Acentra

HEALTH

WEST VIRGINIA

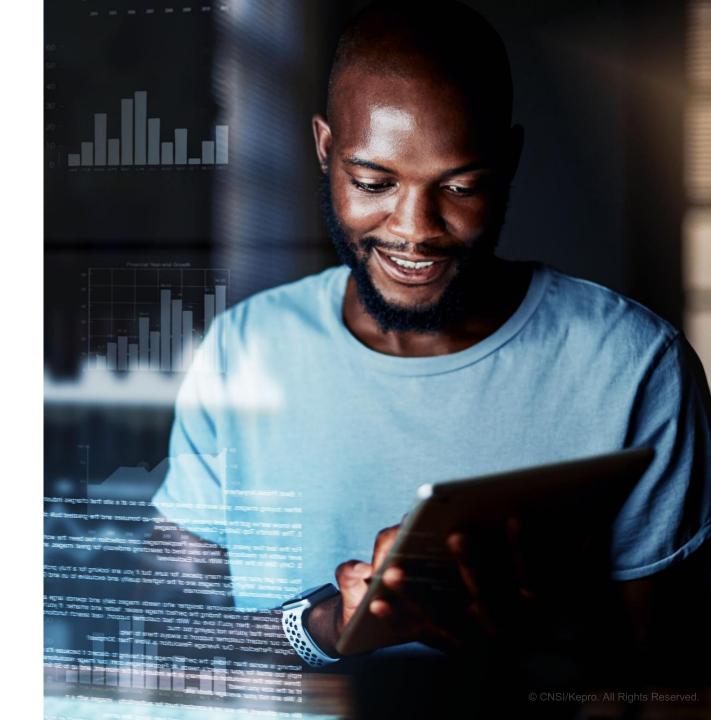
DRUG UTILIZATION REVIEW
BOARD MEETING

September 24, 2025





- Rachele Poissant, PharmD
 Clinical Account Manager
- Cory Chambliss
 Operations Director
- Scott Donald, PharmD
 Director of Clinical Services



	Target Intervention	Profiles Reviewed	Letters Sent	Response Rate
Apr	Antihyperlipidemic Drug Interactions	605	642	7%
May	Increased Risk of Serotonin Syndrome	602	734	4%
Jun	Increased Risk for QT Prolongation	698	677	4%



April 2025:

Antihyperlipidemic Drug Interactions

Drugs included:

Statins

Fibric Acid Derivatives

Bile-Acid Resins

Contraceptives

Antacids

Macrolides

Azole Antifungals

Anticonvulsants

Calcineurin Inhibitors

Protease Inhibitors

Antiarrhythmics

Anticoagulants

Strong CYP3A4

Inhibitors

 Coadministration of [Selected Drug(s)] should be done with caution. [Selected drug] can inhibit the metabolism of [selected drug] thereby increasing the risk of developing adverse reactions (e.g. myopathy and/or rhabdomyolysis).

The concurrent use of [selected drug] with
 [selected drug] should be avoided, as concurrent
 use may increase the risk of myopathy and
 rhabdomyolysis. [Selected drugs] have been
 shown to cause myopathy and rhabdomyolysis.
 Concomitant use of [selected drugs] has been
 shown to significantly increase rosuvastatin
 exposure.



May 2025:

Increased Risk of Serotonin Syndrome

Drugs included:

Selective Serotonin Reuptake Inhibitors (SSRIs)

Serotonin-Norepinephrine Reuptake Inhibitors (SNRIs)

Monoamine Oxidase Inhibitors (MAOIs)

Tricyclic
Antidepressants
(TCAs)

Vortioxetine

Meperidine

Tapentadol

Stimulants

Triptans

Linezolid

- Coadministration of [selected drugs] should be done with caution. With concomitant administration, it is possible that both agents may inhibit each other's metabolism increasing the serum levels of both drugs, possibly resulting in serotonergic syndrome and/or [selected drug] toxicity.
- [Selected drug] is a reversible non-selective inhibitor of monoamine oxidase and concurrent use with a serotonergic agent (e.g., SSRIs, SNRIs, TCAs, and meperidine) is contraindicated unless the patient is carefully observed for signs and symptoms of serotonin syndrome.
- Concomitant use of [selected drug] and selective 5-HT1
 receptor agonists may cause rapid CNS serotonin
 accumulation. Monitor the patient for signs and symptoms
 of serotonin syndrome, which can present as hyperthermia,
 hypertension, myoclonus, rigidity, mental status changes,
 and coma.



June 2025:

Increased Risk for QT Prolongation

Drugs included:

MAOIs

TCAs

Buprenorphine

lloperidone

Quetiapine

Asenapine

Citalopram

Dronedarone

Ribociclib

Vilanterol

CYP 3A4 Substrates

Pimavanserin

Glycopyrrolate/Formoterol

Deutetrabenazine

Dextromethorphan/quinidi

ne

Dofetilide

Pimozide

Rilpivirine

Posaconazole

Pitolisant

- [Selected Drug] can cause QT interval prolongation. Monitor ECGs in patients with congenital long QT syndrome, heart failure, bradyarrhythmias, or those taking drugs known to prolong the QT interval. Also, monitor and correct any electrolyte abnormalities at baseline and periodically during treatment. Withhold [selected drug] for the development of QTc interval prolongation greater than 500 ms or greater than 60 ms increase from baseline. Resume [selected drug] at a reduced dose (per adjustments in official prescribing information) when QTc prolongation resolves to less than or equal to 480 ms or baseline.
- [Selected Drugs] prolongs the QT/QTc interval.
 Coadministration of [selected drug(s)] with other drugs that prolong the QT interval may further increase the incidence of QT prolongation. Monitor patients more frequently with ECG if coadministration of [selected drug(s)] with drugs known to prolong the QT interval is required.



RDUR Outcomes Report

	Criteria	PRE Targeted Patients	POST Targeted Patients with Hits	POST Targeted Patients with Change
October 2023	Cholesterol Guidelines	344	193	56.10%
November 2023	Underutilization of Antihypertensive	271	259	95.57%
December 2023	Increased Risk of Bleeding	288	156	54.17%
January 2024	Enhanced respiratory depression	353	140	39.66%
February 2024	Multiple cardiovascular risk factors	323	222	68.73%
March 2024	Exacerbation of congestive heart failure	237	145	61.18%
Apr 2024	Antihyperlipidemic Drug Interactions	284	157	55.28%
May 2024	Impaired Antihypertensive Effects	167	102	61.08%
June 2024	Increased Risk of Selected Electrolyte Disturbances	328	223	67.99%
July 2024	Potential adverse fetal/neonatal risk in pregnant or lactating patients	197	145	73.60%
August 2024	Increased risk of osteoporosis and/or fracture	269	185	68.77%
September 2024	Inappropriate use of antidiabetic therapies	270	165	61.11%
Total		3331	2,092	62.80%



Lock-In Interventions

Month	Reviewed	Warning Letter	Locked In
Jan	75	7	0
Feb	75	6	0
Mar	75	8	1
Totals	225	21	1

2025 Second Quarter Newsletter



FDA-Approval Spotlight

Journavx (suzetrigine)

Legislative News

Consolidations Appropriations Act (Sec. 203)

2024 Guideline Update

ACP Guidelines for the
Treatment and Prevention
of Acute Episodic
Migraines

