

Michael Denaburg, Pharm.D.<sup>1</sup>; Earnest Alexander, Pharm.D., FCCM<sup>1</sup>; Maresa Glass, Pharm.D., BCPS<sup>1</sup>; Ren Chen, MD, MPH<sup>2</sup> Tampa General Hospital Department of Pharmacy<sup>1</sup>, University of South Florida<sup>2</sup>, Tampa, Florida

### Background

- Tampa General Hospital (TGH) is a 1,018-bed level 1 trauma, academic medical center
- Surgical site infections (SSIs) are a significant source of morbidity and mortality in post surgical patients<sup>1,2</sup>
- Hyperglycemia is associated with an increase in surgical site infections in addition to morbidity and mortality, particularly in postoperative cardiac surgery patients<sup>3,4</sup>
- The Surgical Care Improvement Project (SIPC) was developed by CMS, the CDC, and the Joint Commission to decrease the incidence of SSIs<sup>5</sup>
- SCIP measure INF-4 requires that cardiac surgery patients have blood glucose less than or equal to 200 mg/dL at 6am on postoperative (POD) days one and two<sup>5</sup>
- Endotool<sup>®</sup> is an FDA-approved glucose management software system that that optimizes insulin therapy by utilizing adaptive algorithms to provide individualized therapy<sup>6</sup>
- Endotool<sup>®</sup> has been utilized at TGH in the Cardiothoracic Intensive Care Units (CTICU) since January 2011 and was implemented intraoperatively in cardiac surgery patients in January 2012

# Objective

□ To analyze the impact of utilizing Endotool<sup>®</sup> for blood glucose management intra-operatively on compliance with SCIP measure INF-4 compared to patients on Endotool<sup>®</sup> post-operatively alone

## **Contact Information**

Michael Denaburg, Jr., Pharm.D. PGY2 Critical Care Pharmacy Resident Tampa General Hospital, Tampa, FL Email: mdenaburg@tgh.org

# Implementation of Glucose Management Software (Endotool<sup>®</sup>) Intra-operatively in Cardiac Surgery Patients to Improve Post-operative Blood Glucose Control and Compliance with SCIP INF-4

#### Methods

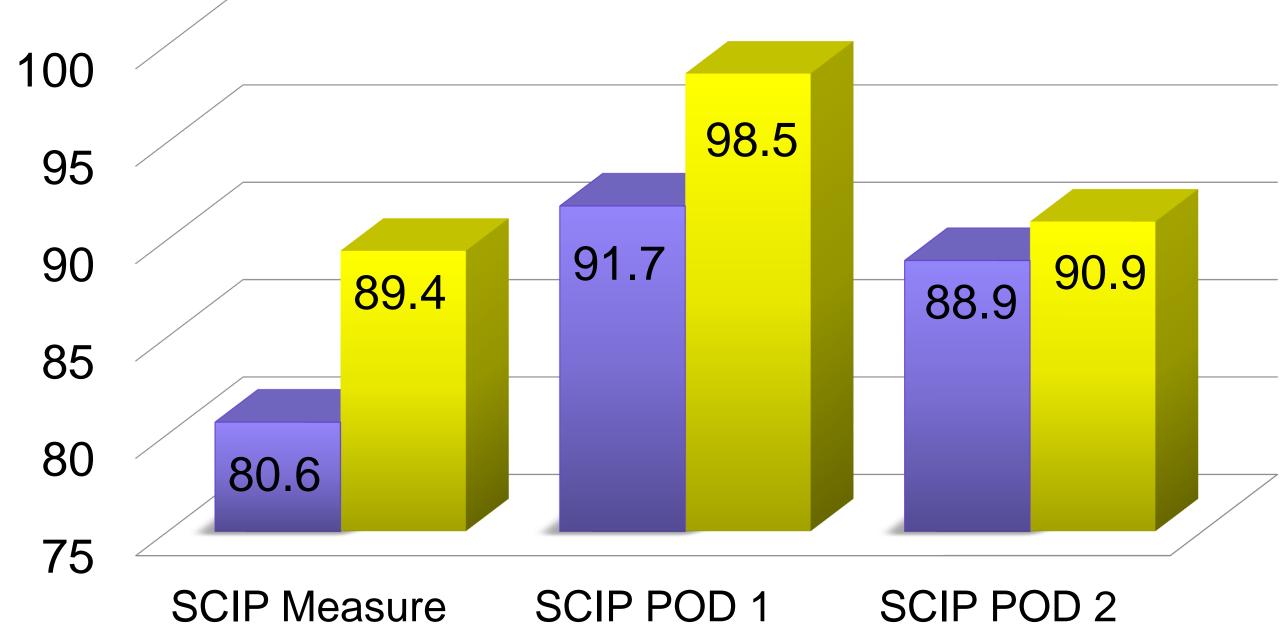
- Single center, retrospective, IRB-approved study
- Summary of patients receiving cardiac surgery were obtained from Data Resources between October 1, 2011 and March 31, 2012.
- Inclusion criteria:
  - All inpatients receiving cardiac surgery and utilizing Endotool<sup>®</sup>
- Exclusion criteria:
  - Less than 18 years of age
  - $\Box$  Length of stay >120 days
  - Diagnosis of preoperative infectious disease
  - Burn and transplant diagnosis
  - **Laparoscopic procedures**
  - Concurrent enrollment in clinical trial
  - Perioperative mortality
  - Patients not utilizing 5Endotool<sup>®</sup>
- Elements of data collection:
  - Anesthesia start and stop date/time
  - □ 6am blood glucose on POD1 and POD2
  - Duration of Endotool<sup>®</sup> (hours)
  - □ Nursing adherence with Endotool<sup>®</sup> recommendations
- Outcomes:
  - Primary:
    - Overall percentage of patients meeting SCIP measure INF-4
  - Secondary:
    - Percentage meeting the SCIP measure INF-4 on POD 1
    - Percentage meeting the SCIP measure INF-4 on POD 2
- **Statistics**:
  - Chi-squared or Fishers exact test, as appropriate

# SCCM 42<sup>nd</sup> Annual Congress – San Juan, Puerto Rico January 18<sup>th</sup> – January 23<sup>rd</sup> 2013

#### Results

- A total of 102 patients were included in the analysis (n=36 control, n=66 intervention)
- □ The overall percentage of patients that met the SCIP measure was 80.5% pre-implementation and 89.4% post-implementation of Endotool® intra-operatively, respectively (p=0.239). See Figure 1
- On POD1, the difference between the pre- and post-implementation was 6.8%; 91.7% and 98.5%, respectively (p=0.125). See Figure 1
- Figure 1.

Percentage of patients compliant with SCIP INF-4 measure overall, POD 1 only, and POD 2 only



Only (p=0.1248) Only (p=0.7390)Met (p=0.2392)

Endotool CTICU Only (Control Group) Endotool in COR and CTICU (Intervention Group)

- 100% of patients in both groups that met the SCIP measure on POD 1 but not POD 2, Endotool<sup>®</sup> was discontinued prior to the POD 2 6am blood glucose
- Number of patients with non-adherence events:
  - □ Control: 6 (16.7%) 1 did not meet SCIP
  - Intervention: 10 (15.2%) 1 did not meet SCIP

## Discussion

Society of

**Critical Care Medicine** 

- Endotool<sup>®</sup> used to manage blood glucose intraoperatively in cardiac surgery patients increased our overall compliance with SCIP measure INF-4 as displayed in Figure 1
- Reasons identified for not meeting SCIP compliance:
  - Discontinuation of Endotool<sup>®</sup> prior to POD 2
  - Decreased time between anesthesia stop date/time and POD 1 blood glucose
- Future Directions:
  - Continue utilization of Endotool<sup>®</sup> in the cardiac operating room and in the cardiothoracic intensive care unit to maintain euglycemia
  - Re-evaluate SCIP measure INF-4 compliance

# Conclusion

Endotool<sup>®</sup> utilized intra-operatively to manage blood glucose in cardiac surgery patients lead to improvements in our overall compliance with SCIP measure INF-4

#### References

- Klevens RM, Edwards JR, Richards CL, et al. *Public* Health Reports (1974-). 2007;112:160-166.
- Ascione R, Rogers CA, Rajakaruna C, et al. Circulation. 2008;118:113-123.
- Zerr KJ, Furnary AP, Grunkemeier GL, et al. Ann Thorac Surg. 1997 Feb;63(2):356-361. PMID: 9033300.
- Latham R, Lancaster AD, Covington JF, et al. Infect *Control Hosp Epidemiol.* 2001 Oct;22(10):607-612. PMID: 11776345.
- Rosenberger LG, Politano AD, Sawyer RG. Surg Infect. 2011;12:163-168
- Endotool website. Available at: http://www.endotool.com. 6. Accessed August 20, 2012.

## Disclosure

Authors of this presentation have nothing to disclose concerning possible financial or personal relationships with commercial entities that may have a direct or indirect interest in the subject matter of this presentation.