



## DEPARTMENT OF HEALTH

Larry Hogan, Governor · Boyd K. Rutherford, Lt. Governor · Dennis R. Schrader, Secretary

Office of Health Care Quality  
Maryland Hospital Safety Program  
7120 Samuel Morse Drive  
Second Floor  
Columbia, MD 21046

Please send encrypted reports and RCAs to [hospital.selfreport@maryland.gov](mailto:hospital.selfreport@maryland.gov)

### **INITIAL REPORT OF AN ADVERSE EVENT**

#### **SECTION I: GENERAL INFORMATION**

Hospital Name: \_\_\_\_\_

Person completing this report: \_\_\_\_\_

Title: \_\_\_\_\_ Phone Number: \_\_\_\_\_ Email: \_\_\_\_\_

Date of Report: \_\_\_\_\_ Date of Event: \_\_\_\_\_

Location of Event: \_\_\_\_\_

Area or Service (e.g. ED, OR, Med/Surg etc.): \_\_\_\_\_

Was TJC notified?    YES                    NO

#### **SECTION II: PATIENT INFORMATION**

**Patient #1** initials or patient number only: \_\_\_\_\_

Date of Admission: \_\_\_\_\_ Age: \_\_\_\_\_

Admitting Diagnosis: \_\_\_\_\_

Current Status: \_\_\_\_\_

Prognosis: \_\_\_\_\_

Was the patient /family informed of the adverse event?    YES                    NO

**Patient #2** initials or patient number only: \_\_\_\_\_

Date of Admission: \_\_\_\_\_ Age: \_\_\_\_\_

Admitting Diagnosis: \_\_\_\_\_

Current Status: \_\_\_\_\_

Prognosis: \_\_\_\_\_

Was the patient /family informed of the adverse event?      YES      NO

**SECTION III: INTENTIONALLY UNSAFE ACTS**

If the event was the result of an intentionally unsafe act such as abuse, please complete the following:

Position/Title: \_\_\_\_\_ License # \_\_\_\_\_

If the staff person was licensed or certified, was the applicable professional board notified?

YES

NO

If the staff is employed through an agency or through a contract company and is not a hospital employee or member of the medical staff, please provide the employer's name.

\_\_\_\_\_

If the police were notified, please provide the jurisdiction and the report number if known: \_\_\_\_\_

**SECTION IV: EQUIPMENT**

If the event was the result of equipment or medical device malfunction or failure please provide the following information.

Equipment or Device: \_\_\_\_\_ Model number: \_\_\_\_\_

Nature of Malfunction: \_\_\_\_\_

\_\_\_\_\_

Was FDA Notified?      YES      NO

**SECTION V: DESCRIPTION OF THE EVENT:** (Briefly describe the event and include the outcome to the patient, use additional pages if necessary)