

# **PSYCHOTROPIC MEDICATION PRESCRIBING GUIDELINES: February 2023**

#### F-758 GUIDELINES: **DOCUMENTATION for INITIATION and CONTINUATION (plus EXCEPTIONS) DURATION OF USE and GDR MEDICATION CLASSES Attempt and document** INITIATION/CONTINUATION **Routine Psychotropics** Medical record must include a validating diagnosis/indication documented for every psychotropic in use per Antipsychotics **GDRs** as clinically CMS SOM Appendix PP §483.45(e)(1) appropriate (unless Antidepressants Anxiolytics contraindicated) to PSYCHOTROPIC GDR CONTRAINDICATION JUSTIFICATION FOR USE determine: Hypnotics Dementia Diagnosis (include but not limited to): \*Other\* lowest effective dose. · Resident's target symptoms returned or worsened after the most recent attempt at a GDR within the or if psychotropic facility; AND Physician documents clinical rationale for why additional dose reduction attempts would likely impair medication can be **CMS PHASE III UPDATES** resident's function or increase distressed behavior. discontinued √ \*Other\* medications **Dose reductions should** Diagnosis OTHER THAN Dementia (Schizophrenia, Bipolar Mania, Depression with Psychotic Features, that affect brain activity occur: or other disorder which may cause Psychosis: (associated with mental in modest increments. The continued use is in accordance with relevant current standards of practice AND processes and and Physician documents the clinical rationale for why any attempted dose reduction would be likely to impair behaviors) ARE subject over adequate periods the resident's function or exacerbate an underlying medical or psychiatric disorder; OR to the psychotropic of time to minimize Resident's target symptoms returned or worsened after the most recent attempt at a GDR within the medication withdrawal symptoms facility; AND requirements if and to monitor for documented use Physician documents the clinical rationale for why any additional attempted dose reduction at that time symptom recurrence. would be likely to impair the resident's function or exacerbate an underlying medical or psychiatric appears to be a For example (not an substitution for another disorder. absolute requirement): psychotropic Compliance with the LONG TERM TREATMENT FOR SPECIFIC DISORDERS WITH PSYCHOTROPIC MEDS medication, rather than requirement to perform a Clinical goal: Minimize/Eradicate symptoms of disorder for the original or GDR may be met if, approved indication. • Even though symptoms have subsided, long term treatment is required so that symptoms do not return. within the 1st year in Reducing or eliminating medication may be contraindicated and must be individualized. which a resident is I.e.: Chronic Depression, Parkinson's Disease Psychosis, Recurrent Seizures, Chronic Psychiatric Illness admitted on a (Schizophrenia, Schizoaffective disorder, Post-Traumatic Stress Disorder); Neurological Disorders

### **ANTIPSYCHOTICS**

psychotropic drug or

after the prescribing practitioner has initiated

a psychotropic drug, a

two separate quarters

the attempts), unless

facility attempts a GDR in

(with > 1 month between

clinically contraindicated.

Require clear documentation of diagnosis and indication for use, multiple attempts at care-planned non-drug interventions that have failed, and ongoing evaluation of these approaches. Diagnoses alone do not warrant use. Indication may be warranted if:

- Behavioral symptoms present danger to self or others,
- Expressions or indications of distress that are significant to the resident,

(Huntington's and Tourette's); Psychosis and Psychotic Episodes.

- If not clinically contraindicated, non-pharmacological approaches previously attempted failed, and/or
- GDR was attempted, but clinical symptoms returned.



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MEDICATION CLASSES	F-758 GUIDELINES: DURATION OF USE and GDR	DOCUMENTATION for INITIATION and CONTINUATION (plus EXCEPTIONS)
PRN Psychotropic (Except Antipsychotics)	<ul> <li>Limited to 14 days         OR</li> <li>Can extend duration         beyond 14 days with         prescribing practitioner's         rationale and duration</li> </ul>	<ul> <li>Effectiveness</li> <li>Ongoing specific diagnosed condition</li> <li>Indication</li> <li>Duration</li> </ul>
PRN Antipsychotic	Limited to 14 days (without exception)	<ul> <li>DIRECT EVALUATION BY PRESCRIBER INCLUDING THE FOLLOWING IN THE RESIDENT'S MEDICAL RECORD WITH EVERY ORDER AND CONTINUATION:</li> <li>Ongoing specific diagnosed condition and indication: Is the antipsychotic medication still needed on a PRN basis?</li> <li>What is the benefit of the medication to the resident?</li> <li>Effectiveness: Have the resident's expressions or indications of distress improved as a result of the PRN medication?</li> <li>Duration cannot exceed 14 days</li> </ul>

### POTENTIAL ADVERSE REACTIONS OF PSYCHOTROPIC MEDICATIONS FOR MONITORING:

- General: anticholinergic effects including flushing, blurred vision, dry mouth, altered mental status, difficulty urinating, falls, excessive sedation, constipation
- Cardiovascular: signs and symptoms of cardiac arrhythmias such as irregular heart beat or pulse, palpitations, lightheadedness, shortness of breath, diaphoresis, chest or arm pain, increased blood pressure, orthostatic hypotension
- Metabolic: increase in total cholesterol and triglycerides, unstable or poorly controlled blood sugar, weight gain
- Neurologic: agitation, distress, EPS, neuroleptic malignant syndrome (NMS), Parkinsonism, tardive dyskinesia, cerebrovascular event.

Emphasis is on seeking lowest effective dose and duration and minimizing risk of adverse consequences.

**HOSPICE:** PRN Psychotropic exceptions are NOT made for residents on Hospice.

**NON-PSYCHOTROPIC INDICATIONS:** PRN antipsychotics used for other non-psychotropic indications such as nausea, hiccups, etc. have not been given exception and are still subject to the 14 day limit.

F757; \$483.45(d) Unnecessary Drugs – General: each resident's drug regimen must be free from unnecessary drugs (any drug when used):

 In excessive dose (including duplicate drug therapy); or for excessive duration; or without adequate monitoring; or without adequate indications for its use; or the presence of adverse consequences which indicate the dose should be reduced or discontinued; or any combinations of the reasons.

CMS SOM – Appendix PP Effective 02.17.2023