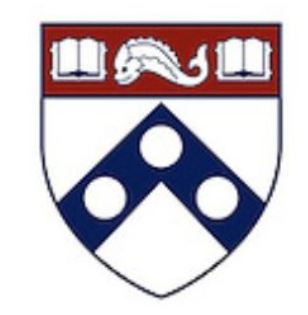


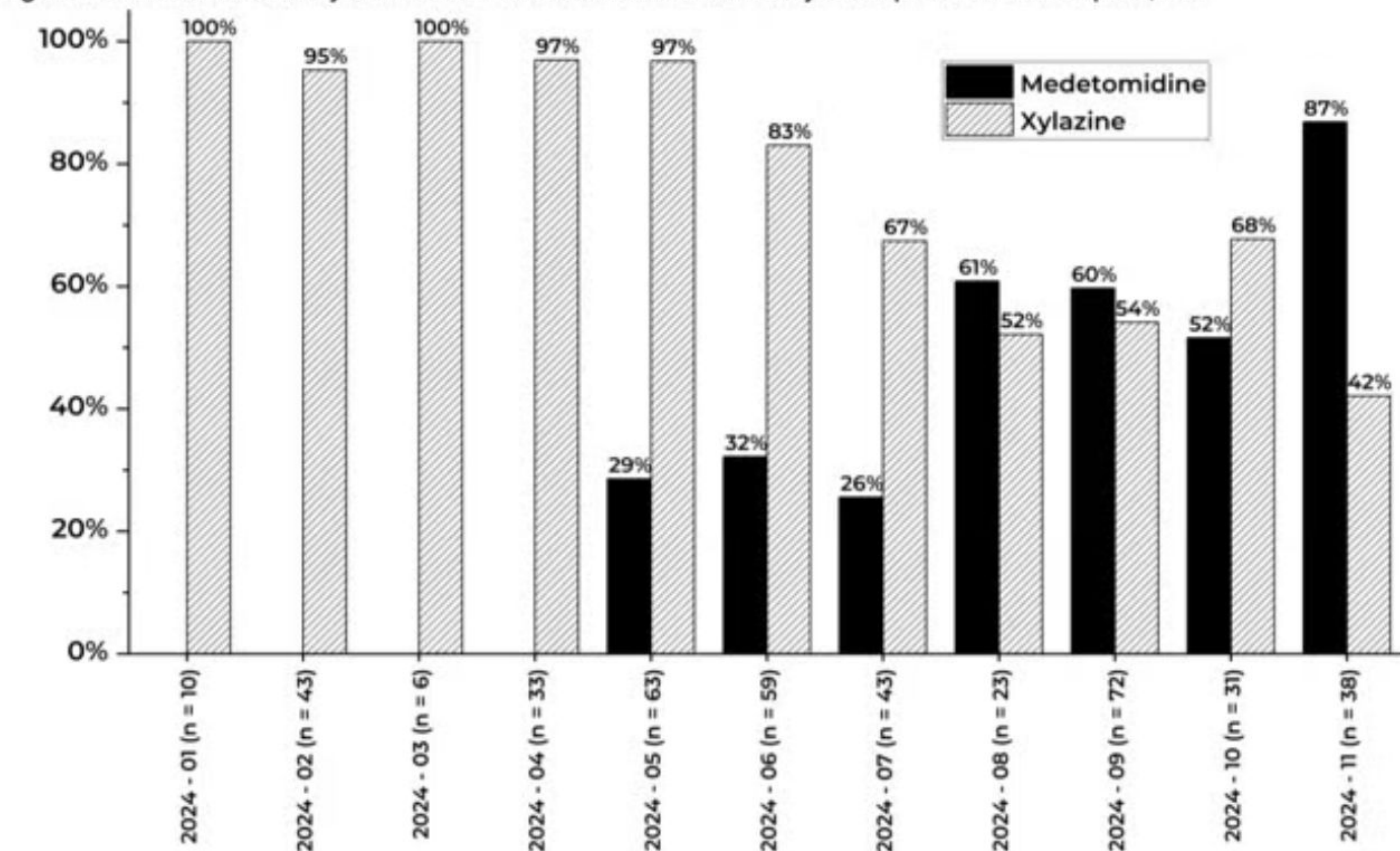


Withdrawal From Medetomidine-Adulterated Fentanyl During Pregnancy: A Case Report

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- Aka “rhino tranq” : α 2-adrenergic agonist racemic mixture of dexmedetomidine and levomedetomidine (inactive)
- Highly selective and potent α 2 agonist (100–200× more potent than xylazine)
- 1 bag of “fentanyl” ~ 15mg of medetomidine (20 infusion vials of dexmedetomidine (400mcg/100mL))
- Toxidrome: naloxone-resistant sedation, sinus bradycardia, hypotension
- Withdrawal: severe sympathetic hyperactivity
 - rapid onset hypertension with systolics >180 mmHg
 - tachycardia
 - vomiting
 - hypoactive delirium
 - tremor without clonus
- Withdrawal often requires ICU admission for severe autonomic hyperactivity leading to end-organ damage including ACS, PRES, and seizures.

Figure 1: Prevalence of Xylazine and Medetomidine in Fentanyl Samples in Philadelphia, PA



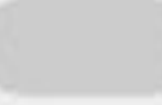
Data source: Center for Forensic Science, Research and Education, PA Groundhogs, Philadelphia Department of Public Health

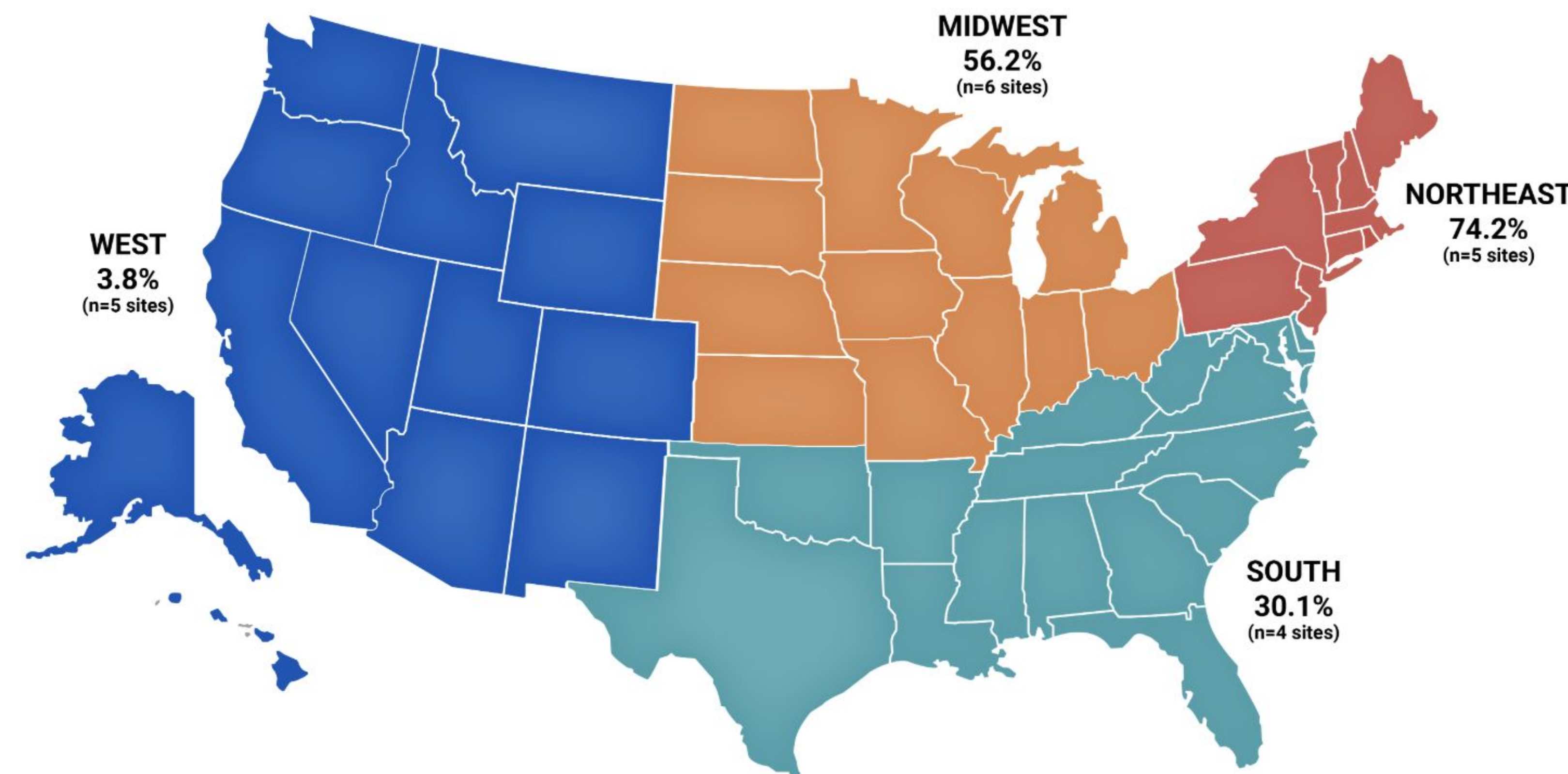
*in 2024, PA made xylazine a scheduled drug

32yo G5P3013 at 39w4d with hypertension, MRSA bacteremia, epilepsy, HIV, HCV, HSV, depression, anxiety, 4 prior cesarean deliveries (CDs), and polysubstance use disorder (opioids, benzos, alcohol, amphetamines) on methadone, presented w/ painful contractions.

Her last use of fentanyl and xylazine was 2 hours prior.

	Day 0 (preop)			Intraop (GA w/ TAP block)			Extubated to NC SICU Day 1			Day 2		
	Anes	Ob	Add. Med / Psych	Anes	Ob	Add. Med / Psych	Anes / ICU	Ob	Add. Med / Psych	Anes / ICU	Ob	Add. Med / Psych
Methadone	40mg PO qd						70mg PO qd (total)			90mg PO qd (total)		
Hydromorphone	4mg IV q2 PRN			2mg IV q1 PRN			4mg IV q2					
Oxycodone	40mg PO q4											
Clonidine	0.6mg PO + 0.4mg q6											
Dexmedetomidine				1 mcg/kg IV + 2.5 mcg/kg/hr (max)			Wean					
Diazepam	10mg IV x1											
Clonazepam	0.5 - 1mg PO q4 PRN											
Ketamine				0.2 mg/kg/hr IV			Wean infusion			0.375 mg/kg PO q6		

 managed by
 start or dose change
 continued same dose

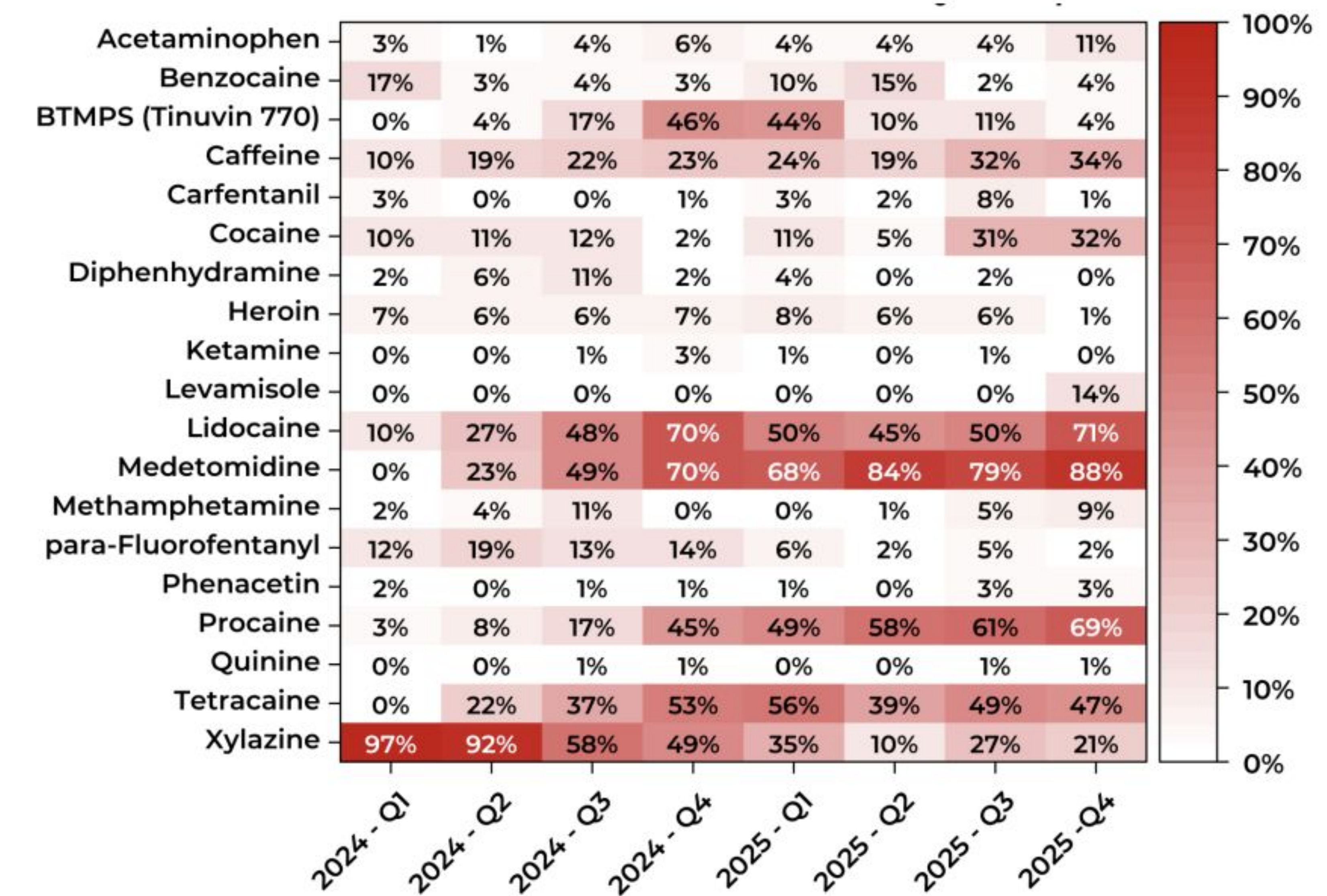


Percentage of opioid-positive drug product and paraphernalia samples positive for medetomidine across 20 sentinel sites: US region, July 2025 – December 2025

- Multidisciplinary coordination is essential and should involve Obstetrics, Anesthesiology, Addiction Medicine, Neonatology, Infectious Disease, Psychiatry, and Bioethics Services
- There is limited evidence-based guidance regarding management of medetomidine withdrawal in parturients.
- Our institution is working on better coordination with nursing and pharmacy to allow dexmedetomidine infusions on acute care floors.
- Clinicians need to be vigilant of the constantly-evolving landscape of novel and synthetic additives to the opioid supply

- Medetomidine-laced fentanyl is on the rise, and withdrawal syndromes are growing increasingly complex. The percentage of fentanyl with medetomidine has been increasing continuously over the past two years and is now in nearly every tested sample.
- Failure to recognize $\alpha 2$ -agonist exposure may result in misdiagnosis and suboptimal management.
- Withdrawal management includes $\alpha 2$ -agonist substitution or bridging therapy, such as dexmedetomidine or clonidine, balanced against risks of hypotension and bradycardia.

FIGURE 5: COOCCURRENCE OF (Y) IN FENTANYL OVER TIME



Selected co-occurring analytes shown from GC-MS analysis of fentanyl-primary samples in Mid-Atlantic 2024 - 25