

Dexmedetomidine in autonomic hyperactivity states: management of two benzodiazepine-refractory cases in Neurology Unit

Andrea Porsio, Marco Zoccarato, Carla D'Ascenzo, Francesco Rossato, Luca Santelli and Anna Maria Basile
Department of Neurology, Sant'Antonio Hospital, Padua

Dexmedetomidine

Delirium tremens

Paroxysmal Sympathetic Hyperactivity

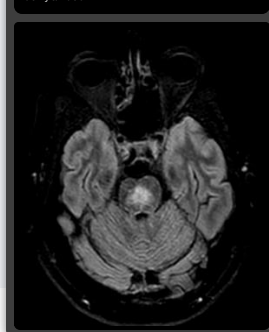
BACKGROUND

- Autonomic hyperactivity states, such as **delirium tremens (DT)** and **paroxysmal sympathetic hyperactivity (PSH)** can be refractory to benzodiazepines (BDZs) and standard sedatives^{1,2}.
- Dexmedetomidine (DEX) is a **high affinity selective α_2 -agonist**, providing powerful sympatholytic action, mild sedation and minimal respiratory depression^{1,2}

CASE 1 — DELIRIUM TREMENS (33 y M)

- DT after alcohol-withdrawal seizure → severe agitation, hallucinations, tremors, HR \approx 150–170 bpm and moderate hypertension (\approx 160/90 mmHg)
- Refractory to very high-dose BDZs** (over 100mg of IV diazepam in 2h), and phenobarbital and tiapride.
- DEX 0.7 μ g/kg/h → **rapid control, arousable sedation without intubation**
- Relapse during a one-day taper → re-start and wean over 3 days;
- MRI: pontine T2/FLAIR hyperintensity** consistent with prior osmotic demyelination (FIGURE 1).

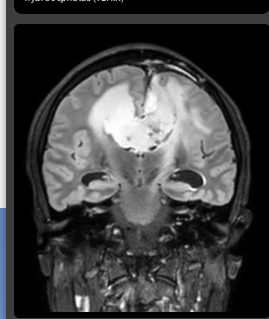
FIGURE 1 — T2/FLAIR central pontine osmotic demyelination



CASE 2 — PSH DUE TO BRAIN TUMOR (22 y M)

- WHO grade-4 callosal, bifrontal astrocytoma with hydrocephalus (FIGURE 2)
- Recurrent PSH paroxysms** (diffuse asymmetric tremors, tachycardia, hypertension, hyperthermia, dystonic posturing); EEG excluded underlying epileptic activity.
- Recurrence despite a combination of anti-seizure medications (ASMs), steroids, maximal anti-edema therapy, β -blockers, and high-dose of BDZs.
- DEX 0.7 → 1.4 μ g/kg/h achieved control.
- Attempts $<$ 0.5 μ g/kg/h → recurrence aborted by 10-minute boluses of 0.5–1 μ g/kg.
- No respiratory or hemodynamic instability observed.

FIGURE 2 — bifrontal callosal astrocytoma with hydrocephalus (FLAIR)



PRACTICAL DOSING

- Start 0.2–0.7 μ g/kg/h, titrate to effect (commonly 0.4–1.4 μ g/kg/h).
- Target cooperative/arousable sedation.
- Optional rescue bolus 0.5–1 μ g/kg over 10 minutes for breakthrough sympathetic crises.
- Once stable, avoid abrupt down-titration—taper gradually to reduce relapse/rebound risk with α_2 -agonists.
- Concurrent β -blockers use**—particularly β_1 -selective agents (e.g., esmolol)—**is generally safe** when used as needed, with monitoring for bradycardia and hypotension

CONCLUSIONS & KEY MESSAGES

- Sympatholytic action and **arousable sedation** → **no respiratory depression** → **facilitates neuro exam**
- Probably superior to benzodiazepines** for autonomic storms, yet non-etiological—appropriate as adjunctive therapy, **not as standalone**.
- Fast titration for paroxysmal storms; **useful outside ICU** when intubation is not indicated/desired.
- Although labeled for ICU sedation, DEX's manageability and minimal respiratory depression make it feasible in high-dependency/step-down units.**

REFERENCES

- Borgu ndvaag B, Belloio F, Miles I, et al. Guidelines for Reasonable and Appropriate Care in the Emergency Department (GRADE-4): Alcohol use disorder and cannabinoid hyperemesis syndrome management in the emergency department. Acad Emerg Med. 2024;31(5):425-455. doi:10.1111/acem.14911
- Meyfroidt G, Baguley JJ, Menon DK. Paroxysmal sympathetic hyperactivity: the storm after acute brain injury. Lancet Neurol. 2017;16(9):721-729. doi:10.1016/S1474-4422(17)30259-4

Contact: andrea.porsio@aopd.veneto.it



24-28 Ottobre 2025
Padova Congress

55° CONGRESSO
SOCIETÀ ITALIANA
DI NEUROLOGIA