

Parkview Health

Parkview Health Research Repository

Pharmacy Residency

Pharmacy Research

2018

Implementation of a pharmacist-guided procalcitonin protocol in a community hospital

Jasmine Coatie PharmD

Follow this and additional works at: <https://researchrepository.parkviewhealth.org/pharmresidency>



Part of the Pharmacy and Pharmaceutical Sciences Commons

Implementation of a Pharmacist-Guided Procalcitonin Protocol in a Community Hospital

Jasmine Coatie, PharmD

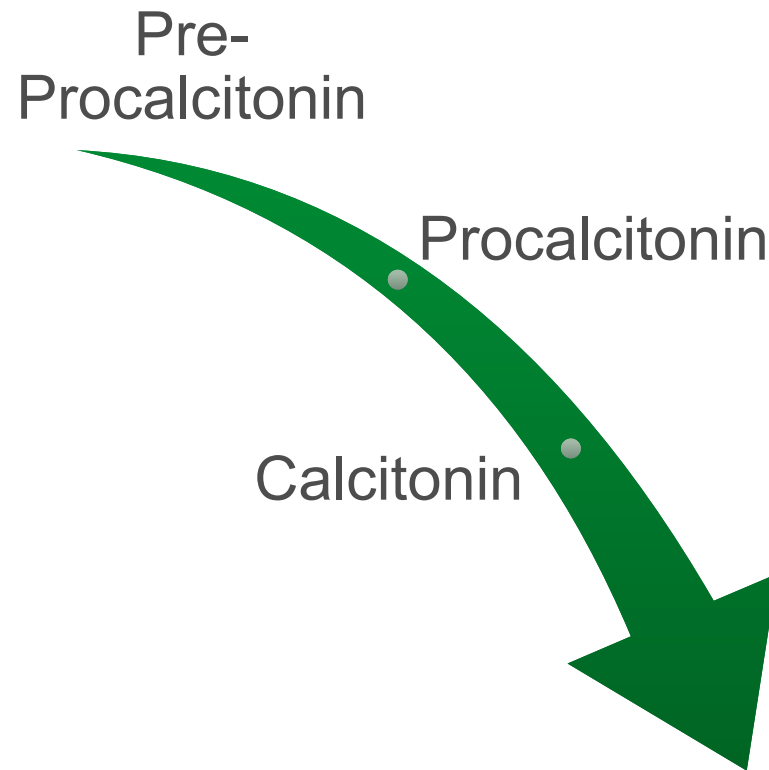
PGY1 Resident

Parkview Health

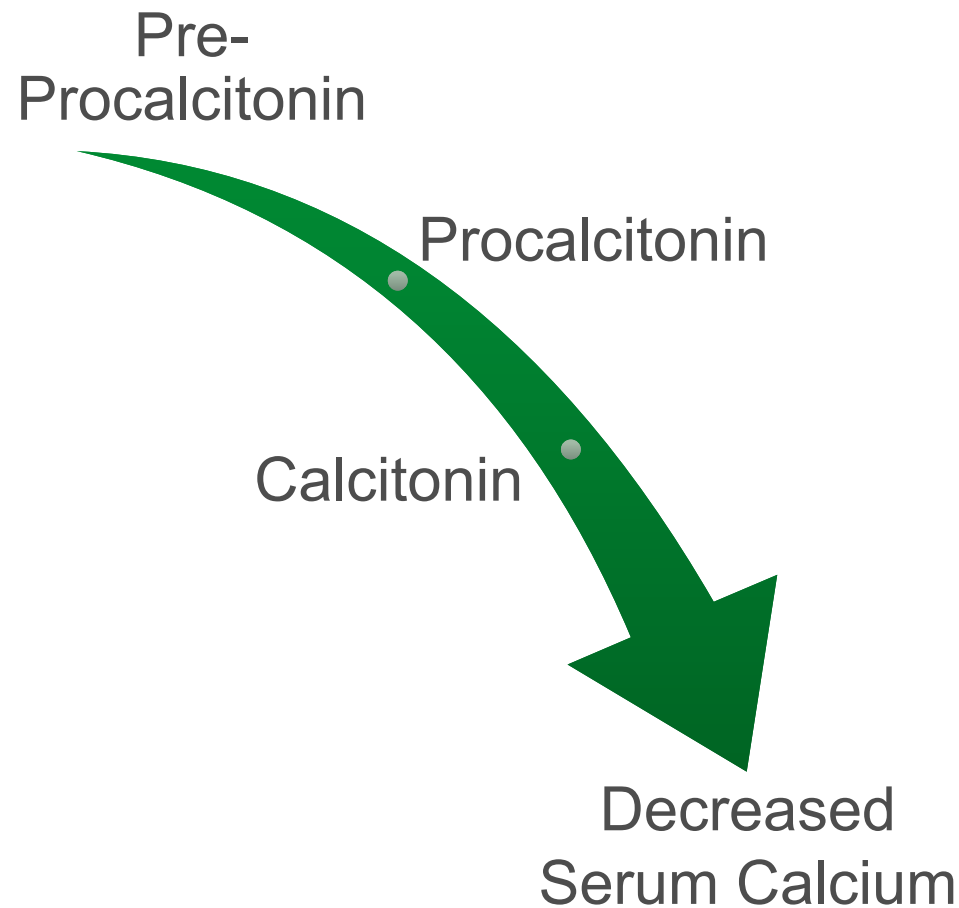


The speaker has no actual or potential conflict of interest in relation to this presentation

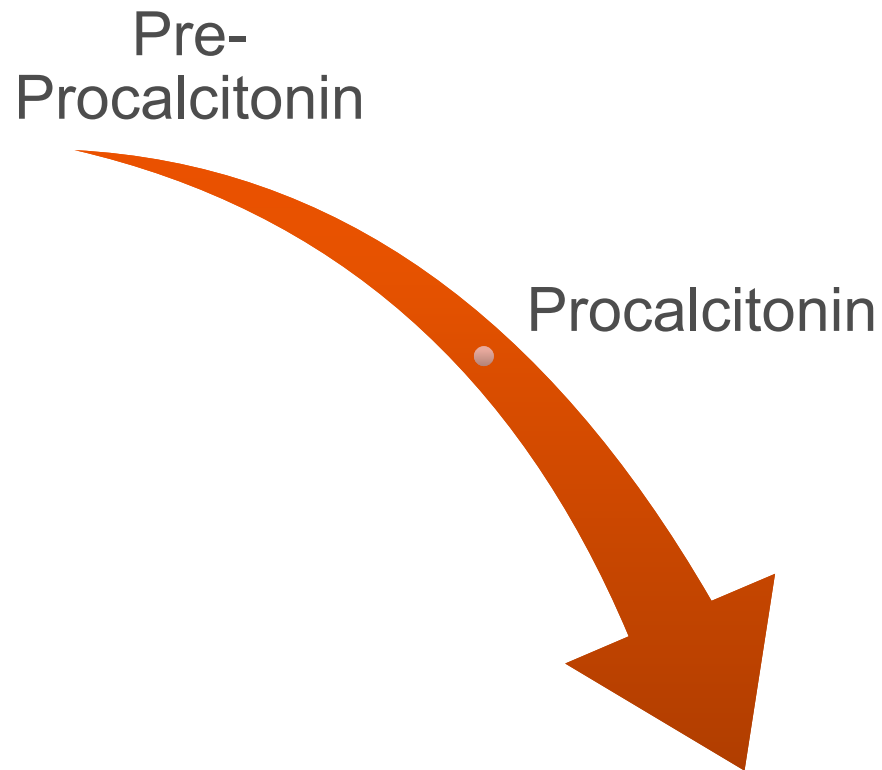
Procalcitonin Pathophysiology - Normal



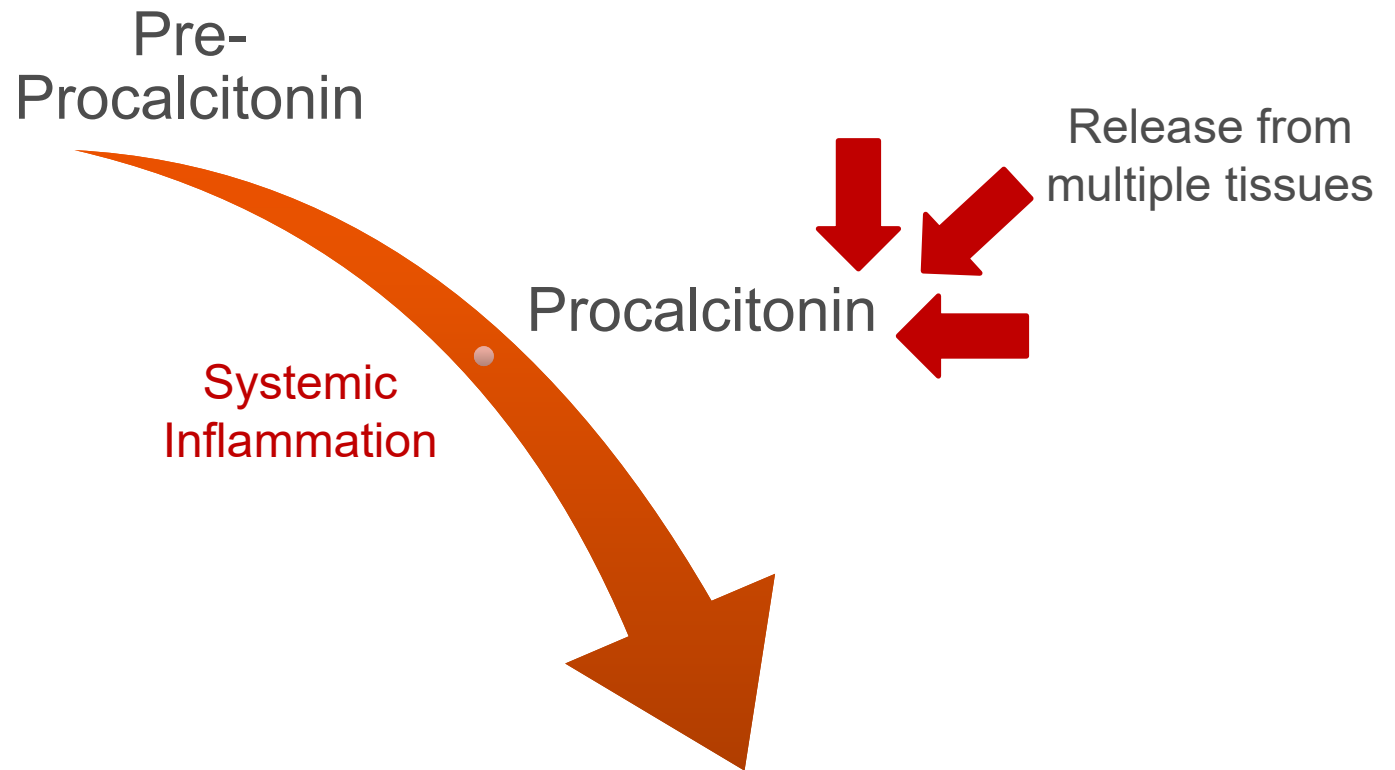
Procalcitonin Pathophysiology - Normal



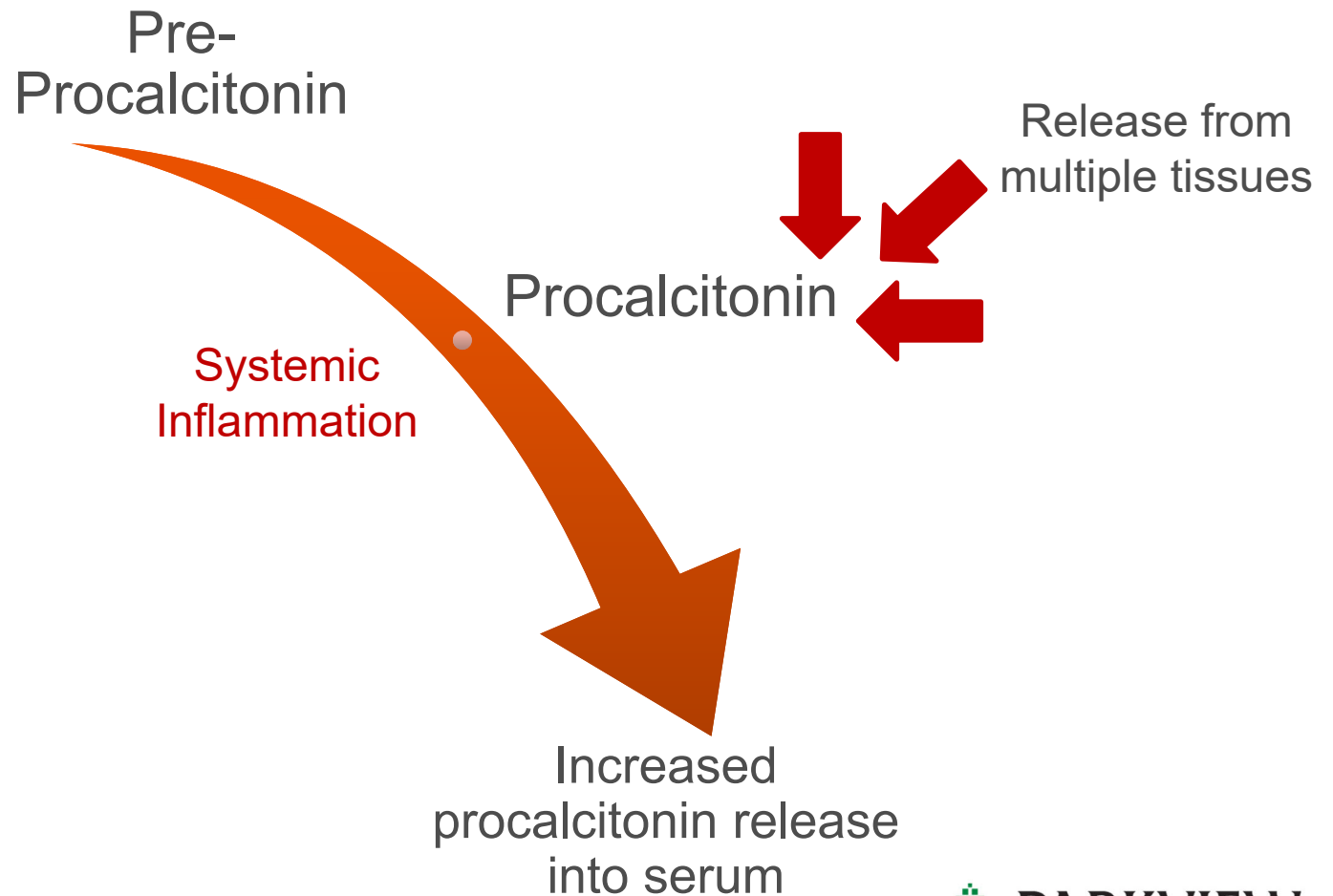
Procalcitonin Pathophysiology - Infection



Procalcitonin Pathophysiology - Infection



Procalcitonin Pathophysiology - Infection



Potential Causes of Procalcitonin Elevation

Infection

Surgery

Cardiac
Shock

Burns

Trauma

ESRD/HD

Assessment Question

The use of procalcitonin as a predictive biomarker has most extensively been studied in which disease states?

- A. Bacteremia and upper respiratory tract infection
- B. Sepsis and lower respiratory tract infection
- C. Cellulitis and sepsis
- D. Meningitis and UTI

Role in Therapy

- Bacterial infections
 - Sepsis
 - Septic shock
 - Lower respiratory tract infections (LRTI)
 - Community-acquired pneumonia (CAP)
 - Healthcare-associated pneumonia (HCAP)
 - Ventilator-associated pneumonia (VAP)

De Jong, et al.

- Prospective, multi-center, randomized, controlled trial
- Utilized non-binding advice
 - Discontinue antibiotics if procalcitonin decreased by 80% or more of its peak value or to 0.5 mcg/L or lower.
- Results
 - Median duration of treatment: 5 days vs. 7 days
 - Mortality at day 28: 20% vs. 25%

Lam, et al.

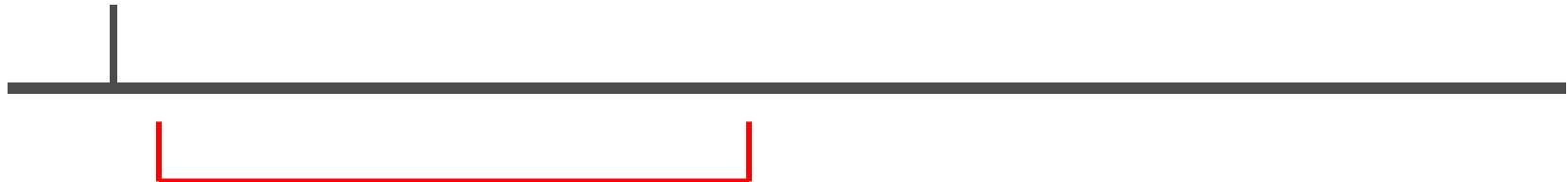
- Systematic review, meta-analysis
- Included 15 randomized controlled trials
- Results
 - Mortality: RR 0.93
 - Difference in antibiotic duration:
 - Cessation trials: - 1.26 days
 - Mixed strategy trials: - 3.10 days
 - No difference in hospital or ICU length of stay

Use of Pharmacist-Guided Protocols at Other Institutions

- Multiple institutions have instituted protocols for LRTI and sepsis
 - Nebraska Medicine and Washington Providence Medical Center
 - Align with other trials that have been done in this area

Procalcitonin Journey at Parkview

June 2016
Availability of
procalcitonin
assay

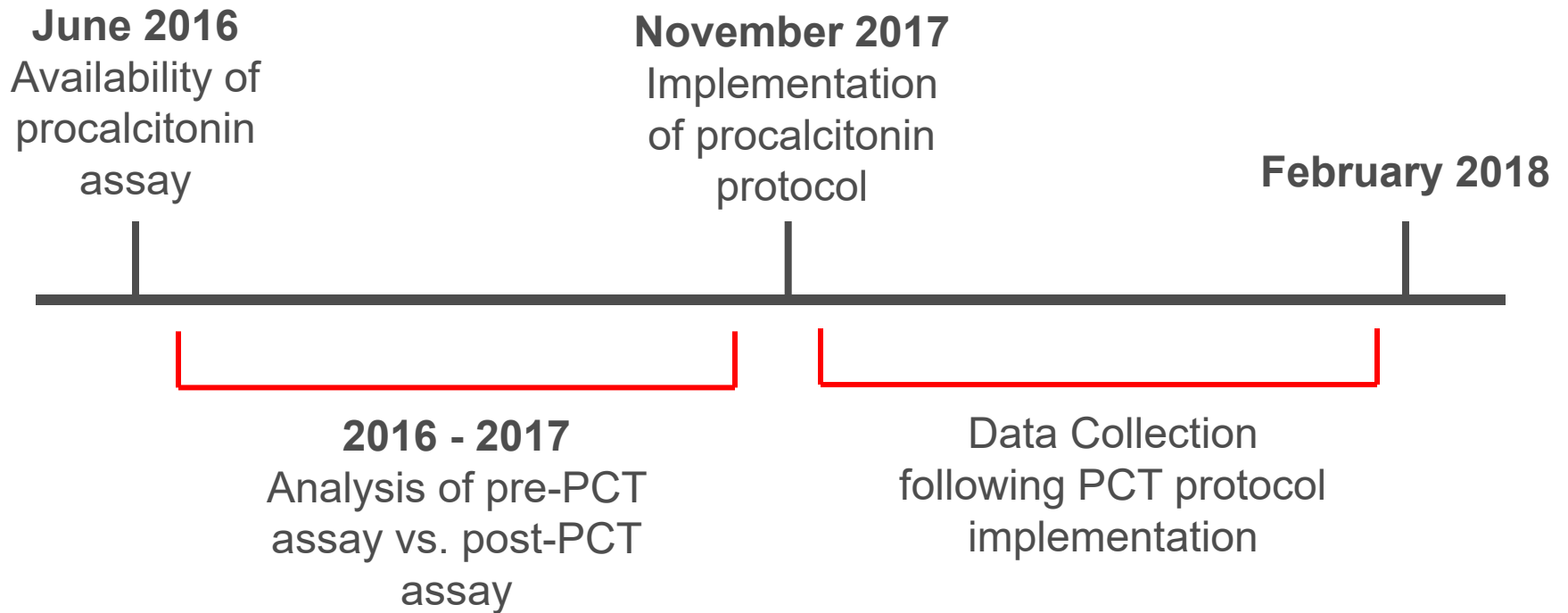


2016 - 2017
Analysis of pre-PCT
assay vs. post-PCT
assay

Procalcitonin Journey at Parkview

- Pre-PCT assay vs. Post-PCT assay analysis showed:
 - Decreased length of antibiotic therapy
 - Decreased hospital length of stay
 - Decreased ICU length of stay

Procalcitonin Journey at Parkview



Assessment Question

Which of the following could result from using procalcitonin levels to assist in guiding therapy?

- A. Increased cost
- B. Decreased need for physician monitoring of microbiology cultures
- C. Increased ICU length of stay
- D. Decreased length of antibiotic therapy

Purpose

- To evaluate patient outcomes following implementation of a pharmacist-guided procalcitonin protocol
 - Antibiotic length of therapy
 - Hospital length of stay
 - ICU length of stay

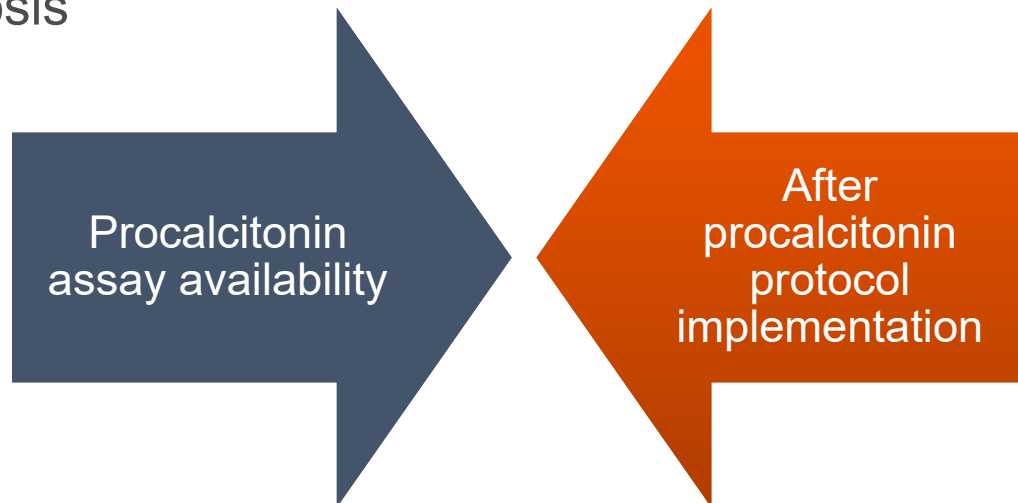
Parkview Regional Medical Center

- Parkview Health
- Community hospital
- Level II trauma center
- 6 intensive care units
- Included units
 - Medical ICU (MICU)
 - Surgical Trauma ICU (STICU)
 - Cardiovascular ICU (CVICU)
 - Cardiac ICU (CICU)
 - Progressive Care Unit



Study Design

- Retrospective chart review following protocol implementation
- Patients were matched on the following criteria:
 - Simplified Acute Physiology Score (SAPS II)
 - Age
 - Diagnosis



Protocol Implemented – Sepsis

PCT < 0.25



**Discontinuation
strongly
encouraged**

**PCT 0.25 – 0.49
Or
Peak decrease
by $\geq 80\%$**



**Discontinuation
encouraged**

**PCT ≥ 0.5
And
Peak decrease
by < 80%**



**Continuation
encouraged**

**PCT ≥ 0.5
And
Rising/not
decreasing**



**Continuation
strongly
encouraged**

Consider continuing therapy if clinically unstable or positive cultures with known source of infection

Protocol Implemented – LRTI

**PCT < 0.1
or
Peak decrease
by > 90%**



**Discontinuation
strongly
encouraged**

**PCT 0.1 – 0.24
Or
Peak decrease
by > 80%**



**Discontinuation
encouraged**

PCT \geq 0.25 - 0.5



**Continuation
encouraged**

PCT > 0.5



**Continuation
strongly
encouraged**

Consider continuing therapy if clinically unstable or positive cultures with known source of infection

Inclusion Criteria

- Age \geq 18 years old
- Diagnosis of sepsis, septic shock, CAP, HCAP or VAP
- Admitted to a critical care unit for \geq 48 hours
- Pharmacy-to-dose consult for management of antibiotics

Exclusion Criteria

- Death during admission
- Infectious Disease consult for antibiotic therapy management
- Known concomitant infection other than those listed in inclusion criteria
- Receiving antibiotics for > 24 hours prior to first drawn PCT level
- Positive culture or known source of infection

Objectives

- Primary Objective
 - Length of antibiotic therapy
- Secondary Objectives
 - Length of hospital stay
 - Length of ICU stay

Study Population

PCT Assay Availability

Included subjects from
assay availability group
(n = 137)

Matched subjects enrolled
into control arm
(n = 12)

Post-PCT Protocol Implementation

Subjects screened from
protocol documentation
(n = 23)

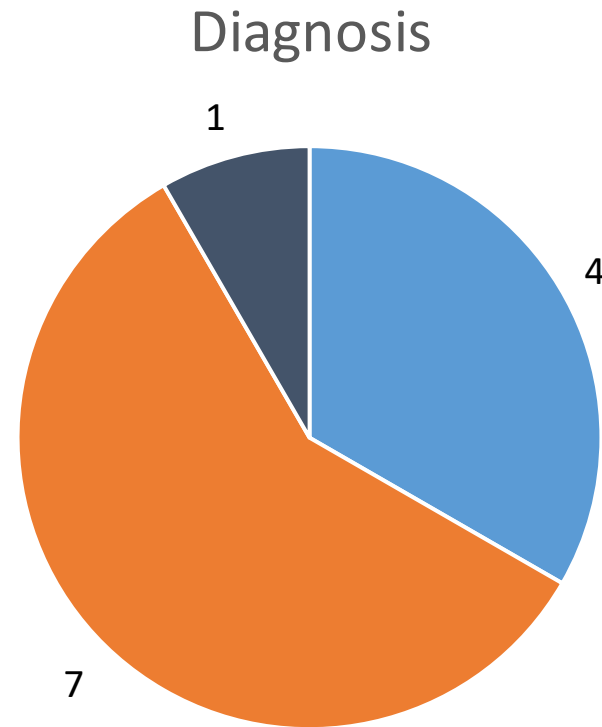
11 Excluded subjects
5 Died
2 Concomitant infection
2 ID consult
1 Antibiotics > 24h
before first PCT
1 Unmatched

Matched subjects enrolled
into treatment arm
(n = 12)

Baseline Characteristics

		PCT assay availability (n = 12)	Post-PCT protocol implementation (n = 12)
Age (years)	18-24:	0	0
	25-34:	0	0
	35-60:	5	5
	61-75:	4	4
	≥76:	3	3
Mean age (age ± SD)		62.8 ± 13.5	64.7 ± 15.1
SAPS II	≤29:	4	4
	30-40:	3	3
	41-52:	3	3
	53-64:	2	2
	≥65:	0	0
Mean SAPS II (score ± SD)		38 ± 13	38.2 ± 12.8

Baseline Characteristics



■ Sepsis ■ HCAP ■ CAP

Note: No patients matched were diagnosed with septic shock or VAP

Results – Primary Outcome

	PCT assay availability	Post-PCT protocol implementation	Significance P-value (95% CI)
Antibiotic length of therapy (days \pm SD)	10.2 \pm 5.7	5.2 \pm 3.7	0.020 (0.89, 9.16)

Results – Secondary Outcomes

	PCT assay availability	Post-PCT protocol implementation	Significance P-value (95% CI)
Hospital length of stay (days)	13.5 ± 5.8	12.1 ± 8.3	0.612 (-4.59, 7.59)
ICU length of stay (days)	6.2 ± 3.6	5.5 ± 3.8	0.659 (-2.46, 3.81)

Intervention Acceptance Rate

- Most interventions resulted in de-escalation with eventual discontinuation
- 2 resulted in immediate discontinuation
- 2 were denied without further attempts

Conclusions

- Pharmacy monitoring of PCT resulted in:
 - Decreased antibiotic therapy by 5 days
 - Decreased hospital length of stay by 1.4 days
 - Decreased ICU length of stay 0.8 days
- Pharmacists role
 - Monitoring and assessing PCT level trends
 - Making recommendations for discontinuation when necessary

Limitations

- Retrospective chart review
- Small sample size
- High exclusion rate
- Lack of consistent staffing schedule

Further Direction

- Continue data collection to obtain similar sample size to previous group
- Expand protocol use to other hospital areas
- Integrate PCT monitoring into daily work flow for clinical pharmacists
 - Potentially develop a monitoring flag on the electronic medical record
- Publication

Acknowledgements

- Jim Roy, PharmD, BCCCP
- Scott Girt, PharmD, BCPS
- Jared Netley, PharmD, BCPS
- Sarah Pfaehler, PharmD, MBA, BCPS

References

- Gilbert DN. Use of procalcitonin levels as an adjunct to clinical microbiology. *J Clin Microbiol.* 2010;48(7):2325-2329.
- Lam SW, Bauer SR, Fowler R, Duggal A. Systematic review and meta-analysis of procalcitonin-guidance versus usual care for antimicrobial management in critically ill patients: focus on subgroups based on antibiotic initiation, cessation or mixed strategies. *Crit Care Med.* 2018. doi: 10.1097/CCM.0000000000002953.
- Linscheid P, Seboek D, Nysten ES, et al. In vitro and in vivo calcitonin I gene expression in parenchymal cells: a novel product of human adipose tissue. *Endocrinology.* 2003;144(12):5578-84.
- De Jong E, Van Oers JA, Beishuizen A, Vos P, Vermeijden WJ, Haas LE, et al. Efficacy and safety of procalcitonin guidance in reducing the duration of antibiotic treatment in critically ill patients: a randomized, controlled, open-label trial. *Lancet Infect Dis.* 2016;16:819-27.
- Procalcitonin algorithm for guidance in antibiotic therapy decisions in respiratory tract infections and sepsis. Washington Providence Medical Center.
- Procalcitonin guidance. Nebraska Medicine.

Implementation of a Pharmacist-Guided Procalcitonin Protocol in a Community Hospital

Jasmine Coatie, PharmD

PGY1 Resident

Parkview Health

jasmine.coatie@parkview.com