | | |
| | Age at time of onset(yrs)* | | | |
| | Weight (kg) | | | |
| | Patient Number | | | |
| | Health Facility | | | |
| | District | | | |
| | Sex: M/F* | | | |
| | Last Menstrual Period | | | |
| | Trimester (if pregnant) | | | |
| В. | SUSPECTED DRUG (S) DETAILS | | | |
| | Generic Name* | | | |
| | Brand Name | | | |
| | Dose ,Route Frequency | | | |
| | Date* started | | | |
| | Date stopped | | | |
| | Prescribed for Expiry date | | | |
| | Batch No | | | |
| C. | SUSPECTED REACTIONS | | | |
| Pleas | se describe the reaction as observed and any treatment given to manage the reaction; | | | |
| i icac | to describe the reaction as observed and any treatment given to manage the reaction, | | | |
| | | | | |
| | | | | |
| Outc | ome: | | | |
| | | | | |
| Recov | vered | | | |
| Recov | vering | | | |
| Contir | nuing | | | |
| Death | Death due to reaction | | | |
| | | | | |

| Date reaction started* | Date reaction stopped | Date of notification |
|------------------------|-----------------------|----------------------|
| | | |

| Patient died | | | | |
|--|--------------------------------|-------------------|-----------------|--------|
| Prolonged inpatient Hospitalization | | | | |
| Involved disability | | | | |
| Life Threatening | | | | |
| Congenital abnormality | | | | |
| D. CONCOMITANT DRUGS | | | | |
| Please give information on the including those taken for chronic Generic Name | | | • | d drug |
| Brand name | | | | |
| Dosage | | | | |
| Date started | | | | |
| Date stopped | | | | |
| Indication(prescribed or OTC) | | | | |
| | | | | _ |
| Relevant laboratory tests including | ng dates | | | |
| Additional relevant information (mo | edical history, allergies,fail | ure of efficacy) | | |
| Name/designation* | Telephone and Email Address | Date of reporting | Health facility | |
| | | | | |

SERIOUSNESS OF THE REACTION