



Northeast Georgia Medical Center

BACKGROUND

- Dexmedetomidine is commonly utilized in the intensive care unit for sedation, although optimal dosing and duration of use are unknown
- Literature supports higher doses and longer duration than the FDA label of < 24 hours at a maximum infusion rate of 1 mcg/kg/hr
- Emerging studies suggest hepatic dysfunction and obesity may impact dexmedetomidine pharmacokinetics, but clinical application has yet to be established

OBJECTIVE

• To evaluate the utilization of dexmedetomidine among critically ill patients

METHODS

Retrospective, single center chart review of adult patients who received dexmedetomidine in the medical or pulmonary intensive care unit at Northeast Georgia Medical Center, a 557-bed community teaching hospital from January 1, 2022, to July 31, 2022



EVALUATING THE USE OF DEXMEDETOMIDINE IN CRITICALLY ILL PATIENTS AT A COMMUNITY TEACHING HOSPITAL

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RESULTS

Table 1. Baseline Characteristics

	N=100
Age (years)	63 (24)
Male	55 (55%)
Body Mass Index (kg/m ²) Non-obese (< 30) Obesity Class I (\geq 30 to 34.9) Obesity Class II (\geq 35 to 39.9) Obesity Class III (\geq 40)	62 (62%) 15 (15%) 13 (13%) 10 (10%)
Liver dysfunction Documented liver cirrhosis Total bilirubin ≥ 1	28 (28%) 9 (9%) 27 (27%)
Target Richmond Agitation Sedation Scale (RASS) goal -2 to +1 -3 to -5 Other	92 (92%) 2 (2%) 6 (6%)
Mechanical ventilation	56 (56%)
Use of concurrent sedation Propofol Fentanyl Benzodiazepine Ketamine	58 (58%) 31 (53%) 57 (98%) 16 (28%) 5 (9%)
Use of concurrent continuous paralytic Values reported as number (%) and median (interquartile	4 (4%)

Figure 1. Duration of Dexmedetomidine Infusion



RESULTS (continued)

Table 2. Dosing Characteristics

Dose (mcg/kg/hr) Initial dose (mcg/kg/hr) Maximum dose (mcg/kg Volume per day (mL)

Cumulative volume (mL) Values reported as median (interguartile range)

Figure 2. Dosing in Specific Patient Populations

- prolonged infusions (> 7 days)
- clinical application of these findings

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The authors of this presentation have nothing to disclose concerning possible financial or personal relationships with commercial entities.

	N=100
	0.56 (0.62)
	0.20 (0.30)
g∕hr)	1.50 (0.65)
	247 (290)
	599 (1150)

CONCLUSIONS

• Overall utilization of dexmedetomidine (dosing, indication, duration, target RASS goal) was appropriate

• Potential to educate providers regarding inability to achieve deep levels of sedation, inappropriate use with continuous paralytic therapy, and risks associated with

• Patients with liver dysfunction or BMI \geq 30 required higher doses, but more studies are needed to determine

REFERENCES

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4. Hughes, C.G, et al. MENDS2 investigators. Dexmedetomidine or propofol for sedation in mechanically ventilated adults with