Implementation of a Pharmacist-Driven Pain Management Service for Patients at High Risk for Respiratory Depression



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Background: Pain Management

Significant disparities exist in safe and effective pain management

Pain is subjective

Numerous classifications of pain

Multidisciplinary and multimodal pain management

Background: Opioid Analgesics & Safety

Opioid analgesics rank among the drugs most frequently associated with adverse drug events²

- Lack of knowledge about potency differences
- Improper prescribing and administration
- Inadequate monitoring

Drug	Equianalgesic Doses Parenteral (IV, IM, SQ)	Equianalgesic Doses [Oral]	Dosing Interval (hours)
Morphine	10 mg	30 mg	3 – 4
Fentanyl	0.1 mg	Transdermal 25 mcg/hr ≈ 45 mg of oral SR morphine	IV: 1 TD: 72
Hydrocodone		30 mg	3 – 4
Hydromorphone	1.5 mg	7.5 mg	3 – 4
Oxycodone		20 mg	4
Chycodone		201116	~
Tramadol		100 mg	4 – 6
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Tapentadol		100 mg	4 – 6

The Joint Commission Sentinel Event Alert #49

- Addresses safe use of opioids in hospitals
- Various patient populations at higher risk for development of respiratory depression
- Actions suggested Create and implement policies and procedures for the second-level review of pain management plans with high-risk opioids by pain specialists or pharmacists^{2,3}

Purpose

To evaluate the effect of an inpatient pharmacist-driven pain management service and opioid stewardship on:

- Amount and types of opioids prescribed
- Pharmacist interventions
- Overall patient outcomes and quality of life

Methods

Study Design



- Approved by Institutional Review Board (IRB)
- Single-centered, prospective study
- Data Collection: August 2015 March 2016
- Study Population: Control (n = 52); Intervention (n = 50)

Patient Selection and Enrollment

Inclusion Criteria	Exclusion Criteria
Age: Greater than or equal to 70 years	Age: Less than 18 years
Weight: Greater than or equal to 150 kg	Cancer diagnosis
SCr: Greater than or equal to 1.4 mg/dL	Decisionally impaired
Presence of sleep apnea	Intensive Care Unit (ICU) stay
Having undergone general anesthesia	Patient Controlled Analgesia (PCA)
Consult request, high severity of illness	

Intervention

Patient Assessment

 Daily computergenerated report of "high risk" patients

Follow-up

Evaluate effect(s) of interventions made

Medication Reconciliation

 Evaluation of pain medication regimens

Interdisciplinary pain medication therapy management

 Make appropriate, evidence-based recommendations

Outcome Measures

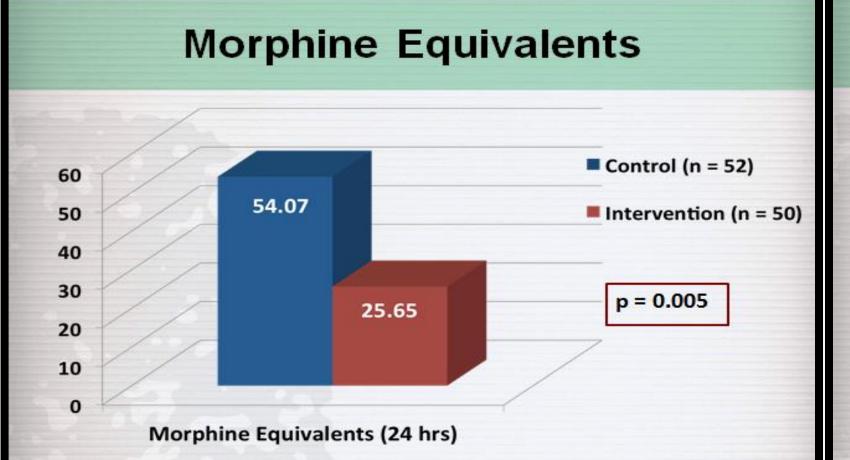
- **Primary Endpoint: Oral morphine equivalents**
- Secondary Endpoints: Number and types of opioids prescribed and administered; Pharmacist interventions; Length of stay

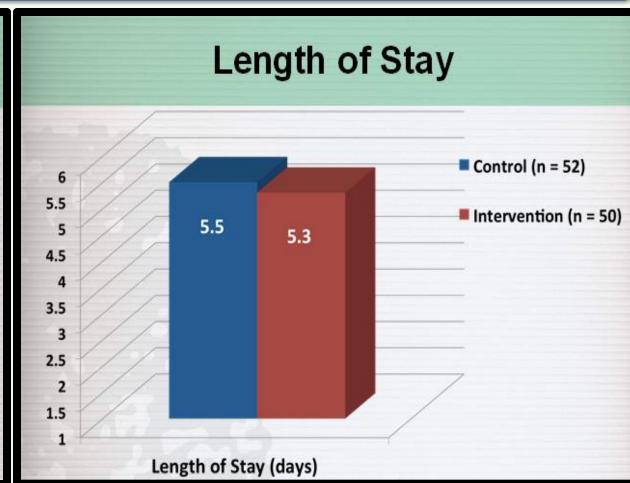
Results

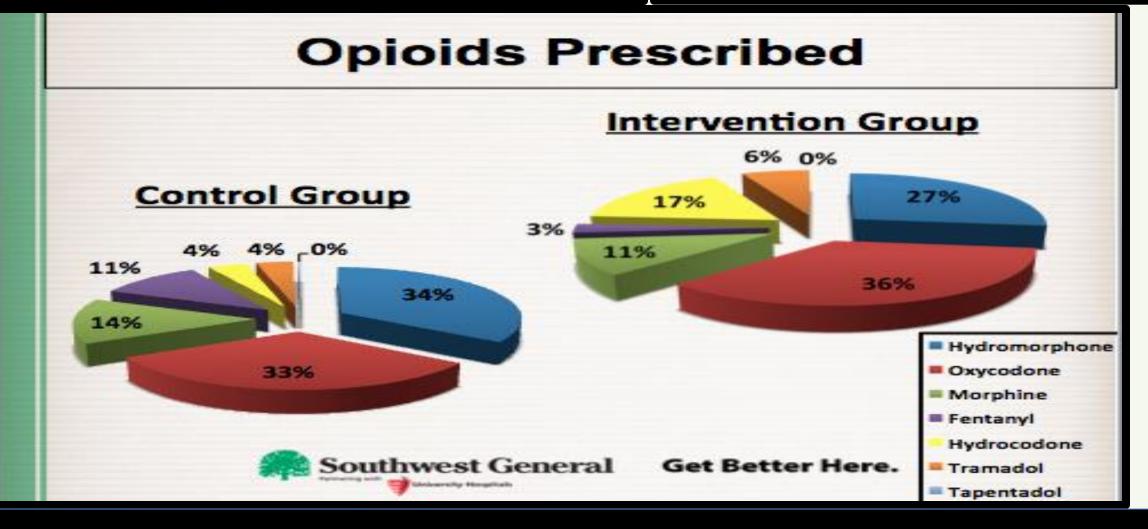
Baseline Characteristics

Patient Demographic	Control Group (n = 52)	Intervention Group (n = 50)
Age	69.65 years	62.98 years
Male Sex	50 % (26/52)	48% (26/50)
Weight (kg)	81.63	87.08
Serum Creatinine (SCr)	1.44	1.06
Sleep Apnea	12% (6/52)	24 % (12/50)
General Anesthesia	90% (47/52)	68 % (34/50)
Other	4% (2/52)	24% (12/50)
Total Number of Risk Factors	1.79	0.99

Primary and Secondary Endpoints







Moving Forward

- Failure Modes and Effects Analysis (FMEA)
- Hospital-wide capnography monitoring
- Opioid stewardship
- Pharmacogenomics testing
- **Collaborative Practice Agreements**

References

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