



Participant ID:

Participant Initials:

Clinical Center:

Site:

Visit Number:

CRF Date:

RC ID:

URGENT ALERT

RC/PI completes upon receiving report containing alert values..

Completion of Urgent Alert CRF may require an EVENT investigation/confirmation and completion of EVENT CRF.

1. Date of Alert Value (s):

___/___/___
MM DD YYYY

Type of Alert Value(s):

2. **LAB:**

- a. Potassium ≥ 6 mEq/L or ≤ 3.0 mEq/L: ₁ Yes ₀ No
- b. Sodium <125 mEq/L or >155 mEq/L: ₁ Yes ₀ No
- c. Total Bicarbonate <15 mEq/L or > 40 mEq/L: ₁ Yes ₀ No
- d. Calcium <6.5 or >13.5 mg/dL: ₁ Yes ₀ No
- e. Glucose < 50 mg/dL or > 350 mg/dL: ₁ Yes ₀ No
- f. Creatinine doubling from last value: ₁ Yes ₀ No
- g. CBC Hb < 7 gm/dL: ₁ Yes ₀ No
- h. Other abnormal lab value. **Specify.** _____ ₁ Yes ₀ No

3. **ECG:**

- a. Date of Reading; _____ ₉₈ Not Done
MM DD YYYY

Skip to Question #4, if ECG was not ordered.

- b. Reading: ₁ Local ₂ Central
 - i. Bradycardia (<45 beats/min): ₁ Yes ₀ No ₂ False Positive
 - ii. Tachycardia (>120 beats/min): ₁ Yes ₀ No ₂ False Positive
 - iii. Acute Myocardial Infarction or acute ischemia: ₁ Yes ₀ No ₂ False Positive
 - iv. Ventricular Tachycardia: ₁ Yes ₀ No ₂ False Positive
 - v. Atrial Fibrillation: ₁ Yes ₀ No ₂ False Positive
 - vi. Atrial Flutter: ₁ Yes ₀ No ₂ False Positive
 - vii. Mobitz Type II 2nd degree Heart Block: ₁ Yes ₀ No ₂ False Positive
 - viii. 3rd degree Heart Block: ₁ Yes ₀ No ₂ False Positive
 - ix. Complete Left Bundle Branch Block: ₁ Yes ₀ No ₂ False Positive

4. **Action Taken:**

- a. Primary MD notified: ₁ Yes ₀ No
- b. Report sent to primary MD: ₁ Yes ₀ No

5. **Participant notified of outcome?**

- ₁ Yes ₀ No ₉₉ N/A