

## HALF-PINT Form 2: Tracking and Randomization

**Note:** Complete this form using information recorded on the paper log "Consented Subject Tracking Sheet".

1. Was tracking discontinued because the subject became ineligible (for a reason other than off IV vasopressors/inotropes or extubated) after consent was obtained?

**If the subject became ineligible for one of these reasons, STOP HERE.**

☐ Yes

If Yes, why did the subject become ineligible?

- ☐ (1) Discharged from ICU  
☐ (2) Family/team have decided to limit/redirect from aggressive ICU technological support  
☐ (3) Had new cardiac surgery  
☐ (4) Reached Study Day C-28  
☐ (99) Other, specify:

☐ No

2. Record the blood glucose (BG) measurements  $\geq 150$  mg/dL that made the subject eligible for randomization

First BG  $\geq 150$  mg/dL:

 mg/dL

Date/time of first BG  $\geq 150$  mg/dL:

|  / |  / |   
|  : |  24-hour clock

Confirmatory BG  $\geq 150$  mg/dL:

 mg/dL

Date/time of confirmatory BG  $\geq 150$  mg/dL:

|  / |  / |   
|  : |  24-hour clock

3. Was the subject ever randomized?

**If the subject was randomized, continue to Forms 3, 4, and the daily forms.**

☐ Yes

If Yes, what was the group assignment?

- ☐ TGC-1 (80-110 mg/dL)  
☐ TGC-2 (150-180 mg/dL)

Date/time of randomization:

|  / |  / |   
|  : |  24-hour clock

☐ No

If No, why was the subject not randomized?

- ☐ (1) All study equipment in use for other randomized subjects  
☐ (2) Withdrawal of consent  
☐ (3) Clinician request, specify:

- ☐ (4) Subject became ineligible after confirmatory BG  $\geq 150$  mg/dL

☐ (5) > 24 hours between confirmatory BG and randomization

☐ (99) Other, specify: