Pharmacotherapy Options for Alcohol Withdrawal

	Benzodiazepines					Gabapentin			
Usage	Inpatient management is recommended for all pregnant patients. However, if inpatient service is not an option due to patient preference or lack of availability, outpatient treatment with close daily monitoring may be offered to those who are at low risk of severe complications. See outpatient resource for details.								
	PAWSS ≥ 4 or	patient is hig	h risk for seve	ere withdrawal	PAWSS < 4 or pat	tient is low risk f	or severe withdra	wal complications.	
Dosing	Prescriptions should be short-term and tapered. Adjust doses daily for outpatients or more frequently for inpatients (q 1h) based on					For immediate-release tablets			
					ired, consult an	Symptoms	Regular Dose	PRN	HS
	addiction me	•	ist and consid	ler inpatient ac	dmission.	If CIWA-Ar is 10- 14 or SAWS ≥ 12	300mg TID. Titrate up to 600mg TID if symptoms are not responding	300mg PRN - leave 2 hrs btwn regular and PRN doses	300-600mg HS PRN
		Day 1	Day 2	Day 3	Day 4				
	Symptom- based (in or	5-10mg q4-6 h	5-10mg q6-8 h	5-10mg q12 h	5-10mg HS	If CIWA-Ar is < 10 or SAWS < 12		300mg q4 h PRN	300-600mg HS PRN
	outpatient) Fixed schedule for outpatients	5-10mg QID	5-10mg TID	5-10mg BID	5-10mg HS	When acute symptoms resolve and CIWA < 10 or SAWS < 12 consistently (e.g., 3 measurements), taper over 3-5 days, reducing dose by 600mg each day Max daily dose is 3600mg Hold doses if patient shows drowsiness, ataxia, or slurred speech			
	Day 1-2 Day 3-4	ate 3rd trimes octentially less 1-2mg q4 h 0.5-1mg q4 h	s impact on ne		·			· · ·	·
	If medication to q2d.	is needed be	yond day 4, co	ontinue to tape	er dose down				
Contra-indications	1. Severe respiratory insufficiency 2. Sleep apnea 3. Myasthenia gravis 4. Narrow angle glaucoma 5. History of allergy or hypersensitivity					1. History of allergy or hypersensitivity			

	Benzodiazepines	Gabapentin					
Cautions	1. Hepatic impairment 2. Renal impairment	1. Renal impairment					
Safety	A careful assessment of benefits of medication vs. risks of continued alcohol use should inform decision-making. Frequent monitoring of fetus or infant is advised.						
	If used with alcohol, can lead to serious safety risks, incl. over sedation, falls, delirium, respiratory depression	If used with alcohol, risk of additive CNS-depressive effects					
	Pregnancy Controversial evidence to suggest an increased risk for cleft lip and palate and "floppy infant syndrome" in human studies. Risk of neonatal withdrawal syndrome Breastfeeding Present in breastmilk. Can cause sedation and inability to suckle. Shorter-acting lorazepam preferred and has better outcomes than diazepam. Monitor infant for drowsiness, decreased feeding, or low weight gain	Pregnancy Limited data indicates minimal adverse effects Breastfeeding No adverse effects reported. Monitor infant for drowsiness, low weight gain, gastrointestinal side effects, and developmental milestones					
Side Effects	Drowsiness, dizziness	Doses greater than avg therapeutic levels may cause ataxia, slurred speech, or drowsiness					
Other Considerations	Risk for non-medical use, diversion, and dependence. Risk for drugdrug interactions leading to excess sedation, impaired psychomotor and cognitive functioning Lorazepam is the preferred option for patients with advanced liver disease Use cautiously in outpatients. Consider blister packing or daily dispensing	Risk for non-medical use, diversion, and dependence Toxicity profile parallels that of alcohol Easy to transition from WDM to long-term relapse prevention					
	Thiamine: Thiamine deficiency is common. For patients planning withdrawal management, offer 100-200 mg oral or parenteral thiamine daily. For patients with suspected Wernicke's encephalopathy, offer 200-300 mg parenteral thiamine for 3-5 days followed by oral therapy.						

Abbreviations: WDM – withdrawal management, PRN – as needed/when necessary, QID – four times per day, TID – three times per day, BID – two times per day, HS – at bedtime