

Using Blood Glucose Meters in the Hospital: Defining "Critically III" and Addressing Accreditation Issues



Stephen E. Kahn, PhD, DABCC, FACB Professor and Vice Chair, Clinical Services, Pathology Loyola University Health System Mawwood, IL

April 28, 2016

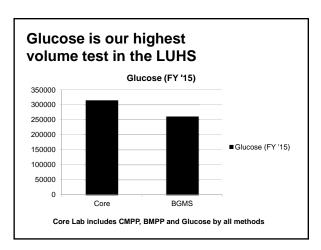
Speaker Financial Disclosure Information

- Potential COI to disclose relevant to the topic and issues of this presentation:
 - Grant/Research Support: None
 - Salary/Consultant Fees: None
 - Board/Committee/Advisory Board Membership: None
 - Stocks/Bonds: None
 - Honorarium/Expenses: AACC for this presentation
 - Intellectual Property/Royalty Income: None

Presentation Objectives

- Following this presentation, audience members will be able to:
 - List 3 analytical, clinical or regulatory challenges faced by U.S. hospitals in the use of BGMS*
 - Describe 3 strategies that could enhance effective use of BGMS in hospitals (esp. your hospital) by increasing patient safety and/or regulatory compliance
 - Recognize the relevance of distinguishing between developing a definition of 'critically ill' and specifying criteria for acceptable use of capillary finger stick specimens (esp. as it applies to your hospital)
- * BGMS: Blood glucose monitoring systems, aka, bedside glucose meters

The Hospital BGMS Practice Environment: Significant Issues



Measuring glucose – Some common approaches

- Non-BGMS* Glucose CAP 2015 C-B Chem/Tox Survey
 - 5 general enzymatic categories, 39 method peer groups, ~5500 participants
- BGMS Glucose CAP 2015 A WBG Survey
 - 8 manufacturers, 22 different BGMS devices or glucose strip categories,
 45000 participants
- BGMS report "Plasma-equivalent glucose"
- * BGMS: Blood glucose monitoring systems, aka bedside glucose meters

BGMS Use in Hospitals in the 21st Century

- Hospitals use multiple protocols for 'managing' glycemic control
- Ongoing concerns patient safety and medical errors
 - Numerous published studies and clinical practice guidelines
 - Studies as large as NICE-SUGAR not done easily
 - Have led to growth in modeling and simulation studies e.g.,
 - Insulin dosing models
 - Impact of BGMS testing frequency for GC monitoring
- Specific concerns are <u>causes and sources of errors</u> in BGMS measurement, especially glycemic control protocols
 - Focus of regulatory and accreditation agency actions
 - Followed through institutional and national quality metrics
 - Can be addressed by unique practices, e.g., insulin dosing software

Clinical Concerns in the Hospitalized and ICU Patient

- Sick dehydrated, in shock, often on oxygen, impaired peripheral circulation
- Fluctuating hematocrits not always known at time of testing
- Take multiple drugs how do drugs affect a specific BGMS?
- Often have rapidly changing glucose values
- Can be tested using multiple meters by multiple operators
- Some operators may be unaware of the limitations
- Potential to test patients where BGMS actual use may be different from mfr product labelling for intended use

Potential SOEs in BGMS* Factors affecting bedside glucose testing User Error User Error e.g., Improper sampling, expired strips, wrong strips, coding errors Environmental Factors e.g., Altitude, temperature, humidity Exogenous Interferences e.g., Hematocrit Exogenous Interferences e.g., Hematocrit e.g., Maltose, galactose, xylose, ascorbate, acetaminophen

Who You Gonna' Call? The 'Usual Suspects' Regulatory Accreditation CMS

Additional Resources





The Joint Commission

Regulatory Actions Since 2014

- Jan '14: FDA draft guidance to BGMS mfrs
 - BGMS must be approved for use in the intended population including 'critically ill'
 - Mounting national concerns and confusion on many issues
- Sept '14: FDA approves first BGMS for use in the 'critically ill' patient
- Nov '14: CMS directive to state surveyors use of BGMS not approved for use in 'critically ill' patients may be 'off label'
 - Use of capillary finger stick specimens in critically ill patients is (and presently remains) 'off label' for all BGMS
- March '15: CMS follow-up temporarily withdraws directive, adds clarifications, requests comments and holds off citations
- April '16: Most agencies and mfrs will not define 'critically ill' although the state of Illinois has done this ('sort of')

Why is FDA concerned about 'Plasma Equivalent Glucose'?



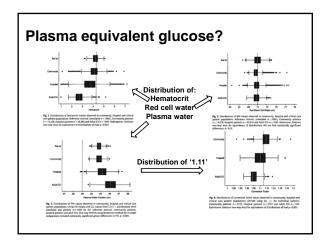
Patient acuity exacerbates discrepancy between whole blood and plasma method through error in molality to molarity conversion: "Mind the gap!"

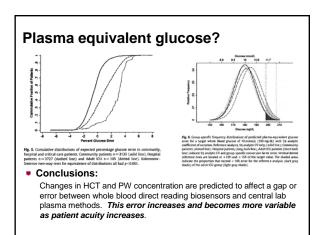
Martha E Lyon **** Andrew W. Lyon *****

**** Patients of Points of Lineary Names (Johns Count)

**** Patients of Points of Lineary Names (Johns Count)

- To report 'Plasma equivalent glucose,' BGMS are pre-set to use a constant molality to molarity conversion factor of 1.11
- In hospitalized patients, this constant is not a constant
- Other key analytical variables taken as a constant in BGMS that aren't constant are hematocrit (43%), plasma water (0.93 kg water/L) and RBC water (0.71 kg water/L)





LUHS Actions Since 2014

- 2014: Using a BGMS not cleared for use in 'critically ill' patients
 - Growing concerns on the FDA draft guidance to mfrs
- Dec '14: Notification of all key LUHS leaders and stakeholders of CMS directive regarding 'off label' use of BGMS
 - Initiate discussions on potential options for response
 - IP glycemic mgt team, ICU medical directors, MEC and nursing
- Feb '15: Lab leaders meet with CMO, CNO, and hospital administration
 - Agree to further explore options to achieve regulatory compliance
- Feb '15: LUHS parent, Trinity Health, selects BGMS cleared for use in 'critically ill' patients
- March '15: CMS follow-up directive 'buys' LUHS some time
- April '16: LUHS plans to convert to new BGMS in 2 months

The 'Usual Suspects' Regulatory (CMS Accreditation CMS The Joint Commission Additional Resources 🙎 **Accreditation Issues** (CMS Address the issues if 'off-label' use · Implement other options for glucose testing in 'critically ill' patients or with capillary finger stick specimens • Otherwise, high complexity testing requirements Validate BGMS in non-ICU and ICU (or 'critically ill') pts The Joint Commission To be determined **Relevant CLSI Guidelines** Several guidelines related to POCT glucose measurement and BGMS ■ POCT06: Effects of Different Sample Types ■ POCT12 - A3: POCT Glucose Testing in • POCT17 - ED1: Use of Glucose Meters for Critically III Patients Recent effort of the CAP Consensus Cmte on POCT Details options for hospitals to consider Outlines key issues and necessary studies for hospitals to address with 'off-label' use

CLSI POCT17 - ED1

- Intended use versus off-label use
- Important elements of mfr's instructions
 - Sticking to stated limitations of the specific BGMS
 - Approved specimen types and in what patient groups
- Issues to consider in developing a definition of 'critically ill'
- Off-label use
 - Specific regulatory requirements for high complexity
 - High complexity performance specifications
- Alternatives to off-label use



CLSI POCT17 - ED1

- Options to address the critically ill limitations in specific BGMS labeling and technical information
 - If a hospital defines 'critically ill,' it should state BGMS shouldn't be used in this patient group
 - Or use a different method without this limitation
 - Or if off-label use, perform validation studies required in CLIA to meet high complexity testing requirements



Every Hospital (or Multi-Hospital System) Is Unique

Analyzing Your Facility's Needs – Defining 'Critically III'

- Decision on creating a definition is a <u>local</u> issue and hospital (health system network) responsibility
 - Must decide whether to develop a definition regardless of BGMS used
 - Decision includes whether the BGMS is approved for use in 'critically ill' or it isn't
 - Overall effort must involve all key stakeholders
 - Other issues may impact your approach e.g.,
 - Applicable state regulations
 - Assessment of risk management/patient safety
 - IT capabilities of the hospital or hospital network

Analyzing Your Facility's Needs – Stakeholders

- Ongoing communication and collaboration between Lab personnel/POCT team and other stakeholders
 - Physician, nursing and administrative leaders
 - ICU medical directors
 - IP glycemic management team
 - Nursing education
 - Information technology
 - Institutional quality groups and leaders
 - Medical executive committee
 - Risk management/office of patient safety
 - Other stakeholders as determined by a specific hospital

Analyzing Your Facility's Needs – The BGMS Used

- Know the analytical limitations of the BGMS and shape policies/procedures accordingly
- Regardless of the BGMS used in a hospital
 - Some will choose to further develop an explicitly detailed definition of 'critically ill'
 - Others may decide to not have any definition at all
- But if a definition is developed, it should be used and clinical practices impacted as it describes

_		

Options for Clinical Criteria

- Impaired peripheral circulation, hypotension, peripheral edema,
 - SBP < 70 100 mmHg or may be age adjusted (e.g., lower in neonates)
 Can incorporate a specific delta decrease in SBP from baseline
 - MAP < 60 65 mmHg
 - Receiving ionotropic and/or vasopressor agents to support BP
 May also use a significant dose change in past 24 hours
 - Need for fluid resuscitation of some length in past 24 hours
 - Serum osmolality > 310 320 mOsm/kg
 - · 'Cold or clammy' skin
- Other clinical criteria
 - Diabetic ketoacidosis
 - Hypoxemia
 - Extreme hypoglycemia or hyperglycemia extremes defined
 - Clinical signs of dehydration

Other Considerations

- Whether a definition for 'critically ill' is developed or not, also must address appropriate patient groups where capillary finger stick specimens can and can't be used
- Any 'off-label' use should be addressed as described in CLSI POCT17 - ED1
- Clinical algorithms and/or critical care paths can be developed for use in the hospital EMR
- Some hospitals may choose to embed the criteria for when BGMS (and/or capillary finger stick specimens) can be used in their FMR
 - Electronically documents appropriateness of orders and provides electronic audit trail

BGMS Use in Hospitals: What Have We Learned

_			
-			
-			
_			
_			
_			
-			
-			
-			
-			
-			
_			
_			
-			
-			
_			

Conclusions

- Appropriate BGMS use is a top patient safety and medical error reduction priority
- Present BGMS use, especially for glycemic control, likely to continue until improved BGMS or different technologies
- Limitations and intended use for all BGMS are explicitly clear
- Patients that a BGMS and/or capillary finger stick specimen shouldn't be used on should also be explicitly clear
- Each hospital (or system) must weigh all relevant factors in taking the decided actions for their use of a specific BGMS
- Key regulatory and accreditation bodies will continue to provide guidance that direct the appropriate actions there is more work to be done

BGMS Use in Hospitals Should Work This Way Harmonizing the spectrum of POC performance, decision making and safety* Manufacturers and Technology Developers Optimizing POC Performance, Decision Making and Pattent Safety Healthcare Professionals *Louie RF et al. Point of Care 2014