

2019 Medicare Part D Opioid Policies: Information for Prescribers

The Centers for Medicare and Medicaid Services (CMS) <u>finalized new opioid policies</u> for Medicare drug plans starting on January 1, 2019. Providers are in the best position to identify and manage potential opioid overutilization in the Medicare Part D population. The new policies include improved **safety alerts** when opioid prescriptions are dispensed at the pharmacy and **drug management programs** for patients determined to be at-risk for misuse or abuse of opioids or other frequently abused drugs.

Residents of long-term care facilities, those in hospice care, patients receiving palliative or end-of-life care, and patients being treated for active cancer-related pain are exempt from these interventions. These policies should not impact patients' access to medication-assisted treatment (MAT), such as buprenorphine.

Opioid Safety Alerts

Part D plans are expected to implement safety alerts (pharmacy claim edits) for pharmacists to review at the time of dispensing the medication to prevent the unsafe utilization of drugs. CMS encourages prescribers to respond to pharmacists' outreach in a timely manner and give the appropriate training to on-call prescribers when necessary to resolve opioid safety edits expeditiously and avoid disruption of therapy.

Opioid Safety Alert	Prescriber's Role
Seven-day supply limit for opioid naïve patients ("hard edit")	Patient may receive up to a 7 days supply or request a coverage determination for full days supply as written.
Medicare Part D patients who have not filled an opioid prescription recently (such as within the past 60 days) will be limited to a supply of 7 days or less.	The physician or other prescriber has the right to request a coverage determination on patient's behalf, including the right to request an expedited or standard
Limiting the amount dispensed with the first opioid prescription may reduce the risk of a future	coverage determination in advance of prescribing an opioid.
dependency or overuse of these drugs.	Prescriber only needs to attest to plan that the days supply is the intended and medically necessary amount.
Important Note: This alert should not impact patients who already take opioids.	Subsequent prescriptions written by prescribers are not subject to the 7 days supply limit, as the patient will no longer be considered opioid naïve.
Opioid care coordination alert at 90 morphine milligram equivalent (MME) This policy will affect Medicare patients when they present an opioid prescription at the pharmacy and	Regardless of whether individual prescription(s) are written below the threshold, the alert will be triggered by the fill of the prescription that reaches the cumulative threshold of 90 MME or greater.
their cumulative MME per day across all of their opioid prescription(s) reaches or exceeds 90 MME. <i>Some plans use this alert only when the patient uses</i>	The prescriber who writes the prescription will trigger the alert and will be contacted even if that prescription itself is below the 90 MME threshold.
<i>multiple opioid prescribers and/or opioid dispensing pharmacies.</i>	Once a pharmacist consults with a prescriber on a patient's prescription for a plan year, the prescriber will
The prescriber will be contacted to resolve the alerts and to be informed of other opioid prescribers or increasing level (MME) of opioids.	not be contacted on every opioid prescription written for the same patient after that unless the plan implements further restrictions.

<u>Important Note:</u> <i>This is not a prescribing limit.</i> Decisions to taper or discontinue prescription opioids are individualized between the patient and prescriber.	On the patient's behalf, the physician or other prescriber has the right to request a coverage determination for a drug(s), including the right to request an expedited or standard coverage determination in advance of prescribing an opioid.
Concurrent opioid and benzodiazepine use or duplicative long-acting opioid therapy ("soft edits") The alerts will trigger when opioids and benzodiazepines are taken concurrently or if on multiple duplicate long-acting opioids.	The pharmacist will conduct additional safety reviews to determine if the patient's opioid use is safe and clinically appropriate. The prescriber may be contacted.
Optional Safety Alert at 200 MME or more ("hard edit") Some plans may implement a hard safety alert when a patient's cumulative opioid daily dosage reaches 200 MME or more. Some plans use this alert only when the patient uses multiple opioid prescribers and/or opioid dispensing pharmacies.	This alert stops the pharmacy from processing the prescription until an override is entered or authorized by the plan. On the patient's behalf, the physician or other prescriber has the right to request a coverage determination for a drug(s), including the right to request an expedited or standard coverage determination in advance of prescribing an opioid.
Important Note: This is not a prescribing limit. Decisions to taper or discontinue prescription opioids are individualized between the patient and prescriber.	In the absence of other approved utilization management requirements, once the prescriber(s) attests that the identified cumulative MME level is the intended and medically necessary amount, the medication will be dispensed to the patient.

Drug Management Programs (DMPs)

Medicare Part D plans may have a DMP that limits access to opioids and benzodiazepines for patients who are considered to be at-risk by the plan for prescription drug abuse.

The goal of a DMP is better care coordination for safer use. Coverage limitations under a DMP can include requiring the patient to obtain these medications from a specified prescriber and/or pharmacy, or implementing an individualized POS edit that limits the amount of these medications that will be covered for the patient. The coverage limitation tools may be put in place for 12 months and extended for an additional 12 months (total of 24 months).

Potential at-risk patients are identified by their opioid use which involve multiple doctors and pharmacies. After the plan conducts case management with prescribers, and before implementing a coverage limitation tool, the plan will notify the patient in writing. Plans are required to make reasonable efforts to send the prescriber a copy of the letter. After this 30 day time period, if the plan determines based on its review that the patient is at-risk and implements a limitation, it must send the patient a second written notice confirming the specific limitation and its duration.

If the plan decides to limit coverage under a DMP, the patient and their prescriber have the right to appeal the plan's decision. The patient or prescriber should contact the plan for additional information on how to appeal.