Characterization of Psychotropic PRN Medications in a Psychiatric ICU



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Background

- Pro re nata (PRN) antipsychotics and benzodiazepines routinely used to rapidly stabilize acutely agitated patients¹
- Limited evidence to guide pharmacological treatment²
- Non-pharmacological de-escalation strategies should be considered before administering medications³
- Incomplete documentation of efficacy and safety of PRN administration¹
- PRN antipsychotics increase risk of adverse effects secondary to polypharmacy and high daily dose exposure⁴
- PRN benzodiazepines have potential for tolerance, sedation and dependence³

Objectives

Primary:

- Describe the prescribed medication, dosing range, route, frequency, indication, and documentation practices for PRN psychotropic medications
- Secondary:
- Describe non-pharmacological interventions used for acute stabilization
- Describe whether total daily PRN dose of antipsychotic drugs is within Health Canada (HC) limits
- Describe adverse effects and events

Methods

- Design: Retrospective cohort chart review
- Inclusion criteria:
 - Age ≥18 years
 - Admitted to psychiatric intensive care unit (ICU) at Surrey
 Memorial Hospital (SMH) between June and September 2018
 - Received PRN antipsychotics and benzodiazepines
- Exclusion criteria:
- Admitted to SMH Emergency Department (ED) at any point during psychiatric ICU admission
- Sample size:
 - Convenience sample, selected in reverse chronological order
- Definitions:
- Documentation: within 4 hours of dose given in nursing notes
- Adverse effect: Naranjo scale (documented if ≥4 "possible")
- Adverse event: unintended patient harm or staff harm occurring within 4 hours prior to administration of PRN medication

Results

Table 1: Patient Characteristics	
	N=32
Male, n (%)	22 (69)
Age (years), median (IQR)	34 (27-50)
Psychiatric ICU length of stay (days), median (IQR)	14 (9-20)
Admitting Diagnosis, n (%)	
Schizoaffective related disorder	11 (34)
Schizophrenia	8 (25)
Bipolar 1 disorder	4 (13)
Drug induced psychosis	3 (9)
Other	6 (19)
Illicit substance use, n (%)	19 (59)
Adverse events*, n (%)	6 (19)
Total number of events	14

Table 2: Characteristics of Psychotropic PRN Orders		
Total number of psychotropic PRN orders	N=123	
Number of PRN orders per patient, median (IQR)	3 (2-3)	
24 hour maximum prescribed, n (%)	121 (98)	
Frequency prescribed, n (%)	119 (97)	
Prescriber indication, n (%)	41 (33)	
Agitation and/or anxiety	25 (20)	
Agitation/anxiety or insomnia	7 (6)	
If refuse regular medication	5 (4)	
Other	4 (3)	
Antipsychotic PRN orders exceeding total daily HC	10 (8)	
dose limits, n (%)		

Table 3: Characteristics of Psychotropic PRN Administrations		
Total number of psychotropic PRN administrations	N=1179	
Number of PRNs administered per patient, median (IQR)	27 (12-53)	
PRNs documented for effectiveness, n (%)	795 (67)	
PRNs documented for nursing rationale, n (%)	1164 (99)	
Non-pharmacological method documented prior to PRN	559 (47)	
administration, n (%)		
Two or more techniques	161 (14)	
Redirection	119 (10)	
Clear mutual expectations	92 (8)	
Education/teaching	61 (5)	
Reassurance	41 (3)	
Listening techniques	37 (3)	
Other	48 (4)	

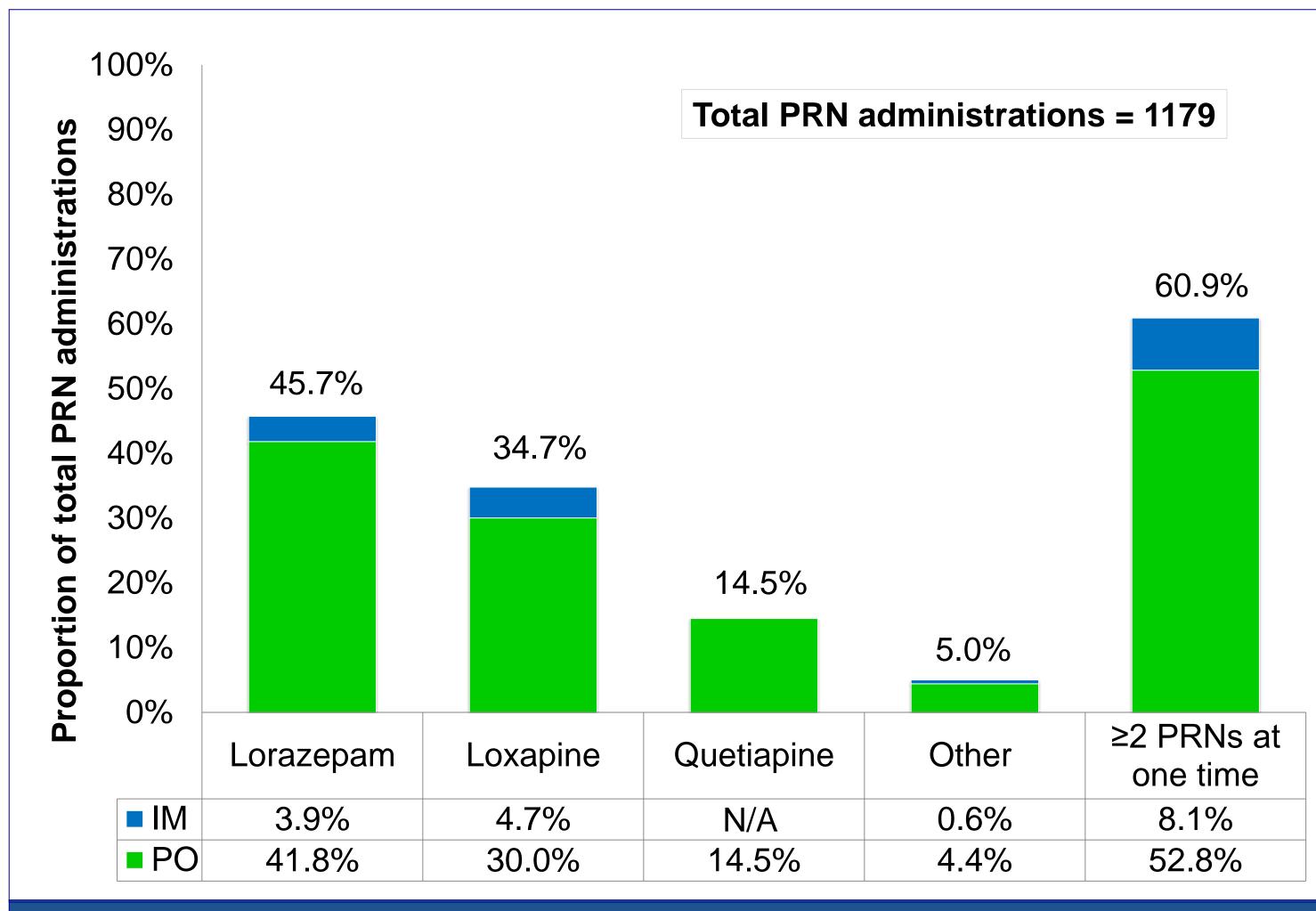


Figure 1: Proportion of Psychotropic PRN Administrations

Limitations

- Single centre, retrospective design
- Total daily dose that included regular antipsychotic administrations in addition to PRN antipsychotics was not assessed
- De-escalation strategies that did not result in PRN administration were not assessed
- Exclusion of patients who were admitted to ED may have led to missed adverse effects

Conclusions

- PRN psychotropic medications administered throughout the psychiatric ICU admission
- There is potential to cause harm due to exceeding 24 hour Health Canada dose maximums for antipsychotics
- Non-pharmacological de-escalation methods and documentation of effectiveness following PRN administration could be optimized
- Further education needed regarding the benefits of prescribing specific indications with each PRN order
- Further studies are needed in assessing appropriateness of psychotropic PRN medications

References

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