

Improving adherence to a severe alcohol withdrawal protocol through targeted order-set interventions Jessica Neal, Pharm.D., Jenni Catlin, Pharm.D. BCPS, Christina Stafford, Pharm.D.

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Background

Alcohol dependence accounts for approximately 16-31% of all intensive care unit (ICU) admissions. Patients who subsequently develop alcohol withdrawal syndrome are at an increased risk of sepsis, nosocomial infections, length of hospital stay, and mortality.² Currently, no evidencebased guidelines exist for the prevention or treatment of alcohol withdrawal in ICU patients.

Historically, benzodiazepines (BZD) have been used as the cornerstone of therapy for both treatment and prevention of alcohol withdrawal . BZDs have been shown to:

- Reduce the incidence of withdrawal seizures
- > Prevent the development of delirium tremens (DT)
- Mitigate symptoms of autonomic hyperactivity such as agitation or anxiety³

Previous research conducted at this institution reported inconsistent utilization of the current protocol, resulting in inappropriate use of dexmedetomidine (DEX) and underutilization of BZDs.⁴ Patients were advanced to DEX before an adequate trial of BZDs, which resulted in:

- > Increased length of ICU and overall hospital stay
- > Increased risk of over-sedation (defined as SAS score of 1 to 2)
- > Increased cost to hospital per patient
- > Inadequate therapy with BZDs while receiving DEX, potentially masking withdrawal symptoms

Purpose and Objectives

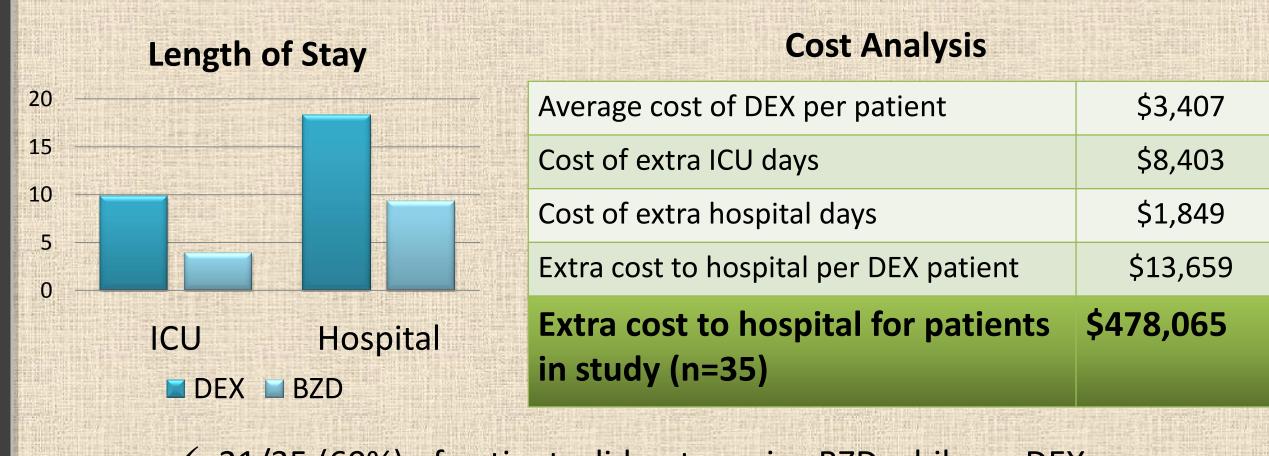
Purpose:

> Improve adherence to a severe alcohol withdrawal protocol by implementing (1) targeted orderset interventions, (2) enhanced electronic safeguards, and (3) healthcare provider education.

Objectives

- > Primary:
 - ✓ Protocol adherence
- > Secondary:
 - ✓ BZD, DEX, and phenobarbital use
 - ✓ ICU and hospital length of stay
 - ✓ Time at target SAS

Historical Results



- ✓ 21/35 (60%) of patients did not receive BZD while on DEX
- √ 6/35 (17%) of patients received DEX at rate > 1.5mcg/kg/hr
- ✓ 9/35 (26%) of patients on DEX had SAS scores of 1 to 2

Methods

- >A single-center, retrospective observational cohort study
- ➤ WIRB approval received on 11/16/2015
- ➤ Order-set available in CPOE 2/19/2016
- ➤ Phase 1 (pre-protocol) compared to Phase 2 (post-protocol) for primary and secondary objectives

Process for Revisions

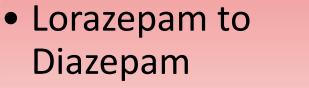
Evaluation of data from prior research conducted at study site

Literature search of medications used for EtOH withdrawal (efficacy & safety)

Collaboration with addiction medicine specialist and nurse educators for revisions

Finalized protocol submitted to appropriate committees for system-wide approval

Protocol Revisions



• Exception: hepatic impairment

BZD of Choice

New DEX Order-Set

comment

Ordering

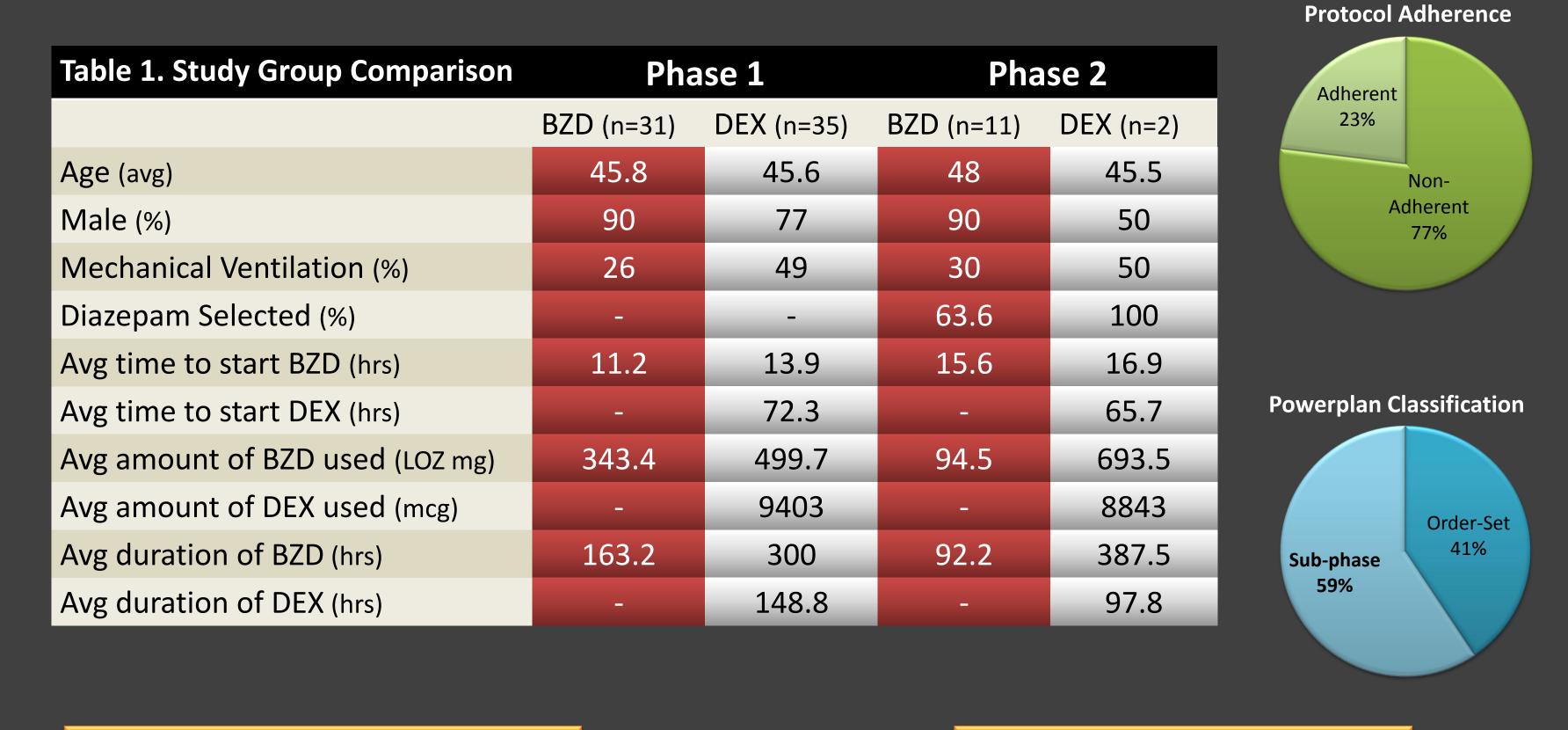
Scheduled BZDs

Nursing

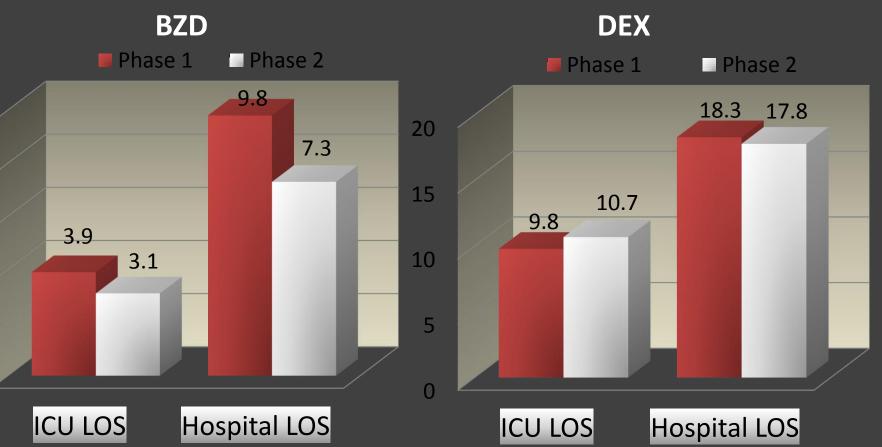
Providers Pharmacists

> System-wide Education

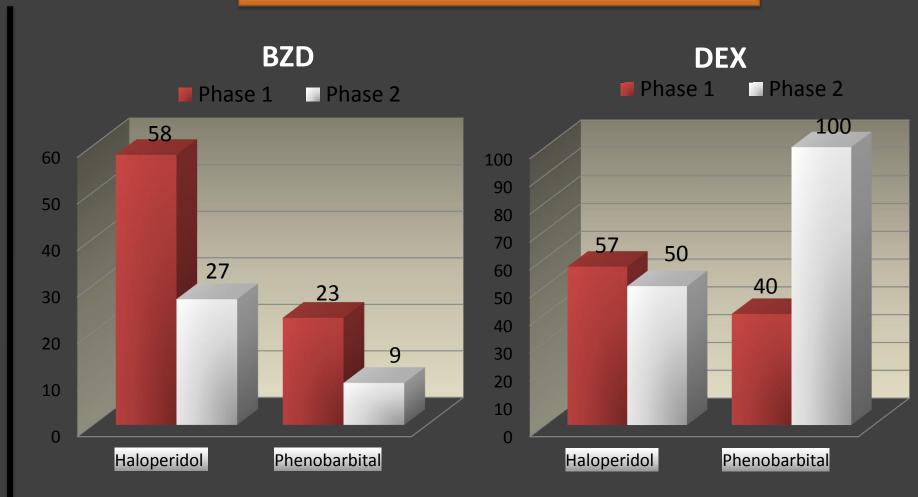
Results



ICU and Hospital Length of Stay



Use of Adjunctive Therapies



Conclusion

Areas for Improvement

- Increased time to BZD administration in both study groups
- Protocol dosing algorithm inconsistently followed (23% adherence)
- Subphase orders without parent power plan (41% correctly ordered)

Areas of

Success

NO rates of DEX >

1.5mcg/kg/hr

- ALL patients received concomitant BZD while on DEX
- ALL patients received phenobarbital prior to DEX
- Duration of DEX decreased by ~2 days on average

Future Direction

Earlier recognition of alcohol withdrawal

Improve phenobarbital availability for repeat administration

Education: adjunctive therapies, protocol ordering, appropriate use of DEX

Discussion

Limitations:

- Small number of patients
- Observational study
- > Time constraints
- Confounding variables

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Disclosures

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.

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